

Q2 FY24 Quarterly Investor Presentation

Microba Life Sciences Limited (ASX: MAP) (“Microba” or the “Company”) is pleased to provide its Q2 FY24 Investor Presentation to be delivered at the Quarterly Investor Webinar held with CEO Dr Luke Reid and SVP, Therapeutics Prof Trent Munro at 10:00am AEST / 11.00am AEDT today.

Presenting: Microba CEO, Dr Luke Reid and SVP Therapeutics Prof. Trent Munro

Time: 10:00am AEST / 11:00am AEDT today, Tuesday 30 January 2024

To register for the Quarterly Investor Webinar please click the link below. Investors can submit questions prior to the webinar to investor@microba.com or do so via the Zoom Q&A function during the webinar.

[Microba Q2 FY24 Investor Webinar Registration](#)

This announcement has been authorised for release by the Board of Directors

For further information, please contact:

Dr Luke Reid

Chief Executive Officer

E: Luke.Reid@microba.com

Investor / Media Relations

E: investor@microba.com

W: <https://ir.microba.com/>

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About Microba Life Sciences Limited

Microba Life Sciences is a precision microbiome company driven to improve human health. With world-leading technology for measuring the human gut microbiome, Microba is driving the discovery and development of novel therapeutics for major chronic diseases and delivering gut microbiome testing services globally to researchers, clinicians, and consumers. Through partnerships with leading organisations, Microba is powering the discovery of new relationships between the microbiome, health and disease for the development of new health solutions.

For more information visit: www.microba.com

Microba encourages all current investors to go paperless by registering their details with the designated registry service provider, Automic Group.

MICROBA™

Transforming health through the human microbiome

Q2 FY24 Results

30 January 2024

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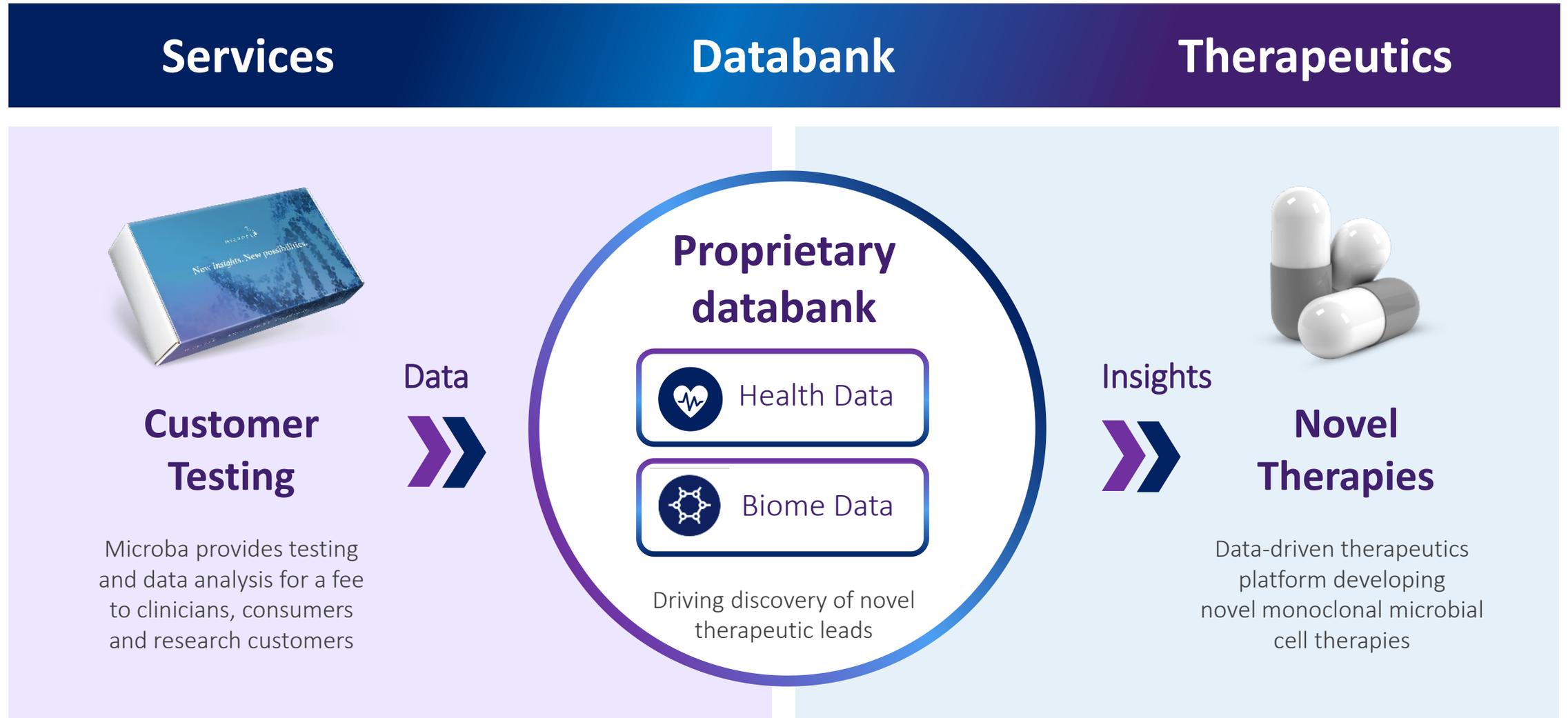
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Two core business segments driven by a proprietary databank



Company Snapshot



Continuous YoY revenue growth over 5 years of operations



Partnerships with market leaders incl. **Sonic Healthcare** (ASX: SHL), **Ginkgo Bioworks** (NYSE: DNA), **SYNLAB** (GR:SYAB) + more



Large, unique, proprietary microbiome databank



3 key therapeutic programs with successful **Phase I clinical trial** for lead IBD



World-leading technology in the emerging and rapidly growing **US\$4.9 billion** microbiome sector¹



Globally **leading microbiome expertise** complemented by drug discovery experts including Prof. Ian Frazer

Recent Milestones



SUCCESSFUL UK ACQUISITION

invivo[®]

Successful acquisition of United Kingdom microbiome testing company, Invivo Clinical.

- *a leading microbiome test provider for healthcare professionals in the UK*
- *established customer base of over 1,700 active healthcare professionals*
- *more than 20,000 microbiome tests sold since 2020*
- *evidence based supplement portfolio*
- *Unaudited sales in H1 FY24 totalled \$4.39m AUD*
- *has been self-funded from cashflow with no external capital required and has been consistently operating cashflow positive*



SUCCESSFUL PHASE I CLINICAL TRIAL FOR IBD LEAD DRUG CANDIDATE

Phase I Clinical Trial of Microba's lead therapeutic candidate MAP 315 successfully completed, demonstrating it is safe and well tolerated.

