Developing RNA-based therapies for the treatment of childhood cancer

October 2024 FORO FIDBAN



Roadmap







Aptadel Lab set up (PCB)



2022

Patent 2 filing

2023



2024

Regulatory preclinic

Series A 5M€



2029

Dec 2020



2021



 Aptadel incorporation

- Patent 1 license
- Interim Management



Scientific team consolidation









Patent 1 granted in China



2025

Clinical Trials



Lead program:

candidate selection

Bridge round closing 1.5M€

















PROBLEM: Lack of therapies developed specifically for children

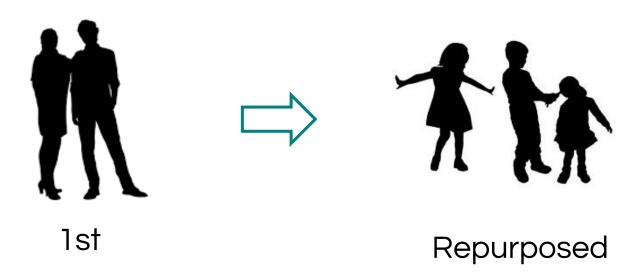


PROBLEM: Lack of therapies developed specifically for children



Childhood cancer is rare but remains the leading cause of death by disease for children under 15.

More than 70% of these children will have chronic health problems



Only 12 new treatments specifically developed for paediatric cancer have been approved by the FDA in the last 25 years (vs >200 for adults)

PROBLEM: Cancer therapeutics are unspecific and highly toxic



- The primary cause of cancer death is metastatic disease
- Metastasis management is performed by chemotherapeutics that kills both cancer cells and healthy cells

TRADITIONAL APPROACH

Chemotherapy

Radiation therapy

- Poor efficacy
- Non-specific
- Serious side-effects
- Low quality of life

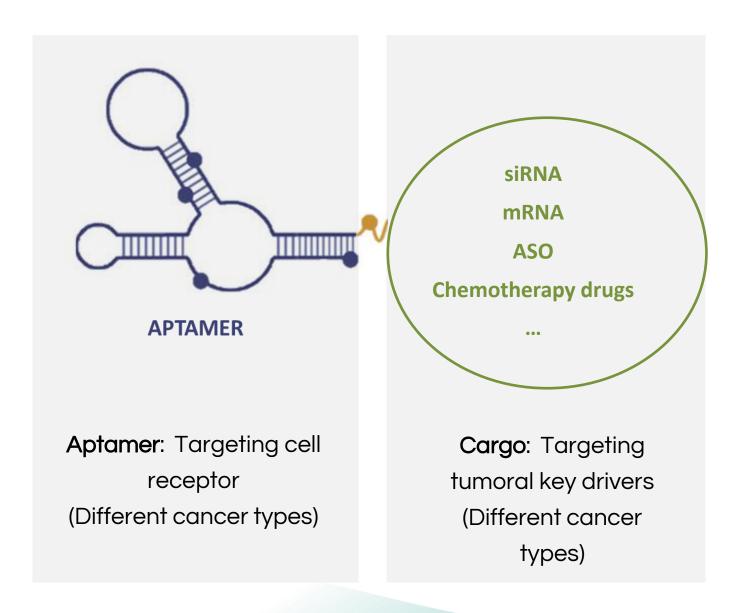
What if cancer cells can be effectively killed while leaving healthy cells unharmed?

