

Kazia Therapeutics

Q123 trading update

Q123 cash flows in line with our estimates

Pharma and biotech

1 November 2022

Price **\$0.83**
Market cap **\$13m**

ADR/Ord conversion ratio 1:10

Net cash (US\$m) at end-September 2022 3.4

ADRs in issue 15.11m

ADR code KZIA

ADR exchange Nasdaq

Underlying exchange ASX

Depository BNY

ADR share price performance



Kazia Therapeutics' Q123 cash flow report provided an update on the company's financial position and business progress. While the quarter was dominated by lead asset paxalisib hitting a roadblock (failing to graduate to stage two of the GBM AGILE study), the period was also marked by clinical progress across other serious indications such as pediatric brain cancers and brain metastases (BMs). With multiple studies expected to read-out in CY23, the next few quarters will be crucial for the company. Period-end cash balance of AU\$5.3m (c US\$3.4m) was supported by an AU\$3.7m equity injection and should be sufficient to extend the runway to end CY22 at current burn rates (A\$6.1m in Q123). Further support is expected from drawing down on the outstanding at-the-market funding facility. We anticipate the capital requirements to come down materially as the GBM study approaches completion in H2 CY23. Our estimates and valuation remain unchanged at US\$146.6m or US\$9.79 per basic ADR.

Year end	Revenue (US\$m)	PTP* (US\$m)	EPADR (US\$)	DPADR (US\$)	P/E (x)	Gross yield (%)
06/21	10.5	(3.1)	(0.25)	0.0	N/A	N/A
06/22	0.0	(14.6)	(1.08)	0.0	N/A	N/A
06/23e	0.0	(18.6)	(1.23)	0.0	N/A	N/A
06/24e	10.6	(16.8)	(1.11)	0.0	N/A	N/A

Note: *Converted at 1.45/US\$. Dividend yield excludes withholding tax. Investors should consult their tax advisor regarding the application of any domestic and foreign tax laws.

In addition to the company's cash flow position, management has highlighted several encouraging developments for Kazia over Q123, notwithstanding the somewhat disappointing announcement from the [Phase III GBM AGILE](#) study in August 2022. The international [expansion of the PNOC022 study](#), investigating paxalisib in combination with ONC201 in diffuse intrinsic pontine glioma (DIPG) and diffuse midline gliomas, hints at paxalisib's potential utility in these serious orphan diseases. Early clinical data in BMs, which appear to show a 100% response rate to paxalisib (albeit in a small patient population), further support the PI3K inhibitor's potential impact in the brain cancer. In addition, the [recent presentation of positive preclinical data](#) for paxalisib in melanoma is potentially encouraging for the drug's use in other, non-central nervous system oncology indications. Several of these studies are anticipated to report data in CY23 and we foresee a number of potential inflection points for the company in the coming quarters.

R&D related expenses (A\$4.3m) made up the majority of cash expenses during the period although with enrollment for the GBM AGILE study now complete, we expect future expenditure to be significantly reduced (management has disclosed that the remaining part of the trial is fully funded). Cash used in operations for the quarter was A\$6.1m and the company received A\$3.7m in proceeds from the issue of equity during the period, resulting in a net cash outflow (including FX movements) of AU\$2.1m. At current burn rates, the company has sufficient cash to last to end CY22, although management's internal projections suggest the funds lasting into Q1 CY23. In light of the current release, our forecasts and valuation of Kazia Therapeutics remains unchanged, pending disclosure of full accounts.

Business description

Kazia Therapeutics is a late-stage clinical pharmaceutical company with lead asset paxalisib (a PI3K inhibitor that can cross the blood-brain barrier, licensed from Genentech) in a pivotal study for GBM and in early-stage studies in childhood brain cancers, DIPG and AT/RT. The other asset is the Phase I drug EVT801, an inhibitor of VEGFR3.

Next events

Interim data from EVT801 Phase I study	H1 CY23
Phase II GMB AGILE top-line data	H2 CY23
Interim data from Phase II PNOC22	CY23

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