

## **Xencor Inc**

### **COMPANY AND PIPELINE OVERVIEW REPORT**

Coverage of the company and a summary of the drug pipeline portfolio.

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i.e. where the development status for the drug(s) is one of the following: No Development Reported, Discontinued, or Withdrawn.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

## COMPANY OVERVIEW

Company Name	Xencor Inc
Parent Company Name	Xencor Inc
Website	<a href="http://www.xencor.com/">http://www.xencor.com/</a>
Country	US
Number of Drugs in Active Development	17
Number of Inactive Drugs	5
Number of Patents as Owner	55
Number of Patents as Third Party	1
Number of Deals	22
Key Indications	Cancer, Rheumatoid arthritis, Autoimmune disease, Inflammatory disease, Multiple myeloma, Non-Hodgkin lymphoma, Breast tumor, Multiple sclerosis, Systemic lupus erythematosus, Allergy, Chronic lymphocytic leukemia
Key Target-based Actions	B-lymphocyte antigen CD19 inhibitor, TNF antagonist, Immunoglobulin gamma Fc receptor IIB antagonist, TNF alpha ligand inhibitor, Immunoglobulin E antagonist, CD40 ligand inhibitor, B-lymphocyte antigen CD20 modulator, Bone marrow stromal antigen 2 modulator, CD30 antagonist, CD80 antagonist, CD86 antagonist, Cytotoxic T-lymphocyte protein-4 stimulator, EpCAM inhibitor, Epidermal growth factor ligand inhibitor, ErbB2 tyrosine kinase receptor modulator, Folate receptor alpha antagonist, Immunoglobulin gamma Fc receptor IIB modulator, Osteoclast differentiation factor ligand inhibitor, T-cell surface glycoprotein CD5 inhibitor, Thrombopoietin receptor agonist, VEGF ligand inhibitor
Key Technologies	Biological therapeutic, Parenteral formulation unspecified, Monoclonal antibody humanized, Protein recombinant, Small molecule therapeutic, Monoclonal antibody, Monoclonal antibody human, Chimeric monoclonal antibody, Peptide, Immunoglobulin-G

## COMPANY PROFILE

### SUMMARY

Xencor Inc is a privately held biotechnology company founded in October 1997 by Stephen Mayo and Bassil Dahiyat with \$5 million in venture capital. It was set up to create, develop and commercialize designed proteins as biotechnology products and research tools for the human therapeutic, high throughput screening assay and industrial enzyme markets. Xencor's proprietary tools are ProCode and Protein Design Automation (PDA). ProCode allows the simultaneous analysis of the entire proteome of a cell by creating protein expression libraries where each protein is covalently linked to its corresponding cDNA.

### TECHNOLOGY

Xmab technology is the company's proprietary platform to produce a portfolio of drug candidates from an antibody with high selectivity for a target antigen by enhancing the antibody-dependent cell cytotoxicity.

### LICENSING AGREEMENTS

In November 2009, Centocor Research & Development to use Xencor's XmAb and Xtend technologies to optimize its antibody drug candidates. Xencor was to receive an upfront payment and was eligible to receive milestones and royalties on product commercialization.

In March 2009, Xencor granted Merck & Co an exclusive license to its Xtend technology for the development of antibodies towards an undisclosed Merck drug target. Xencor would receive an upfront license fee of \$3 million, a payment on selection of an Xtend variant, milestone payments and royalties.

In March 2009, Xencor signed an agreement with Pfizer to optimize the performance of therapeutic monoclonal antibodies.

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In February 2009, Xencor licensed access to its XmAb technology to CSL, for use in advancing the antibody-dependent cell cytotoxicity (ADCC) effector function of CSL's own therapeutic antibodies. The initial phase of the alliance would allow CSL broad access to the technology, to apply towards its complete antibody portfolio. CSL also had commercial licenses in place for the advanced development and commercialization of any antibodies incorporating XmAb; Xencor would receive an upfront payment, plus milestone and royalty payments on each of these antibodies.

In February 2008, Xencor agreed to use its XmAb technology to optimize Human Genome Sciences's monoclonal antibodies against selected targets. Xencor would receive an upfront payment, and was eligible to receive additional milestone payments and royalties on any resulting products that Human Genome commercializes.

In February 2007, Xencor agreed to use its XmAb technology to optimize Boehringer Ingelheim's monoclonal antibodies against selected targets. Xencor would receive an upfront payment, and was eligible to receive additional licence fees, milestones and a royalty payment on any resulting products that Boehringer commercializes.

