



INVESTOR PRESENTATION

COFFEE MICROCAPS

2 MAY, 2023

Steven Lydeamore - CEO

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.

YTD FY2023 results in this presentation are subject to audit review.



EXECUTIVE SUMMARY



Immuron Ltd (ASX:IMC) (NASDAQ:IMRN) is a globally integrated biopharmaceutical company focused on developing, and commercialising, oral immunotherapeutics for the treatment of gut mediated diseases

Company Overview

- Two commercially available oral immunotherapeutic products Travelan® and Protectyn®
- 4 clinical programs Travelan® (IMM-124E), CampETEC, IMM-529, Travelan® (consumer product)
- Market capitalisation of A\$18.67m as of 20 April 2023 with cash & cash equivalates balance of A\$18.5m as of 31 Dec 2022

Recent Business Updates

- Travelan® (IMM-124E) IND filed and received FDA Authorization to proceed under IND 029087a
- CampETEC toxicology study report completed and submitted to FDA
- Travelan® Uniformed Health Services University P2TD IMM-124E field clinical trial recruitment reaches 300
 - 35% of target 868 participants recruited
- Immuron's own shopfront on amazon.com for Travelan® set up for USA market
 - Launch pending establishing necessary safety stock level
- Working towards relaunch of Travelan® into Canadian retail pharmacy in FY24
 - Submitted for range review to major pharmacy chain

Results & Outlook

- FY23 YTD 14 April 2023 revenue of A\$1.78m, up 233% YTD on pcp Australian sales contributing A\$1.18m
- Evaluating options to enter Asian and European markets through distributors
- Evaluating options to add to marketed products portfolio in FY24



DISTRIBUTION CAPABILITY

	Australia	USA	Canada
Retail Pharmacy	©		©
B2B	©	©	
E-commerce	©	©	©





Key Commentary

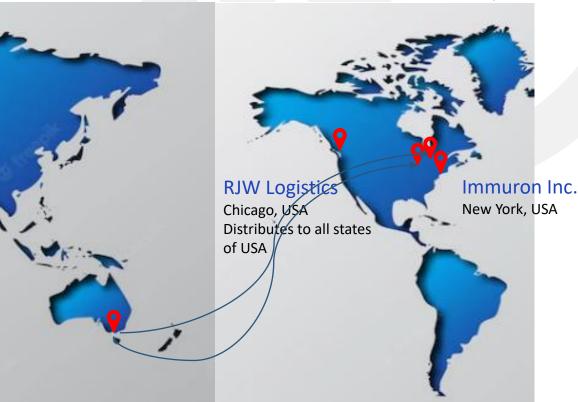
- The Australian retail network includes over 3,500 pharmacies
- In the USA, the key B2B customer is Passport Health the largest network of travel medicine clinics
- Immuron's own shopfront on amazon.com will be introduced in the USA during FY23
- Re-entry into retail pharmacies in USA will be explored in FY24
- In Canada, we are working towards a relaunch into retail pharmacy in FY24
- We are exploring options to expand B2B business in airlines, cruise ships, health & wellness segments
- We are evaluating options to enter Asian and European markets through distributors

Immuron Limited

Vancouver, BC, Canada

ANB Canada

Ontario, Canada Distributes to all provinces of Canada



Immuron Limited

Melbourne, Australia
Distributes to all states of Australia and to distributors in USA
and Canada



ADDRESSABLE MARKET & INDUSTRY OVERVIEW

~\$15b+

Immuron's products are a subset of the global digestive health market, which a multi-billion-dollar market*

~7% CAGR

Travelers diarrhea treatment market is large and growing at a CAGR of ~7% over 2019-2022*



Travelan® has large market potential given that acute diarrhea affects millions of travelers each year

\$83m

Based on US annual travel numbers and a penetration rate of 15%, the market potential is estimated at \$83m**

\$50m

Based on EU travel numbers and a penetration rate of 15%, the market potential is estimated at \$50m**

\$1.7b

Clostridioides difficile infections (CDIs) to grow to almost \$1.7 billion by 2026, according to GlobalData



