

Redx Pharma

Voluntary delisting from AIM to go private

- Redx is proposing to delist from AIM and to re-register as a private limited company. A General Meeting seeking shareholder approval for the cancellation of ordinary shares, re-registration, and adoption of New Articles will occur on 19 April. If approved, which requires >75% of shareholders, the ordinary shares will be cancelled effective 1 May. Redx has already received irrevocable undertakings to vote in favour of the proposed delisting from certain shareholders representing c 84.64% of shares.
- The shift to being a private company is based on: (1) the limited liquidity and high share price volatility; (2) a market valuation that is not reflective of the drug development track record or future potential; (3) the constraints on raising capital and ease of future funding; (4) a more flexible regulatory regime enabling rapid corporate and strategic decision making; and (5) a reduction in administrative costs associated with an AIM listing.
- To briefly summarise Redx's portfolio, its lead ROCK asset is zelasudil, a selective ROCK2 inhibitor, currently in a Phase IIa trial in IPF (idiopathic pulmonary fibrosis), with topline data expected H124. These should confirm safety and tolerability, and could potentially include some early efficacy insights. Data will inform the design of a subsequent Phase IIb programme, expected to include broader interstitial lung diseases (ILDs).
- The second ROCK programme is RXC008, which recently started a Phase I trial. RXC008 is a first-in-class GI-targeted pan-ROCK inhibitor for fibrostenotic Crohn's disease which is expected to be used in combination with standard-of-care anti-inflammatories. The first part of the Phase I study includes healthy volunteers and will assess safety, with initial data expected by end-2024; the second part will include fibrostenotic Crohn's patients. Complete fibrosis reversal has been seen in a therapeutic preclinical model.
- Phase II data for RXC004 (zamaporvint), which has been earmarked for partnering, are also expected in H124. RXC004 is currently in development as a targeted therapy for Wnt-ligand dependent cancers and upcoming data in combination with a checkpoint inhibitor (CPI) will be key for partnering.
- Redx's core medicinal chemistry expertise to discover novel drug candidates provides optionality for pipeline expansion or future partnerships, and this was recently demonstrated with a third collaboration with Jazz Pharmaceuticals. Redx also has a partnership deal with AstraZeneca.

Trinity Delta view: The plan to delist from AIM is disappointing but, in our view, understandable. The shares have failed to reflect the value inherent in the business and, importantly, the future potential. As we have stated in previous notes, Redx has a solid track record of delivery, with six molecules in the clinic and four major partnering deals over the past five years. The announcement prompts us to suspend our Redx Pharma forecasts and valuation with immediate effect. For context, our last published valuation was £386m/\$463m.

2 April 2024

Price	18.50p
Market Cap	£71.96m
Primary exchange	AIM
Sector	Healthcare
Company Code	REDX
company couc	

Corporate client Yes

Company description:

Redx Pharma specialises in the discovery and development of small molecule therapeutics, with an emphasis on oncology and fibrotic diseases. It aims to initially progress them through proof-ofconcept studies, before evaluating options for further development and value creation.

Analysts

Lala Gregorek

lgregorek@trinitydelta.org +44 (0) 20 3637 5043

Philippa Gardner pgardner@trinitydelta.org +44 (0) 20 3637 5042



Philippa Gardner

Lala Gregorek

Franc Gregori

pgardner@trinitydelta.org +44 (0) 20 3637 5042

lgregorek@trinitydelta.org +44 (0) 20 3637 5043

fgregori@trinitydelta.org +44 (0) 20 3637 5041

Disclaimer

Trinity Delta Research Limited ("TDRL"; firm reference number: 725161), which trades as Trinity Delta, is an appointed representative of Equity Development Limited ("ED"). The contents of this report, which has been prepared by and is the sole responsibility of TDRL, have been reviewed, but not independently verified, by ED which is authorised and regulated by the FCA, and whose reference number is 185325.

ED is acting for TDRL and not for any other person and will not be responsible for providing the protections provided to clients of TDRL nor for advising any other person in connection with the contents of this report and, except to the extent required by applicable law, including the rules of the FCA, owes no duty of care to any other such person. No reliance may be placed on ED for advice or recommendations with respect to the contents of this report and, to the extent it may do so under applicable law, ED makes no representation or warranty to the persons reading this report with regards to the information contained in it.

In the preparation of this report TDRL has used publicly available sources and taken reasonable efforts to ensure that the facts stated herein are clear, fair and not misleading, but make no guarantee or warranty as to the accuracy or completeness of the information or opinions contained herein, nor to provide updates should fresh information become available or opinions change.

Any person who is not a relevant person under section of Section 21(2) of the Financial Services & Markets Act 2000 of the United Kingdom should not act or rely on this document or any of its contents. Research on its client companies produced by TDRL is normally commissioned and paid for by those companies themselves ('issuer financed research') and as such is not deemed to be independent, as defined by the FCA, but is 'objective' in that the authors are stating their own opinions. The report should be considered a marketing communication for purposes of the FCA rules. It has not been prepared in accordance with legal requirements designed to promote the independence of investment research and it is not subject to any prohibition on dealing ahead of the dissemination of investment research. TDRL does not hold any positions in any of the companies mentioned in the report, although directors, employees or consultants of TDRL may hold positions in the companies mentioned. TDRL does impose restrictions on personal dealings. TDRL might also provide services to companies mentioned or solicit business from them.

This report is being provided to relevant persons to provide background information about the subject matter of the note. This document does not constitute, nor form part of, and should not be construed as, any offer for sale or purchase of (or solicitation of, or invitation to make any offer to buy or sell) any Securities (which may rise and fall in value). Nor shall it, or any part of it, form the basis of, or be relied on in connection with, any contract or commitment whatsoever. The information that we provide is not intended to be, and should not in any manner whatsoever be, construed as personalised advice. Self-certification by investors can be completed free of charge at <u>www.fisma.org</u>. TDRL, its affiliates, officers, directors and employees, and ED will not be liable for any loss or damage arising from any use of this document, to the maximum extent that the law permits.

Copyright 2024 Trinity Delta Research Limited. All rights reserved.

More information is available on our website: <u>www.trinitydelta.org</u>

Trinity Delta, 80 Cheapside, London, EC2V 6EE, United Kingdom. Contact: info@trinitydelta.org