The applicability of rapid serological and molecular testing for Corona SARS-2 in a Dutch population during the COVID-19 outbreak.

Published: 11-05-2020
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nvt

Ethical review	Not approved
Status	Will not start
Health condition type	Viral infectious disorders
Study type	Observational invasive

Summary

ID

NL-OMON49670

Source
ToetsingOnline

Brief title
Research performance point of care diagnostics Corona SARS-2

Condition

- Viral infectious disorders

Synonym
Corona SARS-2, COVID-19

Research involving
Human

Sponsors and support

Primary sponsor: Ministerie van Defensie
Source(s) of monetary or material Support: Ministerie van Defensie

Intervention

Keyword: COVID-19, Molecular diagnostics, Rapid test, Serology

Outcome measures

Primary outcome

nvt

Secondary outcome

nvt

Study description

Background summary

nvt

Study objective

nvt

Study design

nvt

Study burden and risks

nvt

Contacts

Public
Ministerie van Defensie

Korte molenweg 3
Amsterdam 3941PW
NL
Scientific

Ministerie van Defensie

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NL

Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age
Adults (18-64 years)
Elderly (65 years and older)

Inclusion criteria

zie onder

Exclusion criteria

zie onder

Study design

Design

Study type:	Observational invasive
Intervention model:	Other
Allocation:	Non-randomized controlled trial
Masking:	Open (masking not used)
Control:	Active

Primary purpose:	Health services research

Recruitment

NL

Recruitment status:	Will not start
Enrollment:	300
Type:	Anticipated

Ethics review

Not approved

Date:	11-05-2020
Application type:	First submission
Review commission:	METC NedMec

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
CCMO	NL73721.041.20