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# **AGENDA**

















### STRATEGIC REVIEW UPDATE

### Strategic review commenced in July 2023, including consideration of:

- Investment and realisation strategies;
- Capital allocation and shareholder returns policies;
- A tax-efficient wind-down of the Company

- Prevailing unfavourable market conditions and sustained discount to NAV lead to strategic review
- A number of options remain under active consideration and the Board is evaluating these in conjunction with its advisers
- A conclusion to the process will be announced soon





Peregrine Moncreiffe Robert Lyne

Non-Executive Chairman









Debra Barker MD Senior Independent Director

**Andrew Smith** Non-Executive Director

## ARIX BOARD OF DIRECTORS



### >> Peregrine Moncreiffe

Non-Executive Chairman

30+ years experience in investment, most recently Chairman of North Atlantic Smaller Companies Investment Trust alongside Director at Metage Funds



### Robert Lyne

Chief Executive Officer

Previously COO & General Counsel at Arix. 10 years' experience in listed venture capital and worked on 80+ VC financings and transactions. Broad experience of public company governance.



### >> Debra Barker MD

Senior Independent Director

More than 25 years' senior experience in major pharmaceutical companies. Currently NED of public biotech companies: Destiny Pharma, BergenBio and CureVac.



### >> Isaac Kohlberg

Non-Executive Director

Senior Associate Provost and Chief Technology Development Officer at Harvard University. Distinguished career protecting and commercialising IP for leading universities and research institutions



### Maureen O'Connell

Non-Executive Director

Global business executive, formerly Senior Independent Director of Sucampo Pharmaceuticals and prior to that CFO of Scholastic Corporation



### >> Andrew Smith

Non-Executive Director

Executive leader with over 30 years' international experience within the biopharma sectors. Current CFO of Santhera Pharma and previously CFO of Allecra Therapeutics and Sucampo Pharmaceuticals

**Arix Investment Committee Members** 

## **KEY TEAM MEMBERS**

Investment Team



Tassos Konstantinou

#### **Managing Director**

Previously an investment manager at IP Group, where he led several new investments and served on the board of privately held biotech companies



### Felix Breyer PhD

#### Associate

Felix completed his PhD in biochemistry at University College London and joined Arix in 2021 where he now holds several Board Observer roles



>>> Roy Bartel-Taverner

#### Analyst

Financial and scientific background having worked in equity research at UBS as well as being a trained medical doctor



### Laura Craig

#### **Investor Relations Officer**

Responsible for the shareholder relations and communications alongside internal operations of Arix



### Tom Davidson

#### **Finance Director**

Tom spent nearly a decade at Oakley Capital, a pan-European PE and VC group where he was most recently Finance Director with oversight of the internal finance and accounting operations teams

Finance & Operations

ARIX



### **INTERIM RESULTS HIGHLIGHTS 2023**

#### PERFORMANCE **CAPITAL PORTFOLIO** Clinical and operational NAV of £239m - 185p per share £101m cash at 30 June 2023 progress across portfolio's Arix co-led \$50m Series B round Disc Medicine raised total gross Increase in NAV of £13m for Evommune, investing £6.6m proceeds of \$220m providing from £226m at YE Dec 2022 runway into 2026 Harpoon announced \$25m £16.8m invested into new listed private placement in which Arix Closing of TwelveBio acquisition investments in the period participated with a £2.9m by Ensoma in February as well as \$50m Series B extension in May investment Value of listed portfolio took Ensoma's total round to increased to £68m (31 Dec Successful progress of Sorriso \$135m 2022: £45m) enabled £4.0m draw down on Reverse merger of Enliven with £13.5m invested into unlisted second tranche of Series A Imara to create Nasdag listed portfolio in the period company alongside \$165m raise Unrealised gain of £3.5m in POP Value of unlisted portfolio, against cost to date Interim data readouts across £66.2m (31 Dec 2022: £54.8m) the portfolio showing progress is on track

# GROSS PORTFOLIO PERFORMANCE IN THE PERIOD





# £36.5m deployed in the period

- £20.1m invested into Core Portfolio: Ensoma, Evommune, Harpoon, Sorriso and Disc
- Public Opportunities Portfolio (POP) accounted for £16.4m of investment

# £17.7m realised in the period

- Trading of listed assets including Disc and POP
- £6.2m of Ensoma equity in exchange for Twelve Bio

