# Evaluation of the effect of the diagnostic and therapeutic advices given by an Astma/COPD-service on the referring general practitioners and their patients.

No registrations found.

**Ethical review** Not applicable

**Status** Recruitment stopped

Health condition type -

**Study type** Interventional

## **Summary**

#### ID

NL-OMON27336

Source

NTR

**Brief title** 

N/A

Intervention

#### **Outcome measures**

#### **Primary outcome**

Do general practitioners assess better diagnoses, according tot the NHG-standards, in Asthma/COPD-patients: do more patients have a correct diagnosis, do less patients have a wrong or incomplete diagnosis or no diagnosis at all?

Do general practitioners give their patients a better treatment, according to the NHGstandards: do more patients get the right medication, do less patients get wrong or unnecessary medication. Do patients receive more information about their disease and about selfmanagement

Do patients have a better compliance, do they better follow the advices about stop smoking, exercise, do they have fewer complaints and a better quality of life.

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## **Secondary outcome**

- 1. Validity and reliability of the assessment and the reports of the Asthma/COPD-service;
- 2. Epidemiological data about the prevalence of Asthma/COPD in general practices, the (incorrect) use of medication of in general practice;
- 3. Costs of a support as given by the Asthma/COPD-service, related to the diagnostic and therapeutic gain in patient care.

## **Study description**

#### **Background summary**

N/A

#### Study objective

N/A

## Study design

N/A

#### Intervention

The intervention in the trial was the diagnostic and therapeutic advice offered to the general practitioners by the Asthma/COPD-service in Eindhoven. These advices are a new but already introduced facility for general practitioners and their patients. This facility is in the process of implementation. General practitioners in the research project who, after randomisation, are eligible to first use the facility are considered to be the intervention group. (N=17). General practitioners who don't get the full support of the Asthma/COPD-service yet (N=17) are considered to be the control group.

Each general practice participates in the research project for two years. In this period the patients of the intervention group receive two yearly follow-up consultations at the Asthma/COPD service, on request of their general practitioner. Medical history and spirometry is performed. A lung specialist assesses by protocol the written results of these measurements and sends a structured report to the general practitioner. This report includes a diagnosis or an advice for further diagnostic examinations, and advices for treatment.

In the control group the general practitioners can have spirometry performed for their patients in the way they are used to. However, they don't get the full report of the Asthma/COPD-service, only the description of the lung function.

## **Contacts**

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## **Eligibility criteria**

## **Inclusion criteria**

All general practices in and around the city of Eindhoven could be included as long as they used the facilities of the Diagnostic Centre, which the Asthma/COPD-service is a part of. Although, they should not have any experience with the support of the Asthma/COPD-service.

They also were excluded in case they had employed a "praktijkondersteuner" (specialized nurse or assistant) for the Asthma/COPD diseasemanagement.

In the intervention as well as in the control practices, all patients 12 years and older that have airway complaints could participate as long as they were not treated by a lung specialist.

### **Exclusion criteria**

All general practices in and around the city of Eindhoven could be included as long as they used the facilities of the Diagnostic Centre, which the Asthma/COPD-service is a part of. Although, they should not have any experience with the support of the Asthma/COPD-service.

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In the intervention as well as in the control practices, all patients 12 years and older that have airway complaints could participate as long as they were not treated by a lung specialist.

# Study design

## **Design**

Study type: Interventional

Intervention model: Parallel

Allocation: Randomized controlled trial

Masking: Open (masking not used)

Control: Active

### Recruitment

NL

Recruitment status: Recruitment stopped

Start date (anticipated): 01-01-2003

Enrollment: 1000

Type: Actual

## **Ethics review**

Not applicable

Application type: Not applicable

# **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

RegisterIDNTR-newNL286NTR-oldNTR324

Other : Intern budget number 30.95.01.04.b

ISRCTN ISRCTN45174826

# **Study results**

## **Summary results**

N/A