SAHMRI
South Australian Health & Medical Research Institute

Clinical Trials Platform

Exceptional Participant-Centred Clinical Trial Support and Services

SAHMRI's Clinical Trials Platform offers comprehensive support and services for industry funded and investigator led clinical research. We bring a strong history of successful conduct and independent oversight to **Phase 1b to Phase IV** trials (local, national, international) spanning diverse therapeutic areas. We are also a preferred outpatient clinical trial site renowned for its consistently high recruitment rates.

Situated in the heart of Adelaide BioMed City, the southern hemisphere's premier health and medical precinct, SAHMRI is a leading institute cultivating a collaborative ecosystem. Our institute unites over 700 world-class researchers and clinicians dedicated to translating groundbreaking research into tangible community health benefits.

Your Trusted Research Partner

Academic Research Organisation (ARO)

We're an ARO providing a comprehensive suite of services comparable to a Contract Research Organisation (CRO). Our academic approach follows Good Clinical Practice guidelines closely, with meticulous detail.

Experienced Outpatient Clinical Trial Site

We can also operate as a site to expertly assist you to meet your recruitment targets, provide quality data and sample collection, and adhere to regulations and timelines for your trial's success.

This **dual functionality** allows us to offer seamless integration of services, whether you require comprehensive ARO support or a high-performing site for your outpatient study.

We proudly support a diverse range of stakeholders, including Academics/Researchers (both internal and external), Clinicians, Industry Sponsors, Biotech and Pharma.

For more information on the SAHMRI Clinical Trials Platform scan the QR code

Contact SAHMRI Clinical Trials Platform

Our dedicated team is ready to assist with a comprehensive range of clinical research support

clinicaltrials@sahmri.com
(08) 8128 4570

Capabilities

SAHMRI's Clinical Trials Platform provides dedicated, professional support for clinical trials of all sizes and complexities, underpinned by a strong commitment to quality and strict regulatory compliance.

Expert Project Management & Participant-Centred Trial Coordination

Our experienced team delivers end-to-end project management for trials ranging from single site to complex multi-site trials and international collaborations. We seamlessly integrate at any stage – from initial project conceptualisation and design to navigating ethics, governance approvals and regulatory pathways, effective recruitment and high-quality coordination right through to close out. We are also adept at managing the intricacies of decentralised and hybrid trial designs, ensuring efficient and compliant execution.

Our in-house Study Coordinators and Research Nurses expertly provide participant-centred support, strictly adhering to ICH-GCP guidelines and can coordinate trials of all phases. Their compassionate approach also enhances participant engagement and experience.

Site Capabilities

Inside SAHMRI's iconic building, we have eight multi-purpose well-equipped and easily accessible outpatient clinic rooms, ideally suited for Phase II-IV studies. We collaborate with a strong network of experienced General Practitioner investigators, in-house Principal Investigators, and co-located Specialists across a broad spectrum of therapeutic areas. For participant convenience, onsite parking can be arranged, and our site is readily accessible via public transport options, including tram, train, and bus.

Targeted Digital Marketing Campaigns

Dedicated personnel are able to develop and deliver highly effective, project specific recruitment campaigns across Facebook, Instagram, and Google Ads, precisely targeting relevant participant populations with proven success in various trial designs and populations. In addition, our team manages an in-house Clinical Research Participant Register, INVOLVED, which we leverage to enhance the recruitment of high-quality trial participants.

Efficient Onsite Sample Processing

SAHMRI's onsite laboratory provides streamlined biological sample processing, including protocol-specific collection, processing, and short-term storage for central laboratory shipment. Our experienced team excels in pharmacokinetic (PK) sampling, capable of performing serial blood draws and processing samples for analysis.

Robust Data Management & Biostatistics

Supported by SAHMRI's Biostatistics Unit, our team provides comprehensive data management and biostatistics to develop Statistical Analysis Plans, generate randomisation schedules, assist with database design and construction, develop data quality checks and support report preparation for Data Monitoring Committees.

Comprehensive Monitoring and Auditing Services

Our skilled team of Clinical Research Associates deliver expert monitoring services for both investigator-led and commercially sponsored trials, prioritising participant safety and data integrity in accordance with respective guidelines. Monitoring is tailored based on individual study needs and risk assessments, offering flexible approaches with enhanced focus on quality aspects like protocol endpoints, ICH-GCP compliance, and audit readiness.

Timelines: Efficiency in Trial Initiation

We prioritise efficiency and proactively manage the necessary paperwork to expedite clinical trial initiation.

Our typical timelines are:

- Local Research Governance Office Approval: Expect approval within 2 weeks.
- Contract/Agreement Finalisation: We aim to finalise Agreements within 4 weeks from the conclusion of budget negotiations.