mgc pharma



Developing innovative medicines for unmet medical needs

MGC Pharma is developing innovative medicines for unmet medical needs

"Our vision is to have a global impact on unmet medical needs."

Wielding the power of plant-inspired medicines in novel combinations by using innovative medical technologies to harness knowledge and science towards achieving our "Nature to Medicine" strategy and creating the next-generation of medications."

Roby Zomer

Managing Director and CEO

Established pharma platform

- Plant-inspired ingredients and pharmaceutical formulations
- Advanced clinical programs with innovative formulations
- Core drug development in progress with a strong R&D pipeline
- Integrated global research hub
- Experienced corporate and pharmaceutical team
- Malta GMP standard manufacturing facility
- ESG focussed corporate culture

Invested in a highly qualified team focused on moving products

through clinical trials



Roby Zomer Managing Director & CEO



Brett Mitchell Non-Executive Chairman







Yifat SteuerChief Operational
Officer



Tom Cairns
Chief Accounting
Officer



Robert Clements
Chief Commercial
Officer



Itay Nissim
Chief Manufacturing
and Supply Chain
Officer



Dr. Nadya LisodoverChief Medical Officer



Amir Polak
Chief Pharmaceutical
Development Officer















MGC Pharma is a pharmaceutical company, accelerating medicine for clinical trial

- Robust development pipeline targeting widespread medical conditions, focusing on CannEpil® and CimetrA® (in the short term)
- Develop and supply high-quality, plant-inspired medicines for the medical markets globally
- Growing patient base in Australia, the UK, Brazil and Ireland and a global distribution footprint via an extensive network of commercial partners
- Partnerships established with renowned institutions and academia to optimise the development of targeted plant-inspired medicines
- EU-GMP certified manufacturing facilities

Research and manufacturing

Integrated research hub

- Currently manages and runs clinical trials both in-house* and with third party CROs
- Israeli CRO ('MediCaNL') delivers significant cost savings to the Company
- Opportunity for third party revenue generation

Fully built GMP pharma standard manufacturing facility

- Two, high-quality, European production facilities to manufacture and distribute MGC's proprietary IMP products CannEpil®, CimetrA®, and CogniCann®
- Slovenia production facility EU-GMP since 2018
- Malta production facility EU-GMP certification in early 2023





^{*} In accordance with the European Medicines Agency, Federal Drug Administration, ICH Good Clinical Practice, and Israeli health regulations.

2022/23 achievements

During the past 24 months the Company refocussed into a pharmaceutical business

Drug development

- 2 core drugs in clinical development
 - CannEpil®
 - CimetrA®
- 5 further drugs in development pipeline
- Completion of two successful Phase II studies on CogniCann® and CimetrA®
- Appointment of Ingenu as CRO to run Clinical Trials in Australia

Strategic development

- CannEpil® is now available to patients in the UK by Named Patient Request, to be prescribed by clinicians in the UK who are listed on the GMC Specialist Register.
- CannEpil® accepted by the Irish Health Product Regulatory Authority (HPRA) and obtaining full health insurance coverage by LTI or GMS scheme.
- ArtemiC[™] (CimetrA® supplement) has now been listed as an over-the-counter (OTC), non-prescription drug in the USA, following the listing on the FDA National Drug Code Database (NDC).
- Slovenian Ministry of Health approved MGC to carry out scientific research and development on psychedelic compound Psilocybin
- Construction completed of two pharma-grade production sites (Malta and Slovenia)

Looking ahead (2023- 2024) Key strategic milestones

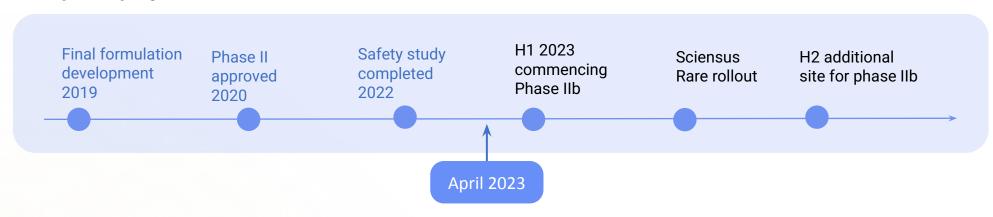
- Ramp up of CannEpil® 'early access' program with Sciensus Rare
- Progress on CannEpil® Phase IIb
- Completion of large animal study for CimetrA®
- CimetrA® IND submission to the US FDA
- Commissioning and GMP approval of Maltese CannEpil® and CimetrA® facility



