

PHAXIAM

Investors call

March 21, 2024

PHAXiAM



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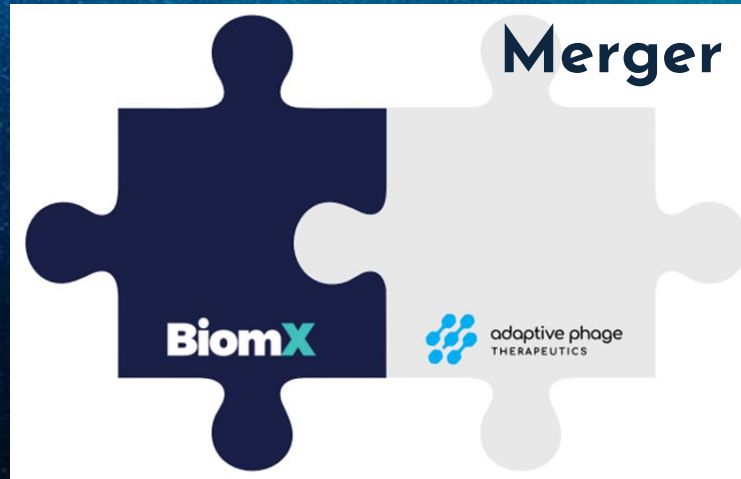
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Agenda

1. Market highlights
2. Business highlights
3. 2023 Financial Results
4. Conclusion

Reinforced interest in Phage Therapy

Major Commitment from Top Tier US investors in March 2024



Purchase agreement with a \$50m financing led by Top tier investor base, incl. Deerfield and Orbimed



Armata Pharmaceuticals has recently announced a \$35m refinancing with Innoviva

Growing attractiveness of the phage-therapy field to leading investors

An Ambitious Clinical Development Strategy

Target High-Value Indications

Severe Resistant Infections with High unmet medical needs
high mortality rate / high budget impacts → claim high pricing

Accelerate the Path to Global Registration in PJI

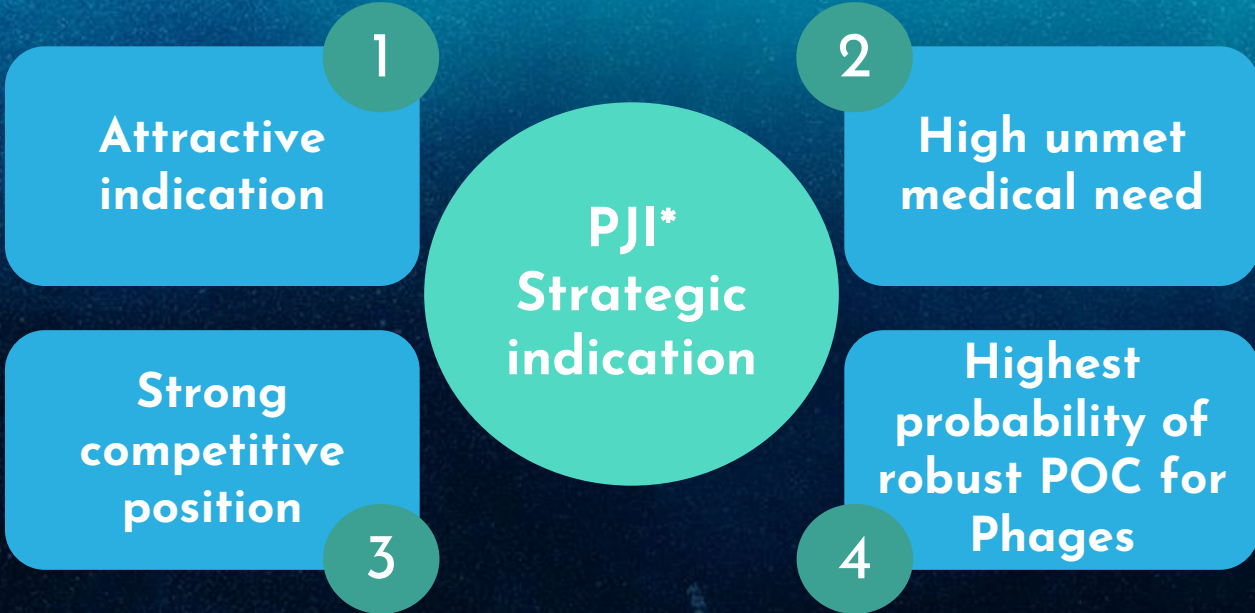
Launch the 1st global randomized Phase 2 study at international scale
Leverage on potential Early access pathway (after the phase 2 part)

