

PRESENTATION AT BIOTECH SHOWCASE INVESTOR CONFERENCE

Event held as part of annual J.P. Morgan Healthcare Conference in San Francisco, USA

Adelaide, Australia, 14 January 2020: LBT Innovations Limited (ASX: LBT) (LBT or the Company), a leader in medical technology automation using artificial intelligence, announces that the Company's CEO and Managing Director, Brent Barnes, has been selected to present at the prestigious Biotech Showcase event in San Francisco on Tuesday 14 January 2020.

The Biotech Showcase event, held as part of the annual J.P. Morgan Healthcare Conference, is one of the industry's leading investor conferences with presentations from over 400 private and public companies. Presentation at the event provides a great opportunity to build the profile of the Company with over 1,000 investors and potential partners participating from around the world.

The J.P. Morgan Healthcare Conference is the largest healthcare investment conference in the world, bringing together a broad range of companies spanning the entire healthcare landscape, including pharmaceutical firms, healthcare service providers and medical device companies. Running since 1983, the conference is held in San Francisco and will take place this year from the 13 - 16 January 2020.

The Company's presentation for the conference is attached.

- ENDS -

About LBT Innovations

LBT Innovations (LBT) improves patient outcomes by making healthcare more efficient. Based in Adelaide, South Australia, the Company has a history of developing world leading products in microbiology automation. Its first product, MicroStreak®, was a global first in the automation of the culture plate streaking process. The Company's second product, the Automated Plate Assessment System (APAS®) is being commercialised through LBT's 50% owned joint venture company Clever Culture Systems AG (CCS) with Hettich Holding Beteiligungs- und Verwaltungs-GmbH. The APAS® Independence is based upon LBT's intelligent imaging and machine learning software and remains the only US FDA-cleared artificial intelligence technology for automated imaging, analysis and interpretation of culture plates following incubation. The Company has regulatory cleared analysis modules able to support microbiology laboratories with the automated analysis of urine culture plates and for MRSA infection control.

CONTACTS

LBT Innovations	Investor Enquiries
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Disruptive Al technology for healthcare Ready for launch

January 2020

ASX code: LBT Ibtinnovations.com

Disclaimer

This document contains certain forward-looking statements that involve risks and uncertainties. Although we believe that the expectations reflected in the forward-looking statements are reasonable at this time, we can give no assurance that these expectations will prove to be correct.

Given these uncertainties, readers are cautioned not to place undue reliance on any forward-looking statements. Actual results could differ materially from those anticipated in these forward-looking statements due to many important factors, risk and uncertainties including, without limitation, risks associated with estimating potential quantity and timing of sales, risks associated with medical device development and manufacture, risks inherent in the extensive regulatory approval processes mandated by regulatory authorities, delays in clinical trials, future capital needs, general economic uncertainly and other risks detailed from time to time in the Company's announcements to the ASX.

Moreover, there can be no assurance that others will not independently develop similar products or processes or design around patents owned or licensed by the Company, or that patents owned or licensed by the Company will provide meaningful protection or competitive advantages.

All reasonable efforts have been made to provide accurate information, but the Company does not undertake any obligation to release publicly any revisions to any "forward-looking statement" to reflect events or circumstances after the date of this presentation, except as may be required under applicable laws. Recipients should make their own enquiries in relation to any investment decisions from a licensed investment advisor.



Agenda

- 1. Global Market Opportunity
- 2. APAS® Value Proposition
- 3. Future Outlook & Conclusions



Disruptive AI technology – commercial ready

APAS® Independence - First and only automated culture plate reader



Patent protected



Global regulatory clearances



Innovative platform technology



Product launched - early sales







Challenges for microbiology laboratories

Around the world scientists still read plates **manually**

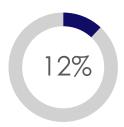
2.4 billion plates per annum and growing





Challenges for microbiology laboratories

Challenges



Inconsistent results

Variability in plate reading Error rates up to 12%



Workplace safety issues

Strain injuries caused by repetitive manual processes
Staff sick days and annual leave

Market Need



High demand

70% of clinical decisions based on *in vitro* diagnostic results



Inefficient use of resources

Up to 85% of plates read are negative



Qualified Microbiologists

9% vacancy rate in the US

Microbiologist average age is high (51 AUS / 42 US)



