T +613 9389 1911 F +613 9389 1434 www.csl.com.au



### **ASX Announcement**

#### For immediate release

29 May 2020

### Webinar Presentation – Australian Shareholders' Association

Please find attached a presentation used for a lunchtime Webinar presentation with the Australian Shareholder's Association today.

This presentation is provided to the ASX for the information of all CSL's shareholders and contains no new or price sensitive information.

Authorised by **Fiona Mead**Company Secretary

#### For further information, please contact:

#### Media:

Christina Hickie Senior Manager Communications Telephone: +61 429 609 762

Email: Christina.Hickie@csl.com.au

#### Investors:

Bernard Ronchi Senior Manager Investor Relations Telephone: +61 3 9389 3470

Email: Bernard.Ronchi@csl.com.au

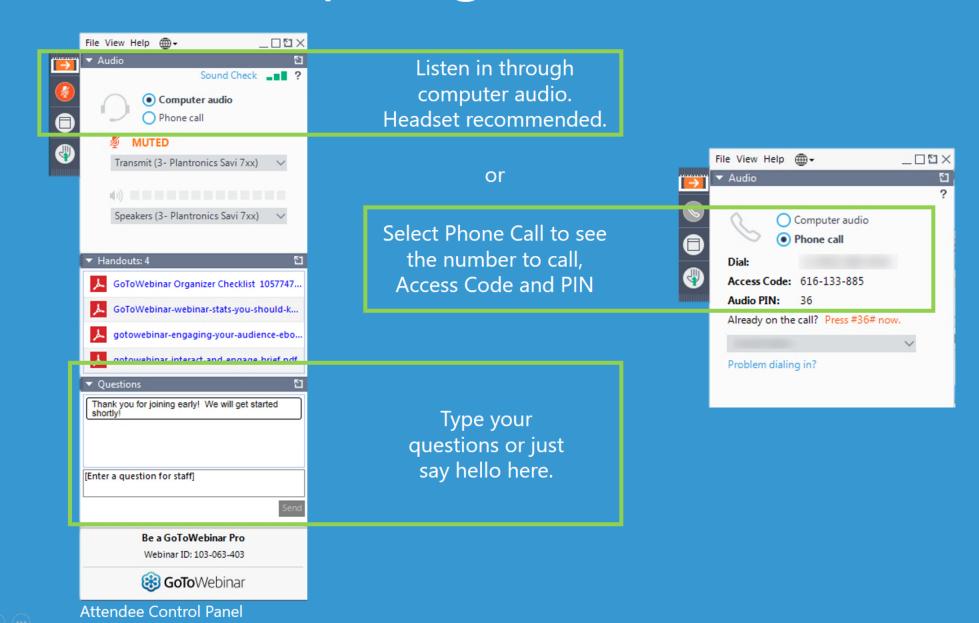


CSL Limited
Update on operations and future growth

David Lamont, CFO

ASA Webinar, 29 May 2020

## Participating in the Webinar



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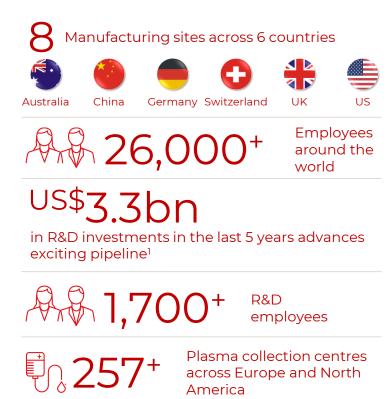




### **Overview**

CSL is a leading global biotech company that develops and delivers innovative biotherapies and influenza vaccines that save lives and help people with life-threatening medical conditions live full lives







### **CSL Today**

### **3rd Largest Global Biotech**



#### Global #1 in plasma therapies

\$30 billion industry



#### Global #2 in influenza vaccines

\$6 billion industry



#### **Strong Market Position**

- Revenue FY19 ~\$8.5bn
- 8 major manufacturing sites in 6 countries
- Major capacity expansion underway
- Deep R&D pipeline fueling future growth



#### **Solid Financial Position**

- Net debt/EBITDA 1.4x
- A3 / A- credit rating (stable / stable)

