

The LentiVector® Platform Company
A leader in gene and cell therapy

Interim results for six months ended 30 June 2018





Forward-looking statements

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Operational highlights

Novartis' commercialised product Kymriah®

 Novartis collaboration progressing well, with Kymriah approved in the US to treat adults with r/r DLBCL and in the EU for children and young adults with r/r ALL and r/r DLBCL

Leading LentiVector® delivery platform for gene and cell therapy partnerships

- \$105 million collaboration and licence agreement with Bioverativ (now part of the Sanofi Group)
- Partnership formed with the UK Cystic Fibrosis Gene Therapy Consortium and Boehringer Ingelheim to develop an inhaled gene therapy treatment

Progress with proprietary product development

- \$842.5 million exclusive worldwide agreement signed with Axovant Sciences for OXB-102 (now known as AXO-Lenti-PD) for Parkinson's disease
 - Phase I/II clinical study for OXB-102 will start before the end of 2018
- Group continues to allocate appropriate value enhancing investment in its proprietary programmes
- Discussions are ongoing for further out-licencing deals or spin out structures for our proprietary programmes

Financial highlights

Revenue

- Gross income growth of 118% to £36.0 million (H1 2017: £15.7 million)
- Licence income of £18.3 million (Axovant and Bioverativ), segmented by Product (£10.2 million) and Platform (£8.1 million)

Profits

Operating profit of £9.4 million compared to an operating loss of £2.2 million in H1 2017

Cash

- Cash inflow before financing activities, of £12.2 million compared to an outflow of £2.2 million in H1 2017
- Cash at 30 June 2018 was £44.0 million (31 December 2017: £14.3 million), reflecting significantly improved trading performance and placing to raise £20.5 million (gross)
- £3 million capital expenditure grant received from Innovate UK
- Gross proceeds of £20.5 million raised from new and existing investors through a placing to fund proposed expansion and fit-out of additional bioprocessing activities

Other

Share consolidation completed in May 2018



Novartis CAR-T partnership progress



Novartis partnership in place since 2014, with new agreement in 2017

Clinical and commercial supply of vector

Kymriah® (tisagenlecleucel)/CTL019 and second undisclosed CAR-T programme

Non-exclusive IP licence

Potential to receive in excess of \$100m over the next 3 years for supply of vector

OXB receives royalties on sales

NHS England

News release (05 Sept 2018) Simon Stevens, Chief Executive NHS England said:

"CAR-T therapy is a true game changer, and NHS cancer patients are now going to be amongst the first in the world to benefit. Today's approval is proof-positive that, in our 70th year, the NHS is leading from the front on innovative new treatments. This constructive fast-track negotiation also shows how responsible and flexible life sciences companies can succeed - in partnership with the NHS - to make revolutionary treatments available to patients."

Current status and expectations

- In Sept 2018, NHS England approved funding for Kymriah for r/r ALL indication
- In Sept 2018, Kymriah received approval for r/r ALL and r/r DLBCL indications by Health Canada
- Sales estimate \$1.4bn¹ in 2023

		US 30 Aug 2017	Europe 27 Aug 2018	
r/r ALL (paedi	atric)	*APPROVED	*APPROVED	
r/r ALL (paedi	dult)	• APPROVED	• APPROVED	
		US 01 May 2018	Europe 27 Aug2018	

Developing the LentiVector® platform

Expansion of capacity to meet long-term demand (March 2018)

- £20.5 million (gross) equity fundraise in March 2018 to fund the expansion
- Group leased a new facility in Oxford
 - Four clean room suites and two fill and finish suites, as well as offices, warehousing and QC laboratories
- Allow the Group to exploit the immediate market opportunity and meet the expected long-term demand

Innovate UK collaboration (January 2018)

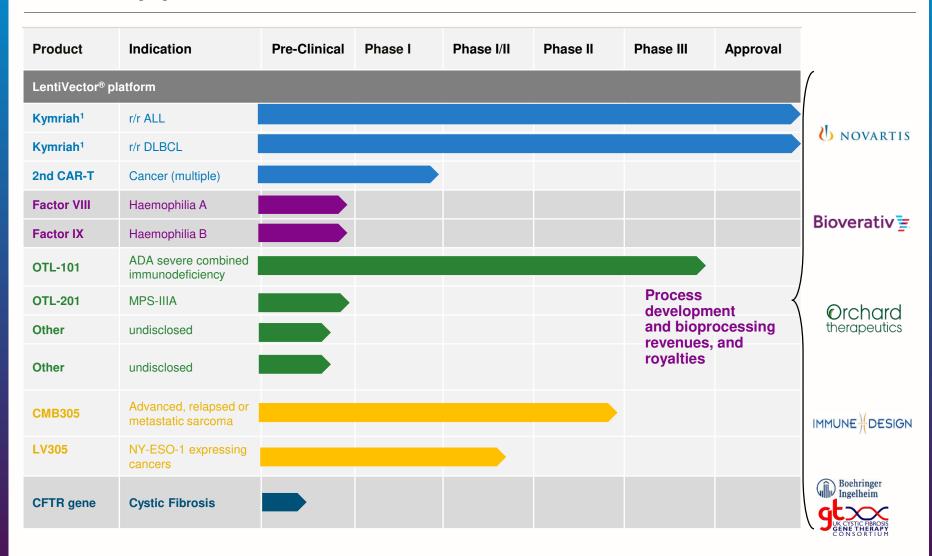
£3 million capital grant to support UK efforts to produce viral vectors

