

Open Label, Dose Escalation, Repeat Dose Study Evaluating YCT-529 in Healthy Males

NCT06542237

Status	RECRUITING
Phase	Phase 1, Phase 2
Sponsor	YourChoice Therapeutics, Inc.
Enrollment	88 participants

Key Eligibility Criteria

Inclusion (16)

- Participant in good health as confirmed by physical examination, medical history, and clinical laboratory tests.
- Participant must provide written informed consent.
- Participant must be willing and able to communicate and participate in the whole study.
- Participant is 28 to 70 years of age (inclusive) at the time of consent.
- Participant has decided to have a vasectomy and is waiting for the procedure or participant, in the opinion of the investigator, has made a firm decision not to father children in the future.

... and 11 more (see full listing online)

Exclusion (24)

- Men participating in another clinical study involving an investigational drug within the last 30 days prior to the first dosing or less than 5 elimination half-lives prior to first dosing, whichever is longer.
- Clinically significant abnormal physical and/or laboratory findings at Screening
- Abnormal serum chemistry values at screening or admission, that indicate liver or kidney dysfunction or that may be considered clinically significant as determined by the PI, except for bilirubin $>24 \mu\text{mol/L}$ and ALT, AST, GGT and ALP 2-fold above the upper limit of normal. Volunteers with known Gilbert's syndrome will be excluded if total bilirubin is $\geq 1.5 \times \text{ULN}$.
- Evidence of renal impairment at screening, as indicated by an estimated eGFR of $<80 \text{ mL/min/1.73 m}^2$ using the 2021 CKD-EPI Creatinine Equation (https://www.kidney.org/professionals/kdoqi/gfr_calculator).
- Use of androgens and selective androgen receptor modulators (SARMs) within 90 days before first screening visit.

... and 19 more (see full listing online)

Locations (1 total)

New Zealand Clinical Research (NZCR), Grafton, Auckland, New Zealand

<https://clinicaltrials.gov/study/NCT06542237>

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