



Source: Refinitiv

| Market data | |
|-------------------------|-------|
| EPIC/TKR | ARIX |
| Price (p) | 176 |
| 12m High (p) | 180 |
| 12m Low (p) | 59 |
| Shares (m) | 135.6 |
| Mkt Cap (£m) | 239 |
| NAV/share (p) | 274 |
| Premium/discount to NAV | -36% |
| Free Float | 71% |
| Market | Main |

Description

ARIX is a publicly listed biotechnology venture capital company. It provides an opportunity for all investors to participate in a balanced portfolio of diverse biotech innovation via a single stock. With a global portfolio of 16 companies and five IPOs achieved since launch in 2016, Arix is a dynamic and modern approach to life sciences venture capital investing.

Company information

Exec. Chairman Naseem Amin MD Jonathan Tobin MD Christian Schetter COO Robert Lyne Finance Director Marcus Karia

+44 20 7290 1050 www.arixbioscience.com

| Key shareholders | |
|------------------|-------|
| Directors | 0.1% |
| Acacia Research | 19.1% |
| Fosun | 8.2% |
| Ruffer | 6.1% |
| Takeda Ventures | 5.5% |
| Diary | |

Mar'21

| Analyst | |
|-------------|---------------|
| Martin Hall | 020 7194 7622 |

mh@hardmanandco.com

2020 results

ARIX BIOSCIENCE

Artios attracts Merck KGaA in deal up to \$7bn

Arix Bioscience (ARIX) is a listed global venture capital (VC) company that presents an opportunity for institutional and retail investors to participate in the high risk-return profile of early-stage biotech investing. ARIX minimises risk through a combination of an expert investment team and portfolio diversification. Along with its 2020 interim results, management provided the market with some aspirational targets for the next three years, which would see the NAV double to ca.£500m. Last month, Merck & Co declared the acquisition VelosBio for \$2.75bn, now Merck KGaA has announced a collaboration with Artios, potentially worth up to \$7bn.

- ▶ **Strategy:** ARIX sources investments from an established network and a strong scientific reputation. The portfolio is diversified by therapeutic area, treatment modality, stage of discovery/development and geography to balance the risk-reward profile. Value is realised when ARIX successfully exits its investments.
- ▶ Artios Pharma: Merck KGaA has announced a strategic collaboration with Artios to gain access to its novel DNA Damage Response (DDR) technology in oncology. Merck will pay \$30m in upfront and short-term milestones in return for rights for the exclusive development of drugs for up to eight targets. If Merck exercises this option, in addition to double-digit option fees, Artios would be eligible for milestones of up to \$860m per target, giving a potential headline deal size of \$7bn.
- ▶ ARIX holds 14.6%: ARIX is the biggest shareholder in Artios, with 14.6% of the outstanding share capital (12.7% on a fully diluted basis). Since inception, ARIX has invested in every tranche of Artios's two funding rounds, making a total investment of £13.8m. The holding value of this investment at 30 June 2020 was £19.1m, representing 9.4% of the gross portfolio value.
- ▶ Market reaction: As expected, the market responded favourably to this news, with the shares rising 18.5p, or 12%, adding £25m to the market capitalisation, reflecting the potential NAV uplift, additional credibility in the ARIX story, and a narrowing of the NAV discount. In the event that just one of the collaboration targets is successful, Merck will have to make a decision about whether it will be cheaper in the long term to own the whole of Artios.
- ▶ Investment summary: Two major deals in quick succession have demonstrated that ARIX has invested in interesting technologies that are attracting keen interest from the pharmaceutical majors. Having set itself some ambitious targets for the next three years, these deals suggest that the targets should be met comfortably, so the market should be narrowing the discount to NAV.

