



Biophytis[®]

LIVE HEALTHIER LONGER



**A pioneering biotechnology company
focused on developing transformative
therapies that impact longevity**



Founded on pioneering
science from researchers at



Forward Looking Statements



This presentation contains forward-looking statements. Forward-looking statements include all statements that are not historical facts. In some cases, you can identify these forward-looking statements by the use of words such as «**outlook** », «**believes**», «**expects**», «**potential**», «**continues**», «**may**», «**will**», «**should**», «**could**», «**seeks**», «**predicts**», «**intends**», «**trends**», «**plans**», «**estimates**», «**anticipates**» or the negative version of these words or other comparable words. These forward-looking statements include statements regarding Biophytis' anticipated timing for its various BIO101 (20-hydroxyecdysone) clinical trials and expectations regarding commercialization. Such forward-looking statements are based on assumptions that Biophytis considers to be reasonable.

However, there can be no assurance that the statements contained in such forward-looking statements will be verified, which are subject to various risks and uncertainties including, without limitation, delays in patient recruitment or retention, interruptions in sourcing or supply chain, its ability to obtain the necessary regulatory authorizations, COVID-19-related delays, and the impact of the current pandemic on the Company's clinical trials. The forward-looking statements contained in this presentation are also subject to risks not yet known to Biophytis or not currently considered material by Biophytis.

Accordingly, there are or will be important factors that could cause actual outcomes or results to differ materially from those indicated in these statements. Please refer to the «Risk Factors» section of the Company's 2023 Full Year Financial Report available on BIOPHYTIS website (www.biophytis.com) and to the risks discussed in the Company's registration statement on Form F-1 and other reports filed with the Securities and Exchange Commission (the "SEC"). We undertake no obligation to publicly update or review any forward-looking statement, whether as a result of new information, future developments or otherwise, except as required by law.

Biophytis[®]

Mission & Vision

Advancing Longevity

by discovering new
targets and drug
candidates addressing
fundamental needs
in human aging





Our leadership team has a proven track record across drug discovery, development, and commercialization

Management Committee



Stanislas Veillet
Chief Executive Officer



Edouard Bieth
Chief Strategy Officer



Pierre Dilda
Chief Scientific Officer



Christophe Courtillat
Chief Financial Officer (Ad Interim)



Chiara Baccelli
Chief Pharmaceutical Officer



Waly Dioh
Chief Clinical Operations Officer



Jean Mariani
Chief Medical Officer

Cumulative years of experience

176y **126y**
in R&D in BioPharma

Nb of IND **20+**

Nb of Rx products
registered /
commercialized **40+**

Broad Experience in BioPharma and beyond



MONSANTO



Our Longevity Research Platform

A unique platform combining several technologies accelerated by AI for the discovery of new drug candidates impacting longevity

- Exploration and identification of the **fundamental mechanisms of aging** to develop 21st-century therapeutic solutions
- A platform leveraging **different synergistic technology modalities**
- **Systematic use of AI** through our partnership with Lynx Analytics.

BIO101 in Sarcopenia

The world's most advanced clinical program in sarcopenia

- **In sarcopenia, no approved treatment exists**, and it is estimated that up to 1 out of 4 people aged over 60 worldwide are affected ⁽¹⁾
- The phase 2 clinical trial demonstrated **BIO101's efficacy in improving mobility** in patients with sarcopenia ⁽²⁾
- For the first time, a phase 3 study in sarcopenia is planned in Asia and Europe

BIO101 in Obesity

A unique program targeting mobility in patients suffering from obesity

- **Obesity is booming, affecting 1 billion patients globally ⁽³⁾, and 96% report experiencing mobility issues ⁽⁴⁾**
- **A phase 2 clinical trial** is planned in Europe and Brazil to test **BIO101's efficacy in combination with GLP1**.
- **A License agreement** in LATAM has been signed with Blanver Pharma and the trial has obtained public **co-funding** in Brazil.



