Arno Therapeutics, Inc
Form S-1
November 08, 2010

As filed with the Securities and Exchange Commission on November 8, 2010 Registration No. 333-

UNITED STATES SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM S-1

REGISTRATION STATEMENT UNDER THE SECURITIES ACT OF 1933

ARNO THERAPEUTICS, INC.
(Exact name of registrant as specified in its charter)

Delaware	2834	52-2286452
(State or other jurisdiction of incorporation or organization)	(Primary Standard Industrial Classification Code Number)	(I.R.S. Employer Identification No.)

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4 Campus Drive, 2nd Floor
Parsippany, New Jersey 07054
(862) 703-7170
(Address, including zip code, and telephone number, including area code, of registrant's principal executive offices)

David M. Tanen President Arno Therapeutics, Inc. 4 Campus Drive, 2nd Floor Parsippany, NJ 07054 (862) 703-7170 (Name, address, including zip code, and telephone number, including area code, of agent for service)	Copies to: Christopher J. Melsha, Esq. Sean M. Nagle, Esq. Fredrikson & Byron, P.A. 200 South Sixth Street, Suite 4000 Minneapolis, MN 55402-1425 Telephone: (612) 492-7000 Facsimile: (612) 492-7077

Approximate date of commencement of proposed sale to the public: From time to time after the effective date of this registration statement, as shall be determined by the selling stockholders identified herein.

If any of the securities being registered on this Form are to be offered on a delayed or continuous basis pursuant to Rule 415 under the Securities Act of 1933, as amended, check the following box. þ

If this Form is filed to register additional securities for an offering pursuant to Rule 462(b) under the Securities Act, check the following box and list the Securities Act registration number of the earlier effective registration statement for the same offering. ¨

If this Form is a post effective amendment filed pursuant to Rule 462(c) under the Securities Act, check the following box and list the Securities Act registration statement number of the earlier effective registration statement for the same offering. ¨

If this Form is a post effective amendment filed pursuant to Rule 462(d) under the Securities Act, check the following box and list the Securities Act registration statement number of the earlier effective registration statement for the same offering. ¨

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, or a smaller reporting company. See the definitions of "large accelerated filer," "accelerated filer" and "smaller reporting company" in Rule 12b-2 of the Exchange Act.

Large accelerated filer ¨	Accelerated filer ¨
Non-accelerated filer ¨ (Do not check if a smaller reporting company)	Smaller reporting company þ

CALCULATION OF REGISTRATION FEE

Title of Each Class of Securities to be Registered	Amount to be Registered (1)	Proposed Maximum Offering Price Per Share (2)	Proposed Maximum Aggregate Offering Price (2)	Amount of Registration Fee
Common stock, par value $0.0001 per share	18,121,901	$ 0.51	$ 9,242,169.51	$ 658.97
Common stock, par value $0.0001 per share (3)	8,693,930	$ 0.51	$ 4,433,904.30	$ 316.14
Total	26,815,831		$ 13,676,073.81	$ 975.10

(1) There is also being registered hereunder an indeterminate number of additional shares of common stock as shall be issuable pursuant to Rule 416 to prevent dilution resulting from stock splits, stock dividends or similar transactions.

(2) Estimated solely for the purpose of calculating the registration fee in accordance with Rule 457(c) of the Securities Act of 1933, as amended, based upon the average of the high and low price of our common stock on April 9, 2010, the most recent date on which shares were traded on the Pink Sheets.

(3) Represents shares of common stock issuable upon exercise of outstanding warrants.

THE REGISTRANT HEREBY AMENDS THIS REGISTRATION STATEMENT ON SUCH DATE OR DATES AS MAY BE NECESSARY TO DELAY ITS EFFECTIVE DATE UNTIL THE REGISTRANT SHALL FILE A FURTHER AMENDMENT WHICH SPECIFICALLY STATES THAT THIS REGISTRATION STATEMENT SHALL THEREAFTER BECOME EFFECTIVE IN ACCORDANCE WITH SECTION 8(A) OF THE SECURITIES ACT OF 1933 OR UNTIL THIS REGISTRATION STATEMENT SHALL BECOME EFFECTIVE ON SUCH DATE AS THE COMMISSION, ACTING PURSUANT TO SUCH SECTION 8(A), MAY DETERMINE.

A registration statement relating to these securities has been filed with the Securities and Exchange Commission. These securities may not be sold nor may offers to buy be accepted prior to the time the registration statement becomes effective. This prospectus shall not constitute an offer to sell or the solicitation of an offer to buy nor shall there be any sale of these securities in any state in which such offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such state.

Subject to completion, dated November 8, 2010

OFFERING PROSPECTUS

26,815,831 Shares
Common Stock

The selling stockholders identified beginning on page 20 of this prospectus are offering on a resale basis a total of 26,815,831 shares of our common stock, of which 15,655,844 shares are issuable upon the conversion of our outstanding Series A Convertible Preferred Stock (including up to 381,844 shares of common stock that may be issuable as payment of accrued dividends upon conversion of our Series A Convertible Preferred Stock) and 8,693,930 shares are issuable upon the exercise of outstanding warrants. We will not receive any proceeds from the sale of these shares by the selling stockholders.

Our common stock is quoted on the "Pink Sheets" quotation system under the symbol "ARNI.PK." On , 2010, the last sale price of our common stock as reported on the Pink Sheets was $.

The securities offered by this prospectus involve a high degree of risk.
See "Risk Factors" beginning on page 7.

Neither the Securities and Exchange Commission nor any state securities commission has approved or disapproved of these securities or determined that this prospectus is truthful or complete. A representation to the contrary is a criminal offense.

The date of this prospectus is , 2010.

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PROSPECTUS SUMMARY

This summary highlights information contained elsewhere in this prospectus. Because it is a summary, it may not contain all of the information that is important to you. Accordingly, you are urged to carefully review this prospectus in its entirety, including the risks of investing in our securities discussed under the caption "Risk Factors" and the financial statements and other information that is contained in or incorporated by reference into this prospectus or the registration statement of which this prospectus is a part before making an investment decision. Unless the context otherwise requires, hereafter in this prospectus the terms the "Company," "we," "us," or "our" refer to Arno Therapeutics, Inc., a Delaware corporation.

Company Overview

We are a development stage company focused on developing innovative products for the treatment of cancer. We currently have the exclusive worldwide rights to commercially develop three oncology product candidates:

- AR-12 – Our lead clinical product candidate is a potentially first-in-class, orally available, targeted anti-cancer agent that inhibits phosphoinositide dependent protein kinase-1, or PDK-1, a protein in the PI3K/Akt pathway involved in the growth and proliferation of cells, including cancer cells. We believe AR-12 may also cause cell death through the induction of stress in the endoplasmic reticulum. In May 2009, the FDA accepted our investigational new drug application, or IND, for AR-12. We are currently conducting a multi-centered Phase I clinical study of AR-12 in adult patients with advanced or recurrent solid tumors or lymphoma. The Phase I study of AR-12 is being conducted in two parts. The first part is a dose-escalating study, which we refer to as the Escalation Phase, primarily designed to evaluate the compound's safety in order to identify the maximum tolerated dose, or MTD, or a recommended dose, or RD, for future studies of AR-12. We anticipate that the Escalation Phase will be completed by the third quarter of 2011. Following the Escalation Phase, we plan to initiate the second part of the study, which involves enrolling an expanded cohort of additional patients at the MTD or RD in multiple tumor types. We refer to this second part of the study as the Expansion Phase. The purpose of the Expansion Phase is to further evaluate and confirm the PD effects, potential anti-tumor activity, and safety of AR-12 at the MTD or RD. We anticipate that the Expansion Phase will be fully enrolled within one year.

- AR-42 – We are also developing AR-42, an orally available, broad spectrum inhibitor of both histone and non-histone deacetylation proteins, which play an important role in the regulation of gene expression, cell growth and survival. In preclinical studies, AR-42 has demonstrated greater potency and activity in solid and liquid tumors when compared to vorinostat (also known as SAHA and marketed as Zolinza® by Merck) and other deacetylase inhibitors. These data demonstrate the potent and differentiating activity of AR-42. Additionally, pre-clinical findings presented at the 2009 American Society of Hematology Annual Meeting and Exposition showed that AR-42 potently and selectively inhibits leukemic stem cells in acute myeloid leukemia, or AML. AR-42 is currently being studied in an investigator initiated Phase I/IIa clinical study in adult patients with relapsed or refractory multiple myeloma, chronic lymphocytic leukemia, or CLL, or lymphoma. Once the MTD is defined, the study is designed so that additional patients can be added to investigate efficacy in a particular disease and help guide future Phase II programs. Up to an additional 10 patients may be enrolled at the MTD dose in each of the multiple myeloma, CLL and lymphoma.

- AR-67 – We are also developing AR-67, a novel, third-generation campothecin analogue that inhibits Topoisomerase I activity. In 2008, we completed a multi-centered, ascending dose Phase I clinical trial of AR-67 in patients with advanced solid tumors. AR-67 is currently being studied in a Phase II clinical trial in patients with glioblastoma multiforme, or GBM, a highly aggressive form of brain cancer. We anticipate having interim data from this Phase II study by the second quarter of 2011. Thereafter, if data permits, we may elect to initiate larger Phase II studies or advance AR-67 into a registration-enabling Phase III study.

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In June 2008, we were acquired by Laurier International, Inc., a Delaware corporation, in a "reverse" merger whereby a wholly-owned subsidiary of Laurier merged with and into Arno Therapeutics, with Arno Therapeutics remaining as the surviving corporation and a wholly-owned subsidiary of Laurier. In accordance with the terms of this merger, stockholders of Arno Therapeutics exchanged all of their shares of common stock of Arno Therapeutics for shares of Laurier common stock at a rate of 1.99377 shares of Laurier common stock for each share of Arno Therapeutics common stock. As a result of the issuance of the shares of Laurier common stock to the former Arno Therapeutics stockholders, following the merger the former stockholders of Arno Therapeutics held 95 percent of the outstanding common stock of Laurier, assuming the issuance of all shares underlying outstanding options and warrants. Upon completion of the merger, all of the former officers and directors of Laurier resigned and were replaced by the officers and directors of Arno Therapeutics. Additionally, following the merger Laurier changed its name to Arno Therapeutics, Inc.

In May 2009, we voluntarily filed a Form 15 with the Securities and Exchange Commission in order to terminate the registration of our common stock under the Securities and Exchange Act of 1934, as amended, or the Exchange Act. As a result, our obligation to file periodic and other reports under the Exchange Act was suspended. Following the effective date of the registration statement of which this prospectus forms a part, we will again be required to file periodic and other reports under the Exchange Act.

Our executive offices are located at 4 Campus Drive, 2nd Floor, Parsippany, New Jersey 07054. Our telephone number is (862) 703-7170. Our website is www.arnothera.com. Information contained in, or accessible through, our website does not constitute a part of this prospectus.

Risk Factors

As with most pharmaceutical product candidates, the development of our product candidates is subject to numerous risks, including the risk of delays in or discontinuation of development from lack of financing, inability to obtain necessary regulatory approvals to market the products, unforeseen safety issues relating to the products and dependence on third party collaborators to conduct research and development of the products. Because we are a development stage company with a very limited history of operations, we are also subject to many risks associated with early-stage companies. For a more detailed discussion of some of the risks you should consider before purchasing shares of our common stock, you are urged to carefully review and consider the section entitled "Risk Factors" beginning on page 7 of this prospectus.

The Offering

The selling stockholders identified beginning on page 20 of this prospectus are offering on a resale basis a total of 26,815,831 shares of our common stock, of which 15,655,844 shares are issuable upon the conversion of our outstanding Series A Convertible Preferred Stock (including up to 381,844 shares of common stock that may be issuable as payment of accrued dividends upon conversion of our Series A Convertible Preferred Stock) and 8,693,930 shares are issuable upon the exercise of outstanding warrants.

Common stock offered	26,815,831 shares
Common stock outstanding before the offering(1)	20,412,024 shares
Common stock outstanding after the offering(2)	44,761,798 shares
Use of Proceeds	We will receive none of the proceeds from the sale of the shares by the selling stockholders, except for

	the warrant exercise price upon exercise of the warrants, which would be used for working capital and other general corporate purposes
Pink Sheets Symbol	ARNI.PK

(1) Based on the number of shares outstanding as of October 31, 2010, not including 2,258,555 shares issuable upon exercise of various warrants and options to purchase our common stock or any Shares of Series A Preferred Stock or warrants to purchase Series A Preferred Stock.

(2) Assumes the issuance of all shares offered hereby that are issuable upon the conversion of our outstanding Series A Convertible Preferred Stock or upon exercise of warrants. Also assumes six months of dividend accrual at the rate of 5% per annum prior to the automatic conversion of the Series A Convertible Preferred Stock into common stock upon the effectiveness of the registration statement of which this prospectus is a part, and our election to pay such accrued dividends in the form of additional shares of common stock in lieu of cash. See "Description of Capital Stock – Series A Convertible Preferred Stock."

Recent Developments

On September 3, 2010, we entered into a Securities Purchase and Registration Rights Agreement, or the Purchase Agreement, with a number of institutional and accredited investors pursuant to which we sold in a private placement an aggregate of 15,274,000 shares of our newly-designated Series A Convertible Preferred Stock, par value $0.0001 per share, or Series A Preferred Stock, at a per share purchase price of $1.00. In accordance with the Purchase Agreement, we also issued to each investor a two-and-one-half-year warrant to purchase a number of additional shares of Series A Preferred Stock equal to 8% of the number of shares purchased by such investor at an initial exercise price of $1.00 per share. We collectively refer to these warrants as the Class A Warrants. In addition to the Class A Warrants, each investor also received a five-year warrant to purchase a number of additional shares of Series A Preferred Stock equal to 42% of the number of shares purchased by such investor at an initial exercise price of $1.15 per share, which we collectively refer to as the Class B Warrants. Pursuant to the Purchase Agreement, we issued to the investors Class A Warrants to purchase an aggregate of 1,221,920 shares of Series A Preferred Stock, and Class B Warrants to purchase an aggregate of 6,415,080 shares of Series A Preferred Stock. The sale of the shares and warrants resulted in aggregate gross proceeds of approximately $15.2 million, before deducting expenses.

Pursuant to the terms of the Purchase Agreement, we agreed to file a registration statement under the Securities Act of 1933, as amended, covering the resale of the shares of our common stock issuable upon conversion of the Series A Preferred Stock sold in the private placement, including the shares issuable upon exercise of the Class A and Class B Warrants. We further agreed to use our reasonable best efforts to cause such registration statement to be declared effective within 180 days following the initial closing under the Purchase Agreement, or by March 8, 2011. If such registration statement is not declared effective by the SEC by such date, we agreed to pay liquidated damages to the investors in the amount of 1% of each investor's aggregate investment amount for each 30-day period until the registration statement is declared effective. The registration statement of which this prospectus is a part registers the shares of our common stock issuable upon conversion of the Series A Preferred Stock sold in the private placement and upon exercise of the Class A and Class B Warrants following their conversion into warrants to purchase shares of common stock. Upon the effectiveness of such registration statement, all outstanding shares of Series A Preferred Stock will automatically convert into shares of our common stock, and all outstanding warrants to purchase shares of Series A Preferred Stock will automatically convert into warrants to purchase shares of our common stock.

In connection with the private placement, we engaged Riverbank Capital Securities, Inc., or Riverbank, to serve as placement agent. In consideration for its services, we paid Riverbank a placement fee of $789,880, and we paid I-Bankers Securities, Inc., or I-Bankers, Riverbank's sub-agent, a placement fee of $267,050. In addition, we issued to designees of Riverbank and I-Bankers five-year warrants to purchase an aggregate of 664,880 and 392,050 shares, respectively, of Series A Preferred Stock at an initial exercise price of $1.10 per share. The warrants issued to Riverbank and I-Bankers are in substantially the same form as the Class A and Class B Warrants issued to the investors, except that they do not include certain anti-dilution provisions contained in the Class A and Class B Warrants. David M. Tanen, our President, Secretary, and a member of our Board of Directors, Peter M. Kash, also a member of our Board of Directors, and Joshua A. Kazam, who served as a director until September 2010, are each officers of and collectively control Riverbank.

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The Purchase Agreement provides that the three co-lead investors in the private placement each have the right to designate one individual to be appointed to our board of directors. Accordingly, following the completion of the private placement, we appointed Tomer Kariv, Yacov Reizman, and Steven Ruchefsky to our board, each of whom was designated by one of the three co-lead investors in the private placement.

RISK FACTORS

Investment in our common stock involves significant risk. You should carefully consider the information described in the following risk factors, together with the other information appearing elsewhere in this prospectus, before making an investment decision regarding our common stock. If any of these risks actually occur, our business, financial conditions, results of operations and future growth prospects would likely be materially and adversely affected. In these circumstances, the market price of our common stock could decline, and you may lose all or a part of your investment in our common stock. Moreover, the risks described below are not the only ones that we face. Additional risks not presently known to us or that we currently deem immaterial may also affect our business, operating results, prospects or financial condition.

Risks Relating to Our Business

We currently have no product revenues and will need to raise substantial additional capital to operate our business.

To date, we have generated no product revenues. Until, and unless, we receive approval from the FDA and other regulatory authorities for our product candidates, we cannot sell our drugs and will not have product revenues. Currently, our only product candidates are AR-12, AR-42 and AR-67, and none of these products are approved for sale by the FDA. Therefore, for the foreseeable future, we will have to fund all of our operations and capital expenditures from cash on hand and, potentially, future offerings. After giving effect to our September 2010 private placement, we believe we have cash on hand to fund our operations through 2011. We will require substantial additional funds in addition to the proceeds from this offering to support our continued research and development activities, and the anticipated costs of preclinical studies and clinical trials, regulatory approvals and eventual commercialization. There can be no assurance that such additional financing can be obtained on desirable terms, if at all. In addition, changes may occur that would consume our available capital before that time, including changes in and progress of our development activities, acquisitions of additional product candidates and changes in regulation. Accordingly, we will need additional capital to fund our continuing operations. Since we do not generate any recurring revenue, the most likely sources of such additional capital include private placements of our equity securities, including our common stock, debt financing or funds from a potential strategic licensing or collaboration transaction involving the rights to one or more of our product candidates. To the extent that we raise additional capital by issuing equity securities, our stockholders will likely experience dilution, which may be significant depending on the number of shares we may issue and the price per share. If we raise additional funds through collaborations and licensing arrangements, it may be necessary to relinquish some rights to our technologies, product candidates or products, or grant licenses on terms that are not favorable to us. If we raise additional funds by incurring debt, we could incur significant interest expense and become subject to covenants in the related transaction documentation that could affect the manner in which we conduct our business.

However, we currently have no committed sources of additional capital and our access to capital funding is always uncertain. This uncertainty is exacerbated due to the current global economic turmoil, which has severely restricted access to the U.S. and international capital markets, particularly for small biopharmaceutical and biotechnology companies. Accordingly, despite our ability to secure adequate capital in the past, there is no assurance that additional equity or debt financing will be available to us when needed, on acceptable terms or even at all. If we fail to obtain the necessary additional capital when needed, we may be forced to significantly curtail our planned research and development activities, which will cause a delay in our drug development programs and may severely harm our business.

We are a development stage company.

We have not received any operating revenues to date and are in the development stage. You should be aware of the problems, delays, expenses and difficulties encountered by an enterprise in our stage of development, and particularly for companies engaged in the development of new biotechnology or biopharmaceutical product candidates, many of which may be beyond our control. These include, but are not limited to, problems relating to product development, testing, regulatory compliance, manufacturing, marketing, costs and expenses that may exceed current estimates and competition. No assurance can be given that our existing product candidates, or any technologies or products that we may acquire in the future will be successfully developed, commercialized and accepted by the marketplace or that sufficient funds will be available to support operations or future research and development programs.

We are not currently profitable and may never become profitable.

We expect to incur substantial losses and negative operating cash flows for the foreseeable future, and we may never achieve or maintain profitability. For the years ended December 31, 2009 and 2008, we had a net loss of $6,936,705 and $12,913,566, respectively, and for the period from our inception on August 1, 2005 through December 31, 2009, we had a net loss of $23,580,861. Even if we succeed in developing and commercializing one or more of our product candidates, we expect to incur substantial losses for the foreseeable future, as we:

- continue to undertake pre-clinical development and clinical trials for our product candidates;
- seek regulatory approvals for our product candidates;
- in-license or otherwise acquire additional products or product candidates;
- seek patent protection for our product candidates;
- implement additional internal systems and infrastructure; and
- hire additional personnel.

Further, for the years ended December 31, 2009 and 2008, we had negative cash flows from operating activities of $7,310,308 and $8,883,956, respectively, and since inception on August 1, 2005 through December 31, 2009, we have had negative cash flows from operating activities of $18,370,053. We expect to continue to experience negative cash flows for the foreseeable future as we fund our operating losses and capital expenditures. As a result, we will need to generate significant revenues in order to achieve and maintain profitability. We may not be able to generate these revenues or achieve profitability in the future. Our failure to achieve or maintain profitability could negatively impact the value of our common stock.

We have a limited operating history upon which to base an investment decision.

We are a development stage company and have not demonstrated our ability to perform the functions necessary for the successful commercialization of any of our product candidates. The successful commercialization of our product candidates will require us to perform a variety of functions, including:

- continuing to undertake pre-clinical development and clinical trials for our product candidates;
- participating in regulatory approval processes;
- formulating and manufacturing products; and
- conducting sales and marketing activities.

Our operations have been limited to organizing our company, acquiring, developing and securing our proprietary technologies and performing pre-clinical and clinical trials of our product candidates. These operations provide a limited basis for you to assess our ability to commercialize our product candidates and the advisability of investing in our securities.

We may not successfully manage our growth.

Our success will depend upon the expansion of our operations and the effective management of our growth, which will place a significant strain on our management and on our administrative, operational and financial resources. To manage this growth, we may need to expand our facilities, augment our operational, financial and management systems and hire and train additional qualified personnel. If we are unable to manage our growth effectively, our business would be harmed.

The relationships between Two River Consulting, Riverbank Capital Securities and certain of our officers and directors may present potential conflicts of interest.

Arie S. Belldegrun and David M. Tanen, each of whom are currently directors of our company, and Joshua A. Kazam, a co-founder and director of our company until September 2010, are the managing members of Two River Consulting, LLC, or Two River. Since June 2009, Mr. Tanen has also served as our President. In June 2009, we entered into a services agreement with Two River pursuant to which it performs various management, clinical development, operational and administrative activities and services for us. As consideration for these services, we pay Two River a monthly cash fee of $55,000. Each of Messrs. Kazam and Tanen, as well as Peter M. Kash, a director of our company, are also officers and directors of Riverbank, a registered broker-dealer, which served as placement agent in connection with our September 2010 private placement of Series A Preferred Stock. Scott L. Navins, the Financial and Operations Principal of Riverbank, serves as our Treasurer.

Generally, Delaware corporate law requires that any transactions between us and any of our affiliates be on terms that, when taken as a whole, are substantially as favorable to us as those then reasonably obtainable from a person who is not an affiliate in an arms-length transaction. We believe that the terms of the agreements that we have entered into with Two River and Riverbank satisfy the requirements of Delaware law, but in the event one or more parties challenges the fairness of such terms we may have to expend substantial resources in resolving such challenges and can make no guarantees of the result. Further, none of our affiliates or Two River is obligated pursuant to any agreement or understanding with us to make any additional products or technologies available to us, nor can there be any assurance, and the investors should not expect, that any biomedical or pharmaceutical product or technology identified by such affiliates or Two River in the future will be made available to us. In addition, certain of our current officers and directors or certain of any officers or directors hereafter appointed may from time to time serve as officers or directors of other biopharmaceutical or biotechnology companies. There can be no assurance that such other companies will not have interests in conflict with our own.

We are substantially dependent on the services of Two River and other consultants.

We have only two employees. We currently rely heavily on Two River to render various management, clinical development, regulatory, operational and administrative activities and services for us. We also rely in substantial part, and for the foreseeable future will continue to rely, on certain independent organizations and consultants to provide other important services, including substantially all aspects of regulatory approval, clinical management, and manufacturing. There can be no assurance that the services of independent organizations, advisors and consultants will continue to be available to us on a timely basis when needed, or that we can find qualified replacements.

Our President provides his services on a part-time basis and significant other services are currently being rendered by outside consultants. If we are unable to hire additional qualified personnel in the future, our ability to grow our business may be harmed.

Although we currently engage Two River to provide personnel to perform a variety of management, clinical development and other services on our behalf on a consulting basis, we expect to directly hire employees, including at

the senior management level, in the future as we further the development of our clinical programs. In addition, David Tanen, our current President, provides his services to us on a part-time, non-employee basis. As we further the development of our product candidates, we intend to hire a full-time chief executive officer and other employees to perform the services currently being rendered by Two River. Accordingly, our ability to attract and retain qualified personnel will be critical to managing and growing our business in the future, especially the hiring and retention of key executive personnel and scientific staff. There is intense competition and demand for qualified personnel in our area of business and no assurances can be made that we will be able to retain the personnel necessary for the development of our business on commercially reasonable terms, if at all.

We may incur substantial liabilities and may be required to limit commercialization of our products in response to product liability lawsuits.

The testing and marketing of medical products entail an inherent risk of product liability. If we cannot successfully defend ourselves against product liability claims, we may incur substantial liabilities or be required to limit commercialization of our products. Our inability to obtain sufficient product liability insurance at an acceptable cost to protect against potential product liability claims could prevent or inhibit the commercialization of the pharmaceutical products we develop, alone or with corporate collaborators. We currently do not have product liability insurance, but do maintain clinical trial insurance coverage. Even if our agreements with any future corporate collaborators entitle us to indemnification against losses, such indemnification may not be available or adequate should any claim arise.

We are controlled by current directors and principal stockholders.

Our executive officers, directors and principal stockholders, which include the persons affiliated with Two River discussed above, beneficially own approximately 70% of our outstanding voting securities. Accordingly, our executive officers, directors, principal stockholders and certain of their affiliates will have the ability to exert substantial influence over the election of our board of directors and the outcome of issues submitted to our stockholders.

We may be required to implement additional finance and accounting systems, procedures and controls in order to satisfy requirements under the securities laws, including the Sarbanes-Oxley Act of 2002, which will increase our costs and divert management’s time and attention.

We are in a continuing process of further establishing and documenting controls and procedures that will allow our management to report on, and our independent registered public accounting firm to attest to, our internal controls over financial reporting if and when required to do so under Section 404 of the Sarbanes-Oxley Act of 2002, or the Sarbanes-Oxley Act. As a company with limited capital and human resources, we anticipate that more of management’s time and attention will be diverted from our business to ensure compliance with these regulatory requirements than would be the case with a company that has well established controls and procedures. This diversion of management’s time and attention may have a material adverse effect on our business, financial condition and results of operations.

In the event we identify significant deficiencies or material weaknesses in our internal controls over financial reporting that we cannot remediate in a timely manner, investors and others may lose confidence in the reliability of our financial statements. If this occurs, the trading price of our common stock, if any, and our ability to obtain any necessary financing could suffer. In addition, in the event that our independent registered public accounting firm is unable to rely on our internal controls over financial reporting in connection with its audit of our financial statements, and in the further event that it is unable to devise alternative procedures in order to satisfy itself as to the material accuracy of our financial statements and related disclosures, we may be unable to file our Annual Reports on Form 10-K with the SEC. This would likely have an adverse affect on the trading price of our common stock, if any, and our ability to secure any necessary additional financing, and could result in the delisting of our common stock if we are listed on an exchange in the future. In such event, the liquidity of our common stock would be severely limited and the market price of our common stock would likely decline significantly.

We will experience increased costs as a result of becoming subject to the reporting requirements of federal securities laws.

Upon the effectiveness of the registration statement of which this prospectus is a part, we will again become subject to the reporting requirements of the Exchange Act, including the requirements of the Sarbanes-Oxley Act of 2002. These requirements may place a strain on our systems and resources. The Securities Exchange Act of 1934 requires that we file annual, quarterly and current reports with respect to our business and financial condition. The Sarbanes-Oxley Act requires that we maintain effective disclosure controls and procedures and internal controls over financial reporting, which is discussed above. In order to maintain and improve the effectiveness of our disclosure controls and procedures, significant resources and management oversight will be required. We will continue to be implementing additional procedures and processes for the purpose of addressing the standards and requirements applicable to public companies. In addition, sustaining our growth will also require us to commit additional management, operational and financial resources to identify new professionals to join our firm and to maintain appropriate operational and financial systems to adequately support expansion. These activities may divert management's attention from other business concerns, which could have a material adverse effect on our business, financial condition, results of operations and cash flows. We expect to incur significant additional annual expenses related to these steps and, among other things, additional directors and officers liability insurance, director fees, reporting requirements of the SEC, transfer agent fees, hiring additional accounting, legal and administrative personnel, increased auditing and legal fees and similar expenses.

Risks Relating to the Clinical Testing, Regulatory Approval, Manufacturing and Commercialization of Our Product Candidates

We may not obtain the necessary U.S. or worldwide regulatory approvals to commercialize our product candidates.

We will need FDA approval to commercialize our product candidates in the U.S. and approvals from the FDA equivalent regulatory authorities in foreign jurisdictions to commercialize our product candidates in those jurisdictions. In order to obtain FDA approval of any of our product candidates, we must submit to the FDA a new drug application, or NDA, demonstrating that the product candidate is safe for humans and effective for its intended use. This demonstration requires significant research and animal tests, which are referred to as pre-clinical studies, as well as human tests, which are referred to as clinical trials. Satisfaction of the FDA’s regulatory requirements typically takes many years, depends upon the type, complexity and novelty of the product candidate and requires substantial resources for research, development and testing. We cannot predict whether our research and clinical approaches will result in drugs that the FDA considers safe for humans and effective for indicated uses. The FDA has substantial discretion in the drug approval process and may require us to conduct additional pre-clinical and clinical testing or to perform post-marketing studies. The approval process may also be delayed by changes in government regulation, future legislation or administrative action or changes in FDA policy that occur prior to or during our regulatory review. Delays in obtaining regulatory approvals may:

- delay commercialization of, and our ability to derive product revenues from, our product candidates;
- impose costly procedures on us; or
- diminish any competitive advantages that we may otherwise enjoy.

Even if we comply with all FDA requests, the FDA may ultimately reject one or more of our NDAs. We cannot be sure that we will ever obtain regulatory clearance for our product candidates. Failure to obtain FDA approval of any of our product candidates will severely undermine our business by reducing our number of salable products and, therefore, corresponding product revenues.

In foreign jurisdictions, we must receive approval from the appropriate regulatory authorities before we can commercialize our drugs. Foreign regulatory approval processes generally include all of the risks associated with the FDA approval procedures described above. We cannot assure that we will receive the approvals necessary to commercialize our product candidate for sale outside the U.S.

All of our product candidates are in early stages of clinical trials, which are very expensive and time-consuming. Any failure or delay in completing clinical trials for our product candidates could harm our business.

All three of our current product candidates are in early stages of development and will require extensive clinical and other testing and analysis before we will be in a position to consider seeking regulatory approval to sell such product candidates. Conducting clinical trials is a lengthy, time consuming and very expensive process and the results are inherently uncertain. The duration of clinical trials can vary substantially according to the type, complexity, novelty and intended use of the product candidate. We estimate that clinical trials of our product candidates will take at least several years to complete. The completion of clinical trials for our product candidates may be delayed or prevented by many factors, including without limitation:

- delays in patient enrollment, and variability in the number and types of patients available for clinical trials;
- difficulty in maintaining contact with patients after treatment, resulting in incomplete data;
- poor effectiveness of product candidates during clinical trials;
- safety issues, side effects, or other adverse events;
- results that do not demonstrate the safety or effectiveness of the product candidates;
- governmental or regulatory delays and changes in regulatory requirements, policy and guidelines; and
- varying interpretation of data by the FDA.

In conducting clinical trials, we may fail to establish the effectiveness of a compound for the targeted indication or discover that it is unsafe due to unacceptable side effects or other reasons. Even if our clinical trials are commenced and completed as planned, their results may not support our product candidate claims. Further, failure of product candidate development can occur at any stage of clinical trials, or even thereafter, and we could encounter problems that cause us to abandon or repeat clinical trials. These problems could interrupt, delay or halt clinical trials for our product candidates and could result in FDA, or other regulatory authorities, delaying or declining approval of our product candidates for any or all indications. The results from pre-clinical testing and prior clinical trials may not be predictive of results obtained in later or other larger clinical trials. A number of companies in the pharmaceutical industry have suffered significant setbacks in clinical trials, even in advanced clinical trials after showing promising results in earlier clinical trials. Our failure to adequately demonstrate the safety and effectiveness of any of our product candidates will prevent us from receiving regulatory approval to market these product candidates and will negatively impact our business. In addition, we or the FDA may suspend or curtail our clinical trials at any time if it appears that we are exposing participants to unacceptable health risks or if the FDA finds deficiencies in the conduct of these clinical trials or in the composition, manufacture or administration of the product candidates. Accordingly, we cannot predict with any certainty when or if we will ever be in a position to submit a new drug application, or NDA, for any of our product candidates, or whether any such NDA would ever be approved.

Our products use novel alternative technologies and therapeutic approaches, which have not been widely studied.

Our product development efforts focus on novel therapeutic approaches and technologies that have not been widely studied. These approaches and technologies may not be successful. We are applying these approaches and technologies in our attempt to discover new treatments for conditions that are also the subject of research and development efforts of many other companies.

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Physicians and patients may not accept and use our drugs.

Even if the FDA approves our product candidates, physicians and patients may not accept and use them. Acceptance and use of our products will depend upon a number of factors including:

- perceptions by members of the health care community, including physicians, about the safety and effectiveness of our drugs;
- cost-effectiveness of our products relative to competing products;
- availability of reimbursement for our products from government or other healthcare payers; and
- effectiveness of marketing and distribution efforts by us and our licensees and distributors, if any.

Because we expect sales of our current product candidates, if approved, to generate substantially all of our product revenues for the foreseeable future, the failure of any of these drugs to find market acceptance would harm our business and could require us to seek additional financing.

Because we are dependent on clinical research institutions and other contractors for clinical testing and for research and development activities, the results of our clinical trials and such research activities are, to a certain extent, not within our control.

We depend upon independent investigators and collaborators, such as universities and medical institutions, to conduct our pre-clinical and clinical trials under agreements with us. These parties are not our employees and we cannot control the amount or timing of resources that they devote to our programs. These investigators may not assign as great a priority to our programs or pursue them as diligently as we would if we were undertaking such programs ourselves. If outside collaborators fail to devote sufficient time and resources to our drug development programs, or if their performance is substandard, the approval of our FDA applications, if any, and our introduction of new drugs, if any, will be delayed. These collaborators may also have relationships with other commercial entities, some of whom may compete with us. If our collaborators assist our competitors at our expense, our competitive position would be harmed.

Our reliance on third parties to formulate and manufacture our product candidates exposes us to a number of risks that may delay the development, regulatory approval and commercialization of our products or result in higher product costs.

We have no experience in drug formulation or manufacturing and do not intend to establish our own manufacturing facilities. We lack the resources and expertise to formulate or manufacture our own product candidates. Instead, we will contract with one or more manufacturers to manufacture, supply, store and distribute drug supplies for our clinical trials. If any of our product candidates receive FDA approval, we will rely on one or more third-party contractors to manufacture our drugs. Our anticipated future reliance on a limited number of third-party manufacturers exposes us to the following risks:

- We may be unable to identify manufacturers on acceptable terms or at all because the number of potential manufacturers is limited and the FDA must approve any replacement contractor. This approval would require new testing and compliance inspections. In addition, a new manufacturer would have to be educated in, or develop substantially equivalent processes for, production of our products after receipt of FDA approval, if any.
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Our third-party manufacturers might be unable to formulate and manufacture our drugs in the volume and of the quality required to meet our clinical and/or commercial needs, if any.

- Our future contract manufacturers may not perform as agreed or may not remain in the contract manufacturing business for the time required to supply our clinical trials or to successfully produce, store and distribute our products.

- Drug manufacturers are subject to ongoing periodic unannounced inspection by the FDA and corresponding state agencies to ensure strict compliance with good manufacturing practice and other government regulations and corresponding foreign standards. We do not have control over third-party manufacturers' compliance with these regulations and standards, but we will be ultimately responsible for any of their failures.

- If any third-party manufacturer makes improvements in the manufacturing process for our products, we may not own, or may have to share, the intellectual property rights to the innovation. This may prohibit us from seeking alternative or additional manufacturers for our products.

Each of these risks could delay our clinical trials, the approval, if any, of our product candidates by the FDA, or the commercialization of our product candidates or result in higher costs or deprive us of potential product revenues.