Further partnerships

- \$105 million Bioverativ collaboration & licence agreement (February 2018)
- Process development collaboration agreement with the UK Cystic Fibrosis Gene Therapy Consortium/Boehringer Ingelheim and Imperial Innovations (August 2018)

Platform pipeline

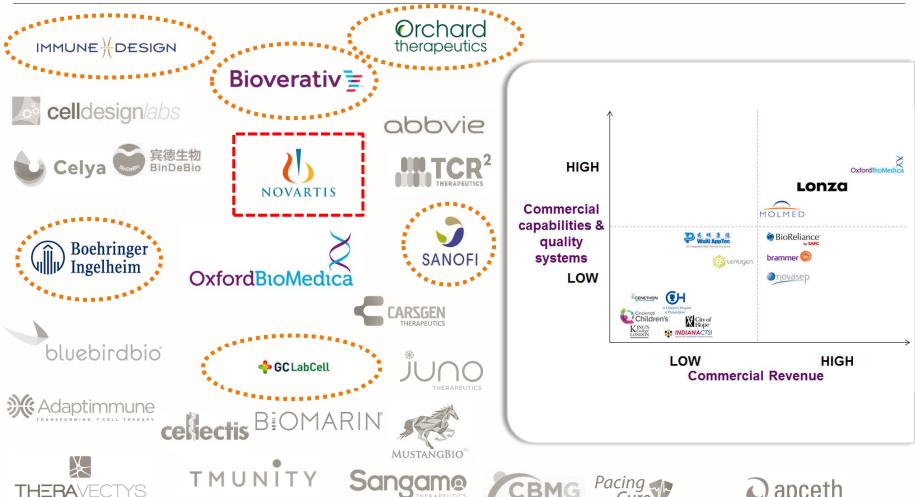




_____ In vivo programmes

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Extensive lentiviral vector clinical/pre-clinical trial activity





















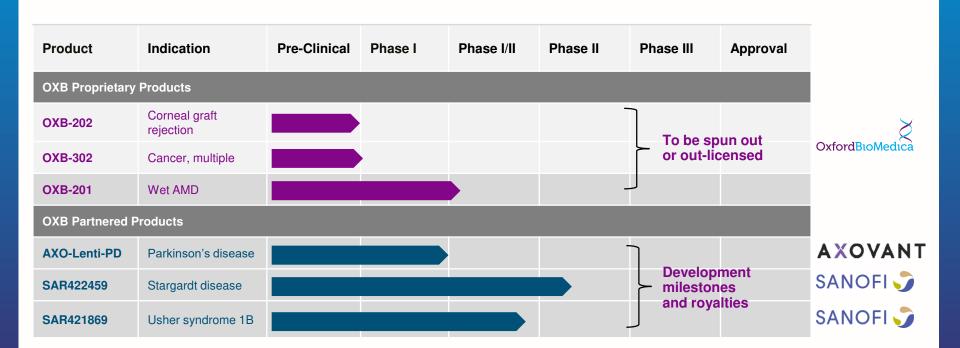


Progress with proprietary product development

- During the period the group continued to prepare the priority programmes for clinical studies
- \$842 million exclusive worldwide agreement with Axovant sciences for OXB-102 (now known as AXO-Lenti-PD) for the treatment of Parkinson's disease
 - Successfully executes on Oxford BioMedica's stated strategy to externalise product development beyond the end of the pre-clinical phase
 - Necessary regulatory filings for the planned Phase I/II study completed
 - Phase I/II clinical study for as AXO-Lenti-PD to start in 2018
- Group continues to allocate appropriate value enhancing investment in its proprietary programmes and discussions are ongoing for further out-licensing or spin out deals for these programmes

