

MacroGenics Inc

COMPANY AND PIPELINE OVERVIEW REPORT

A comprehensive coverage of the company and a summary of the drug pipeline portfolio.

Publication Date: 06-Aug-2014

THOMSON REUTERS

3 Times Square New York, New York 10036 United States

Tel: +1 646 223 4000

thomsonreuters.com



ABOUT COMPANY AND PIPELINE OVERVIEW REPORT

Thomson Reuters provides the knowledge, tools, and expertise to help support drug discovery and development activities, IP portfolio optimization, identification of licensing and partnering opportunities, delivery of successful regulatory submissions, and the ability to keep current with the rapidly-changing pharmaceutical and chemical markets, supporting informed, early decisions.

This report was created by Thomson Reuters, using information from *Thomson Reuters Cortellis™ for Competitive Intelligence*; a comprehensive, proven intelligence solution that leverages the most accurate, complete, and widely respected drug pipeline information. From drug discovery and development activities to patent reports, the latest deals, and partnering opportunities, *Cortellis* can provide the confidence to make the most informed business decisions, faster. *Cortellis for Competitive Intelligence* provides accurate and validated information on pharmaceutical and biotechnology companies globally, their drug pipelines, deals, patents, and clinical trials, plus breaking industry news and conference coverage. All contained in one simple, highly intuitive research platform.

Cortellis Company and Pipeline Overview reports are the first in a series of reports that track pharmaceutical and biotechnology companies worldwide. Further report offerings planned to follow include: Company Detailed Pipeline and Company Competitive Landscape reports. All Cortellis for Competitive Intelligence content is subject to the most comprehensive editorial review process available, conducted by scientists, pharma professionals, regulatory experts, and generics specialists. Featuring timely drug pipeline information expertly uncovered and integrated from a significant number of global meetings each year, you'll always be on top of the latest developments.

Chosen by leading life sciences companies, their executives and investors, *Cortellis for Competitive Intelligence* accelerates your deal-making and gives you timely insights on the development landscape.

Discover undiscovered opportunities in drug development and licensing faster with *Thomson Reuters Cortellis™ for Competitive Intelligence*

DISCLAIMER

The information contained in this report is based on sources believed to be correct but Thomson Reuters does not guarantee the accuracy, timeliness, or completeness of this information. Opinions, if any, are those held by the author of any individual report or article at the time of initial publication and do not necessarily reflect the views of Thomson Reuters.

Information in this report on companies is intended for reference use only, and does not constitute a recommendation to buy or sell any particular security or other investment and does not constitute an offer to buy from or sell to any particular investor. Any company or securities mentioned in this report may not be suitable for any particular investor, depending on that investor's financial position and needs.



GLOSSARY

Number of Drugs in Active Development

Number of drugs associated with the company or subsidiary that are currently in active development, i.e. the development status for the drug(s) is one of the following: Discovery, Clinical, Phase I, Phase II, Phase III, Pre-registration, Registered, Launched, or Suspended.

Number of Inactive Drugs

Number of drugs associated with the company or subsidiary that are currently classified as inactive, i.e. where the development status for the drug(s) is one of the following: No Development Reported, Discontinued, or Withdrawn.

Number of Patents as Owner

Number of patents associated with the company where the company is listed as owner; i.e. the relationship type (or way the patent refers to the company) is: Patent Assignee/Owner, Patent owner (not assignee), Licensee for development and marketing, Licensee – marketing only (Distributor), Patent assignee of family member, Inferred assignee.

Number of Patents as Third Party

Number of patents associated with the company where the company is listed as third party; i.e. the relationship type (or way the patent refers to the company) is: Patent assignee (not owner), Ex-Licensee for development and marketing, Ex-Licensee marketing only (Distributor), Customer of technology, Ex-Customer of technology, Patent opponent or infringer, Affiliate organization of inventor, Owner of underlying technology.

Patents summary table

This table represents a summary of the core patent coverage for this company covering Therapeutic EP, US and WO patents since 1990 only.

Number of Deals

A count of deals where the company or one of its subsidiaries is the primary company.

