



Corporate Presentation

September 2024

Forward-looking statements

This corporate presentation (the "Presentation") has been prepared by ADOCIA S.A. (the "Company" and, together with its subsidiary, the "Group") and is provided for information purposes only. It is not for promotional use. References herein to the Presentation shall mean and include this document, any oral presentation accompanying this document provided by the Group, any question and answer session following that oral presentation and any further information that may be made available in connection with the subject matter contained herein (together with the information, statements and opinions contained in this Presentation, the "Information").

The Information is provided as of the date of the Presentation only and may be subject to significant changes at any time without notice. The Group does not undertake any obligation to update the Information.

The Information has not been independently verified. Subject to applicable law, none of the Group or its advisors accepts any responsibility whatsoever and makes no representation, warranty or undertaking, express or implied, is made as to the fairness, accuracy, completeness or appropriateness of the Information

The Presentation contains information on the Group's markets and competitive position, and more specifically, on the size of its markets. This information has been drawn from various sources or from the Group's own estimates. Investors should not base their investment decision on this information.

The Presentation does not purport to contain comprehensive or complete information about the Group and is qualified in its entirety by the business, financial and other information that the Company is required to publish in accordance with the rules, regulations and practices applicable to companies listed on Euronext Paris including in particular the risk factors described in in the most recent Company's Universal Registration Document filed with the French Financial Markets Authority (Autorité des Marchés Financiers), in any other periodic report and in any other press release, which are available free of charge on the websites of the Group (www.adocia.com) and/or the AMF (www.amf-france.org).

The Information contains certain forward-looking statements. These statements are not guarantees of the Group's future performance. These forward-looking statements relate without limitation to the Group's future prospects, developments and marketing strategy and are based on analyses of earnings forecasts and estimates of amounts not yet determinable. Forward-looking statements are subject to a variety of risks and uncertainties as they relate to future events and are dependent on circumstances that may or may not materialize in the future.

Forward-looking statements cannot, under any circumstance, be construed as a guarantee of the Group's future performance as to strategic, regulatory, financial or other matters and the Group's actual performance, including its financial position, results and cash flow, as well as the trends in the sector in which the Group operates, may differ materially from those proposed or reflected in the forward-looking statements contained in the Information.

Even if the Group's performance, including its financial position, results, cash-flows and developments in the sector in which the Group operates were to conform to the forward-looking statements contained in the Presentation, such results or developments cannot be construed as a reliable indication of the Group's future results or developments. The Group does not undertake any obligation to update or to confirm projections or estimates made by analysts or to make public any correction to any prospective information in order to reflect an event or circumstance that may occur after the date of the Presentation.

The Information does not constitute an offer to sell or subscribe or a solicitation to purchase or subscribe for securities, nor shall there be any sale of these securities in the United States or any other jurisdiction in which such offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such state or jurisdiction. No public offering of securities may be conducted in any member state of the European Economic Area (including France) prior to the publication in the relevant member state of a prospectus that complies with the provisions of Regulation 2017/119.

All persons accessing the Information are deemed to agree to all the limitations and restrictions set out above.

Adocia at a glance

- Mission: Development of innovative formulations of approved peptides and novel cell therapy approaches for diabetes and obesity
- Business model: Licensing-out/Partnering our products and technologies after proof-of-concept in animal and/or human model
- 2 partnerships with large pharma players
 - Tonghua Dongbao: BC Lispro (Ph. 3) out-licensed for Asia
 - Sanofi: Exclusive rights on M1Pram (Ph. 2), in view of a global partnership
- Assets: 3 clinical stage specialty products (Ph. 1 to 3) and 4 proprietary technology platforms, supporting a balanced pipeline
- **€10.3m** cash on-hand as of June 30, 2024

Highly experienced management team













Rosy Eloy
MD
Chief Medical Officer
Inserm Geistlich
NAMSA



Jérémy Benattar
PharmD, Eng
Head of Marketing Strategy

Otsuka
AstraZeneca

- Co-founded by Gérard, Olivier and Rémi Soula in 2005
- IPO on Euronext-Paris in 2012
- 80+ employees mostly dedicated to R&D incl. c. 35 PhDs/MDs
- Located in Lyon, France



Business Strategy: solid partnerships and promising projects

Partnerships:

Agreements with insulin leader Pharma

Proprietary Products to be licensed:

Market potential: multibillion \$USD

Proprietary Technology Platforms:

Business Model: Feasibility study before partnership



M1Pram

Exclusive negotiation rights in view of a global partnership

sanofi

BC Lispro EU/US Ph. 3 ready

AdoShell Islets
Animal POC

AdOral Sema Animal POC

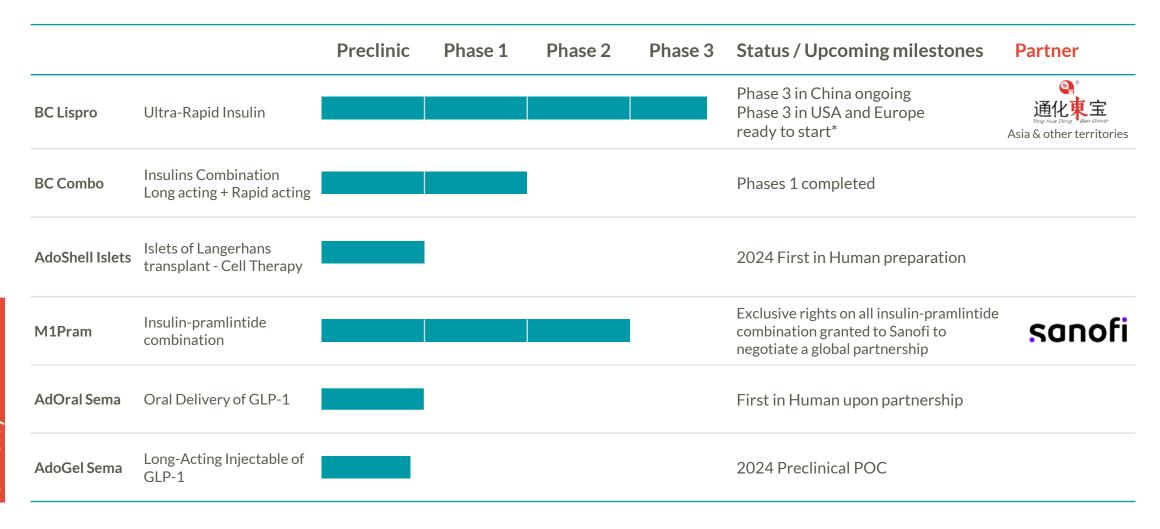
BioChaperone®

AdoShell®

AdOral[®]

AdoGel®

A diversified specialty products pipeline



BC: BioChaperone®; **Lispro**: insulin lispro; **BC Combo**: BC insulin glargine insulin lispro; **M1**: A21G human insulin; **Pram**: pramlintide; **Sema**: semaglutide, **POC**: Proof of Concept * upon partnership signature

Adocia's partnering models are based on proprietary technology platforms with strong IP strategy

BioChaperone[®]

- Pharmaceutical excipient
- Forms a complex with therapeutic peptides
 - → Accelerates absorption
 - → Protects against enzymatic degradation
 - → Improves solubility
 - → Improves stability
- Value:
 - → Improves proteins/peptides efficacy
 - → Combines hormones in one single product

AdOral[®]

- Pharmaceutical excipient
- Enhances peptides oral route of administration
 - → Improves bioavailability
- Value:
 - → Avoids injections and provides unique competitive advantage vs. injectables peptides
 - → Life cycle management of existing injectable products
 - → Avoids large scale sterile manufacturing of injectables

AdoShell®

- Hydrogel scaffold
- Cells encapsulation for cell therapies
 - Protects grafted cells from immune system rejection
 - → Ensures retrievability and easy surgical implantation
- Value:
 - → Avoids immunosuppressive therapies associated to cell therapies

AdoGel®

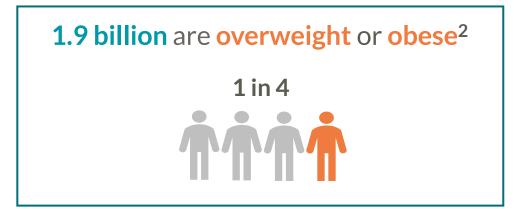
- Biomaterial
- Long-acting drug delivery of small molecules or biologics
 - → Release from 1 to 36 months, without initial burst, for local or systemic use
- Value:
 - → Avoids repetitive drug administrations
 - → Improves compliance

Adocia has a strong track record in preclinical and clinical development up to Phase 3 of specialty products based on innovative technologies



Diabetes and **Obesity**: worldwide chronic pandemics





Diabetes is closely linked to obesity

• In the US, 65% of adults with type 1³ diabetes, and 85% of type 2⁴ are overweight or obese, making **DIABESITY** a new pandemic

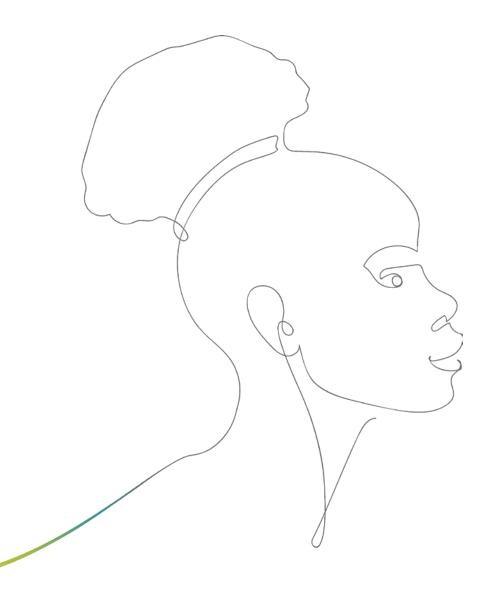


Adocia is developing unique formulations of these key hormones to improve diabetes and obesity treatments

- 1. IDF Atlas, 10th Edition, 2021
- 2. WHO
- 3. Conway et al, Diabetes Med 2010 April; 27(4):398-404. BMI>25, Data for 2004-2007 period
- 4. Epidemiology of Obesity and Diabetes and Their Cardiovascular Complications





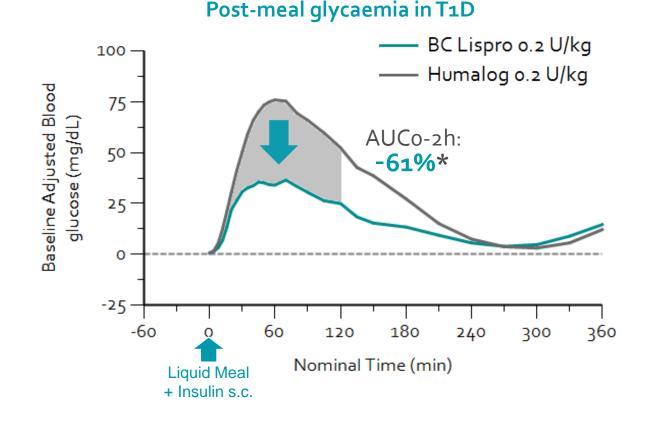


BioChaperone® Lispro

Ultra-Rapid Acting Insulin for a tighter glycemic control

BC Lispro has competitive advantages in the Ultra-Rapid Insulin class

- Better efficacy profile for less hyperglycemia and less hypoglycemia ("Faster-in" / "Fasterout") vs. comparators
- Good tolerance for optimized daily use
- Range of strengths (U100 & U200), adapted to pump miniaturization and patients' requirements