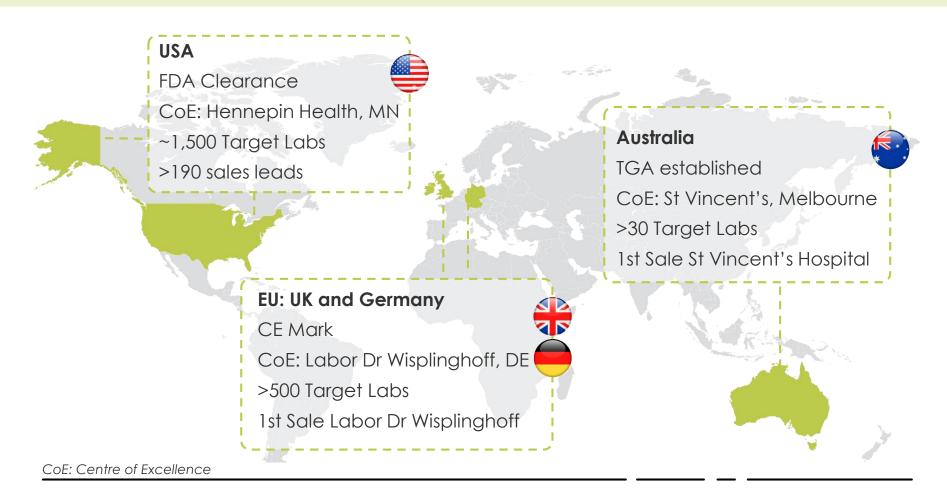
Large addressable market

Compelling value proposition for medium and large laboratories

Number of labs globally **Small Labs** 15,000 <400 plates per day **Medium Labs** 12,000 400 - 1000 plates per day **Delivering payback** in 2-3 years **Large Labs** 1,000 >1000 plates per day **Total Market Opportunity** 13,000

Initial launch markets

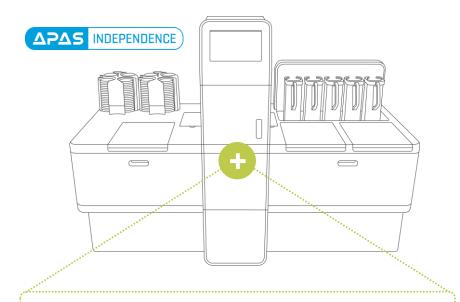
Current launch markets - 2,000 laboratories meeting APAS® target profile





APAS® Independence

The first and only automated culture plate reader - Powered by Al



APAS® analysis modules
Increases number of specimen
types that can be tested
Increases customer utility





in a 10,000-patient clinical trial



3x faster

than manual culture plate reading



Available

regulatory cleared in US, EU, AU



Attractive return on investment

payback between 2 - 3 years



Competitive landscape

Competitors offer large and expensive process automation

- **DO NOT** read plates Automate lab workflow
- **HIGH** capital cost ~USD 2.5m+ cost
- **LARGE** footprint Complex installation
- **LONG** integration period Plates still read manually
- **LOW** penetration ... 150 units in 10 years