# **Current Industry Themes**

Plasma Supply Growth

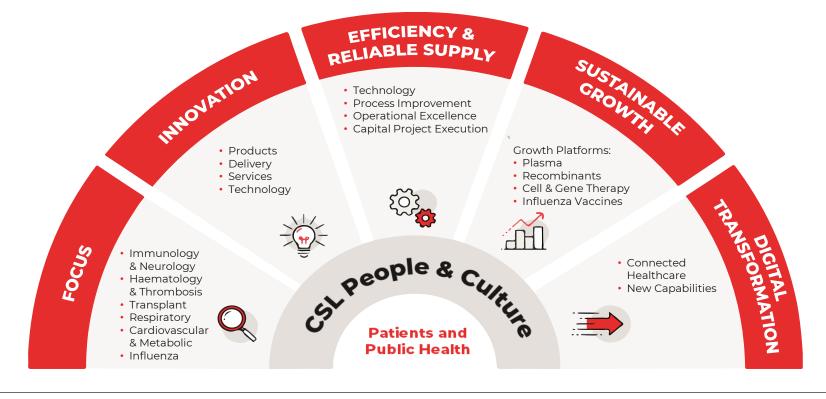
Robust Ig demand

Influenza Vaccine Technology Shift



### **Values and Strategy**

Strategic Overview



Core Values



Patient Focus

We deliver on our promise to patients



Innovation

We turn innovative thinking into solutions



Integrity

We walk the talk



Collaboration

We are stronger together

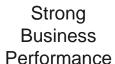


We take pride in our results



### **Business Highlights**







Balanced Regional Growth



Executing to Plan on New Launches



Ig Growth well Above Market



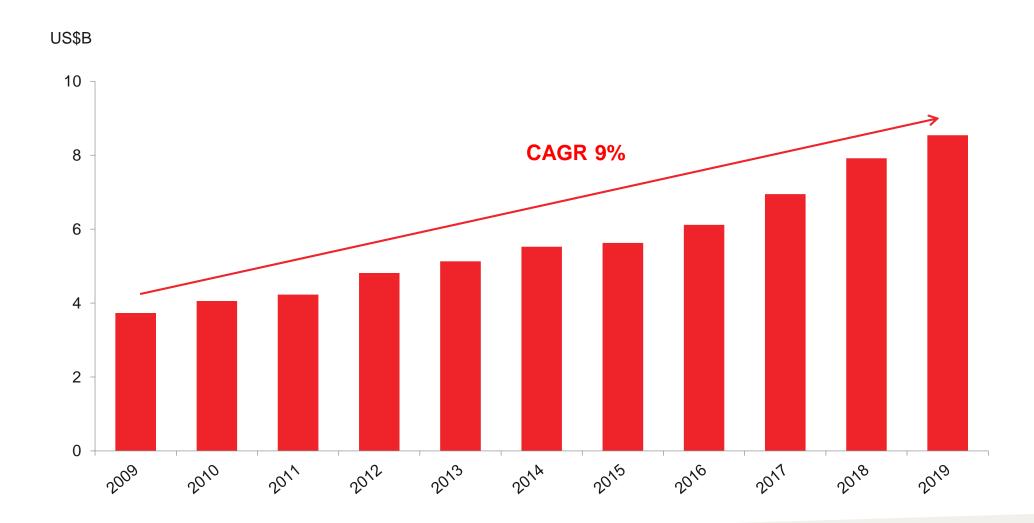
Expanding
Market
Presence
through New
Affiliates



Compelling real-world effectiveness influenza vaccine data



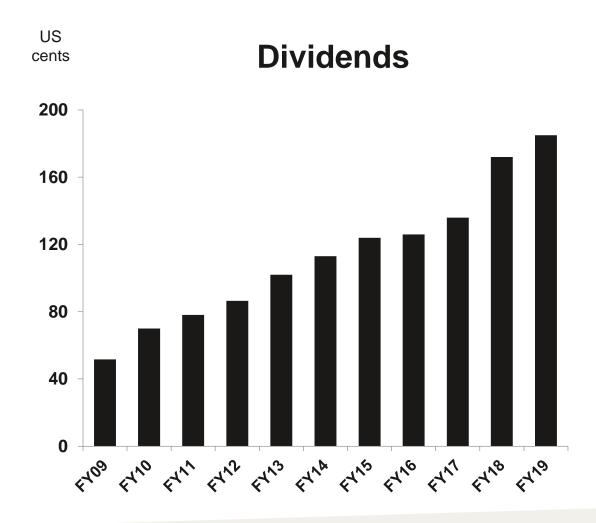
### **Strong and Consistent Revenue Growth**





