Evaluatie van het myAirCoach zelfmanagement ondersteuningssysteem voor mensen met astma in vergelijking met gebruikelijke zorg: een pragmatisch gerandomiseerd onderzoek (myAirCoach: evaluation campaign).

No registrations found.

Ethical review Positive opinion **Status** Recruiting

Health condition type

Study type Interventional

Summary

ID

NL-OMON23774

Source

NTR

Brief title

myAirCoach EC

Health condition

Patients with a doctors-diagnosis of asthma from outpatient clinics and from general practices in the region of London and Manchester in the UK and Leiden in The Netherlands

Sponsors and support

Primary sponsor: Leiden University Medical Center

Source(s) of monetary or material Support: European Union Horizon 2020

Intervention

Outcome measures

Primary outcome

Asthma Control assessed through the Asthma Control Questionnaire (ACQ)

Secondary outcome

Asthma exacerbations, lung function, FeNO, quality of life, patient utilities, costs, user acceptance

Study description

Background summary

DESIGN

Multi-centre, pragmatic randomized control trial

AIMS

To assess the myAirCoach system in a real life environment and determine whether it provides clinical benefit (improvements in asthma control) to patients with asthma.

OUTCOME MEASURES

Primary: Asthma Control (assessed through the ACQ)

Secondary: Lung function, asthma exacerbation, quality of life, Symptoms, costs

POPULATION

Maximal 60 subjects in the Netherlands, minimal 30 subjects in the UK, 90 subjects in total

ELIGIBILITY

Adult patients with uncontrolled asthma

TREATMENT

myAirCoach intervention

DURATION

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3-9 months (variable follow-up length); A sequential phased study so that patients will therefore be involved for a minimum 3 months, up to a maximum 9 months

Study objective

Self-management support by the myAirCoach system as an adjunct to usual care improves asthma control in patients with asthma, compared to usual care.

Study design

Type of study: Multi-centre, pragmatic randomized controlled trial, in which participants will be randomized in a 1:1 ratio to myAirCoach self-management support as an adjunct to usual care (myAirCoach group) or to usual care alone (usual care group).

Duration: 3-9 months (staggered enrolment with variable follow-up duration). A sequential phased study so that patients will therefore be involved for a minimum 3 months, up to a maximum 9 months.

Intervention

Usual care group

Patients in the usual care group will be provided with regular (face-to-face) care with their health care professionals.

myAirCoach group

In addition to usual care patients in the myAirCoach group will be provided with self-management support via the myAirCoach system. This system consists of the several devices and mHealth and web-applications for patients as well as a web-application for health care professionals.

Contacts

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Eligibility criteria

Inclusion criteria

- Age 18+
- · Clinician diagnosis of asthma
- Asthma treatment step 2-5 need for regular controller medication (≥6 months of the year), this equates to 2 or more ICS prescriptions per year
- Poor asthma control (ACQ>0.75) and/or one-or-more exacerbations or hospital visit in the previous year due to asthma
- Ownership of a mobile phone compatible with the myAirCoach system

Exclusion criteria

• Unable to understand English or Dutch, as appropriate

Study design

Design

Study type: Interventional

Intervention model: Parallel

Allocation: Randomized controlled trial

Masking: Open (masking not used)

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Control: N/A, unknown

Recruitment

NL

Recruitment status: Recruiting
Start date (anticipated): 16-02-2018

Enrollment: 90

Type: Anticipated

Ethics review

Positive opinion

Date: 02-05-2018

Application type: First submission

Study registrations

Followed up by the following (possibly more current) registration

ID: 46602

Bron: ToetsingOnline

Titel:

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register ID

NTR-new NL6944 NTR-old NTR7200

CCMO NL62699.058.17 OMON NL-OMON46602

Study results	