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17 February 2016

Australian Securities Exchange

Correction to Half Year Results Announcement and Analyst Briefing

There is an error in the Basic EPS amount for the Half Year ending 31 December 2014 in the Group Results Table. The amount of US$1.43 should have been US$1.46. This then flows through to incorrectly stating the EPS as growing 9% (where as the correct amount is 6%) and 12% (where the correct amount is 9%) in constant currency, on the prior corresponding period.

Please note that the Basic EPS amount for the Half Year ending 31 December 2015 is correctly stated as US$1.55. In addition, the EPS amounts in the Appendix D and Half Year Accounts are correct.

A revised version of the Half Year Announcement and Analyst Briefing is attached (highlighting the changes made).

Edward Bailey
Company Secretary

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ASX Announcement

For immediate release

16 February 2016

Half Year Result 2016

CSL Delivers Exceptional Performance

- **Double-digit sales growth in all plasma therapy groups**
- **1st shipment of Privigen® from new facility**
- **Seqirus formed – No.2 global influenza vaccines manufacturer**

CSL Limited (ASX:CSL; USOTC:CSLLY) today announced a net profit after tax (NPAT) of US$719 million for the six months ended 31 December 2015, up US$27 million or 4% on a reported basis when compared to the prior comparable period (PCP). Earnings per share (EPS) grew 6%. After excluding financials relating to the recently acquired Novartis influenza vaccines business, underlying[1] NPAT grew 7% and EPS grew 9%, at constant currency[2].

HIGHLIGHTS

Financial

- Sales US$3,056 million, up 11% on PCP
  - *Underlying[1] sales up 9% at constant currency[2]*
- NPAT US$719 million, up 4% on PCP
  - *Underlying NPAT up 7% at constant currency*
- Earnings per share US$1.55, up 6% on PCP
  - *Underlying EPS up 9% at constant currency*
- Cashflow from operations US$705 million, up 8% on PCP
- Interim dividend[3] of US$0.58 per share, unfranked for Australian tax purposes, payable on 15 April 2016
  - Converted to Australian currency, the interim dividend increased to approximately A$0.81 per share, up ~10% on PCP.

[1] Underlying excludes financials relating to the Novartis influenza vaccines business (NVS-IV). NVS-IV was acquired on 31 July 2015.

[2] Constant currency removes the impact of exchange rate movements to facilitate comparability of operational performance. See end note (#) for further detail.

[3] For shareholders with an Australian registered address, dividends will be paid in A$ at an amount of A$0.814726 per share (at an exchange rate of A$1.4047/US$1.00), and for shareholders with a New Zealand registered address, dividends will be paid in NZD at an amount of NZ$0.874234 per share (at an exchange rate of NZ$1.5073/US$1.00). The exchange rates used are fixed at the date of dividend determination. All other shareholders will be paid in US$. As a result of the ASX's announced intention to move to a T+2 settlement cycle, CSL's ex-dividend date for its interim dividend will be 23 March 2016 (previously 22 March 2016)

Operational

- Product Portfolio
  - Double digit sales growth in all plasma therapy groups
  - CSL 654 (rIX-FP) – license under review in the U.S. and EU
  - CSL 627 (rVIII-SC) – license under review in the U.S.
  - Respreeza® approved in the EU
  - CSL 362 (AML) – licensee (Janssen) commenced phase 2 study
  - CSL 112 (rHDL) – phase 2b fully enrolled
- Operations
  - New Privigen® manufacturing facility in Broadmeadows, Australia approved by U.S. FDA
    - First Privigen® shipment in December 2015
  - New sales office opened in Russia
- Influenza
  - Novartis influenza vaccines acquisition closed
  - 'Seqirus' launched – No. 2 global influenza vaccine manufacturer
  - FLUAD™ approved by U.S. FDA
  - Quadrivalent influenza vaccines - licenses under regulatory review
- Capital management
  - A$1 billion share buyback[4] underway
  - ~US$500 million private placement completed
  - New US$1.25 billion bank debt facilities negotiated

"CSL delivered an exceptional first half result, led by double-digit sales growth in all of our plasma therapy groups," said CSL Chief Executive Officer and Managing Director Paul Perreault. "In particular we saw strong demand for our immunoglobulin products with subcutaneous immunoglobulin therapy, Hizentra®, growing at 31% and intravenous immunoglobulin therapy, Privigen®, up 13%."

"This year CSL will mark its centenary as a very different organization to the one that was founded in 1916 to ensure Australia had its own supply of sera, antitoxins and vaccines. Today, we are an established and growing global biotherapeutics leader, developing and delivering innovative therapies for patients around the world. Seqirus, our influenza vaccine business, is the second largest provider in the world with a diverse product portfolio, broad global sales reach and manufacturing capabilities in both

[4] CSL reserves the right to suspend or terminate buy-backs at any time.

northern and southern hemispheres. Overall, CSL is well positioned for sustainable growth and continuing to deliver value to shareholders."

OUTLOOK

Commenting on CSL's outlook, Mr. Perreault said, "2016 is an exciting year for CSL. The licenses for our novel recombinant coagulation products are currently under review, and pending approval, we plan to introduce these to the market later this year. We have been investing in our commercial capabilities to support the launch and rollout of these products. We have also continued to invest in our research and development pipeline and our manufacturing spine to ensure we meet growing demand. Notwithstanding this additional expenditure and the current competitive market, I can reconfirm my previous guidance for FY16 of 5% profit growth at constant currency."

Mr Perreault continued, "This guidance does not include financials associated with the acquisition of the Novartis influenza vaccines business, which we anticipate will report a loss in the range of approximately US$90 - 120 million this financial year. However, with the deal now closed a significant multi-year strategy has commenced to integrate this business and turn its performance around."

Earnings per share growth for the Group is expected to exceed profit growth, benefiting from ongoing capital management activity.

In compiling the company's financial forecasts for the year ending 30 June 2016 a number of key variables which may have a significant impact on guidance have been identified and these have been included in the footnote[5] below.

[5] Key variables that could cause actual results to differ materially include: the success of research and development activities, decisions by regulatory authorities regarding approval of our products as well as their decisions regarding label claims; competitive developments affecting our products; the ability to successfully market new and existing products; difficulties or delays in manufacturing; trade buying patterns and fluctuations in interest and currency exchange rates; legislation or regulations that affect product production, distribution, pricing, reimbursement, access or tax; litigation or government investigations, and CSL's ability to protect its patents and other intellectual property.

OPERATING REVIEW

CSL Behring sales of US$2.5 billion grew 10% in constant currency terms when compared to the prior comparable period.

Immunoglobulin product sales of US$1,181 million grew 13% in constant currency terms.

Intravenous immunoglobulin (IVIG) sales growth was underpinned by strong demand for Privigen® with sales growth of 13% over the prior comparable period. Privigen's® expanded indication in Europe to include its use in the treatment of chronic inflammatory demyelinating polyneuropathy (CIDP) was a significant contributor to growth in this region, especially in France and the UK. US sales into the Specialty Pharmacy segment also performed well.

Sales of subcutaneous immunoglobulin product, Hizentra®, was up 31% at constant currency, led by sales in the U.S. and Europe. New patient starts on Hizentra® and those converting from IVIG were key drivers of growth.

Albumin sales of US$376 million grew 10% in constant currency terms, driven by ongoing strong global demand. Demand in China was of particular note with growth supported by the company's ongoing successful sales penetration into Tier 2 and Tier 3 cities.

Haemophilia product sales of US$509 million increased 2% in constant currency terms. Plasma derived haemophilia sales grew 13% following successful tenders for the provision of Beriate® in European countries, including Poland and Russia. Solid Humate® growth in the U.S. was underpinned by expanded use in surgeries and immune tolerance therapy. Biostate® sales lifted in Germany, France and the U.K. A decline in sales of Helixate®, CSL's recombinant factor VIII, to a large extent offset the growth in plasma derived therapies as competition intensifies following the launch of new generation recombinant FVIII products.

Specialty products sales of US$466 million grew 14% in constant currency terms. Sales of Kcentra® (4 factor pro-thrombin complex concentrate) in the U.S. were particularly strong following an increased level of promotion and increasing brand awareness.

Following marketing authorization being granted for Respreeza® in Europe, this product was launched in Germany with plans for rollout in other European countries later this year. Respreeza® is a maintenance treatment for severe Alpha-1 Antitrypsin Deficiency patients and has been shown to slow the progression of emphysema.

Long term investment in a multi-site expansion program to meet future demand for therapies continues. In December the Board approved investment and construction of a new commercial scale manufacturing facility for recombinant coagulation factors in Lengnau, Switzerland. Also in December the first shipment of Privigen was made from a new manufacturing facility in Broadmeadows, Australia. Construction of a new albumin production facility at the same site continues. In Marburg, Germany a new quality control facility together with filling and packaging upgrades is nearing completion. At the Kankakee, U.S. site the construction of significant base fractionation plant is well progressed.

Seqirus sales of US$519 million are reported for the first time, following the combination of CSL's subsidiary bioCSL and Novartis influenza vaccines (NVS-IV) manufacturing business to form CSL's new business unit *Seqirus*. NVS-IV was acquired on 31 July 2015 and Seqirus becomes the second largest manufacturer of influenza vaccines globally. Seqirus sales of influenza vaccine have been impacted by the mild season in the northern hemisphere.

CSL Intellectual Property revenue of US$64 million declined 29% in constant currency terms. The prior comparable period included a payment from CSL's licensee Janssen Biotech Inc to develop and commercialise CSL 362, a product used to treat patients with acute myeloid leukaemia.