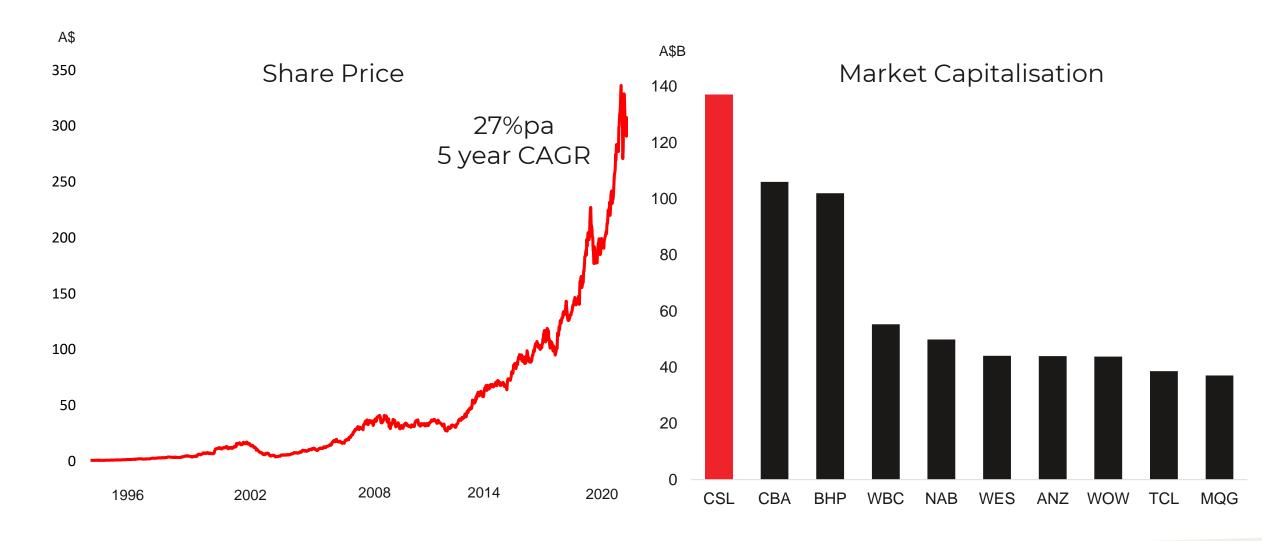
### **Shareholder Returns**





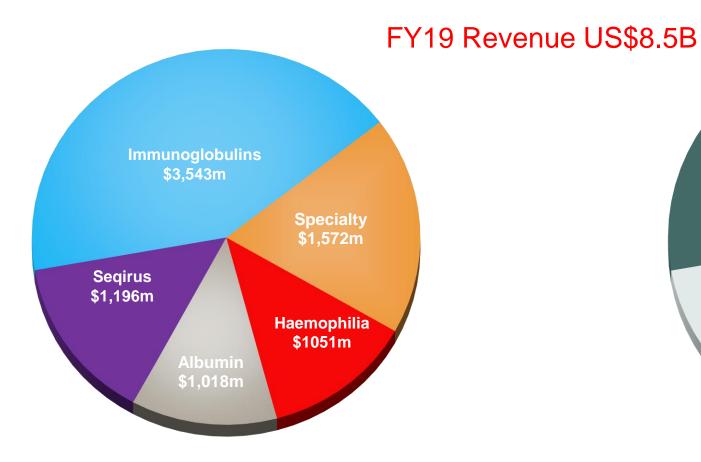


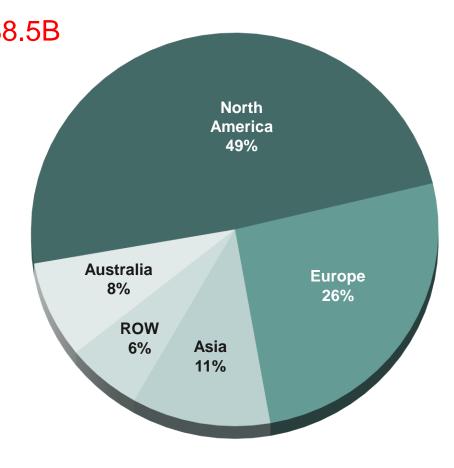
### **Shareholder Value**





### **Global & Diverse Revenue Base**



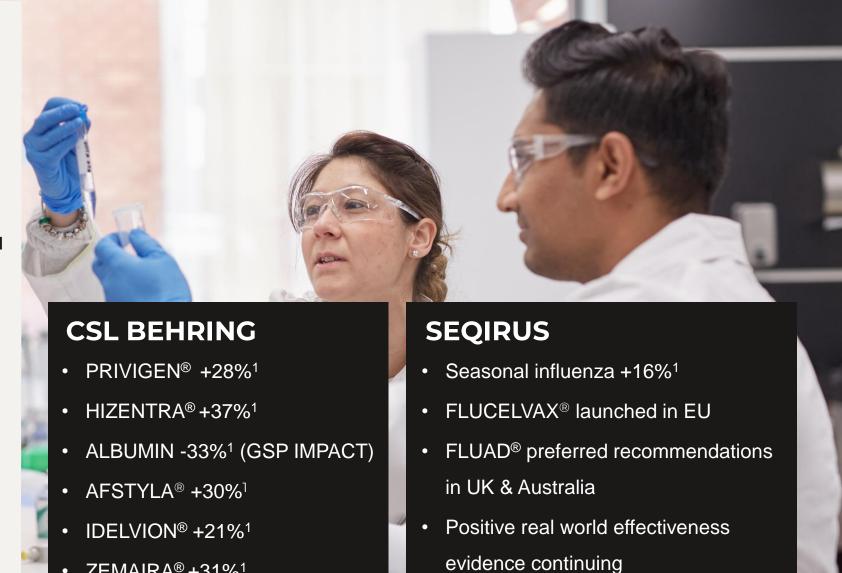




### 1H20 **Performance**

A strong half year for CSL with revenue up 11%<sup>1</sup> and net profit after tax up 11%1 reflecting:

- Strong growth in immunoglobulin portfolio
- Transition to own distribution model in China progressing well
- Continued evolution of the haemophilia portfolio