Copan WASPLab™

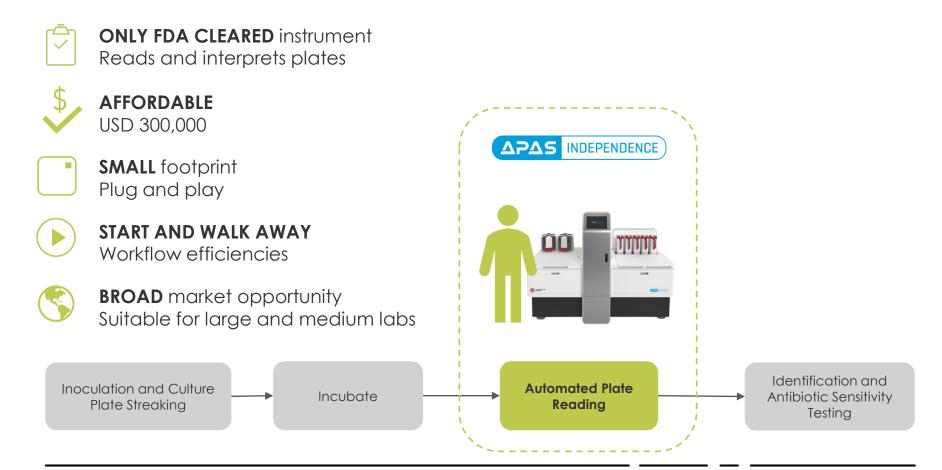


BD Kiestra™ Total Lab Automation



First mover advantage - clinical plate reading

APAS® Independence uses AI for advanced decision making





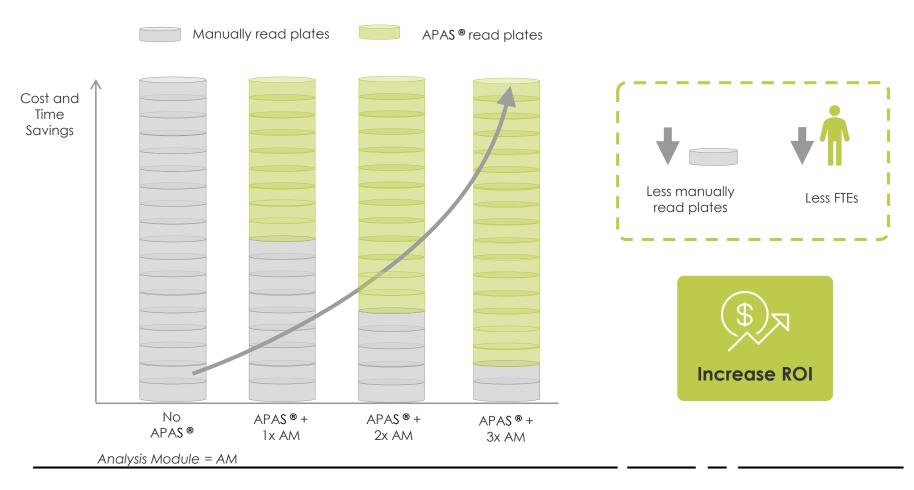
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Building the value proposition

Multiple analysis modules provide increased value for **all** laboratories





Product roadmap - analysis module development

Increasing market opportunity

More analysis modules > More laboratory tests > Increased ROI for labs



APAS® ready for launch

2010 →

2018

2019

2020+

Early technology development

Feasibility completion

Regulatory clearances Establish KOLs

Market awareness

Market launch Early sales

Build sales pipeline

Distributor appointment

Market penetration – US, AU, DE, UK

New market expansion

AM library development









Sales model & market opportunity

Dual strategy of direct to market and distributor appointment

50:50 Joint Venture

Profits shared equally



1. CAPEX: Instrument ~USD 300k

Flexible leasing models and SaaS established



2. Annual Licenses:Analysis Module~USD 20k – 40k



5 year revenue opportunity ~USD 450k per instrument

Launch markets











Market penetration	Instrument*	Analysis Module
20%	~USD 80 - 120 million	~USD 12 million p.a.
30%	~USD 120 - 180 million	~USD 18 million p.a.

^{*} Sales estimate includes expected distributor margin



Comparable products case study

Large market opportunity for laboratory automation

Case Study	Launch	Sales <u>Potential</u>
APAS Independence	2018	Launched with FDA Clearance 1,500 units after 10 years (and still growing) Cumulative instrument sales: ~USD 400 million 50% flows to LBT (after distributor fees & JV costs) Licence fees, building to: ~USD 40 million per annum Majority flows to LBT (after distributor fees)
Comparison Products	Launch	Sales <u>Achieved</u>
MALDI-TOF 2 Manufacturers: Bruker & bioMérieux	2004	~800 units placed after 8 years FDA Clearance after ~9 years 700 units placed in US following FDA Clearance & years of clinical data ~1,500 units placed after 10 years ~12% global penetration
Previ Isola	2009	~500 units placed after 5 years ~4% global penetration bioMérieux terminated agreement with LBT in 2015 - Product no longer sold



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Business and commercialisation strategy

Repeatable process: new opportunities - underpinned by core capability



Developed internal capability

- Technology development
- In-sourced core capability
 - Artificial Intelligence
 - Software Engineering
 - Regulatory Expertise
 - Clinical Science



Appoint distributors & build sales

- Routine clinical use in their market
- Demonstrate customer engagement
- Initial sales pipeline established
- Regulatory cleared product



Outsource non-core activities

- Focus on core capability
- Built trusted relationships with contractors
 - Design engineering services
 - Manufacturing