- *favourable safety and tolerability profile across both low and high dose cohorts.*
- *no clinically significant safety signals*
- *all participants completed the study and all dosing.*
- *faecal kinetics by metagenomic analysis indicates successful delivery into the gastrointestinal tract*
- *results provide strong positive support for continuing to advance the clinical development of MAP 315 for the treatment of Ulcerative Colitis*



2ND ALLERGY PROGRAM AGREEMENT SIGNED

Agreement signed with IFF (NYSE: IFF) to develop novel microbiome-based treatments for multiple forms of allergy

- *second agreement between Microba and IFF*
- *first agreement in 2021 saw the identification of lead species, and IFF has now selected a number of species to move forward into Stage 2.*
- *Stage 2 project will complete isolation and characterisation of strains selected from Stage 1 and characterisation of those strains.*
- *Upfront payment to Microba of AUD \$924k and has exclusive option to license for commercialization, which if executed may result in future royalty payments to Microba.*

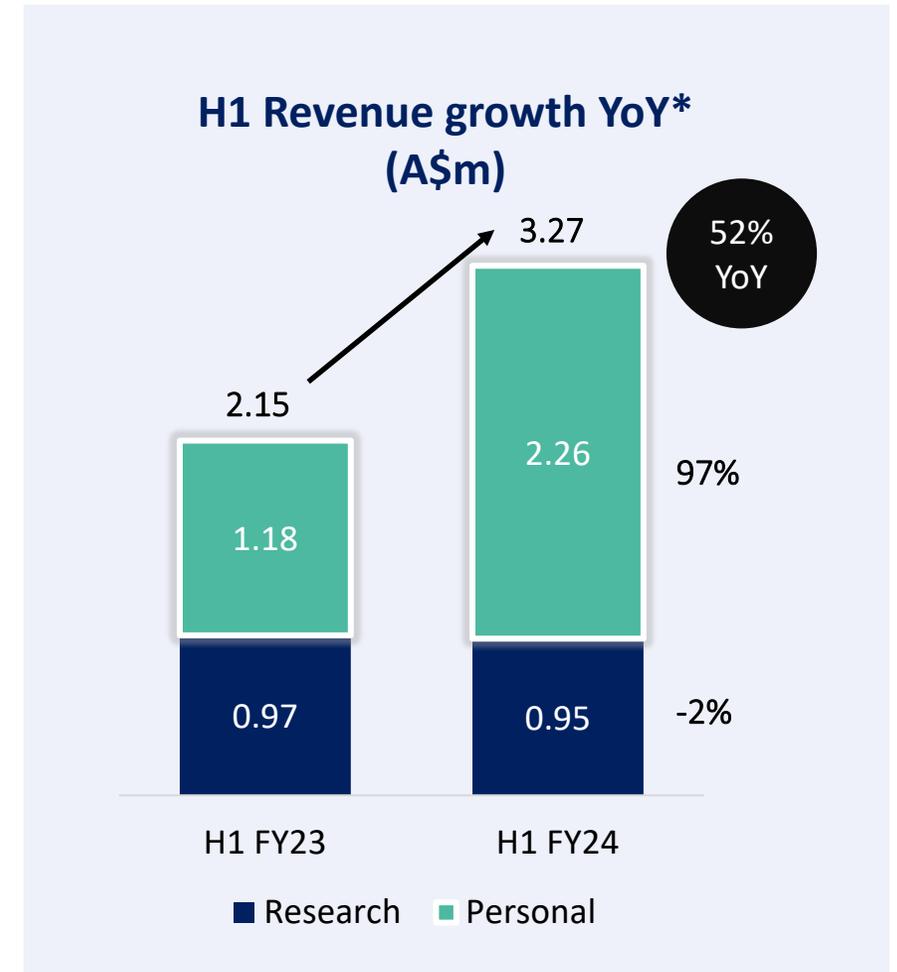
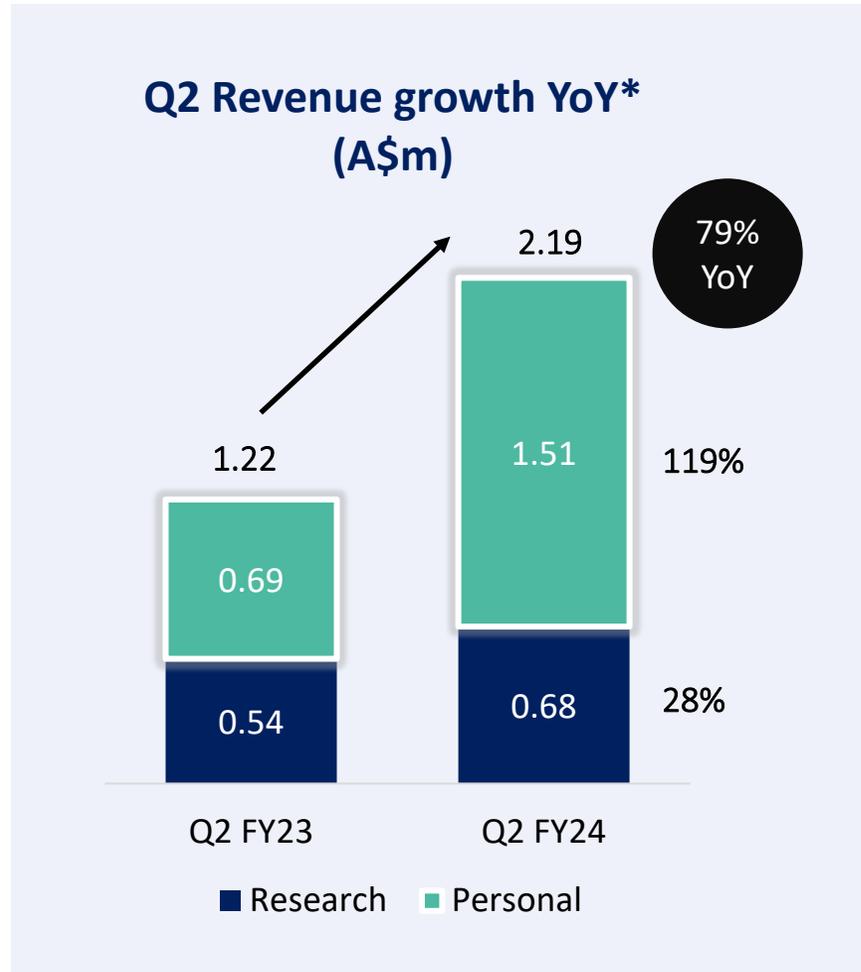


Microbiome Services

Driving global revenues
and data growth

Q2 Financial Highlights – Strong revenue growth

- Growth in MetaXplore translating into strong sales and revenue growth
- Major partner SYNLAB, Europe's largest medical diagnostic company, revenues up 115% YoY
- UK acquisition, Invivo Clinical, revenue contribution from 5 to 31 December



Advanced microbiome testing addressing large markets in healthcare

Addressable Healthcare Markets

Gastrointestinal disorders

MetaXplore™

Is the microbiome contributing to the patients DGBI and how can they manage?

150M est. addressable patients worldwide¹

Well established out of pocket private pay market

Gastrointestinal infection

MetaPanel™

Does the patient have a pathogen and how can I selectively treat it?

