

Interim Results H1 FY20 Highlights



Positive EBITDA of \$0.6 Million for the period ended 31 December 2019

Including a one-time revenue adjustment, IDT generated a positive EBITDA of \$0.6 million representing a year-on-year improvement of 155% (\$1.1 million EBITDA loss reported as at 31 December 2018)

IDT base business results reflect gross margin improvement and ongoing cost control initiatives

Excluding the one-time adjustment, revenue from operations was \$0.26 million lower year-on-year, however the drive for higher value / high margin activities delivered an improved gross margin of \$0.36 million.

Net reported loss from operations after tax was \$1.18 million representing a year-on-year improvement of 51% (\$2.39 million loss reported as at 31 December 2018)

Meaningful revenues starting to flow in from medicinal cannabis manufacturing activities

Revenues starting to flow in from commercial scale-up activities associated with converting medicinal cannabis biomass into high value GMP Active Pharmaceutical Ingredients and Finished Dosage Form products

Cost containment initiatives continuing to deliver year-on-year savings

Total operating expenses (including salaries) were \$0.58 million lower year-on-year; of which \$0.3 million relates to the prior period's U.S. Food and Drug Administration (FDA) Warning Letter remediation activities



Interim Financial Results H1 FY20 Executive Summary

Profitable EBITDA of \$618K due to:

- Reported total revenue up 8% year-on-year
- Revenue and improved gross margins underpinned by a focus on scalable and more profitable income streams
- One time revenue adjustment relating to recognition of previously capitalised temozolomide contract milestone (\$0.9m)
- Other operating expenses down 12% year-on-year due to FDA Warning Letter close-out (FY`18) and reduction in salary expenses

Materially Improved Reported Loss After Tax of \$1.183m

 Loss for the half year improved by \$1.209m (51%) after recognition of one-time asset impairment of temozolomide

Results Support IDT's Push Towards Profitability

| | HYE 31 Dec 2019 \$ 000 | HYE 31 Dec 2018 \$ 000 |
|---------------------------------|------------------------------|------------------------------|
| Total Revenue | 7,213 | 6,672 |
| Raw materials | 768 | 1,216 |
| Other operating expenses | 5,827 | 6,588 |
| EBITDA | 618 | (1,132) |
| Depreciation and Amortisation | 1,155 | 1,262 |
| Asset impairment - temozolomide | 736 | 0 |
| Loss before tax | (1,273) | (2,394) |
| Income tax benefit | 90 | 2 |
| Reported loss for the half year | (1,183) | (2,392) |

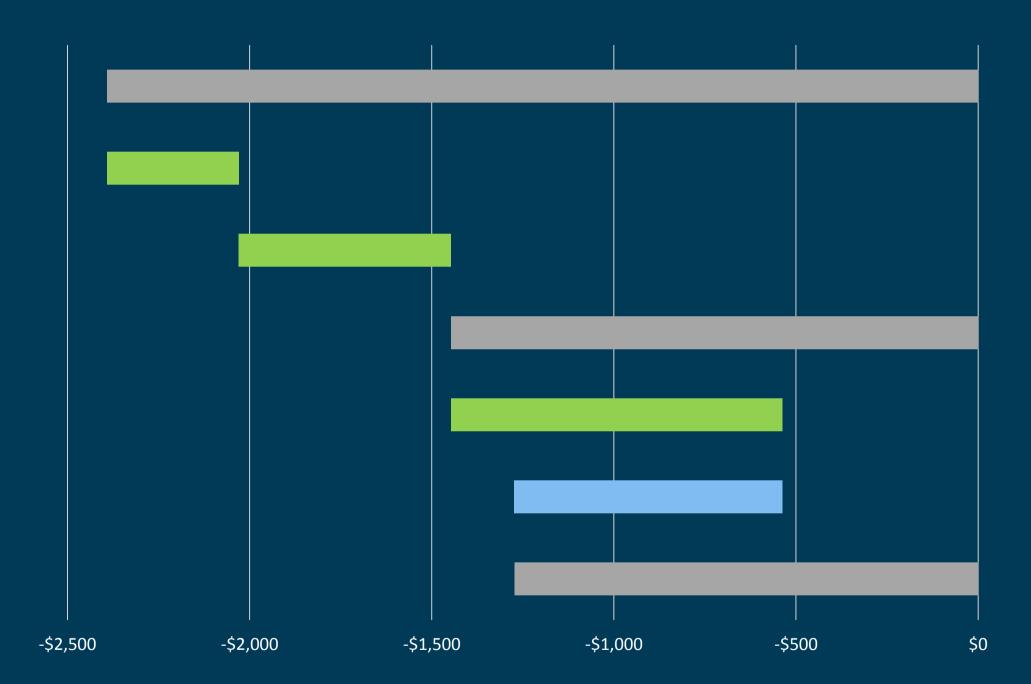


IDT Base Business Operations H1 FY20

Year-on-Year Analysis

51% Year-on-year reduction in net reported loss from operations:

- \$2.392m Reported loss before tax 31 December 2018
 - \$362K Improved gross margin
 - \$583K Cost reduction initiatives
 - \$1.447 Underlying loss before tax
- \$910K One-time revenue adjustment temozolomide
 - \$736K Impairment intangible asset temozolomide
- \$1.273m Reported loss before tax 31 December 2019





The Year Ahead - Half Year Check-in

[Slide presented at IDT's 2019 Annual General Meeting]



Drive and Grow The Business

Year-on-year top-line revenue increased by 8% and gross margin improved by 28%



Continued Focus on Capital Management and Cost Containment

Total operating expenses (including salaries) reduced by \$0.58 million year-on-year



Roll-out Medicinal Cannabis Manufacturing Plan

Commenced medicinal cannabis resin extraction (API) and Finished Dosage Form development and manufacturing activities

Medicinal cannabis biomass supply and commercialisation partnerships

In-progress: Proprietary and partnership medicinal cannabis opportunities under evaluation



Commence commercial scale medicinal cannabis manufacturing

Commercial scale solvent extraction and finished dosage form activities commenced Proprietary/partnered product portfolio expansion

Launch first proprietary/ partnered medicinal cannabis product New product launches of downstream value added products



IDT Australia Limited

Board of Directors

Mr Alan Fisher Chair

Finance specialist with key experience in business restructuring and company turnarounds.

Mr Hugh Burrill

Pharma pipeline portfolio management and product development.
(Ex Hospira/Mayne Pharma)

Ms Mary Sontrop

Biopharmaceutical executive with global experience in quality, manufacturing, regulatory and business integration.

(Ex CSL Behring)

Executive Team

Dr David Sparling CEO

More than 20 years of pharma and diagnostic experience in CEO, Director and corporate/business development roles. (Ex Agenix Limited, GTG Limited)

Joanna Johnson CFO

More than 20 years of pharma experience in finance roles. (Ex F H Faulding, Hospira/Mayne Pharma and Lupin)

Jim Sosic VP Operations Infrastructure

More than 20 years in manufacturing and supply chain roles. (Ex CSL Behring, Hospira/Mayne Pharma)

Daniel Broadhurst Head of Quality

More than 15 years of R&D, operations and quality experience.
(Ex CSL Behring and Catalent)

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