Key Indications

Displays top ten key indications for the company and its subsidiaries based on frequency (indications occurring with high and identical frequency are always included, and this may result in more than ten Key Indications being listed). Includes both indications associated with patents where the company is patent owner and indications associated with drugs in active development. A drug is classified as 'active' if it features on a row (or rows) in the current development status table where the status is one of the following: Discovery, Clinical, Phase I, Phase II, Phase III, Pre-registration, Registered, Launched, or Suspended.

Key Target-based Actions

Displays top ten key target-based actions for the company and its subsidiaries based on frequency (actions occurring with high and identical frequency are always included, and this may result in more than ten Key Target-based Actions being listed). Includes both target-based actions associated with patents where the company patent owner and target-based actions associated with drugs in active development. A drug is classified as 'active' if it features on a row (or rows) in the current development status table where the status is one of the following: Discovery, Clinical, Phase I, Phase II, Phase III, Pre-registration, Registered, Launched, or Suspended. A target-based action is one that is associated with a target.

Key Technologies

Displays top ten key technologies for the company and its subsidiaries based on frequency (technologies occurring with high and identical frequency are always included, and this may result in more than ten Key Technologies being listed). Includes both key technologies associated with patents where the company relationship is patent owner and key technologies associated with drugs in active development. A drug is classified as 'active' if it features on a row (or rows) in the current development status table where the status is one of the following: Discovery, Clinical, Phase I, Phase II, Phase III, Pre-registration, Registered, Launched, or Suspended.

THOMSON REUTERS

PLEASE NOTE: the financials section where present in the report includes: Share Information, Stock Performance chart (including the consensus recommendation), and Major Shareholders. Financial Performance presents graphical and tabular data on Worldwide Sales, Operating Income and Net Income over time, together with a Quarterly earnings update. Balance Sheet lists Assets, Liabilities and Stockholders Equity, and Forecasts includes: EPS Forecast and Other Forecasts. The data reported in all sections (except share information and major stock holders) are correct as of the publication date of the report (and not the download date). For share information and major stock holders the data is correct for the date shown with these sections



TABLE OF CONTENTS

Company Overview	7
Company Profile	7
Subsidiary Companies	8
Product Portfolio Summary	Ĝ
Product Portfolio Drugs	14



MacroGenics Inc

COMPANY OVERVIEW

Company Name	MacroGenics Inc
Parent Company Name	MacroGenics Inc
Website	http://www.macrogenics.com/
Country	US
Number of Drugs in Active Development	13
Number of Inactive Drugs	14
Number of Patents as Owner	56
Number of Patents as Third Party	1
Number of Deals	34
Key Indications	Cancer, Autoimmune disease, Solid tumor, Insulin dependent diabetes, Breast tumor, Inflammatory disease, Rheumatoid arthritis, Systemic lupus erythematosus, Acute myelogenous leukemia, Chikungunya virus infection, Dengue virus infection, Esophagus tumor, Influenza virus infection, Transplant rejection, Variola virus infection
Key Target-based Actions	Immunoglobulin gamma Fc receptor III antagonist,CD276 antigen inhibitor,Erbb2 tyrosine kinase receptor inhibitor,Immunoglobulin gamma Fc receptor II antagonist,Immunoglobulin gamma Fc receptor II modulator,B-lymphocyte antigen CD20 inhibitor,Insulin-like growth factor 1 antagonist,ADAM-9 modulator,B-lymphocyte antigen CD19 modulator,CD79b agonist,CD79b modulator,CDw123 modulator,Carboxypeptidase inhibitor,Cell surface A33 antigen modulator,Epidermal growth factor antagonist,Immunoglobulin gamma Fc receptor II agonist,Oncostatin M
Key Technologies	Biological therapeutic, Parenteral formulation unspecified, Monoclonal antibody humanized, Antibody, Multivalent antibody, Monoclonal antibody, Intravenous formulation, Injectable formulation, Humanized antibody, Chimeric antibody, Drug combination, Immunoglobulin-G

COMPANY PROFILE

SUMMARY

MacroGenics Inc, incorporated in August 2000 and based in Rockville, Maryland, is focused on the development of novel immunotherapeutics. Product candidates include monoclonal antibodies and vaccines for the treatment and prevention of cancer, autoimmune and infectious disease. Programs are derived from both in-house discoveries and inlicensed candidates.