16M est. addressable patients worldwide²

Well established hospital direct-bill and path to reimbursement

¹Estimated based on the prevalence of specific Disorders of the Gut-Brain Interaction across 26 countries (Av prevalence of 32.8% DOI: [10.1111/nmo.14594](https://doi.org/10.1111/nmo.14594)), and the proportion regularly seeking medical support with one or more doctor visit per month (Average 15.4% - DOI: [10.1053/j.gastro.2020.04.014](https://doi.org/10.1053/j.gastro.2020.04.014))

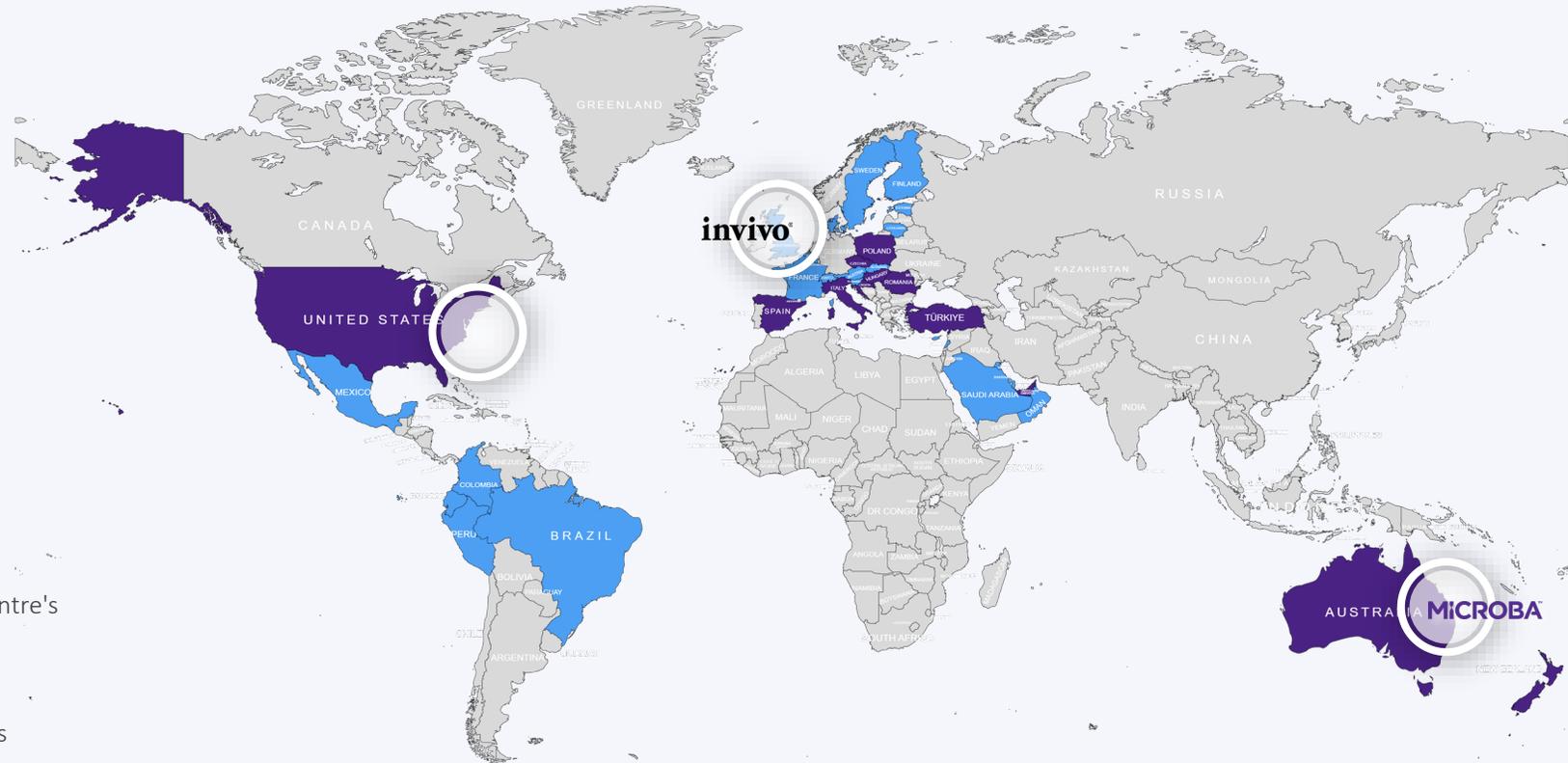
² Estimated based on the global number of immuno-compromised patients and other patients at high risk for gastrointestinal infection (>8.1m chemotherapy treated solid tumor cancer patients and >1.1m haematological cancer patients DOI: [https://doi.org/10.1016/S1470-2045\(19\)30163-9](https://doi.org/10.1016/S1470-2045(19)30163-9)), (>3m dialysis patients DOI: 10.1038/s41581-022-00542-7), (>140k Organ Transplant patients per year <https://www.transplant-observatory.org/>), (>3.5m long stay ICU patients – estimate based on data from <https://ourworldindata.org/grapher/intensive-care-beds-per-100000>)

Significant international distribution

Partners provides access into 35 countries with 17 now operational



Strategic acquisition unlocking a geographical base and laboratory in the UK, supporting the next phase of Microba's international growth strategy.



- Operating Team and/or Laboratory Centre's
- Contracted partner/s operational
- Access via current distribution partners

A scientist in a white lab coat and gloves is working in a laboratory. The scientist is holding a black tray with a grid of wells in their left hand and is using their right hand to interact with a Fisherbrand Bead Mill 24. The machine has a clear protective shield and a control panel with a digital display and several buttons. The background shows a typical laboratory environment with white cabinets and electrical outlets.

| UK Strategic Acquisition

| Unlocking the UK market through a targeted acquisition

invivo®

- UK microbiome company Invivo Clinical acquired in December 2023
- Over 1,700 healthcare professional customers*
- Self-funded with positive operating cashflow
- Positioned for growth acceleration

Laboratory - Bristol

- 10 staff
- PCR laboratory
- CL2 laboratory

Headquarters - Stroud

- 23 staff
- Finance, sales and marketing
- Customer service & clinical support



Multiple growth opportunities and Microba synergies

- Microba's MetaXplore test positioned to be delivered to Invivo customers and expected to fuel growth in GI test uptake
- Continued growth acceleration in Vaginal microbiome testing revenue with 40% growth in FY23.
- The integrative healthcare market is expanding as clinicians move to private practice due to pressure on the NHS.
- Small sales team being expanded with significant opportunity to accelerate growth.



Smith+Nephew

| Strengthening Invivo's position in GI microbiome testing

A platform for UK market entry and growth of Microba's MetaXplore™ test

 **79%**

of Invivo's **testing revenue** is through their GI products*

 **19%**

growth in GI testing revenue in FY23

“From our leading position in market, we expect Microba’s world leading MetaXplore™ will become the most favoured microbiome test for healthcare professionals in the UK” Debbie Cotton – Head of Clinical Innovation, Invivo

- **Current leading GI testing product for UK** healthcare professionals
- Featuring 7 host markers, 62 microbial markers
- Microbiome analysis via multiplex PCR
- Offers static PDF reports



- **Next generation advanced GI test** using Microba’s world leading metagenomic technology
- Includes 7 host markers, 28,000 microbial markers, 19 microbiome functions
- Offers both static PDF and interactive digital reports.



| Invivo Clinical H1 FY24 sales performance

H1 FY24 Key sales metrics

\$4.39m

H1 Sales

49%

H1 Gross Margin

4,208

H1 Tests Sold

62,527

H1 Supplements Sold

Microbiome Therapeutics

Developing novel monoclonal
microbial cell therapies

| Recent Program Highlights



Inflammatory Bowel Disease Program

MAP315 Phase I completed, product safe and well tolerated. Positive MOA data generated.

