



SGS PACE

Health Inspired, Quality Driven.

SGS

Bring your drug to life

A unique, streamlined pathway from preclinical to FIH/POC.

What is SGS PACE?

Drug developers are facing several challenges as they continue to outsource the different steps of their drug development process to different external CDMO's and CRO's to get from preclinical to first in human/proof of concept.

SGS PACE brings together all the phases of drug development that would traditionally be handled by separate organizations and vendors – greatly improving efficiency.

The service integrates bioanalysis, drug development and clinical access, through one point of contact with enabled formulations for toxicology and FIH.

The industry will benefit from a comprehensive scientific, medical, and quality review throughout your project to support decision making and your objectives.

Enhance your drugs potential and show true value, early on in your drug development program.

A tailor-made program

Your bespoke SGS PACE solution provides an exclusive tailor-made program, designed with your objectives and goals in mind for your pace and budget. The customized program will deliver your clinical results effectively and efficiently.



“ SGS PACE offers an improved efficiency, helping to speed up the clinical development process through a single point of contact for all your formulation and early development needs. ”

Doctor Jelle Klein
Medical Director at our Clinical
Pharmacology Unit in Belgium

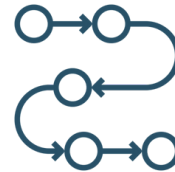
Benefits of SGS PACE



SGS PACE offers a more efficient and effective route to market, starting with preclinical testing right through formulation and manufacturing, to FIH/POC, all provided by one organization. Our tailor-made service provides an expert-directed roadmap to clinic through one Program Manager.



Our solution marries bioanalysis, drug development and clinical access, under one program with enabled formulations for toxicology and FIH, allowing you to maximize your drugs potential and show true value, from early on in your drug development program.



SGS PACE provides a streamlined pathway from preclinical testing right through to Phase II trials and can improve flexibility, efficiency, and the speed of clinical development.



Program Management

SGS PACE provides you with a dedicated program manager and strategic program plan

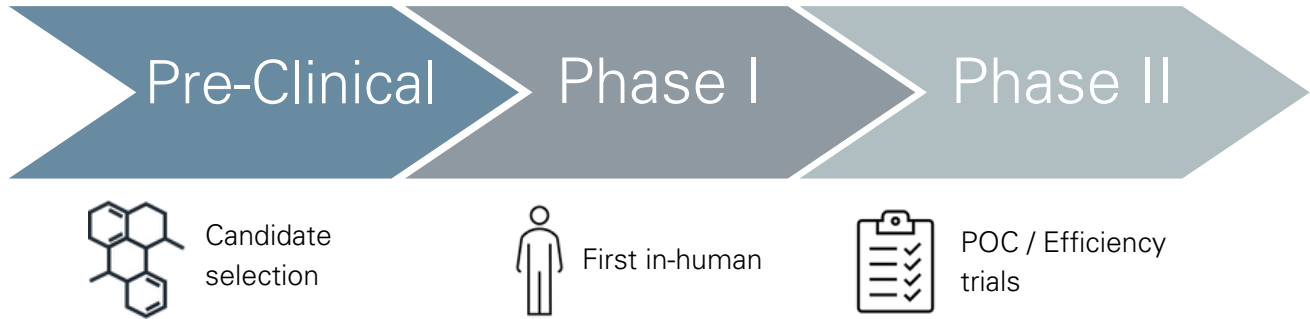
An integrated team of experts to streamline molecules' pathway

SGS PACE offers strategic consultancy and ongoing support from our global network of experts. Our consultants will work with you to build a custom program plan and advise you from formulation stages all the way through to FIH / POC.

Your SGS PACE Manager will provide:

- Coordination and milestone review points
- Clinical development consultancy
- Preclinical formulation and bioanalysis
- Clinical formulation/analysis/manufacturing
- Evaluation and optimization of the formulation in clinic, based on human data
- First in Human (FIH) dosing
- Bioanalysis and data analysis and reporting
- Hybrid FIH studies including POC in small patient population
- Phase II clinical development approach and management

Agile & Flexible Program



OUR PROGRAM

Formulation for Tox

- Bioanalysis
- (FIH formulation)

Dose Development for FIH

- Manufacture and release

Human Dosing

- Clinical trial conduct at our CPU
- Bioanalysis
- Data analysis and reporting

Dose Optimisation

- Manufacture and release
- Clinical trial supply

Patient Dosing – Multi Site

- Clinical trial management
- Bioanalysis
- Data analysis and reporting

Faster project delivery with just in time manufacture and release for tox and clinical studies



“ PACE is a unique, streamlined pathway from preclinical to FIH/POC bringing together world-leading clinical drug development services through one point of contact ”

Mike Frodsham
Chief Technical Officer,
SGS Quay Pharma

SGS PACE

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Contact us

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WHEN YOU NEED TO BE SURE

SGS