CAPITAL MANAGEMENT

Share Buyback

In October 2015, CSL announced its intention to conduct an on-market share buyback of up to A$1 billion. Under the Australian Securities Exchange listing rules this buyback[6] has a 12 month completion window. To date, CSL has repurchased approximately 2.4 million shares for approximately A$235 million, representing about 24% of the intended repurchase program.

[6] CSL reserves the right to suspend or terminate buybacks at any time.

CSL's balance sheet remains very sound and only modestly geared. Cash and cash equivalents totalled US$1,092 million at 31 December 2015.

During the first half of fiscal 2016 the company accessed the private placement market and raised the equivalent of approximately US$500 million as part of the company's overall debt management program. CSL also re-negotiated its major bank facilities, totalling US$1.25 billion with a maturity of 5 years.

CHANGES TO CSL BOARD

Dr Megan Clark AC has been appointed a Director of the Company effective from 16 February 2016. For further information please see separate ASX announcement.

Additional details about CSL's results are included in the company's 4D statement, investor presentation slides and webcast, all of which can be found on the company's website www.csl.com.au A glossary of medical terms can also be found on the website.

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Group Results

US Dollars

Six months ended December US$ Millions	**Dec 2014** Reported	**Dec 2015** Reported	**Dec 2015** NVS-IV[7]	**Dec 2015** Underlying[8]	**Dec 2015** Underlying[8] at CC[#]	**Change** %
Sales	**2,744**	**3,056**	**294**	**2,762**	**2,996**	**9.2%**
Other Revenue / Income	96	80	4	76	79	
Total Revenue / Income	**2,841**	**3,136**	**298**	**2,838**	**3,075**	**8.2%**
Earnings before Interest, Tax, Depreciation & Amortisation	**969**	**848**	**(112)**	**960**	**1,053**	**8.7%**
Depreciation/Amortisation	91	102	9	93	102	
Earnings before Interest and Tax	**878**	**746**	**(121)**	**867**	**952**	**8.3%**
Gain on Acquisition		176	176			
Net Interest Expense / (Income)	21	27	1	26	26	
Tax Expense	165	176	5	171	188	
Net Profit after Tax	**692**	**719**	**50**	**669**	**738**	**6.6%**
Interim Dividend (US$)	0.58	0.58				
Basic EPS (US$)	1.46	1.55			1.59	9%

[7] Novartis influenza vaccines acquisition as from 31 July 2015

[8] Underlying excludes financials relating to the Novartis influenza vaccines business (NVS-IV)

(#) **Constant currency** removes the impact of exchange rate movements to facilitate comparability of operational performance. This is done in three parts: (a) by converting the current period net profit of entities in the group that have reporting currencies other than US Dollars at the rates that were applicable to the prior comparable period ("translation currency effect"); (b) by restating material transactions booked by the group that are impacted by exchange rate movements at the rate that would have applied to the transaction if it had occurred in the prior comparable period ("transaction currency effect"); and (c) adjusting for current year foreign currency gains and losses. The sum of translation currency effect, transaction currency effect and foreign currency gains and losses is the amount by which reported result is adjusted to calculate the operational result.

Summary NPAT

Reported Net Profit after Tax	$718.8m
Translation Currency Effect (a)	$ 64.8m
Transaction Currency Effect (b)	$ (9.8m)
Foreign Currency Gains and Losses (c)	$ 13.7m
Constant Currency Net Profit after Tax *	$787.5m

(a) Translation Currency Effect $64.8m

Average Exchange rates used for calculation in major currencies (six months to Dec 15/Dec 14) were as follows: USD/EUR (0.91/0.77); USD/CHF(0.97/0.93)

(b) Transaction Currency Effect ($9.8m)

Transaction currency effect is calculated by reference to the applicable prior comparable period exchange rates. The calculation takes into account the timing of sales both internally within the CSL Group (ie from a manufacturer to a distributor) and externally (ie to the final customer) and the relevant exchange rates applicable to each transaction.

(c) Foreign Currency Losses (13.7m)

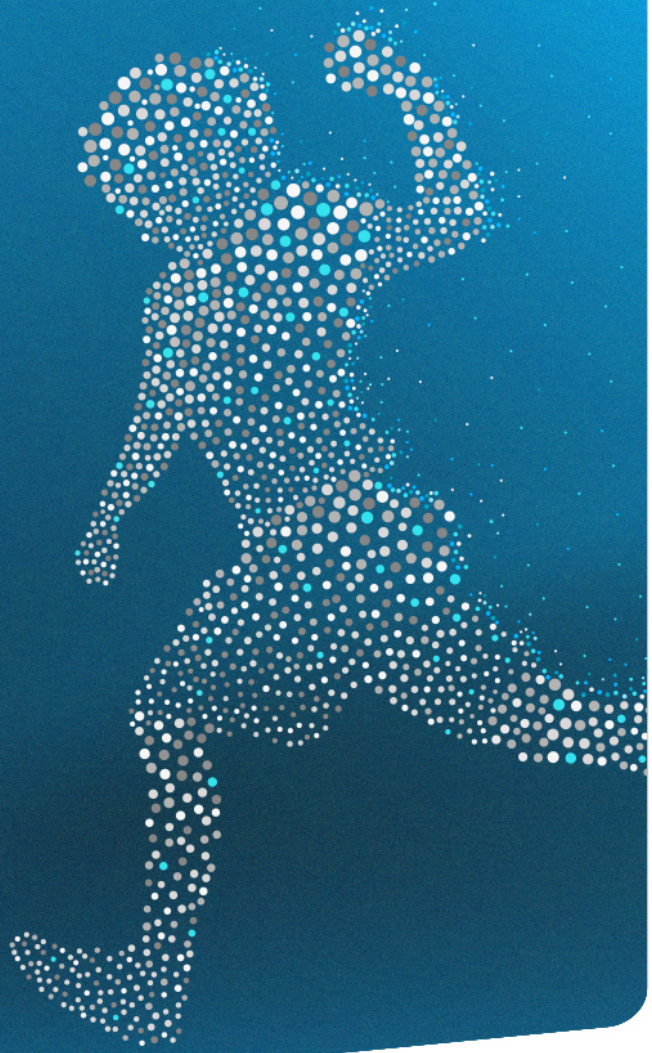
Foreign currency losses recorded during the period

Summary Sales

Reported Sales	$3,056.3m
Currency Effect	$ 234.0m
Constant Currency Sales *	$3,290.3m
Less NVS-IV sales	$ 294.6m
Underlying operational business sales @ CC	$2,995.7m

* Constant Currency Net Profit after Tax and Sales have not been audited or reviewed in accordance with Australian Auditing Standards.

CSL Limited
2016 Half Year Result
16 February 2016



CSL™

Legal Notice

Forward looking statements

The materials in this presentation speak only as of the date of these materials, and include forward looking statements about CSL Limited and its related bodies corporate (CSL) financial results and estimates, business prospects and products in research, all of which involve substantial risks and uncertainties, many of which are outside the control of, and are unknown to, CSL. You can identify these forward looking statements by the fact that they use words such as “anticipate,” “estimate,” “expect,” “project,” “intend,” “plan,” “believe,” “target,” “may,” “assume,” and other words and terms of similar meaning in connection with any discussion of future operating or financial performance. Factors that could cause actual results to differ materially include: the success of research and development activities, decisions by regulatory authorities regarding approval of our products as well as their decisions regarding label claims; competitive developments affecting our products; the ability to successfully market new and existing products; difficulties or delays in manufacturing; trade buying patterns and fluctuations in interest and currency exchange rates; legislation or regulations that affect product production, distribution, pricing, reimbursement, access or tax; litigation or government investigations, and CSL’s ability to protect its patents and other intellectual property. The statements being made in this presentation do not constitute an offer to sell, or solicitation of an offer to buy, any securities of CSL.

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Trademarks

Reported Financials

Sales US$3.1 billion, up 11%

- *Underlying[1] sales up 9% @CC[2]*

NPAT US$719 million, up 4%

- *Underlying NPAT up 7% @CC*

EPS US$1.55, up 6%

- *Underlying EPS up 9% @CC*

Cashflow from operations US$705 million, up 8%

Interim dividend US$0.58, unfranked

- *Converted to A$0.81, up ~10%*

1. *Underlying excludes financials relating to the Novartis influenza vaccines business (NVS-IV) NVS-IV was acquired on 31 July 2015*
2. *Constant Currency (CC) removes the impact of exchange rate movements to facilitate comparability of operational performance. See end note for further detail.*

Operational Highlights

Product Portfolio

- Double digit sales growth in all plasma therapy groups
- CSL 654 (rIX-FP) - license under review in U.S. & EU
- CSL 627 (rVIII-SC) - license under review in U.S.
- Respreeza® approved in EU
- CSL 362 (AML) – licensee (Janssen) commenced Phase II study
- CSL 112 (rHDL) - Phase IIb fully enrolled

Operational Highlights — Business Development

Operations

- New Privigen® manufacturing facility approved by U.S. FDA
  - First shipment December 2015
- New sales office opened in Russia

Influenza

- Novartis influenza vaccines acquisition closed 31 July 2015
- 'Seqirus' Launched – No. 2 global influenza vaccine manufacturer
- FLUAD™ approved by U.S. FDA
- Quadrivalent influenza vaccines – licenses under regulatory review