Trial completed and data confirms that MAP315 is safe and well tolerated. Data also indicates potential engraftment of MAP 315. Multi-modal mechanism of action (MOA) data generated for MAP315.



Immuno-Oncology Program

Confirmed potent anti-tumour activity.

Multiple mouse models have demonstrated that Microba's therapeutic leads can significantly reduce tumour burden. Further, immunological studies demonstrated activity consistent with induction of a specific and targeted immune response. Additional animal studies and immune profiling experiments are continuing to enable selection of a lead therapeutic candidate in H2 FY24.



Autoimmune Disease Program

Primary screening 100% complete.

Stage 1 activity screens complete with 62% of strains demonstrating significant immunomodulatory activity and a further 18% significantly impacting the inflammasome. 35 strains have now progressed into Stage 2 functional screening for expected completion in Q4 FY24 to enable lead candidate selection.

Precision Microbiome Therapeutics

- Precision microbiome analysis platform delivering unparalleled accuracy, coverage & depth.
- Data driven discovery platform utilising one of the worlds **most advanced & highly curated microbiome datasets**.
- Next generation approach to microbiome drug development **identifying and isolating single keystone species to develop Live Biotherapeutics**.



Novel Pipeline of Microbiome Therapies

- ✓ Potent and novel biology mediated by Live Biotherapeutic strategy
- ✓ Scalable GMP manufacturing process
- ✓ Oral delivery and Excellent safety profile

Experienced Leadership



Trent Munro
SVP Therapeutics



Prof Ian Frazer
Director & MAB Chair



Prof Gene Tyson
Co-Founder



Prof Phil Hugenholtz
Co-Founder

Microba Partners



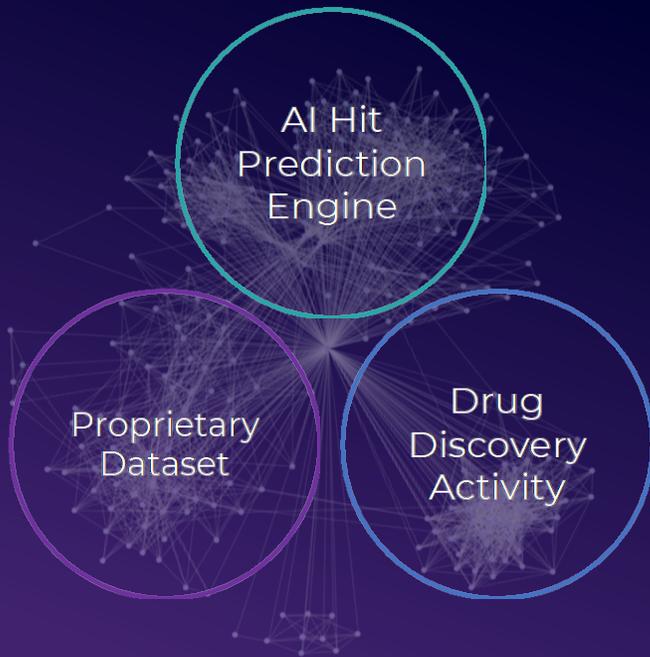
Massachusetts
Institute of
Technology



Queensland
University
of Technology



| Microba is Pioneering an Advanced AI/ML Approach to Development Next Generation Precision Live Biotherapeutics



Worlds most advanced bioinformatic pipeline for metagenomic microbiome analysis

Large human dataset w deep health metadata powering AI/ML* discovery

Repeatable pipeline of single strain live biotherapeutic candidates



Lead candidate MAP 315 safe and well tolerated in Phase I study with unique MOA

MAP 315 is Phase 2 ready for IBD (mild to moderate ulcerative colitis)



Identified oncology leads with checkpoint synergistic anti-tumour activity



Advanced candidate discovery program in auto-immune disease partnered with Ginkgo