- Strong performance from Segirus



Commenced aQIVc development

ZEMAIRA® +31%<sup>1</sup>

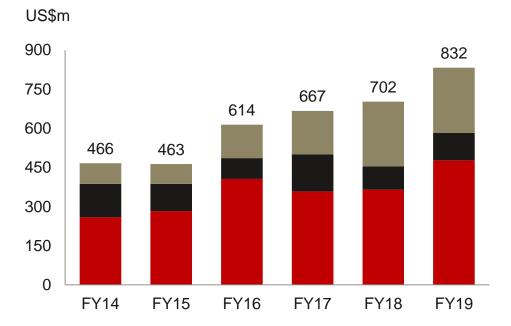
### **1H 20 Financial Highlights**

Half year ended Dec US\$ Millions	1H19 Reported	1H20 Reported	1H20 at CC <sup>1</sup>	Change %
Total Revenue	4,505	4,911	4,980	11% <sup>1</sup>
Gross Profit  GP margin	2,623 58.2%	2,842 57.9%	2,891 58.0%	10% <sup>1</sup>
EBIT	1,553	1,632	1,679	8%1
NPAT	1,161	1,248	1,292	11% <sup>1</sup>
Cashflow from Operations	535	1,245		133%
Capex	539	685		27%
EPS (\$)	2.56	2.75	2.85	11% <sup>1</sup>
DPS (\$)	0.85	0.95 (A\$1.47)		12% 22%