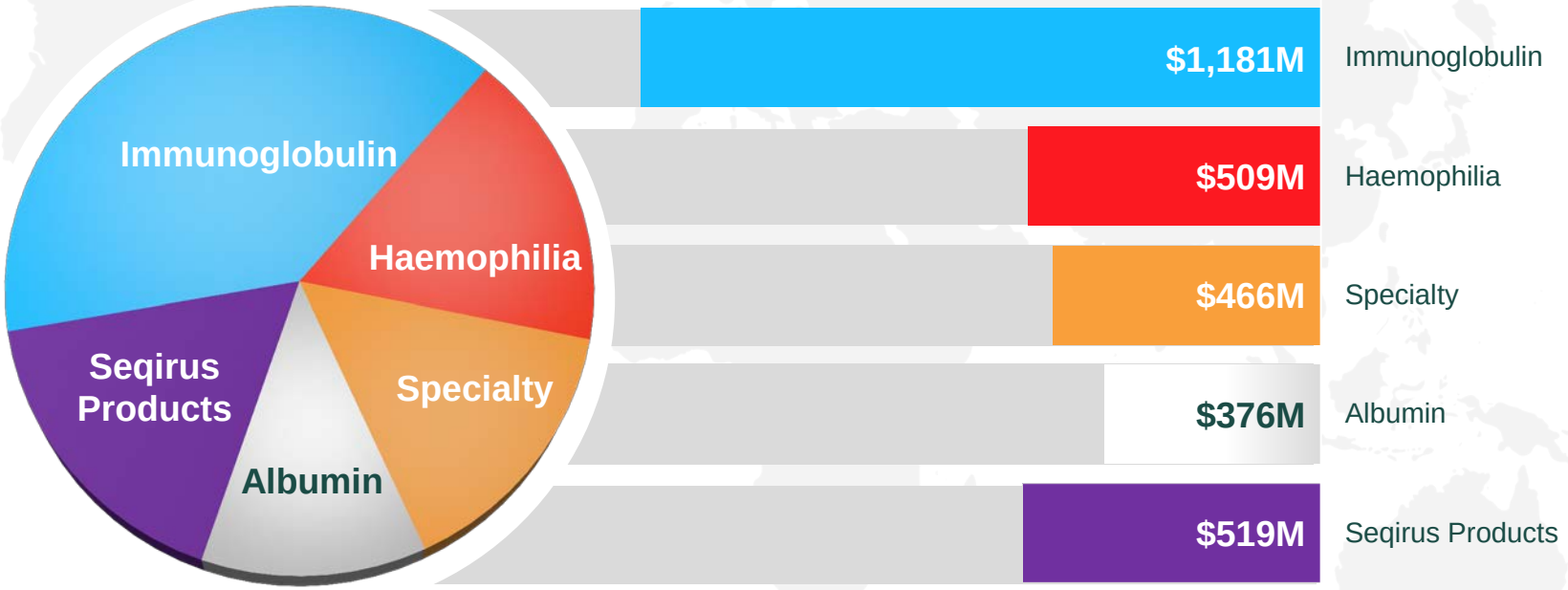
Capital Management

- A$1 billion share buyback[1] underway
- ~US$500 million private placement completed
- New US$1.25 billion bank debt facilities negotiated