We have no experience selling, marketing or distributing products and no internal capability to do so.

We currently have no sales, marketing or distribution capabilities. We do not anticipate having resources in the foreseeable future to allocate to the sales and marketing of our proposed products. Our future success depends, in part, on our ability to enter into and maintain sales and marketing collaborative relationships, the collaborator's strategic interest in the products under development and such collaborator's ability to successfully market and sell any such products. We intend to pursue collaborative arrangements regarding the sales and marketing of our products, however, there can be no assurance that we will be able to establish or maintain such collaborative arrangements, or if able to do so, that they will have effective sales forces. To the extent that we decide not to, or are unable to, enter into collaborative arrangements with respect to the sales and marketing of our proposed products, significant capital expenditures, management resources and time will be required to establish and develop an in-house marketing and sales force with technical expertise. There can also be no assurance that we will be able to establish or maintain relationships with third-party collaborators or develop in-house sales and distribution capabilities. To the extent that we depend on third parties for marketing and distribution, any revenues we receive will depend upon the efforts of such third parties, and there can be no assurance that such efforts will be successful. In addition, there can also be no assurance that we will be able to market and sell our product in the U.S. or overseas.

If we cannot compete successfully for market share against other drug companies, we may not achieve sufficient product revenues and our business will suffer.

The market for our product candidates is characterized by intense competition and rapid technological advances. If our product candidates receive FDA approval, they will compete with a number of existing and future drugs and therapies developed, manufactured and marketed by others. Existing or future competing products may provide greater therapeutic convenience or clinical or other benefits for a specific indication than our products, or may offer comparable performance at a lower cost. If our products fail to capture and maintain market share, we may not achieve sufficient product revenues and our business will suffer.

We will compete against fully integrated pharmaceutical companies and smaller companies that are collaborating with larger pharmaceutical companies, academic institutions, government agencies and other public and private research organizations. Many of these competitors have technologies already approved or in development. In addition, many of these competitors, either alone or together with their collaborative partners, operate larger research and development programs and have substantially greater financial resources than we do, as well as significantly greater experience in:

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- developing drugs;
- undertaking pre-clinical testing and human clinical trials;
- obtaining FDA and other regulatory approvals of drugs;
- formulating and manufacturing drugs; and
- launching, marketing and selling drugs.

Developments by competitors may render our products or technologies obsolete or non-competitive.

The biotechnology and pharmaceutical industries are intensely competitive and subject to rapid and significant technological change. The drugs that we are attempting to develop will have to compete with existing therapies. In addition, a large number of companies are pursuing the development of pharmaceuticals that target the same diseases and conditions that we are targeting. We face competition from pharmaceutical and biotechnology companies in the U.S. and abroad. In addition, companies pursuing different but related fields represent substantial competition. Many of these organizations competing with us have substantially greater capital resources, larger research and development staffs and facilities, longer drug development history in obtaining regulatory approvals and greater manufacturing and marketing capabilities than we do. These organizations also compete with us to attract qualified personnel and parties for acquisitions, joint ventures or other collaborations.

Our ability to generate product revenues will be diminished if our drugs sell for inadequate prices or patients are unable to obtain adequate levels of reimbursement.

Our ability to commercialize our drugs, alone or with collaborators, will depend in part on the extent to which reimbursement will be available from:

- government and health administration authorities;
- private health maintenance organizations and health insurers; and
- other healthcare payers.

Significant uncertainty exists as to the reimbursement status of newly approved healthcare products. Healthcare payers, including Medicare, are challenging the prices charged for medical products and services. Government and other healthcare payers increasingly attempt to contain healthcare costs by limiting both coverage and the level of reimbursement for drugs. Even if our product candidates are approved by the FDA, insurance coverage may not be available, and reimbursement levels may be inadequate, to cover our drugs. If government and other healthcare payers do not provide adequate coverage and reimbursement levels for any of our products, once approved, market acceptance of our products could be reduced.

We may be exposed to liability claims associated with the use of hazardous materials and chemicals.

Our research and development activities may involve the controlled use of hazardous materials and chemicals. Although we believe that our safety procedures for using, storing, handling and disposing of these materials comply with federal, state and local laws and regulations, we cannot completely eliminate the risk of accidental injury or contamination from these materials. In the event of such an accident, we could be held liable for any resulting damages and any liability could materially adversely effect our business, financial condition and results of operations.

In addition, the federal, state and local laws and regulations governing the use, manufacture, storage, handling and disposal of hazardous or radioactive materials and waste products may require us to incur substantial compliance costs that could materially adversely affect our business, financial condition and results of operations.

Risks Related to Our Intellectual Property

If we fail to protect or enforce our intellectual property rights adequately or secure rights to patents of others, the value of our intellectual property rights would diminish.

Our success, competitive position and future revenues will depend in part on our ability and the abilities of our licensors to obtain and maintain patent protection for our products, methods, processes and other technologies, to preserve our trade secrets, to prevent third parties from infringing on our proprietary rights and to operate without infringing upon the proprietary rights of third parties. Additionally, if any third-party manufacturer makes improvements in the manufacturing process for our products, we may not own, or may have to share, the intellectual property rights to the innovation.

To date, we hold certain exclusive rights under U.S. patents and patent applications as well as rights under foreign patent applications. We anticipate filing additional patent applications both in the U.S. and in other countries, as appropriate. However, we cannot predict:

- the degree and range of protection any patents will afford us against competitors including whether third parties will find ways to invalidate or otherwise circumvent our patents;
- if and when patents will issue;
- whether or not others will obtain patents claiming aspects similar to those covered by our patents and patent applications; or
- whether we will need to initiate litigation or administrative proceedings which may be costly whether we win or lose.

If any of our trade secrets, know-how or other proprietary information is disclosed, the value of our trade secrets, know-how and other proprietary rights would be significantly impaired and our business and competitive position would suffer.

Our success also depends upon the skills, knowledge and experience of our scientific and technical personnel, our consultants and advisors as well as our licensors and contractors. To help protect our proprietary know-how and our inventions for which patents may be unobtainable or difficult to obtain, we rely on trade secret protection and confidentiality agreements. To this end, we require all of our employees, consultants, advisors and contractors to enter into agreements which prohibit the disclosure of confidential information and, where applicable, require disclosure and assignment to us of the ideas, developments, discoveries and inventions important to our business. These agreements may not provide adequate protection for our trade secrets, know-how or other proprietary information in the event of any unauthorized use or disclosure or the lawful development by others of such information. If any of our trade secrets, know-how or other proprietary information is disclosed, the value of our trade secrets, know-how and other proprietary rights would be significantly impaired and our business and competitive position would suffer.

If we infringe upon the rights of third parties we could be prevented from selling products, forced to pay damages, and defend against litigation.

If our products, methods, processes and other technologies infringe upon the proprietary rights of other parties, we could incur substantial costs and we may have to:

- obtain licenses, which may not be available on commercially reasonable terms, if at all;

- redesign our products or processes to avoid infringement;
- stop using the subject matter claimed in the patents held by others;

- pay damages; or
- defend litigation or administrative proceedings which may be costly whether we win or lose, and which could result in a substantial diversion of our valuable management resources.

If requirements under our license agreements are not met, we could suffer significant harm, including losing rights to our products.

We depend on licensing agreements with third parties to maintain the intellectual property rights to our products under development. Presently, we have licensed rights from the University of Pittsburgh and The Ohio State University Research Foundation. These agreements require us and our licensors to perform certain obligations that affect our rights under these licensing agreements. All of these agreements last either throughout the life of the patents, or with respect to other licensed technology, for a number of years after the first commercial sale of the relevant product.

In addition, we are responsible for the cost of filing and prosecuting certain patent applications and maintaining certain issued patents licensed to us. If we do not meet our obligations under our license agreements in a timely manner, we could lose the rights to our proprietary technology.

Finally, we may be required to obtain licenses to patents or other proprietary rights of third parties in connection with the development and use of our products and technologies. Licenses required under any such patents or proprietary rights might not be made available on terms acceptable to us, if at all.

Risks Related to Our Securities

We cannot assure you that our common stock will ever be listed on NASDAQ or any other securities exchange.

Our common stock is currently eligible for trading on the “Pink Sheets.” Stocks traded on the Pink Sheets and other electronic over-the-counter markets are often less liquid than stocks traded on national securities exchanges. In fact, the historical trading of our common stock has been extremely limited and sporadic. We plan to seek listing on NASDAQ or the American Stock Exchange in the future, but we cannot assure you that we will be able to meet the initial listing standards of either of those or any other stock exchange, or that we will be able to maintain a listing of our common stock on either of those or any other stock exchange. To the extent that our common stock is not traded on a national securities exchange, such as NASDAQ, the decreased liquidity of our common stock may make it more difficult to sell shares of our common stock at desirable times and at prices.

Our common stock is considered a “penny stock.”

The SEC has adopted regulations which generally define a “penny stock” to be an equity security that has a market price of less than $5.00 per share, subject to specific exemptions. Since trading of our common stock commenced, the market price has been below $5.00 per share. Therefore, our common stock is deemed a “penny stock” according to SEC rules. This designation requires any broker or dealer selling these securities to disclose certain information concerning the transaction, obtain a written agreement from the purchaser and determine that the purchaser is reasonably suitable to purchase the securities. These rules may restrict the ability of brokers or dealers to sell shares of our common stock.

Because we did not become public through an underwritten initial public offering, we may not be able to attract the attention of major brokerage firms.

Additional risks may exist since we did not become public through a an initial public offering underwritten by an investment bank. Security analysts of major brokerage firms may not provide coverage of us since there is no

incentive to brokerage firms to recommend the purchase of our common stock. No assurance can be given that brokerage firms will want to conduct any secondary offerings on behalf of our company in the future. The lack of such analyst coverage may decrease the public demand for our common stock, making it more difficult for you to resell your shares when you deem appropriate.

Because we do not expect to pay dividends, you will not realize any income from an investment in our common stock unless and until you sell your shares at profit.

We have never paid dividends on our common stock and do not anticipate paying any dividends for the foreseeable future. You should not rely on an investment in our common stock if you require dividend income. Further, you will only realize income on an investment in our shares in the event you sell or otherwise dispose of your shares at a price higher than the price you paid for your shares. Such a gain would result only from an increase in the market price of our common stock, which is uncertain and unpredictable.

There may be issuances of shares of blank check preferred stock in the future.

Our amended and restated certificate of incorporation authorizes the issuance of up to 35,000,000 shares of preferred stock, all of which are currently designated as Series A Convertible Preferred Stock. Following the conversion of our Series A Convertible Preferred Stock into common stock upon the effectiveness of the registration statement of which this prospectus is a part, our board of directors will again have the authority to fix and determine the relative rights and preferences of up to 35,000,000 preferred shares, as well as the authority to issue such shares, without further stockholder approval. As a result, our board of directors could authorize the issuance of a series of preferred stock that is senior to our common stock and that would grant to holders preferred rights to our assets upon liquidation, the right to receive dividends, additional registration rights, anti-dilution protection, the right to the redemption to such shares, together with other rights, none of which will be afforded holders of our common stock.

If our results do not meet analysts' forecasts and expectations, our stock price could decline.

In the future, analysts who cover our business and operations may provide valuations regarding our stock price and make recommendations whether to buy, hold or sell our stock. Our stock price may be dependent upon such valuations and recommendations. Analysts' valuations and recommendations are based primarily on our reported results and their forecasts and expectations concerning our future results regarding, for example, expenses, revenues, clinical trials, regulatory marketing approvals and competition. Our future results are subject to substantial uncertainty, and we may fail to meet or exceed analysts' forecasts and expectations as a result of a number of factors, including those discussed above under the sections "Risks Related to Our Business" and "Risks Related to the Clinical Testing, Regulatory Approval, Manufacturing and Commercialization of Our Product Candidates." If our results do not meet analysts' forecasts and expectations, our stock price could decline as a result of analysts lowering their valuations and recommendations or otherwise.

We are at risk of securities class action litigation.

In the past, securities class action litigation has often been brought against a company following a decline in the market price of its securities. This risk is especially relevant for us because biotechnology companies have experienced greater than average stock price volatility in recent years. If we faced such litigation, it could result in substantial costs and a diversion of our management's attention and resources, which could harm our business.

NOTE REGARDING FORWARD-LOOKING STATEMENTS

This prospectus contains "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933, as amended, or the Securities Act, and Section 21E of the Securities Exchange Act of 1934, as amended, or the Exchange Act. The forward-looking statements are only predictions and provide our current expectations or forecasts of future events and financial performance and may be identified by the use of forward-looking terminology, including the terms "believes," "estimates," "anticipates," "expects," "plans," "intends," "may," "will" or "should" or, in each c negative, or other variations or comparable terminology, though the absence of these words does not necessarily mean that a statement is not forward-looking. Forward-looking statements include all matters that are not historical facts and include, without limitation, statements concerning our business strategy, outlook, objectives, future milestones, plans, intentions, goals, future financial conditions, our research and development programs and planning for and timing of any clinical trials, the possibility, timing and outcome of submitting regulatory filings for our product candidates under development, research and development of particular drug products, the development of financial, clinical, manufacturing and marketing plans related to the potential approval and commercialization of our drug products, and the period of time for which our existing resources will enable us to fund our operations.

Forward-looking statements are subject to many risks and uncertainties that could cause our actual results to differ materially from any future results expressed or implied by the forward-looking statements. Examples of the risks and uncertainties include, but are not limited to:

- the risk that recurring losses, negative cash flows and the inability to raise additional capital could threaten our ability to continue as a going concern;

- the risk that we may not successfully develop and market our product candidates, and even if we do, we may not become profitable;

- risks relating to the progress of our research and development;

- risks relating to significant, time-consuming and costly research and development efforts, including pre-clinical studies, clinical trials and testing, and the risk that clinical trials of our product candidates may be delayed, halted or fail;

- risks relating to the rigorous regulatory approval process required for any products that we may develop independently, with our development partners or in connection with any collaboration arrangements;

- the risk that changes in the national or international political and regulatory environment may make it more difficult to gain FDA or other regulatory approval of our drug product candidates;

- risks that the FDA or other regulatory authorities may not accept any applications we file;

- risks that the FDA or other regulatory authorities may withhold or delay consideration of any applications that we file or limit such applications to particular indications or apply other label limitations;

- risks that, after acceptance and review of applications that we file, the FDA or other regulatory authorities will not approve the marketing and sale of our drug product candidates;

- risks relating to our drug manufacturing operations, including those of our third-party suppliers and contract manufacturers;

- risks relating to the ability of our development partners and third-party suppliers of materials, drug substance and related components to provide us with adequate supplies and expertise to support manufacture of drug product for initiation and completion of our clinical studies;
- risks relating to the transfer of our manufacturing technology to third-party contract manufacturers; and
- other risks and uncertainties detailed in “Risk Factors.”

Pharmaceutical and biotechnology companies have suffered significant setbacks in advanced clinical trials, even after obtaining promising earlier trial results. Data obtained from such clinical trials are susceptible to varying interpretations, which could delay, limit or prevent regulatory approval. Except to the extent required by applicable laws or rules, we do not undertake to update any forward-looking statements or to publicly announce revisions to any of our forward-looking statements, whether resulting from new information, future events or otherwise.

USE OF PROCEEDS

We will receive none of the proceeds from the sale of the shares by the selling stockholders, except for the warrant exercise price upon exercise of the warrants, which would be used for working capital and other general corporate purposes.

SELLING STOCKHOLDERS

This prospectus covers the resale by the selling stockholders identified below of 26,815,831 shares of our common stock, as follows:

- 2,466,057 shares that are currently issued and outstanding;
- 15,274,000 shares are issuable upon the conversion of our outstanding Series A Preferred Stock,
- Up to 381,844 shares of our common stock that may be issued in payment of accrued dividends upon the conversion of our Series A Preferred Stock; and
- 8,693,930 shares are issuable upon the exercise of outstanding warrants.

The following table sets forth the number of shares of our common stock beneficially owned by the selling stockholders as of October 31, 2010 and after giving effect to this offering, except as otherwise referenced below.

Selling Stockholder	Shares beneficially owned before offering (1)(2)	Number of outstanding shares offered by selling stockholder	Number of shares offered by selling stockholder upon conversion of preferred stock (2)	Number of shares offered by selling stockholder upon exercise of warrants	Percentage beneficial ownership after offering(1)
3071341 Canada Inc. (3)	79,402	-	25,625	12,500	*
6984321 Canada Inc. (4)	22,467	-	6,150	3,000	*
87111 Canada Limited (5)	100,041	-	25,625	12,500	*
Alan T. Yuasa as Trustee of the Michael J. Shimoko Trust	152,500	-	102,500	50,000	-
Allan Pantuck and Jodi Pantuck (JTWROS)	18,300	-	12,300	6,000	-
Allen Rubin (6)	19,467	-	6,150	3,000	*
Alrac Investments Inc. (7)	19,062	-	12,812	6,250	-
Arie and Rebecka Belldegrun as Trustees of the Belldegrun Family Trust dated February 18, 1994 (8)	1,358,675	-	128,125	62,500	1.3
Benjamin Bernstein (9)	493,691	-	-	75,000	*
Canyon Value Realization Fund (Cayman) Ltd. (10)	1,011,330	687,651	-	-	-
Canyon Value Realization Fund, L.P. (10)	1,011,330	263,069	-	-	-
Canyon Value Realization Mac-18 Ltd. (10)	1,011,330	60,610	-	-	-

Clal Finance Underwriting Ltd. (11)	3,132,227	9,968	-	-	-
Clal Insurance Company Ltd. – Profit Participating Policies (11)	3,132,227	825,578	717,500	350,000	-
Clal Pension & Provident Funds Ltd. – Sapir (11)	3,132,227	412,788	272,650	133,000	-
Clal Pension & Provident Funds Ltd. – Yahalom (11)	3,132,227	206,393	137,350	67,000	-
Commercial Street Capital, LLC (12)	3,355,000	-	2,255,000	1,100,000	-

Selling Stockholder	Shares beneficially owned before offering (1)(2)	Number of outstanding shares offered by selling stockholder	Number of shares offered by selling stockholder upon conversion of preferred stock (2)	Number of shares offered by selling stockholder upon exercise of warrants	Percentage beneficial ownership after offering(1)
DAFNA Life Science Ltd. (13)	152,500	-	24,600	12,000	-
DAFNA Life Science Market Neutral Ltd. (13)	152,500	-	18,450	9,000	-
DAFNA Life Science Select Ltd. (13)	152,500	-	59,450	29,000	-
David M. Tanen (14)	1,634,345	-	66,625	99,618	3.3
Dikla Insurance Company Ltd. (15)	28,620	-	9,528	4,648	*
Esperante AB (16)	762,500	-	512,500	250,000	-
FCC Ltd (17)	763,800	-	307,500	456,300	-
Genesis Capital Advisors, LLC (18)	2,135,000	-	153,750	75,000	-
Genesis Opportunity Fund, LP (18)	2,135,000	-	1,281,250	625,000	-
Georgette Pagano	38,125	-	25,625	12,500	-
Hank C.K. Wuh	38,125	-	25,625	12,500	-
Harel Insurance Company Ltd. (19)	821,010	-	208,446	101,681	*
Harel Pension Fund Management Company Ltd. (19)	821,010	-	53,531	26,113	*
Harel Provident Funds Ltd. (19)	821,010	-	35,993	17,558	*
I-Bankers Securities, Inc. (20)	207,750	-	82,000	40,000	-
IBS Securities Ltd. (20)	207,750	-	-	85,750	-
Ira Kalfus	53,375	-	35,875	17,500	-
Isaac Kier (21)	275,627	-	51,250	25,000	*
Jill Wallace	2,500	-	-	2,500	-
Joia Kazam and Joshua Kazam (JTWROS) (22)	2,059,301	-	256,250	125,000	3.6
Joshua Kazam (22)	2,059,301	-	-	80,728	3.6
Kappa Investors LLC (23)	3,896,788	-	172,210	84,005	4.5
Kardan Israel Ltd. (24)	1,525,000	-	1,025,000	500,000	-
Kenzo Kosuda	76,250	-	51,250	25,000	-
Leumi Overseas Trust Corporation Limited as Trustee of the BTL Trust (8)	1,358,675	-	256,250	125,000	1.3
MDRB Partnership, L.P. (8)	1,358,675	-	128,125	62,500	1.3
Nazy Zomorodian	45,750	-	30,750	15,000	-
Ogier Employee Benefit Trust Limited as Trustees of the MBES Employee Benefit Trust – JD Sub Trust (25)	152,500	-	102,500	50,000	-
Peter Kash (26)	2,034,921	-	-	189,534	3.8
Peter Kash and Donna Kash (JTWROS) (26)	2,034,921	-	102,500	50,000	3.8
Pontifax (Cayman) II L.P. (27)	4,574,998	-	1,503,174	733,256	-
	4,574,998	-	439,540	214,410	-

Pontifax (Israel) II - Individual Investors L.P. (27)					
Pontifax (Israel) II L.P. (27)	4,574,998	-	1,132,284	552,334	-
Primafides (Suisse) SA as Trustees of the Sirius Trust (28)	694,306	-	102,500	50,000	1.2
Ricardo de la Guardia	38,125	-	25,625	12,500	-
Robert I. Falk (29)	397,722	-	102,500	50,000	*
Sabrinco Inc. (30)	39,699		12,812	6,250	*

Selling Stockholder	Shares beneficially owned before offering (1)(2)	Number of outstanding shares offered by selling stockholder	Number of shares offered by selling stockholder upon conversion of preferred stock (2)	Number of shares offered by selling stockholder upon exercise of warrants	Percentage beneficial ownership after offering(1)
Scott Navins (31)	249,532	-	-	100,000	*
Sherry Hyon	5,000	-	-	5,000	-
Steven Blum (32)	164,875	-	-	125,000	*
Susumu Maeda	76,250	-	51,250	25,000	-
Taichi Wakabayashi	76,250	-	51,250	25,000	-
UTA Capital LLC (33)	3,050,000	-	2,050,000	1,000,000	-
Uzi Zucker	457,500	-	307,500	150,000	-
Wexford Spectrum Investors LLC (23)	3,896,788	-	1,098,789	535,995	4.5
Yu Yeung (34)	39,937	-	-	20,000	*
TOTAL		2,466,057	15,655,844	8,693,930	

* denotes less than 1%

(1) Beneficial ownership is determined in accordance with Rule 13d-3 under the Securities Act, and includes any shares as to which the security or stockholder has sole or shared voting power or investment power, and also any shares which the security or stockholder has the right to acquire within 60 days of the date hereof, whether through the exercise or conversion of any stock option, convertible security, warrant or other right. The indication herein that shares are beneficially owned is not an admission on the part of the security or stockholder that he, she or it is a direct or indirect beneficial owner of those shares. Percentage of shares beneficially owned after the resale of all the shares offered by this prospectus assumes there are outstanding 44,761,798 shares of common stock, including all shares offered hereby that are issuable upon the conversion of our outstanding Series A Preferred Stock or upon the exercise of warrants.

(2) Assumes six months of dividend accrual at the rate of 5% per annum prior to the automatic conversion of the Series A Preferred Stock into common stock upon the effectiveness of the registration statement of which this prospectus is a part, and our election to pay such accrued dividends in the form of additional shares of common stock in lieu of cash. See “Description of Capital Stock – Series A Convertible Preferred Stock.”

(3) Ruth Hornstein is the president and sole owner of the selling stockholder. In addition to the shares offered hereby, the selling stockholder beneficially owns 41,277 shares of our common stock.

(4) Danny Ritter is the president of the selling stockholder. In addition to the shares offered hereby, beneficial ownership includes 10,317 shares of our common stock held by Mr. Ritter.

(5) Hershie Schachter, president of the selling stockholder, holds voting and/or dispositive power over the shares held by the selling stockholder. In addition to the shares offered hereby, the selling stockholder beneficially owns 61,916 shares of our common stock.

(6) In addition to the shares offered hereby, the selling stockholder beneficially owns 10,317 shares of our common stock.

(7) Lawrence Stein is the director of the selling stockholder.

(8) Beneficial ownership includes: (i) 128,125 shares of our common stock that are issuable upon the conversion of our outstanding Series A Preferred Stock and 62,500 shares issuable upon the exercise of warrants held by Arie and Rebecka Belldegrun as Trustees of the Belldegrun Family Trust dated February 18, 1994; (ii) 256,250 shares of our common stock that are issuable upon the conversion of our outstanding Series A Preferred Stock, 125,000 shares issuable upon the exercise of warrants and 61,916 shares of our common stock held by Leumi Overseas Trust Corporation Limited as Trustee of the BTL Trust; (iii) 128,125 shares of our common stock that are issuable upon the conversion of our outstanding Series A Preferred Stock and 62,500 shares issuable upon the exercise of warrants held by MDRB Partnership, L.P. (“MDRB”); and (iv) 24,922 shares of our common stock and 509,337 shares issuable upon the exercise of stock options held by Arie Belldegrun, M.D. Dr. Belldegrun, who serves as Chairman of our Board of Directors, is a beneficiary of the BTL Trust and is the managing partner of MDRB. Richard J. Guillaume and Christopher R.P. Lees, directors of Leumi Overseas Trust Corporation Limited (“Leumi”), hold voting and/or dispositive power over the shares held by Leumi as trustee of the BTL Trust.

(9) In addition to the shares offered hereby, the selling stockholder beneficially owns 418,691 shares of our common stock.

(10) John Simpson, Joshua S. Friedman, Mitchell R. Julius and John P. Plaga have voting and/or dispositive power over the shares held by the selling stockholder. The selling stockholder has informed us that it is affiliated with a broker-dealer, and has represented to us that it purchased the shares in the ordinary course of business with no agreement or understanding, directly or indirectly, with any persons regarding the distribution of the shares.

(11) Beneficial ownership includes: (i) 825,578 shares of our common stock held by Clal Insurance Company Ltd. – Profit Participating Policies; (ii) 412,788 shares of our common stock held by Clal Pension & Provident Funds Ltd. – Sapir ("Sapir"); (iii) 206,393 shares of our common stock held by Clal Pension & Provident Funds Ltd. – Yahalom ("Yahalom"); and (iv) 9,968 shares of our common stock held by Clal Finance Underwriting Ltd. Yossi Dori holds voting and/or dispositive power over the shares held by Sapir and Yahalom. Nir Moroz holds voting and/or dispositive power over the shares held by Clal Insurance Company Ltd. – Profit Participating Policies.

(12) Steven Ruchefsky, President of Commercial Street Capital, LLC, is a director of Arno.

(13) Fariba Ghodsian is the managing member of the selling stockholder.

(14) Mr. Tanen is our President and a member of our board of directors. Shares listed as beneficially owned by Mr. Tanen include 149,532 shares of our common stock held by Mr. Tanen's wife as custodian for the benefit of their minor children under the Uniform Gift to Minors Act (UGMA), for which Mr. Tanen disclaims any beneficial ownership. In addition to the shares offered hereby, beneficial ownership also includes 1,307,334 shares of our common stock and 11,236 shares issuable upon the exercise of options and warrants held by Mr. Tanen.

(15) Alfred Rosenfeld and Ofer Nargassi hold voting and/or dispositive power over the shares held by the selling stockholder. In addition to the shares offered hereby, beneficial ownership includes: (i) 4,127 shares of our common stock held by Dikla Insurance Company Ltd. – Nostro; and (ii) 10,317 shares of our common stock held by Dikla Insurance Company Ltd. – Siudi.

(16) Dean Slagel, director of the selling stockholder, holds voting and/or dispositive power over the shares held by the selling stockholder.

(17) Yacov Reizman, chairman and chief executive officer of the selling stockholder, and Rivka Reizman, president of the selling stockholder, hold voting and/or dispositive power over the shares held by the selling stockholder. Mr. Reizman is a director of Arno.

(18) Jaime Hartman is the managing member of the selling stockholder.

(19) Ronen Agassi and Ofer Nargassi hold voting and/or dispositive power over the shares held by the selling stockholder. In addition to the shares offered hereby, beneficial ownership includes: (i) 20,637 shares of our common stock held by Harel Insurance Company Ltd. – Clali; (ii) 115,580 shares of our common stock held by Harel Insurance Company Ltd. – Mishtatefet; (iii) 45,406 shares of our common stock held by Harel Insurance Company Ltd. – Nostro; (iv) 41,277 shares of our common stock held by Harel Pension Fund Management Company Ltd. – Harel Pensia; (v) 28,893 shares of our common stock held by Harel Provident Funds Ltd. – Taoz; (vi) 10,317 shares of our common stock held by Harel Provident Funds, Ltd. – Hishtalmut; (vii) 8,254 shares of our common stock held by Harel Provident Funds, Ltd. – Gmisha; and (viii) 107,324 shares of our common stock held by Harel Provident Funds, Ltd. – Otzma.

(20) Shelley Gluck holds voting and/or dispositive power over the shares held by the selling stockholder. I-Bankers Securities, Inc. ("I-Bankers") is a registered broker-dealer and the shares offered by I-Bankers are issuable upon the

exercise of warrants received as compensation for placement agent services in connection with our September 2010 private placement. IBS Securities Ltd. ("IBS") is an affiliate of I-Bankers and acquired the shares offered hereby in the ordinary course of its business.

(21) In addition to the shares offered hereby, the selling stockholder beneficially owns 199,377 shares of our common stock.

(22) Shares listed as beneficially owned by the selling stockholders include, in addition to the shares offered hereby, (i) 1,129,759 shares of our common stock and 14,946 shares issuable upon the exercise of options and warrants held by Mr. Kazam; (ii) 99,688 shares of our common stock held by Mrs. Kazam as custodian for the benefit of their minor daughter under the UGMA; (iii) 332,293 shares of our common stock held by the Kazam Family Trust; and (iv) 20,637 shares of our common stock held by the Joshua Kazam Trust.

(23) In addition to the shares offered hereby, beneficial ownership includes: (i) 247,345 shares of our common stock held by Kappa Investors LLC ("Kappa"); (ii) a warrant held by Kappa to purchase 24,732 shares of our common stock that is exercisable at $2.42 per share; and (iii) 1,733,712 shares of our common stock held by Wexford Spectrum Investors LLC ("Wexford Spectrum"). Wexford Capital LP, a Delaware partnership ("Wexford Capital"), is a registered Investment Advisor and also serves as an investment advisor or sub-advisor to the members of Kappa and Wexford Spectrum. Wexford GP LLC ("Wexford GP") is the general partner of Wexford Capital. Mr. Charles E. Davidson and Mr. Joseph M. Jacobs are managing and controlling members of Wexford GP.

(24) Eytan Rechter is chief executive officer and a director of the selling stockholder and Asher Elmoznino is chief financial officer of the selling stockholder.

(25) Tania Bearryman and Donna Laverty hold voting and/or dispositive power over the shares held by the selling stockholder.

(26) Peter Kash is a member of our board of directors. Shares listed as beneficially owned by Mr. Kash include 358,876 shares of our common stock held by Mr. Kash's wife as custodian for the benefit of their minor children under the UGMA, for which Mr. Kash disclaims any beneficial ownership. In addition to the shares offered hereby, beneficial ownership also includes 1,327,629 shares of our common stock and 12,472 shares issuable upon the exercise of options and warrants held by Mr. Kash.

(27) Tomer Kariv and Ran Nussbaum hold voting and/or dispositive power over the shares held by the selling stockholder. Mr. Kariv is a director of Arno.

(28) Ari Tatos, Nigel Mifsud, Magali Garcia-Baudin, David Moran, Phillippe De Salis and Ewald Scherrer are directors of Primafides (Suisse) SA, the trustee of the Sirius Trust, and share voting and/or dispositive power over the shares held by the selling stockholder. In addition to the shares offered hereby, the selling stockholder beneficially owns 541,806 shares of our common stock.

(29) In addition to the shares offered hereby, beneficial ownership includes: (i) 49,844 shares of our common stock and warrants to purchase 4,946 shares of our common stock at an exercise price of $2.42 per share held by Falk Family Partners, LP, of which Mr. Falk is General Partner; and (ii) 90,744 shares of our common stock and vested options held by Mr. Falk to purchase 99,688 shares of our common stock at an exercise price of $2.42 per share.

(30) Samuel Gewurz is the president and sole owner of the selling stockholder. In addition to the shares offered hereby, the selling stockholder beneficially owns 20,637 shares of our common stock.

(31) In addition to the shares offered hereby, Mr. Navins, who serves as Arno's Treasurer, beneficially owns 149,532 shares of our common stock.

(32) In addition to the shares offered hereby, the selling stockholder beneficially owns 39,875 shares of our common stock.

(33) YZT Management LLC ("YZT") is the managing member of the selling stockholder. Udi Toledano is the managing member of YZT and holds voting and/or dispositive power over the shares held by the selling stockholder.

(34) In addition to the shares offered hereby, the selling stockholder beneficially owns 19,937 shares of our common stock.

PLAN OF DISTRIBUTION

We are registering the shares offered by this prospectus on behalf of the selling stockholders. The selling stockholders, which as used herein includes donees, pledgees, transferees or other successors-in-interest selling shares of common stock or interests in shares of common stock received after the date of this prospectus from a selling stockholder as a gift, pledge, partnership distribution or other transfer, may, from time to time, sell, transfer or otherwise dispose of any or all of their shares of common stock or interests in shares of common stock on any stock exchange, market or trading facility on which the shares are traded or in private transactions. These dispositions may be at fixed prices, at prevailing market prices at the time of sale, at prices related to the prevailing market price, at varying prices determined at the time of sale, or at negotiated prices. To the extent any of the selling stockholders gift, pledge or otherwise transfer the shares offered hereby, such transferees may offer and sell the shares from time to time under this prospectus, provided that this prospectus has been amended under Rule 424(b)(3) or other applicable

provision of the Securities Act to include the name of such transferee in the list of selling stockholders under this prospectus.

The selling stockholders may use any one or more of the following methods when disposing of shares or interests therein:

- ordinary brokerage transactions and transactions in which the broker-dealer solicits purchasers;
- block trades in which the broker-dealer will attempt to sell the shares as agent, but may position and resell a portion of the block as principal to facilitate the transaction;
- purchases by a broker-dealer as principal and resale by the broker-dealer for its account;
- an exchange distribution in accordance with the rules of the applicable exchange;

- privately negotiated transactions;
- short sales;
- through the writing or settlement of options or other hedging transactions, whether through an options exchange or otherwise;
- broker-dealers may agree with the selling stockholders to sell a specified number of such shares at a stipulated price per share;
- a combination of any such methods of sale; and
- any other method permitted pursuant to applicable law.

The selling stockholders may, from time to time, pledge or grant a security interest in some or all of the shares of common stock owned by them and, if they default in the performance of their secured obligations, the pledgees or secured parties may offer and sell the shares of common stock, from time to time, under this prospectus, or under an amendment to this prospectus under Rule 424(b)(3) or other applicable provision of the Securities Act amending the list of selling stockholders to include the pledgee, transferee or other successors in interest as selling stockholders under this prospectus.

In connection with the sale of our common stock or interests therein, the selling stockholders may enter into hedging transactions with broker-dealers or other financial institutions, which may in turn engage in short sales of the common stock in the course of hedging the positions they assume. The selling stockholders may also sell shares of our common stock short and deliver these securities to close out their short positions, or loan or pledge the common stock to broker-dealers that in turn may sell these securities. The selling stockholders may also enter into option or other transactions with broker-dealers or other financial institutions or the creation of one or more derivative securities which require the delivery to such broker-dealer or other financial institution of shares offered by this prospectus, which shares such broker-dealer or other financial institution may resell pursuant to this prospectus (as supplemented or amended to reflect such transaction).

The aggregate proceeds to the selling stockholders from the sale of the common stock offered by them will be the purchase price of the common stock less discounts or commissions, if any. Each of the selling stockholders reserves the right to accept and, together with their agents from time to time, to reject, in whole or in part, any proposed purchase of common stock to be made directly or through agents. We will not receive any of the proceeds from this offering. Upon any exercise of the warrants by payment of cash, however, we will receive the exercise price of the warrants.

The selling stockholders also may resell all or a portion of the shares in open market transactions in reliance upon Rule 144 under the Securities Act, provided that they meet the criteria and conform to the requirements of that rule.

The selling stockholders might be, and any broker-dealers that act in connection with the sale of securities will be, deemed to be “underwriters” within the meaning of Section 2(11) of the Securities Act, and any commissions received by such broker-dealers and any profit on the resale of the securities sold by them while acting as principals will be deemed to be underwriting discounts or commissions under the Securities Act.

