

A Study to Assess the Safety, Reactogenicity, and Immunogenicity of SK Japanese Encephalitis Messenger Ribonucleic Acid (mRNA) Vaccines (GBP560) in Healthy Adults

NCT06680128

Status	RECRUITING
Phase	Phase 1, Phase 2
Sponsor	SK Bioscience Co., Ltd.
Enrollment	402 participants

Key Eligibility Criteria

Inclusion (13)

- \[Age\]
 - For Stage 1, participant must be 18 to 55 years of age inclusive, at the time of signing the informed consent.
 - For Stage 2, participant must be 18 years of age and older, at the time of signing the informed consent.
 - \[Type of Participant and Disease Characteristics\]
 - Participants who are healthy as determined by medical evaluation including medical history, vital signs, physical examination, clinical laboratory tests and medical judgement of the investigator.
- ... and 8 more (see full listing online)

Exclusion (24)

- \[Medical Conditions\]
 - Any clinically significant respiratory symptoms (e.g., cough, sore throat), febrile illness (tympanic temperature $>38^{\circ}\text{C}$), or acute illness within 24 hours prior to the 1st study vaccination. Prospective participants with these conditions cannot be included until 24 hours after resolution.
 - History of congenital, hereditary, acquired immunodeficiency, or autoimmune disease.
 - Any positive test results for hepatitis B surface antigen, hepatitis C antibody, or human immunodeficiency virus (HIV) at screening.
 - History of bleeding disorder or thrombocytopenia which is contraindicating intramuscular vaccination in the investigator's opinion.
- ... and 19 more (see full listing online)

Locations (3 total)

Nucleus Network - Brisbane (Q Pharm), Brisbane, Australia
Nucleus Network - Melbourne, Melbourne, Australia
New Zealand Clinical Research, Christchurch, Central City, New Zealand

<https://clinicaltrials.gov/study/NCT06680128>

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