



Elvitegravir repurposing for multidrug-resistant Gram-positive bacteria

ISGlobal Barcelona Institute for Global Health

Clínic Barcelona

UNIVERSITAT DE BARCELONA

IrsiCaixa
Institut de Recerca de la Sida

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CENTRO DE INVESTIGACIÓN BIOMÉDICA EN RED



Unmet Medical Need

- Antimicrobial resistance (AMR) is responsible for significant mortality, with drug-resistant bacterial infections leading to an estimated **1.27 million deaths globally in 2019 alone**.
- Methicillin-resistant *Staphylococcus aureus* (MRSA) and vancomycin-resistant *Enterococcus faecium* (VRE_{fm}) are considered **High Priority pathogens by the WHO**.
- There are very **few therapeutic options for MRSA and VRE_{fm}**. In 2023 there were **791,191 MRSA cases** across the 7 Major Markets (7MM).
- Bacteremia due to ***S. aureus*** has been reported to be associated with mortality rates of **15 % to 60 %**, and of **60 % to 70 %** for **VRE infections**.



Technology

- Potent Antibacterial.** Elvitegravir possesses low Minimal Inhibitory Concentration (MIC₅₀ and MIC₉₀) values *in vitro*:

Microorganism (resistant strains)	MIC ₅₀ *	MIC ₉₀ *
<i>Staphylococcus aureus</i> (19 resistant strains, 10 MRSA, 5 resistant to quinolones)	8	8
<i>Enterococcus faecium</i> (22 resistant strains, 11 VRE)	4	16
<i>Enterococcus faecalis</i> (16 resistant strains)	4	16

*in µg/mL

- In vitro Proof-of-Concept.** ELV has been tested against **clinically relevant** multidrug-resistant Gram-positive bacteria.

- Selection of resistant mutants has been investigated in both *S. aureus* and *Enterococcus* spp.

- Killing curves showed a bacteriostatic effect.



Intellectual Property

- ISGlobal** (49 %), **Barcelona Clinic Hospital** (29 %), **University of Barcelona** (12 %), **IRSiCaixa** (5%), and **CIBER** (5 %) share joint ownership.
- European patent application** submitted 17th of October 2023. **The PCT application was filed in October 2024**. No objections were raised in the EESR for the new use of elvitegravir as an antibacterial agent.



Competitive Advantage

- Elvitegravir (EVG) shows activity against multidrug-resistant Gram-positive bacteria, even those highly resistant to quinolone.
- EVG can be administered **orally**, facilitating patient follow-up and **reducing unnecessary hospital admissions and healthcare costs**.
- The repositioning strategy offers **less time to market**, less risk of failure, and **less investment** than traditional drug discovery.
- Very low selection of resistant mutants.



Development

- We are currently completing the Preclinical development stage:



- Currently performing **in vivo efficacy of ELV** combined with other known antibiotics in experimental endocarditis model against VR *E. faecium* and MRSA.

- Studying the MoA** through molecular docking.

- Investigating the activity against biofilm-producing bacteria.**

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

Our Team



Jordi Vila
Research Professor
ISGlobal-Barcelona Clínic Hospital



Josep Maria Miró
Head of Infectious Diseases and AIDS
Barcelona Clínic Hospital



Elisabet Guiral
Project Manager
ISGlobal



Núria Martín
Predoctoral fellow
ISGlobal



Oscar Casado
Innovation Manager
ISGlobal

Get in touch: innovation@isglobal.org