

AEOLUS PHARMACEUTICALS, INC.
Form 10-K
December 20, 2016

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

FORM 10-K

(MARK ONE)

✓ ANNUAL REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934
For the fiscal year ended September 30, 2016

OR
TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF
1934

For the transition period from _____ to _____
Commission File Number 0-50481

AEOLUS PHARMACEUTICALS, INC.
(Exact name of registrant as specified in its charter)

Delaware 56-1953785
(State or other jurisdiction of (I.R.S. Employer
incorporation or organization) Identification No.)

26361 Crown Valley Parkway, Suite 150 92691
Mission Viejo, California (Zip Code)
(Address of principal executive offices)

Registrant's telephone number, including area code:
949-481-9825

Securities registered pursuant to Section 12(b) of the Act:
None

Securities registered pursuant to Section 12(g) of the Act:
Common Stock, \$.01 par value per share
(Title of class)

Indicate by check mark if the registrant is a well-known seasoned issuer, as defined in Rule 405 of the Securities Act.
Yes No ✓

Indicate by check mark if the registrant is not required to file reports pursuant to Section 13 or Section 15(d) of the
Act. Yes No ✓

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the
Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was
required to file such reports), and (2) has been subject to such filing requirements for the past 90 days. Yes ✓ No

Indicate by check mark whether the registrant has submitted electronically and posted on its corporate Web site, if
any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T
(§ 232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required

to submit and post such files). Yes No

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Indicate by check mark if disclosure of delinquent filers pursuant to Item 405 of Regulation S-K (§ 229.405 of this chapter) is not contained herein, and will not be contained, to the best of registrant's knowledge, in definitive proxy or information statements incorporated by reference in Part III of this Form 10-K or any amendment to this Form 10-K. 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 Smaller reporting company

(Do not check if a smaller reporting company)

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). Yes

No

The aggregate market value of the voting common stock held by non-affiliates of the registrant based upon the average of the bid and asked price on the OTC Bulletin Board as of March 31, 2016, the last business day of the registrant's most recently completed second fiscal quarter, was approximately \$10,925,560. Shares of common stock held by each executive officer and director and by each other stockholder who owned 10% or more of the outstanding common stock as of such date have been excluded in that such stockholder might be deemed to be an affiliate of the registrant. This determination of affiliate status might not be conclusive for other purposes.

As of December 20, 2016, the registrant had 152,085,825 outstanding shares of common stock.

DOCUMENTS INCORPORATED BY REFERENCE

Portions of the registrant's definitive Information Statement to be filed pursuant to Regulation 14C in connection with the registrant's Written Consent in Lieu of the 2016 Annual Meeting of Stockholders (the "Information Statement") are incorporated herein by reference into Part III hereof.

AEOLUS PHARMACEUTICALS, INC.
 ANNUAL REPORT ON FORM 10-K
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PART I

NOTE REGARDING FORWARD-LOOKING STATEMENTS

This Annual Report on Form 10-K contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended, that relate to future events or our future financial performance. You can identify forward-looking statements by terminology such as "may," "might," "will," "could," "should," "would," "expect," "plan," "anticipate," "believe," "estimate," "predict," "intend," "potential" or "continue" or the negative of these terms or other comparable terminology. Our actual results might differ materially from any forward-looking statement due to various risks, uncertainties and contingencies, including but not limited to those identified in Item 1A entitled "Risk Factors" beginning on page 42 of this report, as well as those discussed in our other filings with the Securities and Exchange Commission (the "SEC") and the following:

- our need for, and our ability to obtain, additional funds;
- our ability to obtain grants to develop our drug candidates;
- uncertainties relating to non-clinical studies, clinical trials and regulatory reviews and approvals;
- uncertainties relating to our pre-clinical studies and trials and regulatory reviews and approvals;
- uncertainties regarding whether our compounds could inhibit formation of fibrosis in the lungs.
- uncertainties concerning whether we can position our compounds for a pre-Emergency Use Authorization application
- or we can obtain procurements from the Biomedical Advanced Research and Development Authority following any such application;
- our dependence on a limited number of therapeutic compounds;
- the early stage of the drug candidates we are developing;
- the acceptance of any future products by physicians and patients;
- competition with and dependence on collaborative partners;
- loss of key consultants, management or scientific personnel;
- our ability to obtain adequate intellectual property protection and to enforce these rights; and
- our ability to avoid infringement of the intellectual property rights of others.

Although we believe that the expectations reflected in the forward-looking statements are reasonable, we cannot guarantee future results, levels of activity, performance or achievements. We disclaim any intention or obligation to update or revise any forward-looking statements, whether as a result of new information, future events or otherwise. Item 1. Business.

General

Overview

Aeolus Pharmaceuticals, Inc. ("we," "us" or "Aeolus") is a Southern California-based biopharmaceutical company leveraging significant U.S. Government funding to develop a platform of novel compounds for use in biodefense, fibrosis, oncology, infectious disease and diseases of the central nervous system. The platform consists of approximately 180 compounds licensed from the University of Colorado ("UC"), Duke University ("Duke") and National Jewish Health ("NJH").

Our lead compound, AEOL 10150 ("10150"), is being developed under contract with the Biomedical Advanced Research and Development Authority ("BARDA" and the "BARDA Contract"), a division of the U.S. Department of Health and Human Services ("HHS"), as a medical countermeasure ("MCM") against the pulmonary sub-syndrome of acute radiation syndrome ("Pulmonary Acute Radiation Syndrome" or "Lung-ARS") and the delayed effects of acute radiation exposure ("DEARE"). Lung-ARS is caused by acute exposure to high levels of radiation due to a nuclear detonation or radiological event. On January 23, 2014, we announced that 10150 was granted Orphan Drug Designation by the U.S Food and Drug Administration ("FDA") for the Lung-ARS indication.

We are also developing 10150 for the treatment of lung fibrosis, including idiopathic pulmonary fibrosis ("IPF") and other fibrotic diseases. This new development program was created based upon lung protection data generated from animal studies in Lung-ARS and chemical gas lung exposure under the BARDA Contract and National Institutes of Health ("NIH") grants, as well as a company-funded study in a bleomycin lung fibrosis model. On March 17, 2015, we announced that 10150 was granted Orphan Drug Designation by the U.S Food and Drug Administration ("FDA") for the IPF indication. The Company plans to initiate a Phase I safety study in patients with IPF in 2017. After we have completed safety studies, we plan to initiate efficacy studies in patients with fibrosis. AEOL 10150 has previously been tested in two Phase I human clinical trials with no drug-related serious adverse events reported.

We are also developing 10150 for use in combination with radiation therapy for cancer as a treatment to reduce side effects caused by radiation toxicity and improve local tumor control. Pre-clinical studies at Duke, the University of Maryland and Loma Linda University have demonstrated that 10150 protects healthy, normal tissue, while not interfering with the benefit of radiation therapy or chemotherapy in prostate and lung cancer. Additional studies have demonstrated that 10150 enhances the anti-tumor activity of chemotherapy and radiation. A significant portion of the development work funded by the BARDA contract is applicable to the development program for radiation oncology, including safety, toxicology, pharmacokinetics and Chemistry, Manufacturing and Controls ("CMC"). After we have completed safety studies, we plan to initiate studies to demonstrate efficacy in protecting against the toxic side effects related to radiation therapy.

We are also developing 10150 as a MCM for exposure to chemical vesicants (e.g., chlorine gas, sulfur mustard gas and phosgene gas) and nerve agents (e.g., sarin gas and soman gas) with grant money from the NIH Countermeasures Against Chemical Threats ("NIH-CounterACT") program. 10150 has consistently demonstrated safety and efficacy in animal studies of chemical gas exposure and nerve gas exposure.

The Company is developing a second compound, AEOL 11114B ("11114"), as a treatment for Parkinson's disease. Research funded by the Michael J Fox Foundation for Parkinson's disease ("MJFF") demonstrated the neuro-protective activity of 11114 in mouse and rat models of Parkinson's disease. The compounds were invented by Brian J. Day, PhD at NJH and Manisha Patel, PhD at UC in collaboration with the Company. We have obtained worldwide, exclusive licenses to develop the compounds from NJH and the UC. Optimization of the manufacturing and formulation of the compound were finished in 2016, and we plan to complete the remaining work to file an Investigational New Drug ("IND") application with the FDA during 2017.

In April 2015, we announced the discovery of a new compound, AEOL 20415 ("20415"), which has demonstrated anti-inflammatory and anti-infective properties, and could be effective in treating cystic fibrosis and combatting anti-biotic resistant bacteria. The compound was developed under collaboration between Brian J. Day, PhD at NJH and Aeolus Pharmaceuticals. We have obtained a worldwide, exclusive license to develop the rights to the compound from NJH. Animal efficacy studies are underway to confirm the efficacy seen in in-vitro testing against anti-biotic resistant bacteria, and, assuming confirmation, we plan to initiate the work to file an IND application with the FDA during the first half of 2017.

Finally, we have a pipeline of approximately 180 additional compounds. We expect that the development of additional compounds in our portfolio is dependent on our finding non-dilutive capital sources to fund such pipeline opportunities.

We were incorporated in the State of Delaware in 1994. Our common stock trades on the OTCQB under the symbol "AOLS." Our principal executive offices are located at 26361 Crown Valley Parkway, Suite 150 Mission Viejo, California 92691, and our phone number at that address is (949) 481-9825. Our website address is www.aolsrx.com. However, the information on, or that can be accessed through our website is not part of this report. We also make available, free of charge through our website, our most recent annual report on Form 10-K, quarterly reports on Form 10-Q, current reports on Form 8-K, and any amendments to those reports, as soon as reasonably practicable after such material is electronically filed with or furnished to the SEC.

Strategy

Our strategy is to pursue the development of our promising platform of anti-fibrotic, anti-inflammatory, anti-infective and anti-oxidant compounds that address important unmet medical indications of clinical and national strategic importance. Our objective is to use non-dilutive capital whenever possible.

To date, we, and/or our research collaborators, have been awarded more than \$149 million in non-dilutive U.S. government funding in the form of grants and contracts from federal agencies, such as BARDA, NIH-NIAID and NIH-CounterACT. Additional research has been conducted on our compounds with funding from private foundations, such as the MJFF and Citizens United for Research in Epilepsy ("CURE").

The expected benefit of this strategy is threefold. First, safety, toxicology, pharmacokinetic and CMC work funded by the government and foundations is applicable to our traditional commercial development programs. As an example, significant work funded under the BARDA contract for Lung-ARS has generated data that can be used to support our New Drug Applications ("NDA") for pulmonary fibrosis and/or radiation therapy for cancer.

Second, cost-plus development contracts, like our contract with BARDA, include funds for overhead and profit. These overhead and profit streams have significantly reduced our cash burn rate, which reduces our need to raise capital and incur dilution.

Third, some government contracts, such as the Lung-ARS contract with BARDA are designed to lead to the acquisition of the product under development by the US Government for use as a MCM in the Strategic National Stockpile ("SNS" or the "Stockpile"). Government procurement could result in significant revenue to the Company, which could be used to further the development of the product in other indications or for the development of other promising products. Procurements may be made if either the drug meets the requirements for approval by the U.S. FDA under the "Animal Rule" or prior to Animal Rule approval following the filing of a pre-Emergency Use Authorization ("EUA") application. Most of BARDA's procurements to date have been following the filing of a pre-EUA application.

The amount of any potential procurement is undisclosed by BARDA at this time and is unknown to us. Based on publicly available information, as well as other procurements made by the agency after pre-EUA applications, we believe the agency may purchase sufficient courses of therapy to treat a minimum of one hundred thousand people, with options to purchase an additional two hundred thousand courses of treatment. If purchases of such volumes occurred, the revenue to the Company could provide funding to advance numerous clinical studies, including potentially large Phase III programs in lung fibrosis and radiation therapy for cancer. This funding could allow us to fund studies with less dependence on collaborative partnering arrangements and future equity offerings, which is consistent with our strategy to deploy non-dilutive capital wherever possible to develop our compounds for unmet medical indications and thereby generate value for our stockholders. In addition, purchases of such volumes of drug could make the Company profitable.

Business Overview

We are developing a platform of compounds with anti-fibrotic, anti-inflammatory, anti-infective and anti-oxidant activity based on technology discovered and researched at Duke, UC and NJH, developed by Drs. Irwin Fridovich, Brian Day and others. Dr. Day is our Chief Scientific Officer.

Our lead compound, 10150, protects tissue from damage and increases survival in animal models of lung damage after exposure to radiation toxic chemicals, disease and trauma by mitigating and/or preventing cell death, inflammation and fibrosis through its action on oxidative stress (Reactive Oxygen Species, or "ROS") and regulation of growth factors and chemokines including PTEN, TGF- β 1, HIF-1 α , TNF- α and IL-6, as well as impacting subsequent signaling pathways of ROS production, apoptosis and fibrosis such as NOX4, PTEN, PI3K/p-Akt and p53/Bax (for scientific definitions, see Glossary under "AEOL 10150 in Idiopathic Pulmonary Fibrosis"). 10150 is an extremely potent catalytic anti-oxidant that has been shown in scores of animal studies to reduce oxidative stress by neutralizing reactive oxygen and nitrogen species. The neutralization of these species impacts the inflammatory and fibrotic response to insults and reduces or eliminates subsequent tissue damage-signaling cascades that typically result. We are developing 10150 as a MCM for national defense and for use in oncology and treating lung fibrosis.

Our most extensive development program to date is the advanced development of 10150 for Lung-ARS and DEARE. On February 11, 2011, we signed a cost-plus contract with BARDA for the development of 10150 as a MCM against Lung-ARS. BARDA is the government agency responsible for the advanced development and purchase of medical countermeasures for chemical, biological, radiological and nuclear threats. The contract contemplates the advanced development of 10150 through approval by the FDA under 21 CFR Part 314 Subpart I and Part 601 Subpart H (the "Animal Rule.") The Animal Rule allows for approval of drugs using only animal studies when human clinical trials cannot be conducted ethically. The ultimate goal of the BARDA Contract is to complete all of the work necessary to obtain FDA approval for 10150 as a MCM for Lung-ARS. In addition, the BARDA Contract is designed to generate the data that would allow for acquisition of the drug by BARDA prior to FDA approval under a pre-EUA application.

Pursuant to the BARDA Contract, we were awarded approximately \$10.4 million for the base period of the contract (from February 2011 to April 2012). On April 16, 2012, we announced that BARDA had exercised two options under the BARDA Contract worth approximately \$9.1 million. On September 17, 2013, we announced that BARDA had exercised \$6.0 million in additional contract options. On May 7, 2014, we announced that BARDA had exercised a Contract Modification worth approximately \$1.8 million. The Contract Modification allowed Aeolus to reconcile actual costs incurred with billings under the original provisional indirect billing rate. It established a new provisional indirect billing rate and placed a cap on the current and future provisional indirect billing rates. On June 26, 2015, we announced that BARDA had exercised \$3.0 million in additional contract options under its advanced research and development contract for 10150. On February 8, 2016, BARDA exercised a Contract Modification worth approximately \$0.1 million. On May 25, 2016, we announced that BARDA had exercised a Contract Modification worth approximately \$0.4 million. The purpose of the Contract Modification was to provide funding to complete a pharmacometric analysis of data from all completed animal efficacy studies of 10150 to estimate optimal dose, dose frequency and duration of treatment. The total contract value exercised as of September 30, 2016 is approximately \$30.8 million, of which \$30.6 million has been billed. We may receive up to an additional \$87.6 million in options exercisable over the remainder of the BARDA Contract. Options are exercised based on the progress of the development program, including the completion of clinical trials or manufacturing tasks under previously exercised options.

The final goal of the contract is to achieve FDA approval for 10150 and the development of commercial manufacturing capability. In order to achieve these goals, we believe it will be necessary for BARDA to exercise the majority of the options in the contract. We also believe that BARDA is likely to continue to exercise options as long as 10150 continues to demonstrate efficacy in animal testing and safety in humans for Lung-ARS. In the event we begin sales to the U.S. government following the filing of a pre-EUA application, we believe that BARDA is likely to exercise the majority of the remaining options under the contract. One of the requirements of an EUA is that the development program continue towards the goal of FDA approval. If all of the options are exercised by BARDA, the total value of the contract would be approximately \$118.4 million.

There are no existing treatments for Lung-ARS or DEARE and we are not aware of any compounds in development that have shown efficacy in increasing survival when administered after exposure to radiation. 10150 has demonstrated efficacy in two animal models (mouse and non-human primate) when administered after exposure to radiation. The U.S. government's planning scenario for a radiation incident is a 10 kiloton detonation of a nuclear device in a major American city. It is estimated that several hundred thousand civilians would be exposed to high doses of radiation in this scenario.

The BARDA contract is also designed to complete the work necessary for 10150 to be purchased for the Stockpile. BARDA currently acquires drugs for the SNS through a Special Reserve Fund (the "SRF") created under Project BioShield and reauthorized at \$2.8 billion under the Pandemic All-Hazards Preparedness Reauthorization Act of 2013. Although the final goal of the contract is full FDA approval under the Animal Rule, BARDA may purchase product prior to FDA approval following the filing of a pre-EUA application. BARDA has made numerous acquisitions of compounds that were not approved by the FDA, but were the subject of a pre-EUA filing. Procurements from BARDA following a pre-EUA application could result in a significant increase in revenues for

Aeolus and potential profitability.

In August 2014, we filed an Investigational New Drug ("IND") application with the Division of Medical Imaging Products of the U.S. Food & Drug Administration ("FDA") for 10150 as a treatment for Lung-ARS. On September 4, 2014, the Company announced positive results from a study in non-human primates ("NHP") exposed to lethal radiation and treated with 10150. The study demonstrated that administration of 10150 24 hours after exposure to lethal radiation impacted survival at six months post-exposure as follows: survival in the 60 day treatment group was 50%, compared to 25% survival in the radiation only untreated control group. The data from this study, combined with development work completed in manufacturing and human safety data, will form the basis for a pre-EUA application.

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On September 22, 2014, we received a letter from the FDA placing our proposed Phase I study in healthy normal volunteers for 10150 as a treatment for Lung-ARS on clinical hold. On February 19, 2016, we received notice from the FDA that the clinical hold had been lifted, thereby allowing us to proceed with a Phase I single dose study of 10150 in healthy normal volunteers.

We also benefit from research funded by grants from the NIH CounterACT program for the development of 10150 as a MCM for the effects chemical vesicant gasses (e.g., mustard gas, phosgene gas and chlorine gas) exposure. Funding for this indication is provided directly to the research institution and does not flow through our financial statements. Continued funding is generally dependent on continuing evidence of efficacy in animal trials. In October 2011, NJH was awarded a \$12.5 million grant from NIH CounterACT to continue the development of 10150 as a MCM against sulfur mustard and chlorine gas exposure. Also included in the grant was support for research looking at tissue plasminogen activator ("TPA") and Silabilin, which are not Aeolus assets, as MCMs against sulfur mustard gas exposure. Work under this grant was completed in September 2016, and NJH reported in September 2016 that final studies under the grant demonstrated that 10150 treatment significantly improved acute (2 to 3 days post-exposure) and longer term (28 days post-exposure) in a rodent model of sulfur mustard gas lung exposure. Development of a large animal model is currently being funded by the US Government, independent of Aeolus, and once that model is completed, the Company plans to enter into a Material Transfer Agreement with the US Government to test 10150 and confirm the efficacy seen in the rodent model. Aeolus plans to meet with the FDA during the first half of 2017 to discuss filing an IND for the sulfur mustard indication and to discuss what human safety data would be required for making a pre-EUA filing and ultimately for approval under the Animal Rule.

We are also funded by grant money from the NIH CounterACT program and the National Institute of Neurological Disorders and Stroke ("NINDS") for the development of 10150 as a MCM for the effects of nerve gas (e.g., sarin and soman) exposure. NIH-CounterACT awarded a contract on September 24, 2011 worth approximately \$735,000, to UC to develop 10150 as a MCM against nerve agents. Work performed with this initial funding demonstrated that 10150 significantly improved survival when administered with current treatment in a pilocarpine model for nerve gas exposure. In September 2013, we announced that Dr. Manisha Patel at UC had been awarded a \$4.3 million grant from NINDS to further develop as a MCM for exposure to sarin gas and other nerve agents. Efficacy studies in a mouse model of soman gas are ongoing, and results are expected during 2017.

Substantially all of the past costs for the Lung-ARS program have been funded by the BARDA Contract. We may use our own capital to fund any future costs of the Lung-ARS program that are not funded by the BARDA Contract. To date, the chlorine, phosgene, mustard gas and nerve agent programs have been funded by NIH-CounterACT and NINDS through programs at NJH, UC, and the United States Army Medical Research Institute for Chemical Defense ("USAMRICD").

We are also developing 10150 for the treatment of lung fibrosis, including idiopathic pulmonary fibrosis ("IPF") and other fibrotic diseases. Data demonstrating efficacy of 10150 in IPF was initially generated from animal studies in Lung-ARS and chemical gas lung exposure under the BARDA Contract and NIH grants. Additionally, 10150 demonstrated efficacy in the bleomycin model of IPF, a study funded by Aeolus. On March 17, 2015, we announced that 10150 was granted Orphan Drug Designation by the U.S Food and Drug Administration ("FDA"). The Company plans to initiate a Phase I safety study in patients with IPF in 2017. After we have completed safety studies, we plan to initiate efficacy studies in patients with IPF. Although we believe that BARDA may fund a portion of the IPF development work, we may need use our own capital to fund a portion of the future costs of work associated with IPF and other fibrotic diseases.

We are also developing 10150 for use in combination with radiation therapy for cancer as a treatment to reduce side effects caused by radiation toxicity and to improve local tumor control. Pre-clinical studies at Duke and Loma Linda University have demonstrated that 10150 does not interfere with the benefit of radiation therapy or chemotherapy in prostate and lung cancer. Additional studies have shown that 10150 enhances the anti-tumor activity of radiation and chemotherapy. A significant portion of the development work funded by the BARDA contract is applicable to the development program for radiation oncology, including safety, toxicology, pharmacokinetics and Chemistry, Manufacturing and Controls ("CMC"). After we have completed safety studies, we plan to initiate studies to demonstrate efficacy in toxic side effects related to radiation therapy. We expect to fund a portion of the future costs of work related to radiation therapy opportunities from our own capital.

10150 has been tested in two human Phase I safety studies where it was well-tolerated and no adverse events were observed. Efficacy has been demonstrated in animal models for Lung-ARS, chlorine gas exposure, phosgene gas exposure, sulfur mustard gas exposure (lungs and skin) and nerve gas exposure. In both mouse and NHP studies for Lung-ARS, 10150 treated groups showed significantly reduced weight loss, inflammation, oxidative stress, lung damage, and most importantly, mortality. Therapeutic efficacy has been demonstrated when 10150 is administered 24 hours after exposure to radiation, a requirement for consideration as a radiation MCM for the SNS.

Following the events at the Fukushima nuclear plant in Japan in 2011, we performed radiation exposure studies in mice at the request of Japanese researchers to determine how the administration of 10150 would impact the use of leukocyte growth factors ("LGF") used to treat the hematopoietic or bone marrow syndrome of ARS ("H-ARS"). Data showed that 10150 does not interfere with the efficacy of LGF (in this case Amgen's Neupogen®). Additionally, the study demonstrated that administration of Neupogen®, the current standard of care for H-ARS, increased damage to the lungs. When 10150 was administered with Neupogen® this damage was significantly reduced. We believe that this finding may have important implications for the potential procurement of 10150 for the SNS. In September 2013, BARDA announced that it had entered into a procurement and inventory management agreement with Amgen to provide Neupogen® for the SNS. On March 30, 2015, the FDA approved Neupogen® for the treatment of H-ARS.

In addition to our active IND for Lung-ARS, we have an active IND on file with the FDA for 10150 as a potential treatment for amyotrophic lateral sclerosis ("ALS"). At this time, we do not have any plans to continue development of 10150 for ALS.

We have already completed two Phase I safety studies in 50 humans (39 receiving drug and 13 control) demonstrating that 10150 is safe and well tolerated. CMC work has been completed, pilot lots have been prepared and production is being scaled up under the BARDA Contract.

The Company is developing a second compound, 11114, as a treatment for Parkinson's disease. Research funded by MJFF demonstrated the neuro-protective activity of 11114 in mouse and rat models of Parkinson's disease. The compounds were invented by Brian J. Day, PhD at NJH and Manisha Patel, PhD at UC in collaboration with the Company. We have obtained worldwide, exclusive licenses to develop the compounds from NJH and UC. We plan to complete the remaining work to file an IND application with the FDA during 2017.

In April 2015, we announced the discovery of a new compound, 20415, that has demonstrated anti-inflammatory and anti-infective properties and could be effective in treating cystic fibrosis and combatting anti-biotic resistant bacteria.

The compound was developed under a collaboration between Brian J. Day, PhD at NJH and Aeolus Pharmaceuticals. We have obtained a worldwide, exclusive license to develop the rights to the compound from NJH. We plan to initiate the work to file an Investigational New Drug ("IND") application with the FDA during 2017.

Aeolus' Drug Research and Development Program

The focus of our research and development program is to explore and exploit the therapeutic potential of small molecules with anti-fibrotic, anti-inflammatory, anti-infective and neuro-protective properties. We have achieved our initial research objectives and have begun to extend our preclinical accomplishments into IND enabling non-clinical studies, clinical trials and drug development programs, as well as into the exploration of novel approaches to address the urgent need for novel drugs to combat antibiotic resistant bacteria.

Our lead compound, 10150, protects tissue from damage and increases survival in animal models of lung damage after exposure to radiation toxic chemicals, disease and trauma by mitigating and/or preventing cell death, inflammation and fibrosis through its action on oxidative stress (Reactive Oxygen Species, or "ROS") and regulation of growth factors and chemokines including PTEN, TGF- 1, HIF-1 , TNF- and IL-6, as well as impacting subsequent signaling pathways of ROS production, apoptosis and fibrosis such as NOX4, PTEN, PI3K/p-Akt and p53/Bax (for scientific definitions, see Glossary under "AEOL 10150 in Idiopathic Pulmonary Fibrosis"). 10150 is an extremely potent catalytic anti-oxidant that has been shown in scores of animal studies to reduce oxidative stress by neutralizing reactive oxygen and nitrogen species. The neutralization of these species impacts the inflammatory and fibrotic response to insults and reduces or eliminates subsequent tissue damage-signaling cascades that typically result.

Figure 1

AEOL 10150 Overview	<ul style="list-style-type: none"> √ NADPH (NOX4) Inhibitor
Product Type	<ul style="list-style-type: none"> Broad spectrum anti-fibrotic Catalytic antioxidant
Administration Route	<ul style="list-style-type: none"> √ Subcutaneous administration; self-injection possible √ Alternative formulation in development
Indications in Development	<ul style="list-style-type: none"> √ Radiation Oncology √ Idiopathic Pulmonary Fibrosis √ Pulmonary ARS/DEARE √ Sulfur Mustard; Chlorine Gas; Nerve Gas
Technical Readiness Level (TRL)	<ul style="list-style-type: none"> √ TRL 7/8 for Pulmonary Effects of ARS/DEARE √ Active IND for ALS Phase I (3 studies, 50 patients)
Regulatory Status	<ul style="list-style-type: none"> √ Active IND for Lung-ARS Orphan status for ALS, Lung-ARS and IPF

10150 has shown efficacy in a variety of animal models as a protectant against pulmonary damage and diseases including: lung fibrosis, radiation injury, sulfur mustard gas exposure, chlorine gas exposure and phosgene gas exposure, as well as against neurological damage and diseases including: nerve gas exposure, epilepsy, ALS, and stroke and against diabetes. We filed an IND for 10150 in April 2004, under which human safety trials in ALS patients were conducted as more fully described below under the heading "10150 Clinical Program to Date." In February 2016, the FDA Division of Medical Imaging Products ("DMIP") removed the clinical hold on 10150. As a result, we now have an active IND for Lung-ARS. 10150 has been granted orphan drug designation for ALS, Lung-ARS and Idiopathic Pulmonary Fibrosis by the Office of Orphan Products at the FDA. An overview of the activity of 10150 and its potential as a treatment for a variety of traditional medical needs as well for use as a medical countermeasure against radiological and chemical insults follows:

In "Reactive Oxygen Species and Fibrosis: Further Evidence of a Significant Liason", published in Tissue Research; June 2016, Richter, et. al. stated:

There is a common pathway to fibrosis from insults to the lung and other tissue, whether the insult is radiation, chemical, disease or trauma. Lost parenchyma after tissue injury is usually replaced, because of the ability of parenchymal cells to regenerate. However, the opportunity to regenerate usually becomes unavailable upon repetitive insults associated with chronic inflammation, the secretion of chemokines and the release of profibrotic metabolites, among them, ROS. Further, the production of ROS and the concomitant oxidative stress also contribute to the synthesis and activation of various cytokines and growth factors. ROS contribute to fibrosis via feed forward and feedback loops. As the figure illustrates, Insults induce ROS, which causes inflammation. Fibrosis and inflammation feed back into the path and further increase ROS driven production of cytokines and growth factors. Cytokines and growth factors, in turn contribute to additional ROS formation.

The figure above illustrates how 10150 impacts the common pathways to fibrosis. Data from animal studies in models of radiation, chemical and disease, show that 10150 improves survival by reducing inflammation and fibrosis through its action on oxidative stress (Reactive Oxygen Species, or "ROS"), regulation of growth factors and chemokines such as TGF- 1, HIF-1 , TNF- and IL-6, as well as impacting subsequent signaling pathways of ROS production, apoptosis and fibrosis such as NOX4, PTEN, PI3K/p-Akt and p53/Bax. A more detailed discussion of the pathways can be found in the sections entitled: AEOL 10150 in Radiation Therapy and AEOL 10150 in Idiopathic Pulmonary Fibrosis. AEOL 10150 Medical Countermeasure Development Program

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We and our research partners have been awarded in excess of \$149 million for the development of 10150 as a dual-use, broad spectrum medical countermeasure. The table below details the indications currently under development and the major sources of funding from the US Government to date.

Indication	Funding Source	Amount of Grant/Contract	Research Partners
Lung-ARS	BARDA	Up to \$118.4 million	University of Maryland National Jewish Health
Mustard Gas/ Chlorine Gas	NIH CounterACT	\$20.3 million	University of Colorado
Nerve Agents	NIH CounterACT	\$5 million	University of Colorado

10150 as a potential medical countermeasure against the effects of Pulmonary Acute Radiation Syndrome (Lung-ARS)

Overview

The U.S Government's current planning scenario for a nuclear attack is a 10 kiloton detonation in a major American city. For purposes of comparison, the yield of the bomb dropped on Hiroshima in World War II was approximately 16 kilotons. Such an attack would potentially expose hundreds of thousands of citizens to acute, high dose, ionizing radiation and the lethal effects of Acute Radiation Syndrome ("ARS").

ARS is not a single disease, but rather a series of symptoms and progression of injuries that occur following exposure to ionizing radiation. BARDA is pursuing separate development plans for the acute effects of radiation (such as hematopoietic and acute gastro-intestinal effects) as well as for the delayed effects of ARS, among which pulmonary effects are a major component and contributor to mortality to patients surviving acute effects. At present, 10150 is the only compound in advanced development with BARDA for the delayed effects of radiation exposure, and specifically Lung-ARS.

After exposure, the acute syndromes of ARS are the H-ARS and early-onset gastrointestinal ARS ("GI-ARS"), both of which can be lethal. However, depending on the level and location of radiation exposure, the lethal effects of both H-ARS and early-onset GI-ARS may be reduced with proper treatment, including supportive care (fluids and antibiotics) and LGFs like Amgen's Neupogen® or Neulasta®.

In September 2013, BARDA announced that it had entered into a vendor-managed supply agreement with Amgen to supply its LGF, Neupogen®, to the SNS as a treatment for H-ARS. Although Neupogen® is an FDA-approved drug for neutropenia, it was not approved for H-ARS at the time and would have been used under a pre-EUA application. The procurement of Neupogen® for the SNS is significant for 10150 and its potential role in the treatment of ARS. A 2011 murine study conducted at Indiana University at the request of Japanese researchers confirmed that 10150 does not interfere with the positive effects of Neupogen® in H-ARS and the two products in combination were safe and well-tolerated. More importantly, this study also demonstrated that treatment of H-ARS with Neupogen® exacerbates radiation damage to the lung, even at sub-lethal doses of radiation. Treatment with Neupogen® in combination with 10150 significantly reduced the lung damage. We believe that the use of Neupogen® in treating H-ARS makes the use of 10150 crucial in managing the lung effects of acute radiation exposure. On March 30, 2015, the FDA approved Neupogen® for the treatment of H-ARS.

In radiation incidents, victims who have survived H-ARS and GI-ARS, respiratory failure has been the primary cause of death. Research has shown that damage associated with the exposure to upper half body irradiation or total body irradiation is an acute, but delayed, onset of radiation pneumonitis (inflammation of lung tissue) followed by lung fibrosis (scarring caused by inflammation). The incidence of radiation pneumonitis rises very steeply at relatively low radiation doses. This is Lung-ARS, the syndrome that 10150 is being developed to treat.

The figure below illustrates the mechanisms by which radiation exposure damages the lung causing cell death (apoptosis), inflammation and lung fibrosis. Treatment with AEOL 10150 has improved survival in mouse, rat and NHP models of radiation damage by targeting the overproduction of ROS (oxidative stress) and subsequently impacting the signaling pathways for apoptosis, inflammation and fibrosis. The animal studies have not only shown improvement in survival, but protection of lung tissue and improved function in addition to a reduction in inflammation and fibrosis. Analysis of key biomarkers, such as those shown in the figure below, shows that 10150 treatment after radiation exposure returns their levels to, or near baseline levels.

We believe it is in the government's interest in to provide care not only for survival from the short-term effects of radiation exposure following an event (e.g., H-ARS and GI-ARS), but also to provide care for the delayed effects of radiation exposure, such as Lung-ARS. There are no current FDA-approved or pre-EUA therapies for Lung-ARS. We believe 10150 is the only drug in advanced development with BARDA for Lung-ARS.