COMPANY LOCATION

MacroGenics is headquartered in Rockville, Maryland. The company has developed platform technology at three sites. These include protein engineering and animal modeling in Rockville, proteomics and target discovery in Seattle, Washington and genetic immunization and vaccine development in Dallas, Texas. In October 2005, MacroGenics opened a cGMP manufacturing facility in Rockville, MD. In February 2014, MacroGenics planned to expand its manufacturing facility, to increase its production capacity, using approximately US \$5 to 10 million.

ACQUISITIONS AND SPIN-OFFS

In July 2008, MacroGenics acquired Raven Biotechnologies. The company would issue shares of its common stock in the transaction but further financial details were not disclosed.

In June 2002, MacroGenics acquired Eliance Biotechnology Inc, a private vaccine discovery company located in Dallas,TX, which was founded to exploit technology developed at The University of Texas Southwestern Medical Center at Dallas Center for Biomedical Inventions. The company and university also formed an alliance to discover and develop immunotherapeutics to prevent and treat cancer, autoimmune and infectious diseases. In the same month, MacroGenics raised \$12.6 million in financing.



LICENSING AGREEMENTS

In October 2004, MacroGenics acquired exclusive rights to develop compounds against a cancer target identified using OriGene Technologies Inc's Rapid-Scan technology. MacroGenics was to focus on the development of therapeutic monoclonal antibody candidates against this molecule. OriGene would receive option fees, milestones and royalty payments.

In April 2004, Neose Technologies Inc and MacroGenics entered into a monoclonal antibodies research collaboration and license agreement. Neose was to apply its GlycoAdvance and GlycoPEGylation technologies to improve the therapeutic properties of MacroGenics compounds. MacroGenics had the right to take remodeled compounds into development. Following the initial research phase, MacroGenics was to be responsible for funding the further development of these licensed compounds. In exchange, Neose was to be entitled to receive option fee, milestone, and royalty payments as products were developed and commercialized.

In May 2002, MacroGenics licensed a therapeutic target induced by Epstein-Barr virus from EBVax Inc and Tufts University. MacroGenics was to launch a research program to identify treatments to reduce morbidity in patients suffering from the acute and chronic consequences of EBV infection. For a worldwide, exclusive license to the technology, MacroGenics agreed to certain upfront fees, milestone and royalty payments contingent on achieving specific preclinical and clinical objectives in different clinical indications.

In September 2001, MacroGenics reported the completion of several key license agreements relating to its core intellectual property.

EARLY R&D

IN OCTOBER 2010, BOEHRINGER INGELHEIM AND MACROGENICS ENTERED A GLOBAL ALLIANCE TO DISCOVER, DEVELOP AND COMMERCIALIZE ANTIBODY THERAPEUTICS, BASED ON MACROGENICS' DUAL-AFFINITY RE-TARGETING (DART) PLATFORM, WHICH WOULD BE DIRECTED AGAINST UP TO TEN COMBINATIONS OF MOLECULAR TARGETS. THERAPEUTIC AREAS FOR THE DRUG CANDIDATES COULD INCLUDE IMMUNOLOGY, ONCOLOGY, RESPIRATORY, CARDIOMETABOLIC AND INFECTIOUS DISEASES. THE COMPANIES WOULD SHARE DISCOVERY AND CERTAIN PRECLINICAL ACTIVITY RESPONSIBILITY. AFTERWHICH, BOEHRINGER INGELHEIM WOULD BE SOLELY RESPONSIBLE FOR FURTHER PRECLINICAL, CLINICAL, REGULATORY, COMMERCIAL AND MANUFACTURING ACTIVITIES OF ANY RESULTING PRODUCTS. MACROGENICS WAS EXPECTED TO RECEIVE \$60 MILLION, COMPRISED OF AN UPFRONT CASH PAYMENT, ANNUAL MAINTENANCE FEES, R&D FUNDING, AND NEAR-TERM RESEARCH-BASED MILESTONES, DURING THE FIRST THREE YEARS OF THE ALLIANCE. BOEHRINGER INGELHEIM ALSO EXPECTED TO MAKE A FUTURE EQUITY INVESTMENT IN MACROGENICS. FURTHERMORE, MACROGENICS COULD RECEIVE DEVELOPMENT, REGULATORY AND COMMERCIAL MILESTONE PAYMENTS OF UP TO \$210 MILLION FOR EACH OF THE TEN DART PROGRAMS, PLUS TIERED ROYALTIES. MACROGENICS WOULD HAVE AN OPTION TO COPROMOTE CERTAIN DART PRODUCTS IN THE US.