Combines Human data driven leads with advanced high-throughput biodiscovery



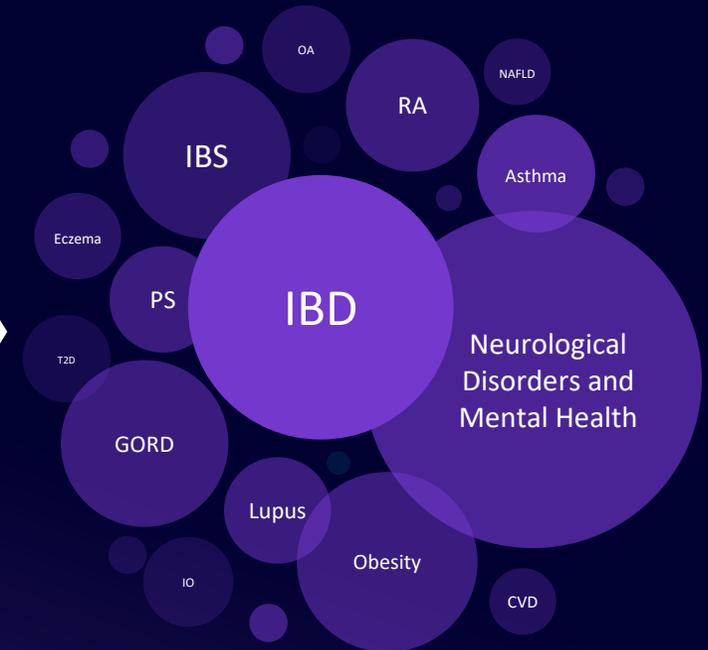
Microbiome Databank

Health Data

Biome Data

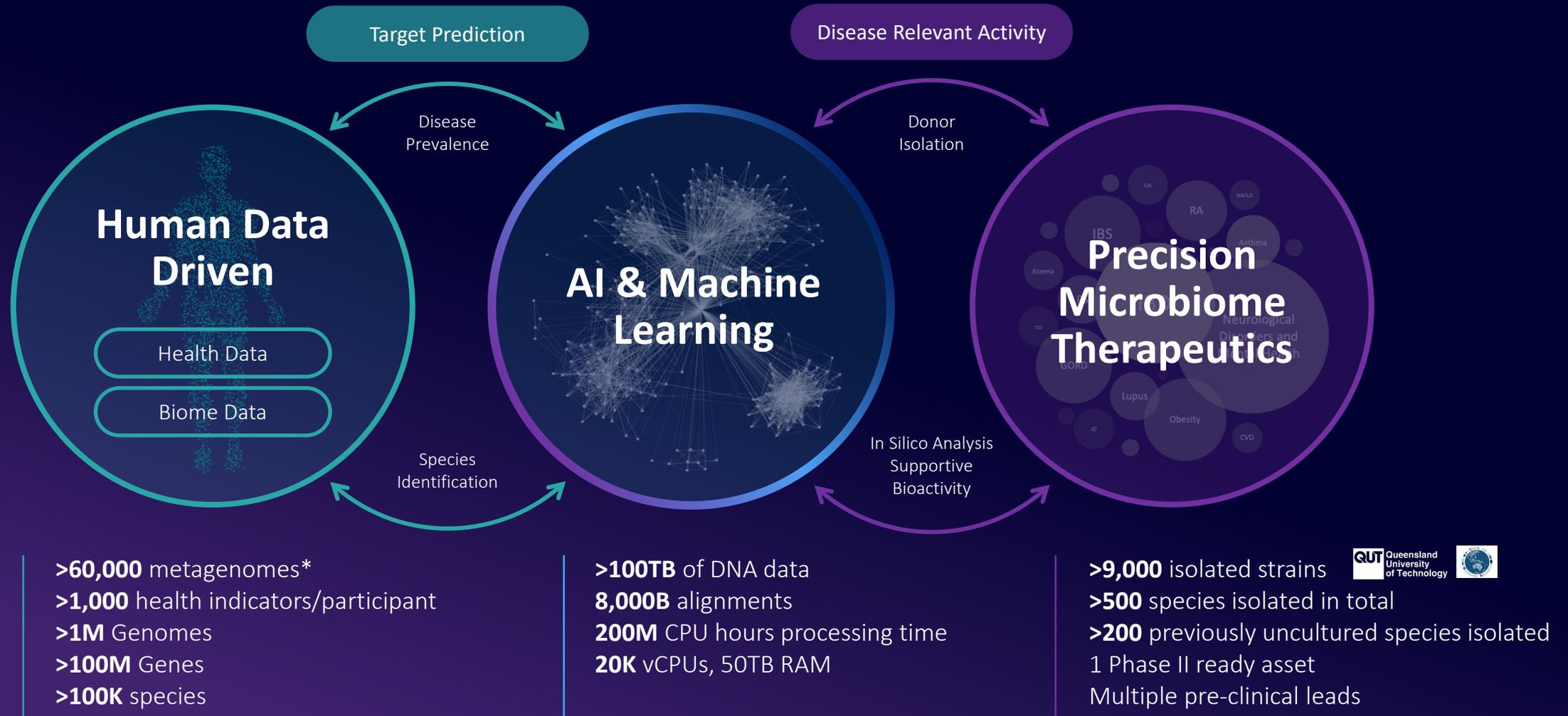


Human data driven insights

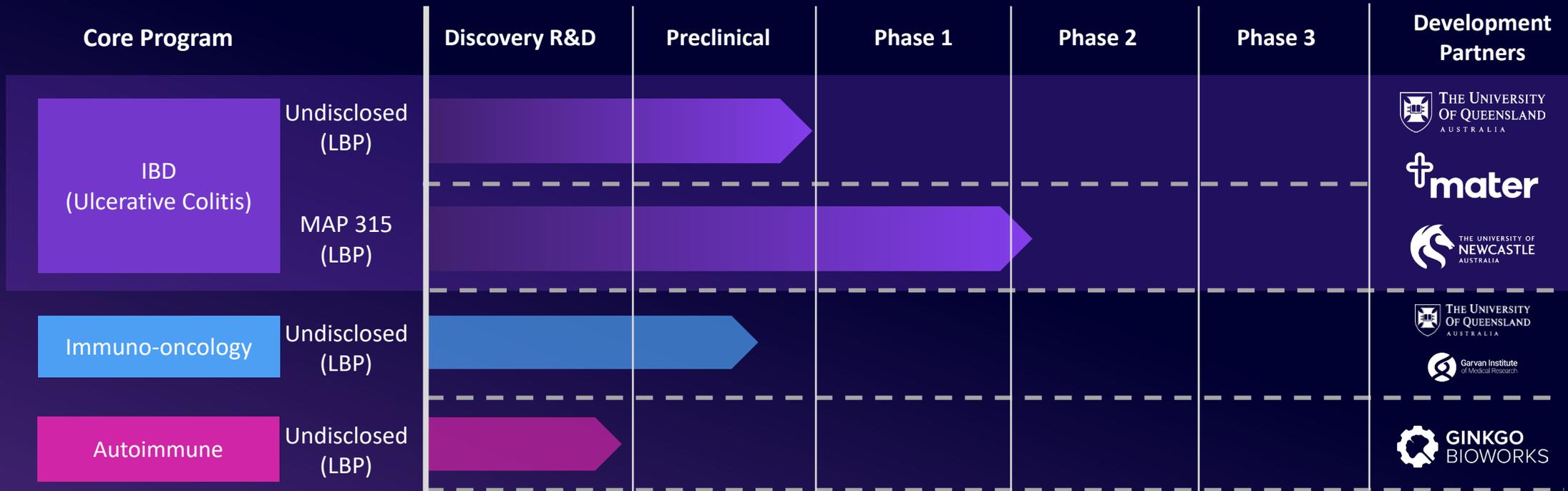


Precision Microbiome Therapeutics

Big data powered therapeutics from the human microbiome



Therapeutic Pipeline



| Advancing allergy treatment - partnership with multi-national probiotics leader IFF

Stage 1 complete discovering therapeutic leads

- ✓ Nov 2021 Microba signed a therapeutic partnership with IFF to develop novel treatments for multiple forms of allergy
- ✓ Stage 1 leveraged Microba's data-driven drug discovery platform to identify multiple leads
- ✓ IFF subsequently executed their option to proceed into a Stage 2 agreement to further pursue those leads

Stage 2 signed to complete isolation and activity assessment

- The parties have now signed a Stage 2 agreement to isolate strains from the identified lead species, and complete primary characterisation of those strains
- IFF will pay €560k (Approx. AUD \$920k) for the Stage 2 work
- IFF have an option to license the strains and intellectual property on commercial terms which may result in future royalties and milestone payments



THE PROVEN
LEADER
IN PROBIOTICS

NYSE IFF | Market cap \$19.6B* | HQ New York

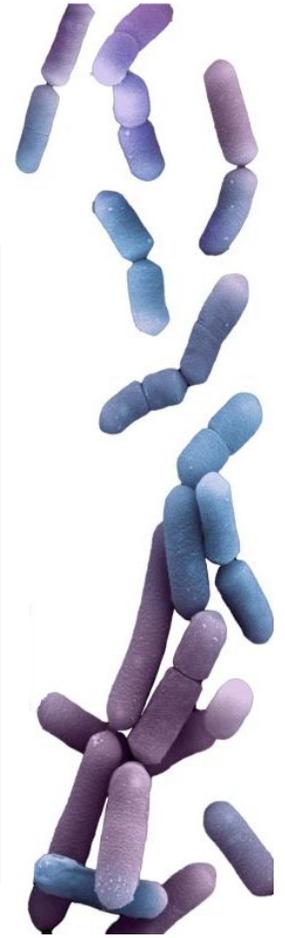
IFF Achieves Industrial Scale Production of a Strict Anaerobic 'Next Generation' Probiotic Strain

Oegstgeest, The Netherlands – Mar. 16, 2023 – IFF—a leading global player in the health and wellbeing industry—today reveals it has achieved the successful production of a strict anaerobic 'next generation' probiotic strain at industrial scale. The remarkable milestone demonstrates IFF's unique capabilities in fermentation, downstream processing and formulation in this space. IFF's advancement in identifying new microbiome solutions and establishing safe and efficient production at industrial scale will ultimately support commercially viable innovation in the dietary supplement market in years to come.