^{*} IQVIA Consumer Health Category QuickView MAT Q1 2019

^{**} IMC Company Report - Travelan Market Analysis 2019





Billion Dollar Market

Travelers diarrhea treatment market is large and growing at a CAGR of ~7%



Industry tailwinds

Travel picking up significantly following COVID lockdowns



Frequent Symptom

30% - 70% of travelers experience traveler's diarrhea***

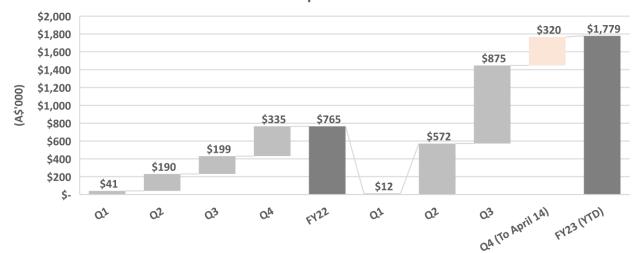




SALES | YEAR TO DATE | 14TH APRIL 2023

A\$000s			
Profit & Loss	FY23 YTD 14 April 2023	FY22YTD 14 April 2022	Var %
Australian Sales	\$ 1,184 \$	137	764%
Rest of World Sales	\$ 595 \$	397	50%
Total Revenue	\$ 1,779 \$	534	233%

Net Sales | FY22 & FY23



*YTD FY2023 results in this presentation are subject to audit review.

Immur@n

Key Commentary

- Strong start to Q4 2023 (to 14 April 2023)
 - \$320k Net Sales; up 233% YTD on pcp
- Supply chain capacity addressed by contracting another packaging supplier
- Strong sales for Travelan®, well above pre-pandemic levels, may results in short term stock outages in some wholesalers and pharmacies over the next month
- Building stock to allow accelerated growth in international markets



BUSINESS POSITIONED FOR ORGANIC GROWTH AND NEW M&A STRATEGY

Organic Growth Strategy

 Focus on commercialised products and nearterm development extensions, including:

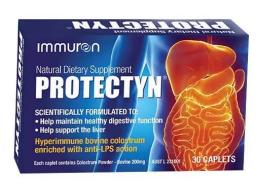
Travelan®:

- Sales expansion across target geographies
- Growth in distribution network and sales
 & marketing initiatives
- Product development (new formulations including once daily dosing) e.g. FDA approval

2 Protectyn®:

- Sales expansion across target geographies
- Growth in distribution network and sales
 & marketing initiatives
- Product development and broader applications





M&A Strategy

- By pursuing growth through M&A of a fragmented market, IMC believes that it will be able to increase market geographies, sales channels and penetration driving revenue growth and ultimately shareholder value
- Our M&A Key Criteria focusses on:
 - Expand market verticals & product offering
 - Expand existing customer base
 - Cost & Earnings Synergies
 - Strength of IP and Management
 - Distribution network and sales & marketing by each product





TECHNOLOGY PLATFORM

Immuron's proprietary technology platform combines the natural human nutrition & health benefits of bovine colostrum with a novel class of specifically targeted oral polyclonal antibodies that offer delivery within the gastrointestinal ("GI") tract and can be used to target viruses or bacteria and neutralize the toxins they produce at mucosal surfaces.



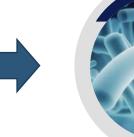
STEP 1 Development of Highly

Specific Vaccines



STEP₂

Isolation of Hyperimmune antibody-rich bovine colostrum



STEP 3

Oral Antimicrobial therapeutics without drawbacks of antibiotics





Toxin Neutralization + Clearance of targeted gut pathogens

- Reduce occurrence and reduce/relieve diarrhoea
- Reduce/relieve abdominal cramping
- Reduce/relieve gastrointestinal pain
- Assists repair of gastrointestinal/gut wall lining
- Enhance/promote immune defence
- Enhance/promote health liver function



Australian Permitted indications; these statements have not been evaluated by the Food and Drug Administration (FDA)



STATUS OF PRODUCT PORTFOLIO AND KEY MILESTONES

- US Department of Defense grant of US\$4.45 million to examine a dosing regimen for Travelan® more suited for use by the military b
 - Market evaluation by <u>Lumanity</u> confirms the Traveler's Diarrhea market opportunity for IMM-124E (Travelan®)^c
 - o IMM-124E (Travelan®) IND was approved by the FDA in 2H2022 a
 - Clinical trials anticipated to commence in 1H2023
- Market evaluation by <u>Lumanity</u> confirms the *Clostridioides difficile* market opportunity for IMM-529 d
 - o 600mg solid dose active formulation development completed
 - Project strategy and budget to be approved by the Immuron Board before initiating clinical trials
- Travelan® Uniformed Services University has recruited more than 35% of participants in a randomized clinical trial with Travelan® to evaluate the effectiveness for prophylaxis during deployment or travel to a high traveler's diarrhea risk region
 - Estimated Study Primary Completion Date 30 July 2023
 - Estimated Study Completion Date 31 December 2023
- Naval Medical Research Center Clinical Trials of CampETEC in campylobacter and enterotoxigenic E.coli (ETEC)^f
 - Toxicology study submitted to FDA 8 March 2023
 - Written Guidance received from FDA 10 March 2023
 - Response letter submitted 5 April 2023
 - Anticipate FDA feedback in May 2023