Diversify Portfolio

Target several Life-Threatening Infections (Endocarditis, VAP*, ...)
Target several Virulent & Resistant Bacteria

PJI, a strategic indication for PHAXIAM

- Relatively High incidence: ~50-60K PJI* (US/EU5; 2027)
- Very High economic burden (cost ~ \$150K in US, €50-70K in EU) → high price
- Most advanced player in EU and US (APT stopped, Armata 18 months behind)
- Clear leadership → potentially 1st to Market

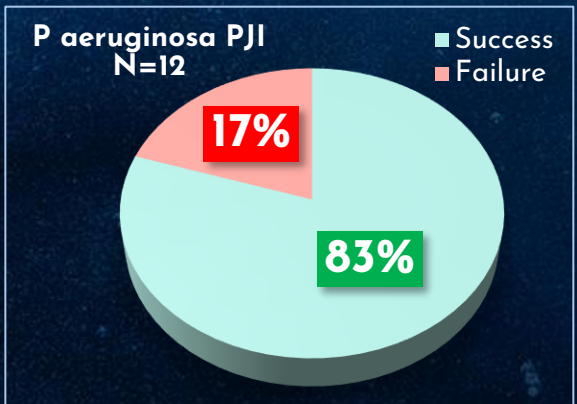
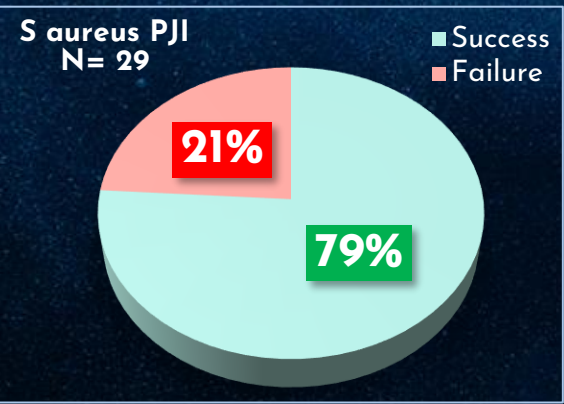


- Rare & devastating complication
- 50% failure rate with DAIR**
- High risk of re-infection (60%), amputation (~11%), mortality (25% at 5 years)
- Derisked by Real-life Compassionate experience
- Local route of administration

Unique and leadership position in the strategic PJI indication

Robust Real-life Clinical Data From ~100 Treated Patients

- STRONG SUPPORT from Regulatory Authorities & over 15 hospitals
- SEVERAL ROUTES OF ADMINISTRATION TESTED, including local, intravenous, nebulisation, ...
- 7 DIFFERENT INDICATIONS TREATED with a majority of PJI

