



## Immuron CEO, Steven Lydeamore presentation at Emerging Growth Conference

Melbourne, Australia, January 23, 2026: Immuron Limited (ASX: IMC; NASDAQ: IMRN) is pleased to advise our Chief Executive Officer, Steven Lydeamore presented virtually at the Emerging Growth Conference on Thursday 22<sup>nd</sup> January 2026 (2:40pm - 2:50pm U.S. Eastern Time).

A copy of the presentation made is included below.

This release has been authorised by the directors of Immuron Limited.

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### COMPANY CONTACT:

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### About Immuron

Immuron Limited (ASX: IMC, NASDAQ: IMRN), is an Australian biopharmaceutical company focused on developing and commercializing orally delivered targeted polyclonal antibodies for the treatment of infectious diseases.

### About Travelan®

Travelan® is an orally administered passive immunotherapy that prophylactically reduces the likelihood of contracting travelers' diarrhea, a digestive tract disorder that is commonly caused by pathogenic bacteria and the toxins they produce. Travelan® is a highly purified tabletized preparation of hyper immune bovine antibodies and other factors, which when taken with meals bind to diarrhea-causing bacteria and prevent colonization and the pathology associated with travelers' diarrhea. In Australia, Travelan® is a listed medicine on the Australian Register for Therapeutic Goods (AUST L 106709) and is indicated to reduce the risk of Travelers' Diarrhea, reduce the risk of minor gastro-intestinal disorders and is antimicrobial. In Canada, Travelan® is a licensed natural health product (NPN 80046016) and is indicated to reduce the risk of Travelers' Diarrhea. In the U.S., Travelan® is sold as a dietary supplement for digestive tract protection.

### Travelers' diarrhea (TD)

TD is generally defined as the passage of  $\geq 3$  unformed stools per 24 hours plus at least one additional symptom (such as nausea, vomiting, abdominal cramps, fever, blood/mucus in the stools, or fecal urgency) that develop while abroad or within 10 days of returning from any resource-limited destinations ([Leung et al., 2006](#)). Diarrhea continues to be the most frequent health problem among travelers to destinations in lower- and middle-income regions ([Steffen, 2017](#)). Deployed US military personnel, essentially representing a long-term traveller population, are particularly affected given their population dynamics and the context in which they seek care and treatment ([Connor et al., 2012](#)). Diarrhea is the leading infectious disease threat to the overall health and preparedness of deployed US armed forces, with diarrheagenic E. coli, Campylobacter spp., and Shigella spp. among the most commonly reported etiologies ([Riddle et al., 2006](#)).

### Immuron Platform Technology

Immuron's proprietary technology is based on polyclonal immunoglobulins (IgG) derived from engineered hyper-immune bovine colostrum. Immuron has the capability of producing highly specific immunoglobulins to any enteric pathogen and our products



are orally active. Bovine IgG can withstand the acidic environment of the stomach and is resistant to proteolysis by the digestive enzymes found in the Gastrointestinal (GI) tract. Bovine IgG also possesses this unique ability to remain active in the human GI tract delivering its full benefits directly to the bacteria found there. The underlying nature of Immuron's platform technology enables the development of medicines across a large range of infectious diseases. The platform can be used to block viruses or bacteria at mucosal surfaces such as the Gastrointestinal tract and neutralize the toxins they produce.

#### **IMM-124E (Travelan®)**

IMM-124E was developed using Immuron's platform technology. IMM-124E is produced from the colostrum of birthing cattle that have been immunised during pregnancy with a vaccine containing the outer antigens of multiple human derived ETEC. A total of 13 ETEC strains are used in the vaccine to produce high levels of antibodies against selected surface antigens from the most common strains of ETEC. ([Otto et al., 2011](#))

The resultant hyperimmune colostrum IMM-124E from ETEC vaccinated cows contains significant levels of polyclonal antibodies specific for ETEC antigens LPS, CFA-I and Flagellin ([Sears et al., 2017](#)).

The antibodies produced in IMM-124E have been found to have a stronger binding and neutralizing activity (than the antibodies of unvaccinated cattle) against a wide range of LPS antigens including both the variable O-polysaccharide region and the preserved oligosaccharide core 'R' region of LPS from the 13 serotypes used in the ETEC vaccine.