| Financial summary and valuation | | | | | | | |
|---------------------------------|-------|-------|-------|--------|-------|-------|--|
| Year-end Dec (£m) | 2017 | 2018 | 2019 | 2020E | 2021E | 2022E | |
| Change in FV of investments | 5.5 | 51.2 | -58.6 | *143.4 | - | - | |
| Other income | 1.9 | 1.3 | 0.5 | 0.2 | 0.2 | 0.0 | |
| Administrative expenses | -11.0 | -11.7 | -9.7 | -7.0 | -5.5 | -5.6 | |
| Operating profit/(loss) | -7.2 | 37.5 | -70.6 | 130.5 | -7.2 | -7.6 | |
| Profit/(loss) before tax | -7.7 | 38.2 | -69.9 | 130.7 | -6.9 | -7.3 | |
| Underlying EPS (p) | -9.5 | 27.2 | -49.9 | 88.9 | -4.7 | -5.0 | |
| Net cash/(debt) | 74.9 | 91.2 | 53.7 | 32.1 | 167.8 | 156.0 | |
| Equity issues | 105.1 | 83.5 | 0.0 | 0.0 | 0.0 | 0.0 | |
| NAV/share (p) | 152.3 | 200.4 | 149.1 | 274.0 | - | - | |

*Based on share prices and forex at close of business on 3 December 2020 Source: Hardman & Co Life Sciences Research



Artios-Merck research collaboration

Merck, ranked 22nd in global pharmaceuticals, and Cambridge-based Artios, have announced a three-year global strategic collaboration, whereby the two groups will conduct research on novel DDR targets in oncology. Merck has the right to opt into exclusive development and commercialisation of compounds for up to eight targets.

Since inception in 2015, Artios has developed expertise in DDR targets with the aim of targeting treatment in cancer more effectively and the promise of delivering precision medicine. The expectation is that this research collaboration will leverage the significant expertise and R&D resources of Merck with the potential of Artios's discovery platform of novel DNA repair nuclease inhibitors and targets that it has been developing.

For clarity, the deal does not include Artios's small-molecule ATR (ataxia telangiectasia and rad3-related kinase) inhibitor (in-licensed from MD Anderson) and Pol Theta (Pol θ) programmes, on which Artios will now focus its internal efforts.

Deal summary

- ► Artios will receive \$30m/£23m in upfront and near-term milestones.
- ▶ Merck has the right to opt into exclusive development of compounds on up to eight targets.
- ▶ If Merck chooses to exercise this option, it will pay double-digit option fees.
- ▶ Artios will be eligible to receive up to \$860m in milestones for each target, in addition to double-digit royalty payments on net sales of each product commercialised by Merck.
- ➤ Subject to specific conditions, Artios has opt-in rights for joint development and commercialisation with Merck KGaA for the programmes.

With the upfront payment and eight potential targets, each with potential milestones of up to \$860m, the deal has a headline value of up to \$7bn. In the event that just one or two of these targets is successful, in our opinion, Merck will face a decision about paying all the associated milestones to Artios, or simply take a view that it will be cheaper in the long term to buy the whole company.

Merck's view

Under the terms of the agreement, the companies will leverage Artios's proprietary nuclease targeting discovery platform to jointly identify multiple synthetic lethal targets for precision oncology drug candidates. During this joint research collaboration, Merck will contribute its significant expertise and resources in the field of DDR and will have exclusive worldwide rights to develop and commercialise selected therapeutics discovered under the collaboration.

"....Targeting DNA damage response has the potential to provide an important therapeutic option for many patients in need of new treatments. We are excited about working with Artios to develop novel precision oncology medicines as we move towards changing the current paradigm in cancer treatment. This collaboration further strengthens our leadership and expertise in the field and discovery of DDR inhibitors and complements our multiple innovative assets currently being evaluated in several Phase Land Phase II clinical studies..."

Source: Andree Blaukat

Head of Translational Innovation Platform Oncology & Immuno-Oncology at Merck

7 December 2020 **2**



Development of Artios

Artios Pharma Limited was incorporated (registration number: 09931309) on 29 December 2015 and has its registered office on a scientific incubator site at Babraham Research Campus, Cambridge, UK.

Artios is focused on the DNA damage response pathways, which are thought to selectively kill cancer cells through a concept called "synthetic lethality". DDR pathways contain three major components (some with overlapping functions): sensors, signal transducers and effectors. Cancer cells often rely on non-primary DDR pathways for survival, which allows them to evolve resistance to therapies. Inhibiting these DDR pathways results in cell death through synthetic lethality.