FUTURE OF CANCER TREATMENT: TARGETED THERAPIES



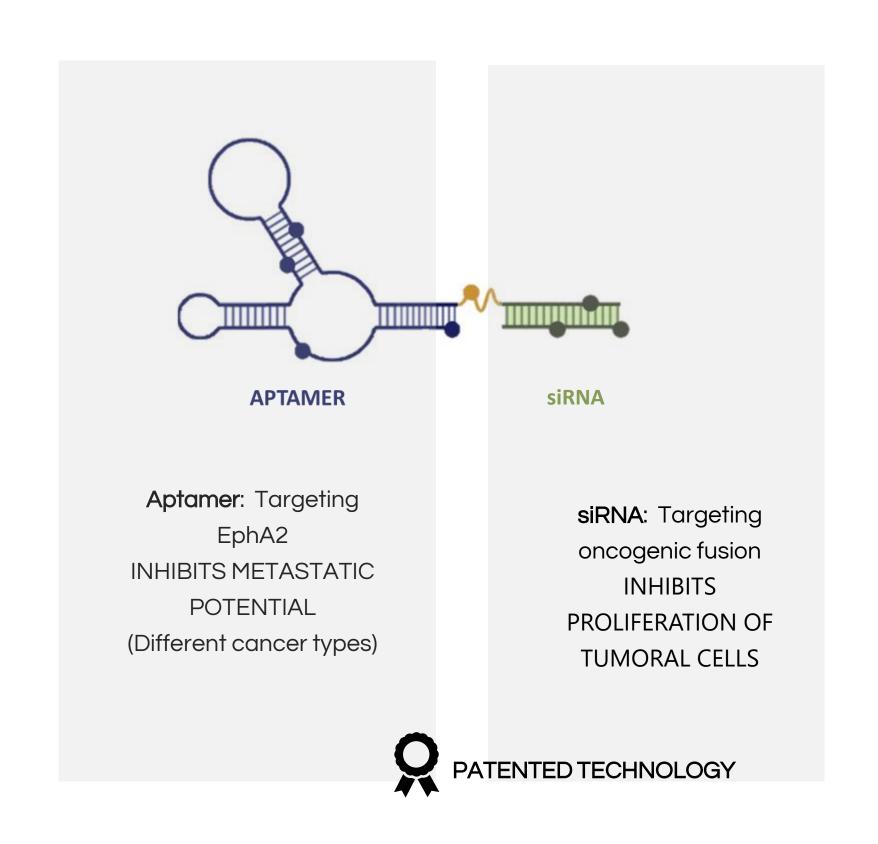
APTAMERS AS THERAPEUTIC AGENTS

- Aptamers are oligonucleotides capable of binding to specific targets with high affinity and specificity and can be conjugated to different drugs for therapeutic delivery.
- They possess intrinsic properties that bring advantages in clinical applicability and drug manufacturing.

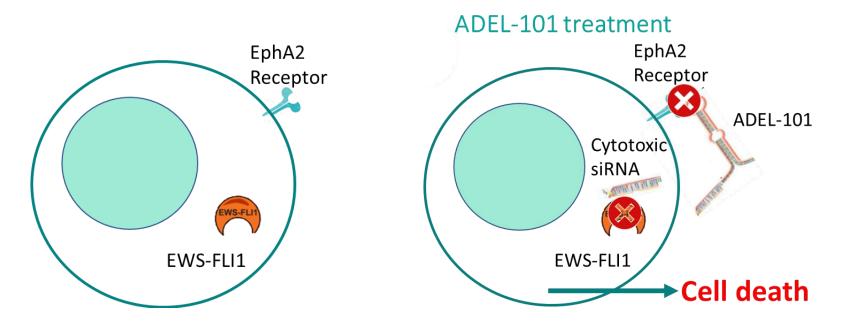


PLATFORM VALIDATION: Ewing Sarcoma and Rhabdomyosarcoma (childhood cancers)

APTADEL has developed a novel RNA aptamer targeting EphA2 cell receptor (overexpressed in cancer cells) as a selective delivery tool to treat cancer









Highly stable



Highly stable

Non toxic



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Non toxic

3x less metastasis



Highly stable

Non toxic

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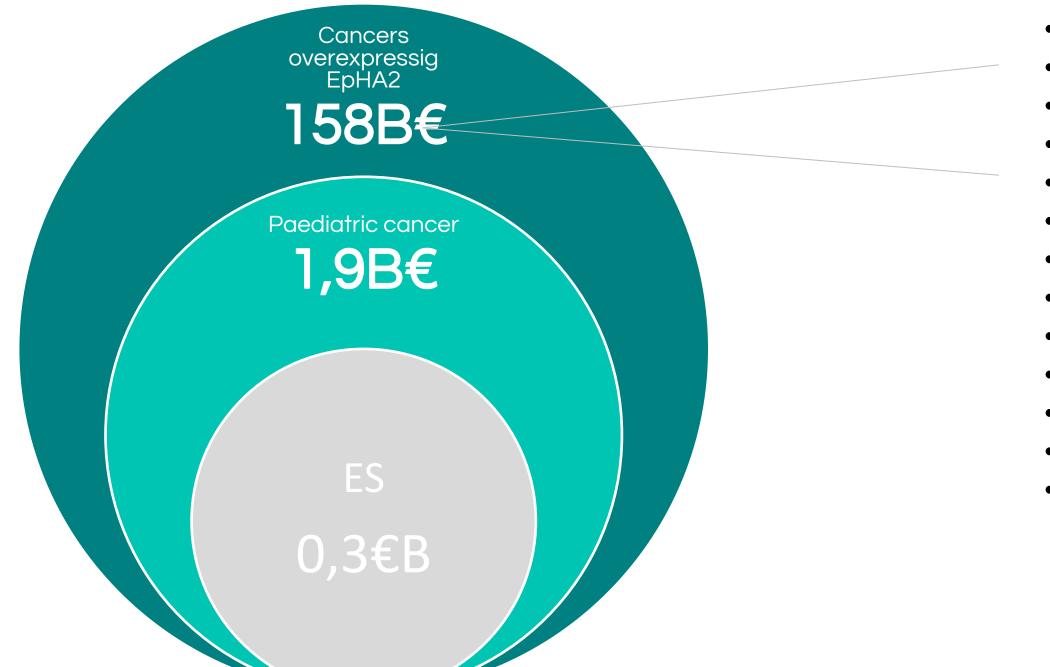
30% less tomour growth



