

## **Argent BioPharma**

## An innovative approach to drug development

Argent BioPharma is a clinical-stage specialty biopharma company developing therapeutics targeting areas of unmet need. Currently focused on CNS conditions and immunology, Argent has a broad pipeline of assets, led by CannEpil (drug-resistant epilepsy), CimetrA (acute lung disease) and CogniCann (CNS disorders). The first two are available under special access schemes in markets such as the UK, US and EU, generating early revenue streams as well as real-world evidence, which should aid regulatory outcomes. Argent has a strong internal IP position with two EU-GMP R&D and manufacturing sites (in Malta and Slovenia) supporting inhouse drug development. These could potentially be monetised via contract manufacturing services. We expect the most significant upcoming catalysts to be the investigational new drug (IND) filings with the FDA for CannEpil and CimetrA, expected by end 2025 according to management.

### A differentiated portfolio and business strategy...

Argent's pipeline, developed fully in-house, targets a broad range of indications, all underserved areas but with significant commercial potential. Leading the pack are CannEpil, targeting drug-resistant/refractory epilepsy (market expected to be worth \$6bn by 2034), currently available in the UK and covered by insurance in Ireland, and CimetrA, a drug in research for acute lung injury and acute respiratory distress syndrome, currently available in the US as an over-the-counter unlicensed drug. CogniCann targets symptoms related to Alzheimer's, while the preclinical pipeline is focused on diverse areas such brain cancer and wound management. Argent's strategic rationale of offering its drugs under special access schemes holds merit in our opinion, not only providing an early revenue stream but also de-risking the development process, given the real-time safety and efficacy data generated.

## ...backed by in-house capabilities

The company's R&D efforts are supported by its two EU-GMP facilities, in Slovenia and Malta (with capacity to manufacture over 20k units/day), which management believes will be able to support the clinical and commercial requirements of its pipeline. We also see monetisation opportunities from offering contract manufacturing services as the company scales internal operations.

## **Next up: US IND filings**

We expect Argent's medium-term focus to be on progressing its pipeline while continuing to implement its early access programmes across the EU, the UK and the US. We view the most significant upcoming catalysts to be the IND filings with the FDA for CannEpil and CimetrA, expected by end 2025 according to management.

Historical financials						
Year end	Revenue (A\$m)	PBT (A\$m)	EPS (c)	DPS (c)	P/E (x)	Yield (%)
06/20	2.1	(18.8)	(1.40)	N/A	N/A	N/A
06/21	3.0	(15.9)	(0.83)	N/A	N/A	N/A
06/22	4.7	(20.8)	(0.79)	N/A	N/A	N/A
06/23	3.4	(21.1)	(0.71)	N/A	N/A	N/A
Source: Compan	y filings					

Pharma and biotech

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#### Share details

Codes RGT.AX (to delist soon), RGT.L
Listings ASX, LSE, OTCQB Venture Market
Shares in issue 48.41m

#### **Business description**

Argent BioPharma (previously MGC Pharma) is a revenue-generating, clinical-stage biopharma company, focused on developing treatments for neurological and immunological conditions. Its pipeline includes three clinical-stage assets (Phase I and II) and one preclinical asset, led by CannEpil (drug-resistant epilepsy) and CimetrA (acute lung disease), both available under special access schemes in the EU, the UK and the US. Management plans to file IND applications for the two assets with the FDA by end 2025.

#### Bull

- Revenue generating, with sales recognised through early access schemes.
- Targeting areas of significant unmet need and commercial potential.
- Growing exposure to the key markets of EU, UK and US with two products on the market (under special access) with IND filings planned by end 2025

#### Bear

- Portfolio in the early stages of clinical development, with long lead times to approval under traditional regulatory pathways.
- Currently funded to Q424 but will need to raise significant capital to advance its development pipeline to the market.
- Potentially more stringent regulatory oversight for some of the assets.

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