# Confocal laser endomicroscopy in patients with non-resolving acute respiratory failure

No registrations found.

Ethical review	Positive opinion
Status	Recruiting
Health condition type	-
Study type	Observational non invasive

## **Summary**

#### ID

NL-OMON25385

Source NTR

**Brief title** CAESAR

**Health condition** 

ARDS, COVID19

#### **Sponsors and support**

**Primary sponsor:** Amsterdam UMC **Source(s) of monetary or material Support:** Research grant from Mauna Kea Technologies, Paris, France

#### Intervention

#### **Outcome measures**

#### **Primary outcome**

Identification of specific CLE patterns of the alveolar compartment in invasively ventilated COVID 19 patients

1 - Confocal laser endomicroscopy in patients with non-resolving acute respiratory f ... 9-05-2025

#### Secondary outcome

Description of how CLE patterns change over time within the same patients, Descriptive correlation of CLE patterns with Chest CT, lung ultrasound, cytology and histology (when available), Time of procedure, Proportion of successful imaging.

# **Study description**

#### **Background summary**

Acute respiratory distress syndrome (ARDS) is an infrequent but severe complication of COVID 19. The clinical syndrome of ARDS includes a heterogeneous group of patients with varying underlying pathophysiology. The innovative probe-based imaging techniques 'Confocal Laser Endomicroscopy' (CLE) is a high-resolution optical technique that, combined with conventional bronchoscopy, has been found to provide real-time, near-histology information about the alveolar compartment. We aim to describe different CLE characteristics of ARDS patients.

#### **Study objective**

CLE provides real-time and high detailed information about the characteristics of the alveolar compartment in mechanically ventilated COVID19 patients.

#### **Study design**

24 hours of follow up after CLE procedure

## Contacts

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# **Eligibility criteria**

## **Inclusion criteria**

-  $\geq$ 18 years of age

- Non-resolving acute respiratory failure mandating a standard diagnostic bronchoscopy with bronchoalveolar lavage

## **Exclusion criteria**

- Inability and willingness to provide informed consent by family-members
- Patients on extra corporal membrane oxygenation (ECMO)

# Study design

## Design

Study type:	Observational non invasive	
Intervention model:	Other	
Allocation:	Non controlled trial	
Masking:	Open (masking not used)	
Control:	N/A , unknown	

#### Recruitment

NL	
Recruitment status:	Recruiting
Start date (anticipated):	21-02-2021
Enrollment:	15
Type:	Anticipated

## **IPD** sharing statement

Plan to share IPD: Undecided

# **Ethics review**

Positive opinion Date: Application type:

21-02-2021 First submission

# **Study registrations**

## Followed up by the following (possibly more current) registration

No registrations found.

## Other (possibly less up-to-date) registrations in this register

No registrations found.

#### In other registers

**Register** NTR-new Other ID NL9281 METC AMC : 2020 294

# **Study results**