Established global platform

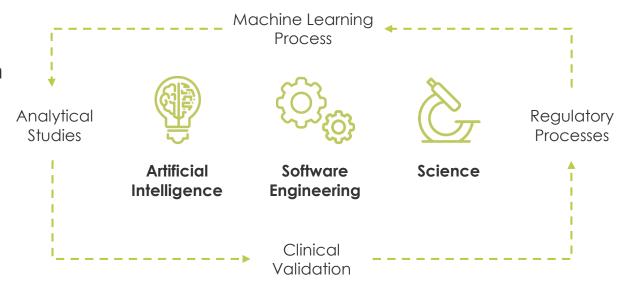
- Regional KOLs
- Regulatory clearances (TGA, FDA and CE Mark)
- Product available for sale
- Delivered early sales



Strategic growth

Opportunity to extend **platform capability** into new applications

- Digital Pathology
- Laboratory Automation
- Wound Care
- Digital Health
- Medical Imaging
- Telehealth





Board and leadership team



Brent BarnesCEO and MD, AU

- 11 years senior executive at Cochlear Ltd (ASX: COH)
- Experience across Class I, II and III medical devices
- Held leadership roles in the US and Asia-Pacific
- Non-Executive Director Connek Ltd



Kate Costello Chairman, AU

- Expertise in legal, corporate governance and company strategy
- Diverse Board experience across multiple industries
- Established leading governance consultancy
- Fellow of the Institute of Company Directors



Damian Lismore NED, AU



Simon Arkell NED, US



Caroline PopperNED, US



Ray Ridge CFO, AU



International board



Healthcare and technology



Public listed company experience (ASX and Nasdaq)



Compliance and financing expertise



Experienced management team



Peter BradleyVP Global Business
Development



Rhys Hill
Research Director
APAS® Inventor



Julie WinsonDirector Quality and
Regulatory Affairs



Steve GiglioScientific Director



Artificial intelligence and software engineering



Medical device product development



Quality and regulatory affairs



Early product commercialisation



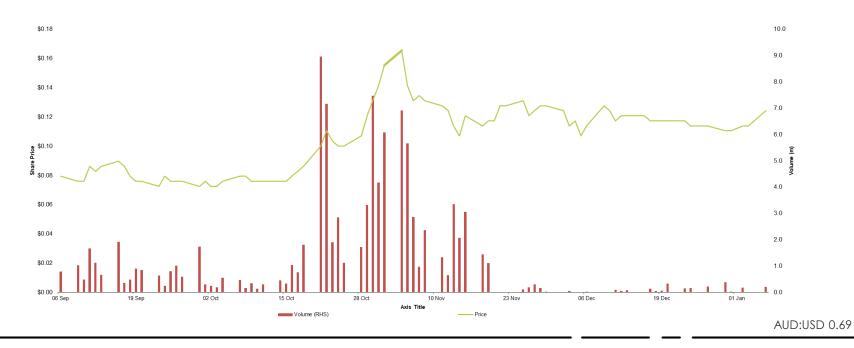
Microbiologist expertise built into APAS ®



Corporate overview

Key Statistics (Closing on 06 January 2020)		
Share Price	\$0.12 per share	
12-month range	\$0.041 - \$0.17	
Number of shares	235.5 million	
Options Issued	6.1 million	
Market Cap	\$29.3 million	

Financials (USD)	
Current Cash	\$6.0 million (30 September 2019)
SAFA Loan Facility	\$2.8 million facility – \$1.7m drawn down Low interest rate, 5-year term
Enterprise value	\$23.3 million
Shareholders	Insto (8%), Industry (7%), Dir + Mgmt (4%)





Investment highlights

Competitive positioning strong

APAS® Independence – the **only FDA and CE cleared** product uses **artificial intelligence** for microbiology plate reading

Clinically proven through 10,000-patient global clinical trial

Large value proposition

Initial launch market - **2,000 labs** meet APAS® target profile

5-year revenue opportunity ~USD 450,000 per instrument

Platform for growth

Closing cash balance of AUD 8.67m (30th September 2019)

Advanced technology + internal capability supports **future opportunities**

Ready for launch

Early sales achieved in Australia and Germany

Regulatory clearances - available to sell in US, EU, AU

Appointment of distributors for market penetration





LBT Innovations

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