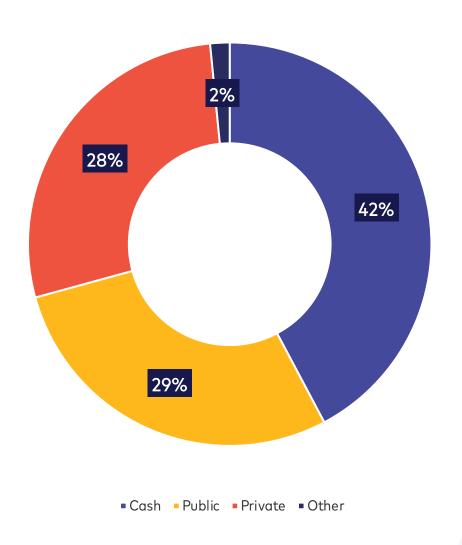
# £19.9m net upward revaluation

• Positive performance in listed portfolio



## NAV BREAKDOWN: CASH / PUBLIC / PRIVATE / OTHER

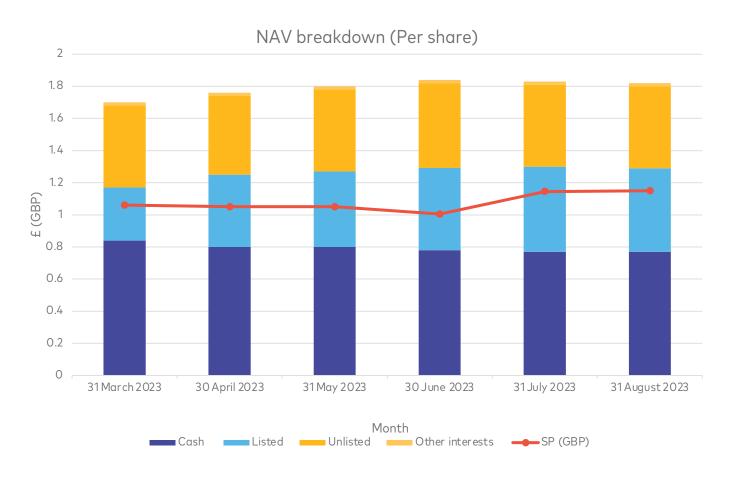
NAV: £239m\* (185p per share)



- >> 42% of the NAV is cash
- 29% of the NAV is in listed companies, including both POP and Core Portfolio companies valued at market prices
- 28% of the NAV is in our Core Portfolio of private companies, valued using The International PE and VC Valuation Guidelines
- 2% of the NAV is in other interests

## **NAV PER SHARE BREAKDOWN**

NAV per share performance against listed share price\*



- Persistent discount to NAV over prolonged period
- Current share price underpinned by cash holdings

# Financial Review NAV of £239m (185p); capital pool of £101m

Investment	31 Dec 2022 £m	Investment in Period £m	Realisations in Period £m	Impairment in Period £m	Change in Valuation £m	FX Movement £m	30 June 2023 £m	Equity Interest %	Fully Funded %
Core Portfolio									
Unlisted									
Artios	24.9	-	-	-	-	-	24.9	8.8%	8.8%
Depixus	8.2	-	-	-	-	(0.2)	8.0	14.2%	14.2%
Ensoma	7.5	6.2	-	-	-	(0.7)	13.0	6.0%	6.0%
Evommune	-	6.6	-	-	-	(0.2)	6.4	3.4%	3.5%
Harpoon Redeemable Preference Shares **	-	2.9	-	-	-	(0.1)	2.8	-	-
Sorriso	6.6	4.0	-	-	-	(0.4)	10.2	26.1%	26.1%
Twelve Bio	5.0	-	(6.2)	-	1.2	-	-	-	-
STipe	1.3	-	-	(1.2)	(0.1)	-	-	-	-
Amplyx	1.3	-	-	-	(0.3)	(0.1)	0.9	-	-
Unlisted total	54.8	19.7	(6.2)	(1.2)	0.8	(1.7)	66.2	-	-
Listed									
Aura	13.1	-	-	-	2.2	(0.6)	14.7	4.0%	4.0%
Disc Medicine	9.0	0.4	(1.1)	-	12.7	(0.3)	20.7	2.6%	2.6%
Harpoon Therapeutics	1.3	-	-	-	-	(0.1)	1.2	5.8%	5.8%
Imara / Enliven ***	7.9	-	-	-	1.9	(0.4)	9.4	1.4%	1.4%
Public Opportunities Portfolio	13.5	16.4	(10.4)	-	3.5	(0.9)	22.1	-	-
Listed total	44.8	16.8	(11.5)	-	20.3	(2.3)	68.1	-	-
Legacy Assets	0.1	-	-	-	-	-	0.1	-	-
Gross Portfolio	99.7	36.5	(17.7)	(1.2)	21.1	(4.0)	134.4	-	-
Other interests	3.0	-	(0.2)	-	-	(0.1)	2.7	-	-
Total Investments	102.7	36.5	(17.9)	(1.2)	21.1	(4.1)	137.1	-	-