FINANCIAL

In February 2014, MacroGenics commenced an underwritten public offering of 2.5 million shares of its common stock, consisting of 1.5 million shares to be offered by MacroGenics and 1 million shares to be offered by existing stockholders. The underwriters were to be granted a 30-day option to purchase up to 375,000 shares of common stock; later that month, the company priced the offering of 3 million shares of its common stock, consisting of 1.8 million shares to be offered by MacroGenics and 1.2 million shares to be offered by existing stockholders at a price of \$36.50 per share. The underwriters were granted a 30-day option to purchase up to 450,000 shares of common stock. The offering was expected to close on February 18, 2014. Later in February 2014, the offering was completed for net proceeds of \$76.7 million, including \$15.4 million of net proceeds from full exercise of underwriters' over-allotment option.

In October 2013, MacroGenics offered an initial public offering of 5 million shares of its common stock, priced at \$16 per share. Underwriters were granted a 30-day option to purchase up to an additional 750,000 shares of common stock to cover any over-allotments. The shares were to be traded on the NASDAQ Global Select Market under the ticker symbol "MGNX". The offering was expected to close on October 16, 2013. Later that month, shares of MacroGenics began trading on the NASDAQ Global Select Market. Later that month, the offering was closed and the company raised \$83.8 million from the offering. The company offered 5,750,000 million shares of its common stock, at a price of \$16 per share. Underwriters were granted a 30-day option to purchase up to an additional 750,000 shares of common stock to cover any over-allotments.

In September 2008, MacroGenics raised \$25 million from a series D-2 financing round.

In May 2006, MacroGenics raised \$45 million from a series C financing round. The proceeds were to be used to fund development of the company's anti-CD3 antibody; in phase II trials for type 1 diabetes and to advance drug programs in autoimmune diseases and cancer.