The combination of a faster release with a good local tolerance will put BC Lispro in a strong position to compete with other Ultra Rapid Insulins



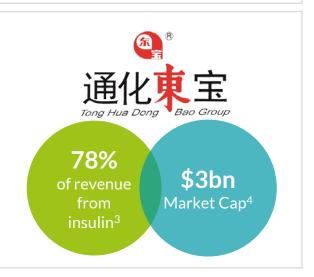
BC Lispro in phase 3 program in China with Tonghua Dongbao

Market

Insulin and analogs in China: c. \$4 billion

Partnership with Tonghua Dongbao (2018)

- Licensed for development & commercialization for China and other Asian territories²:
 - \$10m upfront
 - \$ 5m milestone 1st patient on the Phase 3 trial (Q2 2022)
 - \$30m development and approval milestones
 - Next milestone: end of Phase 3 \$10m
 - Double-digit royalties
- Global supply agreement for GMP lispro for the development and commercialization of BC Lispro outside licensed territories



Project status

✓ Phase 3 clinical program **on track** in China. Completion expected in H2 2024

Phase 3-ready for US/EU, with green light from FDA/EMA

BC Lispro will be the next generation of mealtime insulin in China



2. China and other territories (excluding US, EU, Japan)



BC Combo, glargine-lispro premix

The best insulin premix

BC Combo, a unique combination of glargine and lispro, the gold standards of basal and prandial insulins

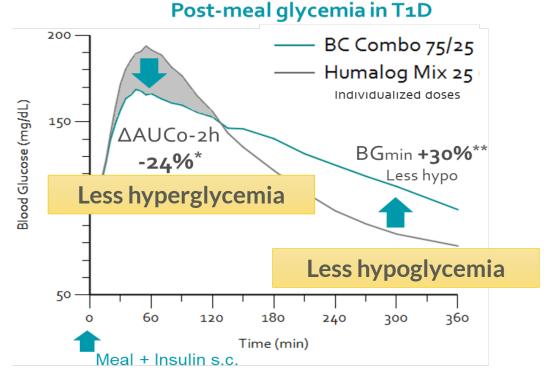
BioChaperone® Combo

1 product
Once or twice a day
U200 fixed ratio (150 U/mL glargine - 50 U/mL lispro)

- ✓ Simple
- ☑ Affordable
- ☑ Limited number of injections
- ☑ Improved glycemic control

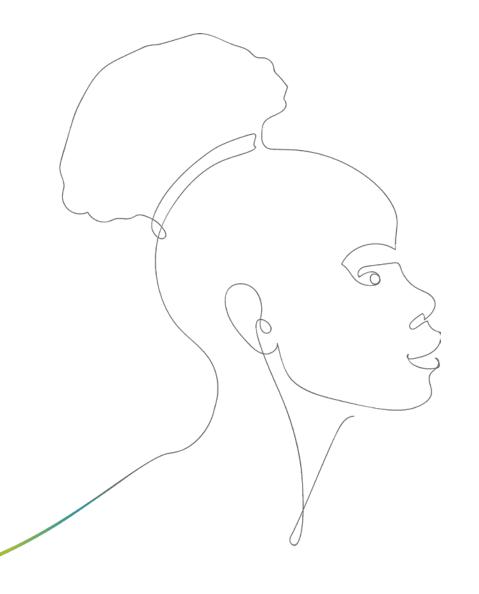
Addressing the premix market¹

\$5bn



Trial in 28 people with T1D (NCT#02514954); *p=3.10⁻³;**p=8.10⁻³

BC Combo offers better performance than Humalog® Mix, current standard of care on the premix market



M1Pram Insulin & Amylin analogs combination

Breakthrough treatment to address the unmet medical need of overweight patients living with diabetes

Agreement with Sanofi on M1Pram

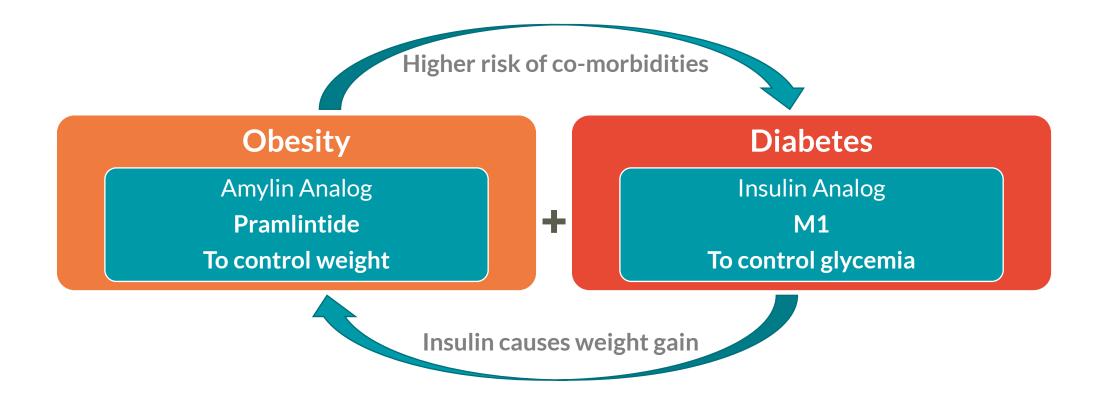


On July 5, 2023,

- Adocia received €10m payment from Sanofi in exchange of an exclusive negotiation right on M1Pram
- Sanofi commitment will contribute to accelerate M1Pram clinical development
- Sanofi agreement covers any programs related to insulin / pramlintide combinations

