



Lumos Diagnostics Holdings Limited Annual General Meeting – CEO Presentation

November 2023

www.lumosdiagnostics.com

Disclaimer and Important Information



This presentation (Presentation) has been prepared solely for informational purposes by Lumos Diagnostics Holdings Limited (Company).

The information contained in this document ("Document") has been prepared by Lumos Diagnostics Holdings Limited (referred to as "Lumos" or "Company"). This Document is current as at the date of this Document and should be read in conjunction with other Lumos periodic and continuous disclosure announcements filed with the Australian Securities Exchange (ASX), available at www.asx.com.au.

The information in this Document is not intended to form the basis of any investment decision in relation to the Company or its assets and should not be considered as a recommendation to the Recipient to acquire securities in the Company. This Document is not a prospectus, profile statement or disclosure document and does not constitute an offer or invitation to acquire securities or otherwise invest in the Company, and no agreement to subscribe for securities will be entered into on the basis of this Document.

No representation or warranty, expressed or implied, is or will be made, and no responsibility or liability is or will be accepted by the Company, any of their respective officers, servants, agents or advisers (collectively "Limited Parties") as to or in relation to the accuracy, reasonableness, completeness or reliability of the information in this Document or any other written or oral information made available to any Recipients or their advisers. Any liability therefore is hereby expressly disclaimed. In particular, no representation or warranty is given as to the achievability or reasonableness of any future projections, management estimates or plans, prospects, returns or forecasts.

To the fullest extent permitted by law, the Limited Parties will not have any responsibility or liability for any loss or damage (whether foreseeable or not), however arising (including as a result of negligence), in relation to or in connection with the provision of this Document, the Recipient's or any other person's purported reliance on this Document, the failure to provide information of which any of the Limited Parties becomes aware or any errors in or omissions from this Document.

None of the Limited Parties makes or gives any representation, warranty or guarantee, express or implied, that the information in this Document is accurate, current, reliable or complete, has been or will be audited or independently verified, or that reasonable care has been taken in compiling, preparing or furnishing it. Various statements in this Document constitute statements relating to intentions, future acts and events including forecast financial information ("Forward Looking Statements"). Forward Looking Statements involve subjective judgment and analysis, known and unknown risks, uncertainties and other important factors that may cause those future acts, events and circumstances to differ from the way or manner in which they are expressly or impliedly portrayed herein.

The Limited Parties do not make or give any representation, warranty or guarantee, express or implied, that any Forward Looking Statements will be achieved or proven correct, or that any assumptions or projections on which the Forward Looking Statements are based are reasonable. No historical financial information, forecast financial information, estimates or projections contained in this Document or any other financial information derived from that information, can be relied upon as a promise or representation, as to the past, present or the future. Past performance is not necessarily a guide to future likelihood of achievement or reasonableness of any Forward Looking Statement, forecast financial information or other forecast. The Limited Parties do not undertake any obligation to (and expressly disclaim any obligation to) provide the Recipients with access to any additional information or to correct any inaccuracies herein which may become apparent or to disseminate any updates or revisions to any Forward Looking Statements in this Document to reflect any change in expectations in relation to any such statements or any change in events, conditions or circumstances on which any such statement is based.

This document also contains statistics, data and other information relating to markets, market sizes, market shares, market positions and other industry data pertaining to the Lumos business and markets. Such information is generally based on independent market and industry data or research. Lumos has not independently verified and cannot give any assurances as to the accuracy and completeness of the information sourced from market and industry data or research contained herein. Accordingly, the accuracy and completeness of such information is not guaranteed. There is no assurance that any of the forecasts or projections contained in the independent market and industry data or research will be achieved. Forecasts and projections involve risks and uncertainties and are subject to change based on various factors. You should note that market data and statistics are inherently predictive and subject to uncertainty and not necessarily reflective of actual market conditions.

Neither the receipt of this Document by any person nor any information contained in it or supplied with it or subsequently communicated to any person in connection with a proposed investment in the Company constitutes, or is to be taken as constituting, the giving of investment or financial product advice (or any other advice) to any such person. Each such person should make their own independent assessment of the merits or otherwise of investing in the Company and should seek their own professional advice in respect of any future investment opportunity and not act on the basis of any matter contained in this Document. In providing this Document, the Company has not considered the objectives, financial position, taxation situation or other needs of any particular Recipient.

The distribution of this document in jurisdictions outside Australia may be restricted by law. Persons who come into possession of this document who are not in Australia, should seek advice on and observe any such restrictions. Any failure to comply with such restrictions may constitute a violation of applicable securities laws. In particular, this document does not constitute an offer to sell, or a solicitation of an offer to buy, any securities in the United States.