SUBSIDIARY COMPANIES

Names

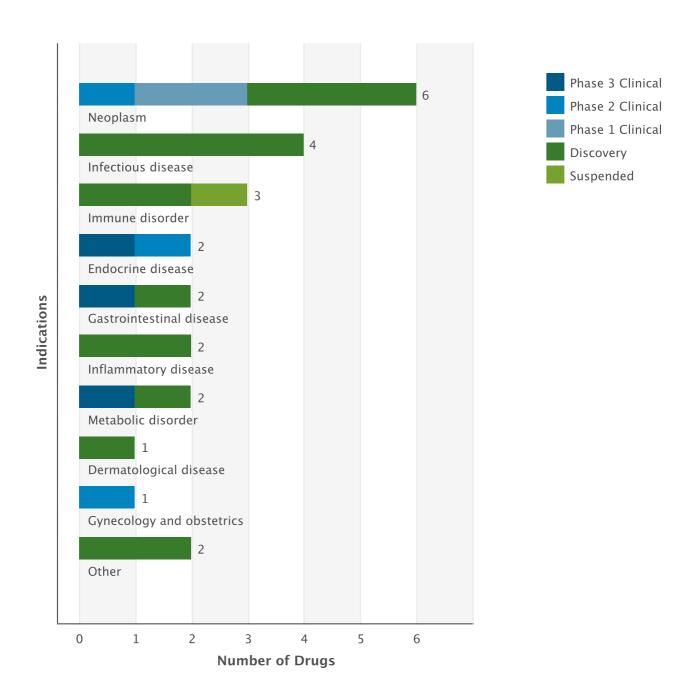
Raven Biotechnologies Inc

PRODUCT PORTFOLIO SUMMARY

DRUGS

Drugs by Indication

Active Drugs by Indication Chart





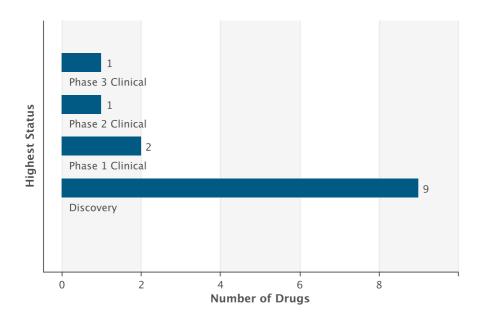
Drugs by Indication Table

Indication	Active	Inactive	Total
Neoplasm	6	10	16
Immune disorder	3	4	7
Infectious disease	4	2	6
Gastrointestinal disease	2	3	5
Respiratory disease	1	3	4
Inflammatory disease	2	1	3
Genitourinary disease	0	3	3
Endocrine disease	2	1	3
Hematological disease	1	2	3
Neurological disease	0	2	2
Dermatological disease	1	1	2
Metabolic disorder	2	0	2
Musculoskeletal disease	1	1	2
Gynecology and obstetrics	1	1	2
Andrology	0	1	1



Drugs by Highest Status

Active Drugs by Highest Status Chart



Drugs by Highest Status Table

Development Status	Number of Drugs
Phase 3 Clinical	1
Phase 2 Clinical	1
Phase 1 Clinical	2
Discovery	9
Discontinued	3
No Development Reported	11



DEALS

Deal Type	Prir	ncipal	Pai	tner	Total
	Active	Inactive	Active	Inactive	
Technology - Other Proprietary	3	0	1	0	4
Drug - Funding	6	0	0	0	6
Drug - Screening/Evaluation	2	0	0	0	2
Drug - Early Research/Development	0	0	3	0	3
Drug - Development/Commercialization License	10	0	4	0	15
Drug - Manufacturing/Supply	0	0	2	0	2
Drug - Development Services	0	0	1	0	1
Technology - Delivery/Formulation	0	0	1	0	1

CLINICAL TRIALS

Trials by Condition Studied

Condition Studied	Ongoing	All
Endocrine disease	1	10
Gastrointestinal disease	0	9
Metabolic disorder	0	7
Neoplasm	2	7
Immune disorder	0	3
Infectious disease	0	3
Neurological disease	0	3
Inflammatory disease	0	3
Gynecology and obstetrics	1	2
Dermatological disease	0	2
Musculoskeletal disease	0	2
Hematological disease	1	1
Genitourinary disease	0	1
Respiratory disease	0	1



Trials by Phase

Phase	Ongoing	All
Phase 3	0	2
Phase 2	1	6
Phase 1	1	11
Phase not specified	0	1

Phase Definitions

Phase 3 Clinical

Includes Phase 3, Phase 3b, Phase 3a, Phase 2/3 (where enrolment count is 300 or over)

Phase 2 Clinical

Includes Phase 2, Phase 2a, Phase 2b, Phase 1/2 (where enrolment count is 100 or over), Phase 2/3 (where enrolment count is under 300 or not specified)

Phase 1 Clinical

Includes Phase 1, Phase 1a, Phase 1, Phase 1/2 (where enrolment count is under 100 or not specified), Phase 0

PATENTS *

Indication	As Owner	As Third Party	Total
Cardiovascular disease	2	0	2
Endocrine disease	19	1	20
Gastrointestinal disease	19	1	20
Genitourinary disease	12	1	13
Hematological disease	9	0	9
Andrology	8	0	8
Immune disorder	31	0	31
Musculoskeletal disease	13	0	13
Neoplasm	43	1	44
Ocular disease	3	0	3
Metabolic disorder	10	0	10
Mouth disease	1	0	1
Neurological disease	10	0	10
Respiratory disease	10	1	11
Infectious disease	18	0	18



Injury	1	0	1
Toxicity and intoxication	1	0	1
Inflammatory disease	23	0	23
Otorhinolaryngological disease	1	0	1
Gynecology and obstetrics	13	1	14
Dermatological disease	13	1	14
Surgical procedure	1	0	1

^{*} This table represents a summary of the core patent coverage for this company covering Therapeutic EP, US and WO patents since 1990 only.