To the extent required, the shares of our common stock to be sold, the names of the selling stockholders, the respective purchase prices and public offering prices, the names of any agents, dealer or underwriter, any applicable commissions or discounts with respect to a particular offer will be set forth in an accompanying prospectus

supplement or, if appropriate, a post-effective amendment to the registration statement that includes this prospectus.

In order to comply with the securities laws of some states, if applicable, the common stock may be sold in these jurisdictions only through registered or licensed brokers or dealers. In addition, in some states the common stock may not be sold unless it has been registered or qualified for sale or an exemption from registration or qualification requirements is available and is complied with.

We have advised the selling stockholders that the anti-manipulation rules of Regulation M under the Exchange Act may apply to sales of shares in the market and to the activities of the selling stockholders and their affiliates. In addition, we will make copies of this prospectus (as it may be supplemented or amended from time to time) available to the selling stockholders for the purpose of satisfying the prospectus delivery requirements of the Securities Act. The selling stockholders may indemnify any broker-dealer that participates in transactions involving the sale of the shares against certain liabilities, including liabilities arising under the Securities Act.

We have agreed to indemnify the selling stockholders against liabilities, including liabilities under the Securities Act and state securities laws, relating to the registration of the shares offered by this prospectus.

We have agreed with the selling stockholders to keep the registration statement that includes this prospectus effective until the earlier of (1) such time as all of the shares covered by this prospectus have been disposed of pursuant to and in accordance with the registration statement or (2) the date on which the shares may be sold without restriction pursuant to Rule 144 of the Securities Act.

Shares Eligible For Future Sale

Upon completion of this offering and assuming the issuance of all shares offered hereby that are issuable upon the conversion of our outstanding Series A Preferred Stock or upon exercise of warrants, there will be 44,584,906 shares of our common stock issued and outstanding. The shares purchased in this offering will be freely tradable without registration or other restriction under the Securities Act, except for any shares purchased by an “affiliate” of our company (as defined in the Securities Act).

The selling stockholders also may resell all or a portion of the shares in open market transactions in reliance upon Rule 144 under the Securities Act, provided they meet the criteria and conform to the requirements of such Rule. Rule 144 governs resale of “restricted securities” for the account of any person (other than us), and restricted and unrestricted securities for the account of an “affiliate” of ours. Restricted securities generally include any securities acquired directly or indirectly from us or our affiliates, which were not issued or sold in connection with a public offering registered under the Securities Act. An affiliate of ours is any person who directly or indirectly controls us, is controlled by us, or is under common control with us. Our affiliates may include our directors, executive officers, and persons directly or indirectly owing 10% or more of our outstanding common stock. In general, under Rule 144, a person (or persons whose shares are aggregated) who is not deemed to have been an affiliate of ours at the time of, or at any time during the three months preceding, a sale, and who has beneficially owned restricted securities for at least six months would be entitled to sell those shares, subject to the requirements of Rule 144 regarding publicly available information about us. Affiliates may only sell in any three month period that number of shares that does not exceed the greater of 1 percent of the then-outstanding shares of our common stock or the average weekly trading volume of our shares of common stock in the over-the-counter market during the four calendar weeks preceding the sale. However, because we were formerly a “shell company,” in order for the holders of our restricted securities to resell their shares in reliance upon Rule 144, we are required to have been subject to the public reporting requirements of the Exchange Act for at least 90 days, and to have filed all reports required to be filed during the 12 months preceding such sale (or such shorter period that we were required to file such reports). Further, Rule 144 will not be available to the selling stockholders until one year has elapsed from the date we have filed “Form 10 information” with the SEC, which generally consists of all the information and disclosures concerning our business, financial information, management and other information that is required to be included in a registration statement filed under the Exchange

Act. Accordingly, the shares held by the selling stockholders will become eligible for sale under Rule 144 one year after the filing of the registration statement of which this prospectus is a part.

Following the date of this prospectus, we cannot predict the effect, if any, that sales of our common stock or the availability of our common stock for sale will have on the market price prevailing from time to time. Nevertheless, sales by existing stockholders of substantial amounts of our common stock could adversely affect prevailing market prices for our stock.

DESCRIPTION OF CAPITAL STOCK

General

Our amended and restated certificate of incorporation authorizes us to issue 115,000,000 shares of capital stock, par value $0.0001 per share, comprised of 80,000,000 shares of common stock, and 35,000,000 million shares of preferred stock.

As of the date of this prospectus, we have issued and outstanding approximately:

- 20,412,024 shares of our common stock,
- 15,274,000 shares of our Series A Preferred Stock,
- warrants to purchase 8,693,930 shares of Series A Preferred Stock,
- options to purchase 1,763,303 shares of our common stock at exercise prices ranging from $0.25 to $3.00 per share, and
- warrants to purchase 495,252 shares of our common stock at exercise prices ranging from $1.00 to $2.42 per share.

Common Stock

The holders of our common stock are entitled to one vote for each share held of record on all matters submitted to a vote of the stockholders and do not have cumulative voting rights. Upon our liquidation, dissolution or winding down, holders of our common stock will be entitled to share ratably in all of our assets that are legally available for distribution, after payment of all debts and other liabilities. The holders of our common stock have no preemptive, subscription, redemption or conversion rights.

Holders of our common stock are entitled to receive such dividends, as the board of directors may from time to time declare out of funds legally available for the payment of dividends. We seek growth and expansion of our business through the reinvestment of profits, if any, and do not anticipate that we will pay dividends in the foreseeable future.

Series A Convertible Preferred Stock

The terms, conditions, privileges, rights and preferences of our Series A Convertible Preferred Stock are described in a Certificate of Designation filed with the Secretary of State of Delaware on September 3, 2010. The following provides only a brief summary of certain of the terms of our Series A Preferred Stock.

Conversion. Each share of our Series A Preferred Stock is initially convertible at the holder's election into one share of our common stock. Upon the effective date of the registration statement of which this prospectus is a part, each share of Series A Preferred Stock will automatically convert into one share of common stock.

Voting. Along with the holders of our common stock, the holders of our Series A Preferred Stock are entitled to one vote on all matters submitted to the holders of common stock for each share of common stock into which the Series A Preferred Stock would be converted as of the record date for such vote based on the conversion ratio then in effect. In addition, the holders of the Series A Preferred Stock are entitled to vote as a separate class with respect to any change in the rights of the Series A Preferred Stock, any amendment to our certificate of incorporation, any increase in the number of shares of Series A Preferred Stock, or the authorization, creation or issuance of any class or series of capital stock ranking senior to or of equal seniority with the Series A Preferred Stock.

Dividends. The holders of our Series A Preferred Stock are entitled to an annual per share cumulative dividend equal to 5% of the original issuance price of $1.00 per share, which dividends are payable upon the conversion of the Series A Preferred Stock into common stock, and which we may elect to pay in the form of additional shares of common stock in lieu of cash. Solely as an example of the foregoing, if the registration statement of which this prospectus is a part is declared effective by the SEC after the Series A Preferred Stock has been issued and outstanding for six months, the total accrued dividend preference on the 15,274,000 shares of Series A Preferred Stock issued in our September 2010 private placement would be $381,850, which we could elect to pay by issuing approximately an additional 381,850 shares of common stock upon the automatic conversion of the Series A Preferred Stock into common stock. The holders of our Series A Preferred Stock are entitled to payment of all accrued dividends prior to the payment of any dividends to the holders of our common stock. Following payment of such accrued dividends, the holders of our Series A Preferred Stock are entitled to participate with the holders of our common stock in any other dividend payment on an as-converted basis.

Liquidation. Upon the liquidation, dissolution or winding up of our company, whether voluntary or involuntary, the holders of our Series A Preferred Stock are entitled to be paid, prior to any payments to the holders of our common stock, an amount per share equal to the sum of (i) 1.5 times the original issuance price of $1.00 per share, plus (ii) any accrued but unpaid dividends on the Series A Preferred Stock.

Authority to Issue Stock

Our board of directors has the authority to issue the authorized but unissued shares of our common stock without action by the shareholders. The issuance of such shares would reduce the percentage ownership held by current shareholders.

Our amended and restated certificate of incorporation authorizes the issuance of up to 35,000,000 shares of preferred stock, all of which are currently designated as Series A Convertible Preferred Stock. Following the conversion of our Series A Preferred Stock into common stock, our board of directors will again have the authority to fix and determine the relative rights and preferences of up to 35,000,000 preferred shares, as well as the authority to issue such shares, without further stockholder approval. As a result, our board of directors could authorize the issuance of a series of preferred stock that is senior to our common stock and that would grant to holders preferred rights to our assets upon liquidation, the right to receive dividends, additional registration rights, anti-dilution protection, the right to the redemption to such shares, together with other rights, none of which will be afforded holders of our common stock.

MARKET FOR COMMON EQUITY AND RELATED STOCKHOLDER MATTERS

Market Information

Our common stock is currently eligible for trading on the “Pink Sheets,” however, there has been only one trade in our common stock since we filed an application to deregister our common stock in May 2009. The historical trading of our common stock has been extremely limited and sporadic. Accordingly, there is not an established public trading market for our common stock.

Prior to May 2009, our common stock traded on the OTC Bulletin Board, or the OTCBB, where the first trade was reported in June 2008. Until July 16, 2008, our common stock traded under the symbol "LRRI.OB." Following our merger with Laurier completed on June 3, 2008, our trading symbol changed to "ARNI.OB" on July 17, 2008. Set forth below are the high and low sales prices for our common stock during each quarter within the last two fiscal years, and during each of the first two quarters of fiscal 2010, as reported by the Pink Sheets or the OTCBB, as applicable. The quotations reflect inter-dealer prices, without retail markup, markdown, or commission, and may not represent actual transactions. Consequently, the information provided below may not be indicative of our common stock price under different conditions.

Quarter Ended	High	Low
March 31, 2008	$ -	$ -
June 30, 2008	$ 2.00	$ 2.00
September 30, 2008	$ 3.50	$ 1.80
December 31, 2008	$ 3.25	$ 1.95
March 31, 2009	$ 1.90	$ 1.90
June 30, 2009	$ 1.50	$ 1.00
September 30, 2009	$ -	$ -
December 31, 2009	$ -	$ -
March 31, 2010	$ -	$ -
June 30, 2010	$ 0.51	$ 0.51
September 30, 2010	$ -	$ -

Stockholders

According to the records of our transfer agent, American Stock Transfer & Trust Company, as of October 31, 2010, we had approximately 275 holders of record of common stock, not including those held in “street name.”

Dividends

We have never declared or paid a dividend on our common stock and do not anticipate paying any cash dividends in the foreseeable future.

Securities Authorized for Issuance under Equity Compensation Plans

We grant stock options and other equity incentive awards pursuant to our Amended and Restated 2005 Stock Option Plan, which has been approved by our stockholders. The following table sets forth certain information as of December 31, 2009 with respect to our Amended and Restated 2005 Stock Option Plan:

Plan category	Number of Securities to be Issued Upon Exercise of Outstanding Options (A)	Weighted-Average Exercise Price of Outstanding Options (B)	Number of Securities Remaining Available for Future Issuance Under Equity Compensation Plans (Excluding Securities Reflected in Column (A)) (C)
Equity compensation plans approved by security holders:			
2005 Stock Option Plan	1,913,241	$ 1.76	1,057,414
Equity compensation plans not approved by stockholders:			
None	—	—	—

Total	1,913,241	$	1.76	1,057,414

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MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS

The following discussion and plan of operations should be read in conjunction with the financial statements and the notes to those statements included in this prospectus. This discussion includes forward-looking statements that involve risk and uncertainties. As a result of many factors, such as those set forth in this prospectus under "Risk Factors," actual results may differ materially from those anticipated in these forward-looking statements.

Company Overview

We are a development stage company focused on developing innovative products for the treatment of cancer. We currently have the exclusive worldwide rights to commercially develop three oncology product candidates:

- AR-12 – Our lead clinical product candidate is a potentially first-in-class, orally available, targeted anti-cancer agent that inhibits phosphoinositide dependent protein kinase-1, or PDK-1, a protein in the PI3K/Akt pathway involved in the growth and proliferation of cells, including cancer cells. We believe AR-12 may also cause cell death through the induction of stress in the endoplasmic reticulum. In May 2009, the FDA accepted our investigational new drug application, or IND, for AR-12. We are currently conducting a multi-centered Phase I clinical study of AR-12 in adult patients with advanced or recurrent solid tumors or lymphoma. The Phase I study of AR-12 is being conducted in two parts. The first part is a dose-escalating study, which we refer to as the Escalation Phase, primarily designed to evaluate the compound's safety in order to identify the maximum tolerated dose, or MTD, or a recommended dose, or RD, for future studies of AR-12. We anticipate that the Escalation Phase will be completed by the third quarter of 2011. Following the Escalation Phase, we plan to initiate the second part of the study, which involves enrolling an expanded cohort of additional patients at the MTD or RD in multiple tumor types. We refer to this second part of the study as the Expansion Phase. The purpose of the Expansion Phase is to further evaluate and confirm the PD effects, potential anti-tumor activity, and safety of AR-12 at the MTD or RD. We anticipate that the Expansion Phase will be fully enrolled within one year.

- AR-42 – We are also developing AR-42, an orally available, broad spectrum inhibitor of both histone and non-histone deacetylation proteins, which play an important role in the regulation of gene expression, cell growth and survival. In preclinical studies, AR-42 has demonstrated greater potency and activity in solid and liquid tumors when compared to vorinostat (also known as SAHA and marketed as Zolinza® by Merck) and other deacetylase inhibitors. These data demonstrate the potent and differentiating activity of AR-42. Additionally, pre-clinical findings presented at the 2009 American Society of Hematology Annual Meeting and Exposition showed that AR-42 potently and selectively inhibits leukemic stem cells in acute myeloid leukemia, or AML. AR-42 is currently being studied in an investigator initiated Phase I/IIa clinical study in adult patients with relapsed or refractory multiple myeloma, chronic lymphocytic leukemia, or CLL, or lymphoma. Once the MTD is defined, the study is designed so that additional patients can be added to investigate efficacy in a particular disease and help guide future Phase II programs. Up to an additional 10 patients may be enrolled at the MTD dose in each of the multiple myeloma, CLL and lymphoma.

- AR-67 – We are also developing AR-67, a novel, third-generation campothecin analogue that inhibits Topoisomerase I activity. In 2008, we completed a multi-centered, ascending dose Phase I clinical trial of AR-67 in patients with advanced solid tumors. AR-67 is currently being studied in a Phase II clinical trial in patients with glioblastoma multiforme, or GBM, a highly aggressive form of brain cancer. We anticipate having interim data from this Phase II study by the second quarter of 2011. Thereafter, if data permits, we may elect to initiate larger Phase II studies or advance AR-67 into a registration-enabling Phase III study.

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We have no product sales to date and we will not generate any product revenue until we receive approval from the U.S. Food and Drug Administration, or the FDA, or equivalent foreign regulatory bodies to begin selling our pharmaceutical product candidates. Developing pharmaceutical products is a lengthy and very expensive process. Assuming we do not encounter any unforeseen safety or other issues during the course of developing our product candidates, we do not expect to complete the development of a product candidate for several years, if ever. To date, a significant amount of our development expenses have related to our lead product candidate, AR-12. As we proceed with the clinical development of AR-12 and as we further develop our other product candidates, AR-42 and AR-67, our research and development expenses will further increase. To the extent we are successful in acquiring additional product candidates for our development pipeline, our need to finance further research and development will continue increasing. Accordingly, our success depends not only on the safety and efficacy of our product candidates, but also on our ability to finance the development of the products. To date, our major sources of working capital have been proceeds from private and public sales of our common stock, and debt financings.

Research and development, or R&D, expenses consist primarily of salaries and related personnel costs, fees paid to consultants and outside service providers for pre-clinical, clinical, and manufacturing development, legal expenses resulting from intellectual property prosecution, costs related to obtaining and maintaining our product license agreements, contractual review, and other expenses relating to the design, development, testing, and enhancement of our product candidates. We expense our R&D costs as they are incurred.

General and administrative, or G&A, expenses consist primarily of salaries and related expenses for executive, finance and other administrative personnel, personnel recruiting fees, accounting, legal and other professional fees, business development expenses, rent, business insurance and other corporate expenses.

Our results include non-cash compensation expense as a result of the issuance of stock, stock options and warrants. We expense the fair value of stock options and warrants over the vesting period. When more precise pricing data is unavailable, we determine the fair value of stock options using the Black-Scholes option-pricing model. The terms and vesting schedules for share-based awards vary by type of grant and the employment status of the grantee. Generally, the awards vest based upon time-based or performance-based conditions. Performance-based conditions generally include the attainment of goals related to our financial performance and product development. Stock-based compensation expense is included in the respective categories of expense in the statements of operations. We expect to record additional non-cash compensation expense in the future, which may be significant.

Results of Operations

General and Administrative Expenses. G&A expenses for the three months ended June 30, 2010 and 2009 were approximately $0.2 million and $0.1 million, respectively. The increase of approximately $0.1 million over 2009 is primarily due to Board of Directors compensation in 2009 versus none in 2008. G&A expenses for the six months ended June 30, 2010 and 2009 were approximately $0.3 million and $0.6 million, respectively. The decrease of approximately $0.3 million over 2009 is primarily due to a reduction in employee headcount from three to zero. G&A expenses for the years ended December 31, 2009 and 2008 were approximately $1.5 million and $2.3 million, respectively. The decrease of approximately $0.8 million over 2008 is primarily due to costs incurred during 2008 related to our June 2008 reverse merger, becoming a publicly traded company and a reduction in employee compensation costs, including stock compensation costs, resulting from the reduced headcount from three to one employee. These reductions in 2009 over 2008 were partially offset by Board of Directors compensation of approximately $0.2 million for 2009 not incurred in 2008 and increased office rent expense.

Research and Development Expenses. R&D expenses for the three months ended June 30, 2010 and 2009 were approximately $1.1 million and $1.2 million, respectively. The decrease of approximately $0.1 million over the three month period in 2009 is due primarily to reduced employee compensation costs from a reduced headcount with one

less employee and from a reduction in clinical trial costs related to AR-67 of approximately $0.1 million offset by an increase in manufacturing costs for AR-12 of approximately $0.1 million. R&D expenses for the six months ended June 30, 2010 and 2009 were approximately $2.1 million and $3.2 million, respectively. The decrease of approximately $1.1 million over the six month period in 2009 was primarily due to reductions in AR-67 clinical, manufacturing, and contractual costs as well as reductions in employee compensation costs due to reduced headcount. Additionally, costs related to AR-42 manufacturing and preclinical activities were reduced by approximately $0.3 million over 2009. R&D expenses for the years ended December 31, 2009 and 2008 were approximately $5.4 million and $9.8 million, respectively. The decrease of approximately $4.4 million over 2008 was primarily due to reduced manufacturing, clinical and nonclinical activities for AR-67 of approximately $1.7 million and reduced nonclinical and contractual activities for AR-42 of approximately $1.7 million. Additional decreases resulted from reduced manufacturing, nonclinical and contractual activities for AR-12 of approximately $0.8 million.

The following table summarizes our R&D expenses incurred for preclinical support, contract manufacturing for clinical supplies, clinical trial services provided by third parties and milestone payments for in-licensed technology for each of our product candidates for the three months ended June 30, 2010 and 2009, the six months ended June 30, 2010 and 2009, the years ended December 31, 2009 and 2008, as well as the cumulative amounts since we began development of each product candidate through June 30, 2010. The table also summarizes unallocated costs, which consist of personnel, facilities and other costs not directly allocable to development programs:

	Three Months Ended June 30,		Six Months Ended June 30,		Years Ended December 31,		Cummulative amounts during development
	2010	2009	2010	2009	2009	2008	
AR-12	$ 383,148	$ 271,489	$ 1,120,981	$ 1,036,443	$ 1,999,469	$ 2,820,442	$ 5,974,269
AR-67	276,876	369,120	201,129	727,819	1,075,141	2,801,368	6,827,847
AR-42	23,081	50,677	37,438	321,697	541,066	2,263,440	2,866,763
Unallocated R&D	384,565	469,226	770,415	1,150,938	1,828,526	1,883,139	4,938,772
Total	$ 1,067,670	$ 1,160,512	$ 2,129,963	$ 3,236,897	$ 5,444,202	$ 9,768,389	$ 20,607,651

AR-12. Our lead clinical product candidate is a potentially first-in-class, orally available, targeted anti-cancer agent that inhibits phosphoinositide dependent protein kinase-1, or PDK-1, a protein in the PI3K/Akt pathway, and may also cause cell death through the induction of endoplasmic reticulum stress. In May 2009, the FDA accepted our investigational new drug application, or IND, for AR-12. We are currently conducting a multi-centered Phase I clinical study of AR-12 in adult patients with advanced or recurrent solid tumors or lymphoma. We expect to complete the Escalation Phase of the study in the first quarter of 2011; however, the Expansion Phase of the study may not be completed until the end of 2011. We then expect to begin Phase II development of AR-12 by the first half of 2012. Based on our current development plans for AR-12, we anticipate spending approximately $1.1 million on external development costs during the second half of 2010.

AR-42. We are also developing AR-42, an orally available, broad spectrum inhibitor of both histone and non-histone deacetylation proteins. AR-42 is currently being studied in an investigator sponsored Phase I/IIa clinical study in adult patients with relapsed or refractory multiple myeloma, chronic lymphocytic leukemia or lymphoma. In preclinical studies, AR-42 has demonstrated greater potency and activity in solid and liquid tumors when compared to vorinostat (also known as SAHA and marketed as Zolinza® by Merck) and other deacetylase inhibitors. These data demonstrate the potent and differentiating activity of AR-42. Additionally, pre-clinical findings presented at the 2009 American Society of Hematology Annual Meeting and Exposition showed that AR-42 potently and selectively inhibits leukemic stem cells in acute myeloid leukemia. Based on our current development plans for AR-42, we anticipate spending approximately $0.1 million on external development costs during the second half of 2010.

AR-67. We are also developing AR-67, a novel, third-generation camptothecin analogue that inhibits Topoisomerase I activity. In late 2008, we completed a multi-centered, ascending dose Phase I clinical trial of AR-67 in patients with advanced solid tumors. AR-67 is currently being studied in a Phase II clinical in patients with glioblastoma multiforme, or GBM, a highly aggressive form of brain cancer. We anticipate having interim Phase II data from each study by the second quarter of 2011, at which point we may elect to initiate larger Phase II studies or advance AR-67 into a registration-enabling Phase III study. If the data is positive and we elect to initiate a larger new trial, we may spend significant additional amounts to conduct such a study. Based on our current development plans for AR-67, we anticipate spending approximately $0.5 million on external development costs during the second half of 2010.

Our expenditures on current and future clinical development programs are expected to be substantial, particularly in relation to our available capital resources, and to increase. However, these planned expenditures are subject to many uncertainties, including the results of clinical trials and whether we develop any of our drug candidates with a partner or independently. As a result of such uncertainties, we cannot predict the duration and completion costs of our research and development projects or whether, when and to what extent we will generate revenues from the commercialization and sale of any of our product candidates. The duration and cost of clinical trials may vary significantly over the life of a project as a result of unanticipated events arising during clinical development and a variety of factors, including:

- the number of trials and studies in a clinical program;
- the number of patients who participate in the trials;
- the number of sites included in the trials;
- the rates of patient recruitment and enrollment;
- the duration of patient treatment and follow-up;
- the costs and timing of manufacturing our drug candidates; and
- the costs, requirements, timing of, and the ability to secure regulatory approvals.

Interest Income. Interest income for the three and six months ended June 30, 2010 and 2009 was $1,925, $3,417, $1,066 and $26,197, respectively. Interest income for the years ended December 31, 2009 and 2008 was $28,145 and $206,054. The decrease in interest income over the earlier period is due to lower interest rates earned on cash in bank accounts, and lower average cash balances levels.

Liquidity and Capital Resources

The following tables summarize our liquidity and capital resources as of June 30, 2010, December 31, 2009 and June 30, 2009 and our net decrease in cash and cash equivalents for the six months ended June 30, 2010 and 2009 and for the year ended December 31, 2009 (the amounts stated are expressed in thousands):

Liquidity and capital resources	June 30, 2010	December 31, 2009	June 30, 2009
Cash and cash equivalents	$ 709	$ 3,087	$ 5,633
Working capital	$ (536)	$ 1,506	$ 4,128
Stockholders' (deficit) equity	$ (472)	$ 1,576	$ 4,207

Cash flow data	Six Months Ended June 30, 2010	Six Months Ended June 30, 2009	Year Ended December 31, 2009
Cash provided by (used in):			
Operating activities	$ (2,379)	$ (4,762)	$ (7,310)
Investing activities	-	-	-
Financing activities	-	-	3
Net decrease in cash and cash equivalents	$ (2,379)	$ (4,762)	$ (7,307)

Our total cash resources as of June 30, 2010 were approximately $0.7 million compared to $3.1 million as of December 31, 2009 and $5.6 million as of June 30, 2009. As of June 30, 2010, we had approximately $1.4 million in liabilities, and $(0.5) million in net working capital. We incurred a net loss of $2.1 million and had negative cash flow from operating activities of $2.4 million for the six months ended June 30, 2010. Since August 1, 2005 (inception) through June 30, 2010, we have incurred an aggregate net loss of approximately $25.7 million, while negative cash flow from operating activities has amounted to $20.8 million. As we continue to develop our product candidates, we expect to continue to incur substantial and increasing losses, which will continue to generate negative net cash flows from operating activities as we expand our technology portfolio and engage in further research and development activities, particularly the conducting of pre-clinical studies and clinical trials.

From inception through June 30, 2010, we have financed our operations through private sales of our equity and debt securities. As we have not generated any revenue from operations to date, and we do not expect to generate revenue for several years, if ever, we will need to raise substantial additional capital in order to continue to fund our research and development, including our long-term plans for clinical trials and new product development, as well as to fund operations generally. We may seek to raise additional funds through various potential sources, such as equity and debt financings, or through strategic collaborations and license agreements. We can give no assurances that we will be able to secure such additional sources of funds to support or operations, or if such funds are available to us, that such additional financing will be sufficient to meet our needs.

During September 2010, we completed a private placement of our Series A Preferred Stock resulting in net proceeds of approximately $13.9 million after cash issuance costs.

Based on our resources at June 30, 2010, and together with the net proceeds from our September 2010 financing, we believe that we have sufficient capital to fund our operations through to the second half of 2012. However, based on the various options for future clinical studies of AR-12 yet to be determined and pending results of our soon to be completed Phase II clinical trial of AR-67, our projected cash needs is difficult to predict. In addition, there are other factor which may also cause our actual cash requirements to vary materially, including the changes in the focus and direction of our research and development programs, including the acquisition and pursuit of development of new product candidates; competitive and technical advances; costs of commercializing any of the product candidates; and costs of filing, prosecuting, defending and enforcing any patent claims and any other intellectual property rights. If we are unable to raise additional funds when needed, we may not be able to market our products as planned or continue development and regulatory approval of our products, we could be required to delay, scale back or eliminate some or all our research and development programs and we may need to wind down our operations altogether. Each of these alternatives would likely have a material adverse effect on our business.

The actual amount of funds we will need to operate is subject to many factors, some of which are beyond our control. These factors include the following:

- the progress of our research activities;
- the number and scope of our research programs;
- the progress of our pre-clinical and clinical development activities;
- the costs and timing of manufacturing our drug candidates;
- the progress of the development efforts of parties with whom we have entered into research and development agreements;
- our ability to maintain current research and development programs and to establish new research and development and licensing arrangements; and
- the cost involved in prosecuting and enforcing patent claims and other intellectual property rights; and the cost and timing of regulatory approvals.

We have based our estimates on assumptions that may prove to be wrong. We may need to obtain additional funds sooner than planned or in greater amounts than we currently anticipate. Potential sources of financing include strategic relationships, public or private sales of equity or debt and other sources. We may seek to access the public or private equity markets when conditions are favorable due to our long-term capital requirements. We do not have any committed sources of financing at this time, and it is uncertain whether additional funding will be available when we need it on terms that will be acceptable to us, or at all. If we raise funds by selling additional shares of common stock or other securities convertible into common stock, the ownership interests of our existing stockholders will be diluted. If we are not able to obtain financing when needed, we may be unable to carry out our business plan. As a result, we may have to significantly limit our operations and our business, financial condition and results of operations would be materially harmed. In such an event, we will be required to undertake a thorough review of our programs and the opportunities presented by such programs and allocate our resources in the manner most prudent.

To the extent that we raise additional funds by issuing equity or convertible or non-convertible debt securities, our stockholders may experience additional significant dilution and such financing may involve restrictive covenants. To the extent that we raise additional funds through collaboration and licensing arrangements, it may be necessary to relinquish some rights to our technologies or our product candidates, or grant licenses on terms that may not be favorable to us. These things may have a material adverse effect on our business.

The continuation of our business beyond the second half of 2012 is dependent upon obtaining further long-term financing, the successful development of our drug product candidates and related technologies, the successful and sufficient market acceptance of any product offerings that we may introduce, and, finally, the achievement of a profitable level of operations. The issuance of additional equity securities by us may result in a significant dilution in the equity interests of current stockholders. Obtaining commercial loans, assuming those loans would be available, on acceptable terms or even at all, will increase our liabilities and future cash commitments.

OUR BUSINESS

Overview

We are a development stage company focused on developing innovative products for the treatment of cancer. We currently have the exclusive worldwide rights to commercially develop three oncology product candidates. The following table summarizes our product development pipeline:

Product Candidate	Indications	Commercial Rights	Ongoing Studies / Status
AR-12	Solid tumors and hematological cancers	Arno	A two part, multi-centered Phase I clinical trial of AR-12 is ongoing in patients with solid tumors and lymphoma is ongoing who have progressed despite treatment with other therapies.
AR-42	Hematological cancers	Arno	Investigator initiated Phase I/IIa clinical study of AR-42 is ongoing at The James Cancer Center at The Ohio State University in patients with advanced or recurrent hematological cancers for which no treatment is available.
AR-67	Glioblastoma multiforme (GBM)	Arno	A two-cohort, multi-center Phase II clinical trial is ongoing in patients with GBM that have progressed with other therapies. The first cohort will enroll up to 26 patients who experienced rapid progression after treatment with bevacizumab (Avastin®,Roche). The second cohort will enroll up to 32 patients who have not been treated with Avastin within the last 90 days.

Corporate History; Merger Transactions

On June 2, 2008, we were acquired by Laurier International, Inc., a Delaware corporation, in a “reverse” merger whereby a wholly-owned subsidiary of Laurier merged with and into Arno Therapeutics, with Arno Therapeutics remaining as the surviving corporation and a wholly-owned subsidiary of Laurier. In accordance with the terms of this merger, stockholders of Arno Therapeutics exchanged all of their shares of common stock of Arno Therapeutics for shares of Laurier common stock at a rate of 1.99377 shares of Laurier common stock for each share of Arno Therapeutics common stock. As a result of the issuance of the shares of Laurier common stock to the former Arno Therapeutics stockholders, following the merger the former stockholders of Arno Therapeutics held 95 percent of the outstanding common stock of Laurier, assuming the issuance of all shares underlying outstanding options and warrants. Upon completion of the merger, all of the former officers and directors of Laurier resigned and were replaced by the officers and directors of Arno Therapeutics. Additionally, following the merger Laurier changed its name to Arno Therapeutics, Inc.

Oncology Overview

Cancer is the second leading cause of death in the United States, surpassed only by heart disease. Since 1990, over 18 million new cancer cases have been diagnosed. According to a 2009 report by the American Cancer Society, the National Institutes of Health estimate direct costs for medical care for cancer related treatments in the United States in

2007 were $93.0 billion. With a 66% 5-year relative survival rate for all cancers from 1996-2004, oncology remains a significant unmet medical need.

Pharmaceutical treatments are widely used to combat cancer and are often used alongside surgery or radiation, when possible. Different types of cancers respond in unique ways to different drugs, and some tumors may not respond at all to particular therapies. In many cases, these treatments extend life by slowing the progression of the disease but become less effective over time as the cancer cells become resistant to a given therapy or a class of compounds with a particular mechanism of action. For this reason, there is a need to develop new agents, particularly those with novel mechanisms that can be added to the current arsenal of treatment options.

Many types of drugs are presently used to treat cancer, including cytotoxics, targeted agents, hormones, and biologics. According to an October 2008 report by Cowen & Co., the global cancer market was roughly $63.0 billion in 2007, of which cytotoxics and targeted agents each accounted for 33%, or $21.0 billion.

Cytotoxics interfere with essential cellular processes in order to kill rapidly dividing cells, an effective approach for destroying cancer cells that remains prevalent despite the fact that these compounds can have significant side effects, particularly in rapidly dividing normal tissues such as those found in bone marrow and the gastrointestinal tract. By contrast, targeted agents attack cellular processes that are more prevalent in cancer cells than in normal tissues, and thus aim to simultaneously reduce side effects and improve efficacy.

Although there are many agents available to treat cancer, a number of factors contribute to determining which particular agent is administered to a patient. There is a considerable amount of overlap in the mechanisms of action of approved therapies, and in many cases, multiple drugs in a class are approved and in clinical use. The choice of a particular agent or class of agents is generally based on the results of empirical clinical trials in specific cancer indications, and a desire to treat the disease aggressively is balanced with considerations for the patient's comfort and quality of life. These considerations highlight the need to develop therapies that not only improve anti-cancer efficacy but also improve patient convenience and reduce side effects.

Product Development Pipeline

AR-12

Overview

We are developing AR-12, a potentially first-in-class, orally available cancer treatment that is currently enrolling patients in a Phase I clinical study. AR-12 is an inhibitor of phosphoinositide dependent protein kinase-1, or PDK-1, that targets the Akt pathway, while also possessing activity in the endoplasmic reticulum stress pathway and other pathways targeting apoptosis. In pre-clinical studies, AR-12 has demonstrated activity in a wide range of tumor types and synergistic effects with several widely used anti-cancer agents, enhancing activity or overcoming drug-resistance when used in combination with Avastin® (Genentech), Herceptin® (Genentech), Gleevec® (Novartis), Tarceva® (Genentech) and tamoxifen. We are currently enrolling patients with advanced or recurrent solid tumors or lymphoma in a Phase I clinical study of AR-12.

Mechanism of Action

AR-12 is an inhibitor of a protein known as PDK-1, a novel target in an important cell growth and proliferation pathway. This pathway has been validated by the approval of therapeutics that target proteins both upstream and downstream of PDK-1. Receptor tyrosine kinases, or RTK, are cell-surface receptors that are involved in cell growth and are upstream of PDK-1. Members of the RTK class are targeted by some of the most successful and widely used targeted oncology agents, including Avastin® (Genentech), Herceptin® (Genentech), Gleevec® (Novartis), Tarceva ® (Genentech), Iressa® (AstraZeneca), Nexavar® (Bayer/Onyx) and tamoxifen. Downstream of PDK-1 is the mammalian target of rapamycin, or mTOR protein. The mTOR inhibitors temsirolimus (Torisel®, Wyeth) and

everolimus (Afinitor®, Novartis) are FDA approved for the treatment of renal cancer, and additional studies are being conducted with mTOR inhibitors in various clinical trials as anti-cancer agents.

Although FDA-approved drugs that target the Akt pathway have shown efficacy in treating cancer, some tumors either do not respond to these drugs or eventually become resistant to therapy. Scientists hypothesize that a combination of drugs that inhibit different targets in this pathway could provide synergistic or additive benefits to increase efficacy and potentially overcome drug resistance. For this reason, there has been particular interest within the biopharmaceutical industry in developing inhibitors of PI3K, PDK-1, and Akt.

PDK-1 can transform normal human cells and may be involved in the invasion and metastasis process. PDK-1 and its downstream target, the protein Akt, are frequently activated in multiple cancer types, and inhibiting PDK-1 facilitates the dephosphorylation and subsequent inactivation of Akt. Activation of the PDK-1/Akt pathway confers resistance to cell death signaling and the apoptotic activity of other cytotoxic agents. Additionally, recent research has demonstrated the importance of PDK-1 in oncology that is independent of its Akt modulation. The inhibition of the PDK-1/Akt pathway in cancer cells where this pathway was previously active has been shown to decrease cell proliferation and increase programmed cell death, which is known as apoptosis. Preclinical data suggests that AR-12 inhibits PDK-1 and data collected from preclinical toxicology studies and tumor distribution studies indicate that AR-12 would be expected to exceed therapeutic concentrations in vivo.

AR-12 has also demonstrated an ability to induce the ER stress mediated apoptosis pathway, which contribute to its unique profile in vitro and in vivo. The ER stress pathway is a cellular mechanism that can either induce cellular protection or apoptosis. AR-12, through the induction of PKR-like Endoplasmic Reticulum Kinase, or PERK, seems to selectively induce the pro-apoptotic response and appears to have a preferential effect on cancer cells.