Non-IFRS financial measures

Recipients should note that certain financial data included in this Document is not recognised under the AAS and is classified as 'non-IFRS financial information' under Regulatory Guide 230 'Disclosing non-IFRS financial information' published by ASIC. The Company believes that this non-IFRS financial information provides useful information to users in measuring the financial performance and condition of Lumos. The non-IFRS financial measures do not have standardised meanings under AAS, and therefore may not be comparable with similarly titled measures presented by other entities, nor should these be interpreted as an alternative to other financial measures determined in accordance with AAS. Investors are cautioned not to place undue reliance on any non-IFRS financial information, ratios and metrics included in this Document.

Board of Directors



Sam Lanyon

Non-Executive
Chair



Doug Ward

CEO and
Managing Director



Bronwyn Le Grice

Non-Executive
Director



Lawrence Mehren

Non-Executive
Director



Catherine Robson

Non-Executive
Director

FY23 - Summary of Achievements



“

The opportunity that Lumos has with its technology, know-how and expertise in the development and production of point-of-care diagnostic tests and its unique reader platform remains as compelling as it was when I decided to join the Company, if not more so.

I feel that we have now turned the corner and are finally in a great position to capitalise on those opportunities. I see a very exciting future ahead for Lumos.

Doug Ward
Chief Executive Officer
Lumos Diagnostics



US\$10.2 million Commercial Services revenue in FY23 ▶ 9% increase vs FY22



Completed reorganisation reducing headcount by >60% & consolidated operations to single site



Results of FebriDx DISRUPT trial published in leading peer reviewed journal JAMA



Secured FDA clearance to market FebriDx in the United States at end of FY2023



Established and expanded strategic partnership with leading women's health company Hologic



New commercial services projects in human health, animal health and food testing

Lumos has a compelling and highly competitive offering



1. Fully-integrated—from design to manufacturing



2. Proprietary reader platform for use in different settings



3. IVD development and manufacturing expertise



4. Clinical validation, trial management, and regulatory



Lumos' POC diagnostic test development engine



Lumos' POC Diagnostics Engine



REVENUE STREAMS

Contract Fees
from services

Margin
from manufacturing

Sales & license payments
from own & other products

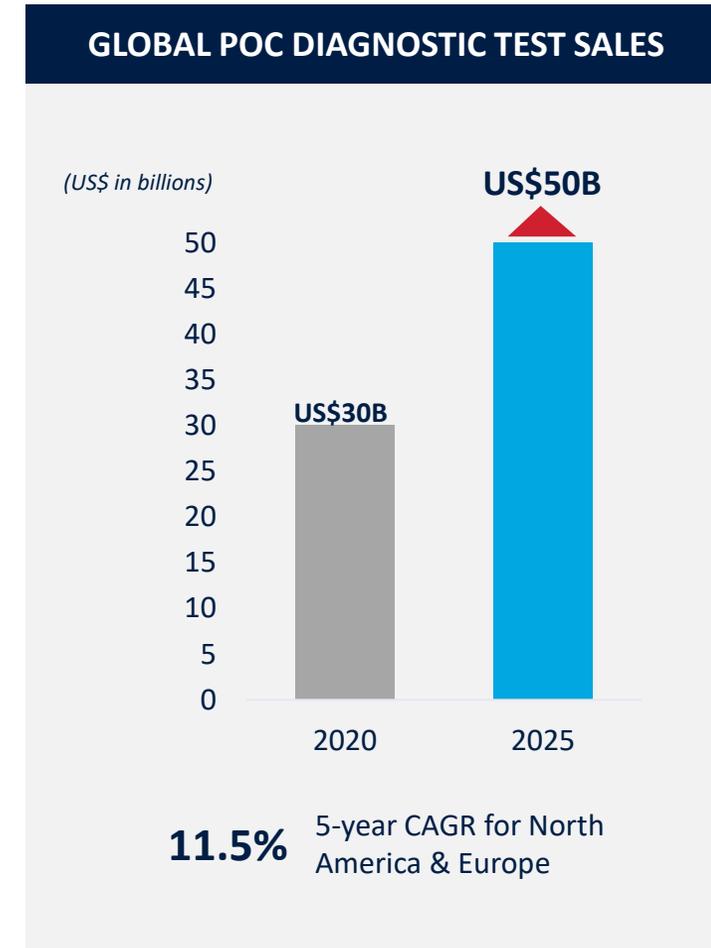
Favourable industry structure and key trends