In February 2006, Xencor granted Centocor a non-exclusive multiyear license to use its ImmunoFilter technology to evaluate the potential immunogenicity of therapeutic proteins, in return for annual license fees.

In January 2006, Xencor granted Eli Lilly a non-exclusive, multiyear license to use its ImmunoFilter technology to evaluate the immunogenicity risk of Lilly's therapeutic proteins.

In December 2005, Xencor gained an exclusive license to technology developed at Cambridge University for making monoclonal antibodies with greatly enhanced potency. The technology included specific Fc variants that complement Xencor's XmAb engineered Fc domains, which could be inserted into antibody candidates against any antigen and may improve effector functions such as enhanced antibody-mediated tumor cell killing, sustained half-life and increased structural stability.

In December 2005, Xencor licensed its XmAb engineered Fc domains to MedImmune for use in the creation of monoclonal antibodies against preclinical tumor targets. Xencor received an upfront payment, as well as license fees, milestones and royalties.

In July 2005, Xencor and Centocor agreed to codevelop monoclonal antibodies with improved anticancer efficacy. Xencor would use its XmAb engineered antibody Fc domains to create drug candidates; Centocor would evaluate these against an oncology target. Xencor was to receive an upfront payment, license fees, and milestone and royalty payments.

In April 2005, Lilly exercised its option to develop one or more therapeutic proteins created by Xencor, which would receive an upfront license fee, milestone payments and royalties. In February 2004, Xencor and Lilly entered a collaboration to optimize the physical and biochemical properties of a protein therapeutic. Xencor would use its PDA technology to create variants of the protein to meet criteria for clinical development. Lilly would have the option to develop the resulting candidates.

In January 2005, Xencor and Roche Holding AG signed a collaboration agreement to use Xencor's XmAb with Roche's antibodies against a cancer target. Xencor would receive technology access and license fees, and future license fees, milestones and royalties if Roche advanced candidates into development; in January 2006, the companies extended the collaboration, allowing Roche to further its research efforts to enhance the therapeutic efficacy of an antibody against a cancer target. Xencor would receive additional fees for the extension.

In January 2005, Xencor and Chugai Pharmaceutical Co Ltd formed a collaboration to use Xencor's XmAb in combination with Chugai's antibodies against a cancer target. Xencor would receive technology access, license fees, milestones and royalties.

In December 2004, Xencor Inc and Genentech entered a collaboration to create next-generation therapeutic antibodies for cancer and autoimmune diseases. Xencor would grant Genentech an exclusive, worldwide license to use its XmAb technology, a suite of engineered antibody Fc domains that can be incorporated into therapeutic candidates, to develop and commercialize products directed against CD20, Her2 and a third undisclosed antigen. In return, Xencor would receive a \$5 million upfront fee, annual licensing fees and milestone payments.

In January 2004, Xencor and Protein Design Labs (PDL) entered into an agreement to generate monoclonal antibodies with enhanced potency. The multiyear collaboration allowed PDL to apply Xencor's XmAb technology to a number of preclinical antibodies against various targets. Xencor was to receive technology access and license fees, development milestone payments and royalties. PDL would be responsible for development and commercialization of the resulting products.

In September 2003, Chromos Molecular Systems Inc and Xencor entered into a non-exclusive research license agreement to develop cell lines that expressed Xencor's monoclonal antibodies and other recombinant protein product candidates. Chromos would receive an upfront payment and annual maintenance fees.

In October 2003, it was reported that Xencor was anticipating partnering a number of its technologies, whilst continuing to develop a number of inhouse projects.

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Xencor has ongoing collaborations with the California Institute of Technology (Caltech) and the Massachusetts Institute of Technology (MIT).

## IP NEWS

In January 2008, Xencor announced that the USPTO had granted US-7317091B1 (WO-03074679 and WO-2004029207) which covered its protein design automation technology. The issuance was the first from a series of applications covering Xencor's XmAb antibody Fc engineering technology.

In September 2003, Xencor was issued US-06627186, covering nucleic acids and protein variants of G-CSF with granulopoietic activity'.

## FINANCIAL

In October 2007, the company raised an additional \$15 million in extended series E financing, bringing the total raised to \$60 million.

In October 2006, Xencor raised \$45 million from a private financing. The funds included a \$6 million bridge financing that was announced in July 2006. Xencor would use the proceeds for the clinical development of two biologics; XPro1595 was to enter the clinic in 2006 for inflammation, and XmAb-2513 was to begin clinical trials for Hodgkin's disease in 2007.