Potential Advantages of AR-12

We believe AR-12's unique mechanisms and ability to improve the efficacy of other approved agents may enable it to become a first-in-class agent with broad applications in oncology and significant sales in the market. In preclinical studies, AR-12 has shown efficacy in a wide range of tumor types, including breast, lung, prostate, pancreatic, brain, and hematological cancers, as both a single-agent as well as in combination with leading oncology therapeutics. AR-12 demonstrated synergy or additive benefit or overcome drug-resistance when used in combination with Avastin® , Herceptin® , Gleevec® , Tarceva® , Iressa® , Nexavar ® and tamoxifen, all of which are widely-prescribed, FDA-approved oncology therapeutics that represented approximately $10 billion in sales in 2008.

We believe AR-12 is a potentially first-in-class molecule and the only PDK-1 inhibitor currently in human clinical development. We also believe that PDK-1 may prove to be highly desirable target for oncology; therapeutic strategies to modulate the Akt pathway are of great scientific, clinical, and financial interest, and there are a few molecules in development that seek to target Akt through the inhibition a protein known as PI3K. PDK-1 is downstream of PI3K and thus may be more likely to impact the desired molecular targets further downstream and less likely to result in off-target toxicity. Inhibition of PDK-1 also seems to be able to regulate other important oncology targets that are not be mediated by PI3K. In addition to targeting PDK-1, we believe that AR-12 also has the ability to induce ER stress and has the potential to become an important agent in a range of cancer indications.

Clinical Development

We are currently enrolling subjects in a multi-site Phase I clinical trial of AR-12 in patients with solid tumors or lymphomas who have progressed despite treatment with other therapies. Subjects in this Phase I study receive an oral daily dose of AR-12. The Phase I study is being conducted at three clinical sites, including The Ohio State University (Columbus, Ohio), Scottsdale Healthcare (Phoenix, Arizona), and The Royal Marsden (London, UK).

Our current Phase I study of AR-12 is being conducted in two parts. The first part is a dose-escalating study, which we refer to as the Escalation Phase, that is primarily designed to evaluate the safety of AR-12 and to identify the maximum tolerated dose, or MTD, or a recommended dose, RD, for future clinical studies of AR-12. The study is also designed to utilize biomarkers and functional imaging to examine the pharmacodynamic effects of AR-12 in modulating certain targets within the PI3K pathway. Secondary objectives for the Escalation Phase include characterizing the phamacokinetics of AR-12 (i.e., how AR-12 is absorbed, distributed, released and eliminated in and from the body) and measuring tumor response. We also anticipate the determination of an RD or MTD with the conclusion of the Escalation Phase in the third quarter of 2011. Following the Escalation Phase, we plan to initiate the second part of the study, which we refer to as the Expansion Phase. We expect that the Expansion Phase will involve enrolling an expanded cohort of additional patients at the MTD or RD for the purpose of further evaluating and confirming the pharmacodynamic effects, potential anti-tumor activity and safety of AR-12 at the MTD or RD. We anticipate that the Expansion Phase will be fully enrolled within one year.

The biomarker selection and evaluation is being led by Johann de Bono, M.D., Ph.D. of The Royal Marsden in London. Dr. de Bono is a prominent researcher in oncology drug development and has been involved in the successful development of novel targeted therapies and compounds in the PI3K/Akt pathway.

We believe that the data generated from the current Phase I study will provide important information to direct future studies, both in terms of safety and exposure and potential efficacy. We also believe that the biomarkers and pharmacodynamic assays planned for the Phase I study may provide deeper understanding of the molecular actions of AR-12 and validate the preclinical hypothesis about AR-12's activity in a clinical setting. The information generated in these studies will also help to guide the Phase II development of AR-12. Arno expects to begin Phase II studies with AR-12 in the first half of 2012, subject to the completion and results of the ongoing Phase I study.

AR-42

We also have exclusive rights to develop and commercialize AR-42, a novel oral cancer therapy in currently in early clinical development. AR-42 is a broad spectrum deacetylase inhibitor of both histone and non-histone proteins, which has demonstrated greater potency and activity in solid and liquid tumors when compared in preclinical studies to vorinostat (also known as "SAHA" or Zolinza®), the first of only two marketed compound in the class. AR-42 may possess additional histone-independent mechanisms, which may contribute to its superior profile in vitro and in vivo. An investigator-initiated Phase I/IIa trial of AR-42 in patients with hematological cancers commenced in June 2010 at The Ohio State University.

Background of HDAC Inhibitors

Histones are proteins that play an important role in the regulation of genes. Histone modification is a key regulator of gene expression, and improper histone acetylation is among the modifications that are linked to expression of a cancerous phenotype. These changes can lead to improper cell growth resulting from altering the expression of important genes involved in cell cycle progression, proliferation, and survival. Histone deacetylases, or HDACs, are a class of enzymes that participate in this form of regulation and have been linked to both solid and hematologic malignancies and thus represent a target for cancer therapy.

HDAC inhibitors are a large emerging class of drug compounds that have demonstrated efficacy primarily in hematological malignancies, also called blood cancers, but are currently being developed in solid tumors as well. It is believed that HDAC inhibitors induce histone hyperacetylation and can cause cell death. The first drug in this class to gain approval is SAHA, which is approved to treat cutaneous T-cell lymphoma, or CTCL, in patients that have failed two previous therapies. Another HDAC inhibitor, romidepsin (Celgene Corporation) was also recently approved in CTCL. These two compounds and other HDACs are currently in late stage development for both hematological

malignancies as well as solid tumors. In preclinical studies, AR-42 has demonstrated activity against a broad spectrum of deacetylase targets and increased potency compared to SAHA.

Potential Advantages of AR-42

AR-42 is a broad spectrum inhibitor of histone and non-histone deacetylase targets that we believe may have advantages over currently approved HDAC inhibitors, including SAHA. As a result, many of our preclinical data compare AR-42 to SAHA. In preclinical models, AR-42 has shown to be more potent or effective than SAHA in various cancer types, including chronic lymphocyte leukemia, or CLL, B-cell lymphoma, prostate and ovarian cancers. Further, preclinical studies suggest that AR-42 has anti-cancer activities that are independent of histone acetylation and include Akt dephosphorylation, Bcl-xL and survivin suppression, and Ku70 acetylation, all of which disrupt the growth and proliferation of cancer cells. We believe that this combination of activity and potency could make AR-42 a more effective treatment for hematological malignancies than currently available HDAC inhibitors.

In addition to its broad activity against hematological malignancies, pre-clinical data presented at the 2009 American Society of Hematology Annual Meeting and Exposition showed that AR-42 potently and selectively inhibits leukemic stem cells in acute myeloid leukemia.

Clinical Development

We are collaborating with The Ohio State University, which commenced an investigator-initiated Phase I/IIa study for AR-42 in June 2010 in patients with advanced or recurrent hematological malignancies for which no treatment is available. The primary goal is to evaluate the safety and tolerability of AR-42 given orally three times per week. Secondary endpoints include characterizing AR-42's pharmacokinetics and its pharmacodynamic profile through the measurement of biomarkers and evaluation of clinical response.

Once the MTD is defined, the study is designed so that additional patients can be added to investigate efficacy in a particular disease and help guide future Phase II programs. Up to an additional 10 patients may be enrolled at the MTD dose in each of the following disease cohorts: CLL/small lymphocytic lymphoma, multiple myeloma, and lymphoma. Cohorts for other indications may be added at this dose with an appropriate protocol amendment.

AR-67

Background on Camptothecins

Camptothecin and its analogues, together referred to as camptothecins, are a class of drugs widely used to treat certain types of cancers, with worldwide annual sales exceeding $660 million. Camptothecins treat cancer by disrupting cell division through the inhibition of topoisomerase I, a critical enzyme in DNA replication. Through this inhibition and additional mechanisms of action, camptothecins target cancer cells preferentially to normal tissues, making them a promising class of drugs in this indication.

All clinically relevant camptothecins react with water and exist in two forms under physiologic conditions: a biologically active "lactone" form and a largely inactive but toxic "carboxylate" form. In human blood, chemical equilibrium converts the active lactone form to the inactive and toxic carboxylate form. Maintaining a therapeutic level of the lactone form in vivo has proven to be a significant challenge in the development of the class.

Second-generation camptothecin analogues focused on improving lactone stability by increasing lipophilicity and modifying binding profiles between the compound and blood proteins. Two second generation therapies, topotecan (Hycamtin®, Glaxo-Smith-Kline) and irinotecan (also known as CPT-11 and marketed as Camptostar® by Pfizer), are approved by the FDA. Topotecan, the first camptothecin to receive marketing approval in the United States, is used as a second-line intravenous therapy in several tumor types including ovarian, small cell lung cancer, and cervical cancers. Irinotecan is a largely inactive intravenous pro-drug for SN-38, a potent but insoluble camptothecin analogue.

Irinotecan is used as a front-line and second-line therapy for colorectal cancer and is by far the leading drug in the class with roughly $400 million in worldwide annual sales in 2009. While these drugs represent a marked improvement compared with the parent compound, their in vivo stability profiles remain suboptimal. Exposure to the active lactone form can be measured by lactone: total area under the curve ratio, or AUC ratio, which measures the ratio of the drug forms over the course of drug exposure. Lactone AUC ratios are 30-40% for topotecan, 40-45% for CPT-11, and 50-75% for SN-38.

AR-67 is a novel, third-generation camptothecin analogue that has demonstrated high potency in pre-clinical studies and improved pharmacokinetic properties in humans as compared with first and second-generation products. In the Phase I study of AR-67, which was completed in 2008, pharmacokinetic data suggest a lactone AUC ratio of approximately 85%.

We believe that this unique profile may translate into superior efficacy in the treatment of a variety of cancers. We believe these advantages could allow AR-67 to become a leading product in the camptothecin market.

Potential Advantages of AR-67

AR-67 has demonstrated potent topoisomerase I inhibition and greatly improved in vivo stability of the active lactone form when compared with topotecan and irinotecan. Structural characteristics make AR-67 highly lipophilic, with pre-clinical evaluation showing 10-fold and 250-fold increases in lipophilicity over SN-38 and topotecan, respectively. Favorable plasma protein binding characteristics also contribute to AR-67's superior lactone AUC ratio compared with marketed camptothecins. In the Phase I study, pharmacokinetic data demonstrated that approximately 85% of AR-67 was present in the lactone form, compared with 30-40% for topotecan, 40-45% for irinotecan, and 50-75% for SN-38. Additionally, gastrointestinal toxicities that are commonly seen with other camptothecins, such as nausea, vomiting and diarrhea, were not observed with AR-67 treatment, an important differentiator when compared to irinotecan.

Pre-clinical studies with AR-67 have demonstrated a unique anti-cancer profile, with in vitro cytotoxicity comparable to topotecan and SN-38 in several tumor lines, including non-small-cell lung and central nervous system cancers. AR-67 was used in pre-clinical xenograft studies and showed particular promise in brain cancers, where the drug significantly inhibited tumor growth and elicited complete responses in subcutaneous and intracranial glioma models. We believe that the pre-clinical evidence of AR-67's potency combined with the preliminary pharmacokinetic data observed in the Phase I study may lead to a superior therapeutic profile.

Clinical Development Program

AR-67 is currently being studied in a Phase II clinical trial in patients with glioblastoma multiforme, or GBM, a highly aggressive form of brain cancer.

GBM is the most prevalent and deadly form of brain cancer. In preclinical in vivo studies, AR-67 has shown significant tumor growth inhibition, including complete tumor regression. Pre-clinical evidence suggests that AR-67's high lipophilicity may promote blood-brain-barrier penetration of therapeutic levels of the lactone form and increase activity relative to other drugs in the class. Arno is currently enrolling patients in a two-cohort Phase II study of AR-67 as a treatment for patients with GBM that have progressed on other therapies. The first cohort will enroll up to 26 patients who have progressed rapidly (within 90 days) after treatment with Avastin, a drug recently approved in this indication. These patients' cancer normally progresses quite aggressively, and the endpoint for this cohort is two months of progression-free survival. The second cohort will enroll up to 32 patients who have not received Avastin treatment in the past 90 days and will look for six months of progression free survival. This multi-centered study includes leading centers from the United States and Canada and will be led by James J. Vrendenburgh, M.D. from Duke University.

While there can be no assurances, demonstrated efficacy in GBM, which is an orphan indication, may provide an accelerated path to approval, increased market protection and expanded sales potential.

Competition

We compete primarily in the segment of the biopharmaceutical market that addresses cancer therapeutics, which is highly competitive. We face significant competition from many pharmaceutical, biopharmaceutical and biotechnology companies that are researching and selling products designed to address the cancer market. Many of our competitors have significantly greater financial, manufacturing, marketing and drug development resources than we do. Large pharmaceutical companies in particular have extensive experience in clinical testing and in obtaining regulatory approvals for drugs. These companies also have significantly greater research capabilities than we do. In addition, many universities and private and public research institutes are active in cancer research. We also compete with commercial biotechnology companies for the rights to product candidates developed by public and private research institutes. Smaller or early-stage companies are also significant competitors, particularly those with collaborative arrangements with large and established companies. In addition to the factors described above under "Risk Factors," our ability to compete in the cancer therapeutics market depends on the following factors:

- our ability to develop novel compounds with attractive pharmaceutical properties and to secure and protect intellectual property rights based on our innovations;
- the efficacy, safety and reliability of our drug candidates;
- the speed at which we develop our drug candidates;
- our ability to design and successfully complete appropriate clinical trials;
- our ability to maintain a good relationship with regulatory authorities;
- the timing and scope of regulatory approvals;
- our ability to manufacture and sell commercial quantities of future products to the market; and
- acceptance of future products by physicians and other healthcare providers.

AR-12

AR-12 is believed to target PDK-1, which is in the PI3K/Akt pathway. Targeting the PI3K/Akt pathway has been of great interest to the pharmaceutical industry, particularly compounds that target PI3K. The approaches for targeting PI3K are either general inhibition or the specific inhibition of the alpha, beta, gamma, or delta subunit of this kinase. Some of these molecules also combine PI3K inhibition with activity against the mammalian target of rapamycin ("mTOR"), a target that is believed to also play a role in the PI3K/Akt pathway. Other approaches to this pathway include targeting Akt directly. Additionally, companies such as Wyeth, Vernalis, GlaxoSmithKline, and Novartis have published data on their preclinical discovery programs to target PDK-1.

Compounds that inhibit PI3K have been the foundation of several recent licensing, acquisition, and financing activities. Despite the great deal of activity in the space, Arno believes that AR-12 can differentiate itself and become an important agent in the treatment of cancer. In multiple preclinical studies, AR-12 has demonstrated the ability to inhibit PDK-1 as well as induce ER stress, a combination that could provide a unique therapeutic profile and differentiate AR-12 from other molecules being developed to inhibit the PI3K/Akt pathway. Arno also believes that AR-12 is the first PDK-1 inhibitor to reach clinical development, which could provide the significant advantage of being first-in-class as well as distinguishing AR-12 from PI3K inhibitors.

AR-42

If approved, AR-42 would compete with other HDAC inhibitors. HDAC inhibitors have displayed efficacy in a broad range of settings as single agents and in combination with other therapeutics. The first HDAC inhibitor to obtain

approval is vorinostat ("SAHA," or Zolinza®), which is approved for the treatment of recurrent cutaneous T-cell lymphoma ("CTCL"). Additionally, in November 2009 Gloucester Pharmaceuticals received FDA approval for its novel HDAC inhibitor, romidepsin, to treat CTCL before being acquired by Celgene Corp. in December 2009. Other compounds are in Phase II and Phase III trials, primarily in hematological malignancies, but also in solid tumors as both single agents and in combination with other oncology therapies.

We believe that AR-42 has a therapeutic profile that will allow it to compete successfully in the crowded class of what are broadly known as HDAC inhibitors. AR-42 is a pan-DAC inhibitor that has demonstrated preclinical activity that compares favorably with SAHA, as well as potentially differentiating activity in targeting cancer stem cells. Additionally, based on AR-42's preclinical toxicology package and dosing schedule, we believe that there is a relatively low risk of cardiac toxicity or fatigue. We also believe that AR-42's potential to selectively target leukemic stem cells in AML may sufficiently differentiate AR-42 from other agents in the class to become an important member of the emerging class of deacetylase inhibitors.

AR-67

If approved, we expect that AR-67 would compete in a class of chemotherapeutic agents known as camptothecins. The annual worldwide sales of camptothecins, which have been used for many years, collectively exceed $660 million. The leading camtothecins on the market today include Hycamitin (topotecan), marketed by GlaxoSmithKline, and Camptostar (irinotecan or CPT-11), which is marketed by Pfizer. If approved, our product candidates may also compete with other cytotoxic, or anticancer, therapies.

We believe that AR-67 can compete successfully with currently available camptothecin derivates as well as those currently in development. Many of the compounds that are currently marketed and in clinical development have experienced limited efficacy. We believe that AR-67's potent topoisomerase I inhibition and greatly improved lactone stability may enable the compound to demonstrate broad clinical utility and future commercial success.

License Agreements and Intellectual Property

General

Our goal is to obtain, maintain and enforce patent protection for our products, formulations, processes, methods and other proprietary technologies, preserve our trade secrets, and operate without infringing on the proprietary rights of other parties, both in the United States and in other countries. Our policy is to actively seek to obtain, where appropriate, the broadest intellectual property protection possible for our current product candidates and any future product candidates, proprietary information and proprietary technology through a combination of contractual arrangements and patents, both in the U.S. and abroad. However, even patent protection may not always afford us with complete protection against competitors who seek to circumvent our patents. If we fail to adequately protect or enforce our intellectual property rights or secure rights to patents of others, the value of our intellectual property rights would diminish. See "Risk Factors – Risks Related to Our Intellectual Property."

We will continue to depend upon the skills, knowledge and experience of our scientific and technical personnel, as well as that of our advisors, consultants and other contractors, none of which is patentable. To help protect our proprietary know-how, which is not patentable, and for inventions for which patents may be difficult to enforce, we currently rely and will in the future rely on trade secret protection and confidentiality agreements to protect our interests. To this end, we require all of our employees, consultants, advisors and other contractors to enter into confidentiality agreements that prohibit the disclosure of confidential information and, where applicable, require disclosure and assignment to us of the ideas, developments, discoveries and inventions important to our business.

AR-12 and AR-42 License Agreements

Our rights to AR-12 and AR-42 are governed by separate license agreements with The Ohio State University Research Foundation, or Ohio State, entered into in January 2008. Pursuant to each of these agreements, we have exclusive, worldwide, royalty-bearing licenses to commercialize certain patent applications, know-how and improvements relating to AR-12 and AR-42 for all therapeutic uses.

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Pursuant to our license agreements for AR-12 and AR-42, we made one-time cash payments to Ohio State in the aggregate amount of $450,000 and reimbursed it for past patent expenses. Additionally, we are required to make performance-based cash payments upon successful completion of clinical and regulatory milestones relating to AR-12 and AR-42 in the U.S., Europe and Japan. In July 2010, we paid Ohio State a milestone payment upon the commencement of the Phase I clinical study of AR-12. The first milestone payment for AR-42 will be due when the first patient is dosed in the first company-sponsored Phase I clinical trial. To the extent we enter into a sublicensing agreement relating to either or both of AR-12 or AR-42, we will be required to pay Ohio State a portion of all non-royalty income received from such sublicensee.

The license agreements with Ohio State further provide that we will indemnify Ohio State from any and all claims arising out of the death of or injury to any person or persons or out of any damage to property, or resulting from the production, manufacture, sale, use, lease, consumption or advertisement of either AR-12 or AR-42, except to the extent that any such claim arises out of the gross negligence or willful misconduct of Ohio State. The license agreements for AR-12 and AR-42, respectively, expire on the later of (i) the expiration of the last valid claim contained in any licensed patent and (ii) 20 years after the effective date of the license. Ohio State will generally be able to terminate either license upon our breach of the terms of the license the extent we fail to cure any such breach within 90 days after receiving notice of such breach or our bankruptcy. We may terminate either license upon 90 days' prior written notice.

AR-67 License Agreement

Our rights to AR-67 are governed by an October 2006 license agreement with the University of Pittsburgh, or Pitt. Under this agreement, we hold an exclusive, worldwide, royalty-bearing license for the rights to commercialize technologies embodied by certain issued patents, patent applications and know-how relating to AR-67 for all therapeutic uses. We have expanded, and intend to continue to expand, our patent portfolio by filing additional patents covering expanded uses for this technology.

Under the terms of our license agreement with Pitt, we made a one-time cash payment of $350,000 to Pitt and reimbursed it for past patent expenses. Additionally, Pitt will receive performance-based cash payments upon successful completion of clinical and regulatory milestones relating to AR-67. We will make the first milestone payment to Pitt following the filing of the first New Drug Application, or NDA, filed with the FDA for AR-67. We are also required to pay to Pitt an annual maintenance fee on each anniversary of the license agreement, and to pay Pitt a royalty equal to a percentage of net sales of AR-67. To the extent we enter into a sublicensing agreement relating to AR-67, we will pay Pitt a portion of all non-royalty income received from such sublicensee.

Under the license agreement with Pitt, we also agreed to indemnify and hold Pitt and its affiliates harmless from any and all claims, actions, demands, judgments, losses, costs, expenses, damages and liabilities (including reasonable attorneys' fees) arising out of or in connection with (i) the production, manufacture, sale, use, lease, consumption or advertisement of AR-67, (ii) the practice by us or any affiliate or sublicensee of the licensed patent; or (iii) any obligation of us under the license agreement unless any such claim is determined to have arisen out of the gross negligence, recklessness or willful misconduct of Pitt. The license agreement will terminate upon the expiration of the last patent relating to AR-67. Pitt may generally terminate the agreement at any time upon a material breach by us to the extent we fail to cure any such breach within 60 days after receiving notice of such breach or in the event we file for bankruptcy. We may terminate the agreement for any reason upon 90 days' prior written notice.

Government Regulation and Product Approval

The FDA and comparable regulatory agencies in state and local jurisdictions and in foreign countries impose substantial requirements upon the testing (pre-clinical and clinical), manufacturing, labeling, storage, recordkeeping, advertising, promotion, import, export, marketing and distribution, among other things, of drugs and drug product candidates. If we do not comply with applicable requirements, we may be fined, the regulatory agencies may refuse to approve our marketing applications or allow us to manufacture or market our products, and we may be criminally prosecuted. We and our manufacturers may also be subject to regulations under other United States federal, state, and local laws.

United States Government Regulation

In the United States, the FDA regulates drugs under the Food, Drug and Cosmetic Act, or FDCA, and implementing regulations. The process required by the FDA before our drug candidates may be marketed in the United States generally involves the following (although the FDA is given wide discretion to impose different or more stringent requirements on a case-by-case basis):

- completion of extensive pre-clinical laboratory tests, pre-clinical animal studies and formulation studies, all performed in accordance with the FDA's good laboratory practice regulations and other regulations;
- submission to the FDA of an IND application, which must become effective before clinical trials may begin;
- performance of multiple adequate and well-controlled clinical trials meeting FDA requirements to establish the safety and efficacy of the product candidate for each proposed indication;
- submission of a new drug application, or NDA, to the FDA;
- satisfactory completion of an FDA pre-approval inspection of the manufacturing facilities at which the product candidate is produced, and potentially other involved facilities as well, to assess compliance with current good manufacturing practice, or cGMP, regulations and other applicable regulations; and
- FDA review and approval of the NDA prior to any commercial marketing, sale or shipment of the drug.

The testing and approval process requires substantial time, effort and financial resources, and we cannot be certain that any approvals for our drug candidates will be granted on a timely basis, if at all. Risks to us related to these regulations are described above under the caption entitled "Risk Factors – Risks Relating to the Clinical Testing, Regulatory Approval, Manufacturing and Commercialization of Our Product Candidates."

Pre-clinical tests may include laboratory evaluation of product chemistry, formulation and stability, as well as studies to evaluate toxicity and other effects in animals. The results of pre-clinical tests, together with manufacturing information and analytical data, among other information, are submitted to the FDA as part of an IND application. Subject to certain exceptions, an IND becomes effective 30 days after receipt by the FDA, unless the FDA, within the 30-day time period, issues a clinical hold to delay a proposed clinical investigation due to concerns or questions about the conduct of the clinical trial, including concerns that human research subjects will be exposed to unreasonable health risks. In such a case, the IND sponsor and the FDA must resolve any outstanding concerns before the clinical trial can begin. Our submission of an IND, or those of our collaboration partners, may not result in the FDA authorization to commence a clinical trial. A separate submission to an existing IND must also be made for each successive clinical trial conducted during product development. The FDA must also approve changes to an existing IND. Further, an independent institutional review board, or IRB, for each medical center proposing to conduct the clinical trial must review and approve the plan for any clinical trial before it commences at that center and it must monitor the study until completed. The FDA, the IRB or the sponsor may suspend a clinical trial at any time on various grounds, including a finding that the subjects or patients are being exposed to an unacceptable health risk. Clinical testing also must satisfy extensive Good Clinical Practice requirements and regulations for informed consent.

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Clinical Trials

For purposes of NDA submission and approval, clinical trials are typically conducted in the following three sequential phases, which may overlap (although additional or different trials may be required by the FDA as well):

- Phase I clinical trials are initially conducted in a limited population to test the drug candidate for safety, dose tolerance, absorption, metabolism, distribution and excretion in healthy humans or, on occasion, in patients, such as cancer patients. In some cases, particularly in cancer trials, a sponsor may decide to conduct what is referred to as a "Phase Ib" evaluation, which is a second safety-focused Phase I clinical trial typically designed to evaluate the impact of the drug candidate in combination with currently FDA-approved drugs or in a particular patient population.

- Phase II clinical trials are generally conducted in a limited patient population to identify possible adverse effects and safety risks, to determine the efficacy of the drug candidate for specific targeted indications and to determine dose tolerance and optimal dosage. Multiple Phase II clinical trials may be conducted by the sponsor to obtain information prior to beginning larger and more expensive Phase III clinical trials. In some cases, a sponsor may decide to conduct what is referred to as a "Phase IIb" evaluation, which is a second, confirmatory Phase II clinical trial that could, if accepted by the FDA, serve as a pivotal clinical trial in the approval of a drug candidate.

- Phase III clinical trials are commonly referred to as pivotal trials. When Phase II clinical trials demonstrate that a dose range of the drug candidate is effective and has an acceptable safety profile, Phase III clinical trials are undertaken in large patient populations to further evaluate dosage, to provide substantial evidence of clinical efficacy and to further test for safety in an expanded and diverse patient population at multiple, geographically dispersed clinical trial sites.

In some cases, the FDA may condition continued approval of an NDA on the sponsor's agreement to conduct additional clinical trials with due diligence. In other cases, the sponsor and the FDA may agree that additional safety and/or efficacy data should be provided; however, continued approval of the NDA may not always depend on timely submission of such information. Such post-approval studies are typically referred to as Phase IV studies.

New Drug Application

The results of drug candidate development, pre-clinical testing and clinical trials, together with, among other things, detailed information on the manufacture and composition of the product and proposed labeling, and the payment of a user fee, are submitted to the FDA as part of an NDA. The FDA reviews all NDAs submitted before it accepts them for filing and may request additional information rather than accepting an NDA for filing. Once an NDA is accepted for filing, the FDA begins an in-depth review of the application.

During its review of an NDA, the FDA may refer the application to an advisory committee for review, evaluation and recommendation as to whether the application should be approved. The FDA may refuse to approve an NDA and issue a not approvable letter if the applicable regulatory criteria are not satisfied, or it may require additional clinical or other data, including one or more additional pivotal Phase III clinical trials. Even if such data are submitted, the FDA may ultimately decide that the NDA does not satisfy the criteria for approval. Data from clinical trials are not always conclusive and the FDA may interpret data differently than we or our collaboration partners interpret data. If the FDA's evaluations of the NDA and the clinical and manufacturing procedures and facilities are favorable, the FDA may issue either an approval letter or an approvable letter, which contains the conditions that must be met in order to secure final approval of the NDA. If and when those conditions have been met to the FDA's satisfaction, the FDA will issue an approval letter, authorizing commercial marketing of the drug for certain indications. The FDA may withdraw drug approval if ongoing regulatory requirements are not met or if safety problems occur after the drug reaches the market. In addition, the FDA may require testing, including Phase IV clinical trials, and surveillance

programs to monitor the effect of approved products that have been commercialized, and the FDA has the power to prevent or limit further marketing of a drug based on the results of these post-marketing programs. Drugs may be marketed only for the FDA-approved indications and in accordance with the FDA-approved label. Further, if there are any modifications to the drug, including changes in indications, other labeling changes, or manufacturing processes or facilities, we may be required to submit and obtain FDA approval of a new NDA or NDA supplement, which may require us to develop additional data or conduct additional pre-clinical studies and clinical trials.

The Hatch-Waxman Act

Under the Hatch-Waxman Act, newly-approved drugs and new conditions of use may benefit from a statutory period of non-patent marketing exclusivity. The Hatch-Waxman Act provides five-year marketing exclusivity to the first applicant to gain approval of an NDA for a new chemical entity, meaning that the FDA has not previously approved any other new drug containing the same active entity. The Hatch-Waxman Act prohibits the submission of an abbreviated NDA, or ANDA, or a Section 505(b)(2) NDA for another version of such drug during the five-year exclusive period; however, submission of a Section 505(b)(2) NDA or an ANDA for a generic version of a previously-approved drug containing a paragraph IV certification is permitted after four years, which may trigger a 30-month stay of approval of the ANDA or Section 505(b)(2) NDA. Protection under the Hatch-Waxman Act does not prevent the submission or approval of another "full" 505(b)(1) NDA; however, the applicant would be required to conduct its own pre-clinical and adequate and well-controlled clinical trials to demonstrate safety and effectiveness. The Hatch-Waxman Act also provides three years of marketing exclusivity for the approval of new and supplemental NDAs, including Section 505(b)(2) NDAs, for, among other things, new indications, dosages, or strengths of an existing drug, if new clinical investigations that were conducted or sponsored by the applicant are essential to the approval of the application. Some of our product candidates may qualify for Hatch-Waxman non-patent marketing exclusivity.

In addition to non-patent marketing exclusivity, the Hatch-Waxman Act amended the FDCA to require each NDA sponsor to submit with its application information on any patent that claims the drug for which the applicant submitted the NDA or that claims a method of using such drug and with respect to which a claim of patent infringement could reasonably be asserted if a person not licensed by the owner engaged in the manufacture, use, or sale of the drug. Generic applicants that wish to rely on the approval of a drug listed in the Orange Book must certify to each listed patent, as discussed above. We intend to submit for Orange Book listing all relevant patents for our product candidates.

Finally, the Hatch-Waxman Act amended the patent laws so that certain patents related to products regulated by the FDA are eligible for a patent term extension if patent life was lost during a period when the product was undergoing regulatory review, and if certain criteria are met. We intend to seek patent term extensions, provided our patents and products, if they are approved, meet applicable eligibility requirements.

Pediatric Studies and Exclusivity

The FDA provides an additional six months of non-patent marketing exclusivity and patent protection for any such protections listed in the Orange Book for new or marketed drugs if a sponsor conducts specific pediatric studies at the written request of the FDA. The Pediatric Research Equity Act of 2003, or PREA, authorizes the FDA to require pediatric studies for drugs to ensure the drugs' safety and efficacy in children. PREA requires that certain new NDAs or NDA supplements contain data assessing the safety and effectiveness for the claimed indication in all relevant pediatric subpopulations. Dosing and administration must be supported for each pediatric subpopulation for which the drug is safe and effective. The FDA may also require this data for approved drugs that are used in pediatric patients for the labeled indication, or where there may be therapeutic benefits over existing products. The FDA may grant deferrals for submission of data, or full or partial waivers from PREA. PREA pediatric assessments may qualify for pediatric exclusivity. Unless otherwise required by regulation, PREA does not apply to any drug for an indication with orphan designation.

Orphan Drug Designation and Exclusivity

The FDA may grant orphan drug designation to drugs intended to treat a rare disease or condition, which generally is a disease or condition that affects fewer than 200,000 individuals in the United States. Orphan drug designation must be requested before submitting an NDA. If the FDA grants orphan drug designation, which it may not, the identity of the therapeutic agent and its potential orphan use are publicly disclosed by the FDA. Orphan drug designation does not convey an advantage in, or shorten the duration of, the review and approval process. If a product which has an orphan drug designation subsequently receives the first FDA approval for the indication for which it has such designation, the product is entitled to seven years of orphan drug exclusivity, meaning that the FDA may not approve any other applications to market the same drug for the same indication for a period of seven years, except in limited circumstances, such as a showing of clinical superiority to the product with orphan exclusivity (superior efficacy, safety, or a major contribution to patient care). Orphan drug designation does not prevent competitors from developing or marketing different drugs for that indication. We may seek orphan drug designation for AR-67 for the treatment of GBM and potentially for certain uses of AR-12 and AR-42.

Under European Union medicines laws, the criteria for designating a product as an “orphan medicine” are similar but somewhat different from those in the United States. A drug is designated as an orphan drug if the sponsor can establish that the drug is intended for a life-threatening or chronically debilitating condition affecting no more than five in 10,000 persons in the European Union or that is unlikely to be profitable, and if there is no approved satisfactory treatment or if the drug would be a significant benefit to those persons with the condition. Orphan medicines are entitled to ten years of marketing exclusivity, except under certain limited circumstances comparable to United States law. During this period of marketing exclusivity, no “similar” product, whether or not supported by full safety and efficacy data, will be approved unless a second applicant can establish that its product is safer, more effective or otherwise clinically superior. This period may be reduced to six years if the conditions that originally justified orphan designation change or the sponsor makes excessive profits.

Fast Track Designation

The FDA’s fast track program is intended to facilitate the development and to expedite the review of drugs that are intended for the treatment of a serious or life-threatening condition and that demonstrate the potential to address unmet medical needs. Under the fast track program, applicants may seek traditional approval for a product based on data demonstrating an effect on a clinically meaningful endpoint, or approval based on a well-established surrogate endpoint. The sponsor of a new drug candidate may request the FDA to designate the drug candidate for a specific indication as a fast track drug at the time of original submission of its IND, or at any time thereafter prior to receiving marketing approval of a marketing application. The FDA will determine if the drug candidate qualifies for fast track designation within 60 days of receipt of the sponsor’s request.

If the FDA grants fast track designation, it may initiate review of sections of an NDA before the application is complete. This so-called “rolling review” is available if the applicant provides and the FDA approves a schedule for the submission of the remaining information and the applicant has paid applicable user fees. The FDA’s review clock for both a standard and priority NDA for a fast track product does not begin until the complete application is submitted. Additionally, fast track designation may be withdrawn by the FDA if it believes that the designation is no longer supported by emerging data, or if the designated drug development program is no longer being pursued.

In some cases, a fast track designated drug candidate may also qualify for one or more of the following programs:

- Priority Review. As explained above, a drug candidate may be eligible for a six-month priority review. The FDA assigns priority review status to an application if the drug candidate provides a significant improvement compared to marketed drugs in the treatment, diagnosis or prevention of a disease. A fast track drug would ordinarily meet the

FDA’s criteria for priority review, but may also be assigned a standard review. We do not know whether any of our drug candidates will be assigned priority review status or, if priority review status is assigned, whether that review or approval will be faster than conventional FDA procedures, or that the FDA will ultimately approve the drug.

- Accelerated Approval. Under the FDA's accelerated approval regulations, the FDA is authorized to approve drug candidates that have been studied for their safety and efficacy in treating serious or life-threatening illnesses and that provide meaningful therapeutic benefit to patients over existing treatments based upon either a surrogate endpoint that is reasonably likely to predict clinical benefit or on the basis of an effect on a clinical endpoint other than patient survival or irreversible morbidity. In clinical trials, surrogate endpoints are alternative measurements of the symptoms of a disease or condition that are substituted for measurements of observable clinical symptoms. A drug candidate approved on this basis is subject to rigorous post-marketing compliance requirements, including the completion of Phase IV or post-approval clinical trials to validate the surrogate endpoint or confirm the effect on the clinical endpoint. Failure to conduct required post-approval studies with due diligence, or to validate a surrogate endpoint or confirm a clinical benefit during post-marketing studies, may cause the FDA to seek to withdraw the drug from the market on an expedited basis. All promotional materials for drug candidates approved under accelerated regulations are subject to prior review by the FDA.

When appropriate, we and/or our collaboration partners intend to seek fast track designation, accelerated approval or priority review for our drug candidates. We cannot predict whether any of our drug candidates will obtain fast track, accelerated approval, or priority review designation, or the ultimate impact, if any, of these expedited review mechanisms on the timing or likelihood of the FDA approval of any of our drug candidates.

