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Screening for Chlamydia trachomatis (CT) with routine Pap smears in general practice: a randomized controlled trial

Submission date	Recruitment status	[_] Prospectively registered
09/09/2005	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
31/10/2005	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
28/10/2021	Infections and Infestations	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof Francis Bowden

Contact details
The Canberra Hospital
P.O. Box 11
Woden
Australia
2606
+61 (0)2 6244 2063
frank.bowden@act.gov.au

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
268058

Study information

Scientific Title
Screening for Chlamydia trachomatis (CT) with routine Pap smears in general practice: a randomized controlled trial

Acronym
GPPaCTS

Study objectives
General practices randomized to CT/Pap screening for women aged 16 to 39 years (intervention) will have an increased participation rate for chlamydia screening compared to those practices randomised to offer 'usual care' (control).

Ethics approval required
Old ethics approval format

Ethics approval(s)
Added as of 30/07/2007:
This study was approved by:
1. The ACT Health Human Research Ethics Committee (ETH.10/03)
2. The Australian National University Ethics Committee (ref: 2003/270)

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Screening

Participant information sheet

Health condition(s) or problem(s) studied
Chlamydia trachomatis

Interventions
Routine offering of testing for Chlamydia trachomatis at the time of regular Pap smear versus usual care.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Participation rates in screening for Chlamydia trachomatis.

Secondary outcome measures
Uptake of Pap smears in women in target age range.

Overall study start date
01/11/2004

Completion date
01/03/2006

Eligibility

Key inclusion criteria
Women between the ages of 16 and 39 years who have ever been sexually active and who present to their general practitioner.

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
1440

Key exclusion criteria
Women who have never been sexually active.

Date of first enrolment
01/11/2004

Date of final enrolment
01/03/2006

Locations

Countries of recruitment
Australia

Study participating centre

The Canberra Hospital
Woden
Australia
2606

Sponsor information

Organisation
Australian National Health and Medical Research Council

Sponsor details
Office of NHMRC (MDP 100)
GPO Box 9848
Canberra
Australia
2601
+61 (0)2 6289 1555
research@nhmrc.gov.au

Sponsor type
Research council

Website
http://www.nhmrc.gov.au/

ROR
https://ror.org/011kf5r70

Funder(s)

Funder type
Research council

Funder Name
Australian National Health and Medical Research Council: 268058

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

Not provided at time of registration

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		21/01/2008	28/10/2021	Yes	No