1. *CSL reserves the right to suspend or terminate buy-backs at any time*

Group Sales

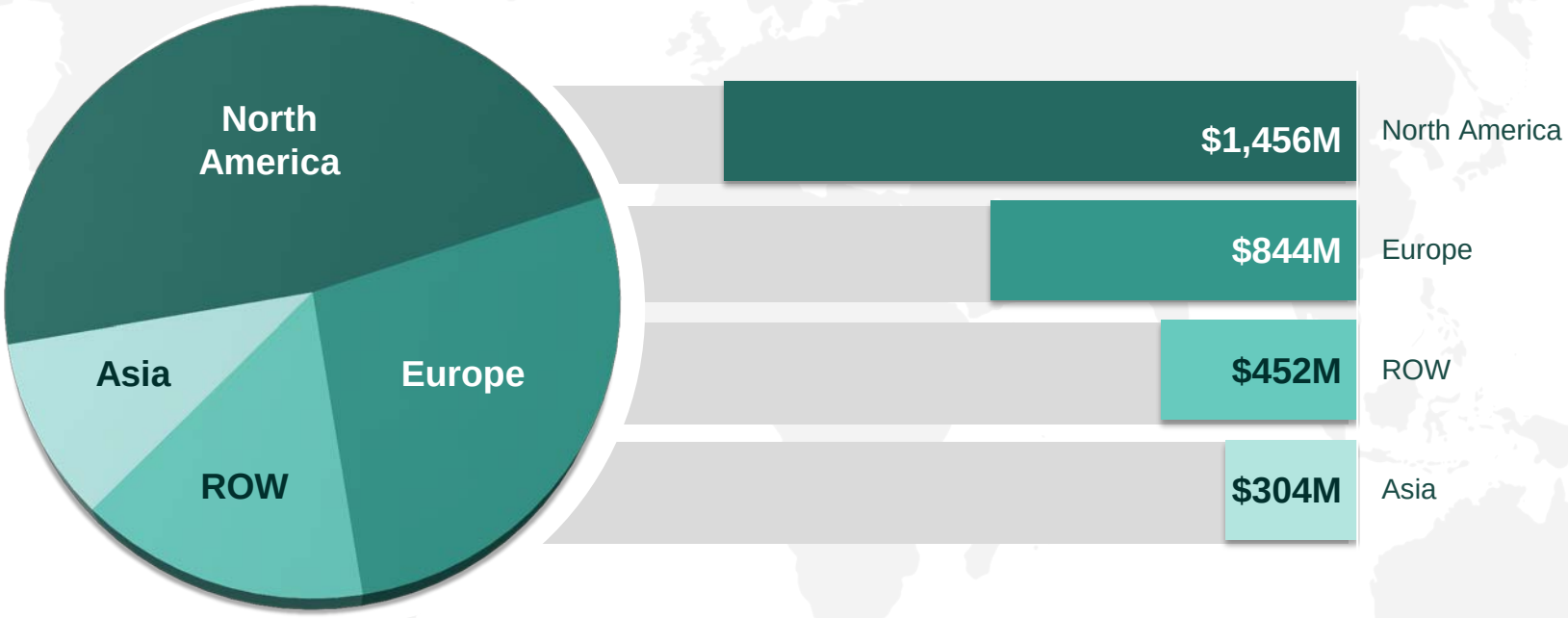
CSL 1H16 Sales US$3.1B



Broad portfolio of products

Broad Sales Reach

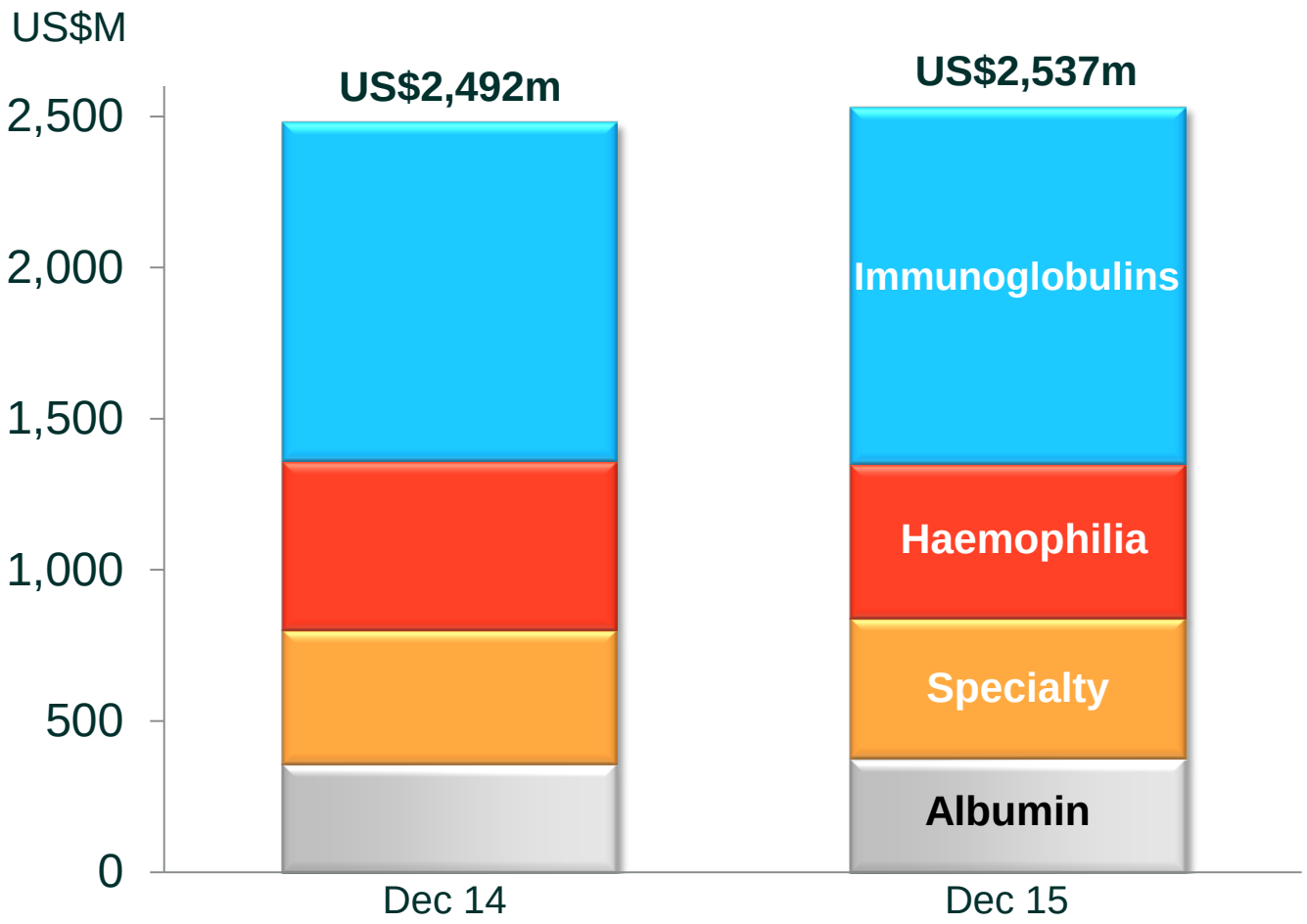
CSL 1H16 Sales US$3.1B



Business Unit Performance

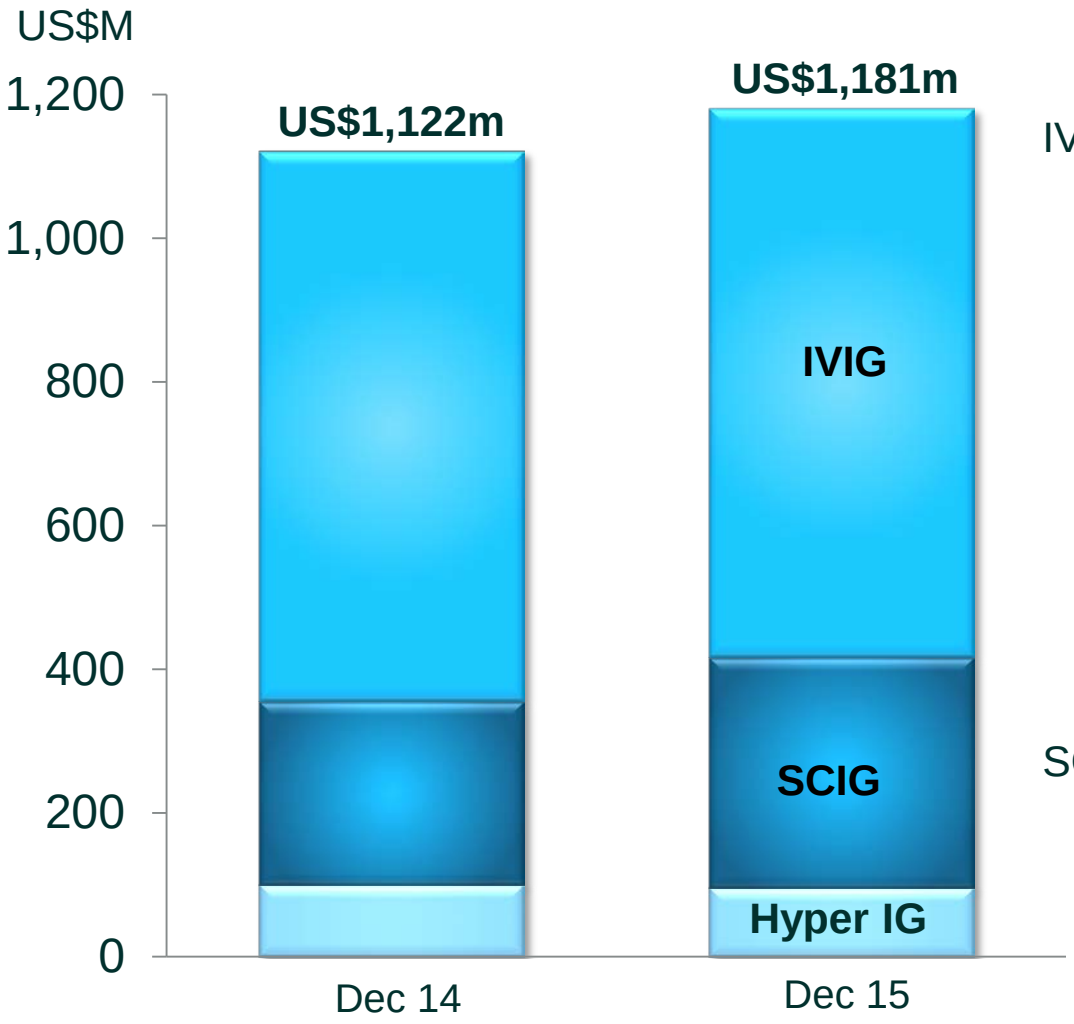
CSL Behring

Product Sales up 10% @CC



Reported sales for the 6 month period

Immunoglobulins Sales up 13% @CC



Reported sales for the 6 month period

Highlights

IVIG

Privigen® up 13%

- Ig IsoLo™ step added to manufacturing process

North America

- Strong demand in the Specialty Pharmacy segment