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Animal Efficacy Studies – Dose Response and Safety Analysis

In September 2016 a "Dose Response and Safety Analysis" of the animal efficacy data from all murine and NHP Studies was completed by the University of Maryland, School of Pharmacy's Center for Translational Medicine. The objective of this report was to identify the optimal dose and regimen of 10150 to improve overall survival after whole thorax lung irradiation in NHP and C57L/J mouse models. Secondary objectives were to explore the relation between 10150 and radiation-induced lung injury biomarkers. The key findings from the report are summarized below: In the C57L/J mice, 10150 administration 24-hour post-exposure provided maximum prolongation of the 180-day survival independent of treatment duration, with a higher median survival rate compared to treatment initiated at a later time after injury. Using a piecewise parametric survival model in NHP, AEOL 10150 treatment for 60 consecutive days starting at 24 hours post-exposure was found to significantly ($p < 0.001$) decrease mortality from acute radiation pneumonitis compared to the control arm. NHP which received 60 day treatment starting 24 hour post exposure died 40% less frequently per unit time during the pneumonitis phase compared to the untreated arm, which is comparable with the estimated hazard ratio of 0.6 from a cox-proportional hazard model. A graph highlighting the NHP results follows:

About 1.4 months delay was seen before any death event occurring across all untreated and treated NHP arms. So the time of latency to onset of pneumonitis was fixed to 1.4 months, and the hazard for this phase was assumed to be not affected by 10150 treatment. It was assumed that 10150 treatment mainly affected the hazard of pneumonitis phase. The hazard (hazard 1, latent phase) before 1.4 months was estimated to be 0.0008, which means NHPs almost have no chance to die during this phase. In pneumonitis phase (1.4 - 6 months in observed data), the hazard ratio in the second phase (hazard 2, pneumonitis phase) between treatment arms and control arm was estimated to be 1.12, 0.6, and 1.13 for the 28 days, 60 days and 1-28 & 60-88 days treatment arms using the parametric constant hazard model in comparison with the hazard ratios of 0.89, 0.49 and 1.09 from cox proportional hazard model. The results show NHPs in 60 days arm may die 40% less frequently per unit time than NHPs in control arm in pneumonitis phase while NHPs in 28 days and 1-28 & 60-88 days arms are 12% and 13% more likely to die compared to Non-Tx arm during pneumonitis phase.

Animal Efficacy studies – Murine Model

In a 2013 study run under the BARDA contract at the University of Maryland at Baltimore, a total of 120, CBA/J mice were exposed to 14.6 Gray of whole thorax irradiation ("WTLI"). Four cohorts of animals were treated with daily doses of 5mg, 10 mg, 25 mg or 40 mg/kg of 10150 beginning 24 hours after exposure for a total for 28 days. The results are shown in the table below. Survival at six months post-exposure in the optimal treatment group of 25mg/kg of 10150 improved to 40 percent, compared to 10 percent survival in the radiation only group. In addition, animals receiving 10150 showed significant protection of the lungs as measured by differences in wet lung weights and breathing frequency. This study confirms previous studies in animals that demonstrate 10150's protection of the lungs from radiation exposure.

Treatment	Survival	Number of Animals
Radiation Only	10%	20
Radiation + 5 mg/kg AEOL 10150	16%	19
Radiation + 10 mg/kg AEOL 10150	16%	19
Radiation + 25 mg/kg AEOL 10150	40%	20
Radiation + 40 mg/kg AEOL 10150	30%	20

A number of other preclinical studies by researchers at the University of Indiana, the University of Maryland and Duke University have demonstrated the efficacy of 10150 in the protection of healthy, normal tissue from damage due to blast and fractionated radiation exposure.

In 2011, we announced positive results from study of 10150 and Neupogen® as combination therapy for treatment of ARS. The study was conducted by Christie Orschell, PhD of Indiana University. The primary endpoint of the study was to determine drug-drug interactions between Neupogen® and 10150, as well as to monitor safety and tolerability of the two treatments given simultaneously. Results of the study confirmed that 10150 does not interfere with the positive effects of Neupogen® on H-ARS and the two products in combination were safe and well tolerated. In 2012, we announced further data from this study, which demonstrated that treatment of H-ARS with Neupogen® exacerbates radiation damage to the lung. The study also confirmed that treatment with 10150 in combination with Neupogen® significantly reduced the lung damage.

The study entitled "Pilot Study to Test the Effects of Aeolus 10150 on Neupogen®-Induced ANC Recovery in Sub-Lethally Irradiated C57Bl/6 Mice" was initiated at the request of Shigetaka Asano, MD of Waseda University and Arinobu Tojo, MD, PhD and Tokiko Nagamura, MD at the Institute of Medical Science at the University of Tokyo to determine whether there would be any interference with the demonstrated efficacy of Neupogen® as a medical countermeasure against the hematopoietic complications of radiation exposure. In previous treatment of radiation accident victims at Tokai-mura, Dr. Asano and others were able to use Granulocyte Colony Stimulating Factor ("G-CSF") and supportive care to enable victims of 8 to 12 Gy exposure to survive the hematopoietic ("heme") syndrome. Unfortunately, these patients later died due to lung and multi-organ complications. As 10150 has shown separate efficacy against lung complications in mice and in Lung-ARS in non-human primates, this study was undertaken to evaluate whether the Neupogen® and 10150 would be beneficial if used in tandem.

The use of Neupogen® or other G-CSFs or Neulasta® or other Granulocyte-Macrophage Colony Stimulating Factor ("GM-CSF") products is recommended by the Radiation Emergency Assistance Center/Training Site (REAC/TS) at radiation exposures greater than 2 to 3 Gy to mitigate damage to the hematopoietic system. REAC/TS is a response asset of the U.S. Department of Energy and provides treatment capabilities and consultation assistance nationally and internationally. In animal studies G-CSF's have been shown to be effective in increasing survival at levels up to 7.5 Gy due to their positive effects on the hematopoietic damage created by radiation exposure. BARDA began procuring Neupogen® and Sanofi-Aventis' drug Leukine® for the Strategic National Stockpile in September 2013 following a pre-EUA filing.

LGFs as a class have not demonstrated an effect on the two other major sub-syndromes -- GI and Lung. 10150 has demonstrated efficacy in treating the GI sub-syndrome in pilot studies conducted by NIH-NIAID, by protecting crypt cells and reducing diarrhea. More extensive studies of the drug in treating the pulmonary effects of radiation at Duke University and the University of Maryland have shown improved survival and enhanced lung function and improved histology at exposures up to 15 Gy in mice and 11.5 Gy in non-human primates. These exposure levels caused death in 100 percent of animals that were not treated with 10150. Studies at Duke University have also shown a significant

survival advantage for animals treated with 10150 after 15 Gy upper half body irradiation, which causes lethal damage to both the GI tract and the lungs.

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In summary, 10150 has consistently shown a survival advantage and protective effect against Lung-ARS and DEARE when administered 24 hours or more after exposure. Additionally, the current standard of care for the acute ARS syndromes, LGF administration, exacerbates damage to the lungs and 10150 has demonstrated efficacy in reducing that damage.

Animal Efficacy Studies – NHP In a 2014 study run under the BARDA contract at the University of Maryland at Baltimore, a total of 80 rhesus macaque monkeys were exposed to 10.74 Gray of WTLI. Three cohorts of animals were treated with 25 mg/kg of 10150 beginning 24 hours after exposure. The three cohorts then received daily doses of 10150 for 28 days, 60 days or 28 days followed by a pause of 32 days and then an additional 28 days of treatment. As we announced in September 2014, survival at six months post-exposure in the 60 day treatment group was 50%, compared to 25% survival in the radiation-only untreated control group. This study confirms previous studies in non-human primate and mouse models that demonstrate 10150's protection of the lungs from radiation exposure. We plan to publish the detailed results of the study with our research collaborators as soon as possible.

In May 2015, we announced secondary endpoints from the study. The data from these endpoints demonstrated that administration of 10150 for 60 days beginning 24 hours after exposure to 10.74 Gy of radiation:

- Increased mean and median overall survival time
- Increased mean and median survival time in subjects that did not survive to 180 days
- Increased time to onset of increased respiratory rate, a clinical measure of lung injury
- Decreased mortality in subjects with elevated respiratory rate
- Decreased wet lung weight in all animals, suggesting less parenchymal damage and edema
- Increased SpO₂, a measure of compensated lung function
- Diminished radiographic evidence of pneumonitis and fibrosis during the later stages of the study (days 90 -180)

In addition, a new approach to investigating lipids, metabolites and proteins in pathophysiological process, matrix-assisted laser desorption/ionization mass spectrometry imaging (MALDI-MSI) was employed in the study to measure potential biomarkers for lung injury in tissue samples from study subject. Analysis using MALDI-MSI showed that the molecular profile for the naïve (un-irradiated) lung is statistically distinct from irradiated lung and treatment with AEOL 10150 shifts the molecular profile back towards the naïve lung. Two prospective biomarkers found in irradiated, damaged lung tissue were not detectable in either naïve or 10150-treated samples.

In 2010, we initiated a study to test the efficacy of 10150 as an MCM to nuclear and radiological exposure in non-human primates ("NHPs"). The study was designed to test the efficacy of 10150 as a treatment for Lung-ARS and to begin establishing an animal model that can be validated and could be utilized by the FDA for approval of an MCM for Pulmonary Acute Radiation Syndrome under the "Animal Rule". The FDA "Animal Rule" enumerates criteria whereby the FDA can rely on animal efficacy data when "evidence is needed to demonstrate efficacy of new drugs against lethal or permanently disabling toxic substances when efficacy studies in humans, ethically cannot be conducted." The criteria are discussed below.

Results from the study were published in the journal Health Physics, Volume 106, Number 1 (January 2014) under the title "A Pilot Study in Rhesus Macaques to Assess the Treatment Efficacy of A Small Molecular Weight Catalytic Metalloporphyrin Antioxidant (AEOL 10150) in Mitigating Radiation-induced Lung Damage." The primary objective of the study was to determine if 10150 could mitigate radiation-induced lung injury and enhance survival in rhesus macaques exposed to whole thorax lung irradiation ("WTLI") and administered supportive care. Two cohorts of NHPs were exposed to 11.5Gy LINAC-derived photon radiation in the WTLI protocol. The control cohort had n=6 and 10150-treated cohort was n=7. This model showed 100% incidence of severe radiation-induced lung damage. 10150 was administered subcutaneously at 5mg/kg beginning at day 1 post WTLI and continued as a single, daily injection for 28 consecutive days. The final results were presented at the 14th International Congress of Radiation Research in Warsaw, Poland in September 2011. Key findings in the study include:

Exposure of the whole thorax to 11.5 Gy resulted in radiation-induced lung injury in all NHPs in the study and proved 100% fatal in the control animals, despite supportive care including dexamethasone. 11.5 Gy is, therefore, equal to or greater than the LD_{100/180} dose for the WTLI model.

10150, as administered in this pilot study (daily for 28 days at a dose of 5mg/kg subcutaneously), demonstrated potential efficacy in mitigating against fatal radiation-induced lung injury. Treatment with the drug resulted in 28.6% survival following exposure to a radiation dose that proved to be 100% fatal in the untreated control group. Serial CT scans demonstrated less quantitative radiographic injury (pneumonitis, fibrosis, effusions) in the 10150-treated cohort, suggesting that the drug reduces the severity of the radiographically detectable lung injury. Dexamethasone administration yielded a transient benefit on both clinical and radiographic evidence of pneumonitis. The 10150 treated cohort required 1/3 less dexamethasone support due to reduced pulmonary injury in the 10150 treated group, resulting in less frequent clinical "triggers" (respiratory rate ≥ 80) to treat with dexamethasone.

The results of this pilot study are encouraging and suggest that treatment with 10150 results in reduced clinical, radiographic and anatomic evidence of radiation-induced lung injury, which also results in improved survival.

10150 merits further study as a post-exposure MCM against radiation-induced lung injury.

In rodents, non-human primates and humans, radiation of the lungs can cause reduced breathing capacity, pneumonitis, fibrosis, weight loss and death and is characterized by oxidative stress, inflammation and elevated macrophage counts. 10150 has proven to be an effective countermeasure to radiation exposure of the lungs in mice and rats in published studies such as Rabbani et al *Int J Rad Oncol Biol Phys* 67:573-80, 2007, Rabbani et al *Free Rad Res* 41:1273-82, 2007 and Gridley et al *Anticancer Res* 27:3101-9, 2007.

Future Development Plans

Our objective is to develop 10150 as an MCM against Lung-ARS via the FDA's "Animal Rule". This development pathway requires demonstration of the key study efficacy parameter of 10150 treatment in two animal models relevant to the human radiation response and its treatment, demonstration of safety in humans, demonstration of relevant dosing and administration in humans, and clear identification of the mechanism of radiation-induced damage to the lung and its amelioration by the drug candidate.

10150 has several distinct advantages as an MCM, including the following:

- Demonstrated survival increase in animal studies of Lung-ARS when administered 24 hours after exposure,
- Demonstrated reduction in lung fibrosis in animal studies when administered up to 24 hours post exposure,
- Demonstrated histological improvement in lung tissue post-radiation exposure,
- Addresses an unmet medical need as an MCM to Lung-ARS,
- Established safety profile in both clinical and pre-clinical studies,
- Subcutaneous self-administration possible by exposed individuals during emergency,
- Rapid administration, allowing large numbers of patients to be treated quickly,
- Original formulation stable for up to 4½ years at 0–8°C and 1 year at room temperature,
- New formulation stability tested in bulk drug for 3 years at room temperature (25°C) and refrigerated conditions (2–8°C); stability testing in bulk drug will continue to three years,
- New formulation stability tested in final drug product to 24 months under room temperature (25°C), accelerated conditions (40°C) and refrigerated conditions (2–8°C); stability testing in final drug product will continue to 5 years,
- Requires no non-standard storage conditions (i.e., not photosensitive),

- Currently in development as an adjunct to radiation therapy and lung fibrosis; if approved will provide a pre-existing distribution and stockpile resource at oncology centers in the event of a radiological emergency,
- Demonstrated advantage when used in combination with Neupogen®,
- Demonstrated potential as both a therapeutic and prophylactic,
 - Demonstrated efficacy against sulfur mustard gas, phosgene gas, chlorine gas and nerve agent exposures,
 - Potential dual use as an adjunct treatment for cancer patients receiving radiation therapy and treatment of idiopathic pulmonary fibrosis, subject to separate FDA approvals for these indications.

We believe that in order to receive approval from the FDA for Lung-ARS, we will need to demonstrate efficacy in animal models and demonstrate product safety. We also plan to request Fast Track status for this indication. If the FDA accepts our Fast Track request, a rolling NDA submission process is enabled, a key step in achieving Priority Review. The FDA determines within 45 days of a company's request, made once the complete NDA is submitted, whether a Priority or Standard Review designation will be assigned.

In August 2014, the Company filed an IND (IND#112103) for 10150 for Lung-ARS with the Division of Medical Imaging Products ("DMIP") at FDA. On September 22, 2014, the Company received a letter from FDA placing the proposed Phase I study in healthy normal volunteers on clinical hold. On February 19, 2016, following the submission of additional data in response to FDA questions, we received notice that the clinical hold had been lifted, allowing us to proceed with a Phase I single-dose study in healthy normal volunteers.

Upon completion of the single dose, Phase I study in healthy, normal volunteers, we plan to initiate studies in patients with cancer and/or idiopathic pulmonary fibrosis, and will use this data together with the existing safety and tolerability data we have in patients with Amyotrophic Lateral Sclerosis ("ALS") to provide the human safety data required for a potential pre-EUA filing and ultimately an NDA filing for approval under the animal rule. As discussed above, we also plan to file separate 10150 IND's with the Respiratory division of the FDA and the Oncology division of the FDA in 2017 and initiate Phase I studies in patients with pulmonary fibrosis and in patients receiving radiation therapy for cancer upon FDA approval of those INDs.

The FDA's "Animal Rule" enumerates criteria whereby the FDA can rely on animal efficacy data when evidence is needed to demonstrate efficacy of new drugs against lethal or permanently disabling toxic substances when efficacy studies in humans cannot be ethically conducted. The criteria are as follows:

- Knowledge of the mechanism of radiation-induced damage to the lung and its amelioration by the candidate drug.
- Pharmacokinetic and pharmacodynamic analysis to provide information on relevant dose and administration schedule.
- Direct correlation of key study parameters (e.g., survival or major morbidity) with the desired clinical benefit in humans.
- Collection of efficacy data in two species relevant to the human radiation response and its treatment unless otherwise justified under GLP-compliant conditions.
 - A Phase I safety trial using the same product and formulation as used in the pivotal trial(s) is required.

Demonstrate Efficacy in Animal Models

Under the BARDA contract, we have developed and validated mouse and NHP models for Lung-ARS. We have also presented these models to the FDA and they have concurred with our design and our development plan for demonstrating efficacy. We believe that the efficacy data produced in pivotal studies using validated models will provide the data required to demonstrate efficacy of 10150 at the dose and schedule proposed for licensure.

Demonstrate Product Safety

For product approval under the "Animal Rule", we will also demonstrate product safety using the same product and formulation used in the animal efficacy trials and proposed for use in humans. Demonstration of safety includes preclinical demonstration of safety via the standard pre-clinical studies and analyses methods and Phase I safety trials sufficient to demonstrate product safety in the target patient population. We believe our proposed safety studies in healthy normal volunteers, planned studies in pulmonary fibrosis and radiation therapy, as well as completed Phase I studies in ALS may be utilized to demonstrate safety for this indication. We believe that BARDA may fund or co-fund safety studies in healthy normal volunteers and other indications to provide the human safety data required for approval under the animal rule.

Competition

Currently there are no FDA-approved drugs for the treatment of Lung-ARS. We are also not aware of any other drug candidates that have demonstrated the ability to protect the lungs from radiation when administered after exposure, which we believe is a critical aspect of the development of an MCM against the effects of acute radiation syndrome. We are also not aware of any drugs with advanced development funding from BARDA for Lung-ARS.

However, in general, we face significant competition for U.S. government funding for both development and procurements of an MCM for biological, chemical and nuclear threats, diagnostic testing systems and other emergency preparedness countermeasures. The U.S. federal government has currently allocated a significant amount of research funding to the development of countermeasures against the effects of radiation exposure. As a result, there are many drug candidates under development as a possible countermeasure against the various effects and sub-syndromes of radiation exposure.

Funding and Funding Options

On February 11, 2011, we signed a five-year, cost-plus contract with BARDA for the development of 10150 as a MCM against Lung-ARS (the "BARDA Contract"). BARDA is the government agency responsible for the advanced development and purchase of medical countermeasures for chemical, biological, radiological and nuclear threats. The contract funds the advanced development of 10150 through approval by the United States Food & Drug Administration ("FDA") under the "Animal Rule." The Animal Rule allows for approval of drugs using only efficacy studies in animal models.

Pursuant to the BARDA Contract we were awarded approximately \$10.4 million for the base period of the contract (from February 2011 to April 2012). On April 16, 2012, we announced that BARDA had exercised two options under the BARDA Contract worth approximately \$9.1 million. On September 17, 2013, we announced that BARDA had exercised \$6.0 million in additional contract options. On May 7, 2014, we announced that BARDA had exercised a Contract Modification worth approximately \$1.8 million. The Contract Modification allowed Aeolus to reconcile actual costs incurred with billings under the original provisional indirect billing rate. It established a new provisional indirect billing rate and placed a cap on the current and future provisional indirect billing rates. On June 26, 2015, we announced that BARDA had exercised \$3.0 million in additional contract options under its advanced research and development contract for 10150. On February 8, 2016, BARDA exercised a Contract Modification worth approximately \$0.1 million. On May 25, 2016, we announced that BARDA had exercised a Contract Modification worth approximately \$0.4 million. The purpose of the Contract Modification was to provide funding to complete a pharmacometric analysis of data from all completed animal efficacy studies of 10150 to determine optimal dose, dose frequency and duration of treatment. The total contract value exercised as of September 30, 2016 is approximately \$30.8 million. We may receive up to an additional \$87.6 million in options exercisable over the remainder of the BARDA Contract. Options are exercised based on the progress of the development program, including the completion of clinical trials or manufacturing tasks under previously exercised options.

Since we have been awarded the BARDA Contract, substantially all of the past costs associated with the research and development of 10150 as a MCM for Lung-ARS have been covered by the BARDA Contract and we expect substantially all future costs would be covered by the BARDA Contract; however, our expectation in this regards is primarily dependent on continued positive results with 10150 in animal studies and the general risks of doing business with the government. See Risk Factors – "Risks Related to Our Dependence on U.S. Government Grants and Contracts." We may need to use our own capital to fund part of the future costs of the Lung-ARS program that are not funded by the BARDA Contract.

10150 as a potential medical countermeasure against the effects of mustard gas

Overview

Sulfur mustards, of which sulfur mustard gas ("SM") is a member, are a class of related cytotoxic, vesicant chemical warfare agents with the ability to form large blisters on exposed skin and cause pneumonitis and fibrosis in the lungs. In their pure form most sulfur mustards are colorless, odorless, viscous liquids at room temperature. When used as warfare agents they are usually yellow-brown in color and have an odor resembling mustard plants, garlic or horseradish. Mustard agents, including sulfur mustard, are regulated under the 1993 Chemical Weapons Convention. Three classes of chemicals are monitored under this Convention, with sulfur and nitrogen mustard grouped in the highest risk class, "schedule 1." However, concerns about its use in a terrorist attack have led to resurgence in research to develop a protectant against exposure.

Mustard gas is a strong vesicant (blister-causing agent). Due to its alkylating properties, it is also strongly mutagenic (causing damage to the DNA of exposed cells) and carcinogenic (cancer causing). Those exposed usually suffer no immediate symptoms. Within 4 to 24 hours the exposure develops into deep, itching or burning blisters wherever the mustard contacted the skin; the eyes (if exposed) become sore and the eyelids swollen, possibly leading to conjunctivitis and blindness. At very high concentrations, if inhaled, it causes bleeding and blistering within the respiratory system, damaging the mucous membrane and causing pulmonary edema. Blister agent exposure over more than 50% body surface area is usually fatal.

In 2011, the NIH awarded a five-year, \$7.8 million Center of Excellence grant to NJH and the UC Health Sciences Center, both in Denver, Colorado. This Center of Excellence was developed to focus on sulfur mustard toxicity in the lung and skin with the long-term goal to develop an effective treatment for mustard gas induced injury in lung and skin. 10150 was identified by the NJH Center of Excellence as a lead compound for its center, and research work there focused on further testing and studies of 10150.

Research in the area of mustard gas-mediated lung injury has provided experimental evidence that the mechanisms of these injuries are directly linked to oxidative stress, inflammation, cell death, DNA damage and airway cast formation. The ability of superoxide dismutase and catalase to reduce oxidative stress by mitigating the formation of reactive oxygen and nitrogen species initially led to 10150 being tested as an MCM against sulfur mustard exposure to the lungs. Subsequent work, funded by the NIH CounterACT grants, has shown that 10150 improves survival in animal models of sulfur mustard lung exposure by reducing oxidative stress, inflammation, cell death and airway cast formation while protecting against DNA damage.

Non-clinical studies

In September 2016, we announced data from studies performed by Brian Day, Ph.D., at NJH in collaboration with USAMRICD and funded by the CounterACT Program utilizing a new sustained-release formulation. 28 rats, 14 control and 14 treated daily with 10150 for 28 days, were followed for 28 days after inhaled exposure to a lethal dose of sulfur mustard gas. The primary endpoint in the study was survival at 28 days and the secondary endpoint was median survival. Survival at 28 days in the 10150-treatment animals was 36 percent compared to 14 percent in the control group. Median survival was 4 days in the control group compared to 18 days in the AEOL 10150 treated group. There was also improvement in blood oxygenation in the 10150 treated group. These positive results build on data from prior acute and longer-term studies noted below.

Also in September 2016, we announced data from a series of studies in a rat model of the acute effects of sulfur mustard gas on survival, breathing and a number of biomarkers for inflammation and fibrosis. In the study, performed by Brian Day, Ph.D., rats were exposed to lethal levels of sulfur mustard gas, which resulted in 64 percent of the untreated animals dying from the effects of the sulfur mustard gas within 48 hours of exposure. Survival for rats treated with 10150 improved dramatically in a dose dependent manner. Animals administered 10150 every four hours, beginning one hour after exposure, had a survival rate of 88 percent at 48 hours. Treatment with 10150 also:

- Increased blood oxygen saturation by more than 10 percent
- Improved clinical scores of lung health by 57 percent
- Decreased airway casts by 69 percent

Restored levels of TNF-a, IL-6, KC/GRO (rat analog of human IL-8) and IL-1b, which are elevated significantly after exposure to sulfur mustard gas, to control levels

Restored levels of TGF- 1 to control levels

In July 2013, we announced that four separate studies conducted at USAMRICD using 85 rats and comparing 2 different 10150 dosing regimens conclusively demonstrated that 10150 improves survival against an LD60-70 sulfur mustard gas exposure. 10150 improved sulfur mustard gas survival up to 82% over 48 hours. The improvement in survival seen with 10150 treated animals after sulfur mustard gas exposure correlated with improvements in clinical scores, blood oxygenation and airway obstruction. The best improvements in survival and lung function occurred with the 10150 dosing regimen of 5 mg/kg body weight given every 4 hours by subcutaneous injection ($p < 0.0004$). The primary endpoints in these studies were survival and blood oxygen saturation. Secondary endpoints included clinical scores, blood gas and histopathology for cast formations. 10150 or placebo was given to rats one hour after sulfur mustard vapor exposure and repeated every 4 or 8 hours. Forty-eight hours after exposure, lung edema was assessed by changes in the Broncho alveolar lavage (BAL) protein levels. At euthanasia, 48 hours after exposure, 10150 significantly improved ($p < 0.0001$) pulse oximetry, and 10150 treated rats had improved blood oxygenation throughout the study period. Treatment with 10150 also restored blood gas parameters to near normal levels, including: pO₂ ($p < 0.001$), pCO₂ ($p < 0.0016$) and pH ($p < 0.0006$). Finally, 10150 treatment reduced airway cast formation by 50% at 24 hours ($p < 0.017$).

In prior studies 10150 reduced lung edema ($p < 0.05$), decreased SM-induced increases in macrophages ($p < 0.05$) and epithelial cells in BAL fluid ($p < 0.05$). In all three measurements 10150 provided approximately 100 percent protection -- with levels approximating that of the control animals in the study. These results indicate that 10150 significantly improves survival and attenuates lung injury from mustard gas exposure and may provide an effective countermeasure against mustard gas-induced lung injury.

The first whole mustard gas study was completed in October 2009. In June 2010, NJH and Lovelace Respiratory Research Institute reported results from a second whole mustard study confirming that 10150 protects lungs from whole mustard gas exposure in rats. The two studies demonstrated that 10150 protects lungs from whole mustard gas exposure in rats. The primary objective of the study was to determine whether administration of 10150, after exposure, reduces the severity of acute lung injury induced by mustard gas. 10150 was given to rats one hour after sulfur mustard exposure and repeated every 6 hours. Twenty-four hours after exposure, lung edema was assessed by changes in the BAL protein levels. 10150 significantly reduced ($p < 0.05$) mustard gas-induced lung edema as measured by BAL protein levels. In addition, 10150 decreased SM-induced increase in the numbers of BAL neutrophils. These results indicate that 10150 can attenuate lung injury from mustard gas exposure and may provide an effective countermeasure against mustard gas-induced lung injury.

Future Development Plans

Development of a large animal model is currently being funded by the US Government, independent of Aeolus, and once that model is completed, the Company plans to enter into a Material Transfer Agreement with the US Government to test 10150 and confirm the efficacy seen in the rodent model. Aeolus plans to meet with the FDA during the first half of 2017 to discuss filing an IND for the sulfur mustard indication and to discuss what human safety data would be required for making a pre-EUA filing and ultimately for approval under the Animal Rule. We will also seek to launch the two pivotal efficacy studies required for approval by the FDA under the "Animal Rule" as well as complete the necessary safety studies as further described under the heading "10150 as a potential medical countermeasure against the effects of Pulmonary Acute Radiation Syndrome – Future Development Plans – Demonstrate Product Safety."

Competition

There are currently no effective treatments for mustard gas exposure.

However, in general, we face significant competition for U.S. government funding for both development and procurements of medical countermeasures for biological, chemical and nuclear threats, diagnostic testing systems and other emergency preparedness countermeasures. The U.S. federal government has currently allocated a significant amount of research funding to the development of countermeasures against bioterrorism. As a result, there are many drug candidates under development as a possible countermeasure against chemical threat agents.

Funding Options

This development program to date has been funded under the NIH-CounterACT Program and we expect that future efficacy studies necessary for approval by the FDA, including the pivotal rat study and second animal model described above, will be funded by either the Department of Defense or BARDA.

10150 as a potential medical countermeasure against the effects of chlorine gas

Overview

Chlorine gas is a toxic gas that inflicts airway injury through primary oxidative stress and secondary inflammation. Chlorine inhalation was recently used in terrorist/insurgent attacks on military and civilian populations and has caused numerous industrial, transportation, swimming pool, and household accidents, as well as deaths to members of the U.S. military in the past. Chlorine gas, also known as bertholite, was first used as a weapon in World War I. Chlorine gas was also used against the local population and coalition forces in the first Iraq War in the form of chlorine bombs. One of the primary goals of the NIH-CounterACT program is to assist in the development of safe and effective medical countermeasures designed to prevent, diagnose, and treat the conditions caused by potential and existing chemical agents of terrorism and which can be added to the SNS. The SNS is maintained by the Centers for Disease Control and Prevention ("CDC"). The SNS now includes CHEMPACKS that are forward-deployed in secure, environmentally controlled areas throughout the United States available for rapid distribution in case of emergency. CHEMPACKS were developed for threats like chemical and nerve gas where treatment must be administered in less than 24 hours and shipment of MCMs from SNS warehouse locations is not practicable. The CDC has established a diagnostic response network for the detection of nerve agents, mustard, cyanide and toxic metals.

Worldwide, independent of warfare and chemical terrorism, chlorine is the greatest single cause of major toxic release incidents (16.Davis DS, Dewolf GB, Ferland KA, et al. Accidental Release of Air Toxins. Park Ridge, New Jersey: NDC; 1989:6-9.). In the U.S., there are about 5-6,000 exposures per year resulting in, on average, about one death, 10 major, 400-500 moderate, and 3-4,000 minor adverse outcomes. Chlorine gas causes damage to upper and lower respiratory tracts. While chlorine is an irritant, its intermediate water solubility may delay emergence of upper airway symptoms for several minutes. No specific beneficial therapies are available. 10150 has demonstrated a decrease in airway injury, inflammation, oxidative damage, hyperreactivity and cell proliferation after acute chlorine gas inhalation in mice, and therefore could be a possible beneficial therapy for chlorine gas inhalation injury to the airways.

Pre-clinical studies

Under a grant from NIH CounterACT, researchers from National Jewish Health and McGill University have completed a series of preliminary studies demonstrating that 10150 protects lungs from chlorine gas exposure in mice and rats. The primary objective of these studies was to determine whether administration of 10150, after exposure, reduces the severity of acute lung injury and asthma-like symptoms induced by chlorine gas. In 2010, 10150 was given to mice at a 5 mg/kg subcutaneous dose one hour after chlorine gas exposure (100 ppm for 5 minutes) and repeated every 6 hours. Twenty-four hours after exposure, lung inflammation was assessed by changes in Broncho alveolar lavage ("BAL") cellularity and neutrophil influx. 10150 significantly reduced ($p < 0.05$, $n = 6$ /group) chlorine gas-induced lung inflammation as measured by BAL fluid cellularity levels by 40% that appeared to be due to limiting neutrophil influx. 10150 also significantly attenuated ($p < 0.05$, $n = 6$) the degree of asthma-like airway reactivity induced by chlorine gas exposure by 40%. These results indicate that 10150 can attenuate lung injury and asthma-like symptoms from chlorine gas exposure and may provide an effective countermeasure against chlorine gas-induced lung injury. The data from this study was published in December 2010 as AEOL10150: A novel therapeutic for rescue treatment after toxic gas lung injury by Toby McGovern, Brian J. Day, Carl W. White, William S. Powell and James G. Martin in the journal Free Radical Biology and Medicine.

National Jewish Health replicated the mice studies previously conducted by McGill University in rats to determine whether 10150 mitigates lung damage due to chlorine gas exposure in multiple studies between 2011 and 2013. In these studies, 10150 significantly reduced protein, white blood cell, red blood cell, macrophage and neutrophil counts in Broncho alveolar lavage fluid. Additional studies are ongoing at National Jewish Health.

Future Development Plans

Development of animal models is currently being funded by the US Government, independently of Aeolus. Once those models are completed, the Company plans to enter into a Material Transfer Agreement with the US Government to test 10150 and confirm the efficacy seen in the rodent model. We believe that the safety and CMC work being done under the BARDA Lung-ARS contract further described under the heading "10150 as a potential medical countermeasure against the effects of Pulmonary Acute Radiation Syndrome – Future Development Plans" will be sufficient to satisfy the safety and CMC requirements for an NDA filing.

Competition

There are currently no effective treatments for chlorine gas exposure.

However, in general, we face significant competition for U.S. government funding for both development and procurements of MCMs for biological, chemical and nuclear threats, diagnostic testing systems and other emergency preparedness countermeasures. The U.S. federal government has currently allocated a significant amount of research funding to the development of countermeasures against bioterrorism. As a result, there are many drug candidates under development as a possible countermeasure against chemical threat agents.

Funding Options

This development program to date has been funded under the NIH-CounterACT Program and we expect that future efficacy studies necessary for approval by the FDA will be funded by the NIH-CounterACT program or BARDA.

[AEOL 10150 as a potential medical countermeasure against the effects of nerve gas](#)

Overview

Nerve agents, such as sarin gas, have been used in Syria, Iraq and Japan and pose a threat to civilian and military personnel. Sarin gas exposure can cause pain, weakness, vomiting, diarrhea and changes in heart rate within minutes to 18 hours after exposure. High levels of exposure can cause convulsions, paralysis, breathing problems and death. BARDA classifies nerve agents as a priority threat.

10150 is the focus of a sponsored research grant awarded by the National Institutes of Health (NIH), National Institute of Neurological Disorders and Stroke (NINDS) and Office of the Director (OD) to the University of Colorado to test its potential efficacy as a MCM against nerve agents.

Pre-clinical studies

In September 2013, we announced that researchers, led by Dr. Manisha Patel at the University of Colorado, completed a pilot study demonstrating that 10150 provides neuroprotection, decreases oxidative stress, and significantly improves survival in rats exposed to pilocarpine – a surrogate for the nerve agents soman and sarin gas. These data were presented at the 6th Annual CounterACT Countermeasures Against Chemical Threats Network Research Symposium in Washington, D.C.

The current standard of care for nerve agent exposure is Atropine® and benzodiazepines. In this study 10150 significantly improved the survival of animals exposed to pilocarpine. Injection of 10150 sixty minutes after pilocarpine in the presence of standard therapy resulted in 100% survival and near complete inhibition of oxidative stress indices in the hippocampus.

Future Development Plans

On September 10th 2013, NIH-NINDS notified Dr. Patel that a total of \$4.3 million had been awarded for her project titled "Neuroprotective Effects of AEOL 10150 against organophosphate toxicity." The research work is being conducted in Dr. Patel's lab at the University of Colorado, at National Jewish Health as well as a large Contract Research Organization. Funding for this research has been awarded under a grant supported by the CounterACT Program, National Institutes of Health Office of the Director (NIH OD), and the NINDS.