Artios has a strong management team with an exceptional track record in this technology, having played key roles in the discovery of Lynparza $^{\tiny\textcircled{\tiny \$}}$ (AstraZeneca). To date, the company has raised £90m through two funding rounds, which were both tranched. ARIX has participated in every tranche of both rounds, making a cumulative investment of £13.8m and giving it a 14.6% stake in the issued share capital (12.7% on a fully diluted basis) and making it the largest shareholder among a strong syndicate of investors, including the venture arms of four major pharmaceutical companies.

| Artios funding history | | | | | | | | |
|------------------------|------------|---------|-------|------------------|--------|--------------|----------|--|
| Date | Round | Tranche | Price | Capital increase | | ARIX holding | | |
| | | | | Shares | Raise | Shares | Invested | |
| Sep'16 | Series A | 1 | 71p | 13.03 | £9.3m | 2.53m | £1.8m | |
| Dec'17 | Series A | 2 | 75p | 12.00 | £9.0m | 2.46m | £1.9m | |
| Aug'18 | Series B | 'Α' | 75p | 9.00 | £6.8m | 2.67m | £2.0m | |
| Aug'18 | Series B | 'B' | 95p | 31.58 | £30.0m | 3.89m | £3.7m | |
| Dec'19 | Series B | 'B' | 95p | 36.84 | £35.0m | 4.53m | £4.3m | |
| Total | | | | *110.5 | £90.0m | | £13.8m | |
| ARIX | 'A' shares | | | | | 7.65m | | |
| ARIX | 'B' shares | | | | | 8.82m | | |
| ARIX | | | | | | 16.07m | | |

*includes Ordinary shares issued at nominal value Source: Hardman & Co Life Sciences Research

Another boost to IRR and track record

At the end of 2019, ARIX's stake in Artios was valued at £15.2m and it represented 10.2% of its gross portfolio value. At its interim results, ARIX raised the value of its holding by £3.9m to £19.1m on the back of progress with its ATR inhibitor and Pol0 programmes, which are in the process of being prepared for IND submissions, with clinical development due to start in 2021. The collaboration announced today between Artios and Merck does not include Artios's lead programmes, Pol0 and ATR inhibitors, for which Artios will retain all rights.

The Artios-Merck collaboration is likely to result in an uplift to ARIX's holding value in Artios when the NAV is recalculated at the end of the year. For example, another Artios shareholder, IP Group (IPO.L), stated that the value of its holding (11.7%) may increase in value by 45%-136% when it is reassessed on 31 December. The 18.5p rise in ARIX shares on this news added £25m to the market capitalisation, which reflects a potential uplift to year-end NAV and a further narrowing of the discount to NAV on greater appreciation of the ARIX story by the market.

This news is the second significant event, over and above any clinical news, at one of its portfolio companies in the past two months, adding further impetus to the IRR from its investments and its development of a strong track record.

7 December 2020 3



Disclaimer

Hardman & Co provides professional independent research services and all information used in the publication of this report has been compiled from publicly available sources that are believed to be reliable. However, no guarantee, warranty or representation, express or implied, can be given by Hardman & Co as to the accuracy, adequacy or completeness of the information contained in this research and they are not responsible for any errors or omissions or results obtained from use of such information. Neither Hardman & Co, nor any affiliates, officers, directors or employees accept any liability or responsibility in respect of the information which is subject to change without notice and may only be correct at the stated date of their issue, except in the case of gross negligence, fraud or wilful misconduct. In no event will Hardman & Co, its affiliates or any such parties be liable to you for any direct, special, indirect, consequential, incidental damages or any other damages of any kind even if Hardman & Co has been advised of the possibility thereof.

This research has been prepared purely for information purposes, and nothing in this report should be construed as an offer, or the solicitation of an offer, to buy or sell any security, product, service or investment. The research reflects the objective views of the analyst(s) named on the front page and does not constitute investment advice. However, the companies or legal entities covered in this research may pay us a fixed fee in order for this research to be made available. A full list of companies or legal entities that have paid us for coverage within the past 12 months can be viewed at $\underline{6}$. Hardman may provide other investment banking services to the companies or legal entities mentioned in this report.

Hardman & Co has a personal dealing policy which restricts staff and consultants' dealing in shares, bonds or other related instruments of companies or legal entities which pay Hardman & Co for any services, including research. No Hardman & Co staff, consultants or officers are employed or engaged by the companies or legal entities covered by this document in any capacity other than through Hardman & Co.