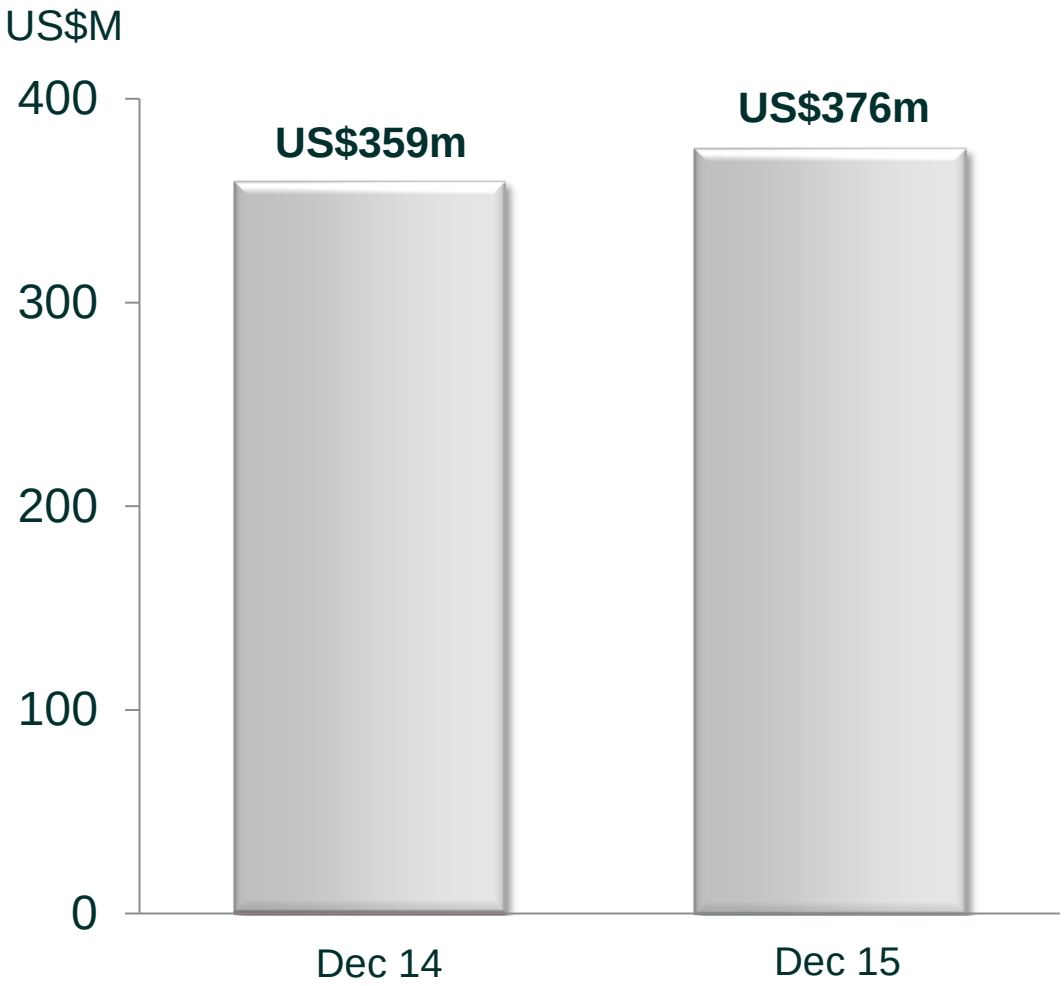
Europe

- CIDP indication driving strong Privigen® demand

SCIG up 31%

Strong growth for Hizentra® from new patient starts and IVIG conversions

Albumin Sales up 10% @CC



Reported sales for the 6 month period

Highlights

China

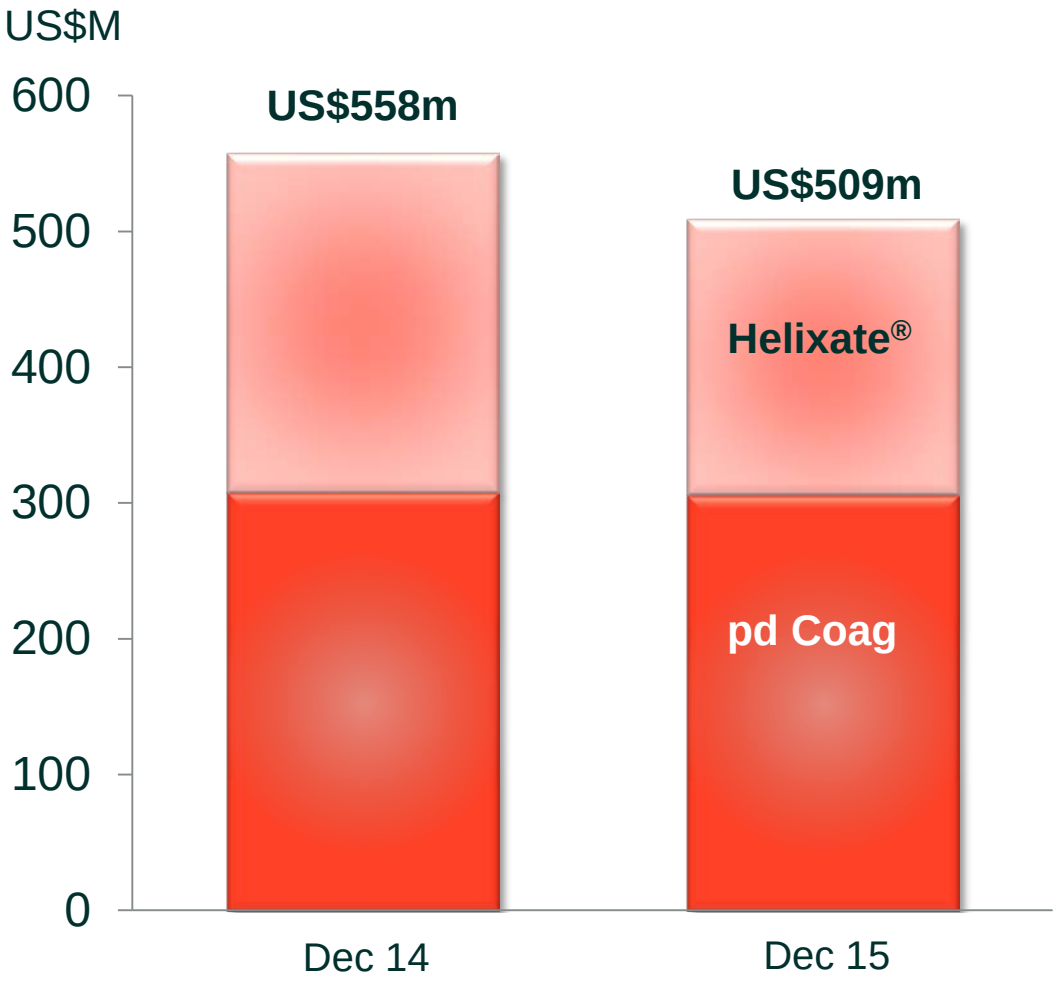
- 17% volume growth
- Successful market penetration into Tier 2 & Tier 3 cities

US

- Solid demand
- Expansion of IDNs and large hospital contracts contributing to majority of growth

Haemophilia

Sales up 2% @CC



Reported sales for the 6 month period

Highlights

Recombinant (Helixate®)