# ARIX

# PORTFOLIO REVIEW

# PORTFOLIO COMPANY OVERVIEW

Breakdown of Core Portfolio\*

Portfolio Company	Therapeutic Area	Company Stage	Expected Milestone	Holding %	Holding £	
Private						
artios	Oncology	Clinical	Safety / efficacy data	8.8%	£24.9m	
Depixus	Genetic Diseases	Development	Proof of application	14.2%	£8.0m	
evommune	Immunology	Clinical	Safety / efficacy data	3.4%	£6.4m	
o ensoma	Genetic Diseases	Pre-clinical	IND Filing	6.0%	£13.0m	
SORRISO	Immunology	Clinical	CTA approval	26.1%	£10.2m	
Public						
aura	Oncology	Clinical	Efficacy data	4.0%	£14.7m	
disc)medicine	Haematology	Clinical	Safety / efficacy data	2.6%	£20.7m	
Enliven	Oncology	Clinical	Safety / efficacy data	1.4%	£9.4m	
HARPOON Therapeutics	Oncology	Clinical	Efficacy data	5.8%	£4.0m	

# CLINICAL AND OPERATIONAL HIGHLIGHTS MADE IN 2023

DNA DAMAGE RESPONSE	Oncology	<ul> <li>Initiated a Ph2 randomised trial for ART0380 plus Gemcitabine in Patients with Platinum Resistant Ovarian Cancer</li> <li>Launched a national project with IUCT-Oncopole, a cancer care, research and training centre in Toulouse, France, to overcome resistance to therapies for familial breast cancer</li> </ul>
्रः evommune	Immunology	> Raised \$50 million Series B in April with Arix investing £6.6 million. In June closed an additional \$7.5m into the round bringing the total raised to \$57.5m
<b>♦</b> e∩soma	Genetic Diseases	<ul> <li>\$85 million financing co-led by Arix for Ensoma along with acquisition of Twelve Bio</li> <li>Closed Series B extension financing, raising a further \$50 million, bringing the total round to \$135 million.</li> </ul>
SORRISO	Immunology	<ul> <li>Good progress preparing for initiation of Ph 1 clinical trial in 2023</li> <li>Progress enabled draw down on second tranche of Series B (£4.0m from Arix)</li> </ul>
aura	Oncology	<ul> <li>Positive interim Ph 2 safety and efficacy data of bel-sar in patients with early-stage Choroidal Melanoma with suprachoroidal (SC) administration. The interim data provides strong confidence to support the launch of the global Phase 3 trial which is on track to dose the first patient this year</li> <li>Bel-sar granted Fast Track Designation by the FDA for the treatment of Choroidal Metastasis</li> </ul>
disc)medicine	Haematology	<ul> <li>Raised \$220 million providing runway into 2026</li> <li>Positive initial data from ongoing Ph 2 trial of Bitopertin in patients with Erythropoietic Protoporphyria</li> <li>Initiation of Ph 1/2 study of bitopertin in patients with Diamond-Blackfan Anemia who have failed corticosteroid treatment</li> <li>Enrolment of patients in two separate Ph 1b/2 studies for DISC-0974, one in patients with NDD-CKD and one in patients with myelofibrosis and anaemia</li> <li>Fast Track Designation for MWTX-003 for the Treatment of Polycythemia Vera</li> </ul>
Enliven	Oncology	<ul> <li>Enliven entered the public market via reverse merger with Imara Inc. (previously Nasdaq: IMRA).</li> <li>Completed a \$165 million private placement with participation from new and existing investors</li> <li>Progress ELVN-001 and ELVN-002, through dose escalation in Ph 1 trials, with initial data for both programs expected in 2024</li> </ul>
HARPOON Therapeutics	Oncology	<ul> <li>Closed private placement to raise \$25 million in which Arix invested £2.9 million to extend cash runway into H2 2024</li> <li>Completed enrolment of patients in Ph 1 study of HPN217 in Relapsed/Refractory Multiple Myeloma</li> <li>Dosed first patients with small cell lung cancer (SCLC) in a Ph 1/2 trial of HPN328 in combination with atezolizumab (Tecentriq®)</li> </ul>