In July 2006, Xencor raised \$6 million in a bridge financing led by Novo Nordisk, which committed to invest a further \$6 million in future financings.

By October 2005, Xencor had raised \$20 million through a series D round of financing. The funds would be used to initiate a phase I trial of XPro-1595 and conduct preclinical development of antibody therapeutics.

## R&D GRANTS

In October 2002, Xencor was awarded a \$2 million advanced technology program grant from the National Institute of Standards and Technology to support the discovery of safer and more effective protein therapeutics. The 3-year federal grant, titled "Rational Design of Non-immunogenic Proteins," would support the development of Xencor's ImmunoPDA technology to create new, non-immunogenic proteins optimized for therapeutic use and to eliminate immunogenicity from known protein drugs.

In June 2001, Xencor was awarded a \$0.5 million SBIR phase II grant from the National Science Foundation. The grant would allow Xencor to further develop its enabling technology for computer directed high-throughput screening of proteins.

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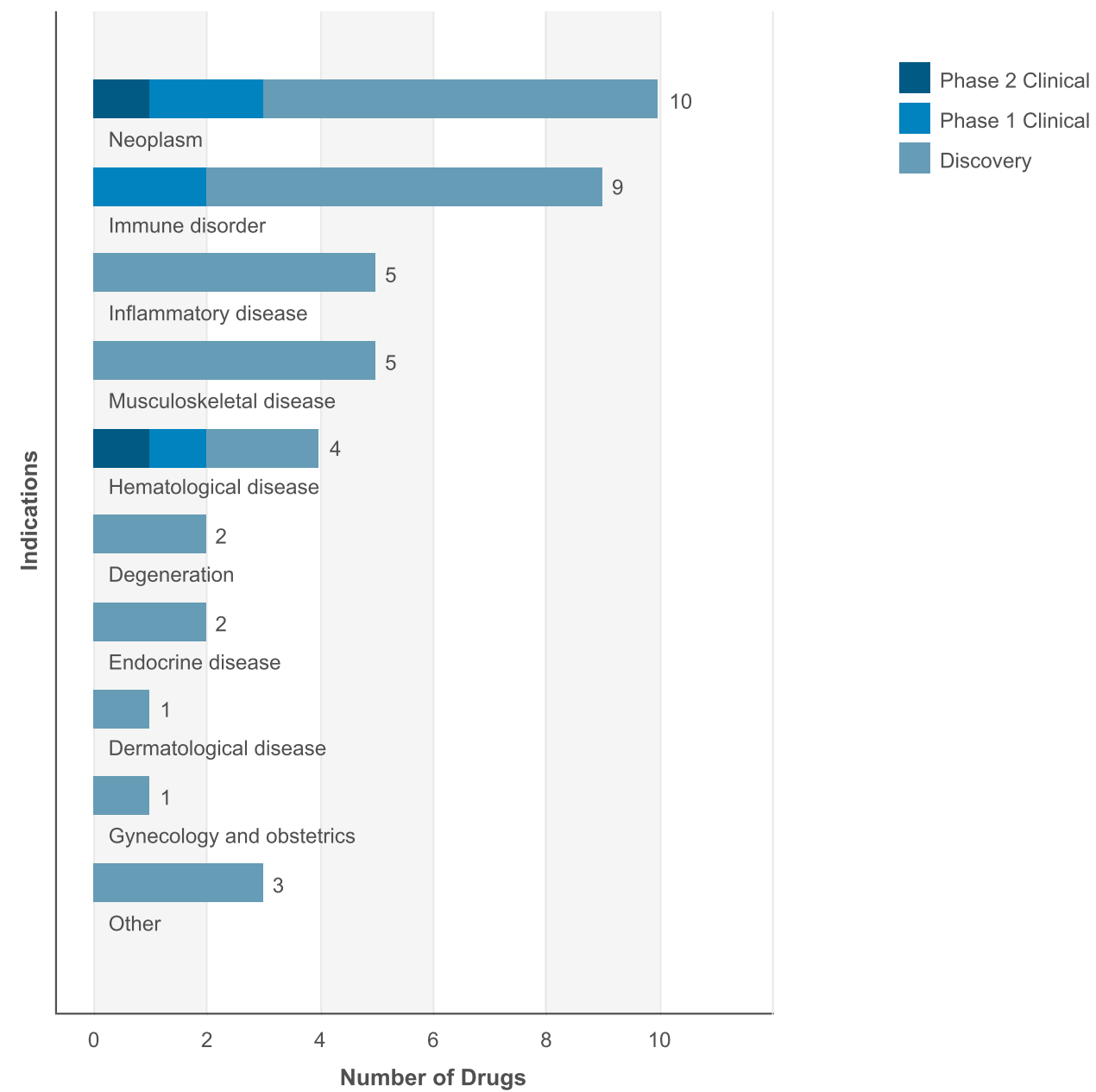