IMM-124E is manufactured into a tablet form referred to as Travelan®.

#### **IMM-529**

Immuron is developing IMM-529 as an adjunctive therapy in combination with standard of care antibiotics for the prevention and/or treatment of recurrent *Clostridioides difficile* infection (CDI). IMM-529 antibodies targeting *Clostridioides difficile* (C. diff) may help to clear CDI infection and promote a quicker re-establishment of normal gut flora, providing an attractive oral preventative for recurrent CDI.

Immuron is collaborating with Dr. Dena Lyras and her team at Monash University, Australia to develop vaccines to produce bovine colostrum-derived antibodies. Dairy cows were immunised to generate hyperimmune bovine colostrum (HBC) that contains antibodies targeting three essential C. diff virulence components. IMM-529 targets Toxin B (TcB), the spores and the surface layer proteins of the vegetative cells.

This unique 3-target approach has yielded promising results in pre-clinical infection and relapse models, including (1) Prevention of primary disease (80% P = 0.0052); (2) Protection of disease recurrence (67%, P < 0.01) and (3) Treatment of primary disease (78.6%, P < 0.0001; TcB HBC). Importantly IMM-529 antibodies cross-react with whole cell lysates of many different human strains of C. diff including hypervirulent strains.

To our knowledge, IMM-529 is, to date, the only investigational drug that has shown therapeutic potential in all three phases of the disease ([Hutton et al., 2017](#)).

#### **ProIBS®**

Immuron has an exclusive distribution agreement with Calmino group AB for the territories of Australia and New Zealand for ProIBS®. ProIBS® - to help patients treat IBS symptoms ProIBS® is a certified medical device for the treatment of IBS symptoms such as abdominal pain, bloating and unsettled bowel movements (diarrhoea and/or constipation). ProIBS® contains AVH200®, derived from the plant *Aloe barbadensis* Mill. AVH200® has gel forming components which support the intestinal mucosal barrier. As IBS is known to affect individuals for a long period of time, it is essential to have a treatment appropriate for long-term use – as ProIBS® is. The product is safe, and no interactions with other medications are known. Science-driven innovative Calmino group AB, the developer of ProIBS®, conducted a usability study among 1,003 users. ProIBS® was helpful for 94% of them. 91% of the users experienced an improvement in daily life and 98% would recommend ProIBS® to someone else. To learn more please check: [www.proibs.eu](http://www.proibs.eu).

Irritable bowel syndrome (IBS) is a common condition where you experience symptoms related to your digestive system. This is sometimes linked to certain foods, lifestyle habits and stress levels or mood. IBS affects around 3 out of every 10 people. Females are more likely than males to be affected. Some key symptoms of IBS include: abdominal pain or discomfort; stomach bloating and wind; chronic diarrhoea or constipation, or alternating between the two. ([healthdirect.gov.au](http://healthdirect.gov.au)) According to available data, the IBS treatment market in Australia is estimated to be a part of the broader "Digestives & Intestinal Remedies" market, generating a revenue of around AU\$221.14 million in 2025, with a projected annual growth rate of 3.28%. ([Statista](https://www.statista.com/statistics/1101111/irritable-bowel-syndrome-market/))

## References

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For more information visit: <https://www.immuron.com.au/> and <https://www.travelan.com>

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This press release may contain "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934, each as amended. Such statements include, but are not limited to, any statements relating to our growth strategy and product development programs and any other statements that are not historical facts. Forward-looking statements are based on management's current expectations and are subject to risks and uncertainties that could negatively affect our business, operating results, financial condition, and stock value. Factors that could cause actual results to differ materially from those currently anticipated include: risks relating to our growth strategy; our ability to obtain, perform under and maintain financing and strategic agreements and relationships; risks relating to the results of research and development activities; risks relating to the timing of starting and completing clinical trials; uncertainties relating to preclinical and clinical testing; our dependence on third-party suppliers; our ability to attract, integrate and retain key personnel; the early stage of products under development; our need for substantial additional funds; government regulation; patent and intellectual property matters; competition; as well as other risks described in our SEC filings. We expressly disclaim any obligation or undertaking to release publicly any updates or revisions to any forward-looking statements contained herein to reflect any change in our expectations or any changes in events, conditions, or circumstances on which any such statement is based, except as required by law.