- Volumes declined as broader market transitions to new generation products

Plasma Derived

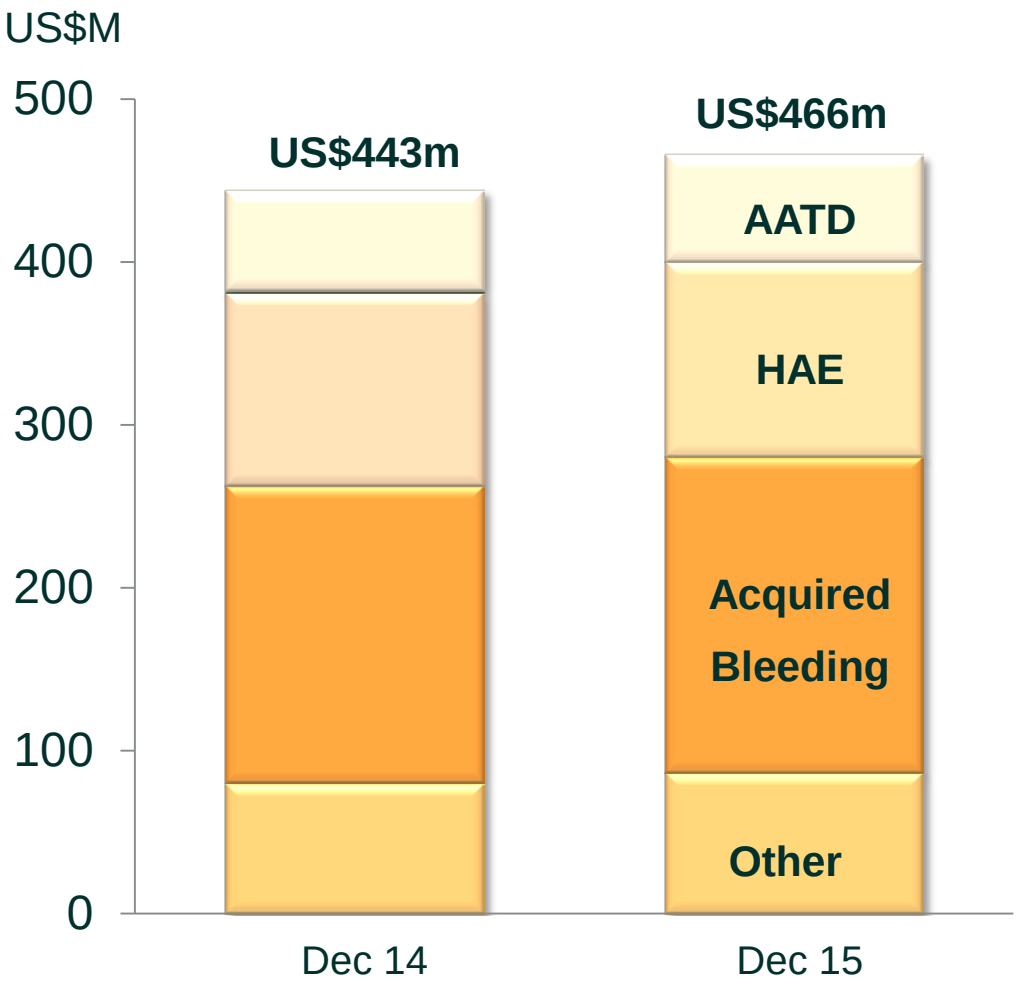
Beriate® volume up 31%

- Growth driven by Poland & Germany
- Successful tenders in Russia & Iran

Volume growth arising in lower priced markets

Specialty

Sales up 14% @CC



Reported sales for the 6 month period

Highlights

Zemaira® / Respreeza®

- Launched in EU

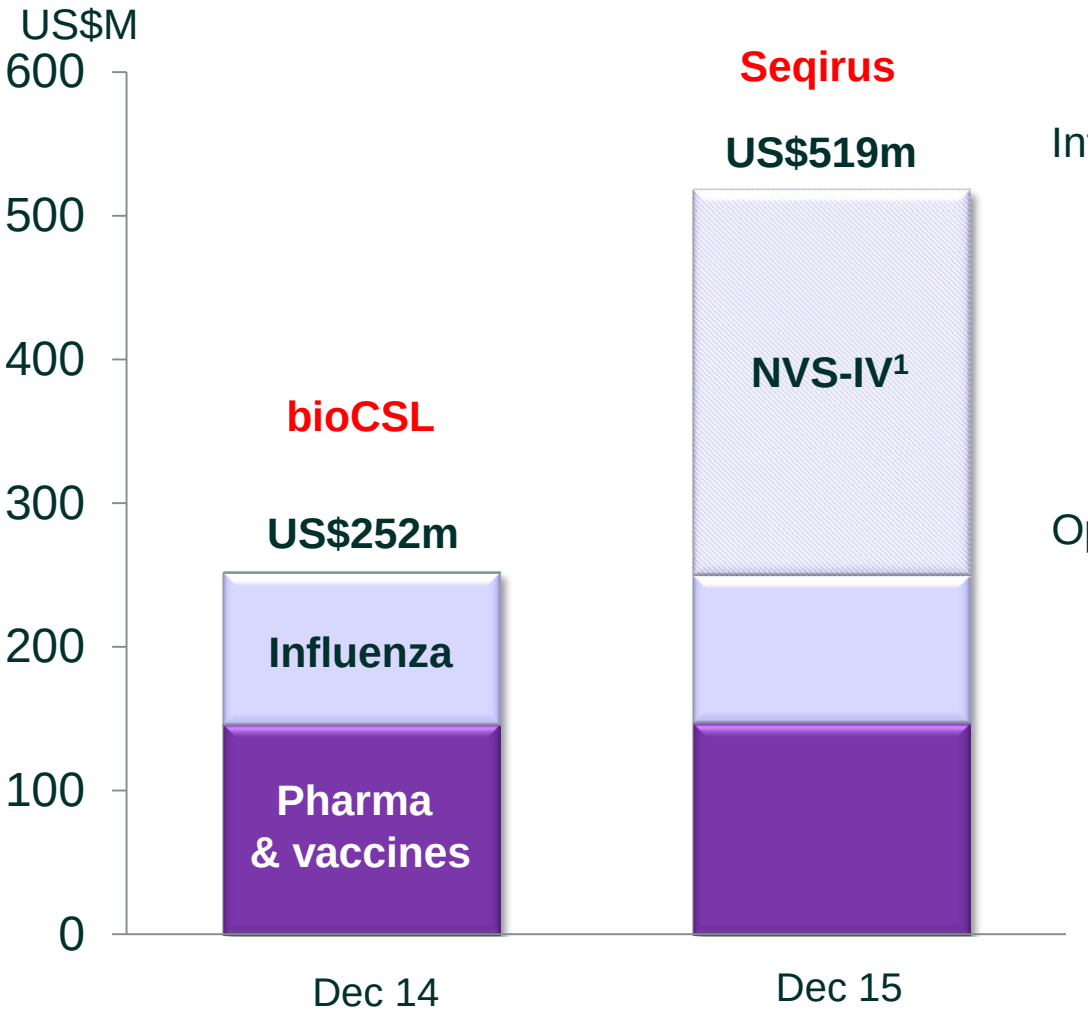
Berinert® P

- New patient starts in the US up 24%

Kcentra® / Beriplex®

- Ongoing demand growth
- Sales up ~40%
- Increasing brand awareness

Seqirus



Reported sales for the 6 month period

Highlights

Integration

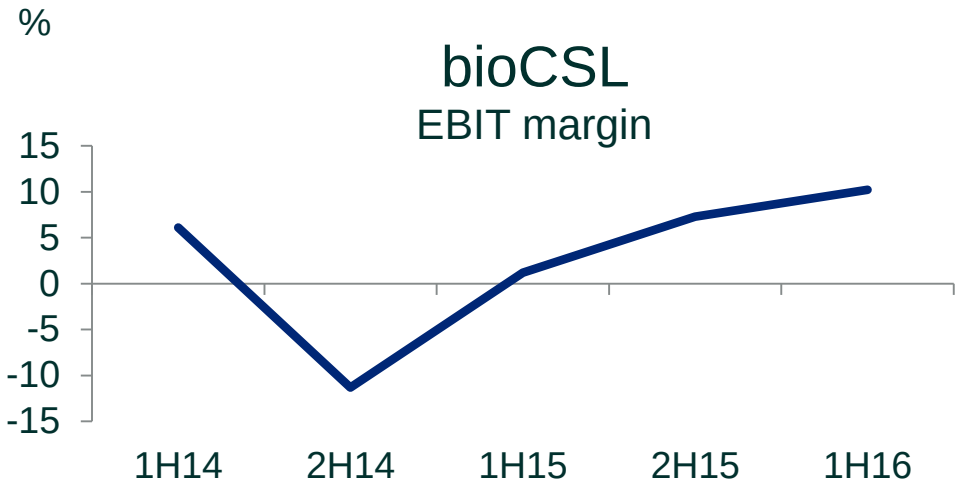
- Novartis influenza vaccines acquisition completed 31 July 2015
- Seqirus launched November 2015
- Corporate office and global leadership team established

Operational

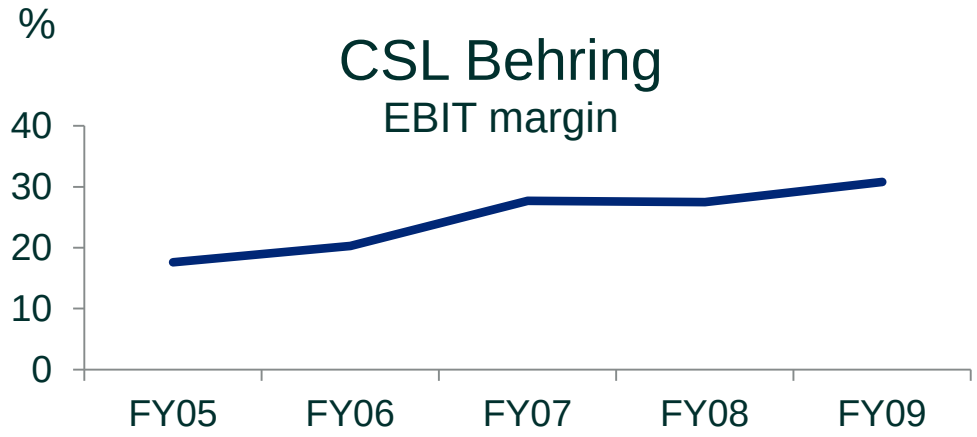
- U.S. FDA approval for FLUAD™
- Cell culture QIV filed with U.S. FDA
- Afluria® QIV filed with U.S. FDA
- Sales impacted by mild northern hemisphere influenza season

1. *Novartis influenza vaccines (NVS-IV) sales for 5 months*

Business Turnarounds



bioCSL was formed in 2013 to improve focus on operational efficiency and drive growth initiatives



CSL Behring was acquired in 2004

CSL is experienced turning around businesses

Business turnarounds take time

CSL Intellectual Property

Segment Revenue $64m, down 29% @CC

- Decline in revenue arising from license payment relating to CSL 362 included in prior period

HPV royalties $63m

- Registration of 9-valent HPV vaccine in US by Merck

CSL 362 (anti-IL-3Rα mAb)

- Exclusive worldwide license with Janssen Biotech Inc to develop and commercialise CSL 362
- Phase 2 AML study commenced by Janssen July 2015
- Commitment to exploratory study in SLE (Lupus) patients

R&D Update

rIX-FP

- rIX-FP Phase III efficacy data supports 7-14 day dosing
- Adult and pediatric indications under review by EMA and FDA
- Health Canada approval received in Jan 16

rVIII-SingleChain

- Phase I/III data supports twice weekly prophylaxis
- BLA accepted for review by FDA in July 15
- MAA submitted to EMA in Dec 15

rVIIa-FP

- Congenital deficiency Phase I/II and Phase II/III in patients with inhibitors continue

Hizentra®

- Hizentra® flexible dosing registration in US
- Hizentra® CIDP pivotal study recruitment completed

R&D Update

Beriplex®

- Phase III study in Japan nearing completion
- Exploring utility in treating bleeding patients receiving NOACs

Berinert®

- CSL 830 (subcut) pivotal Phase III study recruitment completed
- Anti-FXIIa mAb pre-clinical development in HAE completed

Zemaira®/Respreeza® (Alpha1-Proteinase Inhibitor)

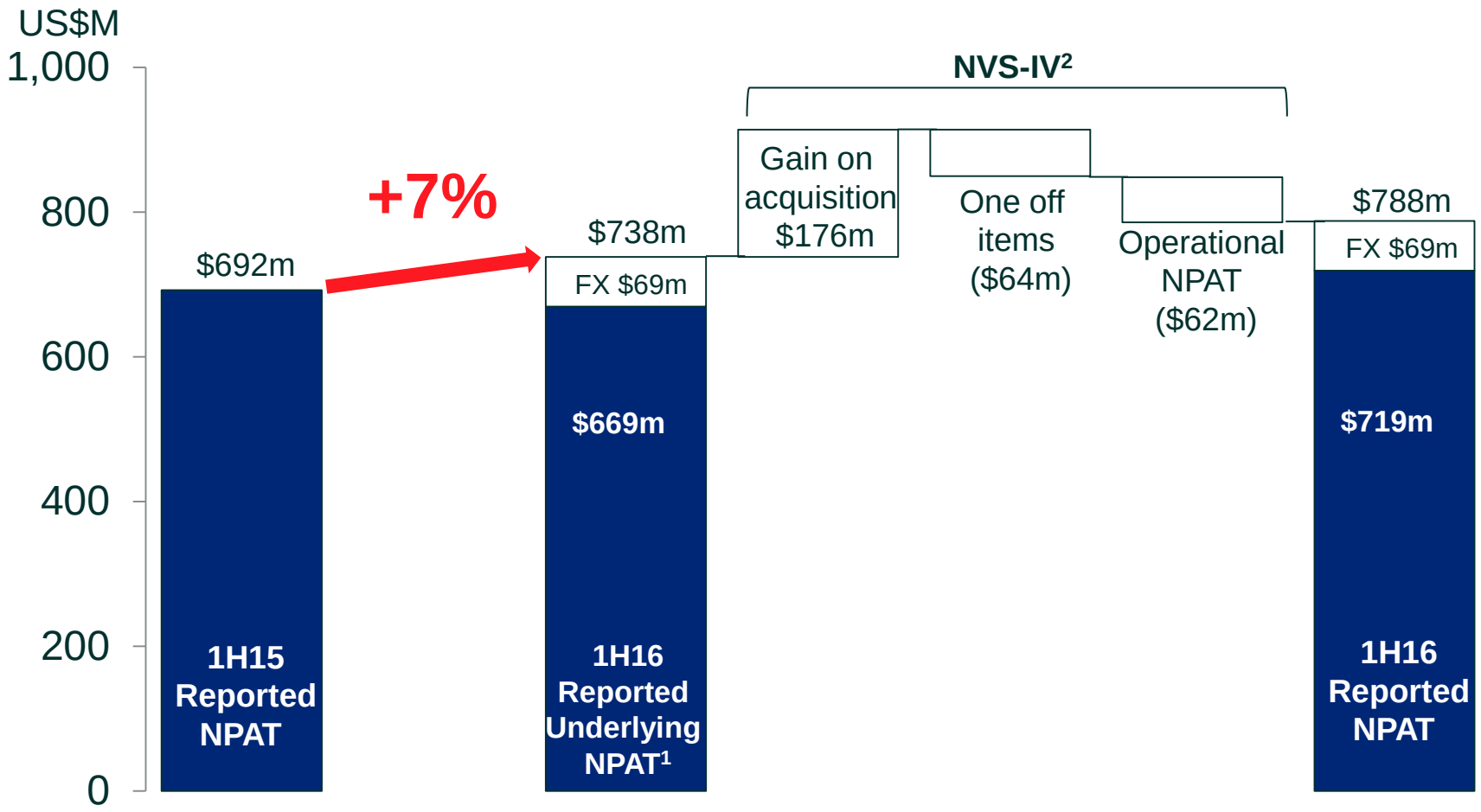
- Patients with AATD treated with Respreeza® have lower annual rate of lung density decline
- Respreeza® approved by EMA in August 15

CSL 112 (reconstituted High Density Lipoprotein)

