

Kinarus Therapeutics

Kinarus secures CHF1.5m financing

Financing update

Pharma and biotech

15 May 2023

Price CHF0.013
Market cap CHF17m

Gross cash (CHFm) at 31 December 2022 1.3
 Shares in issue 1,279m
 Free float 70.7%
 Code KNRS
 Primary exchange SIX Stock Exchange
 Secondary exchange N/A

Share price performance



Business description

Based in Switzerland, Kinarus Therapeutics is a clinical-stage pharmaceutical company focused on advancing lead candidate KIN001 in inflammatory, fibrotic and/or viral infection-related conditions. KIN001 is in Phase II studies for COVID-19 and the company plans to start Phase II studies in the coming months for wet age-related macular degeneration and idiopathic pulmonary fibrosis.

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Kinarus has [announced](#) that it has entered into a CHF1.5m strategic convertible loan agreement with ChaoDian (Hangzhou) Investment Management, an investment company based in China. The CHF1.5m subordinated loan has a three-year fixed term and does not bear any interest, and is convertible to Kinarus shares at a fixed conversion price of CHF0.01 per share. In addition to supporting Kinarus's financial position, the company indicates the loan agreement forms the basis for future discussions on the development and commercialisation of KIN001 for the treatment of idiopathic pulmonary fibrosis (IPF) in China. While discussions are preliminary and no deal or licensing arrangement can be assured at this stage, we are encouraged that KIN001 is attracting interest from potential partners and view this development as supportive of the underlying premise behind the potential anti-fibrotic and anti-inflammatory effects of KIN001.

Year end	Revenue (CHFm)	PBT (CHFm)	EPS (CHF)	DPS (CHF)	P/E (x)	Yield (%)
12/20	0.0	(1.5)	(0.31)	0.00	N/A	N/A
12/21	0.0	(4.9)	(0.00)	0.00	N/A	N/A

Note: 2020 accounts reflect statements of privately held Kinarus and 2021 statements represent pro-forma financials of Kinarus Therapeutics after the reverse merger transaction with Perfect Holding.

Kinarus in January [summarised the preclinical data](#) on lead candidate KIN001 (a proprietary combination of pamapimod and pioglitazone), demonstrating positive clinical activity in a mouse model of IPF both when used alone and in combination with existing approved IPF treatments, such as nintedanib or pirfenidone. This [21-day post-bleomycin injury mouse model data](#) showed pamapimod and pioglitazone each reduced lung fibrosis, although combining them resulted in a greater reduction in fibrosis compared with either drug alone. The combination of KIN001 with pirfenidone showed further decreases in lung fibrosis, suggesting KIN001 could be used as an additive therapy to existing approved treatments. Provided clinical benefit can be shown in human studies, we continue to view this as a significant opportunity for emerging therapeutics such as KIN001 in the IPF market, in light of the [high discontinuation rates](#) associated with existing approved IPF treatments. As a reminder, branded pirfenidone (Esbriet) and nintedanib (OFEV) recorded c US\$4bn in combined 2021 sales. We also note that KIN001 has already shown a favourable safety profile to date in human trials, as shown in the prior [KINETIC study](#) in COVID-19, where more than 130 patients received the drug for four weeks.

Kinarus has disclosed that its planned Phase II study in IPF will be a 52-week, double-blinded placebo-controlled trial assessing oral KIN001 on forced vital capacity, a common endpoint used in IPF studies, in 80 patients with IPF. The study will enrol patients already taking standard-of-care treatment (such as nintedanib or pirfenidone) as well as those taking neither drug. As is the case with Kinarus's lead KIN001 indication ([wet age-related macular degeneration](#)), we believe the company is exploring additional financing and partnering options before starting Phase II studies. The company [reported a CHF1.3m cash position](#) at year-end 2022 and expects to reports audited 2022 financial statements in the coming weeks.

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