# EMERGING GROWTH CONFERENCE

23 JANUARY 2026

NASDAQ: IMRN  
ASX: IMC





# SAFE HARBOR STATEMENT

Certain statements made in this presentation are forward-looking statements and are based on Immuron's current expectations, estimates and projections. Words such as "anticipates," "expects," "intends," "plans," "believes," "seeks," "estimates," "guidance" and similar expressions are intended to identify forward-looking statements.

Although Immuron believes the forward-looking statements are based on reasonable assumptions, they are subject to certain risks and uncertainties, some of which are beyond Immuron's control, including those risks or uncertainties inherent in the process of both developing and commercializing technology. As a result, actual results could materially differ from those expressed or forecasted in the forward-looking statements.

The forward-looking statements made in this presentation relate only to events as of the date on which the statements are made. Immuron will not undertake any obligation to release publicly any revisions or updates to these forward-looking statements to reflect events, circumstances or unanticipated events occurring after the date of this presentation except as required by law or by any appropriate regulatory authority.

FY2026 results in this presentation are subject to audit review.





Immuron Ltd is an Australian integrated biopharmaceutical company with global scale, focused on developing, and commercialising, oral products for the treatment of gut mediated diseases

### Technology Platform

Safe and potentially transformational approach to gut infections

### Research & Development

3 pipeline assets in 2 clinical programmes

### Global footprint

Australia, US, Canada and expanding

### Corporate Research

[Independent Buy recommendation](#)

### Technology Platform

- Proprietary technology based on generating hyper-immune bovine colostrum that is rich in polyclonal antibodies also known as immunoglobulins.
- Bovine antibodies also possess the unique ability to remain active in the human GI tract where they target enteric pathogens.
- The underlying nature of our platform technology enables the development of medicines across a large range of infectious diseases.
- The platform can be used to generate bovine antibodies that block viruses or bacteria at mucosal surfaces in the Gastrointestinal tract and neutralize the toxins they produce.

### Our Products

## Travelan®



- Reduce the risk of Traveller's Diarrhoea
- Sold in pharmacies Australia-wide
- Available in Australia, USA and Canada

NEW

## PROIBS®



- Treatment of symptoms associated with Irritable Bowel Syndrome (IBS)
- To be [sold](#) in pharmacies Australia-wide
- Available in Australia

# FY25 Financial Achievements

**Global Sales  
Revenue**

**\$7.3m**

 **Up 49%**

**Australian Sales  
Revenue**

**\$5.3m**

 **Up 40%**

**North American  
Sales Revenue**

**\$2.0m**


 **Up 76%**

**Gross Profit  
Margin**

**65.4%**

**EBITDA (ex-R&D)<sup>1</sup>**

**\$(3.1)m**

 **\$2.1m  
improvement**

**Cash + Term  
Deposit<sup>2,3</sup>**

**\$5.9m**

<sup>1</sup> ex-R&D: add back research & development, less R&D Tax Incentive and R&D grants;

<sup>2</sup> 30 June 2025, as reported in Immuron's Appendix 4E (Cash \$2.83m; Term Deposit \$3.04m);

<sup>3</sup> Since 30 June 2025, Immuron has raised, cost-effectively, A\$7.15m from the US At-The-Market facility.

# Travelan® continued strong sales growth



## Global

+ FY2025 AUD\$7.3 million **up 49%** on prior year  
+ FY2026 H1 AUD\$4.2 million **up 5%** on FY2025 H1