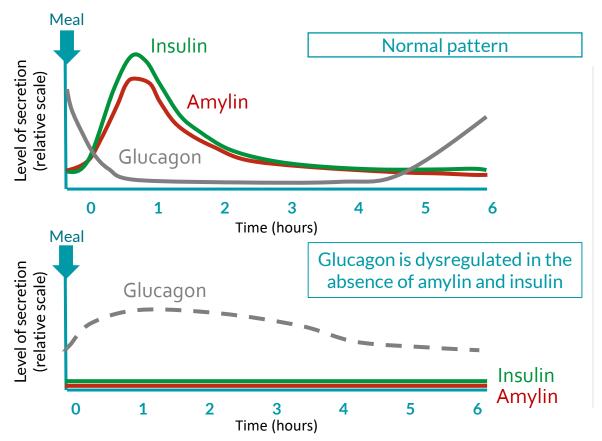
Adocia and Sanofi objective is to establish a global partnership on M1Pram

M1Pram, a bi-hormonal therapy to treat people with obesity and diabetes



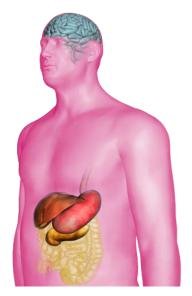
Obesity is a main concern for patients under intensive insulin therapy, affecting 43 million patients globally

Amylin is missing in people with insulin-dependent diabetes, and it contributes to diabetes dysregulations



Amylin exerts important physiological effects on metabolism and weight control

- 1. Activates amylin receptors in different brain areas Satiety, well-being, cognitive functions protection
- 2. Inhibits glucagon secretion
 Better glycemic control, lower PPG rise
- 3. Slows gastric emptying Synchronize insulin arrival with BG rise

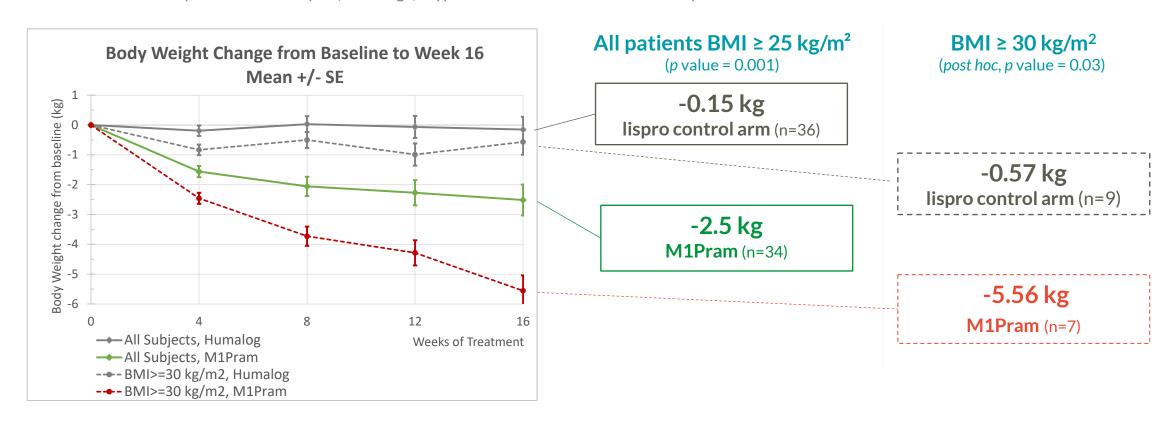


Reestablishing the physiologic equilibrium between insulin and amylin offers strong clinical benefits



M1Pram reduces body weight with high efficacy in T1D with obesity

CTO41: Phase 2 study – M1Pram vs. lispro (Humalog®) - type 1 diabetes - 16 weeks ambulatory



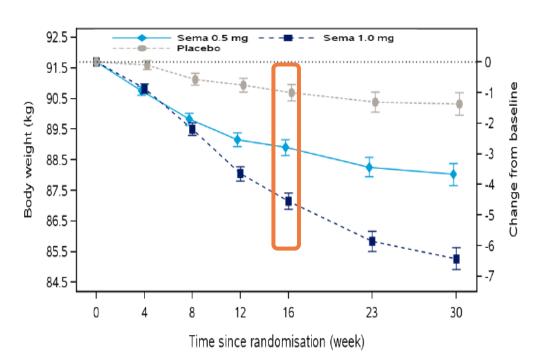
M1Pram could be the answer to the unmet medical need of obesity in T1D

ADOCIA 🛴

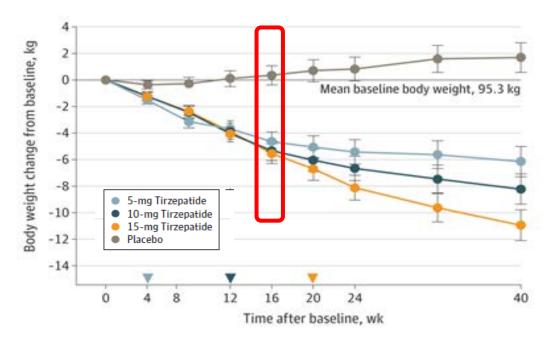
Body weight loss with M1Pram in T1D is comparable with insulin plus Semaglutide or Tirzepatide treatment in T2D

Change from baseline in body weight at W16

Semaglutide - 4.55 kg vs. - 1.08 kg



T2D taking basal insulin Mean BMI 32 kg/m² [19-51] N=396 (3 arms) Novo Nordisk Sustain 5 study Tirzepatide - 5.6 kg vs. + 0.4 Kg



T2D taking glargine Mean BMI 33.4 kg/m² N=475 (4 groups) E. Lilly Surpass 5 study