PRODUCT PORTFOLIO SUMMARY

DRUGS

Drugs by Indication

Active Drugs by Indication Chart



Drugs by Indication Table

Indication	Active	Inactive	Total
Neoplasm	10	4	14

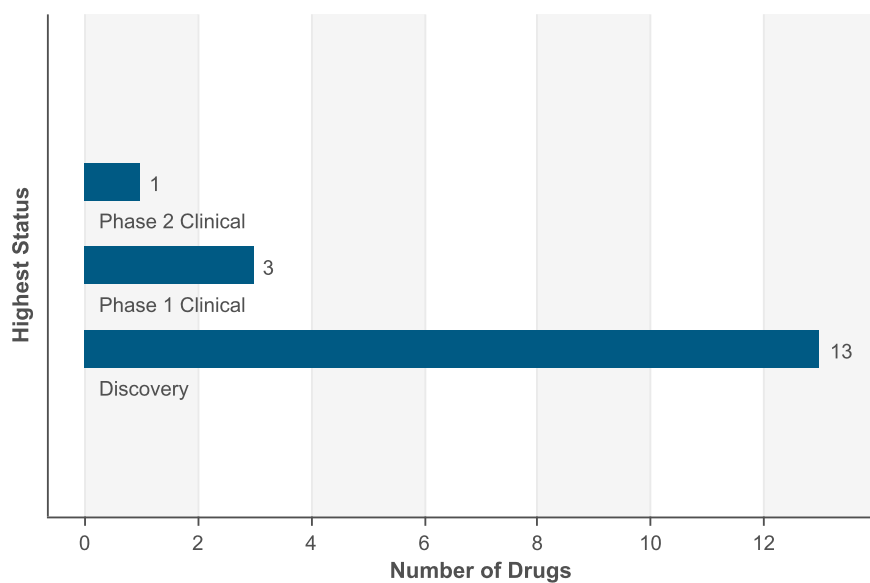
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Immune disorder	9	2	11
Hematological disease	4	2	6
Musculoskeletal disease	5	0	5
Inflammatory disease	5	0	5
Endocrine disease	2	1	3
Degeneration	2	0	2
Gynecology and obstetrics	1	1	2
Dermatological disease	1	0	1
Respiratory disease	1	0	1
Neurological disease	1	0	1
Metabolic disorder	1	0	1
Genitourinary disease	0	1	1
Ocular disease	1	0	1

### Drugs by Highest Status

Active Drugs by Highest Status Chart



Drugs by Highest Status Table

Development Status	Number of Drugs
Phase 2 Clinical	1

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Phase 1 Clinical	3
Discovery	13
Discontinued	2
No Development Reported	3

## DEALS

Deal Type	Principal		Partner		Total
	Active	Inactive	Active	Inactive	
Drug - Discovery/Design	1	0	1	0	2
Technology - Other Proprietary	9	0	1	0	10
Drug - Funding	2	0	0	0	2
Drug - Screening/Evaluation	1	0	0	0	1
Drug - Early Research/Development	1	0	0	0	1
Drug - Development/Commercialization License	3	0	0	0	3
Drug - Manufacturing/Supply	0	0	1	0	1
Technology - Delivery/Formulation	1	0	0	0	1
Technology - Target Validation	1	0	0	0	1

## CLINICAL TRIALS

### Trials by Condition Studied

Condition Studied	Ongoing	All
Immune disorder	1	3
Neoplasm	1	2
Hematological disease	1	2

### Trials by Phase

Phase	Ongoing	All
Phase 1	1	3

### 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)

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**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	38	0	38
Neoplasm	42	0	42
Ocular disease	2	0	2
Genetic disorder	2	0	2
Metabolic disorder	11	0	11
Mouth disease	4	0	4
Neurological disease	18	0	18
Degeneration	2	0	2
Andrology	1	0	1
Cardiovascular disease	10	0	10
Endocrine disease	11	0	11
Genitourinary disease	3	0	3
Dermatological disease	9	0	9
Otorhinolaryngological disease	2	0	2
Gastrointestinal disease	14	0	14
Growth disorder	2	0	2
Hematological disease	13	0	13
Musculoskeletal disease	17	0	17
Nutritional disorder	2	0	2
Respiratory disease	5	0	5
Infectious disease	26	0	26
Injury	2	0	2
Inflammatory disease	28	0	28

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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.