## Australia

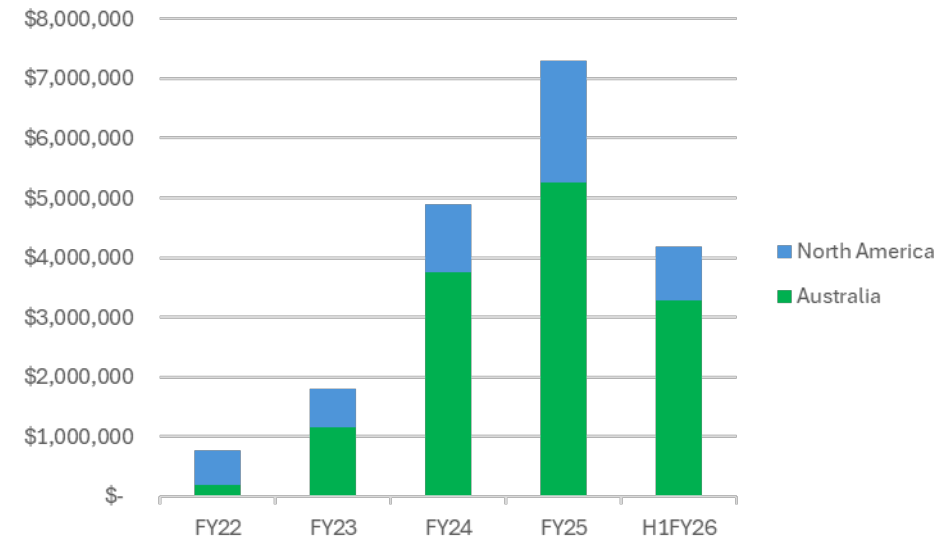
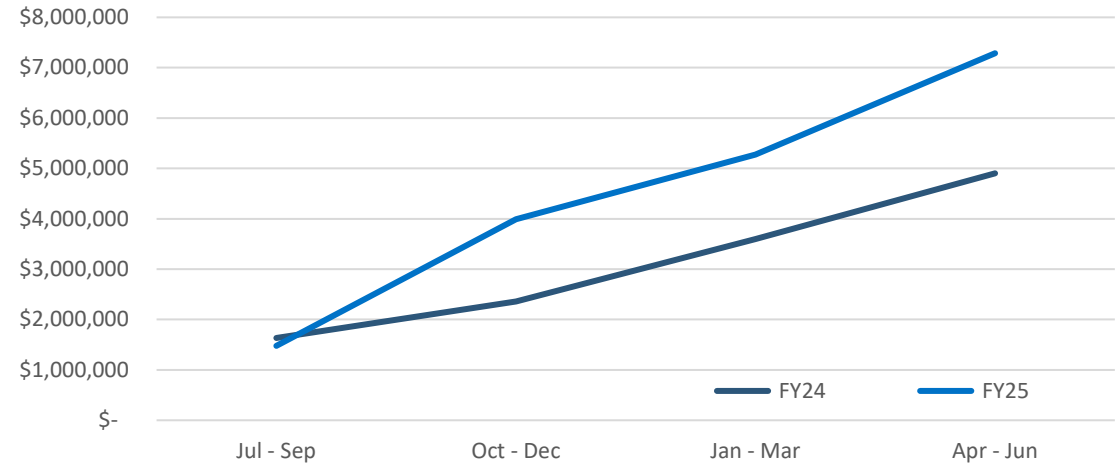
+ FY2025 AUD\$5.2 million **up 40%** on prior year  
+ FY2026 H1 AUD\$3.3 million **up 13%** on FY2025 H1



## North America

+ FY2025 AUD\$2.0 million **up 76%** on prior year  
+ FY2026 H1 AUD\$0.9 million **down 18%** on FY2025 H1  
+ USA FY2026 H1 AUD\$0.9 million **up 17%** on FY2026 H1

Global Year to Date Net Sales (\$AUD)





# FY25 Clinical Achievements and Operational Highlights

FDA approval  
IMM-529  
IND



Travelan®  
n=851 trial  
completed



Travelan® Ph 2  
statistically  
significant  
results<sup>1</sup>



Updated peak  
U.S. sales for  
IMM-529  
US\$400 million<sup>2</sup>

New research  
collaboration  
IMM-986 (VRE)



New product  
launch in  
Australia



1. Immunology results and Microbiome responses; 2. [Lumanity](#) global healthcare consulting

# FY26 Outlook

✓ Continued YoY  
sales growth

Progression  
towards EBITDA  
(ex-R&D)<sup>1</sup>  
breakeven

Travelan®  
End of Phase 2  
Meeting with FDA  
1H 2026

IMM-986 (VRE)  
Pre-clinical data  
1H 2026

IMM-529  
Initiation of Phase 2  
1H 2026

<sup>1</sup> ex-R&D: add back research & development, less R&D Tax Incentive and R&D grants



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