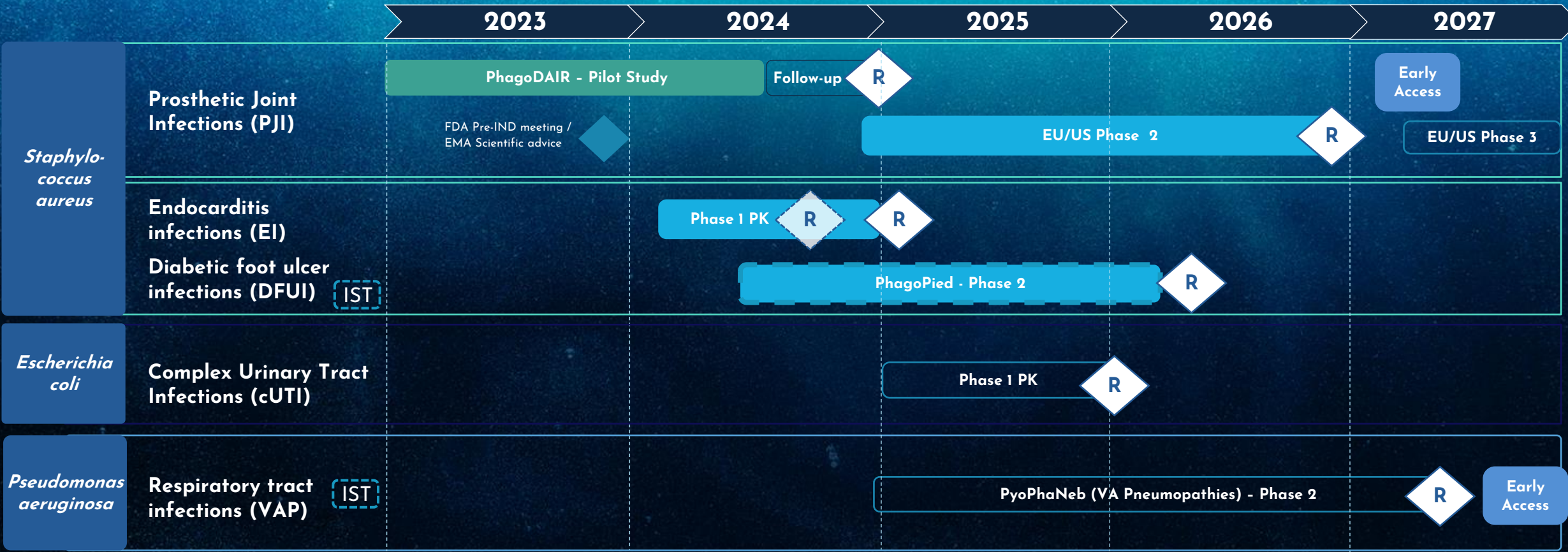


PROMISING CLINICAL ACTIVITY RESULTS (First 77 patients evaluated) 2018-2023

- EXCELLENT RESULTS observed in reported cases: safety + clinical benefit
- Several PUBLICATIONS

~80% CONTROL RATE of infection at 3months for PJI Patients (n=42)
VERY PROMISING Data in « hard to treat » population
(very severe infections - 2nd/3rd line antibiotics)

A Balanced Clinical Portfolio



IST: investigator sponsored study
Recruiting
To be initiated
Scheduled
R Clinical data

First Global (EU/US) PJI Phase 2 Study (GLORIA)

Global Integrated Phase 2 Proof of Concept Study
Multicentric, Randomized
to assess the Efficacy & Safety
of Phage Therapy in Patients (n=100) with Hip or Knee PJI
with open-surgery debridement (DAIR) in combination with antibiotics



Expected launch
Q1 2025



GLORIA Study Preparation

- ◆ **Positive and consistent feedback from both FDA (Pre-IND meeting) and EMA (Scientific Advice)**
 - Value of a clinical development in PJI
 - Non-clinical data and CMC capabilities support a clinical development in EU/US
 - Confirmation of the target population and the Standard of Care to be considered
 - Consistent guidelines for CMC and clinical outputs
- ◆ **Sites and countries selection initiated (FR, ESP, GER, NLD, ITA, SWE, UK, US)**
- ◆ **Regulatory submission to start in US and EU (June/July) then UK (September) → *Expected approval Q4 2024***

FPI expected in Q1 2025

Phase 1 PK in Endocarditis Infections

TRIAL	STATUS & PROGRESS
<p>Endocarditis Infections (EI)</p> <p>Staph. aureus</p> <p>Phase 1 PK</p>	<p>Demonstration for IV indications before Registration Study</p> <ul style="list-style-type: none">● EI patients having resistant infections in the cardiac chambers and valves● Phages administered intravenously (IV)● Expected demonstration of intravenous route of administration for EI and other IV indications before moving to registration study● Conducted in 5 French hospitals to recruit 12 patients <p><u>Key milestones</u></p> <ul style="list-style-type: none">● Agreement to initiate study obtained from ANSM in Q4 2023● First Patient-In expected in March/April 2024 / 4 sites opened

FY 2023 Financial Results - P&L (12 months)

<i>In thousands of euros</i>	FY2023 12 months	FY2022 12 months
Revenues	—	—
Net income from asset sale		24,351
Other income	1,326	6,647
Operating income	1,326	30,998
Research and development	(10,910)	(19,907)
General and administrative	(14,076)	(13,887)
Operating expenses	(24,986)	(33,793)
Operating income (loss)	(23,660)	(2,796)
Financial income	474	4,453
Financial expenses	(511)	(1,364)
Financial income (loss)	37	3,089
Income tax	208	(521)
Net loss	(23,488)	(227)

