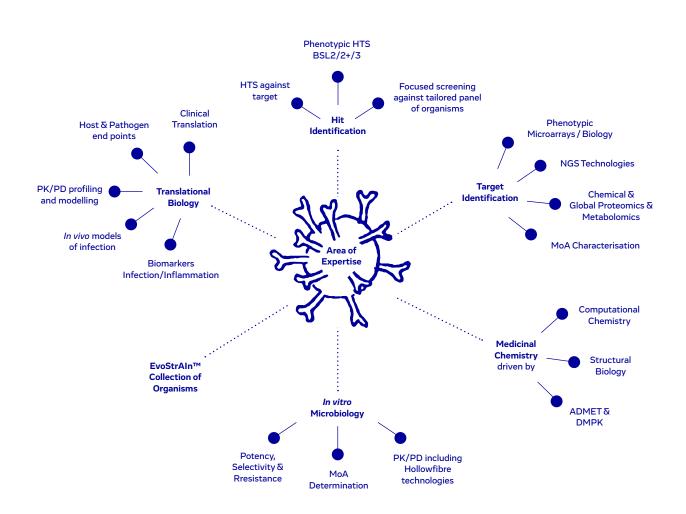


Anti-Infectives: Virology Platform

- Fully integrated platform for fast and efficient identification and progression of novel antiviral therapeutics
- Cutting edge technologies and discovery platforms flexibly tailored around projects' scientific requirements
- Focus on respiratory viruses including SARS-CoV2 as well as hepatitis B virus, capabilities ranging from screening to animal models
- Development of novel tools, assays and models based on project requirements





Viral in vitro assays

- ▶ Viral ToxGlo™ screening assay
 - Single step assay measuring metabolic activity
 - Increase in luminescence signal by inhibition of virus
 - Also suitable for cytotoxicity counter screens
 - Available for a range of viruses including SARS-CoV2
- ▶ Plaque assay
 - Quantification and validation of viral stocks
 - Quantification of viral burden in tissue
 - Generation of resistant virus
 - Mechanistic studies
- ▶ ELISA
 - Quantification of virus in infected cell culture
 - Quantification of virus specific antibodies from infected animals
- Microneutralisation assay
 - Quantification of virus specific neutralising antibodies from infected animals
- ► SARS-CoV2-spike pseudotype infection assay for all variants of concern

Viral animal models – example SARS-CoV2 in hamster model

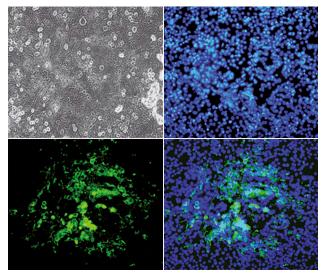
- Gold standard model for development of SARS-CoV2 treatment
- ► Hamster standard laboratory animals for SARS-CoV2 infection
- ▶ Model validated endpoints
 - Viral load in lung, nasal or other tissues homogenate by plaque assay and RT-qPCR
 - -Viral titre in oral swabs by RT-qPCR
 - Antibody titre via ELISA
 - Immunohistochemistry
 - qPCR for viral load under development
- ➤ Transmission model and ACE2 transgenic mouse model in development
- Neutralising antibodies in neutralisation assay
- Blood and tissue cytokine/chemokine quantification

Virology platform: From screening to pharmacodynamic assessment

- Antiviral HTS experience from reporter based replicon read-outs to infected cell assays handled in BSL2+/BSL3
- Medium throughput screening in 96- and 384- well format, in infected cell assays with metabolic or enzyme read-out
- ► SAR screening for integrated programmes antiviral potency vs. cytotoxicity
- MoA work e.g. resistant virus generation, order of addition effects, cell and virus strain specificity
- Access to Evotec platforms, e.g. Target identification by PhotoAffinity Labelling Mass Spectrometry (PALMS) studies, which can be performed in infected and uninfected cells
- ► Routine PK in mouse, rat and hamster, other rodent hosts are possible
- ► Development and performance of relevant rodent models

HEp2 cell infected with RSV-A2 Stained

Stained with DAPI (nucleus)



Stained with anti-RSV-F

Stained with anti-RSV-F (green) and DAPI (blue)