

COOLEY- Study: ACute on ChrOnic Liver FailurE Using the CYtosorb Device

NCT06079021

Status	RECRUITING
Phase	Not Applicable
Sponsor	University Hospital, Antwerp
Enrollment	20 participants

Key Eligibility Criteria

Inclusion (8)

- adult patients (≥ 18 years) admitted to the University Hospital of Antwerp (UZA), Belgium.
- Written informed consent from patient or if not possible due to encephalopathy (> grade 2): legal representative
- acute-on-chronic liver failure (ACLF) grade ≥ 2:
- Acute decompensation event (identifiable trigger)
- Hepatic encephalopathy grade ≥ 2
- ... and 3 more (see full listing online)

Exclusion (4)

- known patient will against participation in the study or against the measures applied in the study
- a decision made prior to inclusion to stop further treatment of the patient within the next 24 hours
- no complete remission of malignancy including hepatocellular carcinoma within the past 12 months
- ongoing intermittent or CRRT before study inclusion

Locations (1 total)

UZA, Edegem, Antwerp, Belgium