Robust underlying category growth forecast

- Significant growth anticipated from growing role of diagnostic tests in healthcare
- Rapid, point-of-care tests increasingly being used – common in US settings
- POC tests increasingly critical for enabling real-time provision of healthcare
- Growing demand for next generation diagnostics which incorporate state-of-the-art reader systems that can move lab-based tests closer to the patient and physician

Significant opportunity for diagnostic tests development and manufacturing services

- Current industry is highly fragmented—dominated by specialist niche players
- Few providers with expertise across development, clinical, regulatory and manufacturing
- Major players increasingly relying on external providers—cost effective access to technology
- Requirement for connected POC tests able to interface with Electronic Medical Records



SOURCE: MarketsandMarkets Report, 2021

FY23 – Financial Result Highlights*

“

While Lumos went into FY2023 facing a number of challenges, I am extremely happy with the successes we achieved during the financial year and believe that we have now put the Company on a firm trajectory for long-term growth.

Doug Ward
Chief Executive Officer
Lumos Diagnostics

-  Revenue US\$10.5 million (US\$11.6 million FY22, incl. CoviDx US\$1.7m), revenue from services up 9%
-  Gross profit margin for FY23 of 56%, an improvement of 12% over FY22
-  Significant reduction in cost base with OPEX for FY23 US\$11.9 million, versus US\$22.8M in FY22
-  Significant reduction in loss after tax to US\$9.0 million (US\$45.7 million loss in FY22)
-  Significant reduction in cash usage for the year, to US\$11.6 million (US\$23.6 million cash usage in FY22)
-  Completed capital raise of A\$5.4 million in July 2023
-  Repaid all Convertible Notes, A\$1.58 million in August 2023

* All amounts are in US\$ unless stated otherwise

Commercial Services - FY23 Operational Summary



CARLSBAD, CA USA

- Lumos commercial services revenue of US\$10.2 million – an increase of 9% from FY22
- Completed major reorganisation which included reduction of headcount by over 60%
- Closed Sarasota FL facility and consolidated operations to single site in Carlsbad CA
- Established strategic partnership with leading women’s health company Hologic that included multiple programs
- Expanded and diversified pipeline of commercial services projects beyond infectious disease applications

Strategic partnerships are a key pillar of Lumos' growth plan



Lumos provides a compelling service offering for leading diagnostics companies

- **Fully-integrated offering**—from concept-to-clinic-to-commercial production
- **Proprietary reader platform**—integrate POC testing with electronic medical records
- **Track record**—successful delivery of products to recognised industry leaders

Strategic partnerships will underpin long-term and durable revenue growth for Lumos

- **Multiple projects**—reduced transaction costs with repeat business
- **Project extensions**—as products migrate through stages of the development process
- **New projects**—creating and developing new products for strategic partners
- **Next gen products**—extending commercial life of partner's products as market evolves
- **Manufacturing**—ongoing revenue stream from commercial-stage products



Lumos has established a strategic partnership with Hologic



- **Hologic is a recognized global leader in women's health based in Massachusetts**
 - NASDAQ: HOLX, Market Capitalization US\$20 billion
 - FY2022 revenue of US\$4.2 billion with net income of US\$1.3 billion
 - Diagnostic products account for >50% of Hologic's revenue
- **Lumos is working with Hologic at multiple levels**
 - Multiple services contracts signed during FY2023
 - Existing, marketed product
 - Development of a new, rapid point-of-care test product
 - Other programs and opportunities currently under consideration
 - US\$4.2 million in non-dilutive funding through sale and leaseback agreement



FebriDx – Lumos' POC test to aid antibiotic prescribing



- **FebriDx offers an aid for healthcare providers to improve patient care and antibiotic stewardship**
 - 211 million outpatient antibiotic prescriptions in the US in 2021¹
 - 40% antibiotics prescribed in for respiratory infections unnecessary (ie. patient had no bacterial infection)²
 - Can result in adverse patient reactions and contribute to antimicrobial drug resistance
- **FebriDx regulatory and commercial update**
 - 708-subject, multicentre clinical trial published in JAMA in 2022 — 98.7% NPV for bacterial infections
 - FebriDx cleared in other markets including Europe, UK, Brazil, Australia and other markets
 - Commercial rollout in cleared markets impacted by pandemic – European orders received late 2022
 - Actively developing sales and partnering opportunities for FebriDx in cleared or pending markets
 - Clearance to market FebriDx in the US awarded in July 2023
 - Henry Schein now distributing FebriDx in UK, Spain, Portugal and the Netherlands