Competition

The current standard of care for nerve agent exposure is Atropine® and benzodiazepines. 10150 would be added to current standard of care to improve outcomes. We believe that other companies are developing potential treatments for nerve gas, although it is difficult to evaluate their success unless data is announced publicly.

However, in general, we face significant competition for U.S. government funding for both development and procurements of medical countermeasures for biological, chemical and nuclear threats, diagnostic testing systems and other emergency preparedness countermeasures. The U.S. federal government has currently allocated a significant amount of research funding to the development of countermeasures against bioterrorism. As a result, there are many drug candidates under development as a possible countermeasure against chemical threat agents.

Funding Options

This development program to date has been funded under the NIH-CounterACT Program and NINDS and we expect that future efficacy studies necessary for approval by the FDA will also be funded by the \$4.3 million grant to Dr.

Patel.

AEOL 10150 in Radiation Therapy

Overview

According to the American Cancer Society, cancer is the second leading cause of death by disease, representing one out of every four deaths in the United States. Approximately 572,000 Americans were expected to die of cancer in 2011. In 2012, about 1.6 million new cancer cases were expected to be diagnosed in the United States. According to the Radiological Society of North America, about 50 to 60 percent of cancer patients are treated with radiation at some time during their disease.

Combinations of surgery, chemotherapy and radiation treatments are the mainstay of modern cancer therapy. Success is often determined by the ability of patients to tolerate the most aggressive, and most effective, treatment regimens. Radiation therapy-induced toxicity remains a major factor limiting radiation doses. The ability to deliver maximal radiation doses for treatment of tumors without injury to surrounding normal tissue has important implications in oncology therapeutic outcomes because higher doses of radiation therapy may improve both local tumor control and patient survival.

Advances in the tools of molecular and cellular biology have enabled researchers to develop a better understanding of the underlying mechanisms responsible for radiation therapy-induced normal tissue injury. For decades ionizing radiation has been known to increase production of free radicals, which is reflected by the accumulation of oxidative damaged cellular macromolecules.

As one example of radiation-induced damage to adjacent normal tissue, radiation therapy may injure pulmonary tissue either directly via generation of ROS or indirectly via the action on parenchymal and inflammatory cells through biological mediators such as TGF- β and pro-inflammatory cytokines. Since the discovery of SOD, it is apparent that these enzymes provide an essential line of defense against ROS. SODs and SOD mimics, such as 10150, act by catalyzing the degradation of superoxide radicals into oxygen and hydrogen peroxide. SODs are localized intra/extracellularly, are widely expressed throughout the body, and are important in maintenance of redox status (the balance between oxidation and reduction). Previous studies have demonstrated that treating irradiated animal models with SOD delivered by injection of the enzyme through liposome/viral-mediated gene therapy or insertion of human SOD gene can ameliorate radiation therapy-induced damage.

For an illustrative example of the radiation therapy reaction see Figure 2.

Figure 2

Figure 2 above shows the mechanisms of radiation injury; 10150 protects against this damage by reducing ROS/RNS and subsequently reducing TGF- 1 and NADPH Oxidase (NOX4) activation resulting in reductions of DNA Damage, Apoptosis, Vascular Dysfunction, Hypoxia, Inflammation and Fibrosis.

In vitro studies have demonstrated that 10150 reduces the formation of lipid peroxides, forms of ROS, and that it inactivates biologically important ROS molecules such as superoxide, hydrogen peroxide and peroxynitrite. 10150 inactivates these ROS by one or two electron oxidation or reduction reactions in which the oxidation state of the manganese moiety in 10150 changes. 10150 is not consumed in the reaction and it continues to inactivate such ROS molecules as long as it is present at the target site. Preclinical models and human safety studies suggest 10150 is not metabolized in the body and is excreted in feces and urine.

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Animal studies have demonstrated that 10150 treatment after exposure to radiation (in radiation therapy and blast radiation models) reduces: oxidative stress, apoptosis, vascular damage, hypoxia, inflammation and fibrosis, as well as returning to or near baseline, levels of TGF- 1, NOX4, PTEN, 8OHdG and 4HNE.

Pre-clinical studies

Figure 3

Figure 3: Relative tumor volumes of human prostate tumor implants in nude mice: Implants of well-vascularized PC3 tumors were grown to substantial size prior to receiving fractionated radiation (5 Gy daily for three days). 10150 (7.5 mg/kg/bid) was administered subcutaneously commencing on the first day of irradiation and continued for 20 days. Other groups of mice received either no irradiation, irradiation only or 10150 without irradiation.

Due to the common mechanism of action in cancer radiation therapy and radiation exposure from nuclear events, we believe that the pre-clinical studies performed for the development of 10150 as a potential medical countermeasure against the effects of Lung-ARS, as described herein, also provide support for the development of 10150 in oncology, to be used in combination with radiation therapy.

We have performed several additional studies specifically for this indication to ensure the use of 10150 in radioprotection of normal adjacent tissue does not interfere with the efficacy of tumor radiotherapy. A number of preclinical, in vivo studies have addressed this issue and have demonstrated that 10150 does not negatively impact tumor radiotherapy.

In one study (Vujaskovic, et al. of Duke) human prostate tumors (PC3) grown in nude mice to substantial size were irradiated with 5 Gy per day for 3 days for a total of 15 Gy. 10150 at 7.5 mg/kg/bid was administered subcutaneously on the first day of radiation and continued for either of two time courses: when tumor volume reached 5 times the initial volume or for twenty days. The receding tumor volume curves for irradiation only and for irradiation plus 10150 demonstrated no difference. Therefore 10150 did not interfere with the radiation effect on prostate tumors.

In another study of prostate cancer tumors (Gridley, et al of Loma Linda University), mouse prostate cancer cell line RM-9 was injected subcutaneously into C57/Bl6 mice, followed by up to 16 days of 10150 delivered intraperitoneally at 6 mg/kg/day. On day seven, a single dose of radiation (10 Gy) was delivered. Therefore, the mice received compound for seven days prior to radiation. The results of this study demonstrated that 10150 does not protect the prostate tumor against radiation, however, 10150 showed a trend towards increasing the effectiveness of the radiation treatment. The primary effect appears to be in down-regulation of radiation induced HIF-1 expression and VEGF and up-regulation of IL-4. Thus, 10150, through its down-regulation of VEGF, may inhibit formation of blood vessels (i.e., angiogenesis) required for tumor re-growth and protects normal tissues from damage induced by radiation and chemotherapy.

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In another study (Vujaskovic, et al. of Duke), mice were implanted with human NSCLC tumors and treated with all potential combinations of paclitaxel, radiation and 10150 to determine the impact on tumor growth. The results showed that 10150 did not impact the effects of either radiation therapy or paclitaxel. Further, the study indicated that the greatest impact in inhibiting tumor growth was with the regimen that included all three courses of treatment, radiation, paclitaxel and 10150.

Figure 4

Figure 4 above measures tumor volume against time after implantation of RM-9 tumor cells and shows that 10150 treatment resulted in inhibition of tumor re-growth in a study performed by Dr. Gridley of Loma Linda University. Daily intraperitoneal injections of 10150 were initiated on day 1. At 12 days, approximately one half of each tumor-bearing group and control mice with no tumor were euthanized for in vitro analyses; remaining mice/group were followed for tumor growth and euthanized individually when maximum allowed tumor volume was attained. Each point represents the mean +/- standard error of the mean. Two-way analysis of the variance for days 8 to 14 revealed that group and time had highly significant main effects ($P < 0.001$) and a group x time interaction was noted ($P < 0.001$).

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Figure 5

Figure 5 above shows the HIF-1 Expression in prostate tumors and the impact of the treatment of 10150 in a study by Dr. Gridley of Loma Linda University.

Figure 6

Figure 6 above shows impact on tumor growth in mice that were implanted with human NSCLC tumors and treated with all potential combinations of paclitaxel (P), Radiation (RT) and 10150 (A).

In summary, the data obtained in these preclinical studies suggest that the post-irradiation, long-term delivery of 10150 may be protective against radiation-induced lung injury, as assessed by histopathology and immunohistochemistry. Oxidative stress, inflammation and hypoxia, which play important roles in the pathogenesis of radiation mediated fibrosis, were shown to be reduced in animals treated with higher doses of 10150. Studies have also shown that 10150 does not adversely impact tumor response to radiation therapy. Thus, treatment with 10150 does not significantly protect tumors from the cell killing effects of radiation therapy. This combined with other studies that have shown that 10150 significantly prevents radiation induced normal tissue injury suggests that 10150 has the potential to achieve protection of normal tissue without protection of tumor tissue. Additionally, it appears the down-regulation of radiation induced HIF-1 expression and VEGF and up-regulation of IL-4 may provide additional anti-tumor effects. Thus, 10150, through its down-regulation of VEGF, may inhibit formation of blood vessels required for tumor re-growth, while protecting normal tissues from damage induced by radiation and chemotherapy.

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Future Development Plans

We are leveraging the significant investment made by U.S. government agencies to develop this promising compound for use in oncology indications, where it would be used in combination with chemotherapy and radiation therapy, and is currently in development for use as both a therapeutic and prophylactic drug. Data has already been published showing that 10150 does not interfere with the therapeutic benefit of radiation therapy in prostate and lung cancer preclinical studies.

In 2017, we expect to initiate a safety study in non-small cell lung cancer patients receiving chemotherapy and radiotherapy. Upon the successful completion of the Phase I study and approval of a protocol by the FDA and the appropriate IRBs, we expect to begin a Phase II study in cancer patients receiving radiation therapy. A specific indication for the Phase II study has not yet been chosen, however, we are considering esophageal, prostate, head and neck and NSCLC patients.

Competition

There are currently three drugs approved for the treatment of the side effects of radiation therapy. We do not believe that any of these drugs directly competes with 10150 in terms of mechanism of action or targeted therapeutic benefit when used in combination with radiation therapy.

Amifostine (Ethyol®) is approved by the FDA as a radioprotector. Amifostine (Ethyol) is marketed by MedImmune, Inc. for use in reduction of chemotherapy-induced kidney toxicity associated with repeated administration of cisplatin in patients with advanced ovarian cancer and radiation-induced xerostomia (damage to the salivary gland) in patients undergoing post-operative radiation treatment for head and neck cancer. MedImmune, Inc. is studying Amifostine in other indications of radiation therapy. Kepivance™ (palifermin) is marketed by Amgen, Inc. for use in the treatment of severe oral mucositis (mouth sores) in patients with hematologic (blood) cancers who are undergoing high-dose chemotherapy followed by bone transplant. Amgen, Inc. is also studying Kepivance as an antimucositis agent in patients with head and neck cancer, non-small cell lung cancer and colon cancer. Salagen Tablets (pilocarpine hydrochloride) is marketed by Eisai Pharmaceuticals in the United States as a treatment for the symptoms of xerostomia induced by radiation therapy in head and neck cancer patients. In addition, there are many drugs under development to treat the side effects of radiation therapy.

Funding Options

Substantially all of our costs associated with the CMC and toxicology necessary for the oncology indications have been covered by the BARDA Contract. We expect such costs to continue to be covered by the BARDA Contract unless BARDA chooses not to exercise its options under the BARDA Contract. In such a circumstance, we would need to raise additional capital, or partner with another firm, in order to complete the non-clinical and safety programs related to Lung-ARS noted above. We may need to use our own capital to fund elements of this program that are not funded by the BARDA Contract, including funds for an initial trial in human patients receiving therapeutic radiation for cancer.

10150 Clinical Development Program to Date

10150 has been tested for safety, tolerability and pharmacokinetics with no serious or clinically significant adverse effects observed. To date, 38 patients have received 10150 in three clinical trials designed to test the safety and tolerability of the drug candidate.

In September 2005, we completed a multi-center, double-blind, randomized, placebo-controlled, Phase I clinical trial. This escalating-dose study was conducted to evaluate the safety, tolerability and pharmacokinetics of 10150 administered by twice daily subcutaneous injections in patients with ALS.

In the Phase Ia study, 4-5 patients diagnosed with ALS were placed in a dosage cohort (3 or 4 receiving 10150 and 1 receiving placebo). Each dose cohort was evaluated at a separate clinical center. In total, seven separate cohorts were evaluated in the study, and 25 ALS patients received 10150. Based upon an analysis of the data, it was concluded that single doses of 10150 ranging from 3 mg to 75 mg were safe and well tolerated. In addition, no serious or clinically significant adverse clinical events were reported, nor were there any significant laboratory abnormalities. Based upon extensive cardiovascular monitoring (i.e., frequent electrocardiograms and continuous Holter recordings for up to 48 hours following dosing), there were no compound-related cardiovascular abnormalities.

The most frequently reported adverse events in this Phase I clinical trial were injection site reactions, followed by dizziness and headache. Adverse events were primarily mild in severity, and approximately one-half of the events were considered to have a possible relationship to the study medication. In addition, no clinically meaningful findings were noted in the safety, laboratory, vital sign, the Unified Parkinson's disease Rating Scale ("UPDRS"), functional ALS, or electro cardiogram ("ECG") data. All cohorts exhibited dose-related peak plasma drug concentrations and consistent disappearance half-lives.

In October 2006, we completed a multi-center, double-blind, randomized, placebo-controlled, Phase Ib clinical trial. This multiple dose study was conducted to evaluate the safety, tolerability and pharmacokinetics of 10150 administered by subcutaneous injection and infusion pump in patients with ALS. Under the multiple dose protocol, three groups of six ALS patients (four receiving 10150 and two receiving placebo) were enrolled.

The Phase Ib study was conducted at five academic clinical ALS centers. Male and female ALS patients, 18 to 70 years of age, who were ambulatory (with the use of a walker or cane, if needed) and capable of orthostatic blood pressure assessments were enrolled in the study. Clinical signs/symptoms, laboratory values, cardiac assessments and pharmacokinetics (PK) were performed.

Based upon an analysis of the data, it was concluded that multiple doses of 10150 for a period of six and one half consecutive days in the amount of 40 mg per day, 60 mg per day and 2 mg per kilogram per day were safe and well tolerated. No serious or clinically significant adverse events were reported or observed. The most frequent adverse events related to study compound were injection site observations related to compound delivery. There were no significant laboratory abnormalities. Based upon extensive cardiovascular monitoring (i.e., frequent electrocardiograms and continuous Holter recordings throughout the six and one half days of dosing), there were no compound-related cardiovascular abnormalities.

Pharmacokinetic findings from the Phase Ib study to data are as follows:

- Increases in C_{max} and AUC (0-8) appear to correlate with increases in dose, but the correlation is not strong. The mean C_{max} for the 40 mg cohort was 1,735 ng/mL; 2,315 ng/mL for the 60 mg cohort and 1,653 ng/mL for the 2 mg/kg cohort.
 - There were probable linear correlations between both C_{max} and AUC(0-8) and dose based on body weight. The terminal half-life (a measurement of the time period for which a compound stays in the body) as determined from Day 7 data was approximately 8 to 9 hours. Steady-state occurs within three days of multiple dosing. There was no evidence for a third longer half-life that would be associated with long term accumulation. Thus, compound accumulation is not expected beyond the third day with multiple dosing.
 - From 48 hours to the end of the infusion, the plasma concentrations of 10150 during the infusion showed little variability, indicating a smoother delivery of the drug than with twice-daily injections.
- During 2008, we completed a follow-on Phase I open label compassionate use multiple dose study of 10150 in a patient diagnosed with progressive and debilitating amyotrophic ALS. The study was conducted at the University of California, Los Angeles by Martina Wiedau-Pazos, M.D., and was designed to evaluate the safety and efficacy of 10150 in an ALS patient over an extended period of time. The patient received a subcutaneous injection of 75mg of 10150 two times each day for 34 days. Efficacy and safety data was monitored for the duration of the study. The primary objective of this study was to assess the clinical efficacy of 10150 with respect to the patient's baseline assessment of functional status. Secondary objectives included the assessments of muscle strength, respiratory function, quality of life and safety. The patient's baseline efficacy results were an ALS Functional Rating Scale ("ALSFRS-R") rating of 19, Muscle strength Manual Muscle Testing Scale ("MMTS") of 68 and a forced vital capacity ("FVC") of 30%. The patient's results after 2 months were an ALSFRS-R rating of 22, a MMTS rating of 86 and an FVC of 28%. It should be noted that the subject began using breathing assistance (BiPAP) approximately two

weeks after the study started. The patient discontinued treatment due to nausea and moderately increased liver transaminases. Other drug-associated adverse events included mild skin irritation at the injection site and mild urine discoloration.

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AEOL 10150 in Idiopathic Pulmonary Fibrosis

Overview

Idiopathic pulmonary fibrosis ("IPF") is a progressive, fibrosing interstitial pneumonia of unknown cause characterized by the development of scar tissue that thickens the lining of the lungs and generally leads to death due to inhibition of lung function. IPF affects approximately 128,000 people in the United States with 48,000 new cases and 40,000 deaths annually. It usually affects older individuals, those in the sixth to seventh decade, and is a severe disease with a variable clinical course and a high mortality rate. Approximately 2/3 of IPF sufferers will die within 5 years of diagnosis. The median survival after diagnosis is 3 to 4 years.

Standard of care for IPF is treatment with anti-inflammatory drugs such as corticosteroids although there is no evidence that these treatments are effective in increasing long-term survival. Supplemental oxygen may improve patient quality of life, but has no effect on the course of the disease. In some instances, lung transplantation may be considered. On October 15, 2014, the FDA approved Esbriet® (pirfenidone), manufactured by Intermune, and Ofev® (nintedanib), manufactured by Boehringer Ingelheim, for the treatment of IPF. We believe that there is still a significant need for additional and more effective treatments for IPF.

As with lung damage from radiation and chemical insults, the fibrosis in IPF patients is believed to be connected to increased levels of ROS, TGF- 1 and NOX4 and their effects on inflammation and apoptosis and the proliferation and differentiation of fibroblasts into myofibroblasts, which damage lung tissue and and lead to Extracellular Matrix (ECM) deposition. The following figure illustrates the pathway from increased TGF- levels to ECM/Fibrosis in the lungs:

Source: Keitzmann, T., e.t al.; Redox Biology, 2015

As noted in the previous discussion on the common pathway to fibrosis from insults to the lung, lost cells after tissue injury are usually replaced. However, normal regeneration typically becomes impossible in the presence of chronic inflammation, TGF- 1 and increased presence ROS levels. ROS are involved in transforming growth factor (TGF) mediated fibrosis.

Keitzmann, et. al. describe the role of ROS, TGF- 1 and NOX4 in generating lung fibrosis, as follows:

"Various cell types such as platelets, parenchymal cells and inflammatory cells (e.g., lymphocytes, macrophages) can release TGF 1. After conversion of the latent to the active form, TGF 1 binds to its receptor and induces SMAD2/3 and/or phosphatidyl inositol 3kinase (PI3K) signaling to express various genes, among them that for NOX4. NOX4 in turn leads to ROS production. Enhanced ROS may activate the proliferation, migration and differentiation of fibroblasts plus epithelial to mesenchymal transition (EMT), apoptosis of epithelial cells and/or excessive extracellular matrix (ECM) deposition. In addition, TGF-β1 contributes to ROS enhancement by decreasing the expression of antioxidant enzymes like glutaredoxin (Grx), catalase (CTL), glutathione peroxidase (GPX), glutathione S transferase (GST), superoxide dismutase (SOD) and the heavy subunit of gamma-glutamylcysteine synthetase (gamma-GCSh).

Targeting TGF- , NOX4 and ROS are widely considered to be valid approaches to treating patients with IPF. AEOL 10150 has demonstrated in animal studies in mice, rats and NHP an ability to reduce the levels of these targets after a variety of lung insults.

Glossary of Scientific Terms

- BAX** Apoptosis regulator BAX - a protein that is encoded by the BAX gene. The expression of this gene is regulated by the tumor suppressor P53 and has been shown to be involved in P53-mediated apoptosis.
- HIF-1** Hypoxia-inducible factor 1-alpha - the master transcriptional regulator of cellular and developmental response to hypoxia. The dysregulation and overexpression of HIF-1 by either hypoxia or genetic alternations have been heavily implicated in cancer biology, as well as a number of other pathophysiologies, specifically in areas of vascularization and angiogenesis, energy metabolism, cell survival, and tumor invasion.
- IL-6** Interleukin 6 – a pro-inflammatory cytokine and an anti-inflammatory myokine encoded by the IL6 gene.
- NOX4** NADPH oxidase 4 - an enzyme that is encoded by the NOX4 gene. The ROS generated by this protein have been implicated in numerous biological functions including signal transduction, cell differentiation and tumor cell growth.
- Akt** Protein kinase B - a protein kinase that plays a key role in multiple cellular processes such as glucose metabolism, apoptosis, cell proliferation, transcription and cell migration.
- PTEN** Phosphatase and tensin homolog - a protein encoded by the PTEN gene. This gene has been identified as a tumor suppressor that is mutated in a large number of cancers at high frequency.
- p53** Tumor protein p53 – a protein encoded by the TP53 gene in humans. It prevents cancer formation and functions as a tumor suppressor.
- PI3K** Phosphatidylinositol-4,5-bisphosphate 3-kinase - a family of enzymes involved in cellular functions such as cell growth, proliferation, differentiation, motility, survival and intracellular trafficking, which in turn are involved in cancer.
- ROS** Reactive oxygen species – chemically reactive chemical species containing oxygen. ROS are formed as a natural byproduct of the normal metabolism of oxygen and have important roles in cell signaling and homeostasis. ROS are also generated by external sources such as ionizing radiation.
- TGF-** Transforming growth factor beta 1 – a cytokine encoded by the TGFB1 gene that controls proliferation, differentiation, and other functions in many cell types.
- TNF-** Tumor necrosis factor alpha – a cell signaling cytokine involved in systemic inflammation. It is produced chiefly by activated macrophages.

Pre-clinical Studies

Whole Thorax Irradiation ("WTLI") is a validated model for IPF in animals. The efficacy studies under the BARDA Contract in monkey and mice were conducted using a WTLI model. Those studies demonstrated a significant reduction in inflammation and fibrosis in animals treated with AEOL 10150. Additionally, during 2015, researchers at National Jewish Health conducted efficacy studies of 10150 using a Bleomycin induced fibrosis model. These studies also demonstrated a significant reduction in fibrosis in animals treated with 10150, and provided further support for the drug's potential as a therapy for patients with pulmonary fibrosis, including IPF.

Development Plans

In March 2015, the Company received notice from the Office of Orphan Products Development at the U.S. Food & Drug Administration (FDA) granting Orphan Drug Designation for AEOL 10150 "for treatment of idiopathic pulmonary fibrosis." Orphan Drug Designation entitles the sponsor to a seven-year marketing exclusivity period, clinical protocol assistance with the FDA, as well as federal grants and tax credits. In October 2015, the Company announced the formation of a Pulmonary Fibrosis Clinical Advisory Board ("PFCAB") led by Kevin Brown, MD, Professor and Vice Chair, Department of Medicine, at National Jewish Health. The Company plans to meet with the Respiratory division of the FDA and file an IND to initiate Phase I studies of 10150 in patients with pulmonary fibrosis.

Competition

Esbriet® and Ofev® were approved by the FDA for the treatment of IPF on October 15, 2014. Esbriet® and Ofev® are also approved for use in Europe, Canada, Japan, South Korea and other countries. The effectiveness of Esbriet® and Ofev® are, however, limited and we believe that there is still a significant need for more effective treatments. There are a number of potential treatments for IPF in advanced development by multiple biotechnology and pharmaceuticals. Many of these potential treatments are monoclonal antibody therapies. We believe that 10150 exploits a novel mechanism of action in the treatment for IPF. In general, however, we believe that there will be substantial, well-funded competition for the development of new IPF treatments.

Funding

Although the initial data supporting a program in IPF was largely generated under the BARDA Contract, we are uncertain as to whether funding for human safety and efficacy studies will be available from that source. However, as with radiation oncology, manufacturing, toxicology and safety work under the contract can also be used in the development of 10150 for IPF. We may use our own capital to fund part of the future costs of this program that are not funded by the BARDA Contract, including funds for an initial trial in patients with IPF. The Company will also seek non-dilutive capital sources and pursue potential partnership opportunities to advance the IPF program.

AEOL 11114B in Parkinsons' Disease

Overview

We have evaluated three compounds, 11114, AEOL 11203 ("11203") and AEOL 11207 ("11207"), for the treatment of Parkinsons Disease with funding from MJFF. From these three candidates, 11114 was selected for further development. Parkinson's disease is a common neurodegenerative disorder, second in occurrence among these disorders only to Alzheimer's disease. According to the National Parkinson Foundation, Parkinson's affects as many as one million people in the United States, with approximately 60,000 new cases diagnosed in the United States each year.

Parkinson's specifically involves the progressive destruction of the nerves that secrete dopamine and control the basal ganglia, an area of the brain involved in the regulation of movement. Dopamine turnover has been shown to elevate the levels of ROS in the brain. In addition, a street-drug contaminant has appeared that can cause Parkinsonism in drug abusers. The compound N-methyl-4-phenyl-1, 2, 3, 6-tetrahydropyridine ("MPTP") has been identified in

underground laboratory preparations of a potent analog of meperidine (Demerol). MPTP-containing powder, sometimes sold as a new "synthetic heroin," can be dissolved in water and administered intravenously or taken by the intranasal route. MPTP has been documented to produce irreversible chronic Parkinson symptoms in drug abusers. Agents such as MPTP overproduce ROS in the basal ganglia. Therefore, ROS mediated neuronal dysfunction may play a key role in the development of Parkinson's disease. Symptoms of this disease include tremors, rigidity and bradykinesia (i.e., slowness of movement). In the more advanced stages, it can cause fluctuations in motor function, sleep problems and various neuro-psychiatric disorders. A biological hallmark of Parkinson's disease is a reduction in brain dopamine levels. Preventing or slowing the destruction of brain cells that lead to the depletion of dopamine levels in the brain is an important therapeutic approach for the treatment of this disease.

Pre-clinical studies

Data developed by our scientists and Dr. Manisha Patel at UC and funded by MJFF, established the maximum tolerated dose, pharmacokinetics, dosing paradigm, drug stability, and drug protein binding and efficacy for 11114. The data from the efficacy work performed show that daily or every other day dosing with the compound protects against oxidative stress, improves performance on behavioral tests, and mitigates dopamine depletion in the 6-hydroxydopamine model of Parkinson's disease in restorative manner. These results validate 11114's efficacy in the MPTP mouse model and suggest that the drug can be given after symptom onset after a 6-hydroxydopamine lesion in rats to achieve neuroprotection. Further, administration of 11114 every other day reduced oxidative stress and protected against behavioral deficits and loss of striatal dopaminergic neurons in the Substantia Nigra Pars Compacta ("SNPc"). Parkinson's disease is characterized by the death of neurons in the SNPc. Currently available treatments for Parkinson's lessen the symptoms for some time, but there remains a great need for more effective therapies, such as 11114, that have shown the potential to protect against the loss of dopaminergic neurons. 11114 readily crosses the blood brain barrier, and daily or every other day treatment prevents oxidative stress and protects against oxidative stress, neuron loss and behavioral deficits in both a rat and mouse model of Parkinson's disease.

Prior to receiving the funding for this program, we filed a new composition of matter and use patent for AEOL11114. In October 2015, the Company received notice of allowance of a composition of matter patent from the European Patent Office. In March 2016, we received notice of allowance of a composition of matter patent from the Japanese Patent Office.

Future Development Plans

We plan to complete the necessary pre-clinical work that will allow us to file an IND and initiate Phase I studies in Parkinson's disease in 2017.

Funding Options

We expect our own capital will be used to fund part of the future costs of this program, including amounts to fund the pre-clinical work necessary to file an IND for 11114. We will also seek non-dilutive capital sources and pursue potential partnership opportunities to fund the human clinical studies of 11114 for Parkinson's disease.

Contracts and Grants

We seek to advance development of our drug candidates through external funding arrangements. We may slow down development programs or place them on hold during periods that are not covered by external funding. We have received external funding awards for the development of 10150 as an MCM for Lung-ARS, GI-ARS, mustard gas and chlorine gas exposure from the NIH.

BARDA Contract; Background and Recent Developments

In December 2009, we were informed by BARDA that we had been chosen to submit a full proposal for funding of our Lung-ARS program from its current stage through FDA approval, based on a summary "white paper" submitted by us earlier in 2009. We submitted a full proposal in February 2010. We were notified in July 2010 that our proposal had been chosen by BARDA, and then entered into negotiations for a development contract with the agency.

On February 11, 2011, we signed a five-year, cost-plus contract with BARDA for the development of 10150 as a MCM against Lung-ARS. BARDA is the government agency responsible for the advanced development and purchase of medical countermeasures for chemical, biological, radiological and nuclear threats. The contract fully funds the advanced development of 10150 through approval by the FDA under 21 CFR Part 314 Subpart I and Part 601 Subpart H (the "Animal Rule.") The Animal Rule allows for approval of drugs using only animal studies when human clinical trials cannot be conducted ethically.

Pursuant to the BARDA Contract we were awarded approximately \$10.4 million for the base period of the contract (from February 2011 to April 2012). On April 16, 2012, we announced that BARDA had exercised two options under the BARDA Contract worth approximately \$9.1 million. On September 17, 2013, we announced that BARDA had exercised \$6.0 million in additional contract options. On May 7, 2014, we announced that BARDA had exercised a Contract Modification worth approximately \$1.8 million. The Contract Modification allowed Aeolus to reconcile actual costs incurred with billings under the original provisional indirect billing rate. It established a new provisional indirect billing rate and placed a cap on the current and future provisional indirect billing rates. On June 26, 2015, we announced that BARDA had exercised \$3.0 million in additional contract options under its advanced research and development contract for 10150. On February 8, 2016, BARDA exercised a Contract Modification worth approximately \$0.1 million. On May 25, 2016, we announced that BARDA had exercised a Contract Modification worth approximately \$0.4 million. The purpose of the Contract Modification was to provide funding to complete a pharmacometric analysis of data from all completed animal efficacy studies of 10150 to determine optimal dose, dose frequency and duration of treatment. The total contract value exercised as of September 30, 2016 is approximately \$30.8 million. We may receive up to an additional \$87.6 million in options exercisable over the remainder of the BARDA Contract. Options are exercised based on the progress of the development program, including the completion of clinical trials or manufacturing tasks under previously exercised options.

Activities under the contract to date include animal efficacy studies, animal model development with radiation survival curve studies, dosing studies, bulk drug manufacturing, bulk drug and final drug product manufacturing, validation testing, compliance studies, stability studies, ADME and metabolic studies, genotoxicity studies and the filing of an orphan drug status application and a fast track designation application with the FDA.

Following the commencement of the BARDA Contract, we entered into a series of agreements with various parties in furtherance of our efforts under the BARDA Contract, which are described below.

On February 14, 2012, we presented the results and deliverables that had been produced during the first twelve months under the base period of the BARDA Contract at an "In-Progress Review" meeting with BARDA, and requested the exercise of additional contract options, which contain additional key items required in the advanced development of 10150.

On February 15, 2012, we announced that we entered into a contract modification and no-cost extension with BARDA. The modification and extension allowed us to continue operating under the base period of the contract awarded in February 2011, and restructured the timing and components of the options that could be awarded under the remaining four years of the agreement. The changes did not impact the total potential value of the contract, which remains at approximately \$118.4 million. The contract restructure was driven by our ability to generate cost savings in the base year contract, and to allow BARDA to better manage contract options to expedite development program.

On April 16, 2012, we announced that BARDA had exercised two contract options worth approximately \$9.1 million. BARDA's exercise of the options was in response to the presentation of the deliverables and progress made under the contract at the meeting on February 14, 2012. Among the key items in the options BARDA exercised are animal efficacy studies, mechanism of action research and manufacturing and process validation work. All of these items build off of work successfully completed during the first twelve months of the contract base period. The contract is designed to produce the data necessary for an approval under the FDA "Animal Rule" and for a potential Emergency Use Authorization (EUA). An approval or EUA would allow the federal government to buy 10150 for the Strategic National Stockpile under Project Bioshield. Project Bioshield is designed to accelerate the research, development, purchase and availability of effective medical countermeasures for the Strategic National Stockpile.

On November 7, 2012, we and the Office of Research and Development of the University of Maryland, Baltimore ("UMB") entered into a Sub-award Agreement, pursuant to which we engaged UMB to, among other things, perform mouse studies supporting the licensure of 10150. Prior to this agreement, our mouse studies had been conducted at

Duke University. In 2012, the research team at Duke responsible for conducting the studies moved to UMB. The Sub-award Agreement is a fixed fee agreement inclusive of all direct and indirect costs. As a result of the contract modification and no-cost extension with BARDA mentioned above, the term of the Sub-award Agreement will continue through at least December 30, 2015.

On July 29, 2013, we presented the results and deliverables that had been produced during the first 28 months of the contract at an "In-Progress Review" meeting with BARDA, and requested the exercise of additional contract options.

On September 17, 2013 we announced that BARDA had exercised \$6.0 million in additional options under the contract. The options that BARDA exercised will fund our IND filing for AEOL-10150 as a treatment for Lung-ARS, additional animal efficacy studies designed to optimize timing and duration of dosing and the continued development of large-scale GMP manufacturing capability to meet potential future demand. When combined with our ongoing studies in non-human primates and our completed work in GMP manufacturing development, these options will help Aeolus meet the requirements for a pre-EUA filing for AEOL-10150 in 2016.

On May 7, 2014, we announced that BARDA had exercised a Contract Modification worth approximately \$1,778,000. The Contract Modification allowed Aeolus to reconcile actual costs incurred with billings under the original provisional indirect billing rate. It established a new provisional indirect billing rate and placed a cap on the current and future provisional indirect billing rates.

On September 4, 2014 we announced positive results from a large study in rhesus macaque monkeys funded by the BARDA Contract. In the study, 10150 was administered beginning 24 hours after exposure to 10.74 Gray of whole thorax radiation. Survival at six months post-exposure in the 60-day treatment group was 50%, compared to 25% survival in the radiation only group. On May 4, 2015, we reported the complete results, including secondary endpoints, from the same study. The study found that administration of 10150 for 60 days beginning 24 hours after exposure to 10.74 Gy of radiation:

- ¶Increased overall survival from 25% in the untreated control group to 50%
- ¶Increased mean and median overall survival time
- ¶Increased mean and median survival time in subjects that did not survive to 180 days
- ¶Increased time to onset of increased respiratory rate, a clinical measure of lung injury
- ¶Decreased mortality in subjects with elevated respiratory rate
- ¶Decreased wet lung weight in all animals, suggesting less parenchymal damage and edema
- ¶Increased SpO₂, a measure of compensated lung function
- ¶Diminished radiographic evidence of pneumonitis and fibrosis during the later stages of the study (days 90 -180)

In addition, a new approach to investigating lipids, metabolites and proteins in pathophysiological process, matrix-assisted laser desorption/ionization mass spectrometry imaging (MALDI-MSI) was employed in the study to measure potential biomarkers for lung injury in tissue samples from study subject. Analysis using MALDI-MSI showed that the molecular profile for the naïve (un-irradiated) lung is statistically distinct from irradiated lung and treatment with 10150 shifts the molecular profile back towards the naïve lung. Two prospective biomarkers found in irradiated, damaged lung tissue were not detectable in either naïve or 10150-treated samples.