Maintained glycemic control, with 21% reduction in prandial insulin dose

| | Change from baseline* Mean (SD) | | Dyoluo |
|--------------------------------|---------------------------------|--------------|-----------|
| | M1Pram | Lispro | P-value |
| HbA _{1c} (%) | 0.14 (0.51) | 0.10 (0.51) | 0.81 (NS) |
| Time-In-Range 70-180 mg/dL (%) | -3.17 (8.76) | -1.54 (8.87) | 0.29 (NS) |
| Prandial insulin dose (U/d) | -5.97(6.18) | -0.61(7.08) | <0.001 |

While M1Pram reduces weight, it maintains same performance than Humalog (standard of care)



^{*} Baseline: 3 week assessment before treatment Results: 3 last week assessment of the 16 week treatment period

Medical value of M1Pram is confirmed

- ✓ Significant weight loss demonstrated & satiety effect
- ✓ Glycemic control maintained, without increase of hypoglycemia
- ✓ Sparing effect on prandial insulin dose
- ✓ Product well tolerated and safe
- ✓ Easy-to-use: one product injected at mealtime
- ✓ Excellent patient satisfaction reported

These attributes will allow M1Pram to serve a multi-billion-dollar market

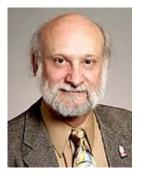
M1Pram generates high expectations from KOLs



"The phase 2 study of M1Pram shows that a single injection with each meal is as easy to use and as efficient as Humalog for glycemic control without increasing the rate of hypoglycemia. In addition, weight control is challenging for T1D patients, potentially limiting glycemic control and adding cardiovascular risk. While reducing insulin requirement, M1Pram improved appetite control and had a beneficial effect on weight, particularly in obese T1D patients. These features support a future role for this combination formulation for T1D."

"The glycemic results with M1Pram (P1b) are quite promising as is the observed weight loss, which is important given the characteristics of the population taking prandial insulin. I look forward to the next series of clinical trials."

Jay S. Skyler, Professor of Medicine, University of Miami Leonard M. Miller School of Medicine



Dr. Matt Riddle, Professor of Medicine, Oregon Health & Science University

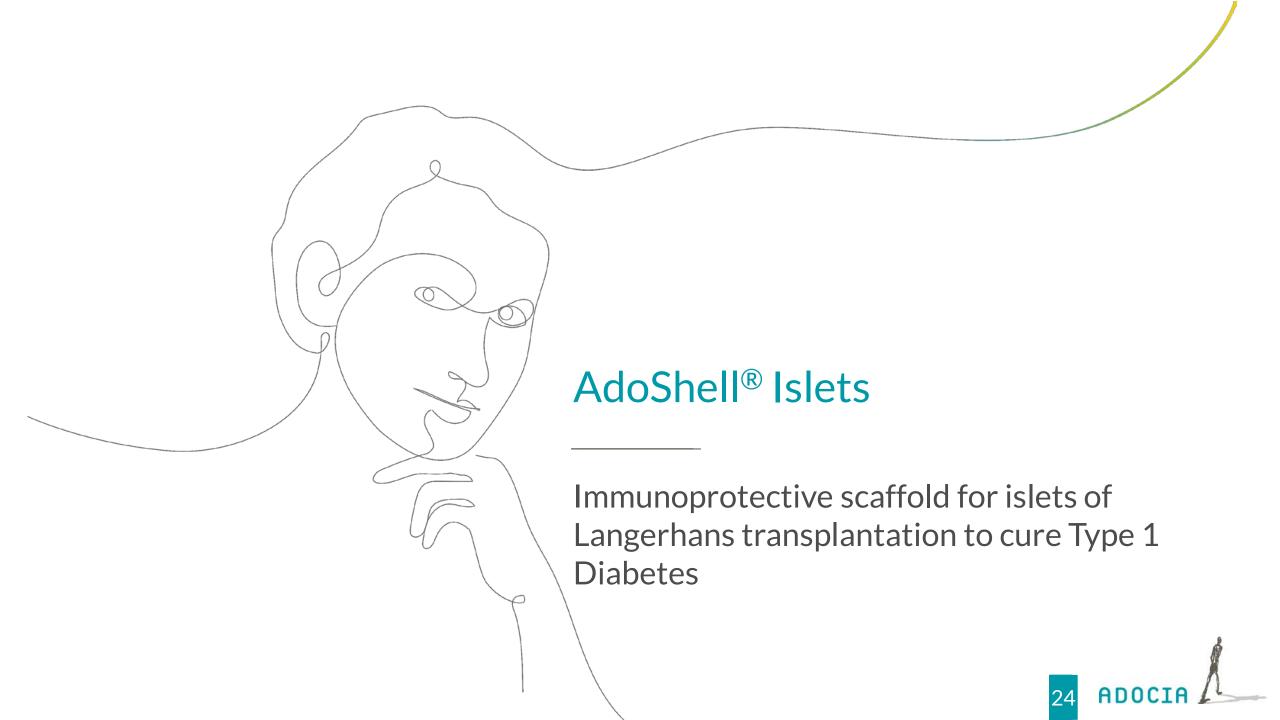
"This combination has the potential to finally deliver on the promise of pramlintide for a large number of patients."

Prof. Robert Ratner, Georgetown University Washington DC



"Remarkably, after only 3 weeks of treatment with M1Pram (P1b), all known pharmacological effects of pramlintide were observed."

