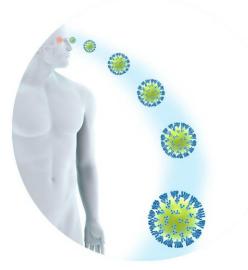


Leaders in gene and cell therapy

JEFFERIES AUTUMN 2015 GLOBAL HEALTHCARE CONFERENCE







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Oxford BioMedica at a Glance



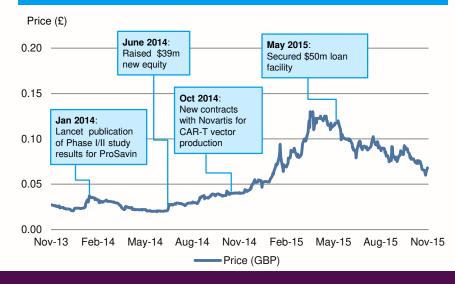
Company Facts

- Spun out of Oxford University in 1996
- IPO on LSE in April 2001 (OXB.L)
- \$260 million raised to date
- Share price 7.95p (3 Nov 2015)
- Current market cap: £204 million / \$314 million (3 Nov 2015)
- 213 employees
- Cash / Debt balance as at 30 October 2015
 - £15 million
 - \$50 million loan facility
 - Headquartered in Oxford, UK

Shareholder Register⁽¹⁾

Investor	Share
M&G Investments	17.9%
Vulpes Investment Management	17.5%
Joy Group	9.1%
Aviva	8.7%
Novartis	2.8%
Others	44.0%

Last 2-Year Share Price Performance



¹ As of 15 October 2015

Investment Highlights



Oxford BioMedica is a leading gene and cell therapy focused biotechnology company with strengths in products, development, manufacturing and IP



- Innovative diversified portfolio of unpartnered gene/cell therapy candidates
 - · 2 in ophthalmology
 - 2 in CNS
 - 2 in oncology



- Validating partnerships
 - Sanofi Licensed 2 ophthalmology gene therapy candidates (both in the clinic)
 - Novartis Manufacturing and process development relationship for CTL-019
 - GSK, Novartis, Sanofi Licence to operate under lentiviral IP



- Established, GMP-qualified, manufacturing facilities
 - · Process and product development expertise



Multiple upcoming inflection points



Experienced team with proven track record of execution

Oxford BioMedica's Business Model and Strategy



Business Model Overview

Product Development Proprietary Partnered SANOFI



OXB Solutions



Lentiviral vector manufacture and process development



IP Ownership



Lentiviral vector patents and know-how

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Revenues

Manufacturing and process development

Licence fees

Milestones

Royalties

Strategy

- Secure additional equity investment and use manufacturing revenue to fund development of product development portfolio in house rather than out-licence
- In-licence complementary product opportunities
- Complete expansion of manufacturing and laboratory capacity during 2016 and continue to develop manufacturing processes
- Sign up more "Novartis-like" process development and manufacturing contracts
- Start generating more significant profits from OXB Solutions business which should help offset Group overheads and, ultimately, to some extent, product development costs
- Patents provide licence-to-operate rights to 3rd parties
- Know-how tie in 3rd parties to long-term relationships using OXB's know how and proprietary materials

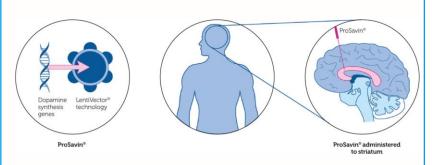
What Is Gene and Cell Therapy?



The use of DNA to treat disease by delivering therapeutic DNA into patients' cells

In vivo development

Example: OXB-102 (Parkinson's disease)



Delivery of the new gene/DNA is achieved using "viral vectors"

 Most commonly used are based on adeno-associated virus (AAV) and lentivirus

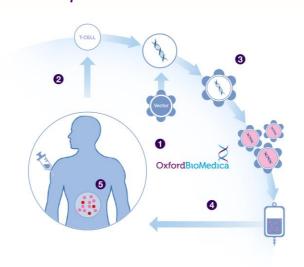
Lentiviral vector advantages over AAV

- · Larger therapeutic payloads
- Permanent modification of dividing cells such as T-cells or stem cells
- No pre-existing immunity
- OXB's lentiviral vector administered directly to over 56 patients
- Cumulative safety data greater than 150 years

Offers potential for single application treatment giving long-term or even permanent efficacy

Ex vivo development

Example: Novartis' CTL019



- OXB produces GMP lentiviral vector encoding CAR targeting CD19
- 2. White blood cells isolated from patients
- 3. Vector used to transduce expanded T-cells
- 4. The modified T-cells are infused back into the patient
- 5. Once inside the patient, the T-cells multiply, 'hunt' cancer cells and destroy them



Current Portfolio of Clinical Pipeline Products



	Field	Product	Indication	Research / Pre- Clinical	Phase I	Phase I/II	Phase II	Next inflection / Comment	Est. date	Rights
	In Vivo Programmes									
ology	CNS	OXB-102	Parkinson's Disease	Phase I/II prepa				Start Phase I/II	H1 2016	Worldwide
chno										
LentiVector® technology	OPHTHALMOLOGY	OXB-201 (RetinoStat®)	Wet AMD	Phase I concluded end point met)	(primary			Decision on next Phase	H1 2016	Worldwide
ntiVe		SAR422459	Stargardt Disease	Phase I/II ongoir	ng				2017 /	5
Le		SAR421869	Usher syndrome Type 1B	Phase I/II ongoi				End of Phase I/II	2018	SANOFI
5T4	ONCOLOGY	OXB-301 (TroVax®)	Cancer (multiple)	Phase I, Phase I/II and Phase II investigator-led studies underway			r-led	End Phase II	2015/16	Worldwide