On September 22, 2014, we received notice from the FDA that the Phase I safety study in healthy normal volunteers proposed in its IND for Lung-ARS had been placed on clinical hold. On February 22, 2016, we announced that the FDA had removed the clinical hold and allowing us to proceed with the planned Phase I study.

On April 23, 2015, the Company executed a Modification of Contract (the "Modification") with BARDA. The purpose of the Modification is to add funding in the amount of \$1,021,000 for the purpose of funding ongoing murine efficacy, pharmacokinetic and ADME studies and project management.

On June 26, 2015 we announced that BARDA had exercised one option under the BARDA Contract worth approximately \$1,989,000 for the purpose of funding laboratory work suggested by the FDA. The funds will also support ongoing project management.

On February 8, 2016, BARDA exercised a Contract Modification worth approximately \$0.1 million. The purpose of the Contract Modification was to provide additional funding for ongoing drug stability studies.

On May 25, 2016, we announced that BARDA had exercised a Contract Modification worth approximately \$0.4 million. The purpose of the Modification was to provide funding to complete a pharmacometric analysis of data from all completed animal efficacy studies of 10150 to determine optimal dose, dose frequency and duration of treatment.

As of September 30, 2016, the total contract value exercised by BARDA under the BARDA Contract is \$30.8 million.

AEOLUS' Anti-Inflammatory/Anti-Infective Program

We established this program to explore to create novel compounds that both reduce inflammation and combat anti-biotic resistant bacteria, with the goal of addressing unmet medical needs in commercial applications, such as cystic fibrosis, while also potentially providing novel approaches to treating major threats to public health from drug resistant bacteria such as burkholderia cepacia ("BC"), pseudomonas aeruginosa ("PA") and others listed as "Biggest Threats" by the Center for Disease Control and Prevention. We have achieved our initial research objectives and begun to extend our preclinical accomplishments into non-clinical studies, clinical trials and drug development programs.

Our anti-inflammatory/anti-infective program is designed to:

- Utilize a selective detoxification pathway that preserves host defense and limits inflammation
- Create and develop stable, small molecules so that they:
 - o Maintain pathogen killing with less off target damage,
 - o Provide faster resolution of inflammation,
 - o Decrease irreversible injury and oxidative stress, and
 - o Produce candidates that are orally available, stable and inexpensive to manufacture.

Our most advanced anti-infective compound, AEOL 20415, is an orally-available, small molecule that has shown the ability to reduce inflammation and to kill bacteria. A patent application for 20415 was filed in April 2015, and in-vitro studies have been completed to demonstrate 20415's activity against BC and PA. A mouse study testing 20415's anti-infective and anti-inflammatory ability should be completed in the first quarter of 2017. If the data confirms the positive in-vitro activity, we intend to initiate the pre-clinical work necessary to file an IND for this compound with an initial indication of Cystic Fibrosis.

Collaborative and Licensing Arrangements

Duke Licenses

Pursuant to our license agreements with Duke, we have obtained exclusive worldwide rights from Duke to products using antioxidant technology and compounds developed by Dr. Irwin Fridovich and other scientists at Duke. The license from Duke covers, among other items, AEOL11203, AEOL11207 and some of the intellectual property related to 10150. We are obligated under the licenses to pay Duke royalties ranging in the low single digits of net product sales during the term of the Duke licenses, and we must make payments upon the occurrence of certain development milestones in an aggregate amount of up to \$2,000,000. In addition, we are obligated under the Duke licenses to pay patent filing, prosecution, maintenance and defense costs. The Duke licenses are terminable by Duke in the event of breach by us and otherwise expire when the last licensed patent expires.

National Jewish Medical and Research Center and National Jewish Health

We have obtained an exclusive worldwide license from the National Jewish Medical and Research Center ("NJMRC") to develop, make, use and sell products using proprietary information and technology developed under a previous Sponsored Research Agreement within the field of antioxidant compounds and related discoveries. The license from NJMRC covers, among other items, the composition of matter for 10150 and some use patents related to alkylating and vesicant agents. We must make milestone payments to the NJMRC in an aggregate amount of up to \$250,000 upon the occurrence of certain development milestones. Our royalty payment obligations to the NJMRC under this license

agreement are in the low single digits of net product sales. We are also obligated to pay patent filing, prosecution, maintenance and defense costs. This NJMRC license agreement is terminable by the NJMRC in the event of breach and otherwise expires when the last licensed patent expires.

In 2009, we obtained an additional exclusive worldwide license from National Jewish Health to develop, make, use and sell products using proprietary information and technology developed at NJH related to certain compounds as an MCM against mustard gas exposure. Under this license agreement, we must make milestone payments to NJH in an aggregate amount of up to \$500,000 upon the occurrence of certain development milestones. In addition, we must make royalty payments to NJH under this license agreement ranging in the low-single digits as a percentage of all sublicensing fees, milestone payments and sublicense royalties that we receive from sublicenses granted by us pursuant to this license agreement. We are also obligated to pay patent filing, prosecution, maintenance and defense costs. This NJH license agreement is terminable by NJH in the event of breach and otherwise expires when the last licensed patent expires.

In April 2015, we obtained an exclusive worldwide license from National Jewish Health to develop, make, use and sell products using proprietary information and technology developed at NJH with funding from the Company for the treatment of infection caused by bacterial, viral and fungal pathogens. Under this license agreement, we must make milestone payments to NJH in an aggregate amount of up to \$1,000,000 upon the occurrence of certain development milestones. In addition, we must make royalty payments to NJH under this license agreement ranging in the low-single digits as a percentage of all net sales, sublicensing fees, milestone payments and sublicense royalties that we receive from sublicenses granted by us pursuant to this license agreement. We are also obligated to pay patent filing, prosecution, maintenance and defense costs. This NJH license agreement is terminable by NJH in the event of breach and otherwise expires when the last licensed patent expires.

University of Colorado

In April 2015, we obtained an exclusive worldwide license from UC to develop, make, use and sell products using proprietary information and technology developed at UC for the treatment of Parkinson's disease or any other indication. There are no milestone payments under the licensing agreement. In addition, we must make royalty payments to UC under this license agreement ranging in the low-single digits as a percentage of all net sales, sublicensing fees, milestone payments and sublicense royalties that we receive from sublicenses granted by us pursuant to this license agreement. We are also obligated to pay patent filing, prosecution, maintenance and defense costs. This UC license agreement is terminable by UC in the event of breach and otherwise expires when the last licensed patent expires.

Research and Development Expenditures

Expenditures for research and development activities were \$2,881,000 and \$3,509,000 during the years ended September 30, 2016 and 2015, respectively. Research and development expenses for fiscal 2016 and 2015 related primarily to the advancement of our lead compound, 10150.

Manufacturing

We currently do not have the capability to manufacture any of our drug candidates on a commercial scale. Materials for non-clinical and clinical studies are produced under contract with third parties. To date, we have partnered with JMPS for the manufacture of our active pharmaceutical ingredients. JMPS is a 200 year-old company that is a global supplier of active pharmaceutical ingredients, fine chemicals and other specialty chemical products and services to a wide range of chemical and pharmaceutical industry customers and industrial and academic research organizations. JMPS is a leader in the manufacture of metal-based pharmaceutical products.

Commercialization

If BARDA elects to procure 10150 following a pre-EUA application, as described above, or after FDA approval, it may be possible for us to generate significant sales revenue without the need to raise significant funds to build a commercial organization. Depending on the size of those procurements, and assuming the successful development and FDA approval of our compounds in other, non-biodefense indications, we may have sufficient financial resources to internally fund the building of a commercial organization. However, in the event procurements from BARDA are not made, and assuming successful development and FDA approval of one or more of our compounds, to successfully

commercialize our catalytic antioxidant programs, we must seek corporate partners with expertise in commercialization or develop this expertise internally. However, we may not be able to successfully commercialize our catalytic antioxidant technology, either internally or through collaboration with others.

Marketing

Our potential products are being developed for large therapeutic markets. We believe these markets are best approached by partnering with established biotechnology or pharmaceutical companies that have broad sales and marketing capabilities. We are pursuing collaborations of this type as part of our search for development partners. However, we may not be able to enter into any marketing arrangements for any of our products on satisfactory terms or at all.

Biodefense Industry

Market Overview

The market for biodefense countermeasures has grown dramatically as a result of the increased awareness of the threat of global terror activity in the wake of the September 11, 2001 terrorist attacks. The U.S. government is the principal source of worldwide biodefense spending. Most U.S. government spending on biodefense programs is in the form of development funding from NIAID, BARDA and the Department of Defense ("DoD") and procurements of countermeasures by BARDA, the CDC and the DoD. The U.S. government is now the largest source of development and procurement funding for academic institutions and biotechnology companies conducting biodefense research or developing vaccines and immunotherapies directed at potential agents of bioterror or biowarfare.

We analyze the biodefense market in three segments; the United States civilian market, United States military market and non U.S. markets, with the U.S. government funding representing the vast majority of the worldwide market.

U.S. Civilian: The U.S. civilian market includes funds to protect the U.S. population from biological agents and is largely funded by the Project BioShield Act of 2004 ("Project BioShield"). Project BioShield is the U.S. government's largest biodefense initiative. Project BioShield was extended through the Pandemic All Hazards and Preparedness Reauthorization Act of 2013, which authorized BARDA to administer a Special Reserve Fund of \$2.8 billion for MCM procurement.

U.S. Military: The DoD is responsible for the development and procurements of countermeasures for the military segment, which focuses on providing protection for military personnel and civilians who are on active duty.

Non-U.S. Markets: Non-U.S. markets address protection against biowarfare agents for both civilians and military personnel in foreign countries. We anticipate that foreign countries will want to procure biodefense products as they are developed and validated by procurements by the U.S. government.

Project BioShield and the Pandemic and All-Hazards Preparedness Act

Project BioShield became law in 2004 and authorizes procurements of countermeasures for chemical, biological, radiological and nuclear attacks for the SNS, which is a national repository of medical assets and countermeasures designed to provide federal, state and local public health agencies with medical supplies needed to treat those affected by terrorist attacks, natural disasters, industrial accidents and other public health emergencies. Project BioShield provided appropriations of \$5.6 billion to be expended over ten years into a Special Reserve Fund.

The Pandemic and All-Hazards Preparedness Act ("PAHPA"), passed in 2006, established BARDA as the agency responsible for awarding procurement contracts for biomedical countermeasures and providing development funding for advanced research and development in the biodefense arena, and supplements the funding available under Project BioShield for chemical, biological, radiological and nuclear countermeasures, and provides funding for infectious disease pandemics. Funding for BARDA is provided by annual appropriations by Congress. Congress also appropriates annual funding for the CDC for procurements of medical assets and countermeasures for the SNS and for NIAID to conduct biodefense research. This appropriation funding supplements amounts available under Project BioShield.

The Pandemic and All-Hazards Preparedness Reauthorization Act ("PAHPRA"), passed in 2013, extended the programs started under Project BioShield and PAHPA. PAHPRA authorized a \$2.8 billion Special Reserve Fund to be administered by BARDA for the purpose of procuring MCMs for the SNS. Currently, the U.S. government may, at its discretion, purchase critical biodefense products for the SNS prior to FDA approval after the filing of a pre-EUA application.

On an ongoing basis we monitor notices for requests for proposal, grants and other potential sources of government funding that could potentially support the development of our drug candidates. Nevertheless, changes in government budgets, priorities and agendas as well as political pressures could result in a reduction in overall government financial support for the biodefense sector in general and/or specifically the drug candidates we are developing. At any time, the U.S. government may be forced or choose to reduce or delay spending in the biodefense field, which could decrease the likelihood of future government contract awards, the likelihood that the government will exercise its right to extend any of its existing contracts and/or the likelihood that the government would procure products from us. (For further information, see "Risk Factors — Risks Related to Our Dependence on U.S. Government Grants and Contracts — Most of our immediately foreseeable future revenues are contingent upon grants and contracts from the U.S. government and we may not achieve sufficient, if any, revenues from these agreements to attain profitability.") As a result, further development of our drug candidates and ultimate product sales to the government, if any, could be delayed or stopped altogether.

Legislation and Regulation Related to Bioterrorism Counteragents

Because some of our drug candidates are intended for the treatment of diseases that may result from acts of bioterrorism, they may be subject to the specific legislation and regulation described below.

Project BioShield

Project BioShield provides expedited procedures for bioterrorism related procurements and awarding of research grants, making it easier for HHS to quickly commit funds to countermeasure projects. Project BioShield relaxes procedures under the Federal Acquisition Regulation for procuring property or services used in performing, administering or supporting biomedical countermeasure research and development. In addition, if the Secretary of HHS deems that there is a pressing need, Project BioShield authorizes the Secretary to use an expedited award process, rather than the normal peer review process, for grants, contracts and cooperative agreements related to biomedical countermeasure research and development activity.

Under Project BioShield, the Secretary of HHS, with the concurrence of the Secretary of the Department of Homeland Security ("DHS"), and upon the approval of the President, can contract to purchase unapproved countermeasures for the SNS in specified circumstances. Congress is notified of a recommendation for a stockpile purchase after Presidential approval. Project BioShield specifies that a company supplying the countermeasure to the SNS is paid on delivery of a substantial portion of the countermeasure. To be eligible for purchase under these provisions, the Secretary of HHS must determine that there are sufficient and satisfactory clinical results or research data, including data, if available, from preclinical and clinical trials, to support a reasonable conclusion that the countermeasure will qualify for approval or licensing within eight years. Project BioShield also allows the Secretary of HHS to authorize the emergency use of medical products that have not yet been approved by the FDA. To exercise this authority, the Secretary of HHS must conclude that:

- the agent for which the countermeasure is designed can cause serious or life-threatening disease;
- the product may reasonably be believed to be effective in detecting, diagnosing, treating or preventing the disease;
- the known and potential benefits of the product outweigh its known and potential risks;
- and
- there is no adequate alternative to the product that is approved and available.

Although this provision permits the Secretary of HHS to circumvent the FDA approval process, its use would be limited to emergency circumstances.

Safety Act

The Support Anti-Terrorism by Fostering Effective Technologies Act enacted by the U.S. Congress in 2002 (the "Safety Act") creates product liability limitations for qualifying anti-terrorism technologies for claims arising from or related to an act of terrorism. In addition, the Safety Act provides a process by which an anti-terrorism technology may be certified as an "approved product" by the DHS and therefore entitled to a rebuttable presumption that the government contractor defense applies to sales of the product. The government contractor defense, under specified circumstances, extends the sovereign immunity of the United States to government contractors who manufacture a product for the government. Specifically, for the government contractor defense to apply, the government must approve reasonably precise specifications, the product must conform to those specifications and the supplier must warn the government about known dangers arising from the use of the product.

Public Readiness and Emergency Preparedness Act

The Public Readiness and Emergency Preparedness Act enacted by Congress in 2005 (the "PREP Act") provides immunity for manufacturers from all claims under state or federal law for "loss" arising out of the administration or use of a "covered countermeasure." However, injured persons may still bring a suit for "willful misconduct" against the manufacturer under some circumstances. "Covered countermeasures" include security countermeasures and "qualified pandemic or epidemic products." For these immunities to apply, the Secretary of HHS must issue a declaration in cases of public health emergency or "credible risk" of a future public health emergency. We cannot predict whether Congress will fund the relevant PREP Act compensation programs; or whether the necessary prerequisites for immunity would be triggered with respect to our drug candidates.

Competition

General

Competition in the pharmaceutical industry is intense and we expect it to increase. Technological developments in our field of research and development occur at a rapid rate and we expect competition to intensify as advances in this field are made. We will be required to continue to devote substantial resources and efforts to research and development activities. Our most significant competitors, among others, are fully integrated pharmaceutical companies and more established biotechnology companies, which have substantially greater financial, technical, sales, marketing and human resources than we do. These companies may succeed in developing and obtaining regulatory approval for competitive products more rapidly than we can for our drug candidates. In addition, competitors may develop technologies and products that are, or are perceived as being, cheaper, safer or more effective than those being developed by us or that would render our technology obsolete.

In addition, we face significant competition for U.S. government funding for both development and procurements of medical countermeasures for biological, chemical and nuclear threats, diagnostic testing systems and other emergency preparedness countermeasures. The U.S. federal government has currently allocated a significant amount of research funding to the development of countermeasures against the effects of radiation exposure. As a result, there are many drug candidates and products under development as a possible countermeasure against the effects of radiation exposure.

We expect that important competitive factors in our potential product markets will be the relative speed with which we and other companies can develop products, complete the clinical testing and approval processes, and supply commercial quantities of a competitive product to the market. With respect to clinical testing, competition might result in a scarcity of clinical investigators and patients available to test our potential products, which could delay development.

Antioxidants

Several companies have explored the therapeutic potential of antioxidant compounds in numerous indications. Historically, most of these companies have focused on engineered versions of naturally occurring antioxidant enzymes, but with limited success, perhaps because the large size of these molecules makes delivery into the cells difficult. Antioxidant drug research continues at a rapid pace despite previous clinical setbacks.

Patents and Proprietary Rights

We currently license rights to our potential products from third parties. We generally seek patent protection in the United States and other jurisdictions for the potential products and proprietary technology licensed from these third parties. The process for preparing and prosecuting patents is lengthy, uncertain and costly. Patents may not issue on any of the pending patent applications owned by us or licensed by us from third parties. Even if patents issue, the claims allowed might not be sufficiently broad to protect our technology or provide us protection against competitive products or otherwise be commercially valuable. Patents issued to or licensed by us could be challenged, invalidated, infringed, circumvented or held unenforceable. Even if we successfully defend our patents for our products, the costs of defense can be significant.

As of December 20, 2013, our catalytic antioxidant small molecule technology base is described in 12 issued United States patents and five United States pending patent applications. These patents and patent applications belong in whole or in part to Duke or the NJH and are licensed to us. These patents and patent applications cover soluble manganic porphyrins as antioxidant molecules as well as targeted compounds obtained by coupling such antioxidant compounds to molecules that bind to specific extracellular elements. The pending U.S. patent applications and issued U.S. patents include composition of matter claims and method claims for several series of compounds. Corresponding international patent applications have been filed, 88 of which have issued, and one of which has been allowed as of December 20, 2013. Our 12 issued U.S. patents will expire between 2015 and 2023.

On November 22, 2013, we filed a provisional patent application with the United States Patent and Trademark Office entitled "Synthesis and Formulation of Porphyrin Compounds." The filing was for brand new patents for synthesis, formulation and pharmaceutical composition of 10150 and other porphyrin compounds. If granted, these new patents could substantially extend the term of patent protection for 10150 beyond the current composition of matter claims.

On November 21, 2014, a patent application was filed under the Patent Cooperation Treaty (PCT) with the United States Receiving Office (US/RO), thereby preserving the right to seek patent protection in all PCT contracting States.

On April 15, 2015, we announced that we had filed a provisional patent application with the United States Patent Office for a new series of compounds demonstrating anti-microbial and anti-inflammatory action. The compounds were developed by Brian J. Day, PhD at NJH as a result of an Aeolus-sponsored research grant. We also announced that we have obtained a worldwide, exclusive license to develop the compounds from NJH.

On November 2, 2015 we announced that we had received notice of allowance of a composition of matter patent for 11114 from the European Patent Office. On April 18, 2016 we announced that the European Patent Office had issued the composition of matter patent for 11114. On the same date, we announced that we had received a notice of allowance from the Japanese Patent Office for 11114. This new compound has demonstrated neuro-protective activity in models of Parkinson's disease and other indications. The compounds were invented by Brian J. Day, PhD at NJH and Manisha Patel, PhD at UC in collaboration with Aeolus Pharmaceuticals, and have been tested and developed by Dr. Patel under two research grants from MJFF. Aeolus has obtained worldwide, exclusive licenses to develop the compounds from NJH and UC.

In addition to patent protection, we rely upon trade secrets, proprietary know-how and technological advances that we seek to protect, in part, through confidentiality agreements with our collaborative partners, employees and consultants. Our employees and consultants are required to enter into agreements providing for confidentiality and the assignment of rights to inventions made by them while in our service. We also enter into non-disclosure agreements to protect our confidential information furnished to third parties for research and other purposes.

Government Regulation

Our research and development activities and the manufacturing and marketing of our future products are subject to regulation by numerous governmental agencies in the United States and in other countries. The FDA and comparable agencies in other countries impose mandatory procedures and standards for the conduct of clinical trials and the production and marketing of products for diagnostic and human therapeutic use. Before obtaining regulatory approvals for the commercial sale of any of our products under development, we must demonstrate through preclinical studies and clinical trials that the product is safe and efficacious for use in each target indication. The results from preclinical studies and early clinical trials might not be predictive of results that will be obtained in large-scale testing. Our clinical trials might not successfully demonstrate the safety and efficacy of any products or result in marketable products.

The United States system of drug approvals is considered to be the most rigorous in the world. It takes an over ten years for a drug candidate to move through the clinical and approval phases in the United States according to a November 2014 study by the Tufts Center for the Study of Drug Development. Only five in 5,000 drug candidates that enter preclinical testing make it to human testing and only one of those five is approved for commercialization. On average, it costs a company \$1.4 billion in out of pocket costs to get one new drug candidate from the laboratory to United States patients according to the report. In addition, time costs, expected returns that investors forego while the drug is in development, averaged \$1.2 billion.

The steps required by the FDA before new drug products may be marketed in the United States include:

- completion of preclinical studies;
 - the submission to the FDA of a request for authorization to conduct clinical trials on an IND, which must become effective before clinical trials may commence;
- adequate and well-controlled Phase I clinical trials, which typically involves normal, healthy volunteers. The tests study a drug candidate's safety profile, including the safe dosage range. The studies also determine how a drug is absorbed, distributed, metabolized and excreted as well as the duration of its action;
 - adequate and well-controlled Phase II clinical trials which typically involve treating patients with the targeted disease with the drug candidate to assess a drug's effectiveness;
- adequate and well-controlled Phase III clinical trials involving a larger population of patients with the targeted disease are treated with the drug candidate to confirm efficacy of the drug candidate in the treatment of the targeted indication and to identify adverse events;
- submission to the FDA of an NDA; and
- review and approval of the NDA by the FDA before the product may be shipped or sold commercially.

In addition to obtaining FDA approval for each product, each product manufacturing establishment must be registered with the FDA and undergo an inspection prior to the approval of an NDA. Each manufacturing facility and its quality control and manufacturing procedures must also conform and adhere at all times to the FDA's current good manufacturing practices ("cGMP") regulations. In addition to preapproval inspections, the FDA and other government agencies regularly inspect manufacturing facilities for compliance with these requirements. Manufacturers must expend substantial time, money and effort in the area of production and quality control to ensure full technical compliance with these standards.

Preclinical testing includes laboratory evaluation and characterization of the safety and efficacy of a drug and its formulation. Preclinical testing results are submitted to the FDA as a part of an IND, which must become effective prior to commencement of clinical trials. Clinical trials are typically conducted in three sequential phases following submission of an IND. Phase I represents the initial administration of the drug to a small group of humans, either patients or healthy volunteers, typically to test for safety (adverse effects), dosage tolerance, absorption, distribution, metabolism, excretion and clinical pharmacology, and, if possible, to gain early evidence of effectiveness. Phase II involves studies in a small sample of the actual intended patient population to assess the efficacy of the drug for a specific indication, to determine dose tolerance and the optimal dose range and to gather additional information relating to safety and potential adverse effects. Once an investigational drug is found to have some efficacy and an acceptable safety profile in the targeted patient population, Phase III studies are initiated to further establish clinical safety and efficacy of the therapy in a broader sample of the general patient population, in order to determine the overall risk-benefit ratio of the drug and to provide an adequate basis for any physician labeling. During all clinical studies, we must adhere to good clinical practices ("GCPs") standards. The results of the research and product development, manufacturing, preclinical studies, clinical studies and related information are submitted in an NDA to the FDA.

The process of completing clinical testing and obtaining FDA approval for a new drug is likely to take a number of years and require the expenditure of substantial resources. If an application is submitted, there can be no assurance that the FDA will review and approve the NDA. Even after initial FDA approval has been obtained, further studies, including post-market studies, might be required to provide additional data on safety and will be required to gain approval for the use of a product as a treatment for clinical indications other than those for which the product was initially tested and approved. Also, the FDA will require post-market reporting and might require surveillance programs to monitor the side effects of the drug. Results of post-marketing programs might limit or expand the further marketing of the products. Further, if there are any modifications to the drug, including changes in indication, manufacturing process, labeling or a change in manufacturing facility, an NDA supplement might be required to be submitted to the FDA.

The rate of completion of any clinical trials will be dependent upon, among other factors, the rate of patient enrollment. Patient enrollment is a function of many factors, including the size of the patient population, the nature of the trial, the availability of alternative therapies and drugs, the proximity of patients to clinical sites and the eligibility criteria for the study. Delays in planned patient enrollment might result in increased costs and delays, which could have a material adverse effect on us.

Failure to comply with applicable FDA requirements may result in a number of consequences that could materially and adversely affect us. Failure to adhere to approved trial standards and GCPs in conducting clinical trials could cause the FDA to place a clinical hold on one or more studies which would delay research and data collection necessary for product approval. Noncompliance with GCPs could also have a negative impact on the FDA's evaluation of an NDA. Failure to adhere to GMPs and other applicable requirements could result in FDA enforcement action and in civil and criminal sanctions, including but not limited to fines, seizure of product, refusal of the FDA to approve product approval applications, withdrawal of approved applications, and prosecution.

Whether or not FDA approval has been obtained, approval of a product by regulatory authorities in foreign countries must be obtained prior to the commencement of marketing of the product in those countries. The requirements governing the conduct of clinical trials and product approvals vary widely from country to country, and the time required for approval might be longer or shorter than that required for FDA approval. Although there are some procedures for unified filings for some European countries, in general, each country at this time has its own procedures and requirements. There can be no assurance that any foreign approvals would be obtained.

In addition to the regulatory framework for product approvals, we and our collaborative partners must comply with laws and regulations regarding occupational safety, laboratory practices, the use, handling and disposition of radioactive materials, environmental protection and hazardous substance control, and other local, state, federal and foreign regulation. The impact of such regulation upon us cannot be predicted and could be material and adverse. In addition, because some of our drug candidates are intended for the treatment of diseases that may result from acts of bioterrorism, they may be subject to the specific legislation and regulation described above under "Project BioShield and the Pandemic and All-Hazards Preparedness Act" and "Legislation and Regulation Related to Bioterrorism Counteragents."

Regulations Regarding Government Contracting

We are a government contractor in the United States and we may become a government contractor elsewhere which would mean that we would be subject to various statutes and regulations that govern procurements of goods and services by agencies of the United States and other countries, including the Federal Acquisition Regulation. These governing statutes and regulations can impose stricter penalties than those normally applicable to commercial contracts, such as criminal and civil damages liability and suspension and debarment from future government contracting. In addition, pursuant to various statutes and regulations, our government contracts may be subject to unilateral termination or modification by the government for convenience in the United States and elsewhere, detailed auditing requirements and accounting systems, statutorily controlled pricing, sourcing and subcontracting restrictions and statutorily mandated processes for adjudicating contract disputes.

Hazardous Materials and Select Agents

Our development and manufacturing processes involve the use of hazardous materials, including chemicals and radioactive materials, and produce waste products. Accordingly, we are subject to federal, state and local laws and regulations governing the use, manufacturing, storage, handling and disposal of these materials. In addition to complying with environmental and occupational health and safety laws, we must comply with special regulations relating to biosafety administered by the CDC, HHS and the DoD.

Other Regulations

In the United States and elsewhere, the research, manufacturing, distribution, sale and promotion of drug and biological products are subject to regulation by various federal, state and local authorities in addition to the FDA, including the Centers for Medicare and Medicaid Services; other divisions of HHS, such as the Office of Inspector General; the U.S. Department of Justice and individual U.S. Attorney offices within the Department of Justice and state and local governments. For example, sales, marketing and scientific and educational grant programs must comply with the anti-kickback and fraud and abuse provisions of the Social Security Act, the False Claims Act, the privacy provisions of the Health Insurance Portability and Accountability Act and similar state laws. Pricing and rebate programs must comply with the Medicaid rebate requirements of the Omnibus Budget Reconciliation Act of 1990 and the Veterans Health Care Act of 1992. All of these activities are also potentially subject to federal and state consumer protection and unfair competition laws.

CPEC, LLC

We were previously developing bucindolol for the treatment of heart failure, but development was discontinued in 1999. Commercial rights to bucindolol are owned by CPEC, LLC, a limited liability company ("CPEC"), of which we own 35% and Endo Pharmaceuticals (formerly Indevus Pharmaceuticals), Inc. owns 65%.

ARCA biopharma, Inc. is developing bucindolol under the trade name Gencaro as a beta-blocker and mild vasodilator for the treatment of atrial fibrillation in patients with heart failure and left ventricular dysfunction. ARCA biopharma, Inc. is headquartered in Broomfield, Colorado. The future development of Gencaro is uncertain. In the event that Gencaro is approved for sale, however, CPEC may be owed royalties on sales. There can be no guarantees, however, that Gencaro will ever be approved or sold or achieve sufficient revenues to generate royalties.

Employees

At December 20, 2016, we had four full-time employees and no part time employees. None of our employees is represented by a labor union. In addition to our employees, we utilize a team of consultants and subcontractors to perform key functions for us.

Executive Officers

Our executive officers and their ages as of December 20, 2016 were as follows:

Name	Age	Position(s)
David Cavalier	47	Chairman of the Board, Chief Financial Officer
John L. McManus	52	President and Chief Executive Officer

David C. Cavalier has been the Chairman of our Board since April 30, 2004, and became our full time employee in November 2009. In June 2013, he became Chief Financial Officer. Since 2001, he has been a Principal and the Chief Operating Officer of Xmark Opportunity Partners, LLC, a manager of a family of private investment funds. From 1995 to 1996, Mr. Cavalier worked for Tiger Real Estate, a \$785 million private investment fund sponsored by Tiger Management Corporation. Mr. Cavalier began his career in 1994 in the Investment Banking Division of Goldman, Sachs & Co. working on debt and equity offerings for public and private real estate companies. He also serves as the Chairman of the New York Advisory Board of Enterprise Community Partners, a national non-profit organization focused on affordable housing. He received a B.A. from Yale University and an M.Phil. from Oxford University.

John L. McManus. Mr. McManus began as a consultant to Aeolus in June 2005 as President. He became employed as our President and Chief Operating Officer in July 2006 and was appointed President and Chief Executive Officer in March 2007. Mr. McManus, who received his degree in business administration from the University of Southern California in 1986, is the founder and president of McManus Financial Consultants, Inc. ("MFC"), which provides strategic, financial and investor relations advice to senior managements and boards of directors of public companies, including advice on mergers and acquisitions. These companies have a combined value of over \$25 billion. He has served as president of MFC since 1997. In addition, Mr. McManus previously served as Vice President, Finance and Strategic Planning to Spectrum Pharmaceuticals, Inc. (NASDAQ: SPPI), where he had primary responsibility for restructuring Spectrum's operations and finances, including the design of strategic and financial plans to enhance Spectrum's corporate focus, and leading the successful implementation of these plans. The implementation of these plans led to an increase in Spectrum's market value from \$1 million to more than \$125 million at the time of Mr. McManus' departure.

Item 1A. Risk Factors.

You should carefully consider the following information about risks described below, together with the other information contained in this annual report on Form 10-K and in our other filings with the SEC, before you decide to buy or maintain an investment in our common stock. We believe the risks described below are the risks that are material to us as of the date of this annual report. If any of the following risks actually occur, our business, financial condition, 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 part of your investment.

Risks Related to Our Business

We have operated at a loss and will likely continue to operate at a loss for the foreseeable future.

We have incurred significant historical losses and had an accumulated deficit of approximately \$190,189,000 as of September 30, 2016. During the years ended September 30, 2016 and 2015, we incurred losses of \$3,558,000 and \$2,628,000, respectively. Our operating losses have been due primarily to our expenditures for research and development on our drug candidates and for general and administrative expenses and our lack of significant, or sufficient, revenues to offset all of the expenditures. We are likely to continue to incur operating losses until such time, if ever, that we generate significant recurring revenues from product sales, whether to the U.S. government for the Strategic National Stockpile or to the general healthcare community for commercial indications, like oncology, epilepsy or Parkinson's disease. We anticipate it will take a minimum of two years (and possibly longer) for us to generate recurring revenues. We expect that it will take at least that long before the development of any of our licensed, or other current potential, products is completed, marketing approvals are obtained from the FDA and commercial sales of any of these products can begin, or that we might receive a procurement from the U.S. Government following a pre-Emergency Use Authorization application or Animal Rule Approval.

We may need substantial additional funding to continue our operations and may be unable to raise capital when needed, or at all, which would force us to delay, curtail or eliminate our clinical programs and our product development programs.

We may need to raise substantial additional capital to fund human clinical trials and continue our research and development, unless and until we receive a procurement of sufficient size from the U.S. Government for the Strategic National Stockpile. In addition, we may need to raise substantial additional capital to enforce our proprietary rights, defend, in litigation or otherwise, any claims that we infringe third party patents or other intellectual property rights; and commercialize, for non-government related indications, any of our products that may be approved by the FDA or any international regulatory authority.

As of September 30, 2016, we had cash of approximately \$3,155,000. During fiscal year 2016, our monthly cash requirements to operate our business that are not reimbursed under the BARDA Contract were approximately \$206,000. However, this historical cash burn does not include amounts described herein whereby we expect that our own capital will be used to fund part of the future costs of ongoing commercial development programs to the extent they are not funded by the BARDA Contract. To the extent we do not have sufficient cash to fund our working capital

requirements, we may not be able to pay our payables timely, which may cause vendors to cease providing services to us, and we may not be able to advance certain, if not all, development programs.

