Clinical validation of lung ultrasound for the diagnosis of COVID-19

No registrations found.

Ethical review Positive opinion **Status** Recruiting

Health condition type -

Study type Observational non invasive

Summary

ID

NL-OMON22931

Source NTR

Brief title

TBA

Health condition

COVID19

Sponsors and support

Primary sponsor: NA

Source(s) of monetary or material Support: NA

Intervention

Outcome measures

Primary outcome

Lung ultrasound sensitivity to predict COVID-19 diagnosis

Secondary outcome

Lung ultrasound specificity to predict COVID-19 diagnosis

1 - Clinical validation of lung ultrasound for the diagnosis of COVID-19 8-05-2025

Study description

Background summary

We will investigate the value of ultrasound as a first-line examination tool in the process of diagnosing COVID-19. We aim to estimate the test characteristics of artificial intelligence methods on lung ultrasound images for diagnosis of COVID-19 on patients entering the hospital emergency department with COVID-19 symptoms. The outcomes of this study will be relevant for hospitals, but also for any other situations or regions where PCR testing or CT scanning is less available.

Study objective

Artificial intelligence methods on lung ultrasound images is able to predict the presence of lung related signs of COVID-19 with a sensitivity around 90%. Artificial intelligence methods on lung ultrasound images is able to predict the presence of lung related signs of COVID-19 with a specificity around 80%.

Study design

after entering the emergency department

Intervention

Lung ultrasound

Contacts

Public

Maastricht University Medical Center Ronald Henry

043-3871562

Scientific

Maastricht University Medical Center Ronald Henry

043-3871562

Eligibility criteria

Inclusion criteria

- patients who enter the emergency department
- 18 years of age or older
- capacitated (able to make a reasonable judgement of their own interests with regards to the study)
- with (newly developed) symptoms of COVID-19 (fever or chills, cough, shortness of breath or difficulty breathing, new loss of taste or smell, sore throat, congestion or runny nose)
- signed informed consent

Exclusion criteria

- pregnancy
- contra-indications for ultrasound

Study design

Design

Study type: Observational non invasive

Intervention model: Other

Allocation: Non controlled trial

Control: N/A , unknown

Recruitment

NL

Recruitment status: Recruiting
Start date (anticipated): 01-02-2021

Enrollment: 150

Type: Anticipated

IPD sharing statement

Plan to share IPD: Undecided

Ethics review

Positive opinion

Date: 04-02-2021

Application type: 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 ID

NTR-new NL9247

Other Not-WMO METC azM: METC2020-2229

Study results