XOMA LTD /DE/ Form FWP January 10, 2006

A Leader in Therapeutic Antibodies January 11, 2006 JP Morgan Conference Filed Pursuant to Rule 433 Registration Nos. 333-130442 January 10, 2006

2 NASDAQ: XOMA

Statements made in this presentation relating to future financial performance or results, the timing of regulatory filings, the timing and results of clinical trials and other aspects of product development, collaborative and other strategic relationships, the regulatory process and approvals, collaboration and licensing

opportunities

and

plans

for

sales

and

marketing,

or

that

Eugai Filling. AOIVIA LTD/DE/ - FOITH FWF
otherwise
relate
to
future
periods,
are
forward-looking
statements
within
the
meaning
of
Section
27A
of
the
Securities
Act
of
1933
and
Section
21E of the Securities Exchange Act of 1934.
These statements are based on assumptions which may not prove accurate. Actual results could differ
materially from those anticipated, due to certain risks inherent
in the biotechnology industry, as well as
for companies engaged in the development of new products in a regulated market.
These risks, including those related to the success of the sales and marketing efforts for our products,
the size and timing of expenditures, whether there are unanticipated expenditures and whether funds are
available on acceptable terms; safety or efficacy of the products being tested; design and progress of
clinical trials; additional time requirements in connection with
regulatory filings for data analysis, filing
preparation, discussions with the FDA, additional clinical studies or manufacturing process
modifications;
action,
inaction
or
delays
by
the EDA
FDA,
European
regulators and/or
their advisory
advisory bodies;
analysis and interpretation by, or submission to, these entities and others of scientific data; results of
pre-clinical
pre-eninear

testing;
changes
in
the
status
of
the
Company's
collaborative
and
other
relationships;
the
ability of partners to meet their obligations; availability of collaboration and licensing opportunities;
success of competitors; market demand for products; uncertainties regarding biotechnology patents;
uncertainties as to the costs of protecting intellectual property; and risks associated with XOMA's
status
as
a
Bermuda
company
are
discussed
in
the
Company's
most
recent
report
on
Form
10-K
and
in
other
SEC filings.
Such risks should be considered carefully in evaluating XOMA's
prospects.
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Forward-Looking Statements

3
NASDAQ: XOMA
XOMA,
Ltd.
has
filed
a
registration
statement
(including

prospectus) with the **SEC** for the offering to which this communication relates. Before you invest, you should read the prospectus in that registration statement and other documents XOMA has filed with the SEC for more complete information about XOMA and this offering. You may get these documents for free by visiting EDGAR on the SEC Web site at www.sec.gov . Alternatively, **XOMA** or the information agent will arrange to send you the prospectus if you

request

it by calling toll-free 1-888-867-6963.

NASDAQ: XOMA
Premier Therapeutic Antibody
Discovery and Development Company
Managing Development and Financial
Risks, Effectively Utilizing Assets
Marketed Product plus Diverse,
Growing Pipeline
XOMA
Right Place, Right Time, By Design

NASDAQ: XOMA
Antibody Technologies
Antibody Technologies
Comprehensive Antibody Platform for
Comprehensive Antibody Platform for
Discovery, Optimization and Manufacturing
Discovery, Optimization and Manufacturing
Multiple Antibody Phage Display Libraries
Proprietary
Human
Engineering TM
Technology
Bacterial Cell Expression

NASDAQ: XOMA

XOMA

Human

Engineering

Technology

Clinically Validated

100% Success Rate To Date

25 mAbs Human Engineered

6 Different Targets

Structural Approach

Applicable to mAbs from any Species

Issued IP

Reduce Immunogenicity

Reduce Immunogenicity

of Non-human mAbs of Non-human mAbs

NASDAQ: XOMA

Therapeutic Antibody Development

Therapeutic Antibody Development

Antibody Lead

Discovery

Preclinical

Development

Clinical

Development

Tech Dev

Mfg

Fully Integrated Development Infrastructure

Fully Integrated Development Infrastructure

Target

Discovery
Functional Biology
Pharmacology (Efficacy, MOA)
Toxicology (IND-enabling safety)
Cell Line and Process Development
Clinical & Regulatory
Pilot Plant and GMP Manufacturing

8
NASDAQ: XOMA
Process Development
Process Development
Cell Line Development
Cell Banking
Pilot Plant Production
Assay Development
Formulation Development
Manufacturing
Manufacturing
cGMP
Production
Scale to 2750 L 3 Trains

Grams to Kilograms 15,000 sq ft Pilot Plant GMP Manufacturing Plant XOMA s Integrated Development and Manufacturing Capability 9 NASDAQ: XOMA Recognized Leader in Technology and Capabilities Leverage Technologies and Capabilities Manage Development and Financial Risks XOMA Strategy

NASDAQ: XOMA

Benefits from Collaborations

Chiron provides validated antibody targets

and leads for oncology products

Goal: 1 IND each year

Lexicon provides validated antibody targets Goal: Minimum of 3 products in 3 years

Bring More Product

Candidates

into

XOMA s Pipeline

Utilize Complementary Capabilities from XOMA and Partners

Manage Financial Risk
Share Development Cost
Utilize XOMA Infrastructure

Provide Other Financial Resources