In order to fund on-going operating cash requirements, or to accelerate or expand our oncology and other programs we will need to raise significant additional funds. We are continuously considering additional strategic and financial options available to us, including public or private equity offerings, debt financings or collaboration arrangements. If we raise additional funds by issuing equity securities, our stockholders will experience dilution of their ownership interest. Debt financings, if available, may involve restrictive covenants and require significant interest payments. If we do not receive additional financing to fund our operations not reimbursed under the BARDA Contract, or if BARDA does not exercise any additional options under the BARDA Contract, and we are unable to raise sufficient capital for operations, we would have to discontinue some or all of our activities, merge with or sell, lease or license some or all of our assets to another company, or cease operations entirely, and our stockholders might lose all or part of their investments.

In addition, if our catalytic antioxidant program shows scientific progress, we will need significant additional funds to move our compounds through continued development and clinical trials. If we are unable to raise the amount of capital necessary, or do not receive a sufficient procurement from the U.S. Government for the Strategic National Stockpile, to complete development and reach commercialization of any of our catalytic antioxidant products, we will need to delay or cease development of one or more of these products or partner with another company for the development and commercialization of these products.

We have a history of operating losses and expect to continue to incur substantial losses and may never become profitable.

We have no products approved for commercialization in the United States or abroad. Our drug candidates are still being developed, and all but 10150 are still in early stages of development. Our drug candidates will require significant additional development, clinical trials, regulatory clearances or approvals by the FDA and additional investment before they can be commercialized in the United States.

Our likelihood of achieving profitability will depend on numerous factors, including success in:

- developing our existing drug candidates and developing and testing new drug candidates;
- protecting our intellectual property;
- establishing our competitive position;
- achieving third-party collaborations;
- receiving regulatory approvals;
- manufacturing and marketing products; and
- receiving government funding and identifying new government funding opportunities.

Many of these factors will depend on circumstances beyond our control. We may not achieve sufficient revenues for profitability. Even if we do achieve profitability, we may not be able to sustain or increase profitability on a quarterly or annual basis in the future. If revenues grow more slowly than we anticipate, or if operating expenses exceed our expectations or cannot be adjusted accordingly, then our business, results of operations, financial condition and cash flows will be materially and adversely affected.

As of September 30, 2016, we had an accumulated deficit of \$190,189,000 from our research, development and other activities. We have not generated material revenues from product sales and do not expect to generate product revenues sufficient to support us for at least several more years. As a result, we may not be successful in obtaining sufficient financing on commercially reasonable terms, or at all. Our requirements for additional capital may be substantial and will be dependent on many factors, including the success of our research and development efforts, our ability to commercialize and market products, our ability to successfully pursue our licensing and collaboration strategy, the receipt of government funding, competing technological developments, costs associated with the protection of our intellectual property and any future change in our business strategy.

Our research and development ("R&D") activities are at an early stage and therefore might never result in viable products.

Our research and development program is in the early stages of development, involves unproven technology, requires significant further R&D and regulatory approvals and is subject to the risks of failure inherent in the development of products or therapeutic procedures based on innovative technologies. These risks include the possibilities that:

- any or all of these proposed products or procedures are found to be unsafe or ineffective or otherwise fail to receive necessary regulatory approvals;
- the proposed products or procedures are not economical to market or do not achieve broad market acceptance;
- third parties hold proprietary rights that preclude us from marketing the proposed products or procedures; and
- third parties market a superior or equivalent product.

Further, the timeframe for commercialization of any product is long and uncertain because of the extended testing and regulatory review process required before marketing approval can be obtained. We may not be able to successfully develop or market any of our proposed products or procedures. If we are not able to successfully market any product, our business will suffer.

Our ability to advance our commercialization objectives may be limited by limitation on our access to capital and the absence of any internal marketing capabilities and external marketing arrangements.

In light of the limited number of employees we currently have, our limited resources, and the early stage of our commercial development efforts, our future success may be limited. In addition, there are significant uncertainties as to our ability to access potential sources of capital, and we may not be able to enter into industry partner collaborations that would provide us with liquidity sources on terms acceptable to us, or at all. These uncertainties may be due to a variety of factors, including evolving or adverse conditions in the pharmaceutical industry or in the economy in general or based on the prospects of our commercial programs. Even if we are successful in obtaining collaboration for our commercial programs, we may have to relinquish rights to technologies, product candidates or markets that we might otherwise develop ourselves. These same risks apply to any attempt to out-license our compounds. Finally, our product candidates are being developed for large therapeutic markets. We believe these markets are best approached by partnering with established biotechnology or pharmaceutical companies that have broad sales and marketing capabilities. We are pursuing marketing collaborations of this type as part of our search for development partners. However, we may not be able to enter into any marketing arrangements for any of our products on satisfactory terms or at all.

If our products are not successfully developed and eventually approved by the FDA, we may be forced to reduce or terminate our operations.

All of our drug candidates are at various stages of development and must be approved by the FDA or similar foreign governmental agencies before they can be marketed. The process for obtaining FDA and foreign regulatory approval is both time-consuming and costly, with no certainty of a successful outcome. This process typically requires extensive preclinical and clinical testing, which may take longer or cost more than we anticipate, and may prove unsuccessful due to numerous factors. Drug candidates that may appear to be promising at early stages of development may not successfully reach the market for a number of reasons. The results of preclinical and initial clinical testing of these drug candidates may not necessarily indicate the results that will be obtained from later or more extensive testing. A number of companies in the pharmaceutical and biotechnology industries have suffered significant setbacks in advanced clinical trials, even after obtaining promising results in earlier trials.

Numerous factors could affect the timing, cost or outcome of our drug development efforts, including the following:

- difficulty in securing research laboratories to conduct research activities;

- difficulty in securing centers to conduct trials;
- difficulty in enrolling patients in conformity with required protocols or projected timelines;
- unexpected adverse reactions by patients in trials;
- difficulty in obtaining clinical supplies of the product;

- changes in the FDA's or other regulatory body's requirements for our testing during the course of that testing;
- inability to generate statistically significant data confirming the efficacy of the product being tested;
- modification of the drug during testing; and
- reallocation of our limited financial and other resources to other clinical programs.

It is possible that none of the products we develop will obtain the regulatory approvals necessary for us to begin commercializing them. The time required to obtain FDA and other approvals is unpredictable but often can take years following the commencement of clinical trials, depending upon the nature of the drug candidate. Any analysis we perform on data from clinical activities is subject to confirmation and interpretation by regulatory authorities, which could delay, limit or prevent regulatory approval. Any delay or failure in obtaining required approvals could have a material adverse effect on our ability to generate revenues from the particular drug candidate and we may not have the financial resources to continue to develop our drug candidates and, as a result, may have to terminate our operations.

If we do not reach the market with our products before our competitors offer products for the same or similar uses, or if we are not effective in marketing our products, our revenues from product sales, if any, will be reduced.

We face intense competition in our development activities. Many of our competitors are fully integrated pharmaceutical companies, biotechnology companies or more established bioterrorism product companies, which have substantially greater financial, technical, sales and marketing and human resources than we do. These companies might succeed in obtaining regulatory approval for competitive products more rapidly than we can for our products. In addition, competitors might develop technologies and products that are less expensive and perceived to be safer or more effective than those being developed by us, which could impair our product development and render our technology obsolete.

We are and expect to remain dependent upon collaborations with third parties for the development of new products, and adverse events involving these collaborations could prevent us from developing and commercializing our drug candidates and achieving profitability.

We currently license from third parties, and do not own, rights under patents and certain related intellectual property for the development of our drug candidates. In addition, we expect to enter into agreements with third parties to license rights to our drug candidates. We might not be able to enter into or maintain these agreements on terms favorable to us, if at all. Further, if any of our current licenses were to expire or terminate, our business, prospects, financial condition and results of operations could be materially and adversely affected.

Our research and development activities rely on technology licensed from third parties, and termination of any of those licenses would result in loss of significant rights to develop and market our products, which would impair our business, prospects, financial condition and results of operations.

We have exclusive worldwide rights to our antioxidant small molecule technology through license agreements with Duke, NJH and UC. Each license generally may be terminated by the licensor if we fail to perform our obligations under the agreement, including obligations to develop the compounds and technologies under license. If terminated, we would lose the right to develop the products, which could adversely affect our business, prospects, financial condition and results of operations. The license agreements also generally require us to meet specified milestones or show reasonable diligence in development of the technology. If disputes arise over the definition of these requirements or whether we have satisfied the requirements in a timely manner, or if any other obligations in the license agreements are disputed by the other party, the other party could terminate the agreement, and we could lose our rights to develop the licensed technology.

If new technology is developed from these licenses, we may be required to negotiate certain key financial and other terms, such as royalty payments, for the licensing of this future technology with these research institutions, and it might not be possible to obtain any such license on terms that are satisfactory to us, or at all.

We now rely, and will continue to rely, heavily on third parties for product and clinical development, manufacturing, marketing and distribution of our products.

We currently depend heavily and will depend heavily in the future on third parties for support in product development, clinical development, manufacturing, marketing and distribution of our products. The termination of some or all of our existing collaborative arrangements, or our inability to establish and maintain collaborative arrangements, could have a material adverse effect on our ability to continue or complete clinical development of our products.

We rely on contract clinical research organizations ("CROs") for various aspects of our clinical development activities including clinical trial monitoring, data collection and data management. As a result, we have had and continue to have less control over the conduct of clinical trials, the timing and completion of the trials, the required reporting of adverse events and the management of data developed through the trial than would be the case if we were relying entirely upon our own staff. Although we rely on CROs to conduct our clinical trials, we are responsible for confirming that each of our clinical trials is conducted in accordance with the investigational plan and protocol. Moreover, the FDA and foreign regulatory agencies require us to comply with GCPs for conducting, recording and reporting the results of clinical trials to assure that the data and results are credible and accurate and that the trial participants are adequately protected. Our reliance on third parties does not relieve us of these responsibilities and requirements.

The third parties on which we rely may have staffing difficulties, may undergo changes in priorities or may become financially distressed, adversely affecting their willingness or ability to conduct our trials. We may experience unexpected cost increases that are beyond our control. Any failure of such CROs to successfully accomplish clinical trial monitoring, data collection and data management and the other services they provide for us in a timely manner and in compliance with regulatory requirements could have a material adverse effect on our ability to complete clinical development of our products and obtain regulatory approval. Problems with the timeliness or quality of the work of a CRO may lead us to seek to terminate the relationship and use an alternate service provider. However, making such changes may be costly and would likely delay our trials, and contractual restrictions may make such a change difficult or impossible. Additionally, it may be difficult to find a replacement organization that can conduct our trials in an acceptable manner and at an acceptable cost.

If BARDA opts not to exercise its options under the BARDA Contract, we would be dependent upon grants from NIH for continued development of 10150 for Lung-ARS, or we would need to curtail our development program in this area significantly and we may be placed at a competitive disadvantage in addressing this market opportunity.

During the fiscal years ended September 30, 2016 and 2015, we received 100% of our revenues from our agreement with BARDA, for the development of 10150 as a MCM against Lung-ARS. These revenues have funded some of our personnel and other R&D costs and expenses. Pursuant to the BARDA Contract we were awarded approximately \$10.4 million for the base period of the contract (from February 2011 to April 2012). On April 16, 2012, we announced that BARDA had exercised two options under the BARDA Contract worth approximately \$9.1 million. On September 17, 2013, we announced that BARDA had exercised \$6.0 million in additional contract options. On May 7, 2014, we announced that BARDA had exercised a Contract Modification worth approximately \$1.8 million. The Contract Modification allowed Aeolus to reconcile actual costs incurred with billings under the original provisional indirect billing rate. It established a new provisional indirect billing rate and placed a cap on the current and future provisional indirect billing rates. On June 26, 2015, we announced that BARDA had exercised \$3.0 million in additional contract options under its advanced research and development contract for 10150. On May 25, 2016, we announced that BARDA had exercised a Contract Modification worth approximately \$0.4 million. The purpose of the Modification was to provide funding to complete a pharmacometric analysis of data from all completed animal efficacy studies of 10150 to determine optimal dose, dose frequency and duration of treatment. The total contract value exercised as of September 30, 2016 is approximately \$30.8 million, of which \$30.6 million has been billed. We may receive up to an additional \$87.6 million in options exercisable over the remainder of the BARDA Contract.

Options are exercised based on the progress of the development program, including the completion of clinical trials or manufacturing tasks under previously exercised options.

Under the terms of the BARDA Contract, BARDA may elect not to exercise some or all of the additional options. Because a significant portion of our current revenues are generated from the BARDA Contract, if BARDA does not exercise its options under the BARDA Contract, our ability to develop 10150 as an MCM for Lung-ARS could be negatively impacted, which could harm our competitive position and materially and adversely affect our business, financial condition and results of operations. In general, we believe that future exercise of options under the contract will depend on successful completion of tasks under exercised options and continued demonstration of efficacy.

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Necessary reliance on the "Animal Rule" in conducting trials is time-consuming and expensive.

To obtain FDA approval for our drug candidate for a bioterrorism indication under current FDA regulations, we are required to utilize animal model studies for efficacy and provide animal and human safety data under the "Animal Rule." For many of the biological and chemical threats, animal models are not yet available, and as such we are developing, or will have to develop, appropriate animal models, which is a time-consuming and expensive research effort. Further, we may not be able to sufficiently demonstrate the animal correlation to the satisfaction of the FDA, as these corollaries are difficult to establish and are often unclear. The FDA may decide that our data are insufficient for approval and require additional preclinical, clinical or other studies, refuse to approve our products, or place restrictions on our ability to commercialize those products. Further, other countries do not, at this time, have established criteria for review and approval of these types of products outside their normal review process; i.e., there is no "Animal Rule" equivalent, and consequently we may not be able to make a submission for marketing approval in foreign countries based on such animal data.

Additionally, few facilities in the U.S. and internationally have the capability to test animals with radiation, nerve agents, or other lethal biotoxins or chemical agents or otherwise assist us in qualifying the requisite animal models. We have to compete with other biodefense companies for access to this limited pool of highly specialized resources. We therefore may not be able to secure contracts to conduct the testing in a predictable timeframe, cost-effectively or at all.

Even if we succeed in commercializing our drug candidates, we may not become profitable and manufacturing problems or side effects discovered at later stages can further increase costs of commercialization.

Any drugs resulting from our research and development efforts may not become commercially available. Even if we succeed in developing and commercializing our drug candidates, we may never generate sufficient or sustainable revenues to enable us to be profitable. Even if effective, a product that reaches the market may be subject to additional clinical trials, changes to or re-approvals of our manufacturing facilities or a change in labeling if we or others identify side effects or manufacturing problems after a product is on the market. This could harm sales of the affected products and could increase the cost and expenses of commercializing and marketing them. It could also lead to the suspension or revocation of regulatory approval for the products.

We and our contract manufacturing organizations ("CMOs") will also be required to comply with the applicable FDA current good manufacturing practice ("cGMP") regulations. These regulations include requirements relating to quality control and quality assurance as well as the corresponding maintenance of records and documentation. Manufacturing facilities are subject to inspection by the FDA. These facilities must be approved to supply licensed products to the commercial marketplace. We and our contract manufacturers may not be able to comply with the applicable cGMP requirements and other FDA regulatory requirements. Should we or our contract manufacturers fail to comply, we could be subject to fines or other sanctions or could be prohibited from marketing any products we develop.

Political or social factors may delay or impair our ability to market our products and our business may be materially adversely affected.

Products developed to treat diseases caused by, or to combat the threat of, bioterrorism will be subject to changing political and social environments. The political and social responses to bioterrorism have been unpredictable. Political or social pressures may delay or cause resistance to bringing our products to market or limit pricing of our products, which would harm our business. Changes to favorable laws, such as Project BioShield, could have a material adverse effect on our business, prospects, financial condition and results of operations.

Legislation limiting or restricting liability for medical products used to fight bioterrorism is new, and we cannot be certain that any such protection will apply to our products or if applied what the scope of any such coverage will be.

The U.S. Public Readiness Act was signed into law in December 2005 (the "Public Readiness Act") and creates general immunity for manufacturers of countermeasures, including security countermeasures (as defined in Section 319F-2(c)(1)(B) of the Public Readiness Act), when the U.S. Secretary of Health and Human Services issues a declaration for their manufacture, administration or use. The declaration is meant to provide general immunity from all claims under state or federal law for loss arising out of the administration or use of a covered countermeasure. Manufacturers are excluded from this protection in cases of willful misconduct. The Secretary of Health and Human Services may not make declarations that would cover any of our other drug candidates or the U.S. Congress may not act in the future to reduce coverage under the Public Readiness Act or it may repeal it altogether.

Upon a declaration by the Secretary of Health and Human Services, a compensation fund would be created to provide "timely, uniform and adequate compensation to eligible individuals for covered injuries directly caused by the administration or use of a covered countermeasure." The "covered injuries" to which the program applies are defined as serious physical injuries or death. Individuals are permitted to bring a willful misconduct action against a manufacturer only after they have exhausted their remedies under the compensation program. A willful misconduct action could be brought against us if an individual(s) has exhausted his or her remedies under the compensation program, which could thereby expose us to liability. Furthermore, the Secretary of Health and Human Services may not issue a declaration under the Public Readiness Act to establish a compensation fund. We may also become subject to standard product liability suits and other third party claims if products we develop which fall outside of the Public Readiness Act cause injury or if treated individuals subsequently become infected or otherwise suffer adverse effects from such products.

Healthcare reform measures and other statutory or regulatory changes could adversely affect our business.

The pharmaceutical and biotechnology industries are subject to extensive regulation, and from time to time legislative bodies and governmental agencies consider changes to such regulations that could have significant impact on industry participants. For example, in light of certain highly-publicized safety issues regarding certain drugs that had received marketing approval, the U.S. Congress is considering various proposals regarding drug safety, including some which would require additional safety studies and monitoring and could make drug development more costly. We are unable to predict what additional legislation or regulation, if any, relating to safety or other aspects of drug development may be enacted in the future or what effect such legislation or regulation would have on our business.

The business and financial condition of pharmaceutical and biotechnology companies are also affected by the efforts of governments, third-party payors and others to contain or reduce the costs of healthcare to consumers. In the United States and various foreign jurisdictions there have been, and we expect that there will continue to be, a number of legislative and regulatory proposals aimed at changing the healthcare system, such as the Affordable Care Act and proposals relating to the reimportation of drugs into the U.S. from other countries (where they are then sold at a lower price) and government control of prescription drug pricing. The roll-out, pendency or approval of such proposals could affect our commercialization efforts and result in a decrease in our share price or limit our ability to raise capital or to obtain strategic collaborations or licenses.

We will need to enter into collaborative arrangements for the manufacturing and marketing of our drug candidates, or we will have to develop the expertise, obtain the additional capital and invest the resources to perform those functions internally.

We do not have the staff or facilities to manufacture or market any of the drug candidates being developed in our catalytic antioxidant program. As a result, we will need to enter into collaborative arrangements to commercialize, manufacture and market products that we expect to emerge from our catalytic antioxidant program, or develop the

expertise within Aeolus. We might not be successful in entering into such third party arrangements on terms acceptable to us, if at all. If we are unable to obtain or retain third-party manufacturing or marketing on acceptable terms, we may be delayed in our ability to commercialize products, which could have a material adverse effect on our business, prospects, financial condition and results of operations. Substantial additional funds and personnel would be required if we needed to establish our own manufacturing or marketing operations. We may not be able to obtain adequate funding or establish these capabilities in a cost-effective or timely manner, which could have a material adverse effect on our business, prospects, financial condition and results of operations.

A failure to obtain or maintain patent and other intellectual property rights would allow others to develop and sell products similar to ours, which could impair our business, prospects, financial condition and results of operations.

The success of our business depends, in part, on our ability to establish and maintain adequate protection for our intellectual property, whether owned by us or licensed from third parties. We rely primarily on patents in the United States and in other key markets to protect our intellectual property. If we do not have adequate patent protection, other companies could develop and sell products that compete directly with ours, without incurring any liability to us. Patent prosecution, maintenance and enforcement on a global basis are time-consuming and expensive, and many of these costs must be incurred before we know whether a product covered by the claims can be successfully developed or marketed.

Even if we expend considerable time and money on patent prosecution, a patent application may never issue as a patent. We can never be certain that we were the first to invent the particular technology or that we were the first to file a patent application for the technology because patent applications in the United States and elsewhere are not typically published for public inspection for at least 18 months from the date when they are filed. It is always possible that a competitor is pursuing a patent for the same invention in the United States as we are and has an earlier invention date. In some jurisdictions outside of the United States, priority of invention is determined by the earliest effective filing date, not the date of invention. Consequently, if a third party pursues the same invention and has an earlier filing date, patent protection outside the United States would be unavailable to us. Also, outside the United States, an earlier date of invention cannot overcome a date of publication that precedes the earliest effective filing date. Accordingly, the patenting of our proposed products would be precluded outside the United States if a prior publication anticipates the claims of a pending application, even if the date of publication is within a year of the filing of the pending application.

Even if patents issue, the patent claims allowed might not be sufficiently broad to offer adequate protection for our technology against competitive products. Patent protection differs from country to country, giving rise to increased competition from other products in countries where patent coverage is either unavailable, weak or not adequately enforced, if enforced at all. Once a patent issues, we still face the risk that others will try to design around our patent or will try to challenge the validity of the patent. The cost of defending against a challenge to one or more of our patents could be substantial and even if we prevailed, there could be no assurance that we would recover damages.

If a third party were to bring an infringement claim against us, we would incur significant costs in our defense; if the claim were successful, we would need to develop non-infringing technology or obtain a license from the successful patent holder, if available.

Our business also depends on our ability to develop and market products without infringing on the proprietary rights of others or being in breach of our license agreements. The pharmaceutical industry is characterized by a large number of patents, patent filings and frequent and protracted litigation regarding patent and other intellectual property rights. Many companies have numerous patents that protect their intellectual property rights. Third parties might assert infringement claims against us with respect to our drug candidates and future products. If litigation were required to determine the validity of a third party's claims, we could be required to spend significant time and financial resources, which could distract our management and prevent us from furthering our core business activities, regardless of the outcome. If we did not prevail in the litigation, we could be required to pay damages, license a third party's technology, which may not be possible on terms acceptable to us, or at all, or discontinue our own activities and develop non-infringing technology, any of which could prevent or significantly delay pursuit of our development activities.

Protection of trade secret and confidential information is difficult, and loss of confidentiality could eliminate our competitive advantage.

In addition to patent protection, we rely on trade secrets, proprietary know-how and confidential information to protect our technology. We use confidentiality agreements with our employees, consultants and collaborators to maintain the proprietary nature of this technology. However, confidentiality agreements can be breached by the other party, which would make our trade secrets and proprietary know-how legally available for use by others. There is generally no adequate remedy for breach of confidentiality obligations. In addition, the competitive advantage afforded by trade secrets is limited because a third party can independently discover or develop something identical to our own trade secrets or know-how, without incurring any liability to us.

In addition, if our current or former employees, consultants or collaborators were to use information improperly obtained from others (even if unintentional), we may be subject to claims as to ownership and rights in any resulting know-how or inventions.

If we cannot retain or hire qualified personnel or maintain our collaborations, our programs could be delayed and may be discontinued.

As of September 30, 2016, we had four full-time employees. We utilize consultants to assist with our operations and are highly dependent on the services of our executive officers. We do not maintain "key person" life insurance on any of our personnel. We also are dependent on our collaborators for our research and development activities. The loss of key executive officers or collaborators could delay progress in our research and development activities or result in their termination entirely.

We believe that our future success will depend in large part upon our ability to attract and retain highly skilled scientific and managerial personnel. We face intense competition for these kinds of personnel from other companies, research and academic institutions, government entities and other organizations. If we fail to identify, attract and retain personnel, we may be unable to continue the development of our drug candidates, which would have a material adverse effect on our business, prospects, financial condition and results of operations.

We face the risk of product liability claims that could exceed our insurance coverage and deplete our cash resources.

The pharmaceutical and biotechnology industries expose us to the risk of product liability claims alleging that use of our drug candidates caused an injury or harm. These claims can arise at any point in the development, testing, manufacture, marketing or sale of pharmaceutical products and may be made directly by patients involved in clinical trials of our products, by consumers or healthcare providers or by organizations selling our products. Product liability claims can be expensive to defend, even if the product did not actually cause the alleged injury or harm.

Insurance covering product liability claims becomes increasingly expensive as a product candidate moves through the development pipeline to commercialization. We have limited product liability insurance coverage for our clinical trials and this coverage may not be sufficient to cover us against some or all potential losses due to liability, if any, or to the expenses associated with defending against liability claims. A product liability claim successfully asserted against us could exceed our insurance coverage, require us to use our own cash resources and have a material adverse effect on our business, financial condition and results of operations.

In addition, some of our licensing and other agreements with third parties require or might require us to maintain product liability insurance. If we cannot maintain acceptable amounts of coverage on commercially reasonable terms in accordance with the terms set forth in these agreements, the corresponding agreements would be subject to termination.

The costs of compliance with environmental, safety and similar laws could increase our cost of doing business or subject us to liability in the event of noncompliance.

Our business is subject to regulation under state and federal laws regarding occupational safety, laboratory practices, environmental protection and the use, generation, manufacture, storage and disposal of hazardous substances. We may be required to incur significant costs in the future to comply with existing or future environmental and health and safety regulations. Our research activities involve the use of hazardous materials, chemicals and radioactive compounds. Although we believe that our procedures for handling such materials comply with applicable state and federal regulations, we cannot eliminate the risk of contamination or injury from these materials. In the event of contamination, we could be liable for any resulting damages, which could have a material adverse effect on our business, financial condition and results of operations.

We are subject to intense competition that could materially impact our operating results.

We may be unable to compete successfully against our current or future competitors. The pharmaceutical, biopharmaceutical, biotechnology and bioterrorism industries are characterized by intense competition and rapid and significant technological advancements. Many companies, research institutions and universities are working in a number of areas similar to our primary fields of interest to develop new products. There also is intense competition among companies seeking to acquire products that already are being marketed. Many of the companies with which we compete have or are likely to have substantially greater research and product development capabilities and financial, technical, scientific, manufacturing, marketing, distribution and other resources than at least some of our present or future strategic partners or licensees.

As a result, these competitors may:

- succeed in developing competitive products sooner than us or our strategic partners or licensees;
- obtain FDA and other regulatory approvals (including EUA approvals) for their products before approval of any of our products;
- obtain patents that block or otherwise inhibit the development and commercialization of our drug candidates;
- develop products that are safer or more effective than our products;
- devote greater resources to marketing or selling their products;
- introduce or adapt more quickly to new technologies or scientific advances;
- introduce products that render our products obsolete;
- withstand price competition more successfully than us or our strategic partners or licensees;
- negotiate third-party strategic alliances or licensing arrangements more effectively; or
- take advantage of other opportunities more readily.

Currently, as discussed under "Competition" there are no drugs approved for Lung-ARS. Two drugs, Ebsriet (Roche) and OFEV (Boehringer-Ingelheim) are approved for use in treating IPF. There is one drug, Amifostine, approved for use in reducing side effects from radiation therapy. However, there are also many companies working to develop pharmaceuticals that act as radiation protection agents and as treatments for IPF.

Acceptance of our products in the marketplace is uncertain, and failure to achieve market acceptance will harm our business.

Even if approved for marketing, our products may not achieve market acceptance. The degree of market acceptance will depend upon a number of factors, including:

- the receipt of regulatory approvals for the indications that we are studying;
- the establishment and demonstration in the medical community of the safety, clinical efficacy and cost-effectiveness of our products and their potential advantages over existing therapeutic products;
- marketing and distribution support;
- the introduction, market penetration and pricing strategies of competing and future products; and
- coverage and reimbursement policies of governmental and other third-party payors such as insurance companies, health maintenance organizations and other plan administrators.

Physicians, patients, payors or the medical community in general may be unwilling to accept, purchase, utilize or recommend any of our products.

We may be required to make milestone payments and other payments relating to the commercialization of our products.

Our agreements by which we acquired rights to our drug candidates provide for milestone payments by us upon the occurrence of certain regulatory filings and approvals related to the acquired products. In the event that we successfully develop our drug candidates, these milestone payments could be significant. In addition, our agreements require us to pay a royalty interest on worldwide sales. Also, any future license, collaborative or other agreements we may enter into in connection with our development and commercialization activities may require us to pay significant milestone, license and other payments in the future.

We are continually evaluating our business strategy, and may modify this strategy in light of developments in our business and other factors.

We continue to evaluate our business strategy and, as a result, may modify this strategy in the future. In this regard, we may, from time to time, focus our development efforts on different drug candidates or may delay or halt the development of our drug candidates. In addition, as a result of changes in our strategy, we may also change or refocus our existing drug discovery, development, commercialization and manufacturing activities.

Our short-term investments, marketable securities and restricted investments, if any, are subject to certain risks which could materially adversely affect our overall financial position.

We invest our cash in accordance with an established internal policy and customarily in instruments which historically have been highly liquid and carried relatively low risk. We maintain a portfolio of investments in short-term investments, marketable debt securities and restricted investments, which are recorded at fair value. Certain of these transactions expose us to credit risk in the event of default of the issuer. To minimize our exposure to credit risk, we invest in securities with strong credit ratings. Should any of our short-term investments, marketable securities or restricted investments lose value or have their liquidity impaired, it could materially and adversely affect our overall financial position by imperiling our ability to fund our operations and forcing us to seek additional financing sooner than we would otherwise. Such financing may not be available on commercially attractive terms or at all.

Our insurance policies are expensive and protect us only from some business risks, which could leave us exposed to significant, uninsured liabilities.

We do not carry insurance for all categories of risk that our business may encounter. We currently maintain general liability, property, auto, workers' compensation, products liability, fiduciary and directors' and officers' insurance policies. We do not know, however, if we will be able to maintain existing insurance with adequate levels of coverage. For example, the premiums for our directors' and officers' insurance policy have increased in the past and may increase in the future, and this type of insurance may not be available on acceptable terms or at all in the future. Any significant uninsured liability may require us to pay substantial amounts, which would adversely affect our cash position and results of operations.

We may have a limitation on the use of net operating loss carryforwards and tax credits.

Our ability to utilize our net operating loss carryforwards, or NOLs, and tax credits may be limited if we undergo or have undergone an ownership change, as defined in Section 382 of the Internal Revenue Code, as a result of changes in the ownership of outstanding stock. An ownership change generally occurs if the percentage of stock owned by one or more stockholders who own, directly or indirectly, 5% or more of the value of our outstanding stock (or are otherwise treated as 5% stockholders under Section 382 and the regulations promulgated thereunder) has increased by more than 50 percentage points over the lowest percentage of our outstanding stock owned by these stockholders at any time during the testing period, which is generally the three-year period preceding the potential ownership change. In the event of an ownership change, Section 382 imposes an annual limitation on the amount of post-ownership change taxable income a corporation may offset with pre-ownership change NOLs.

We are exposed to risks if we are unable to comply with changes to laws affecting public companies, including the Sarbanes-Oxley Act of 2002 and the Dodd-Frank Wall Street Reform and Consumer Protection Act of 2010, and also to increased costs associated with complying with such laws.

Laws and regulations affecting public companies in the U.S., including the provisions of the Sarbanes-Oxley Act of 2002 and the Dodd-Frank Wall Street Reform and Consumer Protection Act of 2010, will cause us to incur increased costs as we evaluate the implications of new rules and respond to new requirements. Delays or a failure to comply

with the new laws, rules and regulations could result in enforcement actions, the assessment of other penalties and civil suits. These laws and regulations make it more expensive for us under indemnities provided by us to our officers and directors and may make it more difficult for us to obtain certain types of insurance, including liability insurance for directors and officers; as such, we may be forced to accept reduced policy limits and coverage or incur substantially higher costs to obtain the same or similar coverage. The impact of these events could also make it more difficult for us to attract and retain qualified persons to serve on our board of directors, or as executive officers. We may be required to hire additional personnel and utilize additional outside legal, accounting and advisory services — all of which could cause our general and administrative costs to increase beyond what we currently have planned.

Our corporate compliance program cannot guarantee that we are in compliance with all potentially applicable regulations.

The development, manufacturing, pricing, sales, coverage and reimbursement of our products, together with our general operations, are subject to extensive regulation by federal, state and other authorities within the United States and numerous entities outside of the United States. While we have developed and instituted a corporate compliance program based on what we believe are the current best practices, governmental authorities may not find that our business practices comply with current or future administrative or judicial interpretations of potentially applicable laws and regulations. If we fail to comply with any of these laws and regulations, we could be subject to a range of regulatory actions, including suspension or termination of clinical trials, the failure to approve a product candidate, restrictions on our products or manufacturing processes, withdrawal of products from the market, significant fines, or other sanctions or litigation.

Risks Related to Our Dependence on U.S. Government Grants and Contracts

Even with the BARDA Contract, we may not be able to fully fund our research and development of 10150 as a MCM for Lung-ARS.

The BARDA Contract is a cost-plus-fixed-fee reimbursement contract that only reimburses certain specified activities that have been previously authorized by BARDA. Additional activities may be needed and, if so, BARDA may not reimburse us for these activities. Performance under the BARDA Contract requires that we comply with appropriate regulations and operational mandates. Our ability to be regularly and fully reimbursed for our activities will depend on our ability to comply and demonstrate compliance with such requirement.

The BARDA Contract award does not guarantee that we will be successful in future clinical trials or that 10150 will be approved by the FDA.

The BARDA Contract provides a cost-plus-fixed-fee reimbursement opportunity for certain specified clinical and development activities, but we remain fully responsible for conducting these activities. The award of BARDA Contract does not guarantee that any of these activities will be successful. Our inability to be successful with certain key clinical or development activities could jeopardize our ability to obtain FDA approval for 10150.

Most of our immediately foreseeable future revenues are contingent upon grants, contracts or purchases from the U.S. government and we may not achieve sufficient, if any, revenues from these agreements to attain profitability.

For the foreseeable future, we believe our main customer, if any, will be national governments, primarily the U.S. government. However, we may not ultimately be successful in receiving any future revenues or grants from such governments. The process of obtaining government contracts is lengthy and uncertain and we may have to compete with other companies for each contract. We may not be awarded any contracts to supply the U.S. or other governments with our drug candidates or products as such awards may be made, in whole or in part, to our competitors. If the U.S. government makes significant future contract awards for the supply to the Strategic National Stockpile of a competing product, our business will be harmed and it is unlikely that we will ultimately be able to supply that particular treatment or product to foreign governments or other third parties. Further, changes in government budgets and agendas, or advances by our competitors, may result in a decreased and de-prioritized emphasis on procuring the biodefense products we are developing.

The U.S. government's determination to award any contracts may be challenged by an interested party, such as another bidder, at the Government Accountability Office ("GAO") or in federal court. If such a challenge is successful, a contract may be terminated.

