Second serosurveillance study in The Netherlands for the evaluation of the Dutch National Immunisation Programme: the PIENTER 2 study

Submission date	Recruitment status No longer recruiting	Prospectively registered		
27/06/2007		☐ Protocol		
Registration date 27/06/2007	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited 18/03/2013	Condition category Infections and Infestations	Individual participant data		

Plain English summary of protocol

Not provided at time of registration

Study website

http://www.pienterproject.nl

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

Acronym

PIENTER 2

Study objectives

The age-specific seroprevalence of a cross-section of the Dutch population for the vaccinations used in the Dutch National Immunisation Programme.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approval received from the local ethics committee (Stichting Therapeutische Evaluatie Geneesmiddelen [STEG]) on the 11th October 2005 (ref: R05-044).

Study design

Observational, cross-section survey

Primary study design

Observational

Secondary study design

Cross-section survey

Study setting(s)

Not specified

Study type(s)

Screening

Participant information sheet

Health condition(s) or problem(s) studied

Vaccination and the National Immunisation Program, infectious diseases

Interventions

A blood sample and a questionnaire regarding health perception, diseases (including sexually transmitted), vaccination data.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

The age-specific seroprevalence (immunity) for diseases included in the National Immunisation Program of the Dutch general population.

As each participant is sampled only once, there will be only one timepoint measurement per person. The timepoint for the complete study population varies from February 2006 to June 2007. Because the participants represent a cross section of the Dutch population aged 0 to 80, we will be able to estimate the seroprevalence in the Dutch population in 2006-2007.

Secondary outcome measures

- 1. The age-specific seroprevalence against other infectious diseases, in particular those that might be vaccine preventable in the near future and against those diseases with a frequent subclinical course
- 2. The age-specific seroprevalence amongst the allochtonous populations (additional sample)
- 3. The age-specific seroprevalence amongst non-vaccinated orthodox reformed individuals (additional sample)

As each participant is sampled only once, there will be only one timepoint measurement per person. The timepoint for the complete study population varies from February 2006 to June 2007. Because the participants represent a cross section of the Dutch population aged 0 to 80, we will be able to estimate the seroprevalence in the Dutch population in 2006-2007.

Overall study start date

06/02/2006

Completion date

19/06/2007

Eligibility

Key inclusion criteria

- 1. Subject is part of the study sample
- 2. Aged 0 79 years old
- 3. Has received a personal invitation for the study
- 4. Subject has given written informed consent before start of the study

Participant type(s)

Patient

Age group

Adult

Sex

Not Specified

Target number of participants

7700

Key exclusion criteria

Subject is an employee of the National Institute of Public Health and Environmental Protection (RIVM).

Date of first enrolment

06/02/2006

Date of final enrolment

19/06/2007

Locations

Countries of recruitment

Netherlands

Study participating centre

National Institute of Public Health and Environmental Protection (RIVM)

Bilthoven Netherlands 3720 BA

Sponsor information

Organisation

National Institute of Public Health and Environmental Protection (RIVM) (The Netherlands)

Sponsor details

P.O. Box 1 Bilthoven Netherlands 3720 BA +31 (0)30 274 9111 info@rivm.nl

Sponsor type

Government

Website

http://www.rivm.nl/en/

ROR

https://ror.org/01cesdt21

Funder(s)

Funder type

Government

Funder Name

The Netherlands Ministry of Health, Welfare and Sport (The Netherlands)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	19/10/2012		Yes	No
Results article	results	01/04/2014		Yes	No