PRODUCT PORTFOLIO DRUGS

PLEASE NOTE: Highest status refers to highest development of that drug for one of the active companies

teplizumab

Drug Name	teplizumab
Key Synonyms	teplizumab
Originator Company	Tolera Therapeutics Inc
Active Companies	MacroGenics Inc
Inactive Companies	Tolera Therapeutics Inc, Eli Lilly & Co
Highest Status	Phase 3 Clinical
Active Indications	Transplant rejection, Insulin dependent diabetes
Target-based Actions	
Other Actions	Anti-inflammatory, Immunosuppressant, Hypoglycemic agent, CD3 antagonist
Technologies	Monoclonal antibody, Subcutaneous formulation, Intravenous formulation, Infusion, Biological therapeutic
Last Change Date	04-Dec-2013

margetuximab

Drug Name	margetuximab
Key Synonyms	margetuximab
Originator Company	MacroGenics Inc
Active Companies	MacroGenics Inc, Green Cross Corp
Inactive Companies	
Highest Status	Phase 2 Clinical
Active Indications	Esophagus tumor, Solid tumor, Breast tumor
Target-based Actions	Erbb2 tyrosine kinase receptor inhibitor
Other Actions	Anticancer protein kinase inhibitor, Anticancer monoclonal antibody
Technologies	Monoclonal antibody, Intravenous formulation, Infusion, Biological therapeutic
Last Change Date	19-Jun-2014

MGA-271

Drug Name	MGA-271
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	Servier, MacroGenics Inc
Inactive Companies	
Highest Status	Phase 1 Clinical
Active Indications	Solid tumor
Target-based Actions	CD276 antigen inhibitor
Other Actions	Immunostimulant, Anticancer monoclonal antibody
Technologies	Monoclonal antibody humanized, Immunoglobulin-G, Intravenous formulation, Biological therapeutic, Protein recombinant
Last Change Date	08-May-2014

MGD-006

Drug Name	MGD-006
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	Servier, MacroGenics Inc
Inactive Companies	
Highest Status	Phase 1 Clinical
Active Indications	Acute myelogenous leukemia
Target-based Actions	CDw123 modulator
Other Actions	CD3 modulator, Anticancer protein kinase inhibitor
Technologies	Multivalent antibody, Humanized antibody, Injectable formulation, Biological therapeutic, Parenteral formulation unspecified
Last Change Date	22-Jul-2014

monoclonal antibody cocktail (DART, smallpox), MacroGenics

Drug Name	monoclonal antibody cocktail (DART, smallpox), MacroGenics
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	MacroGenics Inc
Inactive Companies	
Highest Status	Discovery
Active Indications	Variola virus infection
Target-based Actions	
Other Actions	Antiviral, Unspecified drug target
Technologies	Monoclonal antibody humanized, Drug combination, Biological therapeutic, Parenteral formulation unspecified
Last Change Date	14-Feb-2013

H5N1 influenza mAb therapy, MacroGenics

Drug Name	H5N1 influenza mAb therapy, MacroGenics
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	MacroGenics Inc
Inactive Companies	
Highest Status	Discovery
Active Indications	Influenza virus infection
Target-based Actions	
Other Actions	Unspecified drug target, Antiviral
Technologies	Monoclonal antibody humanized, Biological therapeutic, Parenteral formulation unspecified
Last Change Date	15-Feb-2013