Research/Pre-clinical Pipeline Products



	Field	Product	Indication	Research	Preclinical	Phase I/II	Phase II	Next inflection / Comment	Est. date	Rights
	Ex Vivo Programmes									
(BOIO)	ONCOLOGY	OXB-302 (CAR-T 5T4)	Cancer (multiple)					End preclinical	H2 2016	Worldwide
	OPHTHALMOLOGY		Corneal graft rejection					First patient Phase I/II	2016	Worldwide
	In Vivo Programmes									
	CNS	OXB-103	ALS or Lou Gehrig's Disease or Motor Neuron Disease					End preclinical	H1 2016	Worldwide



Novartis Contract – CTL-019/Other CAR-T Products





Overview

- Non-exclusive licence to OXB's IP
- Initial 3 year manufacturing contract (with minimum offtake commitments) for clinical supply for Novartis CTL019 programme – potential to extend
- Process development collaboration
- Financial terms include:
 - \$14m up front, including an equity investment and IP licence
 - Up to \$76m over 3 year manufacturing and process development
 - Royalties on CTL019 and other CAR-T products

OXB Provides a Key Link in the CTL-019 Supply Chain

- Novartis licensed CAR-T technology from University of Pennsylvania
- Novartis to develop the CTL019 product (and other CAR-T products)
- Complex supply chain / manufacturing process:
 - OXB produces GMP lentiviral vector encoding CAR targeting CD19
 - White blood cells isolated from patients
 - Vector used to transduce expanded T-cells
 - The modified T-cells are infused back into the patient
 - Once inside the patient, the T-cells multiply, 'hunt' cancer cells and destroy them

Facilities



Novartis contract and expectation of further manufacturing and process development led to decision to expand manufacturing and laboratory facilities

Harrow House

(GMP1/GMP2/GMP3 & Fill/Finish)
Owned API manufacturing facility
32,000 sq.ft (2,980 sq.m)
GMP2 and enhanced enabling services
under construction. Expected to be
available for production first few months
of 2016. GMP3 and Fill/Finish yet to be
commissioned



Windrush Court

Corporate Headquarters &
Laboratories (owned)
71,955 sq. ft (6,684 sq m)
Laboratory renovation expected to
complete first few months of 2016



OxfordBioMedica

2015/2016 capital expenditure on committed capacity expansion expected to be in region of £20m

Yarnton (GMP4)

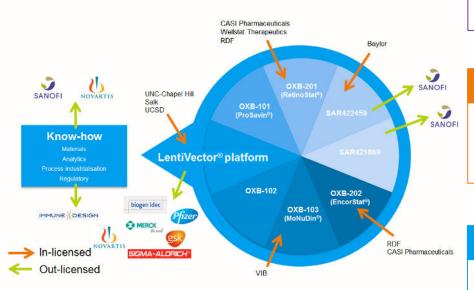
New leased API manufacturing facility 18,300 sq. ft (1,700 sq. m) Handed over by contractors October 2015, validation currently underway. Expected to be in production first few months of 2016





LentiVector® Platform IP & Key Intellectual Property OxfordBioMedical

- Multi-layered IP portfolio
- LentiVector® platform is covered by >100 patents and patent applications



Know-how

 Extensive and deep know-how relating to lentiviral vector manufacturing processes, cell and vector engineering, and proprietary analytics

Product Portfolio Protection

- Data exclusivity
- Market exclusivity relating to orphan products

Patent Portfolio

 Extensive patent estate with out-licences with several major pharmaceutical companies



Near-term Catalysts



Key Upcoming Milestones

Ongoing

• Further IP licences / manufacturing / process development contracts

H1 2016

• H1: First Patient In ("FPI") OXB-102 clinical study

OXB-201 development pathway decision made

Results from OXB-301 Phase II mesothelioma studies

H2 2016

• H2: FPI OXB-202 clinical study

OXB-302 pre-clinical results

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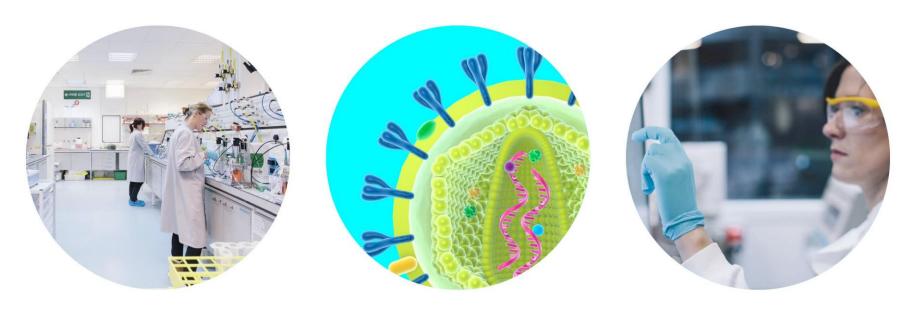
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