Hardman & Co does not buy or sell shares, either for their own account or for other parties and neither do they undertake investment business. We may provide investment banking services to corporate clients. Hardman & Co does not make recommendations. Accordingly, they do not publish records of their past recommendations. Where a Fair Value price is given in a research note, such as a DCF or peer comparison, this is the theoretical result of a study of a range of possible outcomes, and not a forecast of a likely share price. Hardman & Co may publish further notes on these securities, companies and legal entities but has no scheduled commitment and may cease to follow these securities, companies and legal entities without notice.

The information provided in this document is not intended for distribution to, or use by, any person or entity in any jurisdiction or country where such distribution or use would be contrary to law or regulation or which would subject Hardman & Co or its affiliates to any registration requirement within such jurisdiction or country.

Some or all alternative investments may not be suitable for certain investors. Investments in small and mid-cap corporations and foreign entities are speculative and involve a high degree of risk. An investor could lose all or a substantial amount of his or her investment. Investments may be leveraged and performance may be volatile; they may have high fees and expenses that reduce returns. Securities or legal entities mentioned in this document may not be suitable or appropriate for all investors. Where this document refers to a particular tax treatment, the tax treatment will depend on each investor's particular circumstances and may be subject to future change. Each investor's particular needs, investment objectives and financial situation were not taken into account in the preparation of this document and the material contained herein. Each investor must make his or her own independent decisions and obtain their own independent advice regarding any information, projects, securities, tax treatment or financial instruments mentioned herein. The fact that Hardman & Co has made available through this document various information constitutes neither a recommendation to enter into a particular transaction nor a representation that any financial instrument is suitable or appropriate for you. Each investor should consider whether an investment strategy of the purchase or sale of any product or security is appropriate for them in the light of their investment needs, objectives and financial circumstances.

This document constitutes a 'financial promotion' for the purposes of section 21 Financial Services and Markets Act 2000 (United Kingdom) ('FSMA') and accordingly has been approved by Capital Markets Strategy Ltd which is authorised and regulated by the Financial Conduct Authority (FCA).

No part of this document may be reproduced, stored in a retrieval system or transmitted in any form or by any means, mechanical, photocopying, recording or otherwise, without prior permission from Hardman & Co. By accepting this document, the recipient agrees to be bound by the limitations set out in this notice. This notice shall be governed and construed in accordance with English law. Hardman Research Ltd, trading as Hardman & Co, is an appointed representative of Capital Markets Strategy Ltd and is authorised and regulated by the FCA under registration number 600843. Hardman Research Ltd is registered at Companies House with number 8256259.

(Disclaimer Version 8 - Effective from August 2018)

Status of Hardman & Co's research under MiFID II

Some professional investors, who are subject to the MiFID II rules, may be unclear about the status of Hardman & Co research and, specifically, whether it can be accepted without a commercial arrangement. Hardman & Co's research is paid for by the companies, legal entities and issuers about which we write and, as such, falls within the scope of 'minor non-monetary benefits', as defined in the Markets in Financial Instruments Directive II.

In particular, Article 12(3) of the Directive states: 'The following benefits shall qualify as acceptable minor non-monetary benefits only if they are: (b) 'written material from a third party that is commissioned and paid for by a corporate issuer or potential issuer to promote a new issuance by the company, or where the third party firm is contractually engaged and paid by the issuer to produce such material on an ongoing basis, provided that the relationship is clearly disclosed in the material and that the material is made available at the same time to any investment firms wishing to receive it or to the general public...'

The fact that Hardman & Co is commissioned to write the research is disclosed in the disclaimer, and the research is widely available.

The full detail is on page 26 of the full directive, which can be accessed here: http://ec.europa.eu/finance/docs/level-2-measures/mifid-delegated-regulation-2016-2031.pdf

In addition, it should be noted that MiFID II's main aim is to ensure transparency in the relationship between fund managers and brokers/suppliers, and eliminate what is termed 'inducement', whereby free research is provided to fund managers to encourage them to deal with the broker. Hardman & Co is not inducing the reader of our research to trade through us, since we do not deal in any security or legal entity.

7 December 2020 4