CD32BxCD79B DART antibody (autoimmune disease), MacroGenics

Drug Name	CD32BxCD79B DART antibody (autoimmune disease), MacroGenics
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	MacroGenics Inc
Inactive Companies	
Highest Status	Discovery
Active Indications	Inflammatory disease, Autoimmune disease
Target-based Actions	Immunoglobulin gamma Fc receptor II agonist, CD79b agonist
Other Actions	Anti-inflammatory, Anticancer
Technologies	Antibody, Biological therapeutic
Last Change Date	04-Dec-2013

monoclonal antibody (DART, dengue virus infection), MacroGenics

Drug Name	monoclonal antibody (DART, dengue virus infection), MacroGenics
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	MacroGenics Inc
Inactive Companies	
Highest Status	Discovery
Active Indications	Dengue virus infection
Target-based Actions	
Other Actions	Antiviral, Unspecified drug target
Technologies	Immunoglobulin, Monoclonal antibody, Biological therapeutic, Parenteral formulation unspecified
Last Change Date	15-Feb-2013



bispecific antibodies (DART, cancer), MacroGenics/ Pfizer

Drug Name	bispecific antibodies (DART, cancer), MacroGenics/ Pfizer
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	Pfizer Inc, MacroGenics Inc
Inactive Companies	
Highest Status	Discovery
Active Indications	Cancer
Target-based Actions	B-lymphocyte antigen CD19 modulator, Epidermal growth factor antagonist, Insulin-like growth factor 1 antagonist
Other Actions	Anticancer protein kinase inhibitor, Anticancer antibody, CD3 agonist
Technologies	Multivalent antibody, Biological therapeutic
Last Change Date	04-Dec-2013

humanized mAbs (Chikungunya virus infection), MacroGenics/Washington University

Drug Name	humanized mAbs (Chikungunya virus infection), MacroGenics/Washington University
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	Washington University in St Louis, MacroGenics Inc
Inactive Companies	
Highest Status	Discovery
Active Indications	Chikungunya virus infection
Target-based Actions	
Other Actions	Antiviral, Unspecified drug target
Technologies	Monoclonal antibody humanized, Biological therapeutic, Parenteral formulation unspecified
Last Change Date	14-Feb-2013



MGD-007

Drug Name	MGD-007
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	MacroGenics Inc, Servier
Inactive Companies	
Highest Status	Discovery
Active Indications	Cancer
Target-based Actions	Cell surface A33 antigen modulator
Other Actions	CD3 modulator, Anticancer antibody
Technologies	Multivalent antibody, Injectable formulation, Biological therapeutic, Parenteral formulation unspecified
Last Change Date	25-Jul-2014

MGD-010

Drug Name	MGD-010
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	MacroGenics Inc
Inactive Companies	
Highest Status	Discovery
Active Indications	Autoimmune disease, Systemic lupus erythematosus, Rheumatoid arthritis
Target-based Actions	Immunoglobulin gamma Fc receptor II modulator, CD79b modulator
Other Actions	Immunomodulator, Anti-inflammatory
Technologies	Multivalent antibody, Injectable formulation, Biological therapeutic, Parenteral formulation unspecified
Last Change Date	19-Jun-2014

anti-CD3/undisclosed antibodies (DART/injectable, cancer), Macrogenics/Servier

Drug Name	anti-CD3/undisclosed antibodies (DART/injectable, cancer), Macrogenics/Servier
Key Synonyms	
Originator Company	MacroGenics Inc
Active Companies	MacroGenics Inc, Servier
Inactive Companies	
Highest Status	Discovery
Active Indications	Cancer
Target-based Actions	
Other Actions	CD3 antagonist, Anticancer antibody
Technologies	Multivalent antibody, Injectable formulation, Biological therapeutic, Parenteral formulation unspecified
Last Change Date	04-Dec-2013



This report was created by Thomson Reuters, using information from *Thomson Reuters Cortellis*™ *for Competitive Intelligence*; a comprehensive, proven intelligence solution that leverages the most accurate, complete, and widely respected drug pipeline information.

For more information about *Cortellis for Competitive Intelligence*, visit: http://cortellis.thomsonreuters.com/cortellis_for_you/?cid=thomsonone.

For subscription information, e-mail scientific.lifesciences@thomsonreuters.com.

© 2012 Thomson Reuters. All rights reserved. Republication or redistribution of Thomson Reuters content, including by framing or similar means, is prohibited without the prior written consent of Thomson Reuters. 'Thomson Reuters' and the Thomson Reuters logo are registered trademarks and trademarks of Thomson Reuters and its affiliated companies.

THOMSON REUTERS