Satisfaction of the FDA regulations and approval requirements or similar requirements of foreign regulatory agencies typically takes several years, and the actual time required may vary substantially based upon the type, complexity and novelty of the product or disease. Typically, if a drug candidate is intended to treat a chronic disease, as is the case with some of the drug candidates we are developing, safety and efficacy data must be gathered over an extended period of time. Government regulation may delay or prevent marketing of drug candidates for a considerable period of time and impose costly procedures upon our activities. The FDA or any other regulatory agency may not grant approvals for changes in dosage form or new indications for our drug candidates on a timely basis, or at all. Even if a drug candidate receives regulatory approval, the approval may be significantly limited to specific disease states, patient populations and dosages. Further, even after regulatory approval is obtained, later discovery of previously unknown problems with a drug may result in restrictions on the drug or even complete withdrawal of the drug from the market. Delays in obtaining, or failures to obtain, regulatory approvals for any of our drug candidates would harm our business. In addition, we cannot predict what adverse governmental regulations may arise from future United States or foreign governmental action.

Special Protocol Assessment

The FDCA directs the FDA to meet with sponsors, pursuant to a sponsor's written request, for the purpose of reaching agreement on the design and size of clinical trials intended to form the primary basis of an efficacy claim in an NDA. If an agreement is reached, the FDA will reduce the agreement to writing and make it part of the administrative record. This agreement is called a special protocol assessment, or SPA. While the FDA's guidance on SPAs states that documented SPAs should be considered binding on the review division, the FDA has the latitude to change its assessment if certain exceptions apply. Exceptions include identification of a substantial scientific issue essential to safety or efficacy testing that later comes to light, a sponsor's failure to follow the protocol agreed upon, or the FDA's reliance on data, assumptions or information that are determined to be wrong.

Other Regulatory Requirements

Any drugs manufactured or distributed by us or our collaboration partners pursuant to future FDA approvals are subject to continuing regulation by the FDA, including recordkeeping requirements and reporting of adverse experiences associated with the drug. Drug manufacturers and their subcontractors are required to register with the FDA and certain state agencies, and are subject to periodic unannounced inspections by the FDA and certain state agencies for compliance with ongoing regulatory requirements, including cGMP, which impose certain procedural and documentation requirements upon us and our third-party manufacturers. Failure to comply with the statutory and regulatory requirements can subject a manufacturer to possible legal or regulatory action, such as warning letters, suspension of manufacturing, sales or use, seizure of product, injunctive action or possible civil penalties. We cannot be certain that we or our present or future third-party manufacturers or suppliers will be able to comply with the cGMP regulations and other ongoing FDA regulatory requirements. If our present or future third-party manufacturers or suppliers are not able to comply with these requirements, the FDA may halt our clinical trials, require us to recall a drug from distribution, or withdraw approval of the NDA for that drug.

The FDA closely regulates the post-approval marketing and promotion of drugs, including standards and regulations for direct-to-consumer advertising, off-label promotion, industry-sponsored scientific and educational activities and promotional activities involving the Internet. A company can make only those claims relating to safety and efficacy that are approved by the FDA. Failure to comply with these requirements can result in adverse publicity, warning and/or untitled letters, corrective advertising and potential civil and criminal penalties.

Foreign Regulation

In addition to regulations in the United States, we will be subject to a variety of foreign regulations governing clinical trials and commercial sales and distribution of our products. Whether or not we obtain FDA approval for a product, we must obtain approval of a product by the comparable regulatory authorities of foreign countries before we can commence clinical trials or marketing of the product in those countries. The approval process varies from country to country, and the time may be longer or shorter than that required for FDA approval. The requirements governing the conduct of clinical trials, product licensing, pricing and reimbursement vary greatly from country to country.

Under European Union regulatory systems, marketing authorizations may be submitted either under a centralized or mutual recognition procedure. The centralized procedure provides for the grant of a single marketing authorization that is valid for all European Union member states. The mutual recognition procedure provides for mutual recognition of national approval decisions. Under this procedure, the holder of a national marketing authorization may submit an application to the remaining member states. Within 90 days of receiving the applications and assessment report, each member state must decide whether to recognize approval.

In addition to regulations in Europe and the United States, we will be subject to a variety of foreign regulations governing clinical trials and commercial distribution of our future products.

Manufacturing

We do not currently have our own manufacturing facilities. We intend to continue to use our financial resources to accelerate development of our product candidates rather than diverting resources to establish our own manufacturing facilities. We meet our pre-clinical and clinical trial manufacturing requirements by establishing relationships with third-party manufacturers and other service providers to perform these services for us. We rely on individual proposals and purchase orders to meet our needs and typically rely on terms and conditions proposed by the third party or us to govern our rights and obligations under each order (including provisions with respect to intellectual property, if any). We do not have any long-term agreements or commitments for these services. Likewise, we do not have any

long-term agreements or commitments with vendors to supply the underlying component materials of our product candidates, some of which are available from only a single supplier.

Should any of our product candidates obtain marketing approval, we anticipate establishing relationships with third-party manufacturers and other service providers in connection with the commercial production of our products. We have some flexibility in securing other manufacturers to produce our product candidates; however, our alternatives may be limited due to proprietary technologies or methods used in the manufacture of some of our product candidates.

Research and Development Expenses

We spent approximately $5.4 million in fiscal year 2009 and $9.8 million in fiscal year 2008 on research and development activities. These expenses include cash and non-cash expenses relating to the development of our clinical and pre-clinical programs.

Employees

As of the date of this prospectus, we have two full-time employees, none of whom are covered by a collective bargaining agreement. We believe our relations with our employees are satisfactory.

We utilize clinical research organizations and third parties to perform our pre-clinical studies, clinical studies, and manufacturing. We may hire additional research and development staff, as required, to support our product development.

Legal Proceedings

We are not involved in any pending legal proceedings and are not aware of any threatened legal proceedings against us.

Description of Property

Our principal offices are located at 4 Campus Drive, 2nd Floor, Parsippany, New Jersey 07054, where we occupy approximately 5,390 square feet of office space pursuant to the terms of a lease agreement dated October 20, 2008. The lease commencement date was November 14, 2008, with lease payments beginning on January 1, 2009. The lease expiration date is 5 years from the rent commencement date. We provided a security deposit of $44,018, or four months base rent, in the form of a letter of credit. The letter of credit may be reduced by $11,005 on January 1, 2011 and by an additional $11,005 on January 1, 2013, provided we maintain certain conditions described in the lease agreement. We have an early termination option, which provides us the option to terminate the lease on the third anniversary, upon providing the landlord nine months written notice prior to the third anniversary of the lease. If we exercise our termination option, we would be obligated to pay a fee of no more than $53,641 which consists of unamortized costs and expenses incurred by the landlord in connection with the lease. We also have an option to extend the term of the lease for a period of five additional years, provided we give notice to the landlord no later than twelve months prior to the original expiration of the term. We are also responsible for payment of our share of certain charges such as operating costs and taxes in excess of the base year and additional rent. We believe our current facilities in Parsippany, New Jersey will be adequate to meet our needs for the foreseeable future.

MANAGEMENT AND BOARD OF DIRECTORS

Directors and Executive Officers

The following table lists our executive officers, directors and key employees and their respective ages and positions as of the date of this prospectus:

Name	Age	Positions
Arie S. Belldegrun, M.D.	60	Chairman of the Board
David M. Tanen	39	President, Secretary and Director
J. Chris Houchins	47	Chief Operating Officer
Scott L. Navins	39	Treasurer
Stefan Proniuk, Ph.D.	40	Senior Director of Product Development
William F. Hamilton, Ph.D.	70	Director
Tomer Kariv	49	Director
Peter M. Kash	49	Director
Yacov Reizman	59	Director
Steven B. Ruchefsky	48	Director

Arie S. Belldegrun, M.D., FACS has served as the chairman of Arno's board of directors since March 2008. He is currently the Chairman of Two River Group Management, LLC, the managing member of Two River Group Holdings, LLC, and the chairman of the managing member of Two River Consulting, LLC, an organization that provides management, consulting and operational services for development stage biotechnology companies, including Arno. See "Certain Relationships and Related Party Transactions." Dr. Belldegrun is Professor and Chief of Urologic Oncology at the David Geffen School of Medicine at the University of California, Los Angeles, where he holds the Carol and Roy Doumani Chair in Urologic Oncology. He received his medical degree at the Hebrew University Hadassah Medical School, and conducted his post-doctoral studies at the Weizmann Institute of Science in Israel. He completed his Urologic Surgery residency at Harvard Medical School in 1985 and his Surgical Oncology fellowship at the National Cancer Institute/National Institute of Health in 1988. He is certified by the American Board of Urology and is a Fellow of the American College of Surgeons and the American Association of Genitourinary Surgeons. Dr. Belldegrun is also the founder and founding chairman of Agensys Inc., a privately held biotechnology company developing fully human antibody cancer therapeutics based on novel and clinically relevant targets. In December 2007, Agensys was acquired by Astellas Pharma, Inc. in a deal valued at $537 million. Dr. Belldegrun served as Vice-Chairman of the Board of Directors and Chairman of the Scientific Advisory Board of Cougar Biotechnology, an oncology-focused biopharmaceutical company, until its sale to Johnson & Johnson in July 2009. Since October 2009, Dr. Belldegrun has served as a director of Nile Therapeutics, Inc., a publicly-held company focused on the development of biopharmaceutical products for the treatment of cardiovascular diseases. Dr. Belldegrun is on the scientific boards of several biotechnology and pharmaceutical companies and is a reviewer for many medical journals and granting organizations. He served as Chairman of the Molecular and Biological Technology Committee of the American Urological Association and member of its Technology Assessment Council, as a member of the Governor's council on Bioscience for the State of California, and as a biotechnology group leader and member of The Los Angeles Economy and Jobs Committee established in October 2006 by Mayor Antonio Villaraigosa. He is the author of several books on prostate and kidney cancers, holds several biopharmaceutical patents, and has written over 400 scientific publications with an emphasis on urologic oncology.

David M. Tanen co-founded Arno and has been a director and its secretary since its inception and has served as Arno's President since June 2009. In September 2004, Mr. Tanen co-founded Two River Group Holdings, LLC, a venture capital firm that specializes in the creation of new companies that acquire rights to commercially develop biotechnology products, and currently serves as Vice President and Director of Two River's managing member, Two

River Group Management, LLC. Mr. Tanen also serves as an Officer and Director of Riverbank Capital Securities, Inc., a broker dealer registered with FINRA ("Riverbank"), which provided placement agent services for Arno. Mr. Tanen also serves as an officer of the managing member of Two River Consulting, LLC, which provides management, operational and other services for Arno. See "Certain Relationships and Related Party Transactions." Prior to founding Two River, from October 1996 to September 2004, Mr. Tanen was a Director of Paramount BioCapital Investments, LLC, a biotechnology focused venture capital company. Mr. Tanen also served as a member of the General Partner of the Orion Biomedical Fund, LP. Mr. Tanen received his B.A. from The George Washington University and his J.D. from Fordham University School of Law.

J. Chris Houchins has been employed by Arno since September 2007 and has over 16 years of clinical operations and drug development experience focusing in oncology. From 2004 to 2006, Mr. Houchins was the Director of Specialty Care - Clinical Project Management at Schering-Plough, where he was involved with the FDA and European submissions and approvals of Temozolomide, the standard of care for patients with GBM. From 1999 to 2004, Mr. Houchins was on the Searle Celebrex Oncology Team that received FDA approval for familial adenomatous polyposis ("FAP"), a new indication in oncology. After the merger of Searle and Pharmacia & Upjohn, he oversaw the development and clinical operations for the Global Celebrex Oncology Program that grew to over 300 clinical and preclinical trials world-wide within 2 years. When Pfizer, Inc acquired Pharmacia Corp., Mr. Houchins was selected as Director – Team Leader of Oncology Clinical Operations overseeing all eight oncology compounds (Camptosar®, Aromasin®, Ellence®, Celebrex®, Emcyt®, Zavedos®, Trelstar® and Zinecard®) encompassing over 500 clinical and preclinical studies. In addition, he was appointed to the Pfizer Global Oncology Advisory Board. Mr. Houchins also has six years of experience as a Clinical Research Manager at The RUSH Cancer Institute in Chicago where he managed clinical trials across all tumor types including Ovarian Cancer using Topotecan. He is certified by examination through SoCRA and ACRP as a Certified Clinical Research Associate, Coordinator and Professional and holds a BS in Economics from Northern Illinois University.

Stefan Proniuk, Ph.D., M.B.A. has over 12 years of experience in formulation and product development. Prior to joining Arno, he was the Sr. Manager of Pharmaceutical Technologies at Neurocrine Biosciences (2002-2008) where he was responsible for overseeing development programs from Phase I to III. His group was also responsible for the preformulation of NCEs. Prior to his work at Neurocrine, Dr. Proniuk worked as a scientist at Cima Labs (2001-2002) on the development and scale-up of fast dissolving tablet formulations (OraSolv®, DuraSolv®). Throughout his career he has worked on 2 NDAs, 8 INDs, 1 IMPD, 1 CTA, and 3 marketed products. Dr. Proniuk holds a Ph.D. degree in Pharmaceutical Sciences from the University of Arizona, a MBA with emphasis in Entrepreneurship from San Diego State University and a Diplom (FH) in Chemical Engineering from the Polytechnical University Isny in Germany. He is also certified in Intellectual Property Law from the University of California San Diego.

William F. Hamilton, Ph.D. was appointed to Arno's board of directors in October 2008. Dr. Hamilton has served on the University of Pennsylvania faculty since 1967, and is the Landau Professor of Management and Technology, and Director of the Jerome Fisher Program in Management and Technology at The Wharton School and the School of Engineering and Applied Science. He serves as a director of Neose Technologies, Inc. and NovaDel Pharma Inc., both publicly-traded biotechnology companies. Dr. Hamilton also serves on the boards of directors of Yaupon Therapeutics, Inc., a privately-held specialty pharmaceutical company that develops small molecule pharmaceuticals licensed from academic laboratories, Avid Radiopharmaceuticals, Inc., a privately-held clinical-stage product-focused molecular imaging company and Neuro Diagnostic Devices Inc., a privately-held development-stage medical device company. Dr. Hamilton received his B.S. and M.S. in chemical engineering and his MBA from the University of Pennsylvania, and his Ph.D. in applied economics from the London School of Economics.

Tomer Kariv is the Chief Executive Officer of Pontifax Fund, an Israeli based fund that specializes in investing in development stage companies in the pharmaceutical and life sciences industries, and a significant stockholder in Arno. For the past 10 years, Mr. Kariv has played a key role in investing, managing and nurturing technology driven companies and startups. Prior to joining Pontifax, Mr. Kariv served in many senior management positions at top Israeli financial institutions, including STI Ventures, Shrem Fudim Kelner Investment House, and Polaris I, one of Israel's top venture capital performers. Mr. Kariv was also a co-founder of Polaris II (currently Pitango), Israel's largest venture capital fund. Mr. Kariv practiced law with Sullivan & Cromwell, a leading corporate law firm in New York City, and holds a B.A. in Economics from Harvard University and a J.D. from Harvard Law School.

Peter M. Kash co-founded Arno and has been a director since its inception. In September 2004, Mr. Kash co-founded Two River. Mr. Kash is also the President and Chairman of Riverbank. From 1992 until 2004, Mr. Kash was a Senior Managing Director of Paramount BioCapital, Inc., a FINRA member broker dealer, and Paramount BioCapital Investments, LLC, a biotechnology focused venture capital company. Mr. Kash also served as Director of Paramount Capital Asset Management, Inc., the general partner of several biotechnology-related hedge funds (the Paramount companies are collectively referred to as Paramount), and as member of the General Partner of the Orion Biomedical Fund, LP, a private equity fund. Mr. Kash currently serves as a member of Board of Directors of Nile Therapeutics, Inc. (NASDAQ:NLTX), as well as several privately held biotechnology companies. Mr. Kash received his B.S. in Management Science from SUNY Binghamton and his M.B.A. in Banking and International Finance from Pace University. Mr. Kash is currently seeking his doctorate in Jewish education at Yeshiva University.

Yacov Reizman has been the Chairman and Chief Executive Officer of FCC Ltd., a private investment company that he founded in 1987. Over the past decade FCC has invested directly in over 50 publicly traded and privately held companies in a diverse range of industries including: infrastructure; shipping; healthcare; and financial services. FCC also specializes in corporate finance and structured investments. Mr. Reizman was also co-founder and co-CEO of Azimuth Ltd., which traded on the TASE. Previously Mr. Reizman served in the Israeli Air Force (IAF) as a fighter pilot (Major) and led large-scale high-tech projects for the IAF, including joint projects with Israeli and U.S. defense industries. Mr. Reizman holds a B.A. in economics and in psychology from Tel Aviv University.

Steven Ruchefsky is President of Commercial Street Capital LLC, a private investment company and significant stockholder in Arno. For the last decade, Mr. Ruchefsky has been working as an investment manager for the founder and CEO of a multi-billion dollar hedge fund. Mr. Ruchefsky began his career at a prominent New York City law firm where he became a partner, member of management and chair of a specialized litigation group. Upon leaving his law firm and prior to his current employment, Mr. Ruchefsky was a principal of an early stage venture capital operation. Mr. Ruchefsky currently sits on the boards of several public and private companies, including National Investment Managers Inc. (NIVM:OB); Evogene (TASE:EVGN); and Itamar Medical (TASE: ITMR). Mr. Ruchefsky is a graduate of The George Washington University Law School

Scott L. Navins has served as Arno’s Treasurer since its inception, and has been responsible for all of our accounting and financial reporting services since the departure of our former Chief Financial Officer in February 2010. He is also the Vice President of Finance at Two River Group Holdings, LLC and Two River Consulting, LLC, where he is responsible for all accounting, finance and control activities. Mr. Navins joined Two River in 2005. Prior to joining Two River, from 2004 to 2005 Mr. Navins was the Senior Controller at Westbrook Partners, where he managed the accounting for a $560 million real estate private equity fund, including financial and partner reporting, tax coordination, maintaining internal controls and overseeing a $300 million credit facility, among other things. Before that, from 2002 to 2004 Mr. Navins was a Senior Manager at Morgan Stanley, where he managed the accounting for a $2.4 billion real estate private equity fund. Prior to that Mr. Navins was an Associate in the Finance Group at BlackRock, Inc. and the controller for a high-tech venture capital fund. Mr. Navins also serves as the Financial and Operations Principal of Riverbank Capital Securities (member FINRA/SIPC) and has served as Treasurer of Nile Therapeutics, Inc., a publicly-held biopharmaceutical company, since 2005. Mr. Navins graduated with honors from The George Washington University in 1993, where he earned a Bachelor of Accountancy degree. Mr. Navins passed the Uniform Certified Public Accounting examination in 1993.

Experience, Qualifications, Attributes and Skills of Directors

We look to our directors to lead us through our continued growth as an early-stage public biopharmaceutical company. We believe our directors bring their leadership experience from a variety of life science companies and professional backgrounds which we require to continue to grow and bring value to our stockholders. Messrs. Kariv, Kash, Reizman, Ruchefsky and Tanen have venture capital or investment banking backgrounds and offer expertise in financing and growing small companies, particularly small biopharma and life science companies. Each of Drs. Belldegrun and Hamilton and Messrs. Kash and Tanen have significant experience with early stage private and public companies and bring depth of knowledge in building stockholder value, growing a company from inception and navigating significant corporate transactions and the public company process. Dr. Belldegrun's medical background and experience serving as an investigator in clinical trials of oncology drug candidates allows him to contribute significant medical and scientific expertise. Mr. Tanen's current position as our President also allows him to provide a unique insight into our development and growth. As a result of his academic experience and his prior service on the audit committees of several publicly-traded life sciences companies, Dr. Hamilton also bring extensive finance, accounting and risk management knowledge to us.

Independence of the Board of Directors

In determining whether the members of our board of directors and its committees are independent, we have elected to use the definition of "independence" set forth in the listing standards of the NASDAQ Stock Market. After considering all relevant relationships and transactions, our board of directors, in consultation with legal counsel, has determined that Messrs. Kariv, Reizman, Ruchefsky and Dr. Hamilton are "independent" within the meaning of the applicable listing standard of the NASDAQ Stock Market. Messrs. Kash, Tanen and Dr. Belldegrun are not independent, as defined by applicable NASDAQ listing standards.

Board Committees

The Board of Directors has established three standing committees: an Audit Committee, a Compensation Committee and a Nominating & Corporate Governance Committee. The following table provides membership for each of the Board committees:

Committee	Membership
Audit	Dr. Hamilton (Chair) and Mr. Ruchefsky
Compensation	Dr. Belldegrun, Mr. Kariv, Mr. Reizman and Mr. Ruchefsky
Nominating & Governance	Dr. Hamilton, Mr. Reizman and Mr. Ruchefsky

Executive Compensation

The following table sets forth all of the compensation awarded to, earned by or paid to (i) each individual serving as our principal executive officer during the fiscal year ended December 31, 2009; (ii) two other individuals that served as executive officers at the conclusion of the fiscal year ended December 31, 2009 and who received in excess of $100,000 in total compensation during such fiscal year; and one additional individual who received in excess of $100,000 in total compensation during the fiscal year ended December 31, 2009 but who did not serve as an executive officer at the conclusion of such fiscal year. We refer to these individuals as our named executives.

Summary Compensation Table

Name and Principal Position	Year	Salary	Bonus	Option Awards (1)	All Other Compensation(2)	Total
David M. Tanen (3)	2009	$ –	$ –	$ –	$ –	$ –
President						
J. Chris Houchins	2009	$ 207,573	$ 50,000	$ 182,300	$ -	$ 439,873
Senior VP of Clinical Operations						
Roger G. Berlin, M.D. (4)	2009	$ 156,250	$ 76,923	$ -	$ 62,500	$ 295,673
Former Chief Executive Officer	2008	122,356	-	2,025,400	4,330	2,152,086
Scott Z. Fields, M.D. (5)	2009	$ 142,974	$ 100,000	$ 21,900	$ -	$ 264,874
Former President, Chief Medical Officer	2008	340,000	125,000	-	4,244	469,244
Brian Lenz (6)	2009	$ 200,000	$ 28,000	$ 0	$ -	$ 228,000
Former Chief Financial Officer	2008	91,667	25,000	830,500	645	947,812

(1) Amounts reflect the grant date fair value of awards granted under the Company's 2005 Stock Option Plan, computed pursuant to Financial Accounting Standards Board's Accounting Standards Codification 718 "Compensation – Stock Compensation". Assumptions used in the calculation of these amounts are included in Note 10 of the Notes to Audited Financial Statements included in this prospectus.

(2) Except as otherwise noted, amounts reflect premiums paid for life insurance.

(3) Mr. Tanen was appointed President on June 8, 2009. Mr. Tanen, who also serves as a director, does not receive compensation for his service as President, but does receive compensation for his service as a director in accordance with the terms of our non-employee director compensation plan. See "—Director Compensation."

(4) Dr. Berlin's employment terminated on May 31, 2009. In connection with such termination, we continued to pay to Dr. Berlin his annualized base salary through July 31, 2009, which amount totaled $62,500.

(5) Dr. Fields' employment terminated on June 1, 2009.

(6) Mr. Lenz's employment terminated on February 15, 2010.

Employment Agreements, Termination of Employment and Change-in-Control Arrangements

J. Chris Houchins
Chief Operating Officer

Mr. Houchins' employment with us is governed by a letter agreement dated September 12, 2007, as amended on August 26, 2010. The letter agreement provides for Mr. Houchins' employment as our Senior Vice President of Clinical Operations on an at-will basis. Under the letter agreement, Mr. Houchins is entitled to an annual base salary of $180,000, which base salary was subsequently increased to $182,000 on January 1, 2008, to $200,000 on August 1, 2008 and to $207,500 on January 1, 2009. In addition, Mr. Houchins is eligible to receive an annual performance bonus of up to 25% of his base salary upon the successful completion of annual corporate and individual milestones. The letter agreement also provides for the awarding of certain stock options to Mr. Houchins, referred to as Employment Options. On September 17, 2007, Mr. Houchins was granted ten-year Employment Options to purchase 99,689 shares of our common stock at an exercise price of $1.00, with one-fourth vesting after one year and the remainder vesting in 36 equal monthly installments thereafter. On September 29, 2009, Mr. Houchins was granted ten-year Employment Options to purchase 200,000 shares of our common stock at an exercise price of $1.00, which vest in three equal annual installments commencing on the first anniversary of the grant date. Mr. Houchins is also entitled to participate in Arno's employee benefits plans, and to receive other customary benefits. In November 2010, Mr. Houchins was appointed as our Chief Operating Officer.

Roger G. Berlin, M.D.
Former Chief Executive Officer

Dr. Berlin's employment with us was governed by an employment agreement dated August 19, 2008. The employment agreement provided for Dr. Berlin's employment as Chief Executive Officer for a two-year term commencing on September 3, 2008, and for Dr. Berlin's appointment to our board of directors. The agreement provided for an initial annual base salary of $375,000. In addition, Dr. Berlin was eligible to receive an annual performance bonus of up to 50% of his base salary upon the successful completion of annual corporate and individual milestones. The agreement also provided for the awarding of certain stock options to Dr. Berlin, referred to as Employment Options, Performance Options, and Technology Options. Dr. Berlin was also entitled to participate in Arno's employee benefits plans, and to receive other customary benefits.

Dr. Berlin's employment with us terminated on May 31, 2009, pursuant to the terms of a separation agreement. Pursuant to the terms of the separation agreement, we continued to pay Dr. Berlin's annualized base salary and provide health and dental insurance coverage through July 31, 2009. The separation agreement also provides for the extension of the exercise period of Performance Options relating to 71,667 shares of our common stock at an exercise price of $3.00, which vested on December 31, 2008, until May 31, 2014, at which time all unexercised Performance Options shall automatically expire. As of the separation date, Dr. Berlin also held unvested Employment Options relating to 430,000 shares of our common stock, and unvested Performance Options relating to 358,333 shares of our common stock, in both cases at an exercise price of $3.00 per share, all of which automatically terminated as of the separation date. Dr. Berlin was not awarded any Technology Options during his employment with us.

Scott Z. Fields, M.D.
Former President and Chief Medical Officer

Dr. Fields' employment with us was governed by an employment agreement dated June 1, 2007. The employment agreement provided for Dr. Fields' employment as President and Chief Medical Officer for a two-year term commencing on June 1, 2007. The agreement provided for an initial annual base salary of $340,000. In addition, Dr. Fields was eligible to receive an annual performance-based bonus of up to $150,000 upon the successful completion of annual corporate and individual milestones at an exemplary metric (e.g., ahead of schedule, under budget, etc.). The agreement also provided for the awarding of certain stock options to Dr. Fields, referred to as Employment Options, Performance Options, and Technology Options. Dr. Fields was also entitled to participate in Arno's employee benefits plans, and to receive other customary benefits.

In February 2009, Dr. Fields informed us that he would not be continuing his employment with us when the term of his employment agreement expired on June 1, 2007. As of his separation date, Dr. Fields held vested Employment Options to purchase 199,377 shares of our common stock and vested Performance Options to purchase 199,377 shares of our common stock, in both cases at an exercise price of $1.00 per share. Pursuant to his employment agreement, Dr. Fields' vested stock options remained exercisable until their termination 12 months after his separation date and his unvested stock options automatically terminated as of his separation date. On September 29, 2009, Dr. Fields and Arno entered into a Scientific Advisory and Consulting Agreement, or the SAB Agreement. As partial consideration for entering into the SAB Agreement Arno agreed to extend the expiration date of Dr. Field's Employment Options and Performance Options until December 31, 2014.

Brian Lenz
Former Chief Financial Officer

Mr. Lenz's employment with us was governed by an employment agreement dated June 11, 2008, as amended on July 9, 2008. Under the agreement, which provided for a two-year term commencing on July 15, 2008, Mr. Lenz was appointed as our Chief Financial Officer effective August 15, 2008. The agreement provided for an initial annual base salary of $200,000. Mr. Lenz was also eligible to receive an annual performance bonus of up to 30% of his base salary upon the successful completion of annual corporate and individual milestones. In addition, upon the commencement of his employment, Mr. Lenz received a one-time cash bonus in the amount of $25,000 and a ten-year option to purchase 440,000 shares of our common stock at an exercise price of $2.75 per share, with one-fourth vesting after one year and the remainder vesting in 24 equal monthly installments thereafter, subject to Mr. Lenz's continued employment with Arno. Mr. Lenz was also entitled to participate in Arno's employee benefits plans, and to receive other customary benefits. Mr. Lenz voluntarily resigned his employment with us effective February 15, 2010, and all of his options subsequently expired without being exercised.

Outstanding Equity Awards at Fiscal Year-End

The following table sets forth information concerning stock options held by the named executive officers at December 31, 2009:

Name	Number of Securities Underlying Unexercised Options Exercisable	Number of Securities Underlying Unexercised Options Unexercisable	Option Exercise Price ($)	Option Expiration Date
Mr. Tanen	3,333	6,667	1.00	9/29/19 (1)
Mr. Houchins	56,075	43,614	2.00	9/17/17 (2)
	0	200,000	1.00	9/29/19 (3)
Dr. Berlin	71,667	0	3.00	5/31/14
Dr. Fields	398,754	0	2.00	12/31/14
	0	25,000	1.00	9/22/14 (4)
Mr. Lenz	178,750	261,250	2.75	7/16/18 (5)

(1) Option granted on September 29, 2009, as compensation for Mr. Tanen's services as a director prior to his appointment as our President. The option vests in three equal installments on each anniversary of the date of grant.
(2) Option granted September 17, 2017 relating to an aggregate of 99,688 shares, of which 25% vested on the first anniversary of the grant date and the remainder vests in 36 equal monthly installments thereafter.
(3) Option granted September 29, 2009 and vests in three equal annual installments commencing on the first anniversary of the grant date.
(4) Option granted on September 22, 2009, as compensation for Dr. Field's services as a member of the Company's scientific advisory board. The option vests in 3 equal installments on each anniversary of the grant date.
(5) The option expired in its entirety following Mr. Lenz's resignation in February 2010.

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Director Compensation

Pursuant to the non-employee director compensation plan adopted by our Board of Directors, our non-employee directors are entitled to receive the following in consideration for their service on the Board: (1) an annual retainer of $25,000; (2) a stock option grant relating to 30,000 shares of the Company's common stock upon their initial appointment or election to the Board; and (3) an annual stock option grant relating to 10,000 shares of the Company's common stock. In addition, any non-employee director designated as chairman of the Board is entitled to an annual retainer of $10,000, the chair of the Board's audit committee is entitled to an additional annual retainer of $8,000, and the chairs of the Board's compensation and nominating & corporate governance committees are entitled to annual retainers of $4,000. Stock options awarded to our non-employee directors have a 10-year term, vest in three equal annual installments commencing on the first anniversary of the grant date, and have an exercise price equal to the fair market value of the Company's common stock on the grant date.

The following table sets forth the compensation paid to our directors for their service in 2009.

Name (1)	Fees earned or paid in cash	Option Awards (2)	Total
Arie S. Belldegrun, M.D.	$ 39,000	$ 9,100	$ 48,100
William F. Hamilton, Ph.D.	$ 33,000	$ 27,300	$ 60,300
Robert I. Falk (3)	$ 29,000	$ 9,100	$ 38,100
Peter M. Kash	$ 25,000	$ 9,100	$ 34,100
Joshua A. Kazam (3)	$ 25,000	$ 9,100	$ 34,100
David M. Tanen	$ 25,000	$ 9,100	$ 34,100

(1) Roger G. Berlin, our former Chief Executive Officer, has been omitted from this table since he received no additional compensation for serving on our Board; his compensation is described above under "Executive Compensation."

(2) Amounts reflect the grant date fair value of awards granted under the Company's 2005 Stock Option Plan, computed pursuant to Financial Accounting Standards Board's Accounting Standards Codification 718 "Compensation – Stock Compensation". Assumptions used in the calculation of these amounts are included in Note 10 of the audited December 31, 2009 Notes to Financial Statements included in this prospectus.

(3) Messrs. Falk and Kazam resigned as directors in September 2010 upon the completion of our private placement of Series A Preferred Stock.

SECURITY OWNERSHIP OF CERTAIN BENEFICIAL OWNERS AND MANAGEMENT

The following table summarizes certain information regarding the beneficial ownership (as such term is defined in Rule 13d-3 under the Exchange Act) of our common stock and Series A Preferred Stock as of October 31, 2010 by: (i) each of our current directors, (ii) each of our "named executive officers," as defined above under "Executive Compensation," (iii) all of our current directors and executive officers as a group, and (iv) each person known by us to be the beneficial owner of more than 5% of our common stock or Series A Preferred Stock, Except as indicated in the footnotes below, the security and stockholders listed below possess sole voting and investment power with respect to their shares. Except as otherwise indicated, the address of each of our executive officers and directors identified below is 4 Campus Drive, 2nd Floor, Parsippany, New Jersey 07054.

Name of Beneficial Owner	No. Shares of Common Stock Beneficially Owned (1)	Percent of Class (1)	No. Shares of Series A Preferred Stock Beneficially Owned (2)	Percent of Class (2)
Arie S. Belldegrun, M.D. (3)	1,346,175	3.7	750,000	4.9
David M. Tanen (4)	1,632,720	4.6	164,084	1.1
J. Chris Houchins (5)	303,608	*	—	—
Scott L. Navins (6)	249,532	*	100,000	*
Roger G. Berlin, M.D. (5)	71,667	*	—	—
Scott Z. Fields, M.D. (5)	406,987	1.1	—	—
Brian Lenz	4,000	*	—	—
William F. Hamilton, Ph.D. (7)	39,938	*	—	—
Tomer Kariv (8)	4,500,000	12.1	4,500,000	26.8
Peter M. Kash (9)	2,038,511	5.7	339,534	2.2
Yacov Reizman (10)	756,300	2.1	756,300	4.8
Steven B. Ruchefsky (11)	3,300,000	9.0	3,300,000	20.2
All current executive officers and directors as a group (8 persons)	13,863,176	34.7	9,909,918	52.3
Pontifax Ltd. (8)	4,500,000	12.1	4,500,000	26.8
UTA Capital LLC (12) 100 Executive Drive, Ste. 330 Wes Orange, NJ 07052	3,000,000	8.2	3,000,000	18.4
Commercial Street Capital, LLC (11) 800 Westchester Ave. Rye Brook, NY 10573	3,300,000	9.0	3,300,000	20.2
Clal Insurance Co. Ltd. (13) 48 Menachem Begin St. Tel-Aviv 66180, Israel	3,104,727	8.6	1,650,000	10.4
Wexford Capital LP (14) 411 West Putnam Ave. Greenwich, CT 06830	3,865,789	10.6	1,860,000	11.7

* represents less than 1%.

(1) Assumes 35,686,024 shares of our common stock are outstanding, including 15,274,000 shares issuable upon the conversion of our Series A Preferred Stock, but does not include any shares issuable upon the exercise of outstanding warrants and options to purchase shares of our capital stock. Beneficial ownership is determined in accordance with Rule 13d-3 under the Securities Act, and includes any shares as to which the security or stockholder has sole or shared voting power or investment power, and also any shares which the security or stockholder has the right to acquire within 60 days of the date hereof, whether through the exercise or conversion of any stock option, convertible security, warrant or other right. The indication herein that shares are beneficially owned is not an admission on the part of the security or stockholder that he, she or it is a direct or indirect beneficial owner of those shares.

(2) Based upon 15,274,000 shares of Series A Preferred Stock outstanding, excluding warrants to purchase 8,693,930 shares of Series A Preferred Stock.

(3) Common stock beneficially owned include (i) 61,916 shares held in a trust of which Dr. Belldegrun is a beneficiary, and (ii) 509,337 shares issuable upon exercise of stock options. Series A Preferred Stock beneficially owned includes (i) 187,500 shares held in a family trust for which Dr. Belldegrun is a co-trustee, including 62,500 shares issuable upon the exercise of warrants, (ii) 375,000 shares held in a trust of which Dr. Belldegrun is a beneficiary, including 125,000 shares issuable upon the exercise of warrants, and (iii) 187,500 shares held in a family limited partnership of which Dr. Belldegrun is a partner.

(4) Common stock beneficially owned includes 149,532 shares held by Mr. Tanen's minor children and 11,236 shares issuable upon the exercise of options and warrants held by Mr. Tanen. Series A Preferred Stock beneficially owned includes 99,618 shares issuable upon the exercise of warrants.

(5) Represents shares issuable upon the exercise of stock options.