Prof. Thomas Pieber, Medical University of Graz, Austria



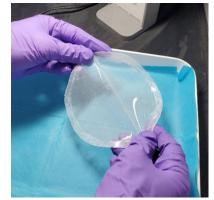
AdoShell® Islets: the promise of cell therapy without immunosuppression



Islet transplantation is already approved and practiced in clinic in several countries



Large clinical application is restricted due to AEs of immunosuppressants, used to avoid rejection



Human size AdoShell® prototype

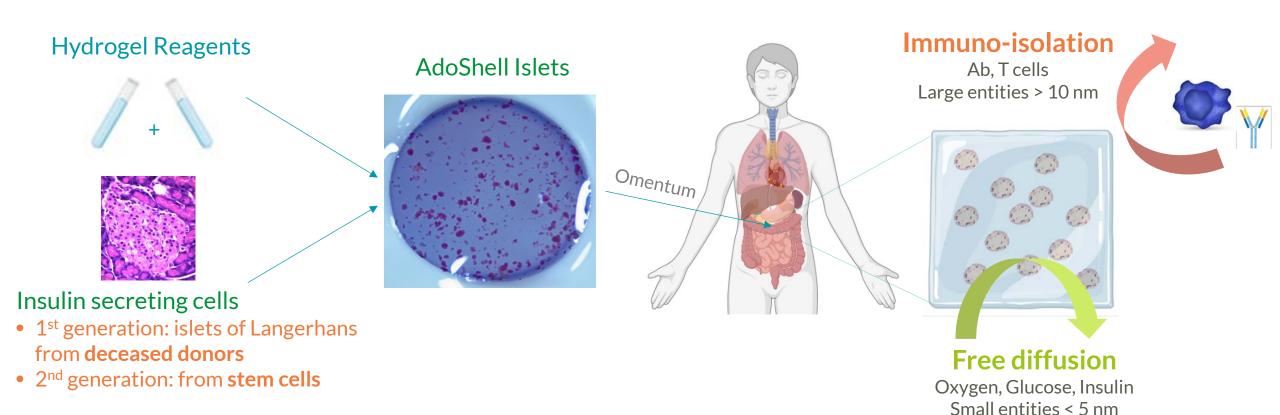
AdoShell Islets:

Designed to overcome the **important challenges in diabetes cell therapy**

Immunoprotective hydrogel encapsulating cells

- ✓ Allows insulin secretion in response of glycemic variation
- ✓ Protects cells from host's immune system
- ✓ Easily implantable & removable by minimally invasive surgery

AdoShell®, an immuno-protective scaffold for islets of Langerhans transplantation for diabetes



AdoShell® could ensure cell engraftment and long-term functionality in the absence of immunosuppression



AdoShell®: An ultra-thin, immuno-protective hydrogel film for cells encapsulation



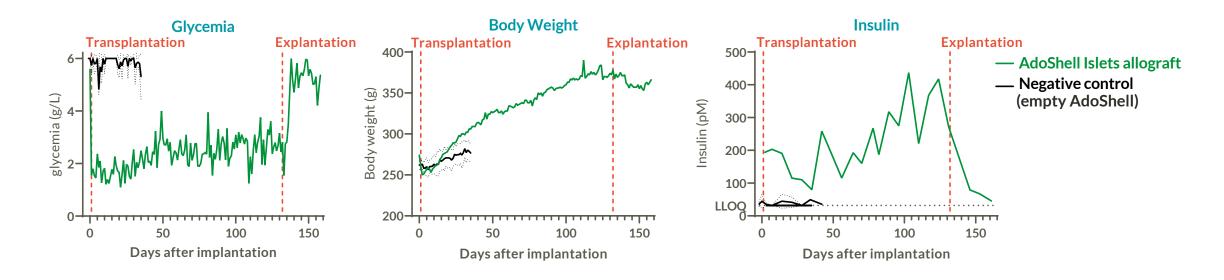
| High water content for cell compatibility | ✓ |
|---|--------------|
| Immuno-isolating gel | \checkmark |
| Ultra-thin aseptic film | ✓ |
| High cell density for miniaturization | \checkmark |
| Biocompatible | ✓ |
| Mechanically resistant | \checkmark |
| Implantable by mini-invasive surgery | ✓ |
| Retrievable by mini-invasive surgery | \checkmark |
| Long term stability | \checkmark |

Technology patented

First AdoShell® application for cell therapy is to provide a cure for diabetes

POC in diabetic rat: glycemia controlled during a 132-day study without immunosuppressor

- 1. Transplantation of AdoShell containing allogenic islets in diabetic rat peritoneum at day 0
- 2. Explantation of the implant at day 132. Sacrifice of control group at day 35