Our Longevity Research Platform

BIO101 in Sarcopenia

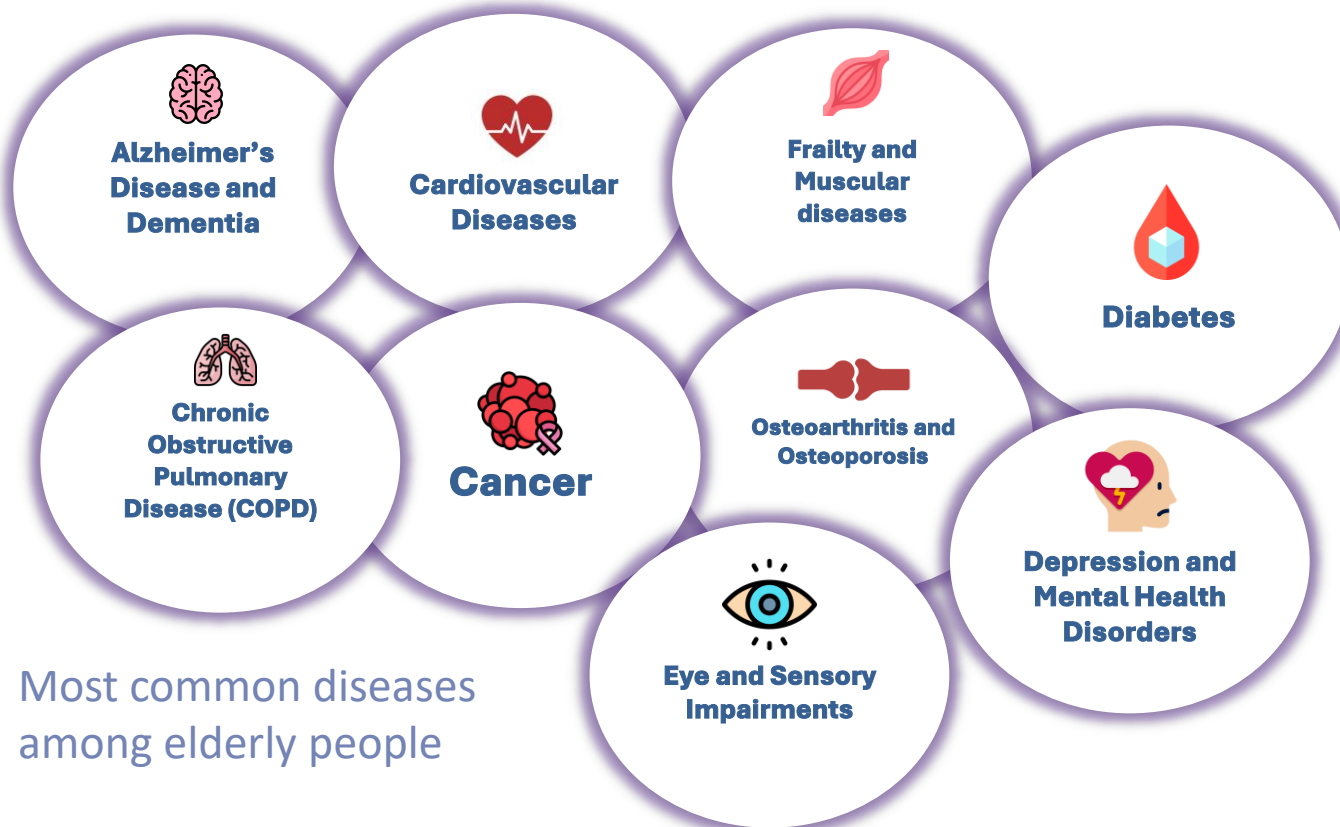
BIO101 in Obesity



Our research explores and identifies the fundamental mechanisms of aging to develop new therapeutic solutions...