(6) Series A Preferred Stock beneficially owned includes 100,000 shares issuable upon the exercise of warrants.

(7) Includes 30,000 shares issuable upon the exercise of stock options.

(8) Represents shares issuable upon conversion of Series A Preferred Stock, including warrants to purchase 1,500,000 shares of Series A Preferred Stock, all of which are held by affiliates of Pontifax Ltd., of which Mr. Kariv is chief executive officer.

(9) Common stock beneficially owned includes (i) 358,876 shares held by Mr. Kash's minor children and (ii) 12,472 shares issuable upon the exercise of stock options and warrants held by Mr. Kash. Series A Preferred Stock beneficially owned includes 239,534 shares issuable upon the exercise of warrants.

(10) Represents shares issuable upon conversion of Series A Preferred Stock, including warrants to purchase 456,300 shares of Series A Preferred Stock, all of which are held by FCC Ltd., of which Mr. Reizman is chairman and chief executive officer.

(11) Represents shares issuable upon conversion of Series A Preferred Stock, including warrants to purchase 1,100,000 shares of Series A Preferred Stock, held by Commercial Street Capital, LLC, of which Mr. Ruchefsky is president.

(12) Represents shares issuable upon conversion of Series A Preferred Stock, including warrants to purchase 1,000,000 shares of Series A Preferred Stock.

(13) Common stock beneficially owned includes 1,650,000 shares of Series A Preferred Stock, including 550,000 shares of Series A Preferred Stock issuable upon the exercise of warrants.

(14) Common stock beneficially owned represents: (i) 247,345 shares of our common stock held by Kappa Investors, LLC ("Kappa"); (ii) a warrant held by Kappa to purchase 24,734 shares of our common stock that are exercisable at $2.42 per share; and (iii) 1,733,712 shares of our common stock held by Wexford Spectrum Investors LLC, a Delaware limited liability company ("Wexford Spectrum"). Series A Preferred Stock beneficially owned represents (i) 1,607,985 shares held by Wexford Spectrum, including 535,995 shares issuable upon the exercise of warrants to purchase Series A Preferred Stock, and (ii) 252,014 shares held by Kappa, including 84,005 shares issuable upon the exercise of warrants. Wexford Capital LP, a Delaware partnership ("Wexford Capital"), is a

registered Investment Advisor and also serves as an investment advisor or sub-advisor to the members of Kappa and Wexford Spectrum. Wexford GP LLC ("Wexford GP") is the general partner of Wexford Capital. Mr. Charles E. Davidson and Mr. Joseph M. Jacobs are each managing and controlling members of Wexford GP.

TRANSACTIONS WITH RELATED PERSONS, PROMOTERS AND CERTAIN CONTROL PERSONS

Dr. Belldegrun and Mr. Tanen, each a current director and substantial stockholder of Arno, and Mr. Kazam, a director until September 2010 and substantial stockholder of Arno, control Two River Consulting, LLC, or TRC. Certain employees of TRC, including Mr. Tanen, our President, and Mr. Scott L. Navins, our Treasurer, perform substantial services for us, including without limitation operational, managerial, financial, clinical and regulatory activities for which we pay TRC a monthly consulting fee of $55,000 pursuant to a services agreement. Other than the payments to TRC, we do not pay any salary or other compensation to Messrs. Tanen and Navins for their services to us.

Mr. Kazam, Mr. Tanen and Mr. Peter M. Kash, who also serves as a director of Arno, are also principals of Riverbank Capital Securities, Inc., a FINRA member broker dealer that acted as our placement agent in connection with our September 2010 and June 2008 private placements. Additionally, Mr. Navins, our Treasurer, is also the Financial and Operations Principal of Riverbank. In consideration for its services in connection with the September 2010 private placement, we paid Riverbank a placement fee of approximately $789,880 and issued to designees of Riverbank five-year warrants to purchase an aggregate of 664,880 shares of Series A Preferred Stock at an initial exercise price of $1.10 per share. Riverbank did not receive any selling commission for its services in connection with the June 2008 private placement, but received a non-accountable expense allowance of $100,000.

Pursuant to a Consulting Agreement entered into between Arno and Fountainhead Capital Management Limited, we paid a $500,000 consulting fee to Fountainhead Capital upon completion of the merger with Laurier. Fountainhead Capital Management was a significant stockholder of Laurier at the time of the merger.

WHERE YOU CAN FIND MORE INFORMATION

Federal securities laws require us to file information with the SEC concerning our business and operations. Accordingly, we file annual, quarterly, and special reports, proxy statements and other information with the SEC. You can inspect and copy this information at the Public Reference Facility maintained by the SEC at Judiciary Plaza, 100 F Street, N.E., Washington, D.C. 20549. You can receive additional information about the operation of the SEC's Public Reference Facilities by calling the SEC at 1-800-SEC-0330. The SEC also maintains a web site at http://www.sec.gov that contains reports, proxy and information statements and other information regarding companies that, like us, file information electronically with the SEC.

VALIDITY OF COMMON STOCK

Legal matters in connection with the validity of the shares offered by this prospectus will be passed upon by Fredrikson & Byron, P.A., Minneapolis, Minnesota.

EXPERTS

The financial statements as of December 31, 2009, and for the year then ended, and for the period from August 1, 2005 (inception) through December 31, 2009, included in this prospectus, have been so included in reliance on the report of Crowe Horwath LLP, independent registered public accounting firm, given on the authority of that firm as experts in accounting and auditing.

The financial statements as of December 31, 2008, and for the year then ended, included in this prospectus, have been so included in reliance on the report of Hays & Company LLP, independent registered public accounting firm, given on authority of said firm as experts in auditing and accounting.

TRANSFER AGENT

The transfer agent for our common stock is American Stock Transfer & Trust Company, and its address is 40 Wall Street, New York, New York, 10005.

DISCLOSURE OF COMMISSION POSITION ON
INDEMNIFICATION FOR SECURITIES ACT LIABILITIES

Insofar as indemnification for liabilities arising under the Securities Act of may be permitted to directors, officers or persons controlling the registrant pursuant to the foregoing provisions, the registrant has been informed that in the opinion of the Securities and Exchange Commission such indemnification is against public policy as expressed in the Securities Act and is therefore unenforceable.

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)

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ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)

Report of Independent Registered Public Accounting Firm

To the Board of Directors and stockholders
Arno Therapeutics, Inc.
Parsippany, New Jersey

We have audited the accompanying balance sheet of Arno Therapeutics, Inc. (a development stage company) as of December 31, 2009, and the related statements of operations, stockholders' equity, and cash flows for the year then ended and for the period from August 1, 2005 (inception) through December 31, 2009. These financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on these financial statements based on our audit. The financial statements of Arno Therapeutics, Inc. for the period from August 1, 2005 (inception) through December 31, 2008 were audited by other auditors whose report dated March 31, 2009 expressed an unqualified opinion. Our opinion on the statements of operations, stockholders' equity, and cash flows for the period from August 1, 2005 (inception) through December 31, 2009, insofar as it relates to the amounts for prior periods through December 31, 2008, is based solely on the report of other auditors.

We conducted our audit in accordance with the standards of the Public Company Accounting Oversight Board (United States). Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement. The Company is not required to have, nor were we engaged to perform, an audit of its internal control over financial reporting. Our audit included consideration of internal control over financial reporting as a basis for designing audit procedures that are appropriate in the circumstances, but not for the purpose of expressing an opinion on the effectiveness of the Company's internal control over financial reporting. Accordingly, we express no such opinion. An audit includes examining, on a test basis, evidence supporting the amounts and disclosures in the financial statements. An audit also includes assessing the accounting principles used and significant estimates made by management, as well as evaluating the overall financial statement presentation. We believe that our audit provide a reasonable basis for our opinion.

In our opinion, the financial statements referred to above present fairly, in all material respects, the financial position of Arno Therapeutics, Inc. (a development stage company) as of December 31, 2009, and the results of its operations and its cash flows for the year then ended and the period from August 1, 2005 (inception) through December 31, 2009, in conformity with U.S. generally accepted accounting principles.

/s/ Crowe Horwath LLP

New York, New York
November 8, 2010

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)

REPORT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

To the Board of Directors and stockholders
Arno Therapeutics, Inc.

We have audited the accompanying balance sheet of Arno Therapeutics, Inc. (a development stage company) as of December 31, 2008 and the related statements of operations, stockholders' equity and cash flows for the year then ended. These financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on these financial statements based on our audit.

We conducted our audit in accordance with the standards of the Public Company Accounting Oversight Board (United States). Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement. The Company is not required to have, nor were we engaged to perform, an audit of its internal control over financial reporting. Our audit included consideration of internal control over financial reporting as a basis for designing audit procedures that are appropriate in the circumstances, but not for the purpose of expressing an opinion on the effectiveness of the Company's internal control over financial reporting. Accordingly, we express no such opinion. An audit also includes examining, on a test basis, evidence supporting the amounts and disclosures in the financial statements, assessing the accounting principles used and significant estimates made by management, as well as evaluating the overall financial statement presentation. We believe that our audit provides a reasonable basis for our opinion.

In our opinion, the financial statements referred to above present fairly, in all material respects, the financial position of Arno Therapeutics, Inc. as of December 31, 2008, and the results of its operations and its cash flows for the year then ended, in conformity with accounting principles generally accepted in the United States of America.

/s/ Hays & Company LLP

March 31, 2009
New York, New York

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
BALANCE SHEETS

	December 31, 2009	December 31, 2008
ASSETS		
Current assets		
Cash and cash equivalents	$ 3,087,299	$ 10,395,007
Prepaid expenses	110,589	315,014
Total current assets	3,197,888	10,710,021
Property and equipment, net	41,567	63,584
Restricted cash	44,018	44,018
Security deposit	-	12,165
Total assets	$ 3,283,473	$ 10,829,788
LIABILITIES AND STOCKHOLDERS' EQUITY		
Current liabilities		
Accounts payable	$ 1,003,030	$ 2,493,658
Accrued expenses	556,204	450,713
Due to related party	132,418	5,616
Total current liabilities	1,691,652	2,949,987
Deferred rent	16,070	17,393
Total liabilities	1,707,722	2,967,380
COMMITMENTS AND CONTINGENCIES		
STOCKHOLDERS' EQUITY		
Preferred stock, $0.0001 par value; 20,000,000 shares authorized, 0 shares issued and outstanding	-	-
Common stock, $0.0001 par value; 80,000,000 shares authorized, 20,412,024 and 20,392,024 shares issued and outstanding	2,041	2,039
Additional paid-in capital	25,154,571	24,504,525
Deficit accumulated during the development stage	(23,580,861)	(16,644,156)
Total stockholders' equity	1,575,751	7,862,408
Total liabilities and stockholders' equity	$ 3,283,473	$ 10,829,788

See accompanying notes to financial statements

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
STATEMENTS OF OPERATIONS

	Year ended December 31, 2009	2008	Period from August 1, 2005 (inception) through December 31, 2009
Operating expenses			
Research and development	$ 5,444,202	$ 9,768,389	$ 18,477,688
General and administrative	1,520,648	2,315,178	4,201,235
Total operating expenses	6,964,850	12,083,567	22,678,923
Loss from operations	(6,964,850)	(12,083,567)	(22,678,923)
Other income (expense)			
Interest income	28,145	206,054	358,161
Interest expense	-	(1,036,053)	(1,260,099)
Total other income (expense)	28,145	(829,999)	(901,938)
Net loss	$ (6,936,705)	$ (12,913,566)	$ (23,580,861)
Net loss per share - basic and diluted	$ (0.34)	$ (0.81)	
Weighted average shares outstanding - basic and diluted	20,399,092	16,022,836	

See accompanying notes to financial statements

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
STATEMENT OF STOCKHOLDERS' EQUITY (DEFICIENCY)
Period from August 1, 2005 (inception) through December 31, 2009

	Common Stock Shares	Amount	Additional paid-in capital	Deficit accumulated during the development stage	Total stockholders' equity (deficiency)
Issuance of common stock to founders at $0.0001 per share	9,968,797	$ 997	$ 4,003	$ -	$ 5,000
Stock based compensation for services	-	-	9,700	-	9,700
Net loss, period from August 1, 2005 (inception) through December 31, 2006	-	-	-	(370,893)	(370,893)
Balance at December 31, 2006	9,968,797	997	13,703	(370,893)	(356,193)
Stock based compensation for services	-	-	88,300	-	88,300
Net loss, year ended December 31, 2007	-	-	-	(3,359,697)	(3,359,697)
Balance at December 31, 2007	9,968,797	997	102,003	(3,730,590)	(3,627,590)
Common stock sold in private placement, net of issuance costs of $141,646	7,360,689	736	17,689,301	-	17,690,037
Conversion of notes payable upon closing of private placement	1,962,338	196	4,278,322	-	4,278,518
Discount arising from note conversion	-	-	475,391	-	475,391
Warrants issued in connection with note conversion	-	-	348,000	-	348,000
Reverse merger transaction - elimination of accumulated deficit	-	-	(120,648)	-	(120,648)
Previously issued Laurier common stock	1,100,200	110	120,538	-	120,648
Warrants issued for services	-	-	480,400	-	480,400
Stock based compensation for services	-	-	1,131,218	-	1,131,218
Net loss, year ended December 31, 2008	-	-	-	(12,913,566)	(12,913,566)
Balance at December 31, 2008	20,392,024	2,039	24,504,525	(16,644,156)	7,862,408

Stock based compensation for services	-	-	647,448	-	647,448
Stock option exercise	20,000	2	2,598	-	2,600
Net loss, year ended December 31, 2009	-	-	-	(6,936,705)	(6,936,705)
Balance at December 31, 2009	20,412,024	$ 2,041	$ 25,154,571	$ (23,580,861)	$ 1,575,751

See accompanying notes to financial statements

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
STATEMENTS OF CASH FLOWS

	Year ended December 31, 2009	2008	Period from August 1, 2005 (inception) through December 31, 2009
Cash flows from operating activities			
Net loss	$ (6,936,705)	$ (12,913,566)	$ (23,580,861)
Adjustments to reconcile net loss to net cash used in operating activities			
Depreciation and amortization	19,337	45,467	77,913
Stock based compensation	647,448	1,131,218	1,876,666
Write-off of intangible assets	-	-	85,125
Warrants issued for services	-	480,400	480,400
Warrants issued in connection with note conversion	-	348,000	348,000
Note discount arising from beneficial conversion feature	-	475,391	475,391
Deferred rent	(1,323)	17,242	16,070
Loss on disposal of assets	2,680	-	2,680
Non cash interest expense	-	98,930	311,518
Changes in operating assets and liabilities			
Prepaid expenses	204,425	(240,922)	(110,589)
Restricted cash	-	(44,018)	(44,018)
Security deposit	12,165	-	-
Accounts payable	(1,490,628)	2,382,184	1,003,030
Accrued expenses	105,491	(669,315)	556,204
Due to related parties	126,802	5,033	132,418
Net cash used in operating activities	(7,310,308)	(8,883,956)	(18,370,053)
Cash flows from investing activities			
Purchase of equipment	-	(37,317)	(77,160)
Cash paid for intangible assets	-	-	(85,125)
Proceeds from related party advance	-	-	525,000
Repayment of related party advance	-	-	(525,000)
Net cash used in investing activities	$ -	$ (37,317)	$ (162,285)

See accompanying notes to financial statements

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
STATEMENTS OF CASH FLOWS (Continued)

	Year ended December 31, 2009	2008	Period from August 1, 2005 (inception) through December 31, 2009
Cash flows from financing activities			
Deferred financing fees paid	$ -	$ (20,000)	$ (45,000)
Proceeds from issuance of common stock in private placement, net	-	17,690,037	17,690,037
Proceeds from issuance of common stock to founders	-	-	5,000
Proceeds from issuance of notes payable	-	1,000,000	1,000,000
Repayment of notes payable	-	(1,000,000)	(1,000,000)
Proceeds from issuance of convertible notes payable	-	-	3,967,000
Proceeds from exercise of stock options	2,600	-	2,600
Net cash provided by financing activities	2,600	17,670,037	21,619,637
Net increase (decrease) in cash and cash equivalents	(7,307,708)	8,748,764	3,087,299
Cash and cash equivalents - beginning of period	10,395,007	1,646,243	-
Cash and cash equivalents - end of period	$ 3,087,299	$ 10,395,007	$ 3,087,299
Supplemental schedule of cash flows information			
Cash paid for interest	$ -	$ 80,000	$ 80,000
Supplemental disclosure of non-cash and financing activities			
Conversion of notes payable and interest to common stock	$ -	$ 4,278,518	$ 4,278,518
Common shares of Laurier issued in reverse merger transaction	$ -	$ 110	$ 110

See accompanying notes to financial statements

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
NOTES TO FINANCIAL STATEMENTS
Years ended December 31, 2009 and 2008 and the period
from August 1, 2005 (inception) to December 31, 2009

NOTE 1 – DESCRIPTION OF BUSINESS

Arno Therapeutics, Inc. ("Arno" or "the Company") develops innovative drug candidates for the treatment of cancer. Arno's lead clinical drug candidate, AR-12, is a potentially first-in-class, orally available, targeted anti-cancer agent that inhibits phosphoinositide dependent protein kinase-1 ("PDK-1"), a protein in the PI3K/Akt pathway, and also causes cell death through the induction of endoplasmic reticulum ("ER") stress. The PI3K/Akt pathway is of increasing interest in oncology, as it is believed to be involved in the survival of cancer cells and their ability to become resistant to therapy. AR-12's ability to both inhibit PDK-1 and cause ER stress may provide a unique therapeutic profile that helps to differentiate this molecule from other PI3K/Akt inhibitors in development. Arno is currently enrolling patients with solid tumors or lymphomas in a Phase I clinical trial who have progressed despite treatment with other therapies. In this study, Arno is measuring biomarkers in order to demonstrate preliminary proof-of-concept of AR-12 as a PDK-1 inhibitor in cancer patients. The Phase I study is also designed to evaluate the safety and tolerability of the drug candidate. Arno plans to complete the AR-12 Phase I study by the end of 2010 and begin Phase II development in the first half of 2011.

Arno's second drug candidate, AR-67, is a novel, third-generation camptothecin analogue that is currently in Phase II clinical development. Camptothecins treat cancer by disrupting cell division through the inhibition of topoisomerase I, a critical enzyme in DNA replication. Through several mechanisms of action, camptothecins preferentially target cancer cells to normal tissues. Two second-generation therapies, topotecan (Hycamtin®, Glaxo-Smith-Kline) and irinotecan (also known as "CPT-11" and marketed as Camptostar® by Pfizer), are approved by the United States Food and Drug Administration (the FDA). In clinical studies, AR-67 has demonstrated activity and an excellent safety profile as well as improved pharmacokinetic properties when compared to these approved therapies. We believe these advantages could allow AR-67 to become a leading product in the camptothecin market. Arno is currently conducting a Phase II study in patients with glioblastoma multiforme (GBM), an aggressive form of brain cancer. Arno expects to have interim data from both of these studies in 2010.

Arno is also developing AR-42, an orally available, targeted broad spectrum histone deacetylase (pan-HDAC) inhibitor. HDAC inhibitors are a growing class of compounds that target histone deactylase, a molecule involved in determining which genes are expressed in a particular cell. This class has two approved agents, vorinostat (SAHA, or Zolinza® by Merck) and romidepsin (Istodax® by Celgene).

In preclinical studies, AR-42 has been shown activity against a broad spectrum of deacetylation targets and increased potency compared to SAHA. This combination of activity and potency could make AR-42 an important member of this emerging class of compounds. Arno plans to initiate a Phase I study of AR-42, in collaboration with The Ohio State University in the first half of 2010.

The Company was incorporated in Delaware in March 2000, at which time its name was Laurier International, Inc. ("Laurier"). Pursuant to an Agreement and Plan of Merger dated March 6, 2008 (as amended, the "Merger Agreement"), by and among the Company, Arno Therapeutics, Inc., a Delaware corporation formed on August 1, 2005 ("Old Arno"), and Laurier Acquisition, Inc., a Delaware corporation and wholly-owned subsidiary of the Company ("Laurier Acquisition"), on June 3, 2008, Laurier Acquisition merged with and into Old Arno, with Old Arno remaining as the surviving corporation and a wholly-owned subsidiary of Laurier. Immediately following this merger, Old Arno merged with and into Laurier and Laurier's name was changed to Arno Therapeutics, Inc. These two merger

transactions are hereinafter collectively referred to as the “Merger.” Immediately following the Merger, the former stockholders of Old Arno collectively held 95% of the outstanding common stock of Laurier, assuming the issuance of all shares issuable upon the exercise of outstanding options and warrants, and all of the officers and directors of Old Arno in office immediately prior to the Merger were appointed as the officers and directors of Laurier immediately following the Merger. Further, Laurier was a non-operating shell company prior to the Merger. The merger of a private operating company into a non-operating public shell corporation with nominal net assets is considered to be a capital transaction in substance, rather than a business combination, for accounting purposes. Accordingly, the Company treated this transaction as a capital transaction without recording goodwill or adjusting any of its other assets or liabilities. All costs incurred in connection with the Merger have been expensed. Upon completion of the Merger, the Company adopted Old Arno’s business plan.

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On May 1, 2009, the Company filed a request with Securities Exchange Commission (the "SEC") to deregister their unissued and unsold shares of common stock previously registered under Form S-1. On May 5, 2009, the Company filed Form 15 with the SEC to effectively terminate their registration.

NOTE 2 – BASIS OF PRESENTATION

The Company is a development stage enterprise since it has not yet generated any revenue from the sale of products and, through December 31, 2009, its efforts have been principally devoted to developing its licensed technologies and raising capital. Accordingly, the accompanying financial statements have been prepared in accordance with the provisions of Accounting Standards Codification (the "ASC") 915, "Development Stage Entities".

For the years ended December 31, 2009 and 2008, the Company reported a net loss of $6,936,705 and $12,913,566, respectively, and the net loss from August 1, 2005 (inception) through December 31, 2009 was $23,580,861. The Company's total cash balance as of December 31, 2009 was $3,087,299 compared to $10,395,007at December 31, 2008. Through December 31, 2009, all of the Company's financing has been through private placements of common stock and debt financing. In June 2008, the Company completed a private placement of its common stock, raising approximately $17,832,000 in gross proceeds. The Company expects to incur substantial and increasing losses and have negative net cash flows from operating activities as it expands its technology portfolio and engages in further research and development activities, particularly the conducting of pre-clinical and clinical trials.

The Company plans to continue to fund operations from its existing cash balance and additional funds raised through various sources, such as equity and debt financing. Based on its current resources at December 31, 2009, and the current plan of expenditure on continuing development of current products, the Company believes that it has sufficient capital to fund its operations through the end of the second quarter of 2010, and will need additional financing in the future until it can achieve profitability, if ever.

The success of the Company depends on its ability to develop its products to the point of FDA approval and subsequent revenue generation and, accordingly, to raise enough capital to finance these developmental effort, or the ability to enter into a strategic transaction with a third party. The Company plans to raise additional capital to finance the continued operating and capital requirements of the Company. Amounts raised will be used to further develop the Company's products, acquire additional product licenses and for other working capital purposes. However, there can be no assurance that the Company will be able to raise additional capital at times or on terms that it desires, if at all, particularly given the current economic conditions, which have made access to the capital markets more difficult. If the Company is unable to raise or otherwise secure additional capital, it will likely be forced to curtail its operations, which would delay the development of its product candidates.

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NOTE 3 – THE MERGER

Description of the Merger and Private Placement Offering

On June 3, 2008, the Company completed the Merger. In accordance with the terms of the Merger, each share of common stock of Old Arno that was outstanding immediately prior to the Merger was exchanged for 1.99377 shares of the Company's common stock. In addition, all securities convertible into or exercisable for shares of Old Arno common stock outstanding immediately prior to the Merger were cancelled, and the holders thereof received similar securities convertible into or exercisable for the purchase of an aggregate of 1,611,760 shares of the Company's common stock. In consideration for their shares of Old Arno's common stock, the shareholders received an aggregate of 19,291,824 shares of the Company's common stock. Immediately prior to the effective time of the Merger, 1,100,200 shares of Laurier's common stock were issued and outstanding. Upon completion of the Merger, the Old Arno shareholders owned approximately 95% of the Company's issued and outstanding common stock.

Following the Merger, the business conducted by the Company is the business conducted by Old Arno prior to the Merger. In addition, the directors and officers of Laurier were replaced by the directors and officers of Old Arno.

As a condition and immediately prior to the closing of the Merger, on June 2, 2008, Old Arno completed a private placement of its equity securities whereby it received gross proceeds of approximately $17,732,000 through the sale of 3,691,844 shares of Old Arno Common Stock to selected accredited investors, which shares were exchanged for 7,360,689 shares of Company Common Stock after giving effect to the Merger. Contemporaneously with the June 2008 private placement, the 6% Notes (Note 8) converted into 984,246 Old Arno Shares and the holders of the 6% Notes received Warrants to purchase an aggregate of 98,409 shares of Old Arno Common Stock at an exercise price equal to $4.83 per share. The Old Arno Shares were exchanged for an aggregate of 1,962,338 shares of the Company's Common Stock and the Conversion Warrants were exchanged for an aggregate of 196,189 five year warrants to purchase our Common Stock at an exercise price equal to $2.42 per share.

All references to share and per share amounts in these financial statements have been restated to retroactively reflect the number of common shares of Arno common stock issued pursuant to the Merger.

Accounting Treatment of the Merger; Financial Statement Presentation

The Merger was accounted for as a reverse acquisition pursuant to the guidance in Appendix B of SEC Accounting Disclosure Rules and Practices Official Text, which provides that the "merger of a private operating company into a non-operating public shell corporation with nominal net assets typically results in the owners and management of the private company having actual or effective operating control of the combined company after the transaction, with the shareholders of the former public shell continuing only as passive investors. These transactions are considered by the Securities and Exchange Commission to be capital transactions in substance, rather than business combinations. That is, the transaction is equivalent to the issuance of stock by the private company for the net monetary assets of the shell corporation, accompanied by a recapitalization." Accordingly, the Merger has been accounted for as a recapitalization, and, for accounting purposes, Old Arno is considered the acquirer in a reverse acquisition.

Laurier's historical accumulated deficit for periods prior to June 3, 2008, in the amount of $120,538, was eliminated against additional-paid-in-capital, and the accompanying financial statements present the previously issued shares of Laurier's common stock as having been issued pursuant to the Merger on June 3, 2008. The shares of common stock of the Company issued to the Old Arno stockholders in the Merger are presented as having been outstanding since August 2005 (the month when Old Arno first sold its equity securities).

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Because the Merger was accounted for as a reverse acquisition under accounting principles generally accepted in the United States ("GAAP"), the financial statements for periods prior to June 3, 2008, reflect only the operations of Old Arno.

NOTE 4 – SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

Use of Estimates

The preparation of financial statements in conformity with GAAP requires that management make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenues and expenses during the reporting periods. Estimates and assumptions principally relate to services performed by third parties but not yet invoiced, estimates of the fair value of stock options issued to employees and consultants, and estimates of the probability and potential magnitude of contingent liabilities. Actual results could differ from those estimates.

Cash and Cash Equivalents

The Company considers all highly liquid investments with a remaining maturity of three months or less at the time of acquisition to be cash equivalents.

Restricted Cash

In October 2008, the Company entered into a five year office lease agreement with an opt-out provision at the end of the third year. In connection with the lease, the Company delivered an irrevocable stand-by and unconditional letter of credit in the amount of approximately $44,000 (or the approximate equivalent of three months rent) as a security deposit with the landlord as the beneficiary in case of default or failure to comply with the lease requirements. In order to fund the letter of credit, the Company deposited a compensating balance of approximately $44,000 into an interest bearing certificate of deposit with a financial institution which shall be reduced by $11,005 on January 1, 2011 and by an additional $11,005 on January 1, 2013, provided the Company maintains certain conditions described in the lease agreement.

Prepaid Expenses

Prepaid expenses consist of payments made in advance to vendors relating to service contracts for clinical trial development and insurance policies. These advanced payments are amortized to expense either as services are performed or over the relevant service period using the straight line method.

Property and Equipment

Property and equipment consist primarily of furnishings, fixtures, leasehold improvements and computer equipment and are recorded at cost. Repairs and maintenance costs are expensed in the period incurred. Depreciation is provided for by the straight-line method over the estimated useful lives of the related assets. Leasehold improvements are amortized using the straight-line method over the remaining lease term or the life of the asset, whichever is shorter.

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Description	Estimated useful life
Office equipment and furniture	5 to 7 years
Leasehold improvements	3 years
Computer equipment	3 years

Fair Value of Financial Instruments

The Company measures fair value in accordance with GAAP. Fair value measurements are applied under other accounting pronouncements that require or permit fair value measurements. The provisions are to be applied prospectively as of the beginning of the fiscal year in which it is initially adopted, with any transition adjustment recognized as a cumulative-effect adjustment to the opening balance of retained earnings. The adoption of this standard had no significant impact on the Company's financial statements. Financial instruments included in the Company's balance sheets consist of cash and cash equivalents, accounts payable, accrued expenses and due to related parties. The carrying amounts of these instruments reasonably approximate their fair values due to their short-term maturities.

Concentration of Credit Risk

Financial instruments which potentially subject the Company to concentrations of credit risk consist principally of cash and cash equivalents. The Company deposits cash and cash equivalents with high credit quality financial institutions and is insured to the maximum limitations. Balances in these accounts may exceed federally insured limits at times, which expose the Company to institutional risk.

Research and Development

Research and development costs are charged to expense as incurred. Research and development includes employee costs, fees associated with operational consultants, contract clinical research organizations, contract manufacturing organizations, clinical site fees, contract laboratory research organizations, contract central testing laboratories, licensing activities, and allocated executive, human resources and facilities expenses. The Company accrues for costs incurred as the services are being provided by monitoring the status of the trial and the invoices received from its external service providers. The Company adjusts its accruals in the period when actual costs become known. Costs related to the acquisition of technology rights and patents for which development work is still in process are charged to operations as incurred and considered a component of research and development expense.

Stock-Based Compensation

Stock-based compensation cost is measured at the grant date based on the value of the award and is recognized as expense over the required service period, which is generally equal to the vesting period.

Common stock, stock options, and warrants or other equity instruments issued to non-employees (including consultants and all members of the Company's Scientific Advisory Board) as consideration for goods or services received by the Company are accounted for based on the fair value of the equity instruments issued (unless the fair value of the consideration received can be more reliably measured). The fair value of any options issued to non-employees is recorded as expense over the applicable service periods.

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Loss per Common Share

Basic loss per share is computed by dividing the loss available to common shareholders by the weighted-average number of common shares outstanding. Diluted loss per share is computed similarly to basic loss per share except that the denominator is increased to include the number of additional common shares that would have been outstanding if the potential common shares had been issued and if the additional common shares were dilutive. For all periods presented, potentially dilutive securities are excluded from the computation of fully diluted loss per share as their effect is anti-dilutive.

Potentially dilutive securities as of December 31, 2009 and 2008 include:

	2009	2008
Warrants to purchase common stock	$ 495,252	$ 495,252
Options to purchase common stock	1,913,241	2,436,511
Total potential dilutive securities	$ 2,408,493	$ 2,931,763

Comprehensive Loss

The Company has no components of other comprehensive loss other than its net loss, and accordingly, comprehensive loss is equal to net loss for all periods presented.

Income Taxes

The Company recognizes deferred tax assets and liabilities for the expected future tax consequences of events that have been included in the financial statements or tax returns. Under this method, deferred income taxes are recognized for the tax consequences in future years of differences between the tax bases of assets and liabilities and their financial reporting amounts at each year-end based on enacted tax laws and statutory tax rates applicable to the period in which the differences are expected to affect taxable income. The Company provides a valuation allowance when it appears more likely than not that some or all of the net deferred tax assets will not be realized.

A tax position is recognized as a benefit only if it is “more likely than not” that the tax position would be sustained in a tax examination, with a tax examination being presumed to occur. The amount recognized is the largest amount of tax benefit that is greater than 50% likely of being realized on examination. For tax positions not meeting the “more likely thank not” test, no tax benefit is recorded.

The Company’s policy is to include interest and penalties related to unrecognized tax benefits within the Company’s provision for (benefit from) income taxes. The Company recognized no amounts for interest and penalties related to unrecognized tax benefits in 2009 and 2008 respectively. In addition the Company had no amounts accrued for interest and penalties as of December 31, 2009 and 2008, respectively.

Reclassification

Certain amounts in the prior period financial statements and related notes have been reclassified to conform to the current period presentation. Reclassifications had no effect on net loss or stockholders' equity.

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Recently Issued Accounting Pronouncements

In June 2009, the Financial Accounting Standards Board ("FASB") issued the ASC. The ASC has become the single source of non-governmental GAAP recognized by the FASB in the preparation of financial statements. The Company adopted the ASC as of July 1, 2009. The ASC does not change GAAP and did not have an effect on the Company's financial position, results of operations or cash flows.

Effective June 30, 2009, the Company adopted a new accounting standard issued by the FASB related to the disclosure requirements of the fair value of financial instruments. This standard expands the disclosure requirements of fair value (including the methods and significant assumptions used to estimate fair value) of certain financial instruments to interim period financial statements that were previously only required to be disclosed in financial statements for annual periods. In accordance with this standard, the disclosure requirements have been applied on a prospective basis and did not have a material impact on the Company's financial statements.

Effective June 30, 2009, the Company adopted a newly issued accounting standard related to accounting for and disclosure of subsequent events in its financial statements. This standard provides the authoritative guidance for subsequent events that was previously addressed only in United States auditing standards. This standard establishes general accounting for and disclosure of events that occur after the balance sheet date but before financial statements are issued or are available to be issued and requires the Company to disclose the date through which it has evaluated subsequent events and whether that was the date the financial statements were issued or available to be issued. This standard does not apply to subsequent events or transactions that are within the scope of other applicable GAAP that provide different guidance on the accounting treatment for subsequent events or transactions. The adoption of this standard did not have a material impact on the Company's financial statements.

In August 2009, the FASB issued an amendment to the accounting standards related to the measurement of liabilities that are recognized or disclosed at fair value on a recurring basis. This standard clarifies how a company should measure the fair value of liabilities and that restrictions preventing the transfer of a liability should not be considered as a factor in the measurement of liabilities within the scope of this standard. The adoption of this standard did not have a material impact on the Company's financial statements

Management does not believe that any other recently issued, but not yet effective, accounting pronouncements, if currently adopted, would have a material effect on the Company's financial statements.

NOTE 5 – PROPERTY AND EQUIPMENT

Property and equipment as of December 31, 2009 and 2008 consist of the following:

	2009	2008
Computer equipment	$ 5,080	$ 12,602
Office furniture and equipment	48,380	53,802
Leasehold improvements	9,144	10,756

Total property and equipment	62,604	77,160
Accumulated depreciation	(21,037)	(13,576)
Total property and equipment, net	$ 41,567	$ 63,584

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Depreciation expense for the years ended December 31, 2009, 2008 and the period from August 1, 2005 (inception) through December 31, 2009 were $19,337, $11,926 and $32,912, respectively.

NOTE 6 - INTANGIBLE ASSETS AND INTELLECTUAL PROPERTY

AR-67 License Agreement

The Company's rights to AR-67 are governed by an October 2006 license agreement with the University of Pittsburgh ("Pitt"). Under this agreement, Pitt granted the Company an exclusive, worldwide, royalty-bearing license for the rights to commercialize technologies embodied by certain issued patents, patent applications and know-how relating to AR-67 for all therapeutic uses. The Company has expanded, and intends to continue to expand, its patent portfolio by filing additional patents covering expanded uses for this technology.

Under the terms of the license agreement with Pitt, the Company made a one-time cash payment of $350,000 to Pitt and reimbursed it for past patent expenses of approximately $373,000. Additionally, Pitt will receive performance-based cash payments upon successful completion of clinical and regulatory milestones relating to AR-67. The Company will make the first milestone payment to Pitt upon the acceptance of the first New Drug Application ("NDA") by the FDA for AR-67. The Company is also required to pay to Pitt an annual maintenance fee of $200,000 upon the third and fourth anniversaries, $250,000 upon the fifth and sixth anniversaries, and $350,000 upon the seventh anniversary and annually thereafter and to pay Pitt a royalty equal to a percentage of net sales of AR-67, pursuant to the license agreement. To the extent the Company enters into a sublicensing agreement relating to AR-67, the Company will pay Pitt a portion of all non-royalty income received from such sublicensee.

