

Tapering off Inhaled Corticosteroids in Asthma patients after Reduction of Allergens

Submission date 19/12/2005	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 19/12/2005	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 25/08/2009	Condition category Respiratory	☐ Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof C.P. Schayck, van

Contact details
University Maastricht
Care and Public Health Research Institute - CAPHRI
Department of General Practice
P.O. Box 616
Maastricht
Netherlands
6200 MD
+31 (0)43 3882446
onno.vanschayck@hag.unimaas.nl

Additional identifiers

Protocol serial number
NTR370

Study information

Scientific Title

Acronym

TICARA

Study objectives

Allergen avoidance allows tapering off inhaled corticosteroids (ICS) in house dust mite allergic asthma patients

Ethics approval required

Old ethics approval format

Ethics approval(s)

Received from local medical ethics committee

Primary study design

Interventional

Study design

Multicentre randomised double blind placebo controlled parallel group trial

Study type(s)

Other

Health condition(s) or problem(s) studied

Asthma, Allergy

Interventions

All patients have been trained to use a self-management plan to adjust the dose of inhaled corticosteroids to symptoms and peak expiratory flow value.

After a run-in period of 3 months the intervention period with placebo controlled allergen avoidance started.

The participants in the intervention group received house dust mite impermeable covers for mattress, pillow and bedding.

The control group received placebo, house dust mite permeable, covers.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Use of inhaled corticosteroids

Key secondary outcome(s)

1. Asthma control
2. Symptoms (dyspnoea, wheezing, coughing)
3. Peak flow parameters (morning peak flow, peak flow variability)

Completion date

01/12/2004

Eligibility

Key inclusion criteria

1. Age 16-60 years
2. Treatment for asthma by the GP
3. Use of inhaled corticosteroids
4. Allergy for house dust mite allergens

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Serious diseases other than asthma with a low survival rate
2. Other diseases, which influence bronchial symptoms and/or lung function
3. Exacerbation within one month before the start of the study
4. The use of oral steroids or inhaled cromoglycates
5. Use of house dust mite impermeable mattress/bedding covers
6. Allergy to cats or dogs while keeping these pets

Date of first enrolment

01/01/1999

Date of final enrolment

01/12/2004

Locations

Countries of recruitment

Netherlands

Study participating centre

University Maastricht

Maastricht

Netherlands

6200 MD

Sponsor information

Organisation

Care and Public Health Research Institute (CAPHRI), University Maastricht (Netherlands)

ROR

<https://ror.org/02jz4aj89>

Funder(s)**Funder type**

Research organisation

Funder Name

Netherlands Organisation for Health Research and Development (ZonMw) (Netherlands)

Alternative Name(s)

Netherlands Organisation for Health Research and Development

Funding Body Type

Private sector organisation

Funding Body Subtype

Other non-profit organizations

Location

Netherlands

Funder Name

Netherlands Asthma Foundation (Netherlands)

Funder Name

Boehringer Ingelheim BV (Netherlands)

Funder Name

AstraZeneca BV (Netherlands)

Results and Publications

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	01/04/2006		Yes	No