

Corporate Structure



Epichem Synthetic Medicinal Chemistry Expected 2019-2020 sales A\$ 4.2m+

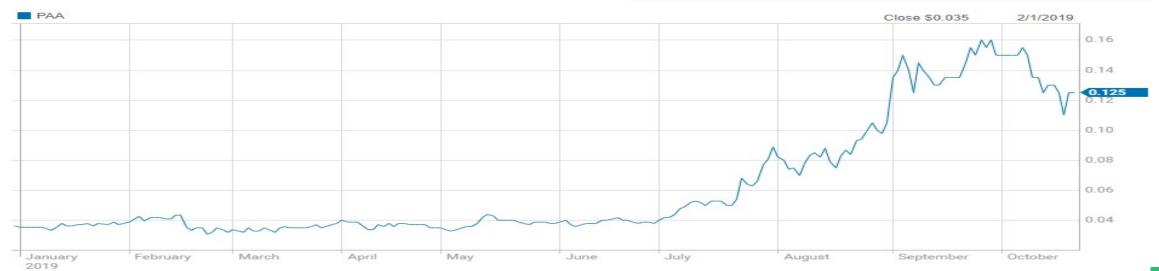
Pitney Pharmaceuticals Focus on Developing and repurposing a registered drug (Monepantel) in oncology



Corporate Snapshot

ASX Code:	PAA
Market Cap at \$0.125	\$35M
Cash (as at 30 June 2019)	\$2.1M
Debt (EFIC)	\$325K
Epichem Revenue Forecast 2019-2020	\$4.2M

Total Shares on Issue	301,814,647
Options (Unlisted)	56,801,956
Top 20 Own	35%
Board/Exec Own	9.3%





Experienced Board & Management Team

Dr. Roger Aston, Executive Chairman

- > 30 years experience in the pharmaceutical and healthcare industries.
- Director or chairman on a number of boards carrying out late stage drug development.

Robert Bishop, Executive Director

- > 30 years experience in corporate finance and equity capital markets
- Lawyer and an investment banker.

Neville Bassett, Non-Executive Director

- Member of the Order of Australia (AM)
- > 35 years working in accounting, finance and stockbroking

Sam Wright, Director & Company Secretary

- > 20 years experience in biotech and healthcare.
- Extensive experience in relation to public company responsibilities, including ASX and ASIC compliance, corporate governance, statutory financial reporting, and shareholder relations.

Dr Richard Mollard, Chief Scientific Officer

- > 20 years experience in biotech and pharma
- Extensive national and international experience.

Colin La Galia, Epichem CEO

 Highly experienced executive in pharmaceuticals, devices and diagnostics, both locally and internationally, and has demonstrated great success in international business development



PharmAust Background

- Lead product is **Monepantel** (MPL) a repurposed drug already approved for Veterinary use by Elanco Animal Health (US \$10.7b)
- PharmAust patented MPL as an anti-cancer drug
- Clinical strategy targeting MPL for vet and human applications
- Option Agreement with Elanco US Inc for veterinary cancer applications
- **Epichem**: profitable business, forecast revenues of \$4.2m in FY2019/20