Highly stable Non toxic 3x less metastasis 30% less tomour growth Highly specific

MARKET OPPORTUNITY

The potential of Aptadel's platform for generating new personalized candidates for different types of cancer is very high, with a significant impact on the metastatic processes of all solid tumors related to EphA2 and numerous \$B oportunities



- Prostate
- Breast
- Colorectal
- Cervical
- Melanoma
- Pancreatic
- Bladder
- Bone Sarcoma
- Glioblastoma
- Ovarian
- Esophageal
- Rhabdomyosarcoma
- Ewing Sarcoma

UNIQUE COMPETITIVE POSITION



BUSINESS MODEL

The business model pursued by Aptadel Therapeutics consists of the preclinical and clinical development of ADEL-101 up to clinical Phase IIa, followed by licensing the solution to a large pharmaceutical company that will make additional investments to bring the product to market.



FEATURED M&A IN THE RNA LANDSCAPE

Astellas acquired Iveric Bio (\$5.9B) FDA-approved aptamer 2023

AstraZeneca development and commercialization deal with Ionis (\$200M upfront payment)

Alnylam acquired Sirna Therapeutics (\$175M upfront payment)

COMPANIES EXPLOITING RNA THERAPEUTICS









APTADEL'S PIPELINE STATUS

NON-REGULATORY REGULATORY LEAD **DISCOVERY** CLINICAL **PROGRAM PRECLINICAL PRECLINICAL OPTIMIZATION** ADEL-001 RNA (Anti-metastatic drug) **Aptamer** ADEL-101 Aptamer-(Ewing sarcoma) conjugate ADEL-102 Aptamer-(Rhabdomyosarcoma) conjugate ADEL-103 Aptamer-(Adult indication) conjugate **ADEL-201** Aptamer-(Diagnostic kit) conjugate 20M€ 1.5 M€ 5 M€ 1.75M€ private investment + ~2M€ national and international competitive calls

















JOIN OUR JOURNEY

We are raising 1.5M€ to initiate the preclinical regulatory stage of Aptadel's childhood program and accelerate our expansion plans to the adult market, validating our RNA platform.

COMMITTED

0.3 M€ current investors

0.5 M€ NEW INVESTORS

COMPETITIVE FUNDING

> 2 M€ submitted (2024)

MILESTONES USE OF FUNDS Business Incorporate specialized investors to further close a 5M€ investment in 2025 to complete preclinical phase. **70%** R&D R+D Finish in vivo PoC and initiate the regulatory preclinical phase. Advance on the adult indication. **Talent 20% PERSONNEL** Incorporate key talent for the next development phase. **10%** G&A

APTADEL'S TEAM IS COMMITED TO FIGHT CHILDHOOD CANCER



Gisela Lorente

CEO

Extensive international experience in tech transfer and innovation with proven track record of portfolio management, IPR negotiation, technology licensing agreements with the pharmaceutical industry, creation of spin-offs, and negotiations with investors.



Nadia Pons

COO

Innovation expert with >10 years of experience managing cutting-edge innovation projects. Specialized in new business development in cross-cultural environments. Solid competence in generating strategic alliances and project management. Previous experience leading operations in oncology start-ups.



Oscar Martinez

Academic Founder

Principal investigator at IDIBELL with >25 years experience in biomedical research and childhood sarcomas. KOL in Ewing sarcoma. Experience in clinical trial design, having participated in two Phase I and one Phase II clinical trials. PhD Biochemistry, Autonomous University of Barcelona.



Adrian Torres
Head of R+D



Jazmine Arevalo
Senior Scientist & PM



Laura Barbera
Lab Technician



Paloma Giangrande PhD

KOL RNA aptamers

drug development



Jaume Reventós PhD, MD Translational medicine - BOD



Juan Martin Liberal MD Medical oncology (sarcoma)



Lluís Valenciano CFO



Silvia Bertran
IPR Management



Judy Carmody PhD CMC Management

Thank you

GISELA LORENTE

Chief Executive Officer

www.aptadeltx.com





Parc Científic de Barcelona (Spain)