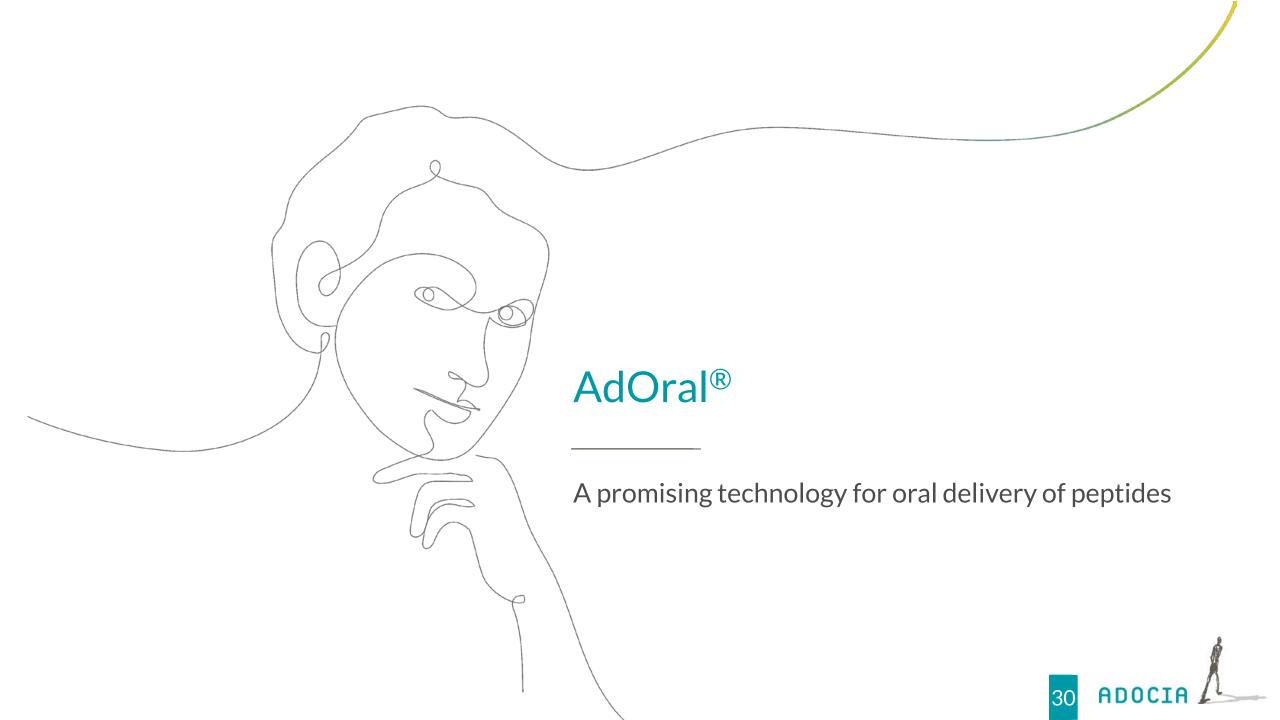
Next step: preparation of regulatory interactions in view of a first in human study in 2025

AdoShell provides a platform, opening to many cell therapy markets

- AdoShell scaffold displays key attributes enabling to progress towards new and more effective cell therapies:
 - ✓ No need of immunosuppressors
 - No need of gene-editing strategy for immune-evasive iPSCs
 - ✓ Safe and efficient cell engraftment
 - ✓ Safely removable and replaceable implant
 - ✓ Containment of proliferative cells
- Application of AdoShell to other pathologies is vast, considering different cell sources:
 - ✓ Stem cells
 - ✓ Engineered cells for therapeutic proteins delivery (cell factory)

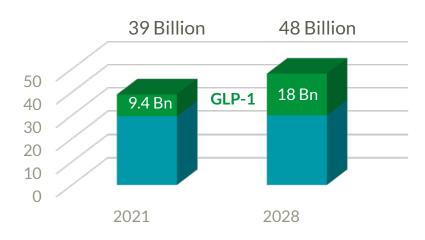
Adocia is looking to partner with companies working in cell therapy



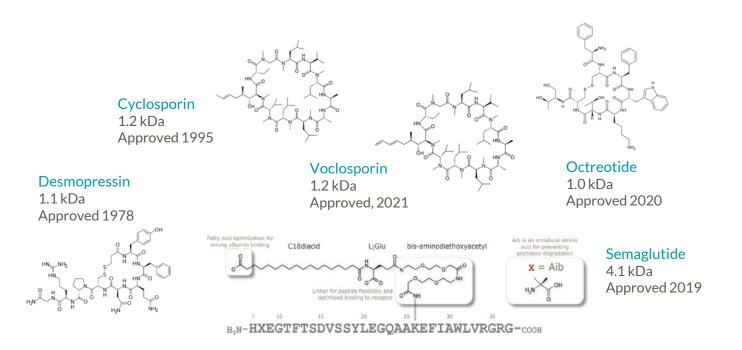


Only 5 peptide therapeutics have been approved for oral delivery in a growing market dominated by injectable forms

Global Peptide Therapeutics Market* (\$) *mAb excluded from the analysis



Data Source: Global Data



There is a huge market opportunity to develop AdOral, a new technology for oral delivery to replace injections of peptides

AdOral: a promising technology for oral delivery of peptides

AdOral is a unique formulation based on a new type of permeation enhancer (PE) combined with peptide protection against degradation

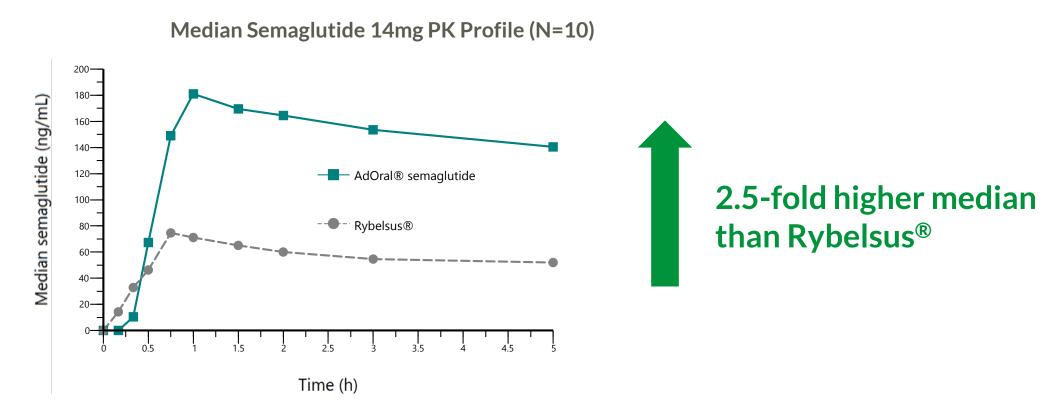
Technology patented until 2042 globally

The technology has been validated with Semaglutide compared to Rybelsus[®], an oral formulation for Type 2 diabetes treatment

Potential applications to other peptides or proteins

Two feasibility studies are ongoing on partners' proteins Other feasibility studies are under discussion

Semaglutide formulated with AdOral has shown significantly improved bioavailability in animal



Adocia seeks to establish proof of concept with peptides from future partners



AdoGel, a biomaterial for ultra long delivery of therapeutic agents

Adocia is developing a soft tissue-like biomaterial for sustained release of small molecules/biologics

- Pharmaceutic forms
 - Implantable & Removable device
 - Injectable (in-situ forming gel)
- Main properties
 - Release Duration: from 1 month and up to 36 months
 - Pharmacokinetics: pseudo zero-order, no burst
 - Local or systemic delivery
- Potential to release different types of APIs
 - Small molecules: contraceptives, anti-HIV, anti-cancer drugs, ...
 - Peptides & Proteins: monoclonal Ab, GLP-1, PTH, ...