Under the license agreement with Pitt, the Company also agreed to indemnify and hold Pitt and its affiliates harmless from any and all claims, actions, demands, judgments, losses, costs, expenses, damages and liabilities (including reasonable attorneys' fees) arising out of or in connection with (i) the production, manufacture, sale, use, lease, consumption or advertisement of AR-67, (ii) the practice by the Company or any affiliate or sublicensee of the licensed patent; or (iii) any obligation of the Company under the license agreement unless any such claim is determined to have arisen out of the gross negligence, recklessness or willful misconduct of Pitt. The license agreement will terminate upon the expiration of the last patent relating to AR-67. Pitt may generally terminate the agreement at any time upon a material breach by the Company to the extent it fails to cure any such breach within 60 days after receiving notice of such breach or in the event the Company files for bankruptcy. The Company may terminate the agreement for any reason upon 90 days prior written notice.

AR-12 and AR-42 License Agreements

The Company's rights to both AR-12 and AR-42 are governed by separate license agreements with The Ohio State University Research Foundation ("Ohio State") entered into in January 2008. Pursuant to each of these agreements, Ohio State granted the Company exclusive, worldwide, royalty-bearing licenses to commercialize certain patent applications, know-how and improvements relating to AR-42 and AR-12 for all therapeutic uses.

Pursuant to the Company's license agreements for AR-12 and AR-42, the Company made one-time cash payments to Ohio State in the aggregate amount of $450,000 and reimbursed it for past patent expenses in the aggregate amount of

approximately $174,000. Additionally, the Company will be required to make performance-based cash payments upon successful completion of clinical and regulatory milestones relating to AR-12 and AR-42 in the United States, Europe and Japan. The first milestone payment for each of the licensed compounds will be due when the first patient is dosed in the first Company sponsored Phase I clinical trial of each of AR-12 and AR-42. In October 2009, the Company remitted a $200,000 milestone payment to Ohio State for the first patient dosed in the first Company sponsored Phase I clinical trial of AR-12. To the extent the Company enters into a sublicensing agreement relating to either or both of AR-12 or AR-42, it will be required to pay Ohio State a portion of all non-royalty income received from such sublicensee.

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The license agreements with Ohio State further provide that the Company will indemnify Ohio State from any and all claims arising out of the death of or injury to any person or persons or out of any damage to property, or resulting from the production, manufacture, sale, use, lease, consumption or advertisement of either AR-12 or AR-42, except to the extent that any such claim arises out of the gross negligence or willful misconduct of Ohio State. The license agreements for AR-12 and AR-42 each expire on the later of (i) the expiration of the last valid claim contained in any licensed patent and (ii) 20 years after the effective date of the license. Ohio State will generally be able to terminate either license upon the Company's breach of the terms of the license to the extent the Company fails to cure any such breach within 90 days after receiving notice of such breach or the Company files for bankruptcy. The Company may terminate either license upon 90 days prior written notice.

NOTE 7 - ACCRUED LIABILITIES

Accrued liabilities as of December 31, 2009 and 2008 consist of the following:

	2009	2008
Accrued compensation and related benefits	$ 250,800	$ 274,268
Accrued research and development expense	305,404	98,761
Accrued other expenses	-	77,684
Total accrued liabilities	$ 556,204	$ 450,713

NOTE 8 - CONVERTIBLE NOTES PAYABLE

During February 2007, the Company completed a private placement offering of 6% convertible promissory notes (the "Notes") for an aggregate principal amount of $3,967,000, due on February 9, 2009. The aggregate principal amount and accrued but unpaid interest on the Notes, which totaled $4,278,518, automatically converted upon the closing of the June 3, 2008 Private Placement into 1,962,338 shares of common stock at a conversion price of $2.42, which was equal to 90% of the per share price of the shares sold in the Financing. Due to the beneficial conversion feature resulting from the discounted conversion price, a discount of $475,391 was recorded as interest expense with a corresponding credit to additional paid-in capital. In addition, in conjunction with the conversion of the convertible debt, the Company issued fully vested warrants to purchase 196,189 shares of common stock to the holders of the Notes. The warrants were valued at $348,000 using the Black-Scholes option-pricing model and the following assumptions: exercise price $2.42, a 3.41% risk-free interest rate, a five year contractual term, a dividend rate of 0%, and 94.30% expected volatility. The cost of the warrants was included in interest expense in the accompanying Statements of Operations, and as an increase in additional paid-in capital.

NOTE 9 – STOCKHOLDERS' EQUITY

Common Stock

In August 2005, the Company issued an aggregate of 9,968,797 shares of common stock to its founders for $5,000.

As a condition to the closing of the Merger, on June 2, 2008, the Company completed a private placement of 7,360,689 shares of its common stock (as adjusted to give effect to the Merger), resulting in gross proceeds of approximately $17,832,000. Issuance costs related to the private placement were approximately $142,000, which were capitalized and charged to stockholders' equity upon the closing of the private placement. In accordance with the terms of the Notes, contemporaneously with the completion of the June 2, 2008 private placement, the outstanding principal and accrued interest of $4,278,518 under the Notes converted into an aggregate of 1,962,338 shares of common stock. Additionally, 1,100,200 shares of common stock that were held by the original stockholders of Laurier prior to the Merger are reflected in the Company's common stock outstanding in the accompanying financial statements.

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Warrants

In connection with the in-licensing of the Company's product candidates AR-12 and AR-42, the Company issued 299,063 fully vested warrants to employees of Two River Group Holdings, LLC (see Note 10) and a consultant for their consultation and due diligence efforts as part of a finder's fee arrangement. The warrants have an exercise price of $2.42 and were valued at $480,400 based upon the Black-Scholes option-pricing model. The assumptions used under the Black-Scholes option-pricing model included a risk free interest rate of 3.27%, volatility of 80.80% and a five year life. None of these warrants have been exercised to date.

In conjunction with the conversion of $4,278,518 of convertible debt, including accrued interest, prior to the Merger, the Company issued fully vested warrants to purchase 196,189 shares of common stock to the holders of such debt. The warrants were issued with an exercise price of $2.42 and expire in June 2, 2013. The fair value of the warrants was determined to be $348,000. None of these warrants have been exercised to date.

NOTE 10 – STOCK OPTION PLAN

The Company's 2005 Stock Option Plan (the "Plan") was originally adopted by the Board of Directors of Old Arno in August 2005, and was assumed by the Company on June 3, 2008 in connection with the Merger. After giving effect to the Merger, there are 2,990,655 shares of the Company's common stock reserved for issuance under the Plan. Under the Plan, incentives may be granted to officers, employees, directors, consultants, and advisors. Incentives under the Plan may be granted in any one or a combination of the following forms: (a) incentive stock options and non-statutory stock options, (b) stock appreciation rights, (c) stock awards, (d) restricted stock and (e) performance shares.

The Plan is administered by the Board of Directors, or a committee appointed by the Board, which determines recipients and types of awards to be granted, including the number of shares subject to the awards, the exercise price and the vesting schedule. The term of stock options granted under the Plan cannot exceed 10 years. Options shall not have an exercise price less than the fair market value of the Company's common stock on the grant date, and generally vest over a period of three to four years.

The weighted-average grant-date fair value of options granted for the years ended December 31, 2009 and 2008 was $0.91 and $2.09 respectively. The total intrinsic value of the options exercised during the years ended December 31, 2009 and 2008 was $121,129 and $0 respectively.

As of December 31, 2009, an aggregate of 1,057,414 shares remained available for future grants and awards under the Plan, which covers stock options, warrants and restricted awards. The Company issues unissued shares to satisfy stock options, warrants exercises and restricted stock awards.

A summary of the status of the options issued under the Plan as of December 31, 2009, and information with respect to the changes in options outstanding is a follows:

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		Options Outstanding		
	Shares Available for Grant	Outstanding Stock Options	Weighted-average exercise price	Aggregate Intrinsic Value
Balance at January 1, 2008	2,302,806	687,849	$ 0.81	
Granted under the Plan	(1,748,662)	1,748,662	2.06	
Exercised	-	-	-	
Surrendered/cancelled	-	-	-	
Forfeited	-	-	-	
Balance at December 31, 2008	554,144	2,436,511	1.71	$ -
Granted under the Plan	(305,000)	305,000	1.00	
Exercised	-	(20,000)	0.13	
Surrendered/cancelled	-	-	-	
Forfeited	808,270	(808,270)	(2.98)	
Balance at December 31, 2009	1,057,414	1,913,241	$ 1.76	$ -
Exercisable at December 31, 2009		1,278,507	$ 1.71	$ -

The Company estimated the fair value of each option award granted using the Black-Scholes option-pricing model and the following assumptions for the years ended December 31, 2009 and 2008:

	2009	2008
Term	5 - 10 Years	5 - 10 Years
Volatility	135%	77 - 123%
Dividend yield	0%	0%
Risk-free interest rate	2.3 - 3.5%	1.5 - 3.2%
Forfeiture rate	0%	0%

As allowed by ASC 718 for companies with a short period of publicly traded stock history, management's estimate of expected volatility is based on the average expected volatilities of a sampling of five companies with similar attributes to the Company, including: industry, stage of life cycle, size and financial leverage.

The Company has no historical basis for determining expected forfeitures and, as such, compensation expense for stock-based awards does not include an estimate for forfeitures.

Stock-based compensation for the year ended December 31, 2009 and 2008 and for the period from August 1, 2005 (inception) through December 31, 2009, are as follows:

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ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
NOTES TO FINANCIAL STATEMENTS
Years ended December 31, 2009 and 2008 and the period
from August 1, 2005 (inception) to December 31, 2009

	Year ended December 31, 2009	2008	Period from August 1, 2005 (inception) through December 31, 2009
General and administrative	$ 270,293	$ 679,948	$ 773,526
Research and development	377,155	451,270	1,103,140
Total	$ 647,448	$ 1,131,218	$ 1,876,666

As of December 31, 2009, the total outstanding, and the total exercisable, options under the Plan were as follows:

Outstanding				Exercisable	
Range of Exercise Prices	Shares	Weighted - Average Remaining Contractual Life	Weighted - Average Exercise Price	Total Shares	Weighted - Average Exercise Price
$ 0.13	129,532	1.86	$ 0.13	129,532	$ 0.13
$ 1.00	823,381	6.84	1.00	501,434	1.00
$ 2.42	428,661	7.61	2.42	387,124	2.42
$ 2.75	440,000	8.52	2.75	178,750	2.75
$ 3.00	91,667	4.28	3.00	81,667	3.00
	1,913,241	6.94	$ 1.76	1,278,507	$ 1.71

As of December 31, 2009 and 2008, there was $768,933 and $1,837,582 of unrecognized compensation costs related to nonvested stock options, respectively. These costs are expected to be recognized over a weighted average period of approximately two years as of December 31, 2009.

NOTE 11 – RELATED PARTIES

On June 1, 2009, the Company entered into a services agreement with Two River Consulting, LLC (“TRC”) to provide various clinical development, operational, managerial, and administrative services to the Company for a period of one year. David M. Tanen, the Company’s President, Secretary and director, Arie S. Belldegrun, the Chairman of the Board of Directors, and Joshua A. Kazam, director are each partners of TRC. The terms of the Services Agreement were reviewed and approved by a special committee of the Company’s Board of Directors consisting of independent directors. None of the members of the special committee has any interest in TRC or the services agreement. As compensation for the services contemplated by the services agreement, the Company will pay to TRC a monthly cash fee of $50,000. On occasion, some of the Company’s expenses are paid by TRC.

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No interest is charged by TRC on any outstanding balance owed by the Company. For the years ended December 31, 2009 and 2008 and for the period from August 1, 2005 (inception) through December 31, 2009, total cash services and reimbursed expenses totaled $382,418, $0 and $382,418, respectively. As of December 31, 2009 the Company has a payable to TRC of $132,418.

Prior to June 1, 2009, some of the Company's expenses were paid by Two River Group Holdings, LLC ("Two River"), a company that is controlled by four of the Company's directors and founders. No interest is charged by Two River on any outstanding balance owed by the Company. For the years ended December 31, 2009 and 2008 and for the period from August 1, 2005 (inception) through December 31, 2009, reimbursable expenses totaled $16,977, $106,085 and $206,039, respectively. In addition, the Company granted fully vested warrants to purchase 299,063 shares of its common stock at an exercise price of $2.42 to the Two River employees who provided consultation and due diligence efforts related to the in-licensing of AR-12 and AR-42. The warrants have a five year life and a fair value of $480,400 based upon the Black-Scholes option-pricing model. As of December 31, 2009 and 2008, the Company had $0 and $5,616 payable to Two River, respectively.

The Company utilized the services of Riverbank Capital Securities, Inc. ("Riverbank"), a FINRA member broker dealer registered with the SEC, for investment banking and other investment advisory services in connection with the June 2008 private placement and the Notes. Riverbank is an entity controlled by several partners of Two River who are also officers and/or directors of the Company. The Company paid a $100,000 non-accountable expense allowance to Riverbank for services related to the June 2008 private placement and is not obligated to Riverbank for any future payments.

During March 2008, the Company issued an unsecured promissory note to an existing shareholder for an aggregate amount of $1,000,000 (the "Promissory Note"). The Promissory Note bore interest at 8% and was due in full on July 3, 2008. The Company repaid the Promissory Note and the full premium on June 3, 2008 when it completed the Merger.

The financial condition and results of operations of the Company, as reported, are not necessarily indicative of results that would have been reported had the Company operated completely independently.

On October 1, 2007, the Company adopted a 401(k) savings plan (the "401(k) Plan") for the benefit of its employees. Under the 401(k) Plan the Company is required to make contributions equal to 3% of eligible compensation for each eligible employee whether or not the employee contributes to the 401(k) Plan. For the years ended December 31, 2009 and 2008 and for the cumulative period from August 1, 2005 (inception) through December 31, 2009, the Company has recorded $2,944, $10,923 and $16,064 of matching contributions to the 401(k) Plan.

NOTE 12 – INCOME TAXES

The Company accounts for income taxes using the liability method, which requires the determination of deferred tax assets and liabilities, based on the differences between the financial statement and tax bases of assets and liabilities, using enacted tax rates in effect for the year in which the differences are expected to reverse. The net deferred tax asset is adjusted by a valuation allowance, if, based on the weight of available evidence, it is more likely than not that some portion of all of the net deferred tax assets will not be realized. The income tax returns of the Company are subject to examination by federal and state taxing authorities. Such examination could result in adjustments to net

income or loss, which changes could affect the income tax liabilities of the Company. The Company's tax returns are open for inspection for the four years ended December 31, 2009.

At December 31, 2009, the Company had no Federal income tax expense or benefit but did have Federal tax net operating loss carry-forwards of approximately $20,234,624, and a research and development credit carry-forward of $1,015,207. The federal net operating loss carry-forwards will begin to expire in 2026, unless previously utilized.

ARNO THERAPEUTICS, INC.
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Deferred income taxes reflect the net effect of temporary differences between the carrying amounts of assets and liabilities for financial reporting purposes and the amounts used for income tax purposes. Significant components of the Company's net deferred tax assets as of December 31, 2009 and 2008 are shown below.

	2009	2008
Research tax credit	$ 1,015,000	$ 702,000
Net operating loss carry forwards	8,900,000	6,111,000
Stock based compensation	735,000	528,000
Depreciation and amortization	-	(6,000)
Total net deferred tax assets	10,650,000	7,335,000
Valuation allowance	(10,650,000)	(7,335,000)
Net deferred tax assets	$ -	$ -

NOTE 13 – COMMITMENTS AND CONTINGENCIES

On October 20, 2008, the Company entered into a lease for office space consisting of 5,390 square feet in Parsippany, New Jersey. The lease commencement date was November 14, 2008, with lease payments beginning on January 1, 2009. The lease expiration date is five years from the rent commencement date. The total five year lease obligation is approximately $713,000.

The Company provided a security deposit of $44,018, or four months base rent, in the form of a letter of credit. The letter of credit may be reduced by $11,005 on January 1, 2011 and by an additional $11,005 on January 1, 2013, provided the Company maintains certain conditions described in the lease agreement. The Company has an early termination option, which provides the Company may terminate the lease on the third anniversary, upon providing the landlord nine months written notice prior to the third anniversary of the lease. If the Company exercises its termination option, the Company would be obligated to pay a fee of $53,641 which consists of unamortized costs and expenses incurred by the landlord in connection with the lease. The Company also has an option to extend the term of the lease for a period of five additional years, provided the Company gives notice to the landlord no later than 12 months prior to the expiration of the original term.

In November 2008, the Company abandoned an office lease for its Fairfield, New Jersey facility that it entered into on August 10, 2007. As a result, the Company recorded a liability of $28,798 on the date of abandonment, in accordance with ASC 420, "Exit or Disposal Cost Obligations."

The Company's remaining gross estimated lease obligation for its Fairfield, New Jersey office space is approximately $0 and $102,000 as of December 31, 2009 and 2008, respectively.

On April 2, 2009, the Company entered into a Termination and Release of Lease Agreement for its Fairfield, New Jersey facility pursuant to which the Company surrendered its $12,165 security deposit and made an additional

one-time payment of $40,500 in exchange for terminating the lease as of this date. The Company recorded additional liability and expense of $23,867 as of March 31, 2009.

The aggregate remaining minimum future payments under these leases at December 31, 2009 are approximately as follows:

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Year ended December 31,		
2010	$	144,000
2011		143,000
2012		143,000
2013		146,000
Total	$	576,000

On August 19, 2008, the Company entered into an employment agreement with its Chief Executive Officer, with an effective commencement date of employment beginning on September 3, 2008. The agreement provides for a term of two years expiring on September 2, 2010, and an initial base salary of $375,000, plus an annual target performance bonus of up to 50% of his base salary or $187,500. Pursuant to the employment agreement, the CEO received a stock option to purchase 430,000 shares of the Company's common stock at an exercise price of $3.00 per share, which shall vest pro rata on each anniversary of his employment. Additionally, pursuant to the employment agreement, the CEO received a stock option to purchase 430,000 performance options contingent upon the successful achievement of performance goals established by the compensation committee of the Board of Directors. The employment stock option grant had an approximate fair value of $896,500 at the date of grant based on the Black-Scholes option-pricing model. The employment agreement also entitles the CEO to certain severance benefits.

On May 14, 2009, the Company entered into a Separation Agreement and General Release (the "Separation Agreement") with its CEO, pursuant to which, the CEO's employment with the Company was terminated effective May 31, 2009. Pursuant to the Separation Agreement, the CEO received salary and benefits through July 31, 2009 and was granted an extended exercise window on his previously vested 71,667, stock options which resulted in additional stock compensation expense of $46,100 as a result of the modification. In addition as a result of the CEO's termination, 788,333 stock options were forfeited and $146,200 of previously recorded stock compensation expense related to these unvested stock options was reversed in the current period.

On June 11, 2008, the Company entered into an employment agreement with its Chief Financial Officer. The agreement provides for a term of two years expiring on July 15, 2010, and an initial base salary of $200,000, plus an annual target performance bonus of up to 30% of his base salary or $60,000.

In addition, the CFO received a one-time cash bonus in the amount of $25,000 and a stock option grant to purchase 440,000 shares of the Company's common stock at an exercise price equal to $2.75 per share. The right to purchase 25% of the shares subject to the stock option vests in July, 2009 and thereafter, the remaining shares vest in equal monthly installments over a 24 month period, subject to his continued employment with the Company. The stock option grant had an approximate fair value of $830,500 at the date of grant based on the Black-Scholes option-pricing model. The employment agreement also entitles the CFO to certain severance benefits.

On February 4, 2010, the CFO tendered his resignation with the Company effective February 15, 2010. As a result of the CFO's resignation, 206,250 stock options were forfeited.

On June 1, 2007, the Company entered into an employment agreement with its President and Chief Medical Officer. The agreement provides for a term of two years expiring on May 31, 2009, and an initial base salary of $340,000, plus an annual target performance bonus of up to $150,000. Pursuant to the employment agreement, the CMO received a stock option to purchase 398,754 shares of the Company's common stock at an exercise price of $1.00. The right to purchase 199,377 shares vests pro rata on the first two anniversaries of his employment, and the right to purchase the remaining 199,377 shares vest upon the achievement of performance milestones, of which one-half, or 99,689 shares vested as of May 31, 2008. The stock option grant had an approximate fair value of $252,800 at the date of grant based on the Black-Scholes option-pricing model. The employment agreement also entitles the CMO to certain severance benefits. However, in February 2009, the CMO informed the Company that he would not be continuing his employment with the Company beyond the expiration of his employment agreement on May 31, 2009.

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The Company has entered into various contracts with third parties in connection with the development of the licensed technology described in Note 6.

The aggregate minimum commitment under these contracts as of December 31, 2009 is approximately $758,000.

In the normal course of business, the Company enters into contracts that contain a variety of indemnifications with its employees, licensors, suppliers and service providers. Further, the Company indemnifies its directors and officers who are, or were, serving at the Company's request in such capacities. The Company's maximum exposure under these arrangements is unknown as of December 31, 2009. The Company does not anticipate recognizing any significant losses relating to these arrangements.

NOTE 14 – SUBSEQUENT EVENTS

On January 20, 2010, the Company received net proceeds of $322,016 through the sale of its New Jersey ("NJ") net operating losses ("NOLS"). In June 2009, the Company submitted an application with the New Jersey Economic Development Authority ("NJEDA") to sell its 2008 NOLS in return for cash consideration to the Company. The NJEDA in conjunction with the State of NJ offers financing and business incentives to NJ based biotechnology and technology companies. As a result of the sale of the NJ NOLS sale, the Company's NJ deferred tax asset carryforward is reduced by $4,000,000 beginning in 2010. The Company's federal net operating losses were not affected by the sale of the NJ NOLS, and thus has the full value of the 2009 deferred tax asset carryforward.

On September 3, 2010, the Company entered into a Securities Purchase and Registration Rights Agreement, or the Purchase Agreement, with a number of institutional and accredited investors pursuant to which the Company sold in a private placement an aggregate of 15,274,000 shares of newly-designated Series A Convertible Preferred Stock, par value $0.0001 per share, or Series A Preferred Stock, at a per share purchase price of $1.00. In accordance with the Purchase Agreement, the Company also issued 1,221,920, two-and-one-half-year Class A warrants to purchase additional shares of Series A Preferred Stock at an initial exercise price of $1.00 per share and 6,415,080, five-year Class B warrants at an initial exercise price of $1.15 per share. The terms of the Class A and Class B warrants contain an anti-dilutive price adjustment provision, such that in the event the Company issues common shares at a price below the current exercise price of the Warrants the exercise price of the Class A and Class B warrants will be adjusted based on the lower issuance price. The sale of the shares and warrants resulted in aggregate gross proceeds of approximately $15.2 million, before expenses.

Issuance costs related to the financing were approximately $1.9 million, including the issuance of warrants ("Placement Warrants") to purchase 1,050,840 shares of the Company's common stock at 110% of the Series A Preferred Stock purchase price per share to designees of Riverbank and I-Bankers Securities, Inc. ("IBS"), that acted as placement agents for the Company in connection with the private placement. The Placement Warrants were valued at $614,400 using the Black-Scholes option-pricing model.

Pursuant to the Registration Rights Agreement, the Company will use its best efforts to cause such registration statement to be declared effective within 180 days following the initial closing under the Purchase Agreement, or by March 8, 2011. If such registration statement is not declared effective by the SEC by such date, the Company will pay liquidated damages to the investors in the amount of 1% of each investor's aggregate investment amount for each

30-day period until the registration statement is declared effective.

On October 29, 2010, the Company was awarded funding of a total of approximately $733,000 under the IRS Qualifying Therapeutic Discovery Project ("QTDP") program, which was created as part of the Patient Protection and Affordable Care Act of 2010 (the "Healthcare Reform Act"). As enacted under the Healthcare Reform Act, the QTDP program provides a tax credit or grant of up to 50% of eligible costs and expenses for the tax years of 2009 and 2010 for qualifying research and development expenses incurred for innovative projects that are determined by the U.S. Department of Health and Human Services to have reasonable potential to result in a new therapy, reduce health care costs, or represent a significant advance in finding a cure for human disease. The Company was awarded approximately $244,000 for R&D expenses incurred for each of its AR-12, AR-42 and AR-67 development programs. The Company expects to receive the amounts granted prior to the end of 2010.

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
CONDENSED BALANCE SHEETS

	June 30, 2010 (unaudited)	December 31, 2009
ASSETS		
Current assets		
Cash and cash equivalents	$ 708,770	$ 3,087,299
Prepaid expenses	169,382	110,589
Total current assets	878,152	3,197,888
Property and equipment, net	35,642	41,567
Restricted cash	44,018	44,018
Total assets	$ 957,812	$ 3,283,473
LIABILITIES AND STOCKHOLDERS' EQUITY (DEFICIT)		
Current liabilities		
Accounts payable	$ 373,467	$ 1,003,030
Accrued expenses and other current liabilities	951,270	556,204
Due to related party	89,605	132,418
Total current liabilities	1,414,342	1,691,652
Deferred rent	15,409	16,070
Total liabilities	1,429,751	1,707,722
Commitments and contingencies		
Stockholders' (deficit) equity		
Preferred stock, $0.0001 par value, 10,000,000 shares authorized, 0 shares issued and outstanding	-	-
Common stock, $0.0001 par value, 80,000,000 shares authorized, 20,412,024 shares issued and outstanding	2,041	2,041
Additional paid-in capital	25,220,874	25,154,571
Deficit accumulated during the development stage	(25,694,854)	(23,580,861)
Total stockholders' (deficit) equity	(471,939)	1,575,751
Total liabilities and stockholders' (deficit) equity	$ 957,812	$ 3,283,473

See accompanying notes to condensed financial statements

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
CONDENSED STATEMENTS OF OPERATIONS

	Three months ended June 30,		Six months ended June 30,		Period from August 1, 2005 (inception) through June 30, 2010
	2010	2009	2010	2009	
Operating expenses:					
Research and development	$ 1,067,670	$ 1,160,512	$ 2,129,963	$ 3,236,897	$ 20,607,651
General and administrative	213,566	115,402	319,060	607,974	4,520,295
Total operating expenses	1,281,236	1,275,914	2,449,023	3,844,871	25,127,946
Loss from operations	(1,281,236)	(1,275,914)	(2,449,023)	(3,844,871)	(25,127,946)
Other income (expense):					
Interest income	1,925	1,066	3,417	26,197	361,578
Interest expense	-	-	-	-	(1,260,099)
Other income	327	-	331,613	-	331,613
Total other income (expense)	2,252	1,066	335,030	26,197	(566,908)
Net loss	$ (1,278,984)	$ (1,274,848)	$ (2,113,993)	$ (3,818,674)	$ (25,694,854)
Net loss per share - basic and diluted	$ (0.06)	$ (0.06)	$ (0.10)	$ (0.19)	
Weighted-average shares outstanding -basic and diluted	21,412,024	20,392,024	21,412,024	20,392,024	

See accompanying notes to condensed financial statements

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
CONDENSED STATEMENT OF STOCKHOLDERS' EQUITY (DEFICIENCY)
Period from August 1, 2005 (inception) through June 30, 2010
(unaudited)

	COMMON STOCK SHARES	COMMON STOCK AMOUNT	ADDITIONAL PAID-IN CAPITAL	DEFICIT ACCUMULATED DURING THE DEVELOPMENT STAGE	TOTAL STOCKHOLDERS' EQUITY (DEFICIT)
Issuance of common shares to founders at $0.0001 per share	9,968,797	$ 997	$ 4,003	$ -	$ 5,000
Stock based compensation for services	-	-	9,700	-	9,700
Net loss, period from August 1, 2005 (inception) through December 31, 2006	-	-	-	(370,893)	(370,893)
Balance at December 31, 2006	9,968,797	997	13,703	(370,893)	(356,193)
Stock based compensation for services	-	-	88,300	-	88,300
Net loss, year ended December 31, 2007	-	-	-	(3,359,697)	(3,359,697)
Balance at December 31, 2007	9,968,797	997	102,003	(3,730,590)	(3,627,590)
Common stock sold in private placement, net of issuance costs of $141,646	7,360,689	736	17,689,301	-	17,690,037
Conversion of notes payable upon closing of private placement	1,962,338	196	4,278,322	-	4,278,518
Note discount arising from note conversion	-	-	475,391	-	475,391
Warrants issued in connection with note conversion	-	-	348,000	-	348,000
Reverse merger transaction - elimination of accumulated deficit	-	-	(120,648)	-	(120,648)
previously issued Laurier common stock	1,100,200	110	120,538	-	120,648

Warrants issued for services	-	-	480,400	-	480,400
Stock based compensation for services	-	-	1,131,218	-	1,131,218
Net loss, year ended December 31, 2008				(12,913,566)	(12,913,566)
Balance at December 31, 2008	20,392,024	2,039	24,504,525	(16,644,156)	7,862,408
Stock based compensation for services	-	-	647,448	-	647,448
Stock option exercise	20,000	2	2,598	-	2,600
Net loss, year ended December 31, 2009				(6,936,705)	(6,936,705)
Balance at December 31, 2009	20,412,024	2,041	25,154,571	(23,580,861)	1,575,751
Stock based compensation for services	-	-	66,303	-	66,303
Net loss, six months ended June 30, 2010	-	-	-	(2,113,993)	(2,113,993)
Balance at June 30, 2010	20,412,024	$ 2,041	$ 25,220,874	$ (25,694,854)	$ (471,939)

See accompanying notes to condensed financial statements

ARNO THERAPEUTICS, INC.
(A DEVELOPMENT STAGE COMPANY)
CONDENSED STATEMENTS OF CASH FLOWS
(unaudited)

	Six months ended June 30,		Period from August 1, 2005 (inception) through June 30, 2010
	2010	2009	
Cash flows from operating activities			
Net loss	$ (2,113,993)	$ (3,818,674)	$ (25,694,854)
Adjustment to reconcile net loss to net cash used in operating activities			
Depreciation and amortization	5,925	9,265	83,838
Stock-based compensation	66,303	163,699	1,942,969
Write-off of intangible assets	-	-	85,125
Warrants issued for services	-	-	480,400
Warrants issued in connection with note conversion	-	-	348,000
Note discount arising from beneficial conversion feature	-	-	475,391
Deferred rent	(661)	(662)	15,409
Loss on disposal of assets	-	2,780	2,680
Noncash interest expense	-	-	311,518
Changes in operating assets and liabilities			
Prepaid expenses	(58,793)	134,395	(169,382)
Restricted cash	-	-	(44,018)
Security deposit	-	12,165	-
Accounts payable	(629,563)	(1,163,218)	373,467
Accrued expenses	395,066	(152,116)	951,270
Due to related party	(42,813)	50,699	89,605
Net cash used in operating activities	(2,378,529)	(4,761,667)	(20,748,582)
Cash flows from investing activities			
Purchase of property and equipment	-	-	(77,160)
Cash paid for intangible assets	-	-	(85,125)
Proceeds from related party advance	-	-	525,000
Repayment of related party advance	-	-	(525,000)
Net cash used in investing activities	-	-	(162,285)
Cash flows from financing activities			
Deferred financing fees paid	-	-	(45,000)
Proceeds from issuance of common stock in private placement, net	-	-	17,690,037
Proceeds from issuance of common stock to founders	-	-	5,000
Proceeds from issuance of notes payable	-	-	1,000,000
Repayment of notes payable	-	-	(1,000,000)
Proceeds from issuance of convertible notes payable	-	-	3,967,000
Proceeds from exercise of stock options	-	-	2,600

Net cash provided by financing activities	-	-	21,619,637
Net (decrease) increase in cash and cash equivalents	(2,378,529)	(4,761,667)	708,770
Cash and cash equivalents at beginning of period	3,087,299	10,395,007	-
Cash and cash equivalents at end of period	$ 708,770	$ 5,633,340	$ 708,770
Supplemental schedule of cash flows information:			
Cash paid for interest	$ -	$ -	$ 80,000
Supplemental schedule of non-cash investing and financing activities:			
Conversion of notes payable and interest to common stock	$ -	$ -	$ 4,278,518
Common shares of Laurier issued in reverse merger transaction	$ -	$ -	$ 110

See accompanying notes to condensed financial statements

ARNO THERAPEUTICS, INC.
(a development stage company)

NOTES TO CONDENSED FINANCIAL STATEMENTS
June 30, 2010
(unaudited)

NOTE 1 – DESCRIPTION OF BUSINESS

Arno Therapeutics, Inc. ("Arno" or "the Company") develops innovative drug candidates for the treatment of cancer. Arno's lead clinical drug candidate, AR-12, is a potentially first-in-class, orally available, targeted anti-cancer agent that inhibits phosphoinositide dependent protein kinase-1, or PDK-1, a protein in the PI3K/Akt pathway involved in the growth and proliferation of cells, including cancer cells. Arno believes that AR-12 may also cause cell death through the induction of stress in the endoplasmic reticulum, which is important to the growth of cells. In May 2009, the FDA accepted Arno's investigational new drug application, or IND, for AR-12. Arno is currently conducting a multi-centered Phase I clinical study of AR-12 in adult patients with advanced or recurrent solid tumors or lymphoma. The Phase I study of AR-12 is being conducted in two parts. The first part is a dose-escalating study, which Arno refers to as the Escalation Phase, primarily designed to evaluate the compound's safety in order to identify the maximum tolerated dose, or MTD, or a recommended dose, or RD, for future studies of AR-12. Arno anticipates that the Escalation Phase will be completed in the second quarter of 2011. Following the Escalation Phase, Arno plans to initiate the second part of the study, which involves enrolling an expanded cohort of up to 50 patients at the MTD or RD in multiple tumor types. Arno refers to this second part of the study as the Expansion Phase. The purpose of the Expansion Phase is to further evaluate and confirm the PD effects, potential anti-tumor activity, and safety of AR-12 at the MTD or RD. Arno anticipates concluding the Expansion Phase by the end of 2011.

Arno is also developing AR-42, an orally available, broad spectrum inhibitor of both histone and non-histone deacetylation proteins, which play an important role in the regulation of gene expression. In preclinical studies, AR-42 has demonstrated greater potency and activity in solid and liquid tumors when compared to vorinostat (also known as SAHA and marketed as Zolinza® by Merck) and other deacetylase inhibitors. These data demonstrate the potent and differentiating activity of AR-42. Additionally, pre-clinical findings presented at the 2009 American Society of Hematology Annual Meeting and Exposition showed that AR-42 potently and selectively inhibits leukemic stem cells in acute myeloid leukemia. AR-42 is currently being studied in an investigator initiated Phase I/IIa clinical study in adult patients with relapsed or refractory multiple myeloma, chronic lymphocytic leukemia, or CLL, or lymphoma. Once the MTD is defined, the study is designed so that additional patients can be added to investigate efficacy in a particular disease and help guide future Phase II programs. Up to an additional 10 patients may be enrolled at the MTD dose in each of the multiple myeloma, CLL and lymphoma.

Arno is also developing AR-67, a novel, third-generation campothecin analogue that inhibits Topoisomerase I activity. In 2008, Arno completed a multi-centered, ascending dose Phase I clinical trial of AR-67 in patients with advanced solid tumors. AR-67 is currently being studied in a Phase II clinical trial in patients with glioblastoma multiforme, or GBM, a highly aggressive form of brain cancer. Arno anticipates having interim data from this Phase II study by the second quarter of 2011. Thereafter, if data permits, Arno may elect to initiate larger Phase II studies or advance AR-67 into a registration-enabling Phase III study.

On May 1, 2009, the Company filed a request with Securities Exchange Commission (the "SEC") to deregister their unissued and unsold shares of common stock previously registered under Form S-1. On May 5, 2009, the Company filed Form 15 with the SEC to effectively terminate their registration.

NOTE 2 – BASIS OF PRESENTATION AND SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

The Company is a development stage enterprise since it has not yet generated any revenue from the sale of products and, through June 30, 2010, its efforts have been principally devoted to developing its licensed technologies and raising capital. Accordingly, the accompanying financial statements have been prepared in accordance with the provisions of Accounting Standards Codification ("ASC") 915, "Development Stage Entities". The Company has experienced net losses since its inception and has an accumulated deficit of approximately $25.7 million at June 30, 2010. The Company expects to incur substantial and increasing losses and have negative net cash flows from operating activities as it expands its technology portfolio and engages in further research and development activities, particularly the conducting of pre-clinical and clinical trials.

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The accompanying unaudited Condensed Financial Statements have been prepared in accordance with generally accepted accounting principles for interim financial information. Accordingly, they do not include all of the information and footnotes required by accounting principles generally accepted in the United States of America for complete financial statements. In the opinion of Arno's management, the accompanying Condensed Financial Statements contain all adjustments (consisting of normal recurring accruals and adjustments) necessary to present fairly the financial position, results of operations and cash flows of the Company at the dates and for the periods indicated. The interim results for the period ended June 30, 2010 are not necessarily indicative of results for the full 2010 fiscal year or any other future interim periods. Because the Merger was accounted for as a reverse acquisition under generally accepted accounting principles, the financial statements for periods prior to June 3, 2008 reflect only the operations of Old Arno.

