

# Comparison of Two sTRAtegies For the Non-Invasive Diagnosis of advanCed Liver Fibrosis in NAFLD

NCT04681573

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Status	RECRUITING
Phase	Not Applicable
Sponsor	University Hospital, Angers
Enrollment	1,045 participants

## Key Eligibility Criteria

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### Inclusion (8)

- Presence of NAFLD as defined by :
  - The presence of liver steatosis as assessed by ultrasonography (bright liver) or magnetic resonance imaging/spectroscopy (fat fraction  $\geq 5.6\%$ ) or Controlled Attenuation Parameter ( $\geq 248$  dB/m)
  - The absence of steatosis-inducing drugs (systemic corticosteroids, methotrexate, amiodarone, tamoxifen)
  - The absence of excessive alcohol consumption ( $< 210$  g/week in men or  $< 140$  g/week in women)
  - The absence of other causes of chronic liver disease (chronic viral hepatitis B or C, hemochromatosis, auto-immune hepatitis, primary biliary cholangitis, primary sclerosing cholangitis, Wilson disease alpha-1-antitrypsin deficiency).
- ... and 3 more (see full listing online)

### Exclusion (9)

- Decompensated cirrhosis (ascites, variceal bleeding, hepatic encephalopathy, liver failure, hepato-renal syndrome)
  - Hepatocellular carcinoma
  - Inability to safely undergo liver biopsy
  - Participation in other intervention study with drug protocol treatment in progress at the time of inclusion or within one month prior to inclusion in the study.
  - Pregnant, breastfeeding or parturient woman
- ... and 4 more (see full listing online)

## Locations (20 total)

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University Hospital of Angers, Angers, France  
University Hospital of Besançon, Besançon, France  
Avicenne Hospital (Greater Paris University Hospitals), Bobigny, France  
... and 17 more locations

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<https://clinicaltrials.gov/study/NCT04681573>

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