The laws and regulations governing procurements of goods and services by the U.S. government provide procedures by which other bidders and other interested parties may challenge the award of a government contract. If we are awarded a government contract, such challenges or protests could be filed even if there are not any valid legal grounds on which to base the protest. If any such protests are filed, the government agency may decide to suspend our performance under the contract while such protests are being considered by the GAO or the applicable federal court, thus potentially delaying delivery of goods and services and payment. In addition, we could be forced to expend considerable funds to defend any potential award. If a protest is successful, the government may be ordered to terminate the contract and reselect bids. The government could even be directed to award a potential contract to one of the other bidders.

Our business may become subject to audit by the U.S. government and a negative audit could adversely affect our business.

U.S. government agencies such as the Defense Contract Audit Agency (the "DCAA"), routinely audit and investigate government contractors. These agencies review a contractor's performance under its contracts, cost structure and compliance with applicable laws, regulations and standards.

The DCAA also reviews the adequacy of, and a contractor's compliance with, its internal control systems and policies, including the contractor's purchasing, property, estimating, compensation and management information systems. Any costs found to be improperly allocated to a specific contract will not be reimbursed, while such costs already reimbursed must be refunded. If an audit uncovers improper or illegal activities, we may be subject to civil and criminal penalties and administrative sanctions, including:

- termination of contracts;
- forfeiture of profits;
- suspension of payments;
- fines; and
- suspension or prohibition from conducting business with the U.S. government.

In addition, we could suffer serious reputational harm if allegations of impropriety were made against us.

Laws and regulations affecting government contracts make it more costly and difficult for us to successfully conduct our business.

We must comply with numerous laws and regulations relating to the formation, administration and performance of government contracts, which can make it more difficult for us to retain our rights under these contracts. These laws and regulations affect how we conduct business with government agencies. Among the most significant government contracting regulations that affect our business are:

- the Federal Acquisition Regulations, and agency-specific regulations supplemental to the Federal Acquisition Regulations, which comprehensively regulate the procurement, formation, administration and performance of government contracts;
- the business ethics and public integrity obligations, which govern conflicts of interest and the hiring of former government employees, restrict the granting of gratuities and funding of lobbying activities and incorporate other requirements such as the Anti-Kickback Act and Foreign Corrupt Practices Act;
- export and import control laws and regulations; and
- laws, regulations and executive orders restricting the use and dissemination of information classified for national security purposes and the exportation of certain products and technical data.

Foreign governments typically also have laws and regulations governing contracts with their respective agencies. These foreign laws and regulations could affect how we conduct business and, in some instances, impose added costs on our business. Any changes in applicable laws and regulations could restrict our ability to obtain contracts, which could limit our ability to conduct our business and materially adversely affect our revenues and results of operations.

Because we depend on clinical research centers and other contractors for clinical and non-clinical testing, including testing under the "Animal Rule", and for certain research and development activities, the results of our clinical trial, non-clinical animal efficacy studies, and research and development activities are largely beyond our control.

The nature of studies, clinical trials and our business strategy of outsourcing substantially all of our research and development and manufacturing work require that we rely on clinical research centers and other contractors to assist us with research and development, clinical and non-clinical testing (including animal efficacy studies under the "Animal Rule"), patient enrollment and other activities. As a result, our success depends largely on the success of these third parties in performing their responsibilities. Although we prequalify our contractors and believe that they are fully capable of performing their contractual obligations, we cannot directly control the adequacy and timeliness of the resources and expertise that they apply to these activities. Furthermore, we have to compete with other biodefense companies for access to this limited pool of highly specialized resources. If our contractors do not perform their obligations in an adequate and timely manner or we are unable to enter into contracts with them because of prior commitments to our competitors, the pace of clinical or non-clinical development, regulatory approval and commercialization of our drug candidates could be significantly delayed and our prospects could be adversely affected.

Data obtained from clinical trials is susceptible to varying interpretations, which could delay, limit or prevent regulatory clearances.

Data already obtained, or obtained in the future, from pre-clinical studies, non-clinical studies and clinical trials does not necessarily predict the results that will be obtained from later pre-clinical studies and clinical trials. Moreover, pre-clinical and clinical data are susceptible to varying interpretations, which could delay, limit or prevent regulatory approval. A number of companies in the pharmaceutical industry have suffered significant setbacks in advanced clinical trials, even after promising results in earlier trials. The failure to adequately demonstrate the safety and effectiveness of an intended product under development could delay or prevent regulatory clearance of the drug candidate, which would result in delays to commercialization and could materially harm our business. Our studies and clinical trials may not demonstrate sufficient levels of safety and efficacy necessary to obtain the requisite regulatory approvals for our drugs, and our proposed drugs may not be approved for marketing.

We may encounter delays or rejections based on additional government regulation from future legislation or administrative action or changes in FDA policy during the period of development, clinical trials and FDA regulatory review. We may encounter similar delays in foreign countries. If any of our products are approved for commercialization, sales of the products outside the U.S. would be subject to foreign regulatory approvals that vary from country to country. The time required to obtain approvals from foreign countries may be shorter or longer than that required for FDA approval, and requirements for foreign licensing may differ from FDA requirements. We may be unable to obtain requisite approvals from the FDA or foreign regulatory authorities, and even if obtained, such approvals may not be on a timely basis, or they may not cover the uses that we request.

Even if we do ultimately receive FDA approval for any of our drug candidates, these drug candidates will be subject to extensive ongoing regulation, including regulations governing manufacturing, labeling, packaging, testing, dispensing, prescription and procurement quotas, record keeping, reporting, handling, shipment and disposal of any such drug. Failure to obtain and maintain required registrations or to comply with any applicable regulations could further delay or preclude development and commercialization of our drugs and subject us to enforcement action.

Unfavorable provisions in government contracts, some of which may be customary, may harm our business, financial condition and operating results.

Government contracts customarily contain provisions that give the government substantial rights and remedies, many of which are not typically found in commercial contracts, including provisions that allow the government to:

- terminate existing contracts, in whole or in part, for any reason or no reason;

- unilaterally reduce or modify contracts or subcontracts, including equitable price adjustments;
- cancel multi-year contracts and related orders if funds for contract performance for any subsequent year become unavailable;
- decline to exercise an option to renew a contract;
- exercise an option to purchase only the minimum amount specified in a contract;
- decline to exercise an option to purchase the maximum amount specified in a contract;
- claim rights to products, including intellectual property, developed under the contract;

- take actions that result in a longer development timeline than expected;
- audit and object to the contractor's contract-related costs and fees, including allocated indirect costs;
- direct the course of a development program in a manner not chosen by the government contractor;
- suspend or debar the contractor from doing business with the government or a specific government agency;
- pursue criminal or civil remedies under the False Claims Act and False Statements Act; and
- control or prohibit the export of products.

Generally, government contracts contain provisions permitting unilateral termination or modification, in whole or in part, at the government's convenience. Under general principles of government contracting law, if the government terminates a contract for convenience, the terminated company may recover only its incurred or committed costs, settlement expenses and profit on work completed prior to the termination.

If the government terminates a contract for default, the defaulting company is entitled to recover costs incurred and associated profits on accepted items only and may be liable for excess costs incurred by the government in procuring undelivered items from another source. Some government contracts grant the government the right to use, for or on behalf of the U.S. government, any technologies developed by the contractor under the government contract. If we were to develop technology under a contract with such a provision, we might not be able to prohibit third parties, including our competitors, from using that technology in providing products and services to the government.

Risks Related to Owning Our Stock

Our principal stockholders own a significant percentage of our outstanding common stock and are, and will continue to be, able to exercise significant influence over our affairs.

As of September 30, 2016, Xmark Opportunity Partners, LLC ("Xmark") possessed voting power over 97,214,694 shares, or 63.9%, of our outstanding common stock as of such date, through its management of Goodnow Capital, L.L.C. ("Goodnow"), Xmark Opportunity Fund, L.P., Xmark Opportunity Fund, Ltd. and Xmark JV Investment Partners, LLC (collectively, the "Xmark Funds"). As a result, Xmark is able to determine a significant part of the composition of our board of directors, holds significant voting power with respect to matters requiring stockholder approval and is able to exercise significant influence over our operations. The interests of Xmark may be different than the interests of other stockholders on these and other matters. This concentration of ownership also could have the effect of delaying or preventing a change in our control or otherwise discouraging a potential acquirer from attempting to obtain control of us, which could reduce the price of our common stock.

David Cavalier, an employee and our Chairman of the board of directors and Mitchell D. Kaye, a Director, are affiliated with Xmark, which possessed voting power of 63.9% of our outstanding common stock as of September 30, 2016. Accordingly, Mr. Cavalier and Mr. Kaye currently have, and will continue to have, a significant influence over the outcome of all corporate actions requiring stockholder approval.

Our executive officers and directors and holders of greater than five percent of our outstanding common stock, together with entities that may be deemed affiliates of, or related to, such persons or entities, beneficially owned greater than 74% of our outstanding common stock as of September 30, 2016. As a result, these stockholders, acting together, may be able to control our management and affairs and matters requiring stockholder approval, including the election of directors and approval of significant corporate transactions, such as mergers, consolidations or the sale of substantially all of our assets. The interests of our current major stockholders may not always coincide with the interests of other stockholders and they may take actions to advance their respective interests to the detriment of other stockholders.

We may need to sell additional shares of our common stock, preferred stock or other securities to meet our capital requirements and these future sales could cause dilution and adversely affect our stock price.

Sales of substantial amounts of capital stock, or the perception that such sales could occur, could adversely affect the prevailing market price of the common stock and our ability to raise capital. We may issue additional common stock in future financing transactions or as incentive compensation for our executive management and other key personnel, consultants and advisors. Issuing any equity securities would be dilutive to the equity interests represented by our then-outstanding shares of common stock. The market price for our common stock could decrease as the market takes into account the dilutive effect of any of these issuances.

In the event of the conversion of our preferred stock and exercises of currently outstanding options and warrants, the ownership interests of our current stockholders could be substantially diluted, which would reduce the market price of our common stock and could make it more difficult for us to raise funds in the future.

As of September 30, 2016, we had 152,085,825 shares of common stock outstanding. We may grant to our employees, directors and consultants, options to purchase shares of our common stock under our 2016 Stock Incentive Plan. In addition, as of September 30, 2016, options to purchase 12,204,000 shares were outstanding at exercise prices ranging from \$0.19 to \$1.19 per share, with a weighted average exercise price of \$0.38 per share, and 25,000,000 shares were reserved for issuance under the 2016 Stock Incentive Plan. In addition, as of September 30, 2016, warrants to purchase 52,947,877 shares of common stock were outstanding at exercise prices ranging from \$0.22 to \$0.49 per share, with a weighted exercise price of \$0.23 per share.

Substantial blocks of our total outstanding common stock may be sold into the market through this registration statement. If there are substantial sales of shares of our common stock, the price of our common stock could decline.

The price of our common stock could decline if there are substantial sales of our common stock, particularly sales by our directors, executive officers and significant stockholders. As of the date of this prospectus, substantially all of our outstanding common stock and substantially all common stock underlying outstanding warrants and Series C Convertible Preferred Stock are covered by this registration statement and/or otherwise freely tradeable without restrictions or further registration under the Securities Act. The market price of our common stock could decline as a result of the sale of a substantial number of our common stock or due to the perception in the market that the holders of a large number of shares intend to sell their common stock.

Our common stock is not listed on a national securities exchange, is illiquid and is characterized by low and/or erratic trading volume, and the intraday per share price of our common stock has fluctuated from \$0.15 to \$0.50 during the last two fiscal years.

Our common stock is quoted on the OTCQB under the symbol "AOLS." An active public market for our common stock is unlikely to develop as long as we are not listed on a national securities exchange. Even if listed, the market for our stock may be impaired because of the limited number of investors, the significant ownership stake of Xmark, and our small market capitalization, which is less than that authorized for investment by many institutional investors.

Historically, the public market for our common stock has been characterized by low and/or erratic trading volume, often resulting in price volatility. For the fiscal year ended September 30, 2016 the average daily trading volume for our common stock was approximately 30,000 shares. In addition, the price of our common stock has been volatile. Our common stock had a closing price of \$0.25 on October 1, 2015 and ended fiscal year 2016 at a closing price of \$0.19. During the twelve month period ended September 30, 2016, our common stock had a low closing price of \$0.15, which occurred on July 5, 2016, and had a high closing price of \$0.30, which occurred on November 30, 2015.

The market price of our common stock is subject to wide fluctuations due to factors that we cannot control, including the results of preclinical and clinical testing of our products under development, decisions by collaborators regarding product development, regulatory developments, market conditions in the pharmaceutical and biotechnology industries, future announcements concerning our competitors, adverse developments concerning proprietary rights, public concern as to the safety or commercial value of any products and general economic conditions.

Furthermore, the stock market has experienced significant price and volume fluctuation unrelated to the operating performance of particular companies. These market fluctuations can adversely affect the market price and volatility of our common stock.

Anti-takeover provisions in our charter documents and under Delaware law could make an acquisition of us, which may be beneficial to our stockholders, more difficult and may prevent attempts by our stockholders to replace or remove our current management.

Provisions in our amended and restated certificate of incorporation and bylaws may delay or prevent an acquisition of us or a change in our management. These provisions include a prohibition on certain actions by written consent of our stockholders and the ability of our board of directors to issue up to 7,150,000 shares of "blank check" preferred stock without stockholder approval. As a result, our board of directors has the power to issue shares without stockholder approval, and such shares can be issued with such rights, preferences, and limitations as may be determined by our board of directors. The rights of the holders of common stock will be subject to, and may be adversely affected by, the rights of any holders of preferred stock that may be issued in the future. In addition, because we are incorporated in Delaware, we are governed by the provisions of Section 203 of the General Corporation Law of the State of Delaware, which prohibits stockholders owning in excess of 15% of our outstanding voting stock from merging or combining with us. These provisions may frustrate or prevent any attempts by our stockholders to replace or remove our current management by making it more difficult for stockholders to replace members of our board of directors, which is responsible for appointing the members of our management.

We do not expect to pay cash dividends on our common stock for the foreseeable future.

We have never paid cash dividends on our common stock and do not anticipate that any cash dividends will be paid on the common stock for the foreseeable future. The payment of any cash dividend by us will be at the discretion of our board of directors and will depend on, among other things, our earnings, capital, regulatory requirements and financial condition. Furthermore, the terms of some of our financing arrangements directly limit our ability to pay cash dividends on our common stock.

Item 1B. Unresolved Staff Comments.

None.

Item 2. Properties.

None.

Item 3. Legal Proceedings.

From time to time we are subject to various lawsuits and claims with respect to matters arising out of the normal course of our business. Due to the uncertainty of the ultimate outcome of these matters, the impact on future financial results is not subject to reasonable estimates. As of December 20, 2016, the Company was not a party to any lawsuits or subject to any claims.

Item 4. Mine Safety Disclosures.

Not applicable.

PART II

Item 5. Market for Registrant's Common Equity, Related Stockholder Matters and Issuer Purchases of Equity Securities.

Price Range of Common Stock

Our common stock is traded on the OTC Bulletin Board under the symbol "AOLS." The following sets forth the quarterly high and low trading prices as reported by the OTC Bulletin Board for the periods indicated. These prices are based on quotations between dealers, which do not reflect retail mark-up, markdown or commissions, and do not necessarily represent actual transactions.

	High	Low
Fiscal Year Ended September 30, 2015		
October 1, 2014 through December 31, 2014	\$0.29	\$0.20
January 1, 2015 through March 31, 2015	\$0.35	\$0.21
April 1, 2015 through June 30, 2015	\$0.41	\$0.28
July 1, 2015 through September 30, 2015	\$0.32	\$0.22

Fiscal Year Ended September 30, 2016

October 1, 2015 through December 31, 2015	\$0.30	\$0.17
January 1, 2016 through March 31, 2016	\$0.25	\$0.16
April 1, 2016 through June 30, 2016	\$0.20	\$0.15
July 1, 2016 through September 30, 2016	\$0.50	\$0.15

Approximate Number of Equity Security Holders

As of December 20, 2016 the number of record holders of our common stock was 125, and we estimate that the number of beneficial owners was approximately 4,200.

Dividends

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

Moreover, any additional preferred stock to be issued and any future credit facilities might contain restrictions on our ability to declare and pay dividends on our common stock. We plan to retain all earnings, if any, for the foreseeable future for use in the operation of our business and to fund future growth.

Recent Sales of Unregistered Securities

The securities issued in the 2015 Securities Placement were offered and sold in reliance upon exemptions from registration pursuant to Section 4(2) under the Securities Act and Rule 506 promulgated thereunder. The securities purchase agreements contained representations to support the Company's reasonable belief that, among other things, the purchasers acquired the securities for their own account and not with a view to the distribution thereof, and that each purchaser was an "accredited investor" as such term is defined in Regulation D promulgated under the Securities Act. The securities issued in the 2015 Securities Placement are deemed to be restricted securities for purposes of the Securities Act and the certificates representing the securities sold in connection with the Private Placement shall bear legends to that effect. Accordingly, the securities sold in connection with the 2015 Securities Placement may not be offered or sold in the United States absent registration or an applicable exemption from registration requirements.

Purchases of Equity Securities by the Issuer and Affiliated Purchasers

None.

Performance Graph

We are a smaller reporting company, as defined by Rule 12b-2 of the Securities Exchange Act of 1934, as amended, and are not required to provide the information required under this item.

Item 6. Selected Financial Data.

Not applicable.

Item 7. Management's Discussion and Analysis of Financial Condition and Results of Operations.

Introduction

You should read the following discussion in conjunction with our consolidated financial statements and the notes appearing elsewhere in this Annual Report on Form 10-K. The following discussion contains forward-looking statements that involve risks and uncertainties. Our actual results could differ materially from those anticipated in the forward-looking statements as a result of various factors, including those discussed in Item 1A - "Risk Factors" and elsewhere in this Annual Report on Form 10-K.

Overview

We are developing a new class of catalytic antioxidant compounds as medical countermeasures ("MCM") against nuclear, radiological and chemical weapons as well as for diseases and disorders of the respiratory system, central nervous system and oncology.

Our lead compound, AEOL 10150 ("10150"), is being developed:

- as a MCM for the pulmonary and delayed effects of acute radiation exposure under contract with the U.S. Government
- as a MCM for exposure to chemical gas and nerve gas exposure with grant funding from the U.S. Government
- as a treatment for idiopathic pulmonary fibrosis
- as a treatment for the side effects of radiation therapy for solid tumors in oncology

Our other development programs include:

- development AEOL 11114B ("11114") as a treatment for Parkinson's disease
- development of AEOL 20415 ("20415") as a treatment for infectious disease

We have reported positive safety results from two Phase I clinical trials of 10150 with no serious adverse events noted.

We had a net loss of \$3,558,000 and \$2,628,000 for the fiscal years ended September 30, 2016 and 2015, respectively. We had an accumulated deficit of approximately \$190,189,000 at September 30, 2016. We have not yet generated any revenue from product sales and we cannot provide assurances that we will receive any product revenue from non-government sales in the foreseeable future, if at all.

We have not had any recurring revenue from product sales. Therefore, we have relied on public or private equity offerings, debt financings, collaboration arrangements and grants to finance our operations.

Corporate Matters

On February 11, 2011, we signed a five-year, cost-plus contract with BARDA for the development of 10150 as a MCM against Lung-ARS (the "BARDA Contract"). BARDA is the government agency responsible for the advanced development and purchase of medical countermeasures for chemical, biological, radiological and nuclear threats. The contract fully funds the advanced development of 10150 through approval by the FDA under 21 CFR Part 314 Subpart I and Part 601 Subpart H (the "Animal Rule.") The Animal Rule allows for approval of drugs using only animal studies when human clinical trials cannot be conducted ethically.

Pursuant to the BARDA Contract we were awarded approximately \$10.4 million for the base period of the contract (from February 2011 to April 2012). On April 16, 2012, we announced that BARDA had exercised two options under the BARDA Contract worth approximately \$9.1 million. On September 17, 2013, we announced that BARDA had exercised \$6.0 million in additional contract options. On May 7, 2014, we announced that BARDA had exercised a Contract Modification worth approximately \$1.8 million. The Contract Modification allowed Aeolus to reconcile actual costs incurred with billings under the original provisional indirect billing rate. It established a new provisional indirect billing rate and placed a cap on the current and future provisional indirect billing rates. On June 26, 2015, we announced that BARDA had exercised \$3.0 million in additional contract options under its advanced research and development contract for 10150. On May 25, 2016, we announced that BARDA had exercised a Contract Modification worth approximately \$0.4 million. The purpose of the Modification was to provide funding to complete a pharmacometric analysis of data from all completed animal efficacy studies of 10150 to determine optimal dose, dose frequency and duration of treatment. The total contract value exercised as of September 30, 2016 is approximately \$30.8 million, of which \$30.6 million has been billed. We may receive up to an additional \$87.6 million in options exercisable over the remainder of the BARDA Contract. Options are exercised based on the progress of the development program, including the completion of clinical trials or manufacturing tasks under previously exercised options.

Research and development activities under the contract to date include animal efficacy studies, animal model development with radiation survival curve studies, dosing studies, bulk drug manufacturing, bulk drug and final drug product manufacturing, validation testing, compliance studies, stability studies and the filing of an orphan drug status application and a fast track designation application with the FDA.

Among the key deliverables accomplished in the program, we delivered a final report on the survival data from our efficacy study in non-human primates, prepared a Fast Track filing for AEOL 10150 in Lung-ARS, completed murine extended natural history, duration of treatment and delayed treatment studies, completed studies to identify biomarkers for use in Lung-ARS studies, completed additional pharmacokinetic pharmacodynamics work and completed an absorption, distribution, metabolism and excretion ("ADME") study in the new formulation for AEOL 10150, manufactured pilot scale batches of bulk drug substance and final drug product, and generated stability data on the new bulk drug substance and new final drug product.

In the event BARDA exercises the remaining options under the contract, we expect to conduct, among other things, bulk drug and final drug product manufacturing, stability studies, animal pivotal efficacy studies, human clinical safety studies and Phase I, Phase II and pre-new drug application ("NDA") meetings and applications with the FDA.

For additional developments under the BARDA Agreement preceding our 2015 fiscal year, please see Item 1 "Business – BARDA Contract; Background and Recent Developments."

Results of Operations

Fiscal Year Ended September 30, 2016 Compared to Fiscal Year Ended September 30, 2015

We had a net loss of \$3,558,000 for the fiscal year ended September 30, 2016, versus a net loss of \$2,628,000 for the fiscal year ended September 30, 2015.

Revenue for the fiscal year ended September 30, 2016 was approximately \$2,076,000, compared to \$3,111,000 revenue for the fiscal year ended September 30, 2015. The revenue is from the collaboration with BARDA announced on February 11, 2011. Since being awarded the BARDA Contract, we generate contract revenue from a cost-plus fee arrangement. Revenues on reimbursable contracts are recognized as costs are incurred, based on allowable costs incurred during the period, plus any recognizable earned fee. We consider fixed fees under cost-plus fee contracts to be earned in proportion to the allowable costs incurred in performance of the contract. Revenue was higher in the prior year primarily due to the timing of work related to the BARDA contract.

Research and Development

Research and development expenses decreased by \$628,000, or 18%, to approximately \$2,881,000 for the fiscal year ended September 30, 2016 from approximately \$3,509,000 for the fiscal year ended September 30, 2015. R&D expenses were lower during the fiscal year ended September 30, 2016 versus September 30, 2015 due to the timing of work related to the BARDA Contract. For the fiscal year ended September 30, 2016, Preclinical fees decreased by \$1,087,000 primarily due to costs associated with the BARDA Contract. Consultant expenses increased about \$118,000 in 2016 over the comparable period in 2015 due to a \$173,000 increase in non-BARDA related work offset by a \$55,000 decrease in costs associated with the BARDA contract. Patent expenses increased about \$269,000. We currently have 8 development programs in progress: studies of 10150 as a medical countermeasure against the effects of radiation on the lungs, against the effects of sulfur mustard gas and chlorine gas on the lungs, against the effects of radiation on the lungs, against the effects of nerve gas exposure, as a treatment for cancer, as a treatment for IPF and studies of 11114B, 11203 and 11207 as potential treatments for Parkinson's disease and epilepsy.

R&D expenses for our antioxidant program have totaled approximately \$65,439,000 from inception through September 30, 2016. Because of the uncertainty of our research and development and clinical studies, we are unable to predict the total level of spending on the program or the program completion date. Future R&D expenses will be determined primarily by the exercise of options under the BARDA contract and the possible initiation of human clinical studies in oncology. We anticipate that much of the R&D expense, except for the possible oncology studies, will be reimbursed by the BARDA Contract.

General and Administrative

General and administrative ("G&A") expenses include corporate costs required to support Aeolus, our employees and consultants and our stockholders. These costs include personnel and outside costs in the areas of legal, human resources, investor relations and finance. Additionally, we include in general and administrative expenses such costs as rent, repair and maintenance of equipment, utilities, information technology and procurement costs that we need to support the corporate functions listed above.

G&A expenses increased approximately \$240,000, or 11%, to approximately \$2,468,000 for the fiscal year ended September 30, 2016 from about \$2,228,000 for the fiscal year ended September 30, 2015. Legal expense increased by about \$102,000 as a result of higher SEC filing and financing costs. Board of Director fees increased by about \$56,000.

Other Income or Expense

Other Expense increased approximately \$283,000 to approximately \$285,000 for the fiscal year ended September 30, 2016, from \$2,000 for the fiscal year ended September 30, 2015 as a result of amortization of discount on note payable to shareholders.

Liquidity and Capital Resources

As of September 30, 2016, we had approximately \$3,155,000 of cash and cash equivalents, an increase of \$3,061,000 from September 30, 2015. The increase in cash was primarily due to our 2015 capital raise, which was offset by cash used in operations. In order to fund on-going operating cash requirements, or to accelerate or expand our oncology and other programs, we may need to raise significant additional funds.

We had a net loss of \$3,558,000 for the fiscal year ended September 30, 2016, compared to a net loss of \$2,628,000 for the fiscal year ended September 30, 2015. For the same periods, we had cash outflows from operations of approximately \$3,109,000 and \$2,443,000, respectively, with the outflows increasing in 2016 due to lower revenue from BARDA.

Our ongoing future cash requirements will depend on numerous factors, particularly the progress of our development programs, clinical trials and our ability to negotiate and complete collaborative agreements or out-licensing arrangements. In order to help fund our on-going operating cash requirements, we intend to seek new collaborations for our antioxidant research program that include initial cash payments and on-going research support. In addition, we might sell additional shares of our stock and/or convertible debentures and explore other strategic and financial alternatives, including a merger with another company, the sale of stock and/or debt, the establishment of new collaborations for current research programs that include initial cash payments and ongoing research support and the out-licensing of our compounds for development by a third party. We expect to incur additional losses and negative cash flow from operations for several more years.

Under the BARDA Contract, substantially all of the costs of the development of 10150 as a medical countermeasure for pulmonary injuries resulting from an acute exposure to radiation from a radiological/nuclear accident or attack, particularly injuries associated with ARS or DEARE could be paid for by the U.S. government through BARDA funding. We recognized approximately \$2,076,000 in revenue during the fiscal year ended September 30, 2016 related to the BARDA Contract.

We do not have any revenues from product sales and, therefore, we rely on investors, grants and collaborations to finance our operations. We generate limited revenue from reimbursable, cost-plus fee R&D contracts and grants. Revenues on reimbursable contracts are recognized as costs are incurred, generally based on allowable costs incurred during the period, plus any recognizable earned fee. We consider fixed fees under cost-plus fee contracts to be earned in proportion to the allowable costs incurred in performance of the contract.

Since the terms of the BARDA Contract include provisions to cover some general corporate overhead as well as a small provision for profit, the result on our liquidity is that our cash burn is related to the level of non-BARDA contract activities that we pursue. In order to fund on-going operating cash requirements, or to further accelerate or expand our programs, we expect to need to raise significant additional funds in order to pursue non-BARDA contract programs, including our oncology program.

There are significant uncertainties as to our ability to access potential sources of capital. We may not be able to enter into any collaboration on terms acceptable to us, or at all, due to conditions in the pharmaceutical industry or in the economy in general or based on the prospects of our catalytic antioxidant program. Even if we are successful in obtaining collaboration for our antioxidant program, we may have to relinquish rights to technologies, product candidates or markets that we might otherwise develop ourselves. These same risks apply to any attempt to out-license our compounds.

Similarly, due to market conditions, the illiquid nature of our stock and other possible limitations on equity offerings, we may not be able to sell additional securities or raise other funds on terms acceptable to us, if at all. Any additional equity financing, if available, could result in substantial dilution to existing stockholders.

Our forecast of the period of time through which our financial resources will be adequate to support our operations is forward-looking information, and actual results could vary.

2015 Capital Raise

On December 10, 2015, we completed the 2015 Securities Placement by entering into securities purchase agreements with certain accredited investors to sell and issue (i) an aggregate of 10,215,275 common units issued at a purchase price of \$0.22 per unit, and (ii) 4,500 preferred stock units issued to existing investors, Biotechnology Value Fund, L.P. and other affiliates of BVF Partners, L.P., for an aggregate purchase price of \$4.5 million, resulting in aggregate gross proceeds to the Company of approximately \$6.75 million. Each common unit consists of one share of the Company's common stock and a five year warrant to purchase one share of the Company's common stock, subject to adjustment. The preferred units collectively consist of (i) 4,500 shares of Series C Convertible Preferred Stock of the Company that are collectively convertible into an aggregate of 20,454,546 common shares and (ii) warrants to purchase an aggregate of 20,454,546 Common Shares, in each case subject to adjustment. The warrants have an initial exercise price of \$0.22 per share. On September 29, 2015, the Company received funding in the form of the BVF Notes. On December 11, 2015, following the completion of the 2015 Securities Placement, the principal and accrued interest amounts under the BVF Notes were converted into 5,414,402 shares of the Company's common stock and warrants to purchase an additional 5,414,402 shares of the Company's common stock at an exercise price per share of \$0.22 subject to adjustment. As a result, the BVF Notes were no longer outstanding as of December 11, 2015. Although the foregoing transactions provide us with additional liquidity, there can be no assurance that such funds will be sufficient to enable us to advance our development objectives. See "Risk Factors -- We may need substantial additional funding to continue our operations and may be unable to raise capital when needed, or at all, which would force us to delay, curtail or eliminate our clinical programs and our product development programs."

Off Balance Sheet Arrangements

We do not have any off-balance sheet arrangements that have or are reasonably likely to have a current or future effect on our financial condition, changes in financial condition, revenues or expenses, results of operations, liquidity, capital expenditures or capital resources as defined under the rules of SEC Release No. FR-67. We do not have any capital leases.

Relationship with Goodnow Capital, LLC and Xmark Opportunity Partners, LLC

In July 2003, we initiated a series of transactions that led to our corporate reorganization and recapitalization. We obtained an aggregate of \$8,000,000 in secured bridge financing in the form of convertible promissory notes we issued to Goodnow Capital, LLC ("Goodnow"). A portion of this financing allowed us to pay our past due payables and become current. We used the remainder for our operations, including a toxicology study for our catalytic antioxidant compounds under development as a treatment for ALS.

We completed our corporate reorganization on November 20, 2003. The reorganization involved the merger of our former parent company into one of our wholly owned subsidiaries. Subsequent to our 2003 reorganization, we completed a number of equity and debt financings, the majority of which included Xmark as investors. As of September 30, 2016, Xmark Opportunity Partners, LLC, through its management of Goodnow and the Xmark Funds, and through the Xmark Voting Trust and options held by David Cavalier, an affiliate of Xmark and the Chairperson of our Board of Directors, had voting power over 63.9% of our outstanding common stock and had beneficial ownership, calculated based on SEC requirements, of approximately 64.1% of our common stock. As a result of this significant ownership, Xmark Opportunity Partners, LLC and its affiliates is able to control future actions voted on by our stockholders.

In addition, under the terms of the warrants to purchase up to 61,822,749 shares of our common stock issued to Xmark on October 6, 2009 as well as subsequent warrant issuances on July 30, 2010, August 11, 2010 and December 28, 2010 (collectively, the "Xmark Warrants"), if we were to pay a dividend on our common stock the exercise price of these warrants would be reset from \$0.28 per share or \$0.50 per share, as applicable, to \$0.01 per share and the warrant holders would also receive any such dividend paid. The Xmark Warrants also contain a provision that provides for the reduction of the exercise price to \$0.01 upon a change of control and anti-dilution provisions in the event of a stock dividend or split, dividend payment or other issuance, reorganization, recapitalization or similar event. In addition, the Xmark Warrants, among other restrictions, prohibit the sale of Aeolus to an entity other than one that is publicly traded.

Effective February 19, 2013, the Company and each of Xmark JV Investment Partners, LLC, Xmark Opportunity Fund, Ltd. and Xmark Opportunity Fund, L.P. (collectively, the "Xmark Entities") entered into a Warrant Repricing, Exercise and Lockup Agreement (the "Xmark Warrant Agreement") pursuant to which the Company agreed to reduce the exercise price of outstanding warrants to purchase an aggregate of up to 59,149,999 shares of Common Stock held by the Xmark Entities (the "Xmark Warrants") to \$0.01 per share. In consideration for the reduction of the exercise price of the Xmark Warrants, each of the Xmark Entities agreed to immediately exercise all of the Xmark Warrants by cashless exercise. The Xmark Warrant Agreement also provides that the Xmark Entities will not transfer the shares issuable upon exercise of the Xmark Warrants (the "Xmark Warrant Shares") until the Company either (i) declares a cash dividend on its common stock or otherwise makes a cash distribution or (ii) effects a Change of Control, subject in each case to the terms of the Xmark Warrant Agreement.

Critical Accounting Policies and Estimates

Our consolidated financial statements have been prepared in accordance with accounting principles generally accepted in the United States of America, which require us to make estimates and judgments that affect the reported amounts of assets, liabilities, revenues, expenses and related disclosure of contingent assets and liabilities. We evaluate our estimates, judgments and the policies underlying these estimates on a periodic basis as the situation changes, and regularly discuss financial events, policies, and issues with our independent registered public accounting firm and members of our audit committee. We routinely evaluate our estimates and policies regarding revenue recognition; clinical trial, preclinical, manufacturing and patent related liabilities; license obligations; inventory; intangible assets; share-based payments; and deferred tax assets.

We generally enter into contractual agreements with third-party vendors to provide clinical, preclinical and manufacturing services in the ordinary course of business. Many of these contracts are subject to milestone-based invoicing and the contract could extend over several years. We record liabilities under these contractual commitments when we determine an obligation has been incurred, regardless of the timing of the invoice. Patent-related liabilities are recorded based upon various assumptions or events that we believe are the most reasonable to each individual circumstance, as well as based upon historical experience. License milestone liabilities and the related expense are recorded when the milestone criterion achievement is probable. We have not recognized any assets for inventory, intangible items or deferred taxes as we have yet to receive regulatory approval for any of our compounds. Any potential asset that could be recorded in regards to any of these items is fully reserved. In all cases, actual results may differ from our estimates under different assumptions or conditions.

