

# Research focus and expertise

- Optimising treatment regimens (monotherapy, combinations) to combat antimicrobial resistance
- **Experimental microbiological** research (including dynamic infection models, multi-OMICS)
  - **Mathematical modelling** (mechanism-based, population pharmacokinetic/ pharmacodynamic, quantitative systems pharmacology modelling, immune response)
- Integrating both to develop better treatments against serious infections (*next slide*)
- Support other projects through mathematical modelling of *in vitro*, *in vivo* and/or clinical data

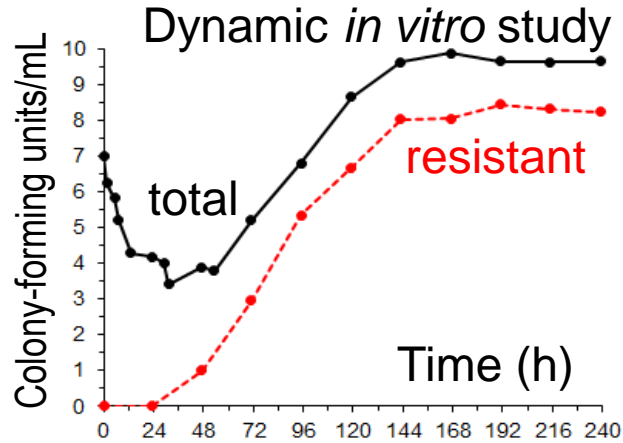
Cornelia Landersdorfer, PhD, Associate Professor



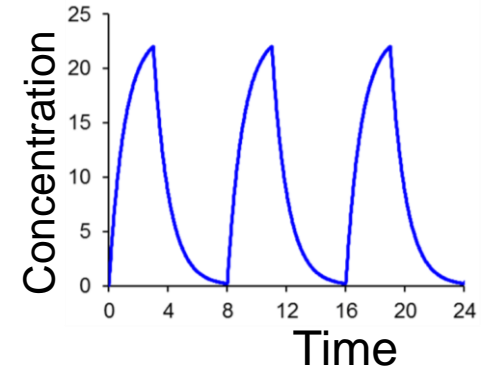
Monash Institute of Pharmaceutical Sciences  
World's Top 2 in Pharmacy and Pharmacology  
2023 (QS World University Rankings)

MONASH  
PHARMACY &  
PHARMACEUTICAL  
SCIENCES

# Combating antimicrobial resistance by integrating dynamic infection models, multi-OMICS and mechanism-based modelling towards personalised therapies



Drug concentration-time profiles in patients representative of the region



Bacterial isolates with resistance mechanisms common in East Asia

Multi-OMICS of resistance

**Pathogen**

**Drug**

**Host**

Bacterial killing  
Resistance emergence

Pharmacokinetics

Mathematical modelling, AI

Infection

Host response

Impact of renal function, body weight, disease state, ethnicity, others

Multi-OMICS on virulence factors, capsule, etc.

*In vivo* infection model, immune response

- Model-optimised monotherapies or synergistic combination regimens, prospectively evaluated *in vitro* and *in vivo*
- Tailored to characteristics of bacteria and patient, to maximise effectiveness and minimise resistance emergence