"We've made this breakthrough achievement thanks to our cutting-edge process development capabilities—which encompass small-scale, high-throughput and high-information tools—paired with our large-scale manufacturing skills and deep investments in our teams, facilities and certifications," said Sebastian Stahl, director for Process R&D at IFF. "Not only does the successful production of Akkermansia at industrial scale demonstrate that we have the know-how and technologies to identify promising microbiome solutions, but we're also experts at establishing their safe and efficient production."

The production of a 'next generation' probiotic at scale is another cornerstone of IFF's strategy to refresh and deliver microbiome-science-based solutions through a mixture of strategic partnerships and investments in research and development and builds on the company's existing leadership in the areas of probiotics, prebiotics and botanical extracts.

"We are proud of this milestone in IFF's history, and we are confident in our ability to commercialize our extensive microbiome pipeline, comprising in-house developments, as well as projects with external partners or customers," said Sebastien Guery, vice president, Health, IFF.



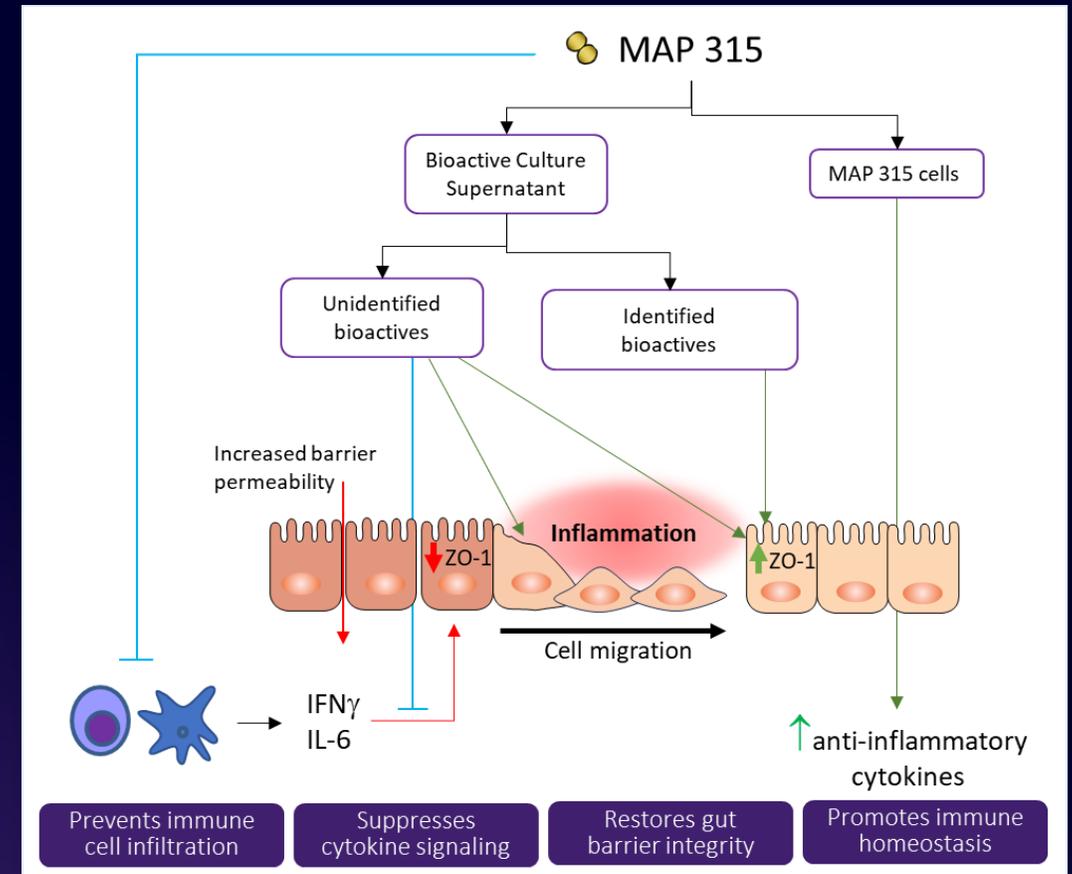
Inflammatory Bowel Disease Program

MAP 315 Proposed Mechanism of Action in IBD

Inflammatory Bowel Disease affects more than 6 million people globally.

Many of the current therapeutics suffer from poor compliance, undesirable side-effects and achieve low rates of mucosal healing.

MAP 315 is a safe, oral live biotherapeutic that promotes mucosal healing and immune homeostasis.



MAP 315 is safe and well tolerated in a Phase I clinical trial

Phase I Clinical trial of MAP 315 completed in Australia

- FDA pre-IND feedback positive
- Lead drug candidate MAP 315, demonstrated it is safe and well tolerated at both low and high doses in the Phase I, healthy volunteer clinical trial.
- Unblinded analysis of trial data demonstrated no clinically significant safety signals from assessments including ECGs and laboratory analysis of haematology, coagulation, clinical chemistry, urinalysis parameters, or impact on inflammatory biomarkers.
- Faecal kinetics assessed by metagenomic analysis indicated presence of MAP 315 at the terminal 28-day analysis timepoint, 14 days after the completion of dosing, confirming the ability to successfully deliver live MAP315 into the gastrointestinal tract.

Phase I trial details

- GMP Material produced with Bacthera
- Placebo controlled, double blind, healthy volunteer study
- Low and High dose via daily oral delivery for 2 weeks

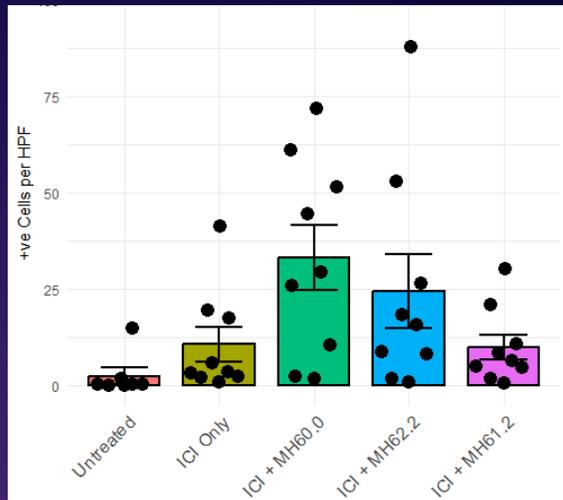
Phase II trial planning Initiated

- Protocol development in final stages
- Study start timing currently being determined - Adaptive Patient Study, N = 140 to 200 examining induction of remission