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## PRODUCT PORTFOLIO DRUGS

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

### MOR-208

<b>Drug Name</b>	MOR-208
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	MorphoSys AG, Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Phase 2 Clinical
<b>Active Indications</b>	Cancer, Chronic lymphocytic leukemia
<b>Target-based Actions</b>	B-lymphocyte antigen CD19 inhibitor
<b>Other Actions</b>	Anti-inflammatory, Anticancer monoclonal antibody
<b>Technologies</b>	Monoclonal antibody humanized, Intravenous formulation, Biological therapeutic
<b>Last Change Date</b>	06-Nov-2012

### XmAb-2513

<b>Drug Name</b>	XmAb-2513
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Phase 1 Clinical
<b>Active Indications</b>	Hodgkins disease, Anaplastic large cell lymphoma
<b>Target-based Actions</b>	CD30 antagonist
<b>Other Actions</b>	Anticancer monoclonal antibody
<b>Technologies</b>	Monoclonal antibody humanized, Infusion, Biological therapeutic, Parenteral formulation unspecified, Protein recombinant
<b>Last Change Date</b>	13-Jun-2012

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### XmAb-5871

<b>Drug Name</b>	XmAb-5871
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Amgen Inc, Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Phase 1 Clinical
<b>Active Indications</b>	Autoimmune disease, Rheumatoid arthritis, Systemic lupus erythematosus
<b>Target-based Actions</b>	Immunoglobulin gamma Fc receptor IIB antagonist, B-lymphocyte antigen CD19 inhibitor
<b>Other Actions</b>	Immunomodulator, Anti-inflammatory
<b>Technologies</b>	Monoclonal antibody humanized, Biological therapeutic, Parenteral formulation unspecified
<b>Last Change Date</b>	30-Nov-2012

### anticancer monoclonal antibody (1), (Xmab), Boehringer

<b>Drug Name</b>	anticancer monoclonal antibody (1), (Xmab), Boehringer
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Boehringer Ingelheim Corp, Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Phase 1 Clinical
<b>Active Indications</b>	Cancer
<b>Target-based Actions</b>	
<b>Other Actions</b>	Anticancer monoclonal antibody, Unspecified drug target
<b>Technologies</b>	Monoclonal antibody, Biological therapeutic, Parenteral formulation unspecified
<b>Last Change Date</b>	02-Oct-2012

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## XPro-1595

<b>Drug Name</b>	XPro-1595
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Inflammatory disease, Multiple sclerosis, Rheumatoid arthritis, Uveitis, Degeneration
<b>Target-based Actions</b>	TNF alpha ligand inhibitor
<b>Other Actions</b>	Anti-inflammatory
<b>Technologies</b>	Subcutaneous formulation, Biological therapeutic, Peptide
<b>Last Change Date</b>	27-Sep-2012

## DN-RANKL

<b>Drug Name</b>	DN-RANKL
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Osteoporosis
<b>Target-based Actions</b>	Osteoclast differentiation factor ligand inhibitor
<b>Other Actions</b>	Anticancer, Bone resorption inhibitor
<b>Technologies</b>	Small molecule therapeutic
<b>Last Change Date</b>	15-Jun-2012

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## DN-BAFF

<b>Drug Name</b>	DN-BAFF
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Autoimmune disease
<b>Target-based Actions</b>	TNF antagonist
<b>Other Actions</b>	Anti-inflammatory
<b>Technologies</b>	Biological therapeutic, Small molecule therapeutic, Parenteral formulation unspecified
<b>Last Change Date</b>	15-Jun-2012

## XmAb-5485

<b>Drug Name</b>	XmAb-5485
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Cancer
<b>Target-based Actions</b>	CD40 ligand inhibitor
<b>Other Actions</b>	Immunomodulator, Anticancer monoclonal antibody
<b>Technologies</b>	Monoclonal antibody humanized, Biological therapeutic, Parenteral formulation unspecified
<b>Last Change Date</b>	15-Jun-2012

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### XmAb-5592

<b>Drug Name</b>	XmAb-5592
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Multiple myeloma
<b>Target-based Actions</b>	Bone marrow stromal antigen 2 modulator
<b>Other Actions</b>	Anticancer monoclonal antibody
<b>Technologies</b>	Monoclonal antibody humanized, Biological therapeutic, Parenteral formulation unspecified
<b>Last Change Date</b>	15-Jun-2012

