A Phase Ib/IIa Study to Evaluate the Safety, Tolerability, Pharmacokinetics and Pharmacodynamics of Oral Ecnoglutide Tablets in Chinese Participants With Overweight or Obesity

NCT07243171

Status	RECRUITING
Phase	Phase 1, Phase 2
Sponsor	Hangzhou Sciwind Biosciences Co., Ltd.
Enrollment	84 participants

Key Eligibility Criteria

Inclusion (2)

- At the time of informed consent, with BMI in the range of 24.0 to 35.0 kg/m2 (including the threshold), body weight \>60.0 kg for male, and weight \>50.0 kg for female;
- Self-declaration of body weight change \<5% within 3 months prior to informed consent ;

Exclusion (7)

- Diagnosis of overweight or obesity due to endocrine disorders , such as Cushing's syndrome;
- Diagnosis of other endocrine disorders with clinical significance, including but not limited to hyperthyroidism or hypothyroidism, thyroid nodules (imaging shows TI-RADSe class 3), thyroid cancer, or personal or family history of type 2 multiple endocrine tumor syndrome (MEN2), etc.
- Diagnosis of cardiovascular or cerebrovascular diseases with clinical significance within 6 months prior to screening, including but not limited to acute stroke, acute coronary syndrome, heart failure, arrhythmia, etc.
- Diagnosis of severe gastrointestinal diseases, including but not limited to inflammatory bowel disease, active ulcers, irritable bowel syndrome, celiac disease, dyspepsia, diabetic gastroparesis, diabetic diarrhea, clinical gastric emptying abnormalities (such as pyloric obstruction), etc.
- History of major gastrointestinal surgery (except cholecystectomy or appendectomy), or metabolic surgery, or plan to undergo major surgery during the study period;

... and 2 more (see full listing online)

Locations (1 total)

China-Japan Friendship Hospital, Beijing, Beijing Municipality, China