Revenue Recognition

We do not currently generate revenue from product sales, but do generate revenue from the BARDA Contract. We recognize revenue from the BARDA Contract in accordance with the authoritative guidance for revenue recognition. Revenue is recognized when all of the following criteria are met: (i) persuasive evidence of an arrangement exists, (ii) delivery (or passage of title) has occurred or services have been rendered, (iii) the seller's price to the buyer is fixed or determinable, and (iv) collectability is reasonably assured. We also comply with the authoritative guidance for revenue recognition regarding arrangements with multiple deliverables.

The BARDA Contract is classified as a "cost-plus-fixed-fee" contract. We recognize government contract revenue in accordance with the authoritative guidance for revenue recognition including the authoritative guidance specific to federal government contracts. Reimbursable costs under the contract primarily include direct labor, subcontract costs, materials, equipment, travel, and indirect costs. In addition, we receive a fixed fee under the BARDA Contract, which is unconditionally earned as allowable costs are incurred and is not contingent on success factors. Reimbursable costs under this BARDA Contract, including the fixed fee, are generally recognized as revenue in the period the reimbursable costs are incurred and become billable. We receive regular updates from our subcontractors regarding estimated completion of individual projects. Management evaluates the status of each project with respect to budgeted work completed, actual work completed, and cost of actual work completed. We are required to provide BARDA with monthly reports in addition to our bi-weekly conference calls with BARDA regarding the progress of each project.

Item 7A. Quantitative and Qualitative Disclosures About Market Risk.

Market Risk

Our exposure to market risk is presently limited to the interest rate sensitivity of our cash and cash equivalents, which is affected by changes in the general level of U.S. interest rates. However, we believe that we are not subject to any material market risk exposure and do not expect that changes in interest rates would have a material effect upon our financial position. A hypothetical 10% change in interest rates would not have a material effect on our Statement of Operations or Cash Flows for the fiscal year ended September 30, 2016. We do not have any foreign currency or other derivative financial instruments as of September 30, 2016.

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Item 8. Financial Statements and Supplementary Data.

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REPORT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

Board of Directors and Shareholders

Aeolus Pharmaceuticals, Inc.

We have audited the accompanying consolidated balance sheet of Aeolus Pharmaceuticals, Inc. (the "Company") as of September 30, 2016, and the related consolidated statements of operations, stockholders' equity (deficit), and cash flows for the year ended September 30, 2016. 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. We were not engaged to perform an audit of the Company's 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 consolidated financial statements referred to above present fairly, in all material respects, the consolidated financial position of Aeolus Pharmaceuticals, Inc. as of September 30, 2016, and the consolidated results of its operations and its cash flows for the year ended September 30, 2016 in conformity with accounting principles generally accepted in the United States of America.

/s/ HASKELL & WHITE LLP

Irvine, California

December 20, 2016

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REPORT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

Board of Directors and Shareholders

Aeolus Pharmaceuticals, Inc.

We have audited the accompanying consolidated balance sheets of Aeolus Pharmaceuticals, Inc. (the "Company") as of September 30, 2015, and the related consolidated statements of operations, stockholders' equity (deficit), 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.

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In our opinion, the consolidated financial statements referred to above present fairly, in all material respects, the financial position of Aeolus Pharmaceuticals, Inc. as of September 30, 2015, and the results of their operations and their cash flows for the year then ended in conformity with accounting principles generally accepted in the United States of America.

/s/ GRANT THORNTON LLP

San Diego, California

December 18, 2015

AEOLUS PHARMACEUTICALS, INC.
CONSOLIDATED BALANCE SHEETS
(DOLLARS IN THOUSANDS)

	September 30,	
	2016	2015
ASSETS		
Current assets:		
Cash and cash equivalents	\$3,155	\$94
Accounts receivable	750	1,585
Deferred subcontractor cost	—	21
Prepaid expenses and other current assets	230	45
Total current assets	4,135	1,745
Investment in CPEC LLC	32	32
Total assets	\$4,167	\$1,777
LIABILITIES AND STOCKHOLDERS' EQUITY (DEFICIT)		
Current liabilities:		
Accounts payable and accrued expenses	\$972	\$1,598
Deferred revenue	—	22
Notes payable to shareholders, net of debt discount of \$273	—	727
Notes payable to shareholders redemption liability	—	275
Total current liabilities	972	2,622
Total liabilities	972	2,622
Commitments and Contingencies (Notes E, J, K, and L)		
Stockholders' equity (deficit):		
Preferred stock, \$0.01 par value per share, 10,000,000 shares authorized:		
Series A nonredeemable convertible preferred stock, 1,250,000 shares authorized as of September 30, 2016 and 2015, respectively; no shares issued and outstanding as of September 30, 2016 and 2015, respectively		
	—	—
Series B nonredeemable convertible preferred stock, 1,600,000 and 1,600,000 shares authorized as of September 30, 2016 and 2015, respectively; zero and 526,080 shares issued and outstanding as of September 30, 2016 and 2015, respectively		
	—	5
Series C nonredeemable convertible preferred stock, 5,000 and zero shares authorized as of September 30, 2016 and September 30, 2015, respectively; 4,500 and zero shares issued and outstanding as of September 30, 2016 and September 30, 2015, respectively		
	—	—
Common stock, \$0.01 par value per share, 200,000,000 shares authorized; 152,085,825 and 135,930,068 shares issued and outstanding at September 30, 2016 and 2015, respectively		
	1,520	1,359
Additional paid-in capital	191,863	184,421
Accumulated deficit	(190,188)	(186,630)
Total stockholders' equity (deficit)	3,195	(845)
Total liabilities and stockholders' equity (deficit)	\$4,167	\$1,777

The accompanying notes are an integral part of these consolidated financial statements.

AEOLUS PHARMACEUTICALS, INC.
CONSOLIDATED STATEMENTS OF OPERATIONS
(IN THOUSANDS, EXCEPT PER SHARE DATA)

	Fiscal Year Ended September 30,	
	2016	2015
Revenue:		
Contract revenue	\$2,076	\$3,111
Costs and expenses:		
Research and development	2,881	3,509
General and administrative	2,468	2,228
Total costs and expenses	5,349	5,737
Loss from operations	(3,273)	(2,626)
Interest expense	285	2
Net loss	(3,558)	(2,628)
Deemed dividend on Series C preferred stock	2,486	—
Net loss attributable to common stockholders	\$(6,044)	\$(2,628)
Net loss attributable to common stockholders – basic	\$(6,044)	\$(2,628)
Net loss attributable to common stockholders – diluted	\$(6,044)	\$(2,628)
Basic net loss per common share	\$(0.04)	\$(0.02)
Diluted net loss per common share	\$(0.04)	\$(0.02)
Weighted average common shares outstanding:		
Basic	148,668	135,883
Diluted	148,668	135,883

The accompanying notes are an integral part of these consolidated financial statements.

AEOLUS PHARMACEUTICALS, INC.

CONSOLIDATED STATEMENTS OF STOCKHOLDERS' EQUITY (DEFICIT)

(DOLLARS IN THOUSANDS)

	Series B Pref. Stock Shares	Par Value	Series C Pref. Stock Shares	Par Value	Common Stock Shares	Par Value	Additional Paid-in Capital	Accumulated Deficit	Stockholders' Deficit (Equity)
Balance at									
September 30, 2014	526,080	\$ 5	—	\$ —	135,850,068	\$1,359	\$184,223	\$(184,002)	\$ 1,585
Exercise of warrants	—	—	—	—	80,000	—	20	—	20
Issuance of warrants to consultants	—	—	—	—	—	—	14	—	14
Stock-based compensation	—	—	—	—	—	—	164	—	164
Net loss for the fiscal year ended									
September 30, 2015	—	—	—	—	—	—	—	(2,628)	(2,628)
Balance at									
September 30, 2015	526,080	5	—	—	135,930,068	1,359	184,421	(186,630)	(845)
Common stock sales, net of issuance costs of \$242	—	—	—	—	10,215,275	102	1,903	—	2,005
Preferred stock sales, net of issuance costs of \$335	—	—	4,500	—	—	—	4,165	—	4,165
Conversion of notes payable to shareholders	—	—	—	—	5,414,402	54	1,233	—	1,287
Conversion of Series B Preferred to common stock	(526,080)	(5)	—	—	526,080	5	—	—	—
Stock-based compensation	—	—	—	—	—	—	141	—	141
Net loss for the fiscal year end September 30, 2016									
September 30, 2016	—	—	—	—	—	—	—	(3,558)	(3,558)
Balance at September 30, 2016	—	\$ —	4,500	\$ —	152,085,825	\$1,520	\$191,863	\$(190,188)	\$ 3,195

The accompanying notes are an integral part of these consolidated financial statements.

AEOLUS PHARMACEUTICALS, INC.
CONSOLIDATED STATEMENTS OF CASH FLOWS
(IN THOUSANDS)

	Fiscal Year Ended September 30,	
	2016	2015
Cash flows from operating activities:		
Net loss	\$(3,558)	\$(2,628)
Adjustments to reconcile net loss to net cash used in operating activities:		
Amortization of discount on notes payable to shareholders	285	—
Accrued interest	—	2
Noncash compensation	141	178
Change in assets and liabilities:		
Accounts receivable	835	(26)
Deferred subcontractor cost	21	405
Prepaid expenses and other current assets	(185)	1
Accounts payable and accrued expenses	(626)	46
Deferred revenue	(22)	(421)
Net cash used in operating activities	(3,109)	(2,443)
Cash flows from financing activities:		
Proceeds from exercise of common stock warrants	—	20
Proceeds from issuance of notes payable to shareholders	—	1,000
Proceeds from issuance of common stock and common stock warrants, net	2,005	—
Proceeds from issuance of preferred stock and preferred stock warrants, net	4,165	—
Net cash provided by financing activities	6,170	1,020
Net increase (decrease) in cash and cash equivalents	3,061	(1,423)
Cash and cash equivalents at beginning of year	94	1,517
Cash and cash equivalents at end of year	\$3,155	\$94
Supplemental disclosure of non-cash financing activities:		
Conversion of notes payable to shareholders for common stock and warrants	\$1,000	\$—
Conversion of accrued interest on notes payable to shareholders for common stock and warrants	\$12	\$—
Issuance of warrants for financing costs	\$266	\$—
Deemed dividend on Series C preferred stock	\$2,486	\$—
Notes payable to shareholders redemption liability	\$—	\$275

The accompanying notes are an integral part of these consolidated financial statements.

AEOLUS PHARMACEUTICALS, INC.
NOTES TO CONSOLIDATED FINANCIAL STATEMENTS
SEPTEMBER 30, 2016 and 2015

A. Organization and Business

Organization

The accompanying audited consolidated financial statements include the accounts of Aeolus Pharmaceuticals, Inc. and its wholly-owned subsidiary, Aeolus Sciences, Inc. (collectively "we," "us," "Company" or "Aeolus"). All significant intercompany accounts and transactions have been eliminated in consolidation. Aeolus is a Delaware corporation. The Company's primary operations are located in Mission Viejo, California.

Business

Aeolus is developing a new class of broad-spectrum, catalytic antioxidant compounds based on technology discovered at Duke University and National Jewish Health. The Company's lead compound, 10150, is a metalloporphyrin specifically designed to neutralize reactive oxygen and nitrogen species. The Company is developing 10150 as a medical countermeasure against the pulmonary effects of radiation exposure under a contract ("BARDA Contract") valued at up to \$118.4 million with the Biomedical Advanced Research and Development Authority ("BARDA"), a division of the Department of Health and Human Services ("HHS"). Additionally, Aeolus receives development support from the National Institutes of Health ("NIH") for development of the compound as a medical countermeasure against radiation and chemical exposure.

B. Summary of Significant Accounting Policies

Basis of Presentation

The consolidated financial statements include the accounts of Aeolus and its wholly owned subsidiary. All significant intercompany accounts and transactions have been eliminated. The Company uses the equity method to account for its 35% ownership interest in CPEC, which is further discussed in Note D.

Use of Estimates

The preparation of financial statements in conformity with accounting principles generally accepted in the United States of America requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosures of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenues and expenses during the reporting period. Such estimates include revenue recognition, allowance for doubtful accounts and stock-based compensation. Actual results could differ from those estimates.

Cash and Cash Equivalents

The Company invests available cash in short-term bank deposits. Cash and cash equivalents include investments with maturities of three months or less at the date of purchase. The carrying value of cash and cash equivalents approximate their fair market value at September 30, 2016 and 2015 due to their short-term nature. Also, the Company maintains cash balances with financial institutions in excess of federally insured limits. The Company does not anticipate any losses with such cash balances.

Significant customers and accounts receivable

For the years ended September 30, 2016 and 2015, the Company's only customer was BARDA. For the years ended September 30, 2016 and 2015, revenues from BARDA comprised 100% of total revenues. As of September 30, 2016 and 2015, the Company's receivable balances were comprised 100% from this customer. Unbilled accounts receivable, included in accounts receivable, totaling \$490,000 and \$589,000 as of September 30, 2016 and 2015, respectively, relate to work that has been performed, though invoicing has not yet occurred. All of the unbilled receivables are expected to be billed and collected within the next year. Accounts receivable are stated at invoice amounts and consists of amounts due from the BARDA Contract. If necessary, the Company records a provision for doubtful receivables to allow for any amounts which may be unrecoverable. This provision is based upon an analysis of the Company's prior collection experience, customer creditworthiness and current economic trends. As of September 30, 2016 and 2015, an allowance for doubtful accounts was not recorded as the collection history from the Company's customers indicated that collection was probable.

Concentrations of credit risk

Financial instruments that potentially subject the Company to concentrations of credit risk consist primarily of cash and cash equivalents and accounts receivable. The Company places its cash and cash equivalents with high quality financial institutions. Management believes that the financial risks associated with its cash and cash equivalents and investments are minimal. Because accounts receivable consist entirely of amounts due from the U.S. federal government agencies, management deems there to be minimal credit risk.

Revenue Recognition

Aeolus recognizes revenue in accordance with the authoritative guidance for revenue recognition. Revenue is recognized when all of the following criteria are met: (i) persuasive evidence of an arrangement exists, (ii) delivery (or passage of title) has occurred or services have been rendered, (iii) the seller's price to the buyer is fixed or determinable, and (iv) collectability is reasonably assured.

The BARDA Contract is classified as a "cost-plus-fixed-fee" contract. Aeolus recognizes government contract revenue in accordance with the authoritative guidance for revenue recognition including the authoritative guidance specific to federal government contracts. Reimbursable costs under the contract primarily include direct labor, subcontract costs, materials, equipment, travel, and indirect costs.

Aeolus estimates subcontractor costs, materials and related revenue for some vendors when invoices are not received timely. Management receives regular updates from our subcontractors regarding estimated completion of individual projects. Management evaluates the status of each project with respect to budgeted work completed, actual work completed, and cost of actual work completed. Management is required to provide BARDA with monthly reports in addition to our bi-weekly conference calls with BARDA regarding the progress of each project.

In addition, Aeolus receives a fixed fee on consultant labor and subcontract labor under the BARDA Contract, which is unconditionally earned as allowable costs are incurred and is not contingent on success factors. Reimbursable costs under this BARDA Contract, including the fixed fee, are generally recognized as revenue in the period the reimbursable costs are incurred and become billable.

Fair Value of Financial Instruments

The carrying amounts of financial instruments, which include cash and cash equivalents, accounts receivable, accounts payable, accrued liabilities, debt, and redemption liability, approximate their fair values due to their short maturities.

Fair Value Measurements

ASC 820 discusses valuation techniques, such as the market approach (comparable market prices), the income approach (present value of future income or cash flow) and the cost approach (cost to replace the service capacity of an asset or replacement cost). The Company utilizes the market approach. The statement utilizes a fair value hierarchy that prioritizes the inputs to valuation techniques used to measure fair value into three broad levels. The following is a brief description of those three levels:

- Level 1: Observable inputs such as quoted prices (unadjusted) in active markets for identical assets or liabilities.
- Level 2: Inputs other than quoted prices that are observable for the asset or liability, either directly or indirectly.
- These include quoted prices for similar assets or liabilities in active markets and quoted prices for identical or similar assets or liabilities in markets that are not active.
- Level 3: Unobservable inputs that reflect the reporting entity's own assumptions.

The redemption liability, discussed further at Note F. Debt, was measured at fair value on a recurring basis as of September 30, 2015 and is summarized below (in thousands):

Fair value at		
September 30,		
2015		
Level	Level	Level
1	2	3
\$—	\$275	\$ —

The Company had no assets or liabilities that were measured at fair value on a recurring basis as of September 30, 2016.

The following table summarizes, as of September 30, 2016, the redemption liability activity subject to Level 2 inputs which are measured on a recurring basis:

Fair value measurement of redemption liability	
Balance at September 30, 2015	\$275
Conversion of convertible promissory notes with redemption feature	(275)
Change in fair value of redemption liability	—
Balance at September 30, 2016	\$—

Research and Development

Research and development costs are expensed in the period incurred.

Leases

The Company leases office space and office equipment under month to month operating lease agreements. For the years ended September 30, 2016 and 2015, total rent expense was approximately \$42,000 and \$42,000, respectively.

Income Taxes

The Company recognizes liabilities or assets for the deferred tax consequences of temporary differences between the tax bases of assets or liabilities and their reported amounts in the financial statements. These temporary differences will result in taxable or deductible amounts in future years when the reported amounts of the assets or liabilities are recovered or settled. A valuation allowance is established when management determines that it is more likely than not that all or a portion of a deferred tax asset will not be realized. Management evaluates the Company's ability to realize its net deferred tax assets on a quarterly basis and valuation allowances are provided, as necessary. During this evaluation, management reviews its forecasts of income in conjunction with other positive and negative evidence surrounding the Company's ability to realize its deferred tax assets to determine if a valuation allowance is required. Adjustments to the valuation allowance will increase or decrease the Company's income tax provision or benefit. Management also applies the relevant guidance to determine the amount of income tax expense or benefit to be allocated among continuing operations, discontinued operations, and items charged or credited directly to stockholders' equity (deficit).

A tax position must meet a minimum probability threshold before a financial statement benefit is recognized. The minimum threshold is a tax position that is more likely than not to be sustained upon examination by the applicable taxing authority, including resolution of any related appeals or litigation process, based on the technical merits of the position. The Company recognizes interest and penalties related to uncertain tax positions in income tax expense.

Net Income (Loss) Per Common Share

The Company computes basic net income (loss) per weighted average share attributable to common stockholders using the weighted average number of shares of common stock outstanding during the period. The Company computes diluted net income (loss) per weighted average share attributable to common stockholders using the weighted average number of shares of common and dilutive potential common shares outstanding during the period. Potential common shares outstanding consist of stock options, warrants and convertible preferred stock using the treasury stock method and are excluded if their effect is anti-dilutive. Diluted weighted average common shares did not include any incremental shares for the fiscal year ended September 30, 2016 and 2015. Diluted weighted average common shares excluded incremental shares from common and preferred shares and warrants of approximately 85,606,000 and 29,536,000 for the fiscal year 2016 and 2015, respectively, due to their anti-dilutive effect.

	In thousands, except per share data	
	Fiscal Year Ended	
	September 30, 2016	2015
Numerator:		
Net loss	\$(3,558)	\$(2,628)
Less deemed dividend on Series C preferred stock	(2,486)	—
Net loss attributable to common stockholders – basic	\$(6,044)	\$(2,628)
Net loss attributable to common stockholders – diluted	\$(6,044)	\$(2,628)
Denominator:		
Weighted-average shares used in computing net loss per share attributable to common stockholders – basic	148,668	135,883
Effect of potentially dilutive securities:		
Common stock warrants	—	—
Convertible preferred warrants	—	—
Convertible preferred stock	—	—
Common stock options	—	—
Non-participating common stock warrants	—	—
Weighted-average shares used in computing net loss per share attributable to common stockholders – diluted	148,668	135,883
Basic net loss per common share	\$(0.04)	\$(0.02)
Diluted net loss per common share	\$(0.04)	\$(0.02)

Accounting for Stock-Based Compensation

The Company recognizes stock-based compensation expense in the statement of operations based upon the fair value of the equity award amortized over the vesting period.

Segment Reporting

The Company currently operates in one segment.

C. Liquidity

As of September 30, 2016, we had approximately \$3,155,000 of cash and cash equivalents, an increase of \$3,061,000 from September 30, 2015. The increase in cash was primarily due to our 2015 capital raise, which was offset by cash used in operations. In order to fund on-going operating cash requirements, or to accelerate or expand our oncology and other programs, we may need to raise significant additional funds.

We had a net loss of \$3,558,000 for the fiscal year ended September 30, 2016, compared to a net loss of \$2,628,000 for the fiscal year ended September 30, 2015. For the same periods, we had cash outflows from operations of approximately \$3,109,000 and cash outflows from operations of approximately \$2,443,000, respectively, with the outflows increasing in 2016 due to lower revenue from BARDA.

Our ongoing future cash requirements will depend on numerous factors, particularly the progress of our development programs, clinical trials and our ability to negotiate and complete collaborative agreements or out-licensing arrangements. In order to help fund our on-going operating cash requirements, we intend to seek new collaborations for our antioxidant research program that include initial cash payments and on-going research support. In addition, we might sell additional shares of our stock and/or convertible debentures and explore other strategic and financial alternatives, including a merger with another company, the sale of stock and/or debt, the establishment of new collaborations for current research programs that include initial cash payments and ongoing research support and the out-licensing of our compounds for development by a third party. We expect to incur additional losses and negative cash flow from operations for several more years.

Under the BARDA Contract, substantially all of the costs of the development of 10150 as a medical countermeasure for pulmonary injuries resulting from an acute exposure to radiation from a radiological/nuclear accident or attack, particularly injuries associated with ARS or DEARE could be paid for by the U.S. government through BARDA funding. We recognized approximately \$2,076,000 in revenue during the fiscal year ended September 30, 2016 related to the BARDA Contract.

D. Investments

Investment in CPEC LLC

The Company uses the equity method to account for its 35% ownership interest in CPEC. CPEC had \$91,000 of net assets at each of September 30, 2016 and 2015. Aeolus' 35% share of CPEC's net assets, which is approximately \$32,000, is included in other assets and CPEC has no operations or activities unrelated to the out licensing of bucindolol. The remaining membership interest in CPEC of 65% is owned by Endo Pharmaceuticals.

E. Commitments

The Company acquires assets still in development and enters into license and research and development arrangements with third parties that often require milestone and royalty payments to the third party contingent upon the occurrence of certain future events linked to the success of the asset in development. Milestone payments may be required, contingent upon the successful achievement of an important point in the development life-cycle of the pharmaceutical product (e.g., approval of the product for marketing by a regulatory agency). If required by the arrangement, the Company may also be required to make royalty payments based upon a percentage of the net sales of the pharmaceutical product in the event that regulatory approval for marketing is obtained. Because of the contingent nature of these payments, they are not included in the table of contractual obligations. No milestones have been met, nor have any payments been made, as of September 30, 2016.

We are also obligated to pay patent filing, prosecution, maintenance and defense costs, if any, for the intellectual property the Company has licensed from National Jewish Health ("NJH"), National Jewish Medical and Research Center (the "NJMRC") and Duke University.

These arrangements may be material individually, and in the unlikely event that milestones for multiple products covered by these arrangements were reached in the same period, the aggregate charge to expense could be material to the results of operations in any one period. In addition, these arrangements often give Aeolus the discretion to unilaterally terminate development of the product, which would allow Aeolus to avoid making the contingent payments; however, Aeolus is unlikely to cease development if the compound successfully achieves clinical testing objectives.

F. Debt

Convertible Promissory Notes

On September 29, 2015, the Company received funding in exchange for issuance of convertible promissory notes (the "Notes") from Biotechnology Value Fund, L.P. and other affiliates of BVF Partners, L.P.

The Notes had an aggregate principal balance of \$1,000,000, accrued interest at a rate of 6% per annum and had a scheduled maturity date of September 28, 2016 (the "Maturity Date"). The outstanding principal and accrued interest on the Notes shall automatically convert into Company equity securities issued in a Qualified Financing (as defined below) at a conversion rate carrying a 15% discount to the lowest price per share (or share equivalent) issued in a Qualified Financing (an "Automatic Conversion"). If, prior to the Maturity Date, the Company enters into an agreement pertaining to a Corporate Transaction (as defined below) and the Note has not been previously converted pursuant to an Automatic Conversion, then, the outstanding principal balance and unpaid accrued interest of the Note shall automatically convert in whole into the right of the holder to receive, in lieu of any other payment and in cancellation of the Note, an amount in cash upon closing of the Corporate Transaction equal to two times the outstanding principal amount of the Note.

For purposes of the foregoing: "Qualified Financing" means a bona fide new money equity securities financing on or before the Maturity Date with total proceeds to the Company of not less than four million dollars; and "Corporate Transaction" means a sale, lease or other disposition of all or substantially all of the Company's assets or a consolidation or merger of the Company with or into any other corporation or other entity or person, or any other corporate reorganization, in which the stockholders of the Company immediately prior to such consolidation, merger or reorganization own less than fifty percent (50%) of the voting power of the surviving entity immediately after such consolidation, merger or reorganization.

As of September 30, 2015, the \$1,000,000 principal balance of the Notes was recorded in the financial statements at face value and was net with a discount of \$273,000 as a result of separating the fair value of the Qualified Financing redemption discount ("Redemption Feature") of 15% on the price per share in the Notes. The Redemption Feature qualified as a derivative and was subject to fair value treatment. The Redemption Feature was being amortized over the expected life of the derivative, and the amortization expense is presented net with the interest expense from the Notes, yielding an effective interest rate of 40% that is different than the 6% stated in the Notes.

On December 11, 2015, following the completion of a Qualified Financing described above, the principal and accrued interest amounts under the Notes were converted into 5,414,402 shares of the Company's common stock and warrants to purchase an additional 5,414,402 shares of the Company's common stock at an exercise price per share of \$0.22 subject to adjustment (Note G). As a result, the Notes were no longer outstanding as of that date.

G. Stockholders' Equity (Deficit)

Basis of Presentation

Preferred Stock

The Certificate of Incorporation of the Company authorizes the issuance of up to 10,000,000 shares of Preferred Stock, at a par value of \$0.01 per share, of which 1,250,000 shares are designated Series A Convertible Preferred Stock and 1,600,000 shares are designated Series B Convertible Preferred Stock and 5,000 shares are designated Series C Convertible Preferred Stock (the "Series C Stock"). The Board of Directors has the authority to issue Preferred Stock in one or more series, to fix the designation and number of shares of each such series, and to determine or change the designation, relative rights, preferences, and limitations of any series of Preferred Stock, without any further vote or action by the stockholders of the Company.

There are no shares of Series A Convertible Preferred Stock issued or outstanding. As of September 30, 2016 and 2015, zero and 526,080 shares of Series B Stock were outstanding, respectively. The decrease in Series B shares outstanding is due to the conversion of the remaining shares into common shares in June 2016. As of September 30, 2016 and 2015, 4,500 and zero shares of Series C Stock were outstanding, respectively.

With respect to dividend rights and rights upon liquidation, winding up and dissolution, the Series B Stock ranks *pari passu* with the common stock. Subject to any rights of senior stock, holders of Series B Stock are entitled to receive dividends or distributions as, when and if declared by the Board of Directors. In the event the Board of Directors declares a dividend or distribution with respect to the outstanding common stock, the holders of Series B Stock are entitled to receive the amount of dividends per share in the same form payable on the common stock based on the largest number of shares of common stock issuable upon conversion of the outstanding Series B Stock. In the event of a liquidation, winding up or dissolution of the Company, subject to any rights of senior stock, the holders of Series B Stock are entitled to receive, *pari passu* with the holders of the common stock, the assets of the Company based on the largest number of shares of common stock issuable upon conversion of the outstanding Series B Stock.

Each share of Series B Stock is convertible into one share of common stock. The Series B Stock can be converted into common stock at any time upon the election of the holders of the Series B Stock except to the extent such conversion would result in the holders of Series B Stock owning in the aggregate more than 9.99% of the outstanding common stock.

The Series B Stock is not entitled to vote on any matter submitted to the vote of holders of the common stock except that the Company must obtain the approval of a majority of the outstanding shares of Series B Stock to either amend the Company's Certificate of Incorporation in a manner that would adversely affect the Series B Stock (including by creating an additional class or series of stock with rights that are senior or *pari passu* to the Series B Stock) or change the rights of the holders of the Series B Stock in any other respect.

The Series C Stock is non-voting stock. Each share of Series C Stock is convertible into 4,545 shares of our common stock except to the extent such conversion would result in such holder of Series C Stock, and its affiliates, owning in the aggregate more than 9.99% of the outstanding common stock. Dividends on the Series C Stock are due whenever dividends are due on the Company's common stock on an as-if-converted basis, but shall be subordinate to any dividends due to holders of the Company's Series B Stock as a result of such common stock dividends. The Series C Stock shall also be junior to the Series B Stock in the event of liquidation of the Company.

On December 10, 2015, the Company entered into securities purchase agreements with certain accredited investors to sell and issue (i) 4,500 preferred stock units issued to existing investors, Biotechnology Value Fund, L.P. and other affiliates of BVF Partners, L.P., for an aggregate purchase price of \$4.5 million, and (ii) an aggregate of 10,215,275 common units issued at purchase price of \$0.22 per unit, resulting in aggregate gross proceeds to the Company of approximately \$6.75 million. The preferred units collectively consist of (i) 4,500 shares of Series C Convertible Preferred Stock of the Company that are collectively convertible into an aggregate of 20,454,546 common shares and (ii) warrants to purchase an aggregate of 20,454,546 Common Shares, in each case subject to adjustment. The warrants have an initial exercise price of \$0.22 per share. The warrants may not be exercised until after 90 days following the date of issuance. The Series C Stock and warrants contain provisions restricting the conversion or exercise of such securities in circumstances where such event would result in the holder and its affiliates to beneficially own in excess of 9.99% of the Company's outstanding common stock.

The fair value of the December 10, 2015 financing warrants issued was estimated to be \$4,476,000 using the Black-Scholes option pricing model with the following assumptions: dividend yield of 0%, expected volatility of 109.74%, risk free interest rate of 1.67%, and an expected life equal to the five year contractual term. The proceeds from the December 10, 2015 financing were allocated based upon the relative fair values of the warrants and preferred shares issued in the transaction.

The allocation of the proceeds based on relative fair values of the instruments resulted in recognition of a discount on the Series C Preferred Stock of \$2,486,000 from a beneficial conversion feature, which is being amortized from the date of issuance to the earliest redemption date of 90 days post issuance. For the nine months ended June 30, 2016 the Company recognized \$2,486,000 of amortization of the discount on Series C Preferred Stock as deemed dividends charged to additional paid in capital. The value of the beneficial conversion feature is calculated as the difference between the effective conversion price of the Series C Preferred Stock and the fair market value of the common stock into which the Series C Preferred Stock are convertible at the commitment date.

Common Stock

On December 10, 2015, the Company entered into securities purchase agreements with certain accredited investors to sell and issue (i) an aggregate of 10,215,275 common units issued at a purchase price of \$0.22 per unit. Each common unit consists of one share of the Company's common stock and a five year warrant to purchase one share of the Company's common stock, subject to adjustment. The warrants may not be exercised until after 90 days following the date of issuance. The warrants contain provisions restricting the conversion or exercise of such securities in circumstances where such event would result in the holder and its affiliates to beneficially own in excess of 9.99% of the Company's outstanding common stock.

On September 29, 2015, the Company received funding in the form of convertible promissory notes (the "BVF Notes") from Biotechnology Value Fund, L.P. and certain other affiliates of BVF Partners, L.P. The BVF Notes have an aggregate principal balance of \$1,000,000, accrue interest at a rate of 6% per annum and have a scheduled maturity date of September 28, 2016. The outstanding principal and accrued interest on the BVF Notes will automatically convert into Company equity securities, provided a Qualified Financing of not less than \$4,000,000 occurs.

On December 11, 2015, following the completion of a Qualified Financing described above, the principal and accrued interest amounts under the BVF Notes were converted into 5,414,402 shares of the Company's common stock and warrants to purchase an additional 5,414,402 shares of the Company's common stock at an exercise price per share of \$0.22 subject to adjustment. As a result, the BVF Notes were no longer outstanding as of that date.

Warrants

As of September 30, 2016, warrants to purchase an aggregate of 52,947,877 shares of common stock were outstanding. Details of the warrants for common stock outstanding at September 30, 2016 were as follows:

Number of Shares	Exercise Price	Expiration Date
1,337,627	\$ 0.40	March 2017
325,000	\$ 0.40	April 2017
300,000	\$ 0.258	June 2017
50,000	\$ 0.26	June 2017
140,000	\$ 0.35	October 2017
12,205,000	\$ 0.25	February 2018
1,242,000	\$ 0.25	March 2018
50,000	\$ 0.49	January 2020
37,298,250	\$ 0.22	December 2020
52,947,877		

As of September 30, 2015, warrants to purchase an aggregate of 15,949,627 shares of common stock were outstanding. Details of the warrants for common stock outstanding at September 30, 2015 were as follows:

Number of Shares	Exercise Price	Expiration Date
50,000	\$ 0.50	May 2016
50,000	\$ 0.50	July 2016
50,000	\$ 1.00	July 2016
50,000	\$ 1.50	July 2016
50,000	\$ 2.00	July 2016
50,000	\$ 2.50	July 2016
1,337,627	\$ 0.40	March 2017
325,000	\$ 0.40	April 2017
300,000	\$ 0.258	June 2017
50,000	\$ 0.26	June 2017

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140,000	\$ 0.35	October 2017
12,205,000	\$ 0.25	February 2018
1,242,000	\$ 0.25	March 2018
50,000	\$ 0.49	January 2020
15,949,627		

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As of September 30, 2015, one warrant with an exercise price of \$0.01 to purchase an aggregate of 896,037 shares of Series B preferred stock was outstanding.

Below is a summary of warrant activity for the last two fiscal years ended September 30:

	Number of Shares	Exercise Price	Weighted Average		Aggregate Intrinsic Value
			Contractual Term (in years)	Remaining	
Outstanding at 9/30/2014	16,925,664	\$0.27	3.1 years		\$215,048
Granted	50,000	\$0.49			\$-
Exercised	(80,000)	\$0.25			\$10,975
Cancelled	(50,000)	\$0.38			\$-
Forfeited	-	\$-			\$—
Outstanding at 9/30/2015	16,845,664	\$0.27	2.2 years		\$206,626
Granted	37,298,250	\$0.22			\$1,864,913
Exercised	—	\$—			\$—
Cancelled	(1,196,037)	\$0.34			\$156,000
Forfeited	—	\$—			\$—
Outstanding at 9/30/2016	52,947,877	\$0.23	3.3 years		\$—
Exercisable at 9/30/2016	52,947,877	\$0.23	3.3 years		\$—

H. Stock-Based Compensation

As an integral component of a management and employee retention program designed to motivate, retain and provide incentive to the Company's management, employees and key consultants, the Board of Directors approved the 2016 Stock Incentive Plan (the "2016 Plan") and reserved 25,000,000 shares of common stock for issuance under the 2016 Plan. As of September 30, 2016, 25,000,000 shares were available to be granted under the 2016 Plan. The exercise price of the incentive stock options ("ISOs") granted under the 2016 Plan must not be less than the fair market value of the common stock as determined on the date of the grant. The options may have a term up to 10 years. Options typically vest immediately or up to one year following the date of the grant.