- €8.8m reduction in Operating expenses (-26%) yoy
- Sharp reduction in R&D expenses (-45%) mostly driven by the closing of Princeton operations and termination of ex-Erytech clinical & regulatory activities
- Integration of ex-Pherecydes activities as of H2 2023
- G&A stable yoy (+1%), including merger-related expenses
- Net loss of €23.5m for the FY 2023

Financial Position

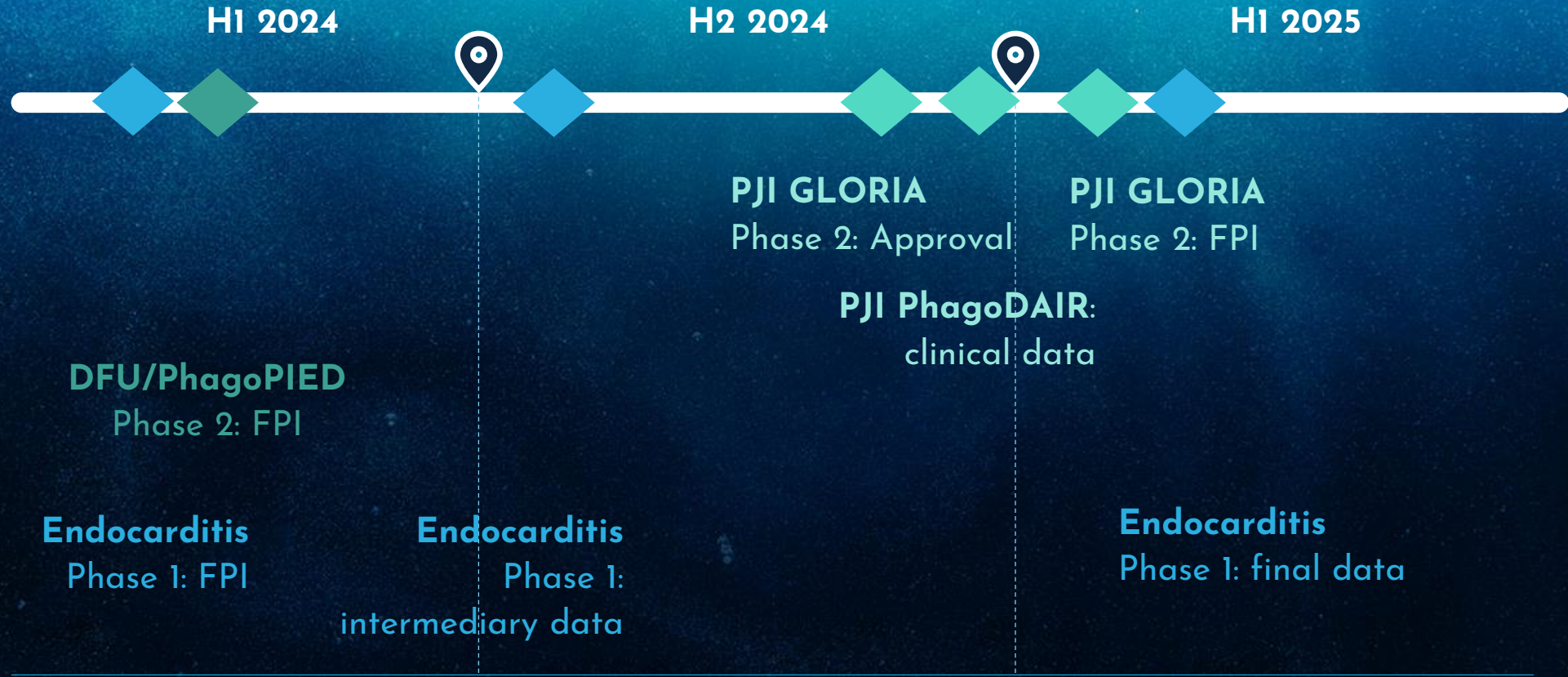
Cash position as of December 31, 2023: €10.5m

Cash runway into September 2024

Reviewing options to further extend cash runway



Expected Major Clinical Catalysts



**CLINICAL
&
REGULATORY**

Key messages

- ◆ **Growing attractiveness of the phage-therapy field to leading investors**
- ◆ **Strategic ambition upgraded to high-value indications in severe resistant infections (PJI, Endocarditis)**
- ◆ **Building on a very strong competitive position in PJI**
- ◆ **Several Clinical & Regulatory catalysts expected in the 12 coming months**

Thank you !

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