MGC Pharma drug development programme

CannEpil®

Development progression



CimetrA®

Development progression



Flagship Product – Refractory Epilepsy

CannEpil®

- IMP based on two active ingredients:
 - Cannabidiol and Delta-9-Tetrahydrocannabinol
 - Delivered by oral mucosal solution
- Approximately 30% of generalised seizures epilepsy patients have Refractory Epilepsy aka "Drug-Resistant Epilepsy" (DRE)¹
- Available to patients in the UK by Named Patient Request, to be prescribed by clinicians in the UK who are listed on the GMC Specialist Register
- ❖ Fully covered under Ireland's Primary Care Reimbursement Service via long term illness and general medical services schemes²
- Real-world data and early regulatory approval (hundreds of patients under early access since 2019)
- Results of Preclinical and Clinical program to date :
 - Shows positive safety assessments
 - Safety study completed CannEpil® was found to be safe for post-treatment driving activities³
- ♦ IP protected (IP Number SIPO 26056)
 - 1. Fattorusso A, et al. (2021) The Pharmacoresistant Epilepsy: An Overview on Existant and New Emerging Therapies. Front. Neurol. 12:674483.
 - 2. https://www2.hse.ie/services/schemes-allowances/medical-cannabis-products-reimbursement-scheme/
 - ASX Announcement 15/8/22
- 4. https://www.biospace.com/article/epilepsy-therapeutics-market-worth-15-1-bn-by-2030-at-a-cagr-of-4-5-percent-/

Phase IIb



Total Addressable Market £15.1bn4

CannEpil® market potential

There are 50m¹ million epilepsy sufferers of which 6.2m¹ in Europe and UK

Approximately **30% of generalised seizures epilepsy patients** have **Refractory Epilepsy** aka "Drug-Resistant Epilepsy" (DRE)²





Treatment Estimate per patient



• 15m

• 1.87m

Treatment cost estimated £70003

https://www.who.int/news-room/fact-sheets/detail/epilepsy

Fattorusso A, et al. (2021) The Pharmacoresistant Epilepsy: An Overview on Existant and New Emerging Therapies. Front. Neurol. 12:674483.

^{3.} Alacrita Report 2019

Flagship Product – COVID-19 and anti-inflammatory

CimetrA®

- ♦ IMP based on two active ingredients (Rollout of ArtemiC™ supplement):
 - Dry extracts of Curcuma longa and Boswellia serrata
 - Delivered via nanoparticle suspension
- Results of Preclinical and Clinical program to date:
 - Demonstrated suppression of cytokine storm in COVID-19 patients¹
 - None of the treatment group patients in the on-going study have required additional oxygen, mechanical ventilation, or admission to intensive care²
 - Pre-clinical in-vitro study indicates wide-ranging application as an anti-inflammatory treatment through the modulation of the production of pro-inflammatory cytokines¹
 - Effective blocking of the IL-32mRNA expression¹, the pro-inflammatory cytokine related to rheumatoid arthritis, inflammatory bowel disease, asthma, psoriasis, and chronic obstructive pulmonary disease
- ♦ IP protected (IP Number SIPO 26055)

Phase IIb and Phase III



Total Addressable Market £13.6bn4

Developed from our food supplement ArtemiC®.

ASX Announcement 27/6/2022

ASX Announcement 15/12/2020

^{3.} https://www.researchandmarkets.com/reports/5733923/coronavirus-covid-19-current-therapy-global

Potentially, CimetrA® can be repurposed to treat inflammatory conditions

Influenza

3-5m

3 to 5 million cases of **severe illness** of influenza worldwide per
annum

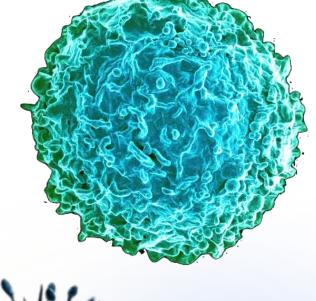
290-650k

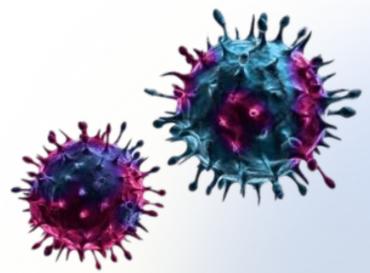
290 000 to 650 000 respiratory **deaths** per annum

https://www.who.int/news-room/fact-sheets/detail/influenza-(seasonal)

Anti Inflammatory market £100b

Total Addressable Market





Investment summary

MGC has successfully completed an equity fundraise of c.£1.83m with the following components:

- Existing shareholder support of c.£913k and new capital committed of c. £292k
- Clinical trial funding Cantheon Capital LLP of c.£630k committed from clinical trial specialist fund for CannEpil® phase IIb*

R&D Tax Credit

The clinical trail run in Australia also allows the Company to apply for a non-dilutive R&D government rebate of up to c.£1m

Broker Option

Broker option available and estimated at £750k

Use of proceeds

	GBP
CimetrA® completion of pre-clinical (large a & IND submission	animal) 2,030,000
CannEpil® preparation of pre-IND (FDA) & site outside of Israel	new PII 1,260,000
Working capital	220,000
Total	3,500,000

This investment round will allow for:

- Submission of IND to the FDA for CimetrA®, triggering MGC's agreement with AMC
- Progressing of CannEpil® through Phase IIb which will be fully funded for 2023 with commitment for 2024 and a ramp up of early access program with our distributor, Sciensus Rare

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