Below is a summary of stock option activity for the last two fiscal years ended September 30:

	Number of Shares	Exercise Price	Weighted Average		Aggregate Intrinsic Value
			Contractual Term (in years)	Remaining	
Outstanding at 9/30/2014	11,596,591	\$0.43	6.1 years		\$ 1
Granted	775,000	\$0.27			\$ -
Exercised	-	\$-			\$ -
Cancelled	(207,000)	\$0.90			\$ -
Forfeited	-	\$-			\$ -
Outstanding at 9/30/2015	12,164,591	\$0.42	5.4 years		\$ 1
Granted	700,000	\$0.23			\$ -
Exercised	-	\$-			\$ -
Cancelled	(660,591)	\$0.82			\$ -
Forfeited	-	\$-			\$ -
Outstanding at 9/30/2016	12,204,000	\$0.38	5.0 years		\$ 1
Exercisable at 9/30/2016	12,041,498	\$0.39	4.9 years		\$ 1

Stock options granted to consultants during fiscal year 2016 and 2015 were fully vested when issued or vested over a twelve month period. Stock option expense for stock options granted to consultants was zero for fiscal year 2016 and 2015, respectively. For the fiscal years 2016 and 2015, all stock options were issued at or above fair market value of a share of common stock. The weighted-average grant-date fair value of options granted during fiscal years 2016 and 2015 was \$0.19 and \$0.24, respectively.

A summary of the status of non-vested shares for the fiscal years ended September 30 was:

	Number of Shares	Weighted Average Grant-Date Fair Value
Unvested at September 30, 2014	168,752	\$ 0.22
Granted	775,000	\$ 0.24
Vested	(781,250)	\$ 0.24
Forfeited	-	\$ -
Unvested at September 30, 2015	162,502	\$ 0.23
Granted	700,000	\$ 0.19
Vested	(700,000)	\$ 0.24
Forfeited	-	\$ -
Unvested at September 30, 2016	162,502	\$ 0.16

The total unrecognized compensation expense for outstanding stock options was \$15,000 as of September 30, 2016, which will be recognized over a weighted average period of five months. The total fair value of shares vested during fiscal years 2016 and 2015 was \$142,000 and \$156,000, respectively.

The details of stock options for the fiscal year ended September 30, 2016 are as follows:

Range of Exercise Prices	Options Outstanding			Options Exercisable		
	Number Outstanding at September 30, 2016	Weighted Average Exercise Price	Weighted Average Remaining Contractual Life (in years)	Number Exercisable at September 30, 2016	Weighted Average Exercise Price	Weighted Average Remaining Contractual Life (in years)
\$0.19-\$0.20	250,000	\$ 0.19	9.42	125,000	\$ 0.19	9.42
\$0.21-\$0.30	3,537,500	\$ 0.27	5.83	3,499,998	\$ 0.27	5.80
\$0.31-\$0.40	6,551,500	\$ 0.39	4.83	6,551,500	\$ 0.39	4.83
\$0.41-\$0.50	502,000	\$ 0.45	5.24	502,000	\$ 0.45	5.24
\$0.51-\$0.60	968,250	\$ 0.59	2.75	968,250	\$ 0.59	2.75
\$0.61-\$0.70	56,000	\$ 0.68	0.04	56,000	\$ 0.68	0.04
\$0.71-\$0.80	80,750	\$ 0.76	4.23	80,750	\$ 0.76	4.23
\$0.81-\$0.90	256,000	\$ 0.90	0.78	256,000	\$ 0.90	0.78
\$0.91-\$1.19	2,000	\$ 1.19	0.58	2,000	\$ 1.19	0.58
	12,204,000	\$ 0.38	4.95	12,041,498	\$ 0.39	4.89

Stock-based compensation expense recognized in the statement of operations is as follows (in thousands):

	For the fiscal year ended September 30,	
	2016	2015
Research and Development Expenses	\$—	\$—
General and Administrative Expenses	141	178
Total Stock-based Compensation Expense	\$141	\$178

The fair value of the options associated with the above compensation expense was determined at the date of the grant using the Black-Scholes option pricing model with the following weighted average assumptions:

	For the fiscal year ended September 30,			
	2016		2015	
Dividend yield	0	%	0	%
Unvested forfeiture rate	7.87	%	5.35	%
Expected volatility	117	%	137	%
Risk-free interest rate	1.42	%	1.70	%
Expected option life after shares are vested	5.27 years		5.27 years	

I. Income Taxes

As of September 30, 2016 and 2015, the Company had federal net operating loss ("NOL") carry-forwards of \$119,215,000 and \$115,515,000, respectively and state operating loss carry-forwards of \$42,615,000 and \$38,916,000, respectively. The use of these federal and state NOL carry-forwards might be subject to limitation under the rules regarding a change in stock ownership as determined by the Internal Revenue Code (the "Code"). The Company may have had a change of control under Section 382 of the Code during fiscal 2004 and 2006; however, a complete analysis of the limitation of the NOL carry-forwards will not be completed until the time the Company projects it will be able to utilize such NOLs. The federal net operating and the state net operating losses began to expire in 2010. Additionally, the Company had federal research and development carry-forwards as of September 30, 2016 and 2015 of \$4,198,000 and \$4,127,000, respectively. The Company had state research and development carry-forwards as of September 30, 2016 and 2015 of \$1,530,000 and \$1,452,000, respectively. Significant components of the Company's deferred tax assets at September 30, 2016 and 2015 consisted of the following (in thousands):

	2016	2015
Accrued payroll related liabilities	\$885	\$1,036
Depreciation and amortization	469	586
Total deferred tax assets	1,354	1,622
State taxes	(94)	(113)
Total deferred tax liabilities	(94)	(113)
Net deferred tax assets	1,260	1,509
Valuation allowance for deferred assets	(1,260)	(1,509)
Net deferred tax asset	\$—	\$—

Due to the uncertainty surrounding the realization of the favorable tax attributes in future tax returns, all of the deferred tax assets have been fully offset by a valuation allowance. The change in the valuation allowance is primarily a result of the net operating loss carry-forwards.

Taxes computed at the statutory federal income tax rate of 34% are reconciled to the provision for income taxes as follows (dollars in thousands):

	2016		2015	
	0	%	0	%
Effective income tax rate				
United States Federal income tax at statutory rate	\$(1,209)		\$(891)	
State income taxes (net of federal benefit)	1		1	
NQSO forfeiture	151		234	
Prior year deferred true up	—		39	
Change in valuation reserves	(212)		(153)	
FIN 48	1,329		883	
Other	(57)		(111)	
Provision for income taxes	\$2		\$2	

At September 30, 2016, we had unrecognized tax benefits of \$5,208,000, all of which would impact the effective tax rate if recognized, however a valuation allowance would be recorded against this amount. During fiscal 2016, unrecognized tax benefits increased to \$5,208,000 as a result of the tax positions taken in prior years and current year. No interest and penalties have been provided for with respect to the unrecognized tax benefits.

A reconciliation of the beginning and ending amount of unrecognized tax benefits is as follows:

	Amount (in thousands)
Unrecognized tax benefits at October 1, 2015	\$ 5,085
Additions for tax positions related to current year	117
Additions/reductions for tax positions taken in prior years	6
Unrecognized tax benefits at September 30, 2016	\$ 5,208

The Company's federal income tax returns for the tax years 2012 to 2014 remain open to examination. The Company's California income tax returns for the tax years 2011 to 2014 remain open to examination.

A reconciliation of the current year tax provision is as follows:

Current – Federal	\$—
Current – State	2
	2
Deferred - Federal	194
Deferred – State	55
	249
Less: Valuation Allowance	(249)
	—
Total current and deferred	\$2

J. Contingencies

Duke Licenses

The Company has obtained exclusive worldwide licenses (the "Duke Licenses") from Duke University ("Duke") to develop, make, have made, use and sell products using certain technology in the field of free radical and antioxidant research, developed by certain scientists at Duke. Future discoveries in the field of antioxidant research from these scientists' laboratories at Duke are also covered by the Duke Licenses. The Duke Licenses require the Company to use its best efforts to pursue development of products using the licensed technology and compounds. These efforts are to include the manufacture or production of products for testing, development and sale. Aeolus is also obligated to use its best efforts to have the licensed technology cleared for marketing in the United States by the U.S. Food and Drug Administration and in other countries in which Aeolus intends to sell products using the licensed technology. Aeolus will pay royalties to Duke on net product sales during the terms of the Duke Licenses, and milestone payments upon certain regulatory approvals and annual sales levels. In addition, Aeolus is obligated under the Duke Licenses to pay all or a portion of patent prosecution, maintenance and defense costs. Unless earlier terminated, the Duke Licenses continue until the expiration of the last to expire issued patent on the licensed technology.

National Jewish Medical and Research Center Agreements

Aeolus has an exclusive worldwide license ("NJH License") from National Jewish Health to develop, make, have made, use and sell products using certain technology developed by certain scientists at NJH. The NJH License requires Aeolus to use commercially reasonable efforts to diligently pursue the development and government approval of products using the licensed technology. Aeolus will be obligated to pay royalties to NJH on net product sales during the term of the NJH License and a milestone payment upon regulatory approval, if obtained. In addition, Aeolus is obligated under the NJH License to pay all or a portion of patent prosecution, maintenance and defense costs. Unless earlier terminated, the NJH License continues until the expiration of the last to expire issued patent on the licensed technology.

K. Agreements

Elan Corporation, plc

In May 2002, the Company entered into a collaboration transaction with affiliates of Elan Corporation, plc for the development of the Company's catalytic antioxidant compounds as a treatment for tissue damage from cancer radiation and chemotherapy. Although Elan and the Company terminated this collaboration in January 2003, the Company will pay Elan a royalty on net sales of its catalytic antioxidant products sold, if any, for the prevention and treatment of radiation-induced and chemotherapy-induced tissue damage.

L. Litigation

From time to time the Company may be subject to various lawsuits and claims with respect to matters arising out of the normal course of our business. Management does not believe these cases will have a material effect on our financial statements.

M. Recently Issued Accounting Pronouncements

From time to time, new accounting pronouncements are issued by the Financial Accounting Standards Board that are adopted by us as of the specified effective date. Unless otherwise discussed, we believe that the impact of recently issued standards that are not yet effective will not have a material impact on our financial position or results of operations upon adoption.

In May 2014, the Financial Accounting Standards Board ("FASB") issued Accounting Standards Update ("ASU") 2015-14, which deferred the effective date of ASU 2014-09, Revenue from Contracts with Customers (ASC 606), which updates the principles for recognizing revenue. ASU 2014-09 also amends the required disclosures of the nature, amount, timing and uncertainty of revenue and cash flows arising from contracts with customers. ASU 2014-09 is now effective for annual reporting periods beginning after December 15, 2017, including interim periods within that reporting period. The Company is evaluating the potential impacts of the new standard on its existing revenue recognition policies and procedures.

In August 2014, the FASB issued ASU 2014-15, Disclosure of Uncertainties about an Entity's Ability to Continue as a Going Concern. ASU 2014-15 requires that an entity's management evaluate whether there are conditions or events that raise substantial doubt about the entity's ability to continue as a going concern within one year after the date that the financial statements are issued. ASU 2014-15 is effective for annual periods ending after December 15, 2016 and for interim periods thereafter. The Company is evaluating the potential impacts of this new standard on its reporting process.

Item 9. Changes in and Disagreements with Accountants on Accounting and Financial Disclosure.

None.

Item 9A. Controls and Procedures.

As of the end of the period covered by this Annual Report on Form 10-K, we carried out an evaluation, under the supervision and with the participation of our management, including our President and Chief Executive Officer (our Principal Executive Officer) and Chief Financial Officer (our Principal Financial and Accounting Officer), of the effectiveness of our disclosure controls and procedures required by Rule 13a-15 under the Securities Exchange Act of 1934, as amended (the "Exchange Act"). Based upon that evaluation, our Principal Executive Officer and Principal Financial and Accounting Officer have concluded that our disclosure controls and procedures were effective as of September 30, 2016 to provide reasonable assurance that information required to be disclosed by us in reports that we file or submit under the Exchange Act is recorded, processed, summarized and reported within the time periods specified in SEC rules and forms.

Management's Report on Internal Control over Financial Reporting

Our management is responsible for establishing and maintaining adequate internal control over financial reporting as defined in Rule 13a-15 (f) and 15d-15 (f) of the Exchange Act. Our internal control over financial reporting is a process designed to provide reasonable assurance to our management and Board of Directors regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles and includes policies and procedures that:

- pertain to the maintenance of records that, in reasonable detail, accurately and fairly reflect the transactions and dispositions of our assets;
- provide reasonable assurance that transactions are recorded as necessary to permit preparation of financial statements in accordance with generally accepted accounting principles, and that our receipts and expenditures are being made only in accordance with authorizations of our management and directors; and
- provide reasonable assurance regarding prevention or timely detection of unauthorized acquisition, use, or disposition of our assets that could have a material effect on our financial statements.

Because of its inherent limitations, internal control over financial reporting may not prevent or detect misstatements, errors or fraud. Also, projections of any evaluations of effectiveness to future periods are subject to the risk that controls may become inadequate because of changes in conditions, or that the degree of compliance with the policies or procedures may deteriorate.

Our management assessed the effectiveness of our internal control over financial reporting as of September 30, 2016. In making this assessment, our management used criteria set forth by the Committee of Sponsoring Organizations of the Treadway Commission ("COSO") in an Internal Control Integrated Framework (2013). Based on the criteria set forth by COSO, management concluded that our internal control over financial reporting was effective as of September 30, 2016.

This Annual Report on Form 10-K does not include an attestation report of our registered public accounting firm regarding internal control over financial reporting. Management's report was not subject to attestation by our registered public accounting firm pursuant to rules of the SEC that permit the Company to provide only management's report in this Annual Report on Form 10-K.

We have instituted new procedures for internal control over financial reporting (as defined in Rules 13a-15(f) and 15d-15(f) under the Securities Exchange Act of 1934, as amended) during the quarter ended September 30, 2016. These procedures consist of the use of external specialists for complex accounting issues. We will continue to monitor the effectiveness of these new control procedures.

Item 9B. Other Information.

None.

PART III

Certain information required by Part III is omitted from this Annual Report on Form 10-K because we expect to file a definitive information statement in connection with our Written Consent in Lieu of 2016 Annual Meeting of Stockholders (the "Information Statement") within 120 days after the end of our fiscal year pursuant to Regulation 14C promulgated under the Exchange Act, and the information included therein is incorporated herein by reference to the extent provided below.

Item 10. Directors, Executive Officers and Corporate Governance.

The information set forth under the headings "Matter No. 1: Election of Directors," "Information Concerning the Board of Directors and its Committees" and "Section 16(a) Beneficial Ownership Reporting Compliance" in our Information Statement is incorporated herein by reference. The information required by this Item 10 concerning our executive officers is set forth under the heading "Executive Officers" located at the end of Part I Item 1 of this Annual Report on Form 10-K.

Code of Ethics

We have adopted a code of ethics that applies to our principal executive officer, principal financial officer, principal accounting officer and persons performing similar functions. We have posted the text of Code of Ethics on our Internet website at www.aolsrx.com. A copy of the Code of Ethics can also be obtained free of charge by writing to David Cavalier, Aeolus Pharmaceuticals, Inc., 26361 Crown Valley Parkway, Suite 150 Mission Viejo, CA 92691.

Item 11. Executive Compensation.

The information set forth under the headings "Director Compensation" and "Executive Compensation" in our Information Statement is incorporated herein by reference.

Item 12. Security Ownership of Certain Beneficial Owners and Management and Related Stockholder Matters.

Equity Compensation Plan and Additional Equity Information as of September 30, 2016

Plan category	(a)Number of securities to be issued upon exercise of outstanding options, warrants and rights	(b)Weighted-average exercise price of outstanding options, warrants and rights	(c)Number of securities remaining available for future issuance under equity compensation plans (excluding securities reflected in column (a))
Equity compensation plans approved by our stockholders:			
2004 Stock Option Plan	12,204,000	\$ 0.38	—
2016 Stock Option Plan	—		25,000,000
Equity compensation plans and securities not approved by our stockholders:			
Warrants to Purchase Common Stock Issued to Dan Delmonico	50,000	\$ 0.49	Not applicable
Warrants to Purchase Common Stock Issued to Roberts Mitani, LLC	300,000	\$ 0.258	Not applicable
Total – Common Stock	12,554,000		25,000,000

Since the foregoing presentation is presented as of September 30, 2015, consistent with SEC rules, it does not reflect the issuance of warrants in connection with the 2015 Securities Placement and the conversion of the BVF Notes.

Description of Equity Compensation Plans and Equity Securities Not Approved by Our Stockholders

The warrant to purchase shares of our Common Stock issued to Roberts Mitani, LLC have not been approved by our stockholders. In June 2012, we entered into an advisory agreement with Roberts Mitani, LLC whereby we engaged Roberts Mitani, LLC to serve as an advisor to provide strategic advisory services to us on a non-exclusive basis. For these services, on June 26, 2012, we issued a warrant to purchase up to 300,000 shares of our Common Stock with a per share exercise price of \$0.258. The warrant is exercisable for seven years from the date of grant and contains standard adjustment provisions in the event we declare a stock dividend or engage in a recapitalization, reclassification or reorganization of our capital stock.

The warrants to purchase shares of our Common Stock issued to Columbia Capital Securities, Inc. and Monarch Bay Associates, LLC have not been approved by our stockholders. In August 2012, we entered into an advisory agreement with Columbia Capital Securities, Inc. and Monarch Bay Associates, LLC whereby we engaged them to serve as an advisor to provide strategic advisory services to us on a non-exclusive basis. For these services, we have agreed to pay each of Columbia Capital Securities, Inc. and Monarch Bay Associates, LLC a monthly retainer in the form of a warrant to purchase up to an aggregate of 17,500 shares of Common Stock, commencing on August 17, 2012 and continuing monthly thereafter during the term of their engagement under the advisory agreement. Each of these warrants has an exercise price equal to the closing price of the Common Stock on the date of issuance, is deemed fully vested upon issuance, is exercisable at any time on or before the five year anniversary of the date of issuance and contains standard adjustment provisions in the event we declare a stock dividend or engage in a recapitalization, reclassification or reorganization of our capital stock. On August 17, 2012, we issued a warrant to purchase up to an

aggregate of 17,500 shares of Common Stock with a per share exercise price of \$0.30 to each of Columbia Capital Securities, Inc. and Monarch Bay Associates, LLC. On September 17, 2012, we issued a warrant to purchase up to an aggregate of 17,500 shares of Common Stock with a per share exercise price of \$0.44 to each of Columbia Capital Securities, Inc. and Monarch Bay Associates, LLC.

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For a description of compensatory warrants issued before June of 2012 and included in the numbers in the table above, please refer to Item 12 in our Annual Report on Form 10-K for the period ended September 30, 2012.

The information set forth under the headings "Security Ownership of Certain Beneficial Owners and Management" in our Information Statement is incorporated herein by reference.

Item 13. Certain Relationships and Related Transactions, and Director Independence.

The information set forth under the headings "Information Concerning the Board of Directors and its Committees" and "Certain Related Transactions" in our Information Statement is incorporated herein by reference.

Item 14. Principal Accounting Fees and Services.

The information set forth under the heading "Independent Registered Public Accounting Firm — Fees" in our Information Statement is incorporated herein by reference.

PART IV

Item 15. Exhibits and Financial Statement Schedules.

(a) The following financial statement schedules and exhibits are filed as part of this report or incorporated herein by reference:

(1) Financial Statement Schedules.

All financial statement schedules for which provision is made in Regulation S-X are omitted because they are not required under the related instructions, are inapplicable, or the required information is given in the financial statements, including the notes thereto.

(2) Exhibits.

Exhibit Number	Description of Document	Incorporated by Reference To		
		Registrant's Form	Date Filed with the SEC	Exhibit Number Filed Herewith
2.1	Agreement and Plan of Merger and Reorganization dated September 16, 2003 between Incara, Inc. and Incara Pharmaceuticals Corporation	S-4	09/19/03	2.1
3.1	Amended and Restated Certificate of Incorporation	10-K	12/31/12	3.1
3.2	Certificate of Designation for Series C Preferred Stock	8-K	12/15/15	10.1
3.3	Bylaws	8-K	10/27/15	3.1
4.1	Form of Common Stock Certificate	10-Q	08/11/04	4.1
4.2	Form of Series B Preferred Stock Certificate	S-4	09/19/03	4.8
4.3	Form of Warrant to Purchase Common Stock dated June 5, 2006.	8-K	06/06/06	10.3
4.4	Registration Rights Agreement dated May 22, 2007 by and among the Company and each of the Purchasers whose names appear on the Schedule attached thereto.	8-K	5/23/07	4.1

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Exhibit Number	Description of Document	Incorporated by Reference To		Exhibit Number	Filed Herewith
		Registrant's Form	Date Filed with the SEC		
4.5	Registration Rights Agreement dated October 6, 2009 by and among the Company and the investors whose names appear on the signature pages thereof.			8-K	10/06/09 4.1
4.6	Form of Warrant to Purchase Common Stock dated May 22, 2007.			8-K	5/23/07 10.2
4.7	Form of Warrant to Purchase Common Stock			8-K	10/06/09 10.2
4.8	Registration Rights Agreement dated September 16, 2003 among Incara Pharmaceuticals Corporation, Incara, Inc. and Goodnow Capital, L.L.C.			S-4	09/19/03 10.101
4.9	Registration Rights Agreement dated August 11, 2010 by and among Aeolus Pharmaceuticals, Inc. and the investors listed therein			8-K	8/12/10 4.1
4.10	Registration Rights Agreement dated March 4, 2013 by and among Aeolus Pharmaceuticals, Inc. and the investors listed therein			8-K	03/06/13 10.2
4.11	Form of Warrant to Purchase Common Stock dated March 4, 2013.			8-K	03/06/13 10.3
4.12	Warrant Repricing, Exercise and Lockup Agreement dated February 19, 2013 by and among the Company, Xmark JV Investment Partners, LLC and affiliates			8-K	02/19/13 10.4
4.13	Form of Series C Preferred Stock Certificate				X
4.14	Form of Warrant			8-K	12/15/15 10.2
10.1*	License Agreement between Duke University and Aeolus Pharmaceuticals, Inc., dated July 21, 1995			S-1	12/08/95 10.4
10.2	Amended and Restated Limited Liability Company Agreement of CPEC LLC dated July 15, 1999, among CPEC LLC, Intercardia, Inc. and Interneuron Pharmaceuticals, Inc.			8-K	07/23/99 10.42
10.3	Assignment, Assumption and License Agreement dated July 15, 1999, between CPEC LLC and Intercardia, Inc.			8-K	07/23/99 10.43
10.4*	License Agreement dated January 19, 2001 between Incara Pharmaceuticals Corporation and Incara Development, Ltd.			10-Q	02/13/01 10.59
10.5*	License Agreement dated January 19, 2001 between Elan Corporation, plc, Elan Pharma International Ltd. and Incara Development, Ltd.			10-Q	02/13/01 10.60
10.6	Intentionally left blank				
10.7	Agreement and Amendment, effective as of January 22, 2001, by and among Incara Pharmaceuticals Corporation, Elan International Services, Ltd. and Elan Pharma International Limited			10-Q	05/14/01 10.64
10.8	Second Agreement and Amendment, effective as of January 22, 2001, by and among Incara Pharmaceuticals Corporation, Elan International Services, Ltd. and Elan Pharma International Limited			10-Q	05/14/01 10.65
10.9	Third Agreement and Amendment, effective as of January 22, 2001, by and among Incara Pharmaceuticals Corporation, Elan International Services, Ltd. and Elan Pharma International Limited			8-K	06/01/01 10.66

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Exhibit Number	Description of Document	Incorporated by Reference To		Exhibit Number	Filed Herewith
		Registrant's Form	Date Filed with the SEC		
10.10	Agreement and Fourth Amendment, effective February 13, 2002, by and among Incara Pharmaceuticals Corporation, Elan International Services, Ltd., Elan Pharma International Limited and Elan Pharmaceutical Investments III, Ltd.		02/14/02	10-Q	10.75
10.11*	License Agreement dated June 25, 1998 between Duke University and Aeolus Pharmaceuticals, Inc.		05/15/02	10-Q	10.82
10.12*	License Agreement dated May 7, 2002 between Duke University and Aeolus Pharmaceuticals, Inc.		05/15/02	10-Q	10.83
10.13*	License Agreement dated November 17, 2000 between National Jewish Medical and Research Center and Aeolus Pharmaceuticals, Inc.		02/13/01	10-Q	10.56
10.14	Exclusive License Agreement, dated January 15, 2009, by and between the Company and National Jewish Health		05/16/11	10-Q	10.7
10.15*	Securities Purchase Agreement dated as of May 15, 2002, among Incara Pharmaceuticals Corporation, Aeolus Pharmaceuticals, Inc., Elan Pharma International Limited and Elan International Services, Ltd.		07/03/02	8-K/A	10.84
10.16*	Development and Option Agreement dated May 15, 2002, among Elan Pharma International Limited, Incara Pharmaceuticals Corporation and Aeolus Pharmaceuticals, Inc.		07/03/02	8-K/A	10.85
10.17	Intentionally left blank				
10.18	Amendment No. 1 to License Agreement dated May 14, 2002, between Aeolus Pharmaceuticals, Inc. and Duke University (amending License Agreement dated July 21, 1995)		07/03/02	8-K/A	10.87
10.19	Amendment No. 1 to License Agreement dated May 14, 2002, between Aeolus Pharmaceuticals, Inc. and Duke University (amending License Agreement dated June 25, 1998)		07/03/02	8-K/A	10.88
10.20	Amendment No. 1 to License Agreement dated May 14, 2002, between Aeolus Pharmaceuticals, Inc. and National Jewish Medical and Research Center (amending License Agreement dated November 17, 2000)		07/03/02	8-K/A	10.89
10.21*	Subaward Agreement, dated March 16, 2011, by and between the Company and the Office of Research and Development of the University of Maryland, Baltimore		05/16/11	10-Q	10.4
10.22	Letter dated May 17, 2004 from Elan International Services, Limited and Elan Pharma International Limited to Incara Pharmaceuticals Corporation		08/11/04	10-Q	10.106
10.23+	Aeolus Pharmaceuticals, Inc. 1994 Stock Option Plan, as amended		08/11/04	10-Q	10.109
10.24+	Aeolus Pharmaceuticals, Inc. Amended and Restated 2004 Stock Incentive Plan		11/16/12	14-C	D
10.25+	Amended and Restated Employment Agreement dated July 30, 2010 between Aeolus Pharmaceuticals, Inc. and John L. McManus		08/02/10	8-K	10.4

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Exhibit Number	Description of Document	Incorporated by Reference To		Exhibit Number	Filed Herewith
		Registrant's Form	Date Filed with the SEC		
10.26+	Letter Agreement dated July 10, 2006 between Aeolus Pharmaceuticals, Inc. and McManus & Company, Inc.			8-K	07/10/06 10.2
10.27+	Form of Indemnity Agreement			10-K	12/27/11 10.27
10.28	Terms of Outside Director Compensation			10-K	12/17/04 10.114
10.29+	Form of Incentive Stock Option Agreement			10-Q	02/08/05 10.115
10.30+	Form of Nonqualified Stock Option Agreement			10-Q	02/08/05 10.116
10.31	Subscription Agreement dated June 5, 2006 by and between the Company and the investors whose names appear on the signature pages thereof.			8-K	06/06/06 10.1
10.32	Board Observer Letter dated June 5, 2006 by and among the Company and Efficacy Biotech Master Fund Ltd.			8-K	06/06/06 10.6
10.33+	Consulting Agreement, dated December 1, 2010, between Aeolus Pharmaceuticals, Inc. and Brian J. Day			8-K	12/03/10 10.1
10.34*	Sponsored Research Agreement (Non-Clinical), dated April 12, 2011, by and between the Company and Duke University			10-Q	05/16/11 10.5
10.35	Securities Purchase Agreement dated August 11, 2010 by and among Aeolus Pharmaceuticals, Inc. and the investors listed therein			8-K	8/12/10 10.1
10.36	Form of Warrant pursuant to Securities Purchase Agreement dated August 11, 2010 by and among Aeolus Pharmaceuticals, Inc. and the investors listed therein			8-K	8/12/10 10.2
10.37	Convertible Promissory Note dated February 7, 2007 issued by Aeolus Pharmaceuticals, Inc. to Elan Pharma International Ltd.			S-1	06/04/07 10.43
10.38	Amendment No. 1 To Convertible Promissory Note dated February 7, 2009 by and between Aeolus Pharmaceuticals, Inc. and Elan Pharma International Limited			8-K	3/16/09 10.1
10.39+	Form of Restricted Share Award Agreement			S-8 POS	3/31/08 99.2
10.40	Securities Purchase and Exchange Agreement dated October 6, 2009 by and among the Company and the investors whose names appear on the signature pages thereof			8-K	10/06/09 10.1
10.41	Amendment Agreement to the Securities Purchase and Exchange Agreement, dated December 24, 2009, by and among the Company and the investors whose names appear on the signature pages thereof			8-K	12/28/09 10.1
10.42+	Intentionally Omitted			8-K	02/16/11 10.1
10.43*	Contract No. HHSO100201100007C, dated February 11, 2011, by and between the Company and the U.S. Department of Health and Human Services Biomedical Advanced Research and Development Authority			10-Q	05/16/11 10.1
10.44*	Research and Manufacturing Agreement, dated February 18, 2011 (the "JMPS Agreement"), by and between the Company and Johnson Matthey Pharmaceutical Materials, Inc. (d/b/a Johnson Matthey Pharma Services).			10-Q	05/16/11 10.2

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Exhibit Number	Description of Document	Incorporated by Reference To		Exhibit Number	Filed Herewith
		Registrant's Form	Date Filed with the SEC		
10.45*	Appendix 2 to the JMPS Agreement, dated February 18, 2011		10-Q 8/14/12		10.4
10.46*	Appendix 3 to the JMPS Agreement, dated April 30, 2012		10-Q 8/14/12		10.5
10.47*	Appendix 4 to the JMPS Agreement, dated April 30, 2012		10-Q 8/14/12		10.6
10.48*	Appendix 5 to the JMPS Agreement, dated April 30, 2012		10-Q 8/14/12		10.7
10.49*	Appendix 6 to the JMPS Agreement, dated April 30, 2012		10-Q 8/14/12		10.8
10.50*	General Management Consulting Assignment, dated February 23, 2011, by and between the Company and Booz Allen Hamilton Inc.		10-Q 05/16/11		10.3
10.51	Form of Securities Purchase Agreement by and among the Company and the investors whose names appear on the signature pages thereof		8-K 4/5/12		10.1
10.52	Form of Registration Rights Agreement by and among the Company and the investors party thereto		8-K 4/5/12		10.2
10.53	Form of Warrant issued to investors in March and April 2012		8-K 4/5/12		10.3
10.54	Amended and Restated Employment Agreement by and between the Company and John L. McManus		8-K 03/05/13		10.1
10.55	Form of Registration Rights Agreement		8-K 12/15/15		10.2
<u>21.1</u>	List of Subsidiaries				X
<u>23.1</u>	Consent of Haskell & White LLP, Independent Registered Public Accounting Firm				
<u>23.2</u>	Consent of Grant Thornton LLP, Independent Registered Public Accounting Firm				X
<u>31.1</u>	Certification of the Principal Executive Officer pursuant to Rule 13a-14(a) and 15d-14(a)				X
<u>31.2</u>	Certification of the Principal Financial and Accounting Officer pursuant to Rule 13a-14(a) and 15d-14(a)				X
<u>32.1</u>	Certification by the Principal Executive Officer and Principal Financial and Accounting Officer pursuant to 18 U.S.C. 1350 as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002				X
101.INS†	XBRL Instance Document				X
101.SCH†	XBRL Taxonomy Extension Schema Document				X
101.CAL†	XBRL Taxonomy Extension Calculation Linkbase Document				X
101.DEF†	XBRL Taxonomy Extension Definition Linkbase Document				X
101.LAB†	XBRL Taxonomy Extension Label Linkbase Document				X
101.PRE†	XBRL Taxonomy Extension Presentation Linkbase Document				X

* The Company has received confidential treatment of certain portions of this agreement which have been omitted and filed separately with the U.S. Securities and Exchange Commission.

+ Indicates management contract or compensatory plan or arrangement.

† Attached as Exhibit 101 to this report are documents formatted in XBRL (Extensible Business Reporting Language). Users of this data are advised that, pursuant to Rule 406T of Regulation S-T, the interactive data file is deemed not filed or part of a registration statement or prospectus for purposes of Sections 11 or 12 of the Securities Act of 1933, as amended, is deemed not filed for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, and is otherwise not subject to liability under these sections.

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SIGNATURES

Pursuant to the requirements of Section 13 or 15(d) of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

AEOLUS PHARMACEUTICALS, INC.

By: /s/ John L. McManus

John L. McManus

President and Chief Executive Officer

Date: December 20, 2016

Pursuant to the requirements of the Securities Exchange Act of 1934, this report has been signed below by the following persons on behalf of the registrant and in the capacities and on the dates indicated.

SIGNATURE	TITLE(S)	DATE
/s/ John L. McManus John L. McManus	President and Chief Executive Officer (Principal Executive Officer)	December 20, 2016
/s/ David C. Cavalier David C. Cavalier	Chief Financial Officer and Secretary (Principal Financial and Accounting Officer)	December 20, 2016
/s/ David C. Cavalier David C. Cavalier	Chairman of the Board of Directors	December 20, 2016
/s/ John M. Clerici John M. Clerici	Director	December 20, 2016
/s/ John M. Farah, Jr. John M. Farah, Jr., Ph.D.	Director	December 20, 2016
/s/ Mitchell D. Kaye Mitchell D. Kaye	Director	December 20, 2016
/s/ Amit Kumar Amit Kumar, Ph.D.	Director	December 20, 2016
/s/ Chris A. Rallis Chris A. Rallis	Director	December 20, 2016
Jeffery Scott	Director	December 20, 2016