

Arecor Therapeutics

AT220 European launch marks first Arestat success

- The first product incorporating Arecor's Arestat technology, AT220, has been launched in Europe with first commercial sale triggering an unspecified milestone payment, with future royalty payments on sales. As the first commercial programme developed with Arestat technology, AT220 effectively validates the approach of employing Arecor's formulation expertise to create differentiated products with enhanced or superior product features and physical properties.
- AT220 is a novel and differentiated formulation of a biosimilar developed with an undisclosed "global pharmaceutical and healthcare company". The deal was struck in late-2017 and two milestone payments have already been received. Reassuringly, the first approval and launch has been achieved within the expected timelines. Neither the product nor financial terms have been made public. However, in our <u>December 2022 Outlook</u> we had identified several potential candidates for the reference biologic.
- A mix of in-house assets and partnered products provides Arecor with an attractive blend of value inflection points, in our view. The next most advanced partnered asset, Inhibrx's INHBRX-101 is in a registration-enabling trial for emphysema due to alpha-1 antitrypsin deficiency (AATD). The milestone received on transfer of RTU injectable AT307 to Hikma for further development suggests progress here is also being maintained. We also anticipate the execution of further licensing agreements (particularly for the Specialty Hospital products), as well as technology partnerships.
- Arecor's diabetes franchise underpins the bulk of the investment case and the clinical data to date from the two key diabetes assets, AT278 (ultrarapid, ultra-concentrated insulin) and AT247 (pump optimised insulin), suggest competitive profiles that are well suited to the changing diabetes landscape in terms of demographics and the technological advances seen with novel insulin pump delivery systems. The next diabetes-related catalyst is topline Phase I AT278 data in Type II diabetes in early 2024.

Trinity Delta view: Commercial launch of AT220 provides further evidence that Arecor's innovative and lower risk pipeline is making tangible progress. Management has created a broad and well-balanced pipeline of innovative products that potentially offer similar milestone and royalty streams to classic drug discovery plays, yet with lower development risks and in a less costly and more rapid manner. The proprietary diabetes programmes, AT278 and AT247, are particularly attractive and underpin a sizeable portion of our valuation. However, due to the commercial sensitivities surrounding the partnered programmes there is limited disclosure, which means the investment appeal of these as well as the partnered and in-house Specialty Hospital product portfolio tends to be overlooked, yet their likely future revenue streams are appealing and their collective contribution could be sizeable. Our valuation, based on conservative assumptions, is £176m, equivalent to 575p a share.

17 November 2023

Price	182.5p
Market Cap	£56.7m
Primary exchange	AIM
Sector	Healthcare
Company Code	AREC
Corporate client	Yes

Company description:

Arecor Therapeutics is a revenuegenerating clinical stage drug developer, with a well-balanced portfolio of inhouse and partnered programmes. Its proprietary Arestat formulation platforms result in enhanced products with lower development risks and less onerous regulatory approvals.

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 2023 Trinity Delta Research Limited. All rights reserved.

More information is available on our website: www.trinitydelta.org