# MULTIPLE CLINICAL CATALYSTS EXPECTED IN H2 2023 & 2024

Company	Programme/Indication	Stage	H2 2023	H1 2024	H2 2024	2024-25
	ART0380 - Advanced or Metastatic Solid Tumours	Ph1			Ph2 data	
artios	ART6043 – Solid Tumour	Ph1	Ph1 Safety & Tolerability Data			
	ART4215 + Talazoparib – Breast Cancer	Ph1			Initial Ph2 Data	Clinical data from multiple studies
evommune	EVO101 – Atopic Dermatitis	Ph2	Ph2 Data			
	AU-011 - Choroidal Melanoma (SC)	Ph2	Ph2 data Initiation of Ph3 study		Interim Ph3 Data	New trial
aura	AU-011 –Choroidal metastasis	Ph2			Initiation of Ph2 study	initiations
	AU-011 – Non-Muscle Invasive Bladder Cancer	Ph1			Ph1 Data	
	Bitopertin – Erythropoietic Porphyrias (EPP and XLP)	Ph2	Ph2 BEACON Data	Ph2 AURORA do	ata	Progress discovery and
disc)medicine	Bitopertin – Diamond-Blackfan Anemia (DBA)	Ph2			Initial Ph2 DBA data	preclinical programmes
	DISC-0974 – Anemias of myelofibrosis (MF) and chronic kidney disease (CKD)	Ph1	Initial Ph1b/2 Data		Ph2 MF & CKD data	
Enliven	ELVN-001 - Chronic Myeloid Leukemia	Ph1		P	h1a Safety / Efficacy Data	
	ELVN-002 – Lung Cancer	Ph1		Р	h1a Safety / Efficacy Data	
HARPOON Therapeutics	HPN328 – Small Cell Lung Cancer	Ph1	Ph1/2 Data			
	HPN217 – Multiple Myeloma	Ph1	Ph1/2 Data			

## PUBLIC OPPORTUNITIES PORTFOLIO

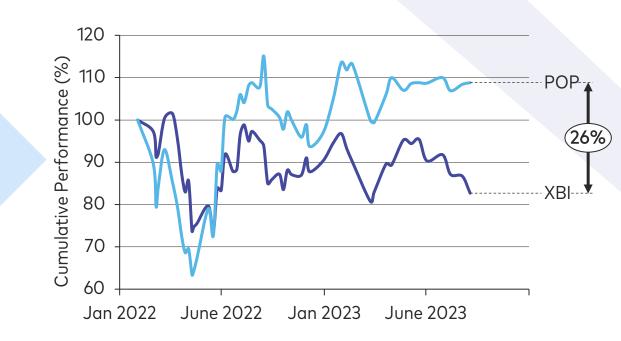
## Taking advantage of opportunities in the public markets

- >> Disconnect in public and private valuations of biotech companies observed in early 2022 lead to the creation of our Public Opportunities Portfolio, "POP"
- >> The investment criteria was the same as our private venture investments but offered a significant value play given unwarranted shift in sentiment against the public biotech markets
- >> Liquid positions in well-funded, clinical-stage biotech companies, with near term milestones, provided a dynamic opportunity

### POP outperforms XBI by 26% since portfolio inception

The mid-September 2023 value of active POP was broadly flat at a value of \$20.8m vs a cost of \$21.6m, which was favourable to the XBI having had a 17.3% drop since POP's inception

As of mid-September we maintained out performance by 26% returning 8.8%, including realised gains and losses, against a drop in the XBI of 17.3% in the same period





## OUTLOOK



Arix

- >> Strategy Review launched in July 2023
  - A number of options remain under active consideration
    - Significant cash balance of £101m at 30 June 2023



