

# Safety, Tolerability, and Efficacy of AXA1125 in NASH With Fibrosis

**Status:** Recruiting

## Eligibility Criteria

**Age:** 18 years and over

This study is NOT accepting healthy

**Healthy Volunteers:** volunteers

### Inclusion Criteria:

- Willing to participate in the study and provide written informed consent.
- Male and female adults aged > 18 years.
- Must have NASH and fibrosis on a liver biopsy sample
- If a historical liver biopsy is used for Screening, obtained within 6 months prior to Screening;
- Subjects may have a diagnosis of T2DM

---

### Exclusion Criteria:

- History or presence of liver disease (other than NAFLD or NASH)
- History or presence of cirrhosis and/or history or presence of hepatic decompensation

## Conditions & Interventions

### Interventions:

Drug: AXA1125, Drug: Placebo

### Conditions:

Non Alcoholic Steatohepatitis (NASH)

### Keywords:

Steatosis, Lobular inflammation, Ballooning, Liver biopsy, Liver fat, Liver stiffness, NASH, Aminio Acids, Fibrosis

## More Information

**Contact(s):** Margaret Koziel, MD - [clinicaltrials@axcellahealth.com](mailto:clinicaltrials@axcellahealth.com)

**Principal Investigator:** Siddiqui, Mohammad, S

**Phase:** Phase 2

**IRB**

**Number:** HM20022234

**System ID:** NCT04880187

---

Thank you for choosing StudyFinder. Please visit <http://studyfinder.cctr.vcu.edu> to find a Study which is right for you and contact [ctrrecruit@vcu.edu](mailto:ctrrecruit@vcu.edu) if you have questions or need assistance.