Paul Perreault CEO & Managing Director



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CSL is a global specialty biotherapeutics company that develops and delivers innovative biotherapies that save lives, and help people with life-threatening medical conditions live full lives



Global #1 in plasma therapies, #2 in influenza vaccines

CSL Behring Recent Highlights

- FY16 product sales up 10% @ constant currency
- IDELVION[®] (rFIX-FP) approved by US FDA, EMA, Japan PMDA and Health Canada
- AFSTYLA® (rFVIII-SC) approved by US FDA and EMA
- Respreeza[®] (AATD) approved by the EMA
- Filed CIDP indication for Privigen[®] in US
- Hizentra[®] CIDP trial on track
- Haegarda[™] filed in US
- New Privigen[®] (IVIG) manufacturing facility approved by US FDA



Group Sales



Broad portfolio of products and geographic reach

Commitment to R&D

Research and Development Investment (US\$ millions)



- New Product Development activities focus on innovative new therapies for life-threatening diseases.
- Market Development strategies seek to bring therapies to new markets and new indications.
- Life Cycle Management ensures continuous improvement of existing products.

R&D investment growing to ~10-11% of sales

• Accelerate CSL's exciting pipeline of assets



Immunoglobulins Global Market

- Global market volume growth projected at 5-7% in 2017
- Demand driven by medical need
- Medical education and brand promotion drive PRIVIGEN[®] and HIZENTRA[®] growth
- Growing patient acceptance of subcutaneous delivery in developed and emerging markets
- Evidence-based opportunities for future indications

Sources: Company 3Q 2016 reports, Markets and Markets Plasma Fractionation Report 2016, based on 2015 data, CSL Actuals FY16



Global IG Franchise: Strategic Objectives

GROW our Current Franchise by: Maximising current indications globally: continue geographic expansion; accelerate subcutaneous growth; launch 5 & 10 ml PFS in 2017

BUILD a Leading Neuro Franchise by: Focusing on CIDP: PRIVIGEN[®] today, HIZENTRA[®] in the near term; new neurology indications such as myositis in the future

EXPAND the Global Franchise by: Continue to invest in a broad range of potential new indications, product innovations and disruptive technologies

Category Leadership



- Orphan/rare diseases
- Unmet medical need
- Often under or misdiagnosed
- Awareness and education
- Significant patient value



Sources: Company annual reports/financial schedules, based on 3Q 2016 data, MRB WW Plasma Fractionation Market 2016 interim report, CSL Actuals FY16

HAE Franchise

Revenue Potential of \$0.75M - \$1B p.a.

HAEGARDA™

BERINERT[®]

<u>Most effective</u> in preventing HAE attacks



<u>Most effective</u> in stopping HAE attacks



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HAEGARDA™

Subcutaneous Dosing Maintains Trough above Protective C1-INH Level

- SC trough remains above predictive 40% threshold
- Reduced attack rate



Haemophilia Global Market

- Recombinants growing in developed markets
- 75% of patients with bleeding disorders are under/un-treated
- Launches of multiple longer-acting products in Hem-A space
- Payers contemplating active category management
- Rapid transition of Hem-B category



Sources: Company 3Q 2016 reports/financial schedules, based on 2016 data, MRB global Coagulation Factors Concentrate Market 2015 & 2016, Hemophilia World, December 2013, Vol 20. No 3, CSL Actuals FY16

Recombinant Coagulation Launches



- Portfolio of preclinical and early-mid stage clinical opportunities consistent with CSL commercial objectives
- Delivery of high quality candidates for clinical development



*Partnered with Janssen Biotech

Unmet Medical Need:

- Approximately 20% of patients that survive a heart attack will experience a recurrent CV event within one year
- About half of these will occur in the first month post index event

Potential Clinical Benefit:

Significant reduction in early, recurrent CV events (CV death, Recurrent MI, stroke) in high-risk ACS patients

MOA:

Rapidly removes cholesterol from atherosclerotic lesions/plaque via significantly enhanced cholesterol efflux

Source: WHO 2013 Update; CDC Heart Disease Fact Sheet August 2014

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Uncontested sub-acute market space



Sources:
1. Figure adapted from Wallentin L, et al. N Engl J Med. 2009;361:1045-1057
2. Figure adapted from Jernberg T, et al. Eur Heart J. 2015;36:1163-1170

Seqirus

No. 2 global influenza vaccine manufacturer

Operations

- Enhanced efficiency, sharing of best practices
- New formulation facility in Liverpool

Product launches

- Recent US launch of Flucelvax Quadrivalent[™] and Fluad[™]
 - Full year impact from FY18
- Afluria Quad[™] approved by Australian TGA
- Seqirus influenza vaccines 1st to market in the US

Clinical trials

- Fluad[™]: quadrivalent in elderly and paediatric
- Afluria Quad™: paediatric

Business turnaround on track



Collaboration and License Agreement with Momenta for recombinant Fc mimetic molecule

- Neurological indications treated by Ig are mediated by the Fc portion of Ig molecule
- M230 is a trimeric Fc construct and a selective immunomodulator of Fc receptors
- Plan to start Phase 1 clinical trial this year
- Research collaboration for additional Fc multimer proteins

Investment in the newly established AU\$230M Biomedical Translation Fund

Largest life sciences fund in Australia, managed by Brandon Capital Partners

Launch of AU\$25 million CSL Centenary Fellowships

CSL Strategy for Profitable Growth

Future Growth Pipeline		•
Growth Drivers	Seqirus	•
	CSL Behring	•
Core Plasma		•

- CSL 112 new treatment paradigm in ACS
- Hizentra[®] expansion into neurology
- CSL 830 (Haegarda) HAE
- Pipeline antibodies
- Targeted business development
- Drive Seqirus business to profitability
- Successfully launch pipeline vaccines
- Launch and grow Idelvion[®] & Afstyla[®]
- Maintain leadership in Ig and albumin
- Grow high-margin specialty products
- · Relentless commitment to lowest cost base
- Remain ahead of the demand curve
- Organic growth of core plasma products

Sustained Financial Performance

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