### cetuximab biosuperior, Xencor

<b>Drug Name</b>	cetuximab biosuperior, Xencor
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Cancer
<b>Target-based Actions</b>	Epidermal growth factor ligand inhibitor
<b>Other Actions</b>	Anticancer monoclonal antibody
<b>Technologies</b>	Chimeric monoclonal antibody, Biological therapeutic, Parenteral formulation unspecified, Protein recombinant
<b>Last Change Date</b>	28-Sep-2012

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### adalimumab biosuperior, Xencor

<b>Drug Name</b>	adalimumab biosuperior, Xencor
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Autoimmune disease
<b>Target-based Actions</b>	TNF antagonist
<b>Other Actions</b>	Immunomodulator
<b>Technologies</b>	Monoclonal antibody human, Biological therapeutic, Parenteral formulation unspecified
<b>Last Change Date</b>	14-Aug-2012

### bevacizumab biosuperior, Xencor

<b>Drug Name</b>	bevacizumab biosuperior, Xencor
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Solid tumor
<b>Target-based Actions</b>	VEGF ligand inhibitor
<b>Other Actions</b>	Anticancer monoclonal antibody
<b>Technologies</b>	Monoclonal antibody human, Biological therapeutic, Parenteral formulation unspecified, Protein recombinant
<b>Last Change Date</b>	28-Sep-2012

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### omalizumab biosuperior, Xencor

<b>Drug Name</b>	omalizumab biosuperior, Xencor
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Allergy, Asthma
<b>Target-based Actions</b>	Immunoglobulin E antagonist, Immunoglobulin gamma Fc receptor IIB modulator
<b>Other Actions</b>	Anti-inflammatory
<b>Technologies</b>	Monoclonal antibody humanized, Biological therapeutic, Parenteral formulation unspecified, Protein recombinant
<b>Last Change Date</b>	28-Sep-2012

### abatacept biosuperior, Xencor

<b>Drug Name</b>	abatacept biosuperior, Xencor
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Rheumatoid arthritis
<b>Target-based Actions</b>	CD80 antagonist, CD86 antagonist, Cytotoxic T-lymphocyte protein-4 stimulator
<b>Other Actions</b>	Anti-inflammatory, Immunosuppressant, IL-2 release inhibitor
<b>Technologies</b>	Protein conjugated, Immunoglobulin-G, Biological therapeutic, Protein recombinant
<b>Last Change Date</b>	28-Sep-2012

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### monoclonal antibody (2) (Xmab, cancer), Xencor/Boehringer Ingelheim

<b>Drug Name</b>	monoclonal antibody (2) (Xmab, cancer), Xencor/Boehringer Ingelheim
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Boehringer Ingelheim Corp, Xencor Inc
<b>Inactive Companies</b>	
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Cancer
<b>Target-based Actions</b>	
<b>Other Actions</b>	Anticancer monoclonal antibody, Unspecified drug target
<b>Technologies</b>	Monoclonal antibody, Biological therapeutic, Parenteral formulation unspecified
<b>Last Change Date</b>	02-Oct-2012

### trastuzumab biosuperior, Xencor

<b>Drug Name</b>	trastuzumab biosuperior, Xencor
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	Genentech Inc
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Breast tumor
<b>Target-based Actions</b>	ErbB2 tyrosine kinase receptor modulator
<b>Other Actions</b>	Anticancer monoclonal antibody
<b>Technologies</b>	Monoclonal antibody humanized, Biological therapeutic, Protein recombinant
<b>Last Change Date</b>	29-Sep-2012

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## rituximab biosuperior, Xencor

<b>Drug Name</b>	rituximab biosuperior, Xencor
<b>Key Synonyms</b>	
<b>Originator Company</b>	Xencor Inc
<b>Active Companies</b>	Xencor Inc
<b>Inactive Companies</b>	Genentech Inc
<b>Highest Status</b>	Discovery
<b>Active Indications</b>	Rheumatoid arthritis, Non-Hodgkin lymphoma
<b>Target-based Actions</b>	B-lymphocyte antigen CD20 modulator
<b>Other Actions</b>	Anti-inflammatory, Anticancer monoclonal antibody
<b>Technologies</b>	Chimeric monoclonal antibody, Biological therapeutic, Protein recombinant
<b>Last Change Date</b>	29-Sep-2012

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