Product pipeline

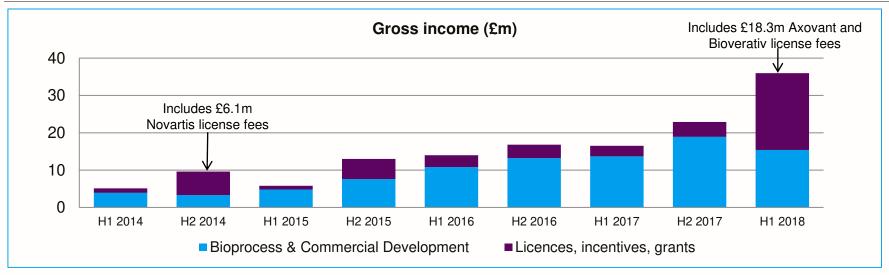


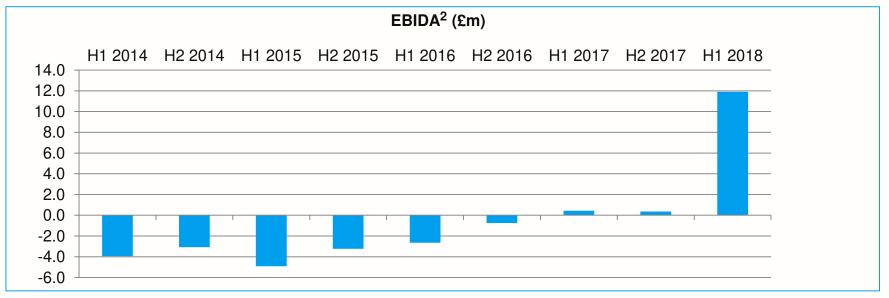


In vivo programmes



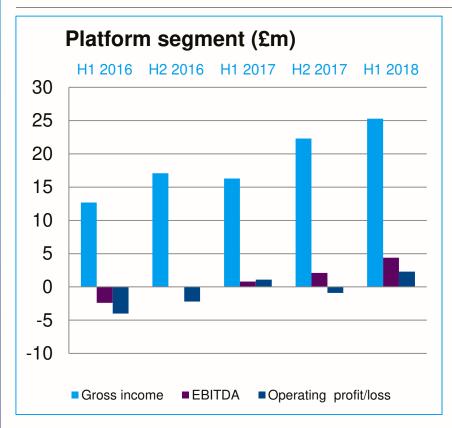
Gross income¹ and EBIDA²

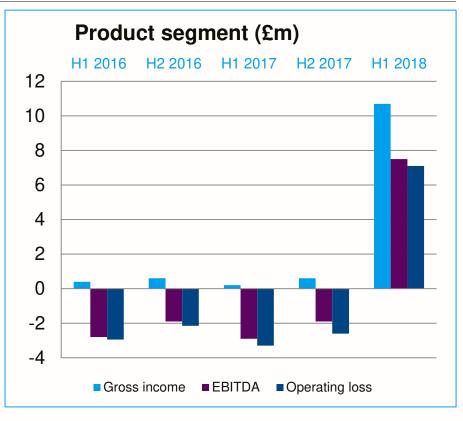




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Segmental analysis





Platform segment

- Includes Gross income received from commercial partnerships and costs of investing in LentiVector[®] technology
- Now generating cash
- Infrastructure strategy in place to support continued growth

Product segment

- Covers product development (discovery, pre-clinical and preparation for clinical studies)
- Costs include employees and directly related internal costs, external project expenditure, and allocation of Group overheads

Expected news flow

Partners' programmes

Novartis

2nd CAR-T programme to enter clinic

Royalty stream from Novartis/CTL019¹ increasing in 2018

Regulatory submissions for paediatric r/r ALL and adult r/r DLBCL in Japan and Australia expected in 2018

Orchard Therapeutics

Intends to file a BLA for ADA-SCID during H2 2018

Bioverativ

Bioverativ gene therapy product for haemophilia A & B progressing towards clinical development material by end of 2018

LentiVector® delivery platform

Further contracts with new and existing partners giving us long-term economic interest in partners' product candidates expected in the next 12 months

Invest in further development of platform to improve the volume and yield from bioprocessing and efficacy of vector transduction of target cells during 2018

In-house products

Advancement of AXO-Lenti-PD (formerly OXB-102) for Parkinson's disease into clinical development by Axovant - expected before year end 2018

Spin out / out-license of at least one in-house product candidate expected in H2 2018

Outlook

- Demand for process development and manufacture of lentiviral vectors remains very strong - further contracts with new partner companies may be concluded over the next twelve months
- These will help the Group maintain sustainable cash generation
- Continue to develop our proprietary products and pre-clinical pipeline whilst seeking to spin-out or out-license appropriate candidates
- Continue to invest in our LentiVector[™] technology platform
- Continue to monitor and prudently manage our cost base required to ensure our future sustainability and enable growth

A world leading gene and cell therapy company



Growing Market



Oxford BioMedica is at the centre of the gene and cell therapy revolution – a growing multi-billion \$ market¹.

Three therapies approved in six months and seven more expected over the next few years.

01

Profitable Platform



02

Developing Products



Developing a pipeline of our own assets in key therapeutic areas - to be either spun out or out-licensed.

Products and patents licensed to Sanofi and Axovant.

03

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