Our Longevity Research Platform

...in partnership with
the most prestigious research academies in France



Most common diseases
among elderly people



Target ID & validation

Technology Assessment and Lead ID

Screening and Candidate selection

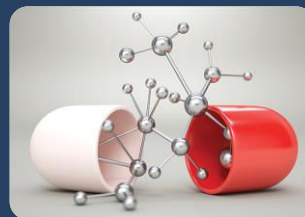
Exploration
and identification
of the
fundamental
mechanisms
of aging ⁽¹⁾



Biology of Aging: Targeting pathways like Mas-related G protein-coupled receptor (MasR), which is part of the renin-angiotensin system, Peroxisome proliferator-activated receptor pathway and key inflammatory and angiogenic signaling routes, including NF- κ B and AP-1 transcription factors.

Expertise across discovery and development with a unique tech-forward approach integrating **various drug Modalities:**

Small Molecules
Precision-targeted,
scalable, orally available



Other technologies
Programmable, tissue-
selective, adapted for
complex targets



We use 2 types of screening Methods :

Focused Screening:
Selection and testing of a
smaller group of compounds
believed to have activity
based on prior knowledge,
literature or known chemical
classes



**Virtual Screening
via AI-Driven Discovery:**
Advanced analytics to reveal
new lead candidates





Our Longevity Research Platform

BIO101 in Sarcopenia

BIO101 in Obesity

BIO101 : Our drug candidate is an oral small molecule potential for best in class acting on the root cause of muscle wasting



First in class molecule

- Validated mechanism of action
- Activation of MAS receptor (protective arm of the renin-angiotensin system)
- Regulation of smooth, cardiac and skeletal muscle metabolism



Ease of administration

- API manufactured at industrial scale
- Advanced CMC
- Oral capsules adapted to obese patients



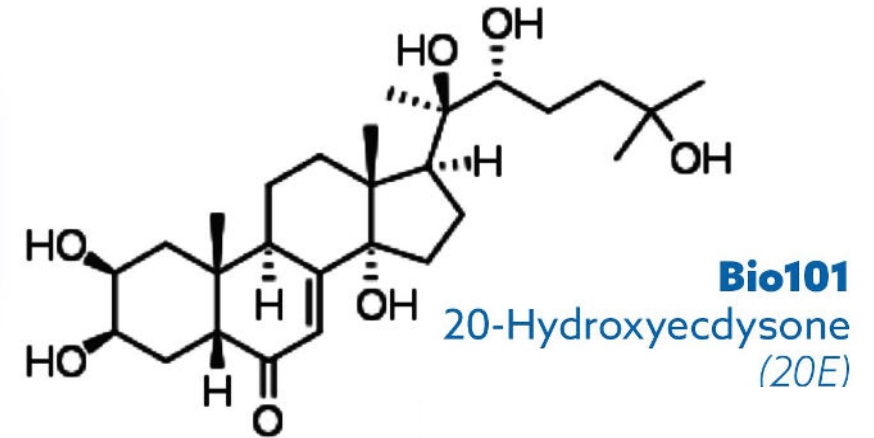
Excellent safety profile

- More than 275 patients exposed to BIO101 in clinical trials
- No major adverse event observed



US IND granted & Rock-solid IP

- 14 patent families granted in key countries
- Running until 2044
- 90% of world Pharma market covered



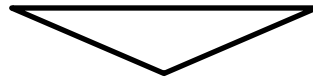
Focused development program for improving mobility

Candidate	Indication	Program	Preclinical	Phase 1	Phase 2	Phase 3	Regulatory	Market
BIO 101 20-hydroxyecdysone	Sarcopenia							
	Obesity							

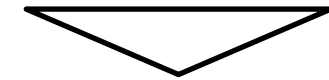
Sarcopenia is a disorder defined consensually by several consortia (FNIH, EWGSOP, SDOC and ESCEO), characterized by progressive and generalized loss of skeletal muscle mass and strength, reduction of physical performance associated with an increased risk of adverse events such as disability, poor quality of life and death



up to 1/4 people aged > 60 are affected⁽¹⁾



No pharmacological treatment has yet been approved, only limited options exist



Vitamin/dietary supplements

It may improve muscle strength and muscle mass
No Clinical evidence



Off label Drugs

The use of off label drugs is based on empiric practice
Data on sub population of large trials exist but not demonstrated