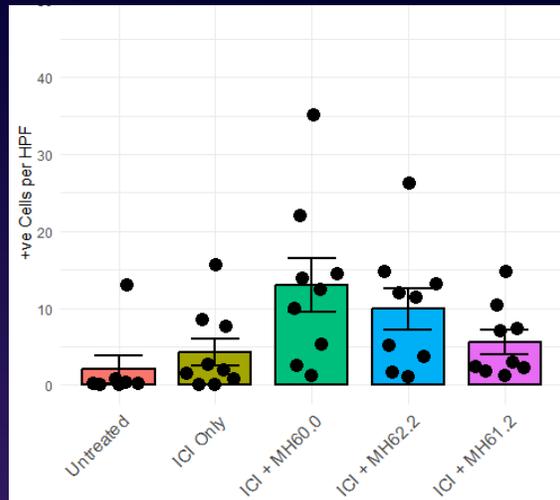
| Oncology Program

Our Oncology Leads Drive Potent Biology and May Turn Cold Tumours Hot

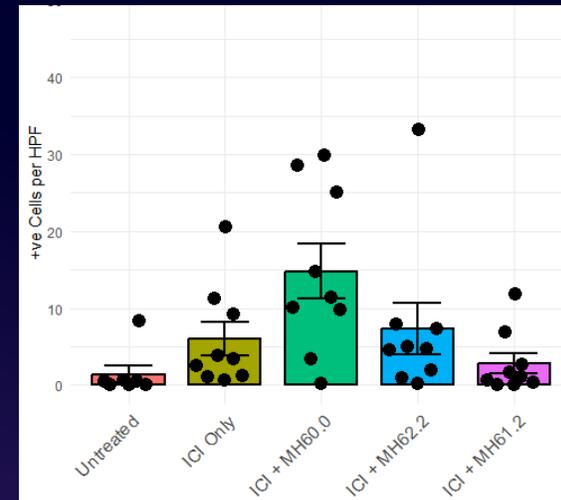
CD3



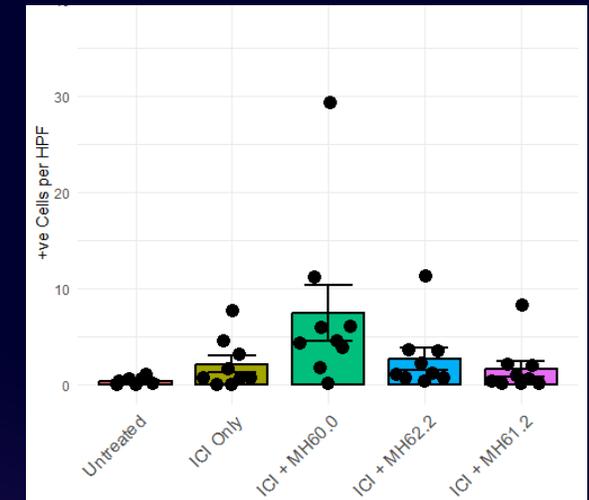
CD4



CD8

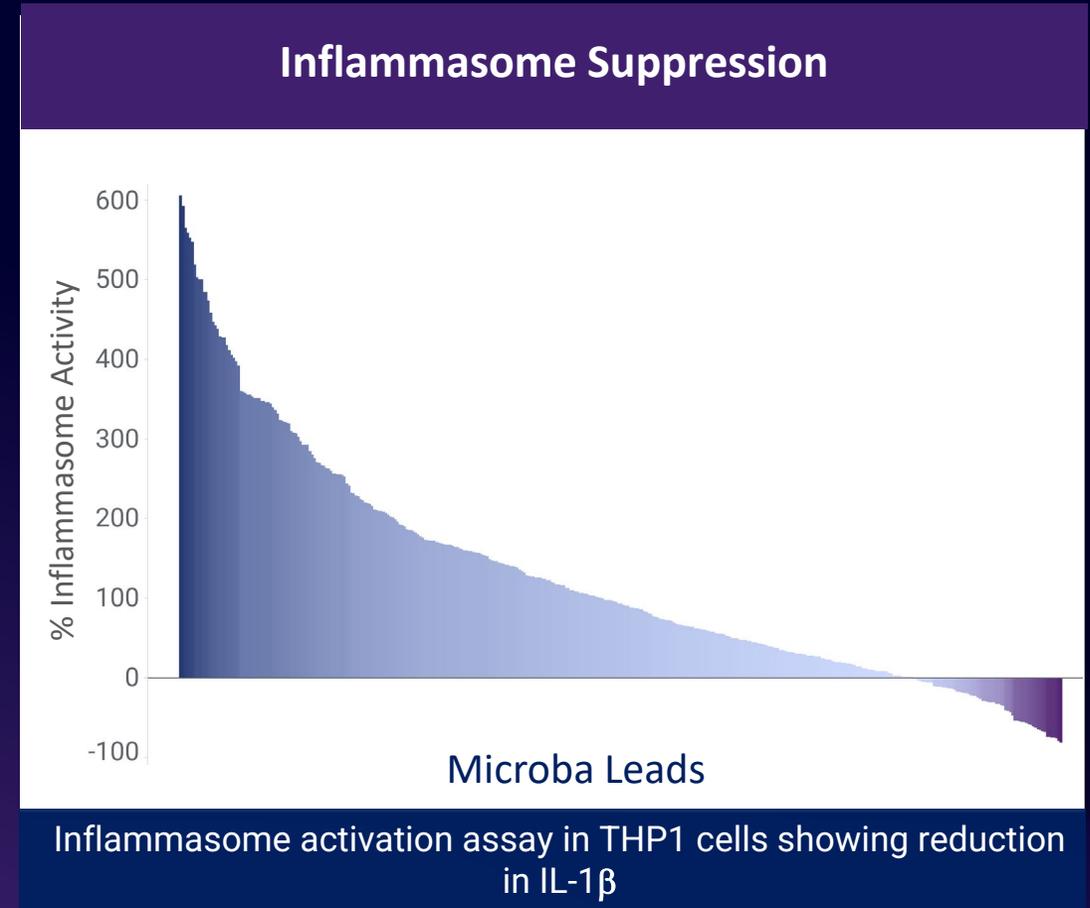
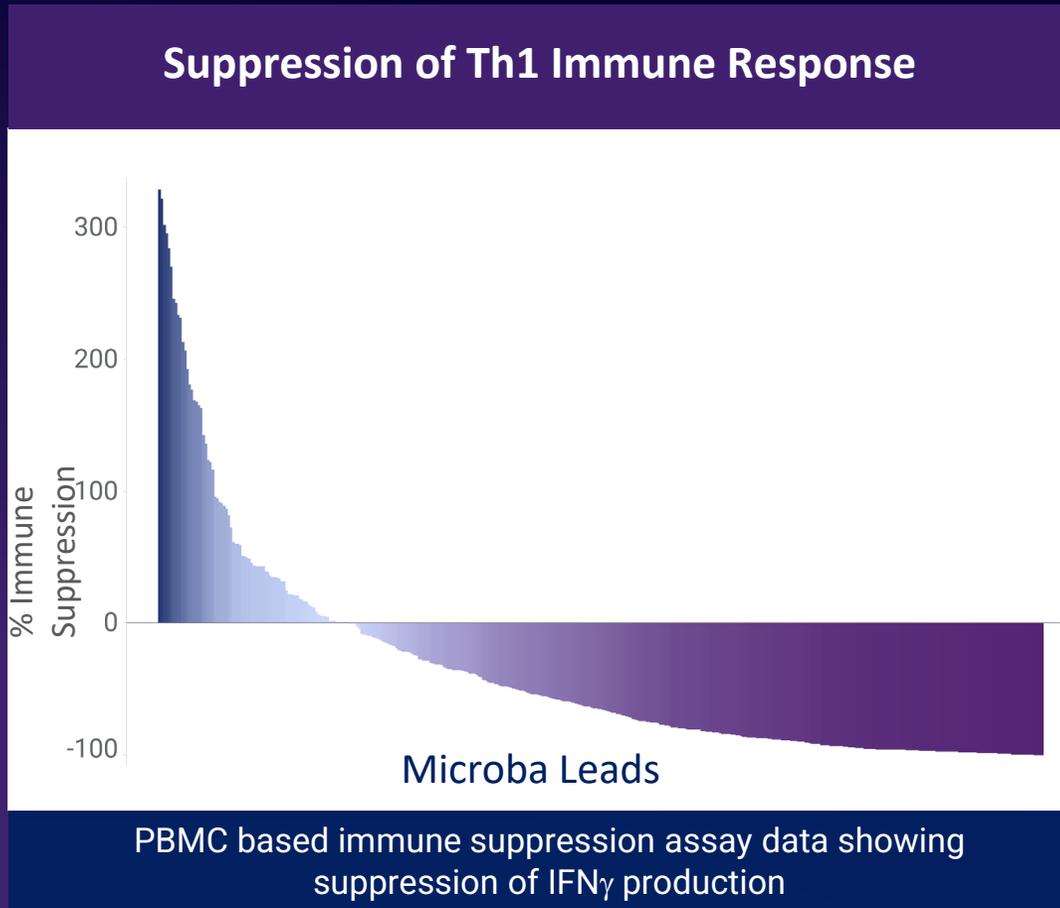