- AEGIS-I Phase IIb study fully recruited
- Planning for Phase III commenced

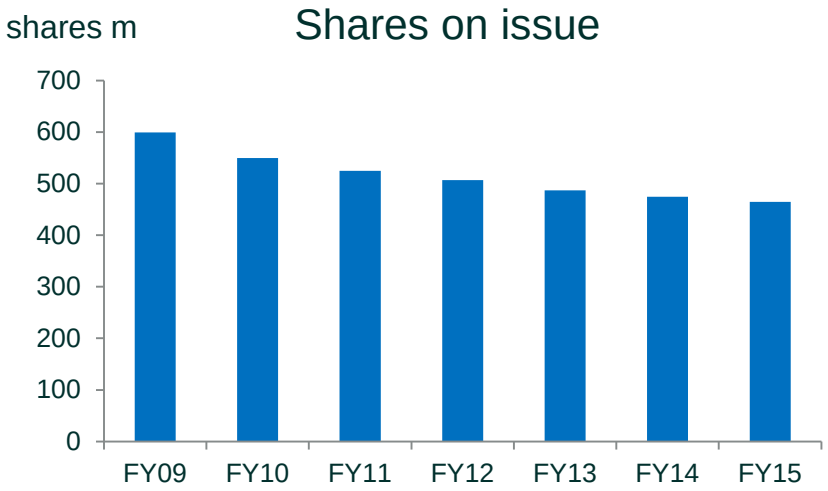
Financials

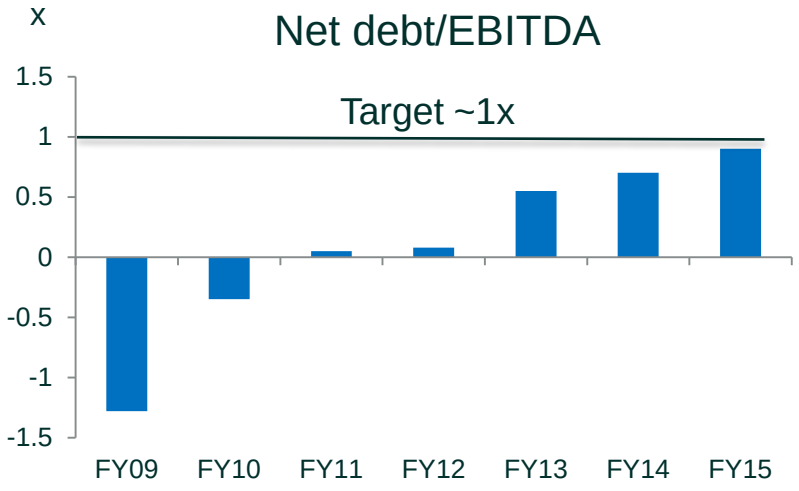
1H16 Profit Growth



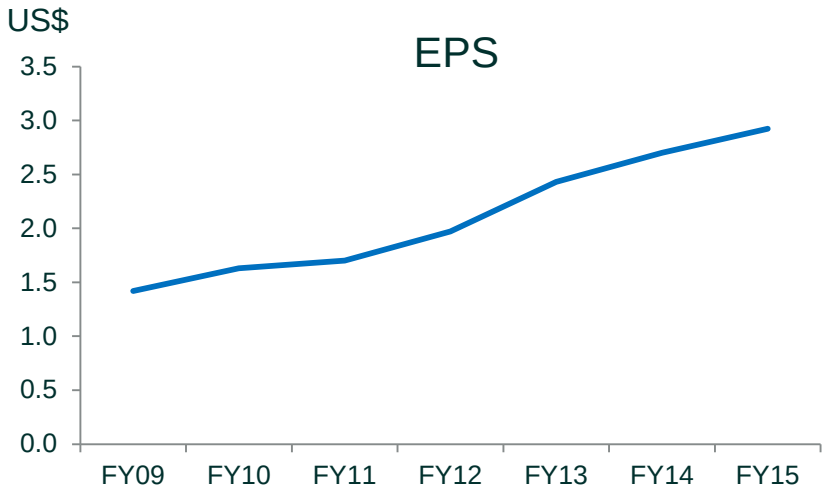
1. *Underlying excludes financials relating to the Novartis influenza vaccines business (NVS-IV)*
2. *NVS-IV was acquired on 31 July 2015*

Effective Capital Management

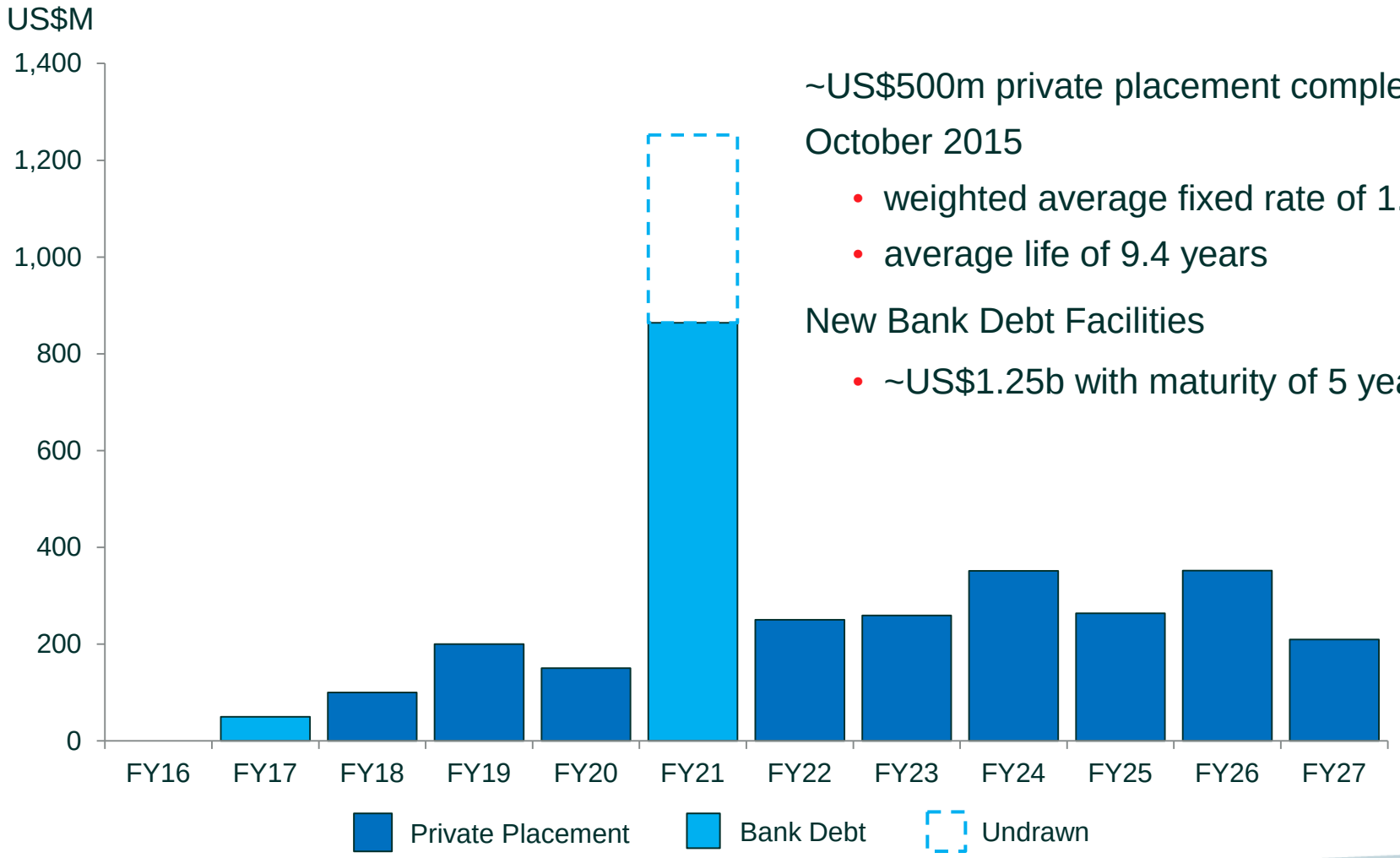




- 27% of shares repurchased over the last 10 years
- Buyback program continues to drive EPS accretion
- WACC continues to reduce