(1) Yuan 2023, Epidemiology of Sarcopenia, Metabolism

(2) Landi F, Liperoti R, Russo A, et al. Sarcopenia as a risk factor for falls in elderly individuals. Results for the iSIRENTE Study. *CI Nut.* 2012;31(5):652-658.

(3) Brown JC, Harhay MO, Harhay MN. Sarcopenia and mortality among a population-based sample of community-dwelling older adults. *J Cachexia Sarcopenia Muscle.* 2016;7(3):290-298

Drug	Company	Technology	Main Endpoints	Safety & Side effects	Administration route	Status
BIO101 (20 hydroxyecdysone)		MAS Receptor activator	High Dose showed promising results on gait speed (Phase 2)	Excellent tolerability profile and no serious adverse events	oral	Phase 3 to start (SARA)
 Bimugrumab, Trevogrumab, MYMD-1, Enobosarm						Development in sarcopenia halted after Phase 2
RJx-01		Combo Metformin + Galantamine	In a Phase 1b clinical trial, RJx - 01 showed promising results in improving muscle strength.	Generally safe and well tolerated with no reported treatment-related serious or severe adverse events	oral	Phase 1b finalized
OC-514		Small molecule AI-driven	Preclinical proof of concept	Favorable safety profile and pharmacokinetic parameters	oral	Phase 1 to start



Trial Design

A total of **233** patients randomized

BIO101 350mg BID

BIO101 175mg BID

Placebo

Study Termination or Extension



Journal of Cachexia, Sarcopenia and Muscle



	1 Sarcopenic obesity	2 Slow walkers (Gait speed $\leq 0,8$ m/s)	3 Mobility frailty (chair stand ≤ 2)	Total SARA-INT population
Gait speed (m/s) 400MWT Change from Baseline PP analysis	+0.06	+0.07	+0,09	+0,09
% Gait Speed improvement	+7%	+8%	+11%	+11%
P value	0,0037	0,0154	0,0037	0,008



BIO101 350mg BID *versus* placebo :
Gait Speed Improvement

+11%



SARA 31 : The first Phase 3 ever in sarcopenia

A phase 3 trial to evaluate efficacy & safety of BIO-101 versus placebo in older patients suffering from sarcopenia: an interventional, randomized double-blind placebo-controlled, clinical trial

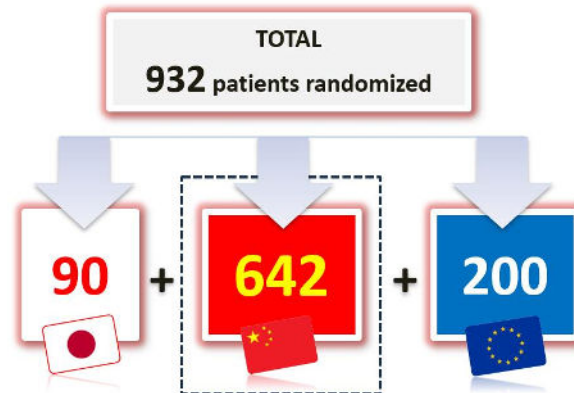
Population:

932 sarcopenic seniors treated for at least 12 months with BIO101 350 mg bid versus placebo

Primary endpoint:

Time (in days) to the first occurrence of **Major Mobility Disability (MMD)** assessed by the inability to complete the 400-meter walk test within 15 minutes

Secondary endpoints: Gait speed 4-m from SPPB, Handgrip Strength (HGS), Sarcopenia – QoL (SarQoL) Patient-Reported Outcome



