

Evaluation of Miricorilant on Liver Fat in Patients With MASLD

NCT06947304

Status	RECRUITING
Phase	Phase 1
Sponsor	Corcept Therapeutics
Enrollment	8 participants

Key Eligibility Criteria

Inclusion (8)

- AST \geq 17 U/L for women and AST \geq 20 U/L for men. The AST inclusion criterion does not apply to participants with an eligible historical liver biopsy performed within 12 months of screening showing one of the following:
 - NAFLD Activity Score (NAS) \geq 4 (with at least 1 point in each subcomponent of steatosis, inflammation, and ballooning) and NASH Clinical Research Network (CRN) fibrosis score of F0 OR
 - NAS \geq 3 (with at least 1 point in each subcomponent of steatosis, inflammation, and ballooning) and NASH CRN fibrosis score of F1 OR
 - NAS \geq 2 (with at least 1 point in subcomponent of ballooning or inflammation) and a NASH CRN fibrosis score of F2-3
 - MRI-PDFF with \geq 8% steatosis; this assessment must be performed within 4 weeks of the Baseline Visit.
- ... and 3 more (see full listing online)

Exclusion (12)

- Participation in another clinical trial for MASH or weight loss (e.g., GLP-1 receptor agonists) within the last 3 months.
 - Participation in any other clinical trial within the last 3 months or 5 half-lives of the treatment, whichever is longer.
 - Women who are pregnant, planning to become pregnant, or lactating.
 - BMI $<$ 18 kg/m² or \geq 45 kg/m².
 - Significant alcohol consumption exceeding 20 g/day for women or 30 g/day for men within 1 year prior to screening.
- ... and 7 more (see full listing online)

Locations (1 total)

University of Missouri, Columbia, Missouri, United States

<https://clinicaltrials.gov/study/NCT06947304>

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