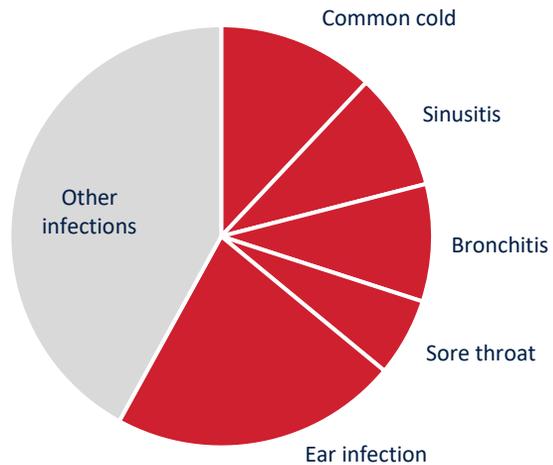


¹ Outpatient Antibiotic Prescriptions—United States 2021: <https://www.cdc.gov/antibiotic-use/data/report-2021.html>

² Tse, J.; Near, A.M. et al; Antibiotics 2022, 11, 1058. <https://doi.org/10.3390/antibiotics11081058>

Current Antibiotic prescribing in the US

ANTIBIOTICS PRESCRIBED IN THE U.S. BY TYPE



Acute respiratory infections may account for **58%** of all antibiotics prescribed ⁴

ANTIBIOTICS PRESCRIBED



211M

antibiotic prescriptions issued in outpatient settings each year ¹

44%

of antibiotic prescriptions are written to treat patients with ARIs ²

40%

of these are unnecessary ³



¹ Outpatient Antibiotic Prescriptions—United States 2021: <https://www.cdc.gov/antibiotic-use/data/report-2021.html>

² Unnecessary Antibiotics for Acute Respiratory Tract Infections: Associations with Care Setting and Patient Demographics, 2016 Tse, J.; Near, A. *et al*; *Antibiotics* 2022, 11, 1058. <https://doi.org/10.3390/antibiotics11081>.

⁴ Centers for Disease Control and Prevention. *MMWR*, 2011, 60:1153-5

FebriDx – preparations for US launch well advanced



- **Scaling up production to meet additional anticipated demand**
 - Materials ordered for increased volume of production
 - Test strip optimisation currently underway
 - On track to have units available for initial orders expected in late CY2024
- **Commenced marketing activities to initial key customers**
 - Initial discussions with urgent care and relevant physician offices through Lumos' US sales channel
 - Identified CPT codes that are expected to provide reimbursement coverage for FebriDx
- **Looking at potential partnering opportunities:**
 - Large national distributors (Henry Schein, McKesson, Medline, Cardinal/Fisher)
 - Add menu to other diagnostic test platforms



ViraDx™ – Lumos’ POC test for key respiratory infections



- **ViraDx highly relevant POC test for post-pandemic environment:**
 - SARS-CoV-2 pandemic increased consumer and healthcare POC testing
 - ViraDx is a 3-in-1 test for COVID-19/flu A/flu B
 - One of two tests available in market that provides visual read-out
- **ViraDx regulatory and commercial update:**
 - US EUA authorisation awarded in September 2023
 - Additional studies may be required to transition for EUA to 510(k) clearance
 - Scale-up of production of ViraDx has commenced
 - Initial sales through Lumo’s US sales channel
 - First orders received with shipping expected to commence in Nov 2023



Lumos' direct US sales channel



- **Low-cost, targeted sales channel**
 - Targeting urgent care and relevant physician offices
 - Two internal FTEs (Head of Sales, Regional Manager)
 - Currently 7 commission-only reps — will increase over time
 - Two distributors signed, ten at final negotiation stage
- **Focus on infectious diseases and women's health**
 - Lumos-branded products: FebriDx and ViraDx
 - Secured US distribution rights for Binx IO point-of-care test
 - Molecular POC test for *chlamydia* and *gonorrhoeae*
 - Cartridge-based test with CLIA waiver
 - Currently evaluating other women's health and infectious disease products



Promising Outlook

“

We have come out of FY2023 a much stronger and more focused company with an exciting pipeline of opportunities in front of us.

I believe in that the foundations we established over the last year will will enable Lumos to accelerate the building and growth of its business.

Doug Ward
Chief Executive Officer
Lumos Diagnostics



Expand and diversify pipeline of commercial services projects and recurring revenue



Continue to build foundation for long-term growth by establishing strategic partnerships



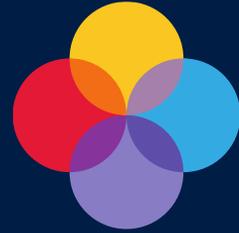
Launch FebriDx and ViraDx in US market using established sales channel



Improve operating cash flow through revenue growth and ongoing cost management



Strong underlying fundamentals from increasing use of rapid point-of-care diagnostic tests



LUMOS
DIAGNOSTICS

www.lumosdiagnostics.com