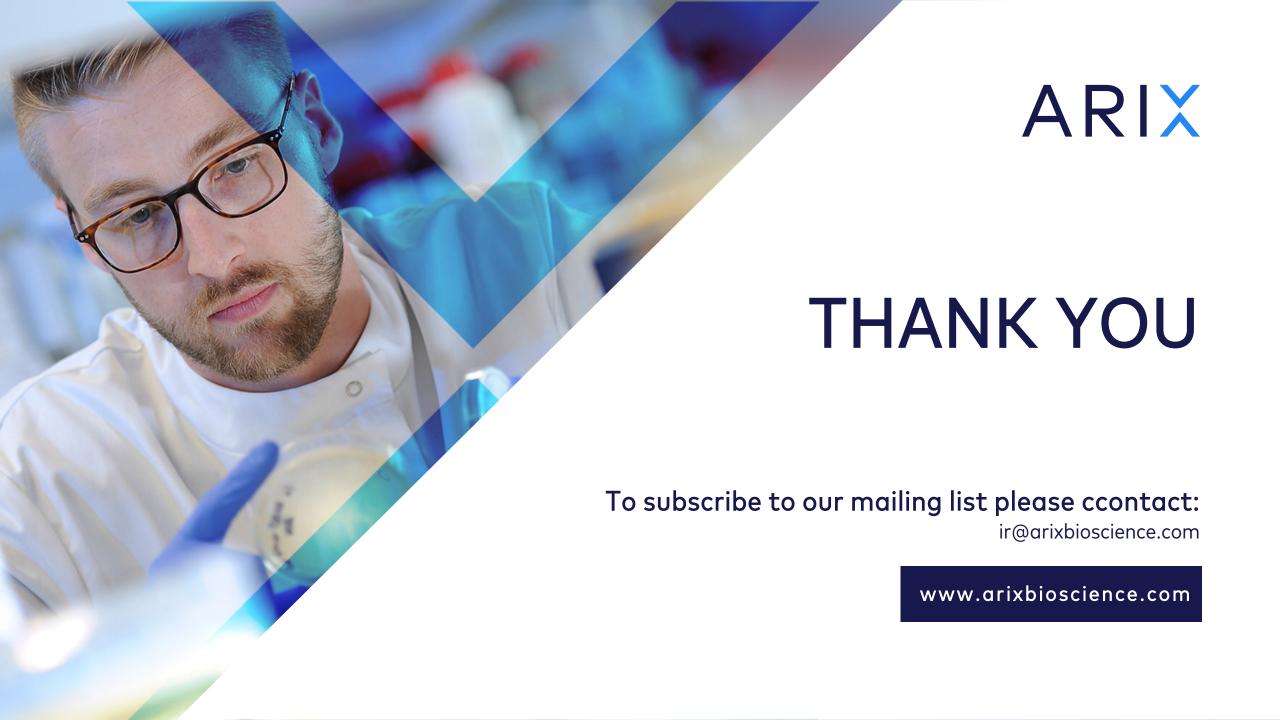
>> Well-funded portfolio with multiple clinical and preclinical catalysts expected over the next 12-18 months





# ARIX

Q&A



# **GLOSSARY OF ABBREVIATIONS**

ADC	Antibody Drug Conjugate			
NDA	New Drug Application			
IBD	Inflammatory Bowl Disease	<b>&gt;&gt;</b>	NDD-CKD	Non-Dialysis Dependent Chronic Kidney Disease
DDR	DNA Damage Response	<b>&gt;&gt;</b>	Bel-sar	Belzupacap Sarotalocan
NAV	Net Asset Value	<b>&gt;&gt;</b>	EPP	Erythropoietic Protoporphyria
NMIBC	Non-Muscle Invasive Bladder Cancer	<b>&gt;&gt;</b>	XLP	X-linked Protoporphyria
EMA	European Medicines Agency	<b>&gt;&gt;</b>	DBA	Diamond-Blackfan Anemia
GPV	Gross Portfolio Value	<b>&gt;&gt;</b>	MF	Myelofibrosis
IND	Investigational New Drug	<b>&gt;&gt;</b>	CKD	Chronic Kidney Disease
Ph1	Phase 1 Trial – studies the safety of drug	<b>&gt;&gt;</b>		
Ph2	Phase 2 Trial – studies the efficacy of drug	<b>&gt;&gt;</b>		
Ph3	Phase 3 Trial – studies the safety, efficacy & dosing	<b>&gt;&gt;</b>		
SCLC	Small Cell Lung Cancer	<b>&gt;&gt;</b>		
	NDA IBD DDR NAV NMIBC EMA GPV IND Ph1 Ph2 Ph3	NDA New Drug Application  IBD Inflammatory Bowl Disease  DDR DNA Damage Response  NAV Net Asset Value  NMIBC Non-Muscle Invasive Bladder Cancer  EMA European Medicines Agency  GPV Gross Portfolio Value  IND Investigational New Drug  Ph1 Phase 1 Trial – studies the safety of drug  Ph2 Phase 2 Trial – studies the efficacy of drug  Ph3 Phase 3 Trial – studies the safety, efficacy & dosing	NDA New Drug Application  IBD Inflammatory Bowl Disease  DDR DNA Damage Response  NAV Net Asset Value  NMIBC Non-Muscle Invasive Bladder Cancer  EMA European Medicines Agency  GPV Gross Portfolio Value  IND Investigational New Drug  Ph1 Phase 1 Trial – studies the safety of drug  Ph2 Phase 2 Trial – studies the efficacy of drug  Ph3 Phase 3 Trial – studies the safety, efficacy & dosing	NDA New Drug Application  IBD Inflammatory Bowl Disease