Debt Maturity Profile



~US$500m private placement completed October 2015

- weighted average fixed rate of 1.43%
- average life of 9.4 years

New Bank Debt Facilities

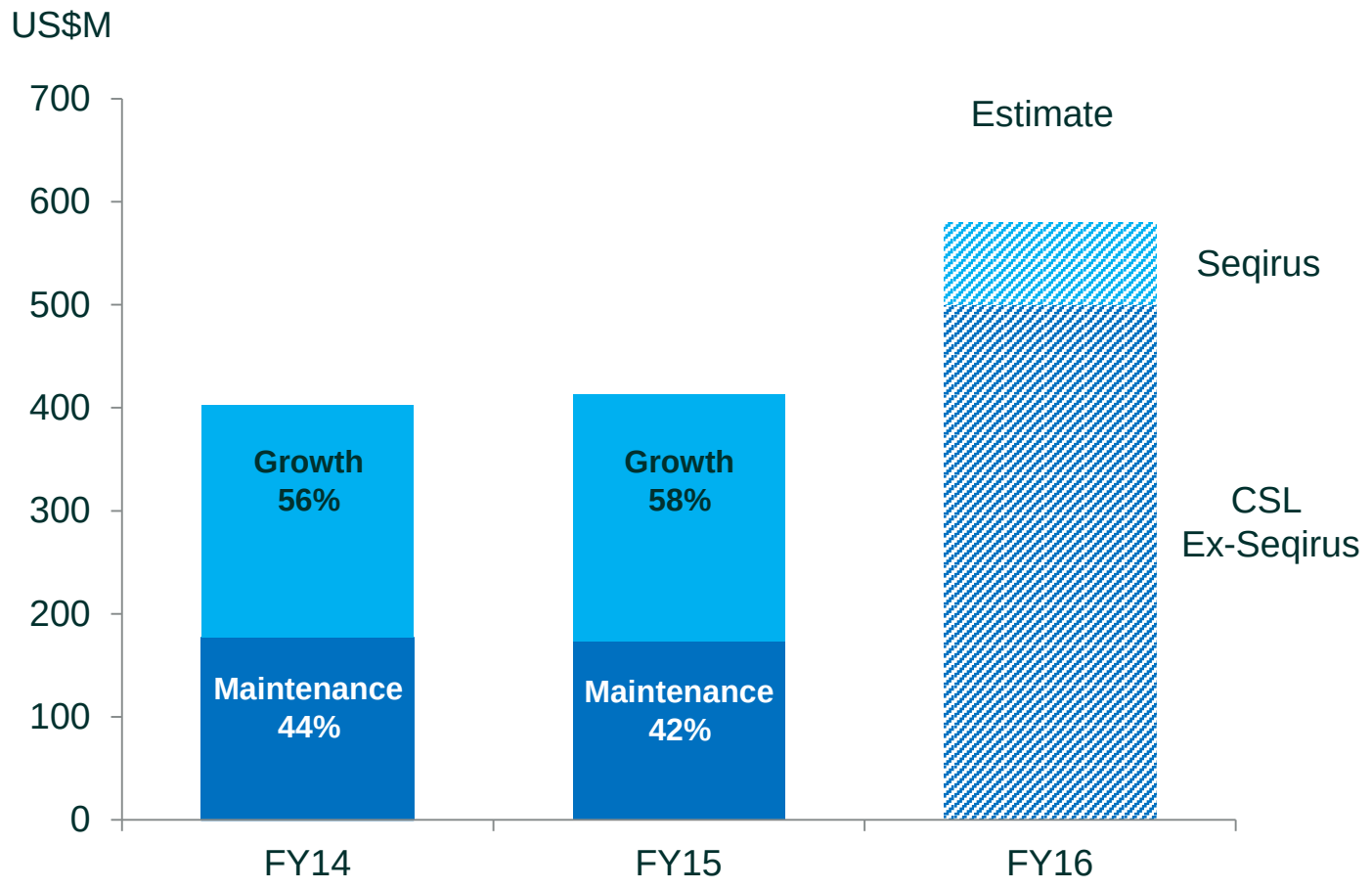
- ~US$1.25b with maturity of 5 years

Continued Investment[1]

Selling & Marketing	up 18% @CC[2] +$43m	• Expansion of commercial operations • Expanding geographic sales reach
Research & Development	Flat @CC	• 1H / 2H phasing
Depreciation & Amortisation	Up 12% @CC +$11m	• Multi-site capacity expansion
Net financing costs	Up 22% @CC	• Private Placement

1. *Reflects movements in the underlying business. Excludes financials relating to the Novartis influenza vaccines business (NVS-IV)*
2. *Constant Currency (CC) removes the impact of exchange rate movements to facilitate comparability of operational performance*

Capital Expenditure



Outlook for FY16[1]

CSL[2]

Revenue growth	~7% @CC[4]
NPAT growth	~5% @CC[4]

NVS-IV[3] ***(11 Months)***

Revenue	~US$360m[5]
NPAT	~(US$90-120m)

- **CSL[2] guidance reaffirmed**
- EPS growth will exceed NPAT growth driven by past and current capital management initiatives

- NVS-IV gain on acquisition less acquisition related one-off costs ~US$90 million

1 For forward looking statements, refer to Legal Notice on page 2
2 Excludes Novartis influenza vaccines business (NVS-IV)
3 Influenza vaccine business acquired from Novartis 31 July 2015
4 Constant Currency (CC) removes the impact of exchange rates movements to facilitate comparability
5 Excludes gain on acquisition ~US$176m

CSL Strategy for Profitable Growth

Future Growth Pipeline

- CSL 112 – new treatment paradigm in ACS
- CSL 830 – HAE
- Pipeline antibodies
- Targeted business development

Growth Drivers

Seqirus

- Drive Seqirus business to profitability
- Successfully launch pipeline vaccines

CSL Behring

- Launch recombinant coagulation factors
- Maintain leadership in Ig and albumin
- Grow high-margin specialty products

Core Plasma

- Relentless commitment to lowest cost base
- Remain ahead of the demand curve
- Organic growth of core plasma products

Sustained Financial Performance

CSL Limited
2016 Half Year Result
16 February 2016

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Group Results

Adjusted for NVS-IV

Half year ended December US$ Millions	Dec 2014 Reported	Dec 2015 Reported	Dec 2015 NVS-IV[2]	Dec 2015 Underlying[1]	Dec 2015 Underlying[1] at CC[3]	Change %
Sales	**2,744**	**3,056**	**294**	**2,762**	**2,996**	**9.2%**
Other Revenue / Income	96	80	4	76	79	
Total Revenue / Income	**2,841**	**3,136**	**298**	**2,838**	**3,075**	**8.2%**
Earnings before Interest, Tax, Depreciation & Amortisation	**969**	**848**	**(112)**	**960**	**1,053**	**8.7%**
Depreciation/Amortisation	91	102	9	93	102	
Earnings before Interest and Tax	**878**	**746**	**(121)**	**867**	**952**	**8.3%**
Gain on Acquisition		176	176			
Net Interest Expense / (Income)	21	27	1	26	26	
Tax Expense	165	176	5	171	188	
Reported Net Profit after Tax	**692**	**719**	**50**	**669**	**738**	**6.6%**
Interim Dividend (US$)	0.58	0.58				
Basic EPS (US$)	1.46	1.55			1.59	9%

1. *Underlying excludes financials relating to the Novartis influenza vaccines business*
2. *Novartis influenza vaccines acquisition as from 31 July 2015*
3. *Constant Currency (CC) removes the impact of exchange rate movements to facilitate comparability of operational performance.*

CSL Behring Sales

Half year ended December US$ Millions	Dec 2014	Dec 2015	Dec 2015 CC[1]	Change %
Immunoglobulins	**1,122**	**1,181**	1,269	**13.2%**
Albumin	**359**	**376**	395	**10.1%**
Haemophilia	**558**	**509**	569	**2.0%**
- Recombinants	250	203	221	(11.5%)
- Plasma	308	306	348	12.9%
Specialty	**443**	**466**	506	**14.2%**
Total Product Sales	**2,482**	**2,532**	2,740	**10.4%**
Other sales (mainly plasma)	10	5		
Total Sales	2,492	2,537		

1. Constant Currency (CC) removes the impact of exchange rate movements to facilitate comparability. See end note for further detail.

Financial Appendix[1]

Full Year ended June US$ Millions	CSL[2] FY15 Actual	CSL[2] FY16 Guidance	NVS-IV[3] (11 months) FY16 Guidance
Total Revenue	5,613	~7% @CC[4]	~360[5]
Reported Net Profit after Tax	1,379		~(90-120)
NVS-IV gain on acquisition less acquisition related one-off costs			~90
Adjusted Net Profit after Tax	1,401	~5% @CC[4]	~(180-210)
FX Impact[6]		~(100)	

1 For forward looking statements, refer to Legal Notice on page 2
2 Excludes Novartis influenza vaccines business (NVS-IV)
3 Influenza vaccine business acquired from Novartis 31 July 2015
4 Constant Currency (CC) removes the impact of exchange rates movements to facilitate comparability
5 Excludes gain on acquisition ~US$176m
6 Assumes current rates remain steady for the remainder of the year, giving rise to the unfavourable full year FX impact

Notes

(#) Constant currency removes the impact of exchange rate movements to facilitate comparability of operational performance. This is done in three parts: (a) by converting the current period net profit of entities in the group that have reporting currencies other than US Dollars at the rates that were applicable to the prior comparable period ("translation currency effect"); (b) by restating material transactions booked by the group that are impacted by exchange rate movements at the rate that would have applied to the transaction if it had occurred in the prior comparable period ("transaction currency effect"); and (c) adjusting for current year foreign currency gains and losses. The sum of translation currency effect, transaction currency effect and foreign currency gains and losses is the amount by which reported result is adjusted to calculate the operational result.

Summary NPAT

Reported Net Profit after Tax	$718.8m
Translation Currency Effect (a)	$ 64.8m
Transaction Currency Effect (b)	$ (9.8m)
Foreign Currency Gains and Losses (c)	$ 13.7m
Constant Currency Net Profit after Tax *	$787.5m

(a) Translation Currency Effect $64.8m

Average Exchange rates used for calculation in major currencies (six months to Dec 15/Dec 14) were as follows: USD/EUR (0.91/0.77); USD/CHF(0.97/0.93)

(b) Transaction Currency Effect ($9.8m)

Transaction currency effect is calculated by reference to the applicable prior comparable period exchange rates. The calculation takes into account the timing of sales both internally within the CSL Group (ie from a manufacturer to a distributor) and externally (ie to the final customer) and the relevant exchange rates applicable to each transaction.

(c) Foreign Currency Losses (13.7m)

Foreign currency losses recorded during the period

Summary Sales

Reported Sales	$3,056.3m
Currency Effect	$ 234.0m
Constant Currency Sales *	$3,290.3m
Less NVS-IV sales	$ 294.6m
Underlying operational business sales @ CC	$2,995.7m

* Constant Currency Net Profit after Tax and Sales have not been audited or reviewed in accordance with Australian Auditing Standards.