Targeted Indication



Treatment of severe sarcopenia in older patients (≥65 years) who are at risk of mobility disability

Partnership Formation in Asia



- **Creation of a joint venture in Hong Kong**
- **Injection of US\$20 million by the consortium of investors**
- **Start of clinical operations in early 2026**

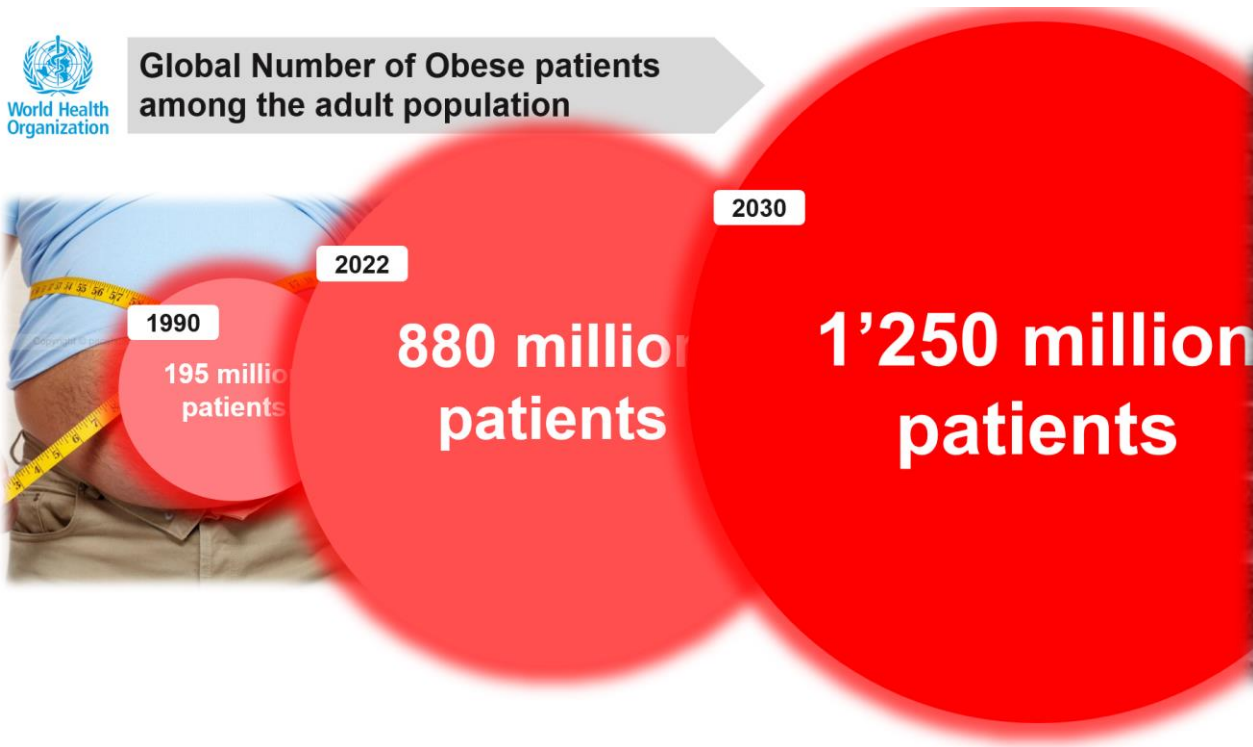


Pr Fielding, TOP Voice in sarcopenia

«In the SARA-INT trial, BIO101 showed a promising improvement of the 400MWT gait speed, a critical outcome for seniors at risk of mobility disability. These important findings reinforce the need to move to a confirmatory Phase 3 clinical program in older sarcopenic patients»