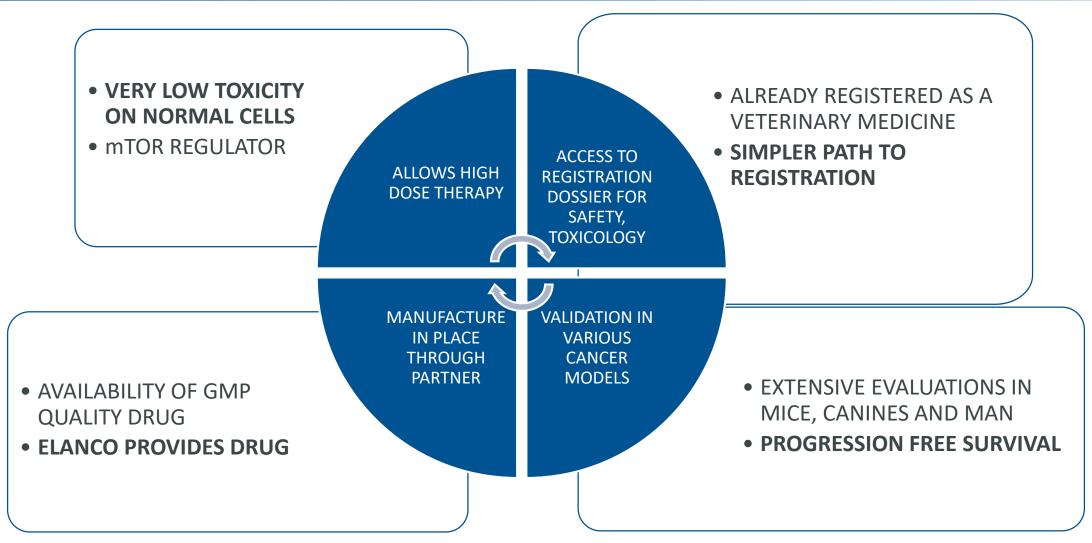


Chemotherapy (Global Sales \$52B) Mainstay Of Cancer Treatment

- Chemotherapy kills rapidly dividing cells
- Typically tumour cells replicate more quickly than normal cells and this difference is harnessed in chemotherapy to selectively kill cancer
- Side Effects during chemotherapy arise because normal cells also need to divide and they become casualties (immune system, digestive system and hair follicle)
- Monepantel, being an mTOR inhibitor acts differently and demonstrates minimal toxicology



Monepantel (MPL) In Oncology A Unique Anti-Cancer Paradigm





Elanco Animal Health – Option Agreement

Elanco Animal Health (NYSE:ELAN) – MC=\$10.7B US



- Monepantel compound is owned by Elanco, approved for the treatment of parasitic infections in sheep (patent protection to 2024)
- Option Agreement with Elanco to exclusive, worldwide royalty bearing license to commercialise MPL for treatment of Cancer in Animals

Commercial Outcome would allow PharmAust to focus on Human Cancer market



Elanco To Become No.2 In Animal Medicines



Elanco strategically enters therapeutic oncology arena by the purchase of Aratana Therapeutics

Elanco agree on \$7.6billion acquisition of Bayer's veterinary drugs unit



RISK MITIGATION IN NEXT STAGE DEVELOPMENT

TUMOUR MARKERS IN
HUMAN PROSTATE, LUNG
AND BOWEL CANCER
SIGNIFICANLY SUPPRESSED BY
MPL

CANINES RECEIVING MPL SHOW SIGNIFICANCE IN PROGRESSION FREE SURVIVAL

MPL

PROGRESSION FREE
SURVIVAL AN FDA APPROVED
ENDPOINT FOR
REGISTRATION

CANINE TRIAL BEGINNING
NOW WILL BE BASIS TO
MOVE TOWARD FULL
LICENCE



Pivotal Phase II Trial in Canine Cancer During 2019

KEY CLINICAL ENDPOINTS

- 1. Safety
- 2. Lack of Toxicity
- 3. Progression Free Survival
- 4. Regression
- 5. Administration by Owner at Home
- 6. Suppression of Tumour Markers





Clinical Outcomes To Commercial Opportunities



Pilot phase II (2017) in lymphoma trial in dogs Shows MPL effective at Progression-Free Survival Using poor formulation



Extensive Reformulation Program: 2018 - 2019 Establishes New, High Dose, Palatable Tablet Extensive Phase I Program During 2019 Establishes
Optimum Dosing for Canine Trials

PIVOTAL PHASE II TRIAL 2019



COMMERCIAL AND CLINICAL OUTCOMES

ACCELERATED HUMAN DEVELOPMENT PROGRAM



>\$2 Billion market for approved mTOR Drugs

Drug	Approved Indications	Company	2016 Sales (US\$m) ¹
Sirolimus	Transplantation	Pfizer	170
Rapalogues (Afinitor/Torisel)	Transplantation Renal Cell Carcinoma Breast Cancer Pancreatic Neuroendocrine Tumours Mantle Cell Lymphoma	Novartis/Pfizer	>2,000





Initial Market -Pets and Cancer

◆ 1 in 4 dogs die of cancer - 6 million diagnosed annually in US



- Pets are living longer (50% dogs > 10 yrs often die of cancer)
- ◆ Significant unmet need for new oncology drugs (US\$500m USD\$1b market)
- Side effects associated with products and treatments are limiting market growth (Monepantel comparatively has little or no side effects)
- Vet therapeutic market dominated by repurposed drugs already approved for use in humans and/or animals (= monepantel)
- Pet Insurance now commonplace
- Canines are a close reflection of human outcomes with MPL



PharmAust – current status and next steps

- Successful \$2.0m Rights Issue to existing shareholders completed in April 2019
- Oversubscribed placement primarily to Australian and Singaporean fund management institutions raises \$2.4m in October 2019
- Sufficient funds to complete Phase II in dogs as well as progression of the human trial, including further development of formulation & manufacture of additional tablets
- Successful micronisation and GMP tablet manufacture for dog trial
- Phase II trial on dogs with lymphoma commenced Sep 2019 at the University of Melbourne's U-Vet Werribee Animal Hospital
- Open label trial interim dossier to be prepared for Elanco Dec 2019
- Elanco Option Agreement covers veterinary uses only
- Canine success would open door to Human use a much larger market
- Epichem new CEO and state of the art laboratories a key to further growth and profitability





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