

Antibacterial cyclic peptide (PLP-3)

ISGlobal Barcelona Institute for Global Health

IRB BARCELONA INSTITUTE FOR RESEARCH IN BIOMEDICINE



Partners

Clínica Barcelona

UNIVERSITAT DE BARCELONA

NIH NIAID



Market Opportunity

• Antibiotic resistance was responsible for **1.27 M deaths** worldwide in 2019. By **2050, 10 M people may die every year** from infections caused by multidrug resistant bacteria, becoming the number 1 cause of global deaths.

• It causes **2.5 M days of extra hospitalization** per year throughout the EU resulting in huge **costs of 900 M€**.

• The global Antibiotics Market Size is estimated at \$63,2bn by 2027, with a **CAGR of 5.1%** from 2020 to 2027.



Technology

• **Potent Antibacterial.** The cyclic peptide possesses low Minimal Inhibitory Concentration (MIC₅₀) values:

Microorganism (resistant strains)	MIC ₅₀ *	MIC ₉₀ *
<i>Acinetobacter baumannii</i>	1	2
<i>Klebsiella pneumoniae</i>	4	16
<i>Pseudomonas aeruginosa</i>	4	8
<i>Staphylococcus aureus</i>	4	8
<i>Enterococcus faecium</i>	2	2

*in µg/mL

• **Stable and active under physiological conditions.** No significant changes in antimicrobial activity of PLP-3 in the presence of human albumin were observed.

• **Toxicity in human cells.** IC₅₀ values of ca. **227 µg/ml** for XTT assays indicating up to >100 folds over MIC values for strains of the tested bacterial species over cells. Maximum Tolerated Dose (MTD) via IV of 2,5 mg/kg.

• IC₅₀ of **48 µg/ml** for haemolysis assays.



Intellectual Property

• **ISGlobal (47 %), Institute for Research in Biomedicine Barcelona (40 %), Barcelona Clinic Hospital (7 %) and University of Barcelona (6 %)** share joint ownership.

• **European patent application** submitted 27th of May 2022. **PCT application** filled in **May 2023**. No objections were raised in the EESR and ISR for all the claims.

• **National phase entry** in the USA and Europe.



Competitive Advantage

• **Novel cyclic peptide** designed and synthesized.

• **Antibacterial activity** against the most problematic **Gram-negative and Gram-positive** resistant strains, including *A. baumannii*, *P. aeruginosa*, *K. pneumoniae*, *S. Aureus* and *E. faecium*

• **Low in vitro toxicity** in human cells.

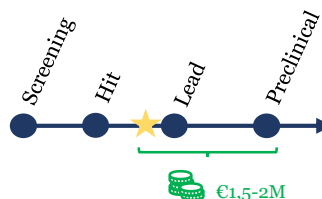
• Presence of the **disulfide bond** within the peptide backbone would facilitate metabolism thus **reducing nephrotoxicity** issues.

• Bactericidal and very **rapid killer**.



Development

• We are currently completing the Hit-to-Lead stage:



• Currently studying the **mechanism of action** through genomic mutant analysis, **biofilm activity**, and analyzing the **activity of three more analogs**.

• Assessing inhaled route. **In vivo cystic fibrosis model** planned for early 2025.

• **OUR ASK: CO-DEVELOPMENT, LICENSE AGREEMENT and/or INVESTMENT.**

Our Team



Jordi Vila
Research Professor
ISGlobal-Clínica Hospital



Clara Ballesté
Associate Research
Professor
ISGlobal-Clínica
Hospital



Elisabet Guiral
Project Manager
ISGlobal



Núria Martín
Predoctoral fellow
ISGlobal



Oscar Casado
Innovation Manager
ISGlobal

Get in touch: innovation@isglobal.org