Immuron's Clinical Programs

Compound or brand name	Indication	Phase I	Phase II	Phase III	Market
IMM-124E Travelan®	Traveler's Diarrhea ETEC challenge	IMMU	ron		
IMM-529	Clostridioides difficile Infection & Recurrence	ımmu	ron		

Our Partners' Clinical Programs

Compound or brand name	Partner	Phase I	Phase II	Phase III	Market
Travelan®	THE PROPERTY OF THE PROPERTY O	Uniformed Se Universit			
CampETEC	AND THE PARTY OF T	Naval Med Research Ce			

^{*}Further information on the clinical programs can be found on slide 11





IMMURON'S CLINICAL PROGRAMS – OPPORTUNITY ASSESSMENT

Lumanity* Opportunity Assessment for IMM-124E b

- Immuron's development of IMM-124E (hyperimmune bovine colostrum) as a prescription medication has the potential to address this unmet need
- Primary care physicians (PCP)s impressed with clinical efficacy endpoint targets demonstrating > 80% protection against the development of diarrhea.
- If base case efficacy targets are reached, IMM-124E would mostly be used by travelers going to the highest risk areas (e.g., rural Central America/Asia/Africa).
- Based on the estimated market size and pricing, the base case yearly revenue in USA for IMM-124E is projected at US\$102M.
- Reaching higher efficacy goals could broaden use.

Lumanity Opportunity Assessment for IMM-529 c

- Infectious disease experts reacted favorably to the IMM-529 MOA, and its unique ability to target three elements of the rCDI infection – the spores, vegetative cells, and Toxin B
- Non-microbiome approaches (such as IMM-529) are still appealing to experts, who noted that clinical trial efficacy (reduction in relapse rate) and cost/access will be the key drivers of clinical use in recurrent patients, not mechanism of action
- Based on the estimated market size, anticipated payer restrictions, pricing, and competition, base case yearly revenue in USA for IMM-529 is conservatively projected at \$93M for the target patient population (limited to 2nd recurrence and later based on trial design and payer coverage)
- Positioning IMM-529 at the point of second relapse and/or efficacy targets could lead to higher uptake.

Compound or brand name	Indication	Phase I	Phase II	Phase III	Market
IMM-124E - Travelan®	Traveler's Diarrhea ETEC challenge	IMMUron			
IMM-529	Clostridioides difficile Infection & Recurrence	IMMUron			



KEY MILESTONES ANTICIPATED TO DRIVE VALUE



		2H 2022		1H 2023		2H 2023		1H 2024
Travelan®	•	FDA IND ¹ approved for single daily dose IMM-124E ETEC ² CHIM ³ clinical trial ^a	•	IRB Approval ⁴ Initiate IMM-124E ETEC CHIM clinical trial	•	Topline results for IMM- 124E ETEC CHIM clinical trial	•	Clinical Study Report
CampETEC	•	Submitted Response Letter to FDA Clinical Hold ^f Immuron sponsored Toxicology study - completed	•	Toxicology Study Report FDA feedback to clinical hold response letter	•	Initiate NMRC ⁵ CampETEC ETEC CHIM clinical trial Initiate NMRC CampETEC Campylobacter CHIM clinical trial	•	Topline results for NMRC CampETEC ETEC CHIM clinical trial Topline results for NMRC CampETEC Campylobacter CHIM clinical trial
IMM-529	•	600 mg solid dose active formulation development	•	Project strategic planning and budget development	•	IMM-529 cGMP manufacture ⁶ IMM-529 IND Pre-IND submission ⁶	•	IMM-529 IND submission ⁶ Initiate IMM-529 phase 2 CDI clinical trial ⁶
Travelan®	•	USU ⁷ P2TD IMM-124E field clinical trial recruitment commencement ^e	•	35% of 868 participants recruited	•	Primary Study Completion	•	Study Completion







