

The Digital Doc in the Emergency Department: The solution for patient care in the Covid-19 pandemic?

No registrations found.

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

Summary

ID

NL-OMON19962

Source

Nationaal Trial Register

Brief title

DigitalDoc

Health condition

Suspicion for Covid-19

Sponsors and support

Primary sponsor: none

Source(s) of monetary or material Support: none

Intervention

Outcome measures

Primary outcome

Number of PPE used per patient

Secondary outcome

Study description

Background summary

Quasi experimental study with before-after design of digital communication in the ED. Contact isolation of patients has negative impact on patient's psychological well-being, safety and satisfaction. Furthermore, it is advised to limit patient contact because the supplies of personal protective equipment (PPE) are limited. Digital consultation is therefore recommended. Digital consultation, or telemedicine, means a healthcare worker has video contact with the patient while the patient is in a contact isolation room. The effect of digital consultation has not been studied yet.

Aim: Does digital consultation reduce PPE use in the ED in patients treated in contact isolation during the Covid-19 pandemic, and does digital consultation improve patient's well-being (e.g. Hospital Anxiety, patient satisfaction)?

Study objective

Telemedicine will reduce personal protection equipment. Furthermore it could benefit patients' well-being

Study design

The primary outcome is measured during ED visit. Physician and nurse note how often they go inside the room in PPE, till the patient is discharged.

The secondary outcome is measured at the end of the ED visit. The patient is asked to fill in the HAD-A anxiety score and Pickers Patient Satisfaction-15 Questionairry

Intervention

Telemedicine by Ipads

Contacts

Public

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Scientific

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Eligibility criteria

Inclusion criteria

Patients treated in the ED in contact isolation.

Exclusion criteria

15L O2/min. cognitive impairment, GCS<15, <18years

Study design

Design

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

Recruitment

NL	
Recruitment status:	Recruiting
Start date (anticipated):	08-04-2020
Enrollment:	50
Type:	Anticipated

IPD sharing statement

Plan to share IPD: Undecided

Ethics review

Positive opinion

Date: 12-04-2020

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	NL8522
Other	METC Máxima MC : N20.042

Study results