Global Number of Obese patients among the adult population

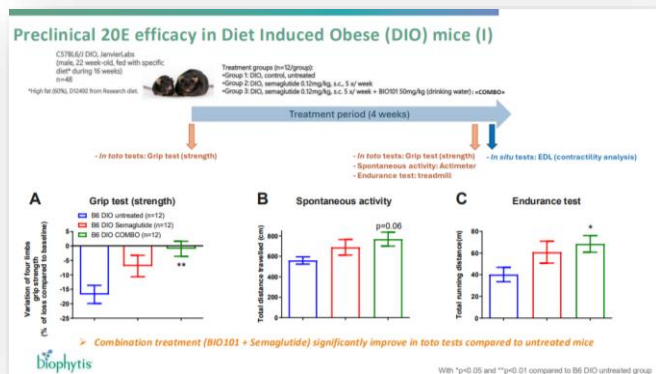
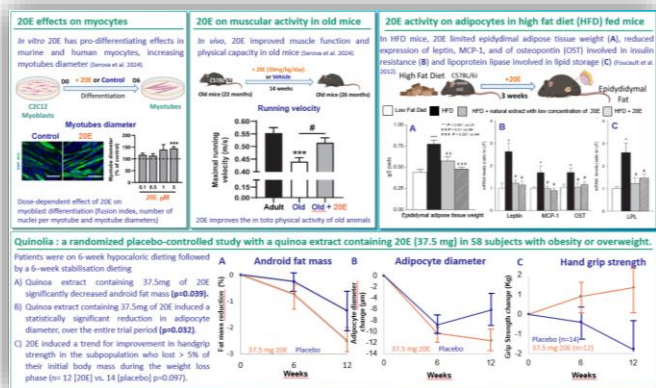


96% of obese patients have
impaired muscle strength⁽²⁾

(1) <https://www.morganstanley.com/insights/articles/weight-loss-medication-market-unstoppable-growth>

(2) Valenzuela et al. BMC Musculoskeletal Disorders (2020) 21:200

Strong Proof of concept ⁽¹⁾



A Phase 2 targeting muscle quality in obesity

Sample size : 164 patients

Target population:

Obese patients BMI ≥ 30 or overweight (BMI ≥ 27) with one or more weight-related sequelae (e.g. dysglycemia, hypertension) who will start treatment with semaglutide (a GLP-1 agonist).



Pr Cornier, President of Obesity Society US

« It is critical for us to study the safety and efficacy of BIO101 designed to reduce the risk of muscle mass loss »

1st Endpoint

Muscle strength (knee extension determined by dynamometry)



2nd Endpoints



6 minutes walking test



5 time Sit-to-Stand test



Hand grip test



Body composition (lean body & fat mass)

Partnership established

License agreement in LATAM signed with BLANVER PHARMA



Valued up to 108m€

Up to 50% of the local clinical operations in Brazil financed by public funding





Major upcoming milestones on our 3 R&D programs

Our Longevity Research Platform

ACHIEVED

- ✓ Partnership signed with LYNX Analytics in AI
- ✓ Protocol to identify new candidates in sarcopenia finalized
- ✓ New therapeutic targets exploration ongoing

TO COME

- Additional modalities to be included
- Partnership with Academic Research Centers in Aging / Longevity

BIO101 in Sarcopenia

ACHIEVED

- ✓ Trial readiness : work preparation with CROs
- ✓ CTA granted by EMA
- ✓ BIOPHYTIS and a Consortium of Investors including RONGHUI RENHE Life Technology Join Forces to Finance and Launch First-Ever Phase 3 Trial in Sarcopenia

TO COME

- Partnership Formation
- IND submission in China (NMPA)
- IND submission in Japan (PMDA)
- Study Start up
- First patient inclusion

BIO101 in Obesity

ACHIEVED

- ✓ BIO101 + GLP-1 animal studies results
- ✓ BLANVER License agreement signed
- ✓ IND granted by FDA
- ✓ CTA Granted by EMA
- ✓ Public funding obtained in Brazil
- ✓ Clinical Centers identified and selected in Europe, Brazil and US

