Monitoring the course of the COVID-19 epidemic in Estonia

Submission date	Recruitment status No longer recruiting	Prospectively registered		
14/03/2021		[X] Protocol		
Registration date	Overall study status Completed	Statistical analysis plan		
16/03/2021		[X] Results		
Last Edited 21/09/2021	Condition category Infections and Infestations	[] Individual participant data		

Plain English summary of protocol

Background and study aims

COVID-19 is a condition caused by the coronavirus (called SARS-CoV-2) that was first identified in late 2019. The ongoing pandemic is an unprecedented global emergency. Epidemic risk management is a highly complex task in which sets of measures need to be put in place and coordinated at local, national, and international levels to minimize health and economic consequences. This complex and monumental task is highly knowledge/evidence-dependent. There are many unanswered questions on the novel SARS-Cov-2 virus including the immune response and pathogenesis at the individual level. However, the answers to very similar questions at the population level (scale of exposure/infection in the population, the effect of community mitigation measures implemented) are crucial for an effective response to the pandemic and to inform future strategies.

The main aim of this study is to estimate the prevalence of symptomatic and asymptomatic COVID-19 in the general population and how this varies over time.

Who can participate?

People selected at random from the Estonian population registry will be invited for participation

What does the study involve?

Participants will be asked to visit a national covid testing center and fill in a web-based questionnaire and undergo nasopharyngeal swabbing (collected from the back of the nose and throat).

What are the possible benefits and risks of participating?

The possible benefits are that participants will get results from tests for SARS-CoV-2. The main disadvantage of taking part is the time and inconvenience of nasopharyngeal swabbing.

Where is the study run from?

University of Tartu (Estonia), and delivered by Kantar-Emor AS, OÜ Medicum Eriarstiabi, and SYNLAB Eesti OÜ (Estonia)

When is the study starting and how long is it expected to run for? March 2020 to December 2023

Who is funding the study? Ministry of Education and Research (Estonia)

Who is the main contact? Unfortunately, this study is not recruiting public volunteers.

Study website

https://www.ut.ee/en/research/study-prevalence-coronavirus-estonia

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

Nil known

IRAS number

ClinicalTrials.gov number

Nil known

Secondary identifying numbers

Nil known

Study information

Scientific Title

COVID-19 active monitoring program in Estonia

Acronym

COVEST

Study objectives

The main aim of this observational study is to estimate the prevalence of symptomatic and asymptomatic SARS-CoV-2 infection in the general population and how this varies over time.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 13/04/2020, Research Ethics Committee of the University of Tartu (Grant Office, University of Tartu, Raekoja plats 9, 51004 Tartu, Estonia; +372 (0)737 6215; eetikakomitee@ut. ee), ref: 310/T-1

Study design

Surveillance study based on repeated cross-sectional surveys of the general population (recruited via stratified random sampling)

Primary study design

Observational

Secondary study design

Cross sectional study

Study setting(s)

Other

Study type(s)

Screening

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

COVID-19 (SARS-CoV-2 infection)

Interventions

Randomly selected consenting adults are asked to visit national testing centers for a nasopharyngeal swab and fill out a web-based questionnaire.

Intervention Type

Other

Primary outcome measure

SARS-CoV-2 prevalence measured using SARS-CoV-2 RNA RT-PCR at each of the cross-sectional study rounds (every month in 2020, six times a year in 2021, 2022/23 to be decided)

Secondary outcome measures

The proportion of symptomatic cases among people testing positive for SARS-CoV-2, measured using a web-based questionnaire at each of the cross-sectional study rounds (every month in 2020, six times a year in 2021, 2022/23 to be decided)

Overall study start date

17/03/2020

Completion date

30/12/2023

Eligibility

Key inclusion criteria

- 1. Adult, male or female
- 2. Willing and able to give informed consent for participation in the study

Participant type(s)

All

Age group

Adult

Sex

Both

Target number of participants

2000 in one cross-sectional study

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

23/04/2020

Date of final enrolment

30/11/2023

Locations

Countries of recruitment

Estonia

Study participating centre University of Tartu

Ravila 19 Tartu Estonia 50411

Sponsor information

University of Tartu

Sponsor details

Ülikooli 18 Tartu Estonia 50090 +372 (0)737 5100 info@ut.ee

Sponsor type

University/education

Website

http://www.ut.ee/et

ROR

https://ror.org/03z77qz90

Funder(s)

Funder type

Government

Funder Name

Ministry of Education and Research (Estonia)

Results and Publications

Publication and dissemination plan

- 1. Publication of positivity rates and behavioural characteristics after each recruitment round to the funding agency and the Estonian governmental task group on COVID-19
- 2. Planned publications in high-impact peer-reviewed journals

Intention to publish date

31/01/2024

Individual participant data (IPD) sharing plan

The data-sharing plans for the current study are unknown and will be made available at a later date

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<u>Protocol file</u>			06/04/2021	No	No
Preprint results	non-peer reviewed 1st year results	13/09/2021	21/09/2021	No	No