First showcase application: AdoGel with Levonorgestrel (LNG)

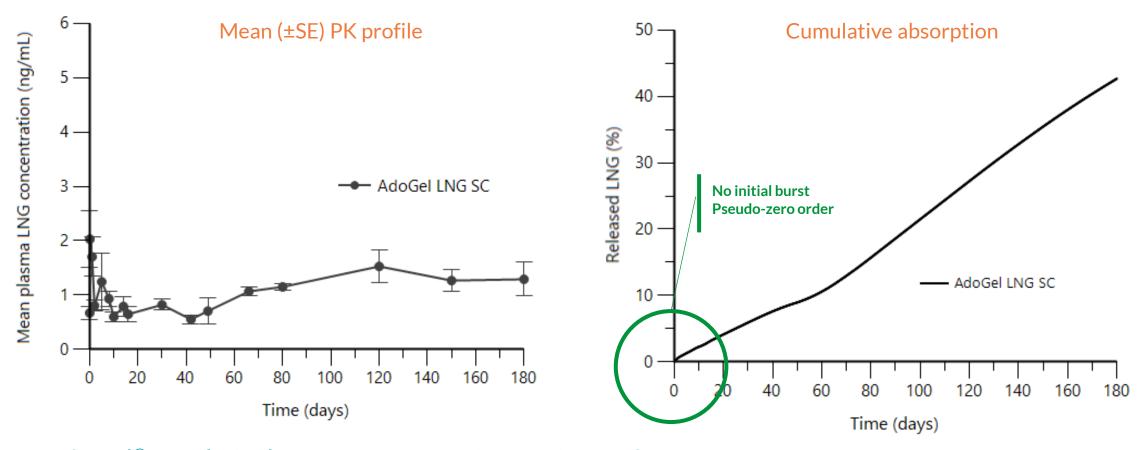


Implantable device



Injection and in-situ forming gel

First application: sustained release of Levonorgestrel (LNG) over 180 days after SC administration in animal model



AdoGel® Preclinical proof of concept is ongoing with LNG Additional showcase studies are also ongoing with antibodies Adocia is offering this technology to partners, for application on their proprietary molecules



Financial & news flow

News flow

BC Lispro

- In China (with partner THDB): Phase 3 ongoing
 - Phase 3 completion expected H2 2024
 - \$10m milestones payment
- In USA/EU: Phase 3 ready to be launched under partnering condition

M1Pram

- Exclusive negotiation rights granted to Sanofi in view of a global partnership
- Preparation of Phase 2b T1D with obesity (U.S.) ongoing

AdoShell® Islets

First-in-human study preparation in 2024

AdoGel® Sema

Preclinical Proof of Concept in 2024

AdOral®

 First-in-human study expected under partnering condition



THDB: Tonghua Dongbao Corporate Presentation

Key financials & shareholder information

As of June 30, 2024

Key financials features

Already received

- €97m raised in direct offerings since inception
- \$135m received from partnerships

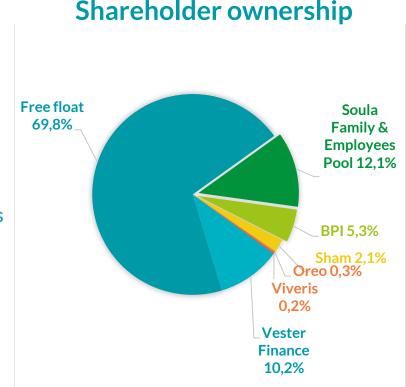
Expected payments

• \$30m as per contract with THDB + royalties

Cash position: €10.3m

Equity line: 845k shares issued, out of max.
 1.7m shares*

Debt (excl. derivatives and IFRS16): €5.6m



Market information

- Listed on Euronext Paris (ADOC)
- 15 million of outstanding shares
- ADR program in the US (ADOCY)





Analyst coverage

- Kepler Chevreux Justine Telliez
- Oddo Oussema Denguir



Summary

- Adocia's projects are based on 4 proprietary technologies: BioChaperone®, AdOral®, AdoShell®, AdoGel®
- Adocia is seeking to license-out the assets from its well-advanced and diversified pipeline:
 - Key products in ongoing clinical trials:
 - o BC Lispro: Phase 3 in China
 - o **BC Combo:** positive results on 3 clinical trials
 - o M1Pram: Exceptional weight loss in Phase 2 Exclusivity right granted to Sanofi in view of a global partnership
 - Proprietary innovative technology platforms, with applications tested at preclinical stage:
 - AdoShell® Islets, an immuno-protective scaffold for cell therapy for diabetes
 - AdOral[®], for oral delivery of peptides (semaglutide)
 - AdoGel®, for long-acting delivery of small molecules (LNG) and peptides (semaglutide)
- \$30m in development milestones to come from Tonghua Dongbao partnership (\$10m expected in 2024)
- Cash position: €10.3m¹





Thank you for your kind interest

115 avenue Lacassagne

69003 Lyon – FRANCE

Ph.:+33 4 72 610 610

contact@adocia.com