The preparation of financial statements in conformity with generally accepted accounting principles requires that management make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenues and expenses during the reporting periods. Estimates and assumptions principally relate to services performed by third parties but not yet invoiced, estimates of the fair value and forfeiture rates of stock options issued to employees and consultants, and estimates of the probability and potential magnitude of contingent liabilities. Actual results could differ from those estimates.

NOTE 3 - LIQUIDITY AND CAPITAL RESOURCES

Cash resources as of June 30, 2010 were approximately $0.7 million, compared to $3.1 million as of December 31, 2009. Based on its resources at June 30, 2010, the financing completed in August 2010 and the current plan of expenditure for continued development of the Company's current product candidates, the Company believes that it has sufficient capital to fund its operations through the second half of 2011. However, the Company will need substantial additional financing in the future until it can achieve profitability, if ever. The Company's continued operations will depend on its ability to raise additional funds through various potential sources, such as equity and debt financing, or to license its product candidates to another pharmaceutical company. The Company will continue to fund operations from cash on hand and through sources of capital similar to those previously described. The Company cannot assure that it will be able to secure such additional financing, or if available, that it will be sufficient to meet its needs.

The success of the Company depends on its ability to discover and develop new products to the point of FDA approval and subsequent revenue generation and, accordingly, to raise enough capital to finance these developmental efforts. Management plans to raise additional equity capital or license one or more of its products to finance the continued operating and capital requirements of the Company. Amounts raised will be used to further develop the Company's product candidates, acquire additional product licenses and for other working capital purposes. While the Company will extend its best efforts to raise additional capital to fund all operations for the next 12 to 24 months, management can provide no assurances that the Company will be able to raise sufficient funds. The accompanying condensed financial statements do not include any adjustments that might result from the outcome of this uncertainty.

In addition, to the extent that the Company raises additional funds by issuing shares of its common stock or other securities convertible or exchangeable for shares of common stock, stockholders may experience additional significant

dilution. In the event the Company raises additional capital through debt financings, the Company may incur significant interest expense and become subject to covenants in the related transaction documentation that may affect the manner in which the Company conducts its business. To the extent that the Company raises additional funds through collaboration and licensing arrangements, it may be necessary to relinquish some rights to its technologies or product candidates, or grant licenses on terms that may not be favorable to the Company. These things may have a material adverse effect on the Company's business.

NOTE 4 – BASIC AND DILUTED LOSS PER SHARE

Basic loss per share is computed by dividing the loss available to common shareholders by the weighted-average number of common shares outstanding. Diluted loss per share is computed similarly to basic loss per share except that the denominator is increased to include the number of additional common shares that would have been outstanding if the potential common shares had been issued and if the additional common shares were dilutive.

For all periods presented, potentially dilutive securities are excluded from the computation of fully diluted loss per share as their effect is anti-dilutive.

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Potentially dilutive securities include:

	June 30, 2010	June 30, 2009
Warrants to purchase common stock	495,252	495,252
Options to purchase common stock	1,453,303	2,436,511
Total potentially dilutive securities	1,948,555	2,931,763

NOTE 5 - INTANGIBLE ASSETS AND INTELLECTUAL PROPERTY

AR-67 License Agreement

The Company's rights to AR-67 are governed by an October 2006 license agreement with the University of Pittsburgh ("Pitt"). Under this agreement, Pitt granted the Company an exclusive, worldwide, royalty-bearing license for the rights to commercialize technologies embodied by certain issued patents, patent applications and know-how relating to AR-67 for all therapeutic uses. The Company has expanded, and intends to continue to expand, its patent portfolio by filing additional patents covering expanded uses for this technology.

Under the license agreement with Pitt, the Company also agreed to indemnify and hold Pitt and its affiliates harmless from any and all claims, actions, demands, judgments, losses, costs, expenses, damages and liabilities (including reasonable attorneys' fees) arising out of or in connection with (i) the production, manufacture, sale, use, lease, consumption or advertisement of AR-67, (ii) the practice by the Company or any affiliate or sublicensee of the licensed patent; or (iii) any obligation of the Company under the license agreement unless any such claim is determined to have arisen out of the gross negligence, recklessness or willful misconduct of Pitt. The license agreement will terminate upon the expiration of the last patent relating to AR-67. Pitt may generally terminate the agreement at any time upon a material breach by the Company to the extent it fails to cure any such breach within 60 days after receiving notice of such breach or in the event the Company files for bankruptcy. The Company may terminate the agreement for any reason upon 90 days prior written notice.

AR-12 and AR-42 License Agreements

The Company's rights to both AR-12 and AR-42 are governed by separate license agreements with The Ohio State University Research Foundation ("Ohio State") entered into in January 2008. Pursuant to each of these agreements, Ohio State granted the Company exclusive, worldwide, royalty-bearing licenses to commercialize certain patent applications, know-how and improvements relating to AR-42 and AR-12 for all therapeutic uses.

The license agreements with Ohio State further provide that the Company will indemnify Ohio State from any and all claims arising out of the death of or injury to any person or persons or out of any damage to property, or resulting from the production, manufacture, sale, use, lease, consumption or advertisement of either AR-12 or AR-42, except to the extent that any such claim arises out of the gross negligence or willful misconduct of Ohio State. The license agreements for AR-12 and AR-42 each expire on the later of (i) the expiration of the last valid claim contained in any licensed patent and (ii) 20 years after the effective date of the license. Ohio State will generally be able to terminate either license upon the Company's breach of the terms of the license to the extent the Company fails to cure any such

breach within 90 days after receiving notice of such breach or the Company files for bankruptcy. The Company may terminate either license upon 90 days prior written notice.

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NOTE 6 – STOCKHOLDERS' EQUITY

Preferred Stock

On August 11, 2010, the Company amended and restated its certificate of incorporation, increasing the number of shares of preferred stock authorized for issuance thereunder from 10,000,000 to 35,000,000, all of which are currently designated as Series A Convertible Preferred Stock.

Common Stock

In August 2005, the Company issued an aggregate of 9,968,797 shares of common stock to its founders for $5,000.

As a condition to the closing of the Merger, on June 2, 2008, the Company completed a private placement of 7,360,689 shares of its common stock (as adjusted to give effect to the Merger), resulting in gross proceeds of approximately $17.8 million. Issuance costs related to the private placement were approximately $142,000, which were capitalized and charged to stockholders' equity upon the closing of the private placement. In accordance with the terms of the Notes, contemporaneously with the completion of the June 2, 2008 private placement, the outstanding principal and accrued interest of $4,278,518 under the Notes converted into an aggregate of 1,962,338 shares of common stock. Additionally, 1,100,200 shares of common stock that were held by the original stockholders of Laurier prior to the Merger are reflected in the Company's common stock outstanding in the accompanying financial statements.

Warrants

In connection with the in-licensing of the Company's product candidates AR-12 and AR-42, the Company issued 299,063 fully vested warrants to employees of Two River Group Holdings, LLC (see Note 10) and a consultant for their consultation and due diligence efforts as part of a finder's fee arrangement. The warrants have an exercise price of $2.42 and were valued at $480,400 based upon the Black-Scholes option-pricing model. The assumptions used under the Black-Scholes option-pricing model included a risk free interest rate of 3.27%, volatility of 80.80% and a five year life. None of these warrants have been exercised to date.

In conjunction with the conversion of $4,278,518 of convertible debt, including accrued interest, prior to the Merger, the Company issued fully vested warrants to purchase 196,189 shares of common stock to the holders of such debt. The warrants were issued with an exercise price of $2.42 and expire in June 2, 2013. The fair value of the warrants was determined to be $348,000. None of these warrants have been exercised to date.

NOTE 7 – STOCK OPTION PLAN

The Company's 2005 Stock Option Plan (the "Plan") was originally adopted by the Board of Directors of Old Arno in August 2005, and was assumed by the Company on June 3, 2008 in connection with the Merger. After giving effect to the Merger, there are 2,990,655 shares of the Company's common stock reserved for issuance under the Plan. Under the Plan, incentives may be granted to officers, employees, directors, consultants, and advisors. Incentives under the

Plan may be granted in any one or a combination of the following forms: (a) incentive stock options and non-statutory stock options, (b) stock appreciation rights, (c) stock awards, (d) restricted stock and (e) performance shares.

The Plan is administered by the Board of Directors, or a committee appointed by the Board, which determines recipients and types of awards to be granted, including the number of shares subject to the awards, the exercise price and the vesting schedule. The term of stock options granted under the Plan cannot exceed 10 years. Options shall not have an exercise price less than the fair market value of the Company's common stock on the grant date, and generally vest over a period of three to four years.

As of June 30, 2010, an aggregate of 1,517,352 shares remained available for future grants and awards under the Plan, which covers stock options, warrants and restricted awards. The Company issues unissued shares to satisfy stock options, warrants exercises and restricted stock awards.

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A summary of the status of the options issued under the Plan as of June 30, 2010, and information with respect to the changes in options outstanding is a follows:

	Shares Available for Grant	Options Outstanding Outstanding Stock Options	Weighted-Average Exercise Price	Aggregate Intrinsic Value
Balance at January 1, 2010	1,057,414	1,913,241	$ 1.76	
Options granted under the Plan	-	-	$ -	
Options exercised	-	-	$ -	
Options forfeited	459,938	(459,938)	$ 2.72	
Balance at June 30, 2010	1,517,352	1,453,303	$ 1.47	$ -
Exercisable at June 30, 2010		1,154,326	$ 1.53	$ -

The Company estimates the fair value of each option award granted using the Black-Scholes option-pricing model. For the three months ended March 31, 2010 and 2009, there were no options granted.

As allowed by ASC 718 for companies with a short period of publicly traded stock history, management's estimate of expected volatility is based on the average expected volatilities of a sampling of five companies with similar attributes to the Company, including: industry, stage of life cycle, size and financial leverage.

The Company has no historical basis for determining expected forfeitures and, as such, compensation expense for stock-based awards does not include an estimate for forfeitures.

Stock-based compensation for the six months ended June 30, 2010 and 2009 and for the period from August 1, 2005 (inception) through June 30, 2010, are as follows:

	Three months ended June 30,		Six months ended June 30,		Period from August 1, 2005 (inception) through June 30, 2010
	2010	2009	2010	2009	
General and administrative	$ 6,300	$ (190,179)	$ (3,000)	$ (22,595)	$ 973,592
Research and development	34,800	80,878	69,300	186,294	969,377
Total	$ 41,100	$ (109,301)	$ 66,300	$ 163,699	$ 1,942,969

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As of June 30, 2010, the total outstanding, and the total exercisable, options under the Plan were as follows:

Outstanding				Exercisable	
Range of Exercise Prices	Shares	Weighted-Average Remaining Contractual Life (in years)	Weighted-Average Exercise Price	Total Shares	Weighted-Average Exercise Price
$ 0.13	129,532	1.4	$ 0.13	129532	$ 0.13
1.00	803,443	6.5	1.00	546034	1.00
2.42	428,661	7.1	2.42	397093	2.42
3.00	91,667	3.8	3.00	81667	3.00
	1,453,303	6.1	$ 1.47	1,154,326	$ 1.53

As of June 30, 2010 and 2009, there was $261,431 and $752,982, respectively, of unrecognized compensation costs related to non-vested stock options. These costs are expected to be recognized over a period of approximately 2.3 years as from June 30, 2010.

NOTE 8 – RELATED PARTIES

On June 1, 2009, the Company entered into a services agreement with Two River Consulting, LLC (“TRC”) to provide various clinical development, operational, managerial, accounting and financial, and administrative services to the Company for a period of one year. David M. Tanen, the Company’s President, Secretary and director, Arie S. Belldegrun, the Chairman of the Board of Directors, and Joshua A. Kazam, director are each partners of TRC. The terms of the Services Agreement were reviewed and approved by a special committee of the Company’s Board of Directors consisting of independent directors. None of the members of the special committee has any interest in TRC or the services agreement. As compensation for the services contemplated by the services agreement, the Company pays TRC a monthly cash fee of $55,000. The monthly fee increased from $50,000 to $55,000 effective February 2010, when the scope of services provided by TRC increased to include accounting and financial services upon the resignation of the Company’s Chief Financial Officer.

On occasion, some of the Company’s expenses are paid by TRC. No interest is charged by TRC on any outstanding balance owed by the Company. For the six months ended June 30, 2010 and 2009, and for the period from August 1, 2005 (inception) through June 30, 2010, total cash services and reimbursed expenses totaled $379,074, $50,000 and $784,074, respectively. As of June 30, 2010 and 2009, the Company had a payable to TRC of $89,605 and $50,000, respectively.

Prior to June 1, 2009, some of the Company’s expenses were paid by Two River Group Holdings, LLC (“Two River”), a company that is controlled by four of the Company’s directors and founders. No interest is charged by Two River on any outstanding balance owed by the Company. For the six months ended June 30, 2010 and 2009 for the period from August 1, 2005 (inception) through June 30, 2010, reimbursable expenses totaled $0, $10,904 and $304,550, respectively. In addition, the Company granted fully vested warrants to purchase 299,063 shares of its common stock

at an exercise price of $2.42 to the Two River employees who provided consultation and due diligence efforts related to the in-licensing of AR-12 and AR-42. The warrants have a five year life and a fair value of $480,400 based upon the Black-Scholes option-pricing model. As of June 30, 2010 and 2009, the Company had $0 and $6,315 payable to Two River, respectively.

The Company utilized the services of Riverbank Capital Securities, Inc. (“Riverbank”), a FINRA member broker dealer registered with the SEC, for investment banking and other investment advisory services in connection with the June 2008 private placement and the Notes. Riverbank is an entity controlled by several partners of Two River who are also officers and/or directors of the Company. The Company paid a $100,000 non-accountable expense allowance to Riverbank for services related to the June 2008 private placement and is not obligated to Riverbank for any future payments.

The financial condition and results of operations of the Company, as reported, are not necessarily indicative of results that would have been reported had the Company operated completely independently.

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NOTE 9 – COMMITMENTS AND CONTINGENCIES

On October 20, 2008, the Company entered into a lease for office space consisting of 5,390 square feet in Parsippany, New Jersey. The lease commencement date was November 14, 2008, with lease payments beginning on January 1, 2009. The lease expiration date is five years from the rent commencement date. The total five year lease obligation is approximately $713,000.

The Company provided a security deposit of $44,018, or four months base rent, in the form of a letter of credit. The letter of credit may be reduced by $11,005 on January 1, 2011 and by an additional $11,005 on January 1, 2013, provided the Company maintains certain conditions described in the lease agreement. The Company has an early termination option, which provides the Company may terminate the lease on the third anniversary, upon providing the landlord nine months written notice prior to the third anniversary of the lease. If the Company exercises its termination option, the Company would be obligated to pay a fee of $53,641 which consists of unamortized costs and expenses incurred by the landlord in connection with the lease. The Company also has an option to extend the term of the lease for a period of five additional years, provided the Company gives notice to the landlord no later than 12 months prior to the expiration of the original term.

The Company has entered into various contracts with third parties in connection with the development of the licensed technology described in Note 4.

The aggregate minimum commitment under these contracts as of June 30, 2010 is approximately $900,000.

In the normal course of business, the Company enters into contracts that contain a variety of indemnifications with its employees, licensors, suppliers and service providers. Further, the Company indemnifies its directors and officers who are, or were, serving at the Company’s request in such capacities. The Company’s maximum exposure under these arrangements is unknown as of June 30, 2010. The Company does not anticipate recognizing any significant losses relating to these arrangements.

NOTE 10 – SUBSEQUENT EVENTS

On September 3, 2010, the Company entered into a Securities Purchase and Registration Rights Agreement, or the Purchase Agreement, with a number of institutional and accredited investors pursuant to which the Company sold in a private placement an aggregate of 15,274,000 shares of newly-designated Series A Convertible Preferred Stock, par value $0.0001 per share, or Series A Preferred Stock, at a per share purchase price of $1.00. In accordance with the Purchase Agreement, the Company also issued 1,221,920, two-and-one-half-year Class A warrants to purchase additional shares of Series A Preferred Stock at an initial exercise price of $1.00 per share and 6,415,080, five-year Class B warrants at an initial exercise price of $1.15 per share. The terms of the Class A and Class B warrants contain an anti-dilutive price adjustment provision, such that in the event the Company issues common shares at a price below the current exercise price of the Warrants the exercise price of the Class A and Class B warrants will be adjusted based on the lower issuance price. The sale of the shares and warrants resulted in aggregate gross proceeds of approximately $15.2 million, before expenses.

Issuance costs related to the financing were approximately $1.9 million, including the issuance of warrants ("Placement Warrants") to purchase 1,050,840 shares of the Company's common stock at 110% of the Series A Preferred Stock purchase price per share to designees of Riverbank and I-Bankers Securities, Inc. ("IBS"), that acted as placement agents for the Company in connection with the private placement. The Placement Warrants were valued at $614,400 using the Black-Scholes option-pricing model.

Pursuant to the Registration Rights Agreement, the Company will use its best efforts to cause such registration statement to be declared effective within 180 days following the initial closing under the Purchase Agreement, or by March 8, 2011. If such registration statement is not declared effective by the SEC by such date, the Company will pay liquidated damages to the investors in the amount of 1% of each investor's aggregate investment amount for each 30-day period until the registration statement is declared effective.

On October 29, 2010, the Company was awarded funding of a total of approximately $733,000 under the IRS Qualifying Therapeutic Discovery Project ("QTDP") program, which was created as part of the Patient Protection and Affordable Care Act of 2010 (the "Healthcare Reform Act"). As enacted under the Healthcare Reform Act, the QTDP program provides a tax credit or grant of up to 50% of eligible costs and expenses for the tax years of 2009 and 2010 for qualifying research and development expenses incurred for innovative projects that are determined by the U.S. Department of Health and Human Services to have reasonable potential to result in a new therapy, reduce health care costs, or represent a significant advance in finding a cure for human disease. The Company was awarded approximately $244,000 for R&D expenses incurred for each of its AR-12, AR-42 and AR-67 development programs. The Company expects to receive the amounts granted prior to the end of 2010.

26,815,831 Shares

Common Stock

PROSPECTUS

, 2010

PART II

INFORMATION NOT REQUIRED IN PROSPECTUS

ITEM 13. OTHER EXPENSES OF ISSUANCE AND DISTRIBUTION.

The following table sets forth all costs and expenses, other than underwriting discounts and commissions, payable by us in connection with the sale of the common stock being registered. All amounts shown are estimates except for the SEC registration fee.

		Amount to be Paid
SEC registration fee	$	975
Legal fees and expenses		50,000
Accounting fees and expenses		20,000
Printing and engraving and miscellaneous expenses		5,000
Total	$	75,975

ITEM 14. INDEMNIFICATION OF DIRECTORS AND OFFICERS.

Section 145 of the General Corporation Law of the State of Delaware provides as follows:

A corporation may indemnify any person who was or is a party or is threatened to be made a party to any threatened, pending or completed action, suit or proceeding, whether civil, criminal, administrative or investigative (other than an action by or in the right of the corporation) by reason of the fact that the person is or was a director, officer, employee or agent of the corporation, or is or was serving at the request of the corporation as a director, officer, employee or agent or another corporation, partnership, joint venture, trust or other enterprise, against expenses (including attorneys' fees), judgments, fines and amounts paid in settlement actually and reasonably incurred by the person in connection with such action, suit or proceeding if the person acted in good faith and in a manner the person reasonably believed to be in or not opposed to the best interest of the corporation, and, with respect to any criminal action or proceeding, had no reasonable cause to believe the person's conduct was unlawful. The termination of any action, suit or proceeding by judgment, order, settlement, conviction or upon a plea of nolo contendere or its equivalent, shall not, of itself, create a presumption that the person did not act in good faith and in a manner which he reasonably believed to be in or not opposed to the best interests of the corporation, and, with respect to any criminal action or proceeding, had reasonable cause to believe that the person's conduct was unlawful.

A corporation may indemnify any person who was or is a party or is threatened to be made a party to any threatened, pending or completed action or suit by or in the right of the corporation to procure a judgment in its favor by reason of the fact that he is or was a director, officer, employee or agent of the corporation, or is or was serving at the request of the corporation as a director, officer, employee or agent of another corporation, partnership, joint venture, trust or other enterprise against expenses (including attorneys' fees) actually and reasonably incurred by the person in connection with the defense or settlement of such action or suit if the person acted in good faith and in a manner the person reasonably believed to be in or not opposed to the best interests of the corporation and except that no indemnification will be made in respect to any claim, issue or matter as to which such person shall have been adjudged to be liable to the corporation unless and only to the extent that the Court of Chancery or the court in which such action or suit was brought shall determine upon application that, despite the adjudication of liability but in view of all the circumstances of the case, such person is fairly and reasonably entitled to indemnity for such expenses which the Court of Chancery or such other court shall deem proper.

Our certificate of incorporation provides that we will indemnify any person, including persons who are not our directors and officers, to the fullest extent permitted by Section 145 of the Delaware General Corporation Law.

In addition, pursuant to our bylaws, we will indemnify our directors and officers against expenses (including judgments or amounts paid in settlement) incurred in any action, civil or criminal, to which any such person is a party by reason of any alleged act or failure to act in his capacity as such, except as to a matter as to which such director or officer shall have been finally adjudged to be liable for negligence or misconduct in the performance of his duty to the corporation or not to have acted in good faith in the reasonable belief that his action was in the best interest of the corporation.

Reference is made to Item 17 for our undertakings with respect to indemnification for liabilities arising under the Securities Act of 1933.

ITEM 15. RECENT SALES OF UNREGISTERED SECURITIES.

The following summarizes all sales of unregistered securities by the Registrant within the last three years. In June 2008, Arno was acquired by Laurier International, Inc., a Delaware corporation, in a "reverse" merger whereby a wholly-owned subsidiary of Laurier merged with and into Arno Therapeutics, with Arno Therapeutics remaining as the surviving corporation and a wholly-owned subsidiary of Laurier. In accordance with the terms of this merger, stockholders of Arno Therapeutics exchanged all of their shares of common stock of Arno Therapeutics for shares of Laurier common stock at a rate of 1.99377 shares of Laurier common stock for each share of Arno Therapeutics common stock.

On February 9, 2007, Arno issued to certain accredited investors 6% Convertible Promissory Notes in the aggregate principal amount of approximately $4,000,000 (the "6% Notes"). The 6% Notes provided that upon the next closing of any equity financing in excess of $5,000,000 (a "Qualified Financing"), the 6% Notes would automatically convert into the same securities issued by Arno in the Qualified Financing ("Conversion Shares"), in an amount determined by dividing the principal amount of the 6% Notes, and all accrued interest thereon, by 90% of the price per share sold in the Qualified Financing (the "Offering Price"). In addition, upon conversion, Arno agreed to issue to the holders of the 6% Notes five-year warrants ("Conversion Warrants"), to purchase a number of shares of Arno common stock equal to 10% of the Conversion Shares at an exercise price equal to the Offering Price.

As a condition and immediately prior to the closing of the merger, on June 2, 2008, Arno completed a private placement of its equity securities whereby it received gross proceeds of approximately $17,732,000 through the sale of approximately 3,691,900 shares of its common stock to accredited investors, which shares were exchanged for 7,360,689 shares of Company common stock after giving effect to the merger. Contemporaneously with the June 2008 private placement, the 6% Notes converted into 984,246 Conversion Shares and the holders of the 6% Notes received Conversion Warrants to purchase an aggregate of 98,409 shares of Arno common stock at an exercise price equal to $4.83 per share. The Conversion Shares were exchanged for an aggregate of approximately 1,962,338 shares of our common stock and the Conversion Warrants were exchanged for an aggregate of approximately 196,189 five year warrants to purchase our common stock at an exercise price equal to $2.42 per share.

On September 9, 2010, we entered into a Securities Purchase and Registration Rights Agreement (the "Purchase Agreement") with a number of accredited investors pursuant to which we agreed to sell in a private placement up to an aggregate of 20,000,000 shares of our newly-designated Series A Convertible Preferred Stock, par value $0.0001 per share (the "Series A Preferred Stock"), at a per share purchase price of $1.00. Pursuant to the terms of the Purchase Agreement, we also agreed to issue to each investor a two-and-one-half-year warrant to purchase a number of additional shares of Series A Preferred Stock equal to 8% of the number of shares purchased by such investor at an initial exercise price of $1.00 per share (collectively, the "Class A Warrants"), and a five-year warrant to purchase a number of additional shares of Series A Preferred Stock equal to 42% of the number of shares purchased by such investor at an initial exercise price of $1.15 per share (collectively, the "Class B Warrants"). Between September 9, 2010 and September 13, 2010, we completed a number of closings under the Purchase Agreement, issuing an aggregate of 15,274,000 shares of Series A Preferred Stock, Class A Warrants to purchase 1,221,920 shares of Series A Preferred Stock, and Class B Warrants to purchase 6,415,080 shares of Series A Preferred Stock. The sale of the shares and warrants resulted in aggregate gross proceeds of approximately $15.2 million, before deducting expenses. In connection with the private placement, we issued five-year warrants to purchase an aggregate of 1,050,840 shares of Series A Preferred Stock at an initial exercise price of $1.10 per share to the placement agents and their designees.

Except as noted above, the sales of the securities identified above were made pursuant to privately negotiated transactions that did not involve a public offering of securities and, accordingly, we believe that these transactions were exempt from the registration requirements of the Securities Act pursuant to Section 4(2) thereof and rules

promulgated thereunder. Each of the above-referenced investors in our stock represented to us in connection with their investment that they were "accredited investors" (as defined by Rule 501 under the Securities Act) and were acquiring the shares for investment and not distribution, that they could bear the risks of the investment and could hold the securities for an indefinite period of time. The investors received written disclosures that the securities had not been registered under the Securities Act and that any resale must be made pursuant to a registration or an available exemption from such registration. All of the foregoing securities are deemed restricted securities for purposes of the Securities Act.

ITEM 16. EXHIBITS AND FINANCIAL STATEMENT SCHEDULES.

(a) Exhibits.

Exhibit Number	Description of Document
2.1	Agreement and Plan of Merger dated March 5, 2008, by and among Laurier International, Inc., Laurier Acquisition, Inc. and Arno Therapeutics, Inc. (incorporated by reference to Exhibit 2.1 of the Registrant's Form 8-K filed on March 6, 2008).
2.2	Amendment No. 1 dated May 12, 2008 to Agreement and Plan of Merger by and among Laurier International, Inc., Laurier Acquisition, Inc. and Arno Therapeutics, Inc. (incorporated by reference to Exhibit 2.2 of the Registrant's Registration Statement on Form S-1 filed July 31, 2008, SEC File No. 333-152660).
2.3	Amendment No. 2 dated May 30, 2008 to Agreement and Plan of Merger by and among Laurier International, Inc., Laurier Acquisition, Inc. and Arno Therapeutics, Inc. (incorporated by reference to Exhibit 2.3 of the Registrant's Registration Statement on Form S-1 filed July 31, 2008, SEC File No. 333-152660).
3.1	Amended & Restated Certificate of Incorporation of Arno Therapeutics, Inc.
3.2	Bylaws of the Registrant (incorporated by reference to Exhibit 3.2 of the Registrant's Registration Statement on Form SB-2 filed on October 2, 2002, SEC File No. 333-100259).
3.3	Certificate of Designation of Series A Convertible Preferred Stock.
4.1	Specimen Common Stock Certificate (incorporated by reference to Exhibit 4.1 of the Registrant's Form 8-K filed June 9, 2008).
4.2	Form of Common Stock Purchase Warrant issued to former note holders of Arno Therapeutics, Inc. (incorporated by reference to Exhibit 4.2 of the Registrant's Form 8-K filed June 9, 2008).
4.3	Form of Class A Warrant issued to investors in September 2010 private placement.
4.4	Form of Class B Warrant issued to investors in September 2010 private placement.
4.5	Form of Placement Agent Warrant issued in September 2010 private placement.
5.1	Opinion of Fredrikson & Byron, P.A.
10.1	Employment Agreement dated June 1, 2007 between Arno Therapeutics, Inc. and Scott Z. Fields, M.D. (incorporated by reference to Exhibit 10.1 of the Registrant's Form 8-K filed June 9, 2008).
10.2	Letter agreement dated September 2, 2007 between Arno Therapeutics, Inc. and J. Chris Houchins (incorporated by reference to Exhibit 10.2 of the Registrant's Form 8-K filed June 9, 2008).
10.3	Arno Therapeutics, Inc. 2005 Stock Option Plan (incorporated by reference to Exhibit 10.3 of the Registrant's Form 8-K filed June 9, 2008).
10.4	Form of stock option agreement for use under Arno Therapeutics, Inc. 2005 Stock Option Plan (incorporated by reference to Exhibit 10.4 of the Registrant's Form 8-K filed June 9, 2008).
10.5	License Agreement dated October 25, 2006 between Arno Therapeutics, Inc. and The University of Pittsburgh (incorporated by reference to Exhibit 10.5 of the Registrant's Form 8-K filed June 9, 2008).+
10.6	License Agreement dated January 3, 2008 between Arno Therapeutics, Inc. and The Ohio State University Research Foundation (incorporated by reference to Exhibit 10.6 of the Registrant's Form 8-K filed June 9, 2008).+
10.7	License Agreement dated January 9, 2008 between Arno Therapeutics, Inc. and The Ohio State University Research Foundation (incorporated by reference to Exhibit 10.7 of the Registrant's Form 8-K filed June 9, 2008).+
10.8	Form of Subscription Agreement between Arno Therapeutics, Inc. and the investors in the June 2, 2008 private placement (incorporated by reference to Exhibit 10.8 of the Registrant's Form 8-K filed June 9, 2008).
10.9	

	Employment Agreement dated June 9, 2008 between Arno Therapeutics, Inc. and Brian Lenz, as amended on July 9, 2008 (incorporated by reference to Exhibit 10.1 of the Registrant's Form 10-Q for the quarter ended June 30, 2008).
10.10	Employment Agreement by and between Arno Therapeutics, Inc. and Dr. Roger Berlin, dated September 3, 2008 (incorporated by reference to Exhibit 10.1 of the Registrant's Form 8-K filed September 3, 2008).
10.11	Services Agreement dated June 1, 2009, between Arno Therapeutics, Inc. and Two River Consulting, LLC.
10.12	Lease Agreement by and between Arno Therapeutics, Inc. and Maple 4 Campus L.L.C., dated October 17, 2008 (incorporated by reference to Exhibit 10.11 of the Registrant's Form 10-K for the fiscal year ended December 31, 2008).
10.13	Form of Securities Purchase and Registration Rights Agreement dated September 3, 2010 among Arno Therapeutics, Inc. and the purchasers identified therein.
10.14	First Amendment to Services Agreement dated September 9, 2010, between Arno Therapeutics, Inc. and Two River Consulting, LLC.
23.1	Consent of Crowe Horwath LLP
23.2	Consent of Hays & Company LLP.
23.3	Consent of Fredrikson & Byron, P.A. (included in Exhibit 5.1).
24.1	Power of Attorney (included on signature page hereof).

+Confidential treatment has been granted as to certain omitted portions of this exhibit pursuant to Rule 406 of the Securities Act or Rule 24b-2 of the Exchange Act.

(b) Financial Statement Schedules.

The Financial Statement Schedules have been omitted either because they are not required or because the information has been included in the financial statements or the notes thereto included in this Registration Statement.

ITEM 17. UNDERTAKINGS.

The undersigned registrant hereby undertakes:

(1) To file, during any period in which offers or sales are being made, a post-effective amendment to this registration statement:

(i) To include any prospectus required by section 10(a)(3) of the Securities Act;

(ii) To reflect in the prospectus any facts or events arising after the effective date of the registration statement (or the most recent post-effective amendment thereof) which, individually or in the aggregate, represent a fundamental change in the information set forth in the registration statement. Notwithstanding the foregoing, any increase or decrease in volume of securities offered (if the total dollar value of securities offered would not exceed that which was registered) and any deviation from the low or high end of the estimated maximum offering range may be reflected in the form of prospectus filed with the Commission pursuant to Rule 424(b) if, in the aggregate, the changes in volume and price represent no more than a 20% change in the maximum aggregate offering price set forth in the "Calculation of Registration Fee" table in the effective registration statement; and

(iii) To include any material information with respect to the plan of distribution not previously disclosed in the registration statement or any material change to such information in the registration statement.

(2) That, for the purpose of determining any liability under the Securities Act, each post-effective amendment shall be deemed to be a new registration statement relating to the securities offered therein, and the offering of the securities at that time shall be deemed to be the initial bona fide offering thereof.

(3) To remove from registration by means of a post-effective amendment any of the securities being registered that remain unsold at the termination of this offering.

(4) That, for the purpose of determining liability under the Securities Act to any purchaser, if the registrant is subject to Rule 430C, each prospectus filed pursuant to Rule 424(b) as part of a registration statement relating to an offering, other than registration statements relying on Rule 430B or other than prospectuses filed in reliance on Rule 430A, shall be deemed to be part of and included in the registration statement as of the date it is first used after effectiveness. Provided, however, that no statement made in a registration statement or prospectus that is part of the registration statement or made in a document incorporated or deemed incorporated by reference into the registration statement or prospectus that is part of the registration statement will, as to a purchaser with a time of contract of sale prior to such first use, supersede or modify any statement that was made in the registration statement or prospectus that was part of the registration statement or made in any such document immediately prior to such date of first use.

Insofar as indemnification for liabilities arising under the Securities Act may be permitted to directors, officers and controlling persons of the registrant pursuant to the foregoing provisions, or otherwise, the registrant has been advised that in the opinion of the SEC this form of indemnification is against public policy as expressed in the Securities Act and is, therefore, unenforceable. In the event that a claim for indemnification against these liabilities (other than the payment by the registrant of expenses incurred or paid by a director, officer or controlling person of the registrant in the successful defense of any action, suit or proceeding) is asserted by a director, officer or controlling person in connection with the securities being registered, the registrant will, unless in the opinion of its counsel the matter has been settled by controlling precedent, submit to a court of appropriate jurisdiction the question whether such indemnification by it is against public policy as expressed in the Securities Act and will be governed by the final adjudication of this issue.

SIGNATURES

Pursuant to the requirements of the Securities Act of 1933, the Registrant has duly caused this Registration Statement to be signed on its behalf by the undersigned, thereunto duly authorized, in the City of Parsippany, State of New Jersey, on November 8, 2010.

ARNO THERAPEUTICS, INC.

By: /s/ David M. Tanen
David M. Tanen, President

KNOW ALL MEN BY THESE PRESENTS, that each person whose signature appears below hereby constitutes and appoints David M. Tanen and Scott L. Navins, and each of them, his true and lawful attorneys-in-fact and agent, with full power of substitution and resubstitution, for him in his name, place and stead, in any and all capacities, to sign any or all amendments to this registration statement and additional registration statements relating to the same offering, and to file the same, with all exhibits thereto, and all other documents in connection therewith, with the Securities and Exchange Commission, granting unto said attorney-in-fact and agent full power and authority to do and perform each and every act and thing requisite and necessary to be done in and about the premises, as fully to all intents and purposes as he might or could do in person, hereby ratifying and confirming all that said attorney-in-fact and agent, or his substitutes, may lawfully do or cause to be done by virtue thereof.

Pursuant to the requirements of the Securities Act of 1933, this Registration Statement has been signed by the following persons in the capacities and on the dates indicated.

Signature	Title	Date
/s/ David M. Tanen David M. Tanen	President, Secretary, and Director (Principal Executive Officer)	November 8, 2010
/s/ Scott L. Navins Scott L. Navins	Treasurer (Principal Financial and Accounting Officer)	November 8, 2010
/s/ Arie S. Belldegrun Arie S. Belldegrun, M.D.	Chairman of the Board	November 8, 2010
/s/ William F. Hamilton William F. Hamilton, Ph.D.	Director	November 8, 2010
Tomer Kariv	Director	November 8, 2010
/s/ Peter M. Kash Peter M. Kash	Director	November 8, 2010
/s/ Yacov Reizman Yacov Reizman	Director	November 8, 2010
/s/ Steven B. Ruchefsky Steven B. Ruchefsky	Director	November 8, 2010

INDEX TO EXHIBITS FILED WITH THIS REGISTRATION STATEMENT

Exhibit Number	Description of Document
3.1	Amended & Restated Certificate of Incorporation of Arno Therapeutics, Inc.
3.3	Certificate of Designation of Series A Convertible Preferred Stock.
4.3	Form of Class A Warrant issued to investors in September 2010 private placement.
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