### **Commitment to Research and Development**

### R&D Spend



- 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

### New R&D Facilities







#### **Bio21 Institute, Melbourne**

- ~ 4100m² of lab and office space
- Parkville precinct
- Melbourne University, MRI's
- · 4 major teaching hospitals

#### SITEM<sup>1</sup>, Bern

- 2000m<sup>2</sup> of lab and office space
- Bern University and Hospital campus

#### **Gene therapy, Pasadena**

- Expanding gene therapy expertise
  - Research, QA, cell processing and manufacture
  - Wet-lab space (non-GMP) tripled from 132 to 480 m<sup>2</sup>
  - GMP space (330 m²) to engineering qualification level



### **R&D Highlights**



Immunology and Neurology

- HIZENTRA® granted Orphan Drug Exclusivity for CIDP
- HIZENTRA® Dermatomyositis (DM) Phase III Study initiated
- Garadacimab (Anti-FXIIa) in Hereditary Angioedema (HAE) Phase II double blind period complete



Haematology and Thrombosis

- CSL200 (CAL-H) in Sickle Cell Disease (SCD) Phase I Study initiated
- CSL889 Hemopexin in SCD Phase I Study initiated



Respiratory

- CSL311 (Anti-Beta Common) Phase I study commenced
- Approval of convenient single-vial dosing for ZEMAIRA® (Alpha1-Proteinase Inhibitor) in the US



Cardiovascular and Metabolic

• CSL112 (ApoA-1) Phase III study (AEGIS-II) progressing well with >7000 patients recruited



**Transplant** 

• CSL964 Alpha-1 Antitrypsin (AAT) for prevention of Graft versus Host Disease (GvHD) after Transplantation of Allogenic Hematopoietic Cell Transplantation (HCT) Phase III study actively recruiting and on track



Influenza Vaccines

- First cell-based quadrivalent seasonal influenza vaccine, FLUCELVAX® TETRA, approved in Europe
- AFLURIA® QUAD (quadrivalent influenza vaccine) granted expanded indication for use in children 6M+ in Australia
- aQIVc (MF59 plus FLUCELVAX® antigen) new product development commenced



### **COVID-19 Update**

### Response

- Primary objective is to ensure safety and wellbeing of our staff, donors and continuity of supply of medicines to patients
- Business continuity plans implemented at all sites
- Expertise, technologies, equipment and materials are being provided to support the response to COVID-19
- Providing additional southern hemisphere influenza vaccine doses to Australia

### **Employees**

- Health and welfare of staff are of primary importance to CSL
- Wherever possible, staff are strongly encouraged to work remotely
- Facilities are 'critical infrastructure/essential facilities' and excluded from most government shutdown processes
- On-boarding of critical staff positions accelerated
- International and domestic travel limited to business critical







**Evolving Pandemic** 

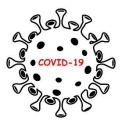




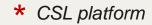




### Focused COVID-19 Response



Prevention		Treatment		
		S e v e	rity	
Population	General public	At-risk healthcare workers and infected patients		
Therapeutic Options	* Vaccines	<ul><li>* Hyperimmunes</li><li>Polyclonal antibodies</li></ul>	<ul><li>Monoclonal antibodies</li></ul>	
CSL Assets & Capabilities	Adjuvant technology Scaled manufacturing	Process development Scaled manufacturing	CSL312; CSL324 IL6; C1inhibitor	
Collaborators	University of Queensland/CEPI	Hyperimmune: Alliance with Takeda and others	Academic clinical researchers	
		Polyclonal: Sab Therapeutic	S	





### **Growth Catalysts**

#### PLASMA PROTEINS

- Ongoing robust demand
- Commercialization of 5 global product launches
- Grow China business
- R&D pipeline
  - CVD, Fc Multimer, transplant, gene therapy

#### **INFLUENZA**

- Product differentiation FLUCELVAX®
- Sales shifting towards FLUAD® and QIV

#### **EFFICIENCY & FLEXIBILITY**

- Harness benefits from new technology investments
- Significant manufacturing capacity expansion
- Opening 40 collection centres in FY20

## **Outperformance Drivers**