TO COME

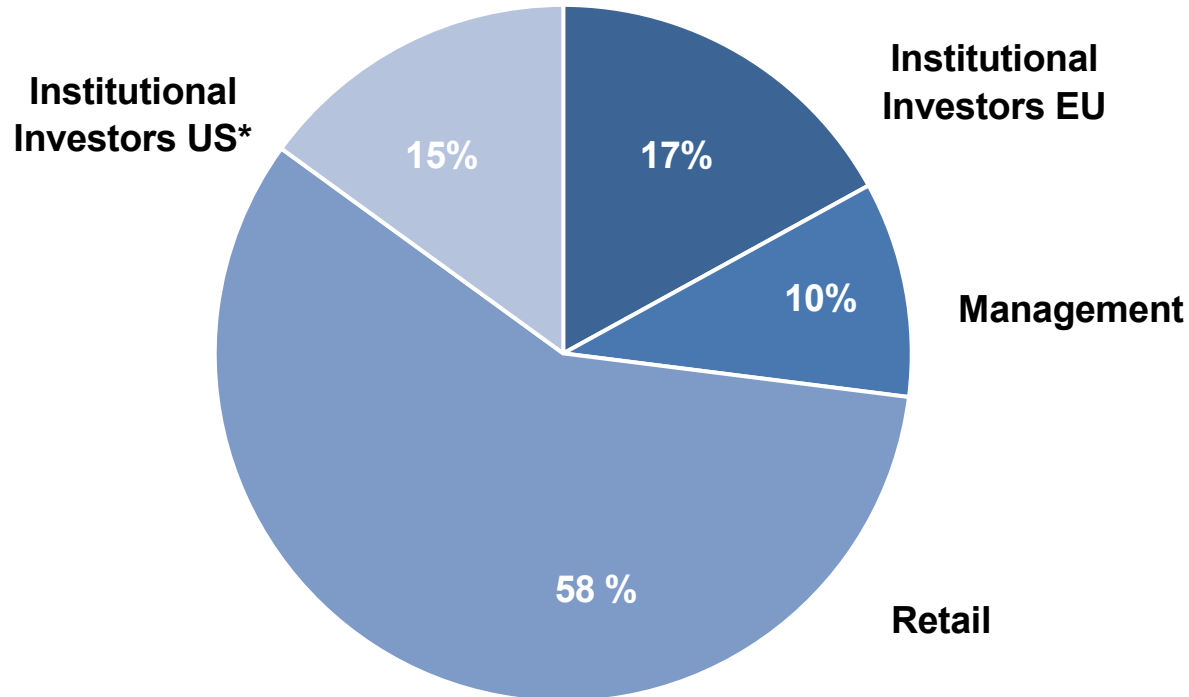
- IND submission in Brazil (ANVISA)
- Study Start up
- First patient inclusion



Our Pipeline: Transformative therapies that impact longevity

	Candidate	Indication	Program	Discovery	Lead Op	IND Enabling	Phase 1	Phase 2	Phase 3	Registration
Strategic Programs	BIO101	Sarcopenia	SARA							
	BIO101	Obesity	OBA							
	Target Discovery	Multiple Targets								
other projects	BIO101	Covid 19	COVA							
	BIO101	DMD	MYODA							
	BIO203	Dry AMD								
	BIO203	Stargardt disease								

Shareholding structure



Noteworthy investors: Atlas, BlackRock and Armistice
Number of shares: 29,766,653 (September 30, 2025)



Key financials

Listing Euronext (ALBPS) and US market (OTC)

- **Cash position:** €78K (December 31st, 2024)
- **Additional cash in FY2025 :**
 - €2.5m in January 2025 (cash contribution)
 - €2.6m in March 2025 (private placement)
- **Debt conversion:** €4m commitment from BlackRock and Atlas



Analyst Coverage



Nicolas Pauillac

Jamila El Bougrini



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by discovering new
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The world's most advanced
clinical program in sarcopenia

BIO101 in Obesity

A unique program targeting
mobility in patients suffering
from obesity