

# Quantitative Systems Pharmacology (QSP) Psoriatic Arthritis Model

Over 30 Phase 2 and Phase 3 clinical trials were used in training this model, spanning 14 distinct therapeutic agents, including TNF inhibitors, IL-17 inhibitors, IL-23 inhibitors, as well as JAK and TYK2 inhibitors in multiple patient populations (ex: naïve to biologic therapies).



## Key Applications

- Support customized pre-trial disease initiation and therapy pipeline failures
- Predict clinical efficacy of novel compounds
- Export data to other programs for ad hoc analysis

## Key Features

- Convenient generation and calibration of virtual populations that include inter-patient variability in pathophysiology as well as clinical endpoints
- Includes qualitative and quantitative data during model training
- Flexibility in trial pipeline specification allows for customized predictions of novel compounds in a virtual population calibrated to hundreds to thousands clinical trial datapoints
- A single, unified platform to generate virtual populations, simulate trials, analyze simulations, and generate informative plots allowing easy comparison to published trials to support pre-clinical decision making

**Validated virtual population** with new biological and pharmacological components can include novel compound predictions while recapitulating and validating against existing clinical trial data.

## Sound Science



### Leverages

domain knowledge in both rheumatoid arthritis and psoriasis to simulate immunological species, their interactions, and therapeutic interventions in both the joint and skin environments specific to psoriatic arthritis patients



### Integration

of biological and clinical trial data to provide predictions of common clinically-relevant biomarkers and trial endpoints for novel compounds, including ACR and PASI scores



### Capability

of physiological modelling and efficacy predictions for both topical and systemic administration routes



### Customization

of population characteristics to meet client-specific criteria or produce common trial populations such as biologic naïve or anti-TNF therapy non-responders

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