NK1.1



- Little or no increase in T cell infiltrate by ICI only (“responder” mice slightly higher)
- MH60 causes higher CD4 and CD8 infiltration
- MH62 causes higher CD4 infiltration

Significant hit rate and potency observed in primary screens



Microba is developing the 3rd Generation of Microbiome Therapeutics using a human data driven approach

1 First Generation



Fecal Doner Derived & FMT

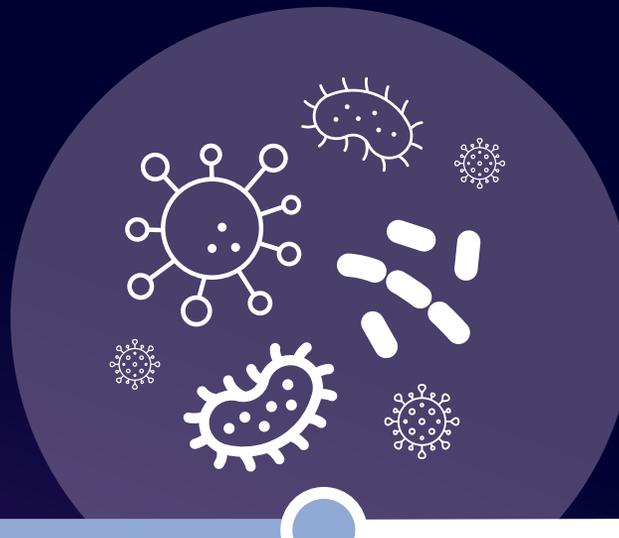


FDA approval Nov 30 2022
Faeacal doner derived enema drug product



FDA approval April 27 2023
Faeacal doner derived oral capsule drug product

2 Second Generation



Bacterial Consortia



Vedanta Biosciences
Vedanta Biosciences Raises \$106.5M in Funding



SER-155
Phase Ib showing engraftment reduction in pathogens

3 Next Generation



Single Strain & Small Molecule



Microba
MAP 315 Phase I IBD
Advancing oncology & autoimmune pipeline

| Recent IBD and microbiome deals

Licensee / Acquiror	Licensor / Target	Date	Type	Deal size	Indication / clinical stage
		October 2023	Acquisition	Total deal size: US\$7.2b	Ulcerative Colitis (Phase 2)
		Apr 2023	Acquisition	Total deal size: US\$10.8b	Ulcerative Colitis (phase 2) Crohn's disease (phase 2A) Autoimmune diseases (phase I or pre clinical)
		July 2022	R&D collaboration and license agreement	Total deal size: Undisclosed Upfront: €40m Sales / regulatory milestones: Undisclosed	Food allergies and IBD (pre clinical)
		Mar 2022	Acquisition	Total deal size: US\$6.7b	Ulcerative Colitis (phase 3), Crohn's disease (phase 2/3), Atopic dermatitis (phase 2 complete)
		Nov 2021	R&D collaboration and license agreement	Total deal size: US\$605m Upfront: US\$15m Additional payments: US\$590m	IBD (Discovery)
		Jul 2021	Licensing agreement in US/Canada	Total deal size: US\$525m Upfront: US\$175m Sales / regulatory milestones: Up to US\$225m	Recurrent Clostridioides difficile infection (Phase 3)

<https://www.roche.com/investors/updates/inv-update-2023-10-23>
<https://www.merck.com/news/merck-strengthens-immunology-pipeline-with-acquisition-of-prometheus-biosciences-inc/>
<https://www.enterome.com/news-events/enterome-signs-major-strategic-rd-collaboration-with-nestle-health-science-to-develop-and-commercialize-new-allermimics-and-endomimics-immunotherapies-for-food-allergies-and-inflam/>

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<https://ir.serestherapeutics.com/news-releases/news-release-details/seres-therapeutics-nestle-health-science-announce-ser-109-co>

Outlook



Upcoming Company Milestones

2023 APR - JUN	2023 JUL - DEC	2024 JAN - MAR
Services New international distribution deals 	Services New country expansion 	Services Invivo Clinical integration
Therapeutics - IBD MAP 315 program HREC Approval for Phase I trial 	Services First Sonic partners operational 	Services Sonic MetaPanel national launch
Therapeutics - IBD MAP 315 program GMP manufacture complete Phase I 	Therapeutics - IBD MAP 315 program Phase I participant dosing complete 	Therapeutics - IBD MAP 315 program Phase II clinical trial ready
Therapeutics - IBD MAP 315 program Phase I trial commencement 	Therapeutics - IBD MAP 315 program Phase I complete 	Therapeutics – IO Program MOA studies to support candidate selection
	Therapeutics – IO Program Immunological pre-clinical results 	Therapeutics - Autoimmune Program Stage 2 screening to support Q4 completion
	Therapeutics - Autoimmune Program Stage 1 activity screening complete 	

| Therapeutic program milestone tracking

Inflammatory Bowel Disease



PHASE I COMPLETE

Recently Completed

- ✓ Phase I HREC approval
- ✓ cGMP manufacturing complete
- ✓ Phase I trial commenced
- ✓ Phase I participant dosing complete
- ✓ Phase I complete

Upcoming milestones

- Phase II readiness



Immuno-Oncology Program



PRECLINICAL

Recently Completed

- ✓ Leads discovered
- ✓ Leads isolated
- ✓ First animal model results
- ✓ Immunological pre-clinical data

Upcoming milestones

- Lead candidate selection



Autoimmune Diseases



STAGE 2 SCREENING

Recently Completed

- ✓ Program commenced
- ✓ Strains provided to Ginkgo
- ✓ First in vitro screening results
- ✓ Stage 1 activity screening complete

Upcoming milestones

- Stage 2 activity screening complete



MICROBA™



Dr Luke Reid

Chief Executive Officer
luke.reid@microba.com



Pasquale Rombola

Chairman
pasquale.rombola@microba.com

CONTACT

Head Office

Level 10, 324 Queen Street
Brisbane QLD Australia

Laboratory

Princess Alexandra Hospital
Woolloongabba QLD Australia