Steven Lydeamore Chief Executive Officer Immuron Limited 2 May 2023

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CORPORATE SNAPSHOT



Shareholder Base:

- Large retail shareholder base in US and Australia
- BNY Mellon Asset Management (American Depository Shares) holds 34% stake
- Board and management aligned with shareholders, holding 3% stake

Financial Snapshot	
Shares on Issue	227,798,346
Total Options	12,879,720
Last Traded Price	\$0.082
52 week High/Low	0.125/0.067
Market Cap (20 Apr 23)	\$18.67m
Cash & Cash Equivalents (as at 31 Dec 22)	\$18.5m

Substantial Shareholders*					
Holder	Units	% of CSO			
BNY Mellon Asset Management	77,903,505	34.2 %			
Management & Board	6,904,554	3.0 %			
Authentics Australia Pty. Ltd.	6,000,000	2.6 %			
Grandlodge	3,846,712	1.7 %			

^{*}As at 21 April 2023



SCIENTIFIC REFERENCES

Travelan® (IMM-124E)	
Travelan® has been shown to reduce both the incidence and severity of ETEC-induced diarrhea in up to 90% of volunteers	Scandinavian Journal of Gastroenterology, 46:7-8, 862-868, DOI: 10.3109/00365521.2011.574726
Travelan as a broad Spectrum anti-bacterial	Immuron Limited, 29 April, 2011
Travelan® demonstrates broad reactivity to Vibrio cholera strains from Southeast Asia indicating broad potential for prevention of traveler's diarrhea	US Department of Defense, Armed Forces Research Institute of Medical Sciences (AFRIM), 4 September, 2019
Travelan® prevented clinical shigellosis (bacillary dysentery) in 75% of Travelan® treated animals compared to placebo and demonstrated a significant clinical benefit	US Department of Defense, Armed Forces Research Institute of Medical Sciences (AFRIM), 5 September, 2018
Travelan® able to bind and was reactive to 60 clinical isolates of each bacteria, Campylobacter, ETEC, and Shigella	US Department of Defense, Armed Forces Research Institute of Medical Sciences (AFRIM), 30 January, 2017
Efficacy of hyperimmune bovine colostrum against shigellosis in rhesus macaque (Macaca mulatta), and bioactivity of HBC against common enteric pathogens	Islam et al., 2020. Submitted to mSphere, American Society for Microbiology
Bioactive Immune Components of Travelan®	Clin Vaccine Immunol 24:e00186-16. https://doi.org/10.1128/CVI.00186-16
Hyperimmune bovine colostrum containing lipopolysaccharide antibodies (IMM-124E) has a non-detrimental effect on gut microbial communities in unchallenged mice	Rachele Gore, Mitra Mohsenipour, Jennifer L Wood, Gayathri K Balasuriya, Elisa L Hill-Yardin, Ashley E Franks
Administration of the Hyper-immune Bovine Colostrum Extract IMM-124E Ameliorates Experimental Murine Colitis	Journal of Crohn's and Colitis, Volume 13, Issue 6, June 2019, Pages 785–797, https://doi.org/10.1093/ecco-jcc/jjy213
IMM-529	
Bovine antibodies targeting primary and recurrent Clostridium difficile disease are a potent antibiotic alternative	Sci Rep 7, 3665 (2017). https://doi.org/10.1038/s41598-017-03982-5



ASX RELEASES REFERENCED

V	

Ref	Slide statement	ASX Release
а	Travelan® (IMM-124E) IND filed with and approved by FDA	Immuron Receives FDA Approval for Travelan IND Application
b	US Department of Defense grant of US\$4.45 million to examine a dosing regimen for Travelan® more suited for use by the military	Immuron awarded A\$6.2 million US DoD funding for Travelan
С	Market evaluation by Lumanity confirms the Traveler's Diarrhea market opportunity for IMM-124E (Travelan®)	AGM Presentation
d	Market evaluation by Lumanity confirms the Clostridioides difficile market opportunity for IMM-529	AGM Presentation
е	Travelan® - Uniformed Services University has recruited more than 20% of participants in a randomized clinical trial with Travelan® to evaluate the effectiveness for prophylaxis during deployment or travel to a high traveler's diarrhea risk region	US DOD Travelan Clinical Recruitment Milestone
f	Naval Medical Research Center Clinical Trials of CampETEC in campylobacter and enterotoxigenic E.coli (ETEC)	Immuron and US NMRC respond to CampETEC FDA Clinical Hold

