

Durata Therapeutics Inc

COMPANY AND PIPELINE OVERVIEW REPORT

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

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ABOUT COMPANY AND PIPELINE OVERVIEW REPORT

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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i.e. where the development status for the drug(s) is one of the following: No Development Reported, Discontinued, or Withdrawn.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.

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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	7
Product Portfolio Summary	7
Product Portfolio Drugs	1.



Durata Therapeutics Inc

COMPANY OVERVIEW

Company Name	Durata Therapeutics Inc
Parent Company Name	Durata Therapeutics Inc
Website	http://www.duratatherapeutics.com/
Country	US
Number of Drugs in Active Development	3
Number of Inactive Drugs	13
Number of Patents as Owner	24
Number of Patents as Third Party	2
Number of Deals	9
Key Indications	Bacterial infection,Bacterial skin infection,Gram positive bacterium infection,MRSA infection
Key Target-based Actions	Peptide deformylase inhibitor,N-acetylglucosamine deacetylase inhibitor,Unspecified enzyme inhibitor
Key Technologies	Small molecule therapeutic,Antibiotic,Oral formulation,Biological therapeutic,Dermatological formulation,Injectable formulation,Parenteral formulation unspecified,Peptide

COMPANY PROFILE

SUMMARY

Durata Therapeutics Inc, a biopharmaceutical company, was formed in 2009 by a 5-member venture capital syndicate for the development of antibiotic programs to treat infectious diseases.

ACQUISITIONS & SPIN-OFFS

In December 2009, Durata acquired Vicuron Pharmaceuticals from Pfizer. The transaction was funded through a stock purchase by New Leaf Venture Partners, Domain Associates, Aisling Capital, Sofinnova Ventures and Canaan Partners. Durata obtained rights to dalbavancin and two undisclosed preclinical antibiotic programs.

FINANCIAL

In March 2012, Durata planned to conduct an IPO; in July 2012, the company priced 7.5 million shares of common stock at \$9 each. The underwritters were granted 30-day option to purchase additional shares up to 1.125 million and the offering was expected to close on July 24, 2012. The company also reported that the shares would be traded on NASDAQ under the ticker symbol 'DRTX'; later that month, the IPO of 8.625 million common stock shares was closed before the underwriting discounts. By that time, a registration statement relating to these securities had been declared effective by the SEC.

SUBSIDIARY COMPANIES

Names

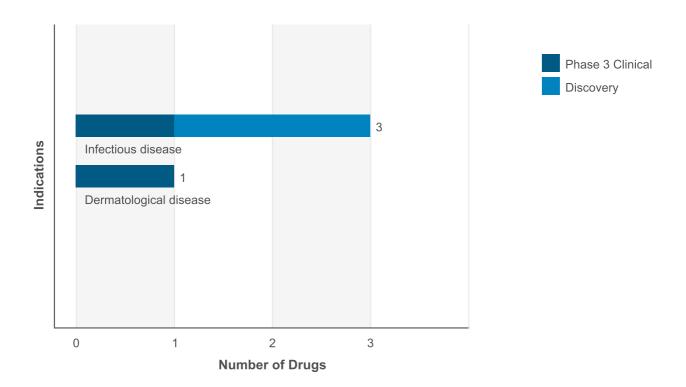
Vicuron Pharmaceuticals Inc

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PRODUCT PORTFOLIO SUMMARY DRUGS

Drugs by Indication

Active Drugs by Indication Chart



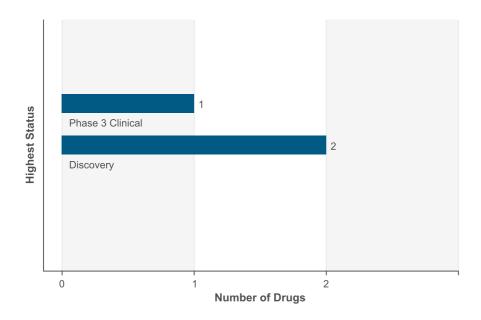
Drugs by Indication Table

Indication	Active	Inactive	Total
Infectious disease	3	11	14
Dermatological disease	1	2	3

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Drugs by Highest Status

Active Drugs by Highest Status Chart



Drugs by Highest Status Table

Development Status	Number of Drugs
Phase 3 Clinical	1
Discovery	2
Discontinued	2
No Development Reported	7

DEALS

Deal Type	Principal		Partner		Total
	Active	Inactive	Active	Inactive	
Technology - Other Proprietary	0	0	1	0	1
Drug - Asset Divestment	1	0	0	0	1
Drug - Early Research/Development	1	0	1	0	2
Drug - Development/Commercialization License	1	0	2	0	4
Drug - Manufacturing/Supply	0	0	1	0	1



CLINICAL TRIALS

Trials by Condition Studied

Condition Studied	Ongoing	All
Infectious disease	2	22
Dermatological disease	2	8

Trials by Phase

Phase	Ongoing	All
Phase 3	2	7
Phase 2	0	7
Phase 1	0	5
Phase not specified	0	7

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
Immune disorder	0	1	1
Genetic disorder	1	0	1
Metabolic disorder	1	0	1
Neurological disease	1	0	1
Degeneration	1	0	1
Dermatological disease	6	0	6
Gastrointestinal disease	2	1	3
Musculoskeletal disease	1	0	1
Respiratory disease	5	1	6
Infectious disease	22	2	24



Inflammatory disease	1	0	1	
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^{*} 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

dalbavancin

Drug Name	dalbavancin
Key Synonyms	dalbavancin, Zeven, Exulett
Originator Company	Biosearch Italia SpA
Active Companies	Durata Therapeutics Inc
Inactive Companies	RaQualia Pharma Inc, Pfizer Inc, Vicuron Pharmaceuticals Inc, Biosearch Italia SpA
Highest Status	Phase 3 Clinical
Active Indications	Gram positive bacterium infection, Bacterial skin infection, MRSA infection
Target-based Actions	
Other Actions	Bacterial cell wall synthesis inhibitor, Systemic dermatological antibacterial product, Antibacterial
Technologies	Intravenous formulation, Antibiotic, Glycoprotein, Infusion, Small molecule therapeutic
Last Change Date	29-Mar-2012

lincosamide antibacterials, Durata

Drug Name	lincosamide antibacterials, Durata
Key Synonyms	
Originator Company	Vicuron Pharmaceuticals Inc
Active Companies	Durata Therapeutics Inc
Inactive Companies	Vicuron Pharmaceuticals Inc
Highest Status	Discovery
Active Indications	Bacterial infection
Target-based Actions	
Other Actions	Bacterial protein synthesis inhibitor, Ribosome binding agent
Technologies	Antibiotic, Small molecule therapeutic
Last Change Date	23-Aug-2011

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pleuromutilins (bacterial infections), Durata

Drug Name	pleuromutilins (bacterial infections), Durata
Key Synonyms	
Originator Company	Vicuron Pharmaceuticals Inc
Active Companies	Durata Therapeutics Inc
Inactive Companies	
Highest Status	Discovery
Active Indications	Bacterial infection
Target-based Actions	
Other Actions	Antibacterial, Protein synthesis inhibitor
Technologies	Antibiotic, Small molecule therapeutic
Last Change Date	23-Aug-2011



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