Expert opinion on the impact of inhaler choice on climate change and personalised healthcare

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Administrative details

EU PAS number	
EUPAS39494	
Study ID	
39495	
DARWIN EU® study	
No	
Study countries	
Australia	
Austria	
Belgium	
Canada	

Denmark
Finland
France
Germany
Greece
Iceland
Ireland
Italy
Japan
Korea, Republic of
Netherlands
Norway
Portugal
Singapore
Spain
Sweden
Switzerland
United Kingdom
United Kingdom (Northern Ireland)

Study description

This project aims to provide an opinion piece on choice of inhaler delivery method and the impact of on climate change and personalised healthcare. It has the following objectives: 1. Identify experience and preferences of patients and healthcare professionals (HCPs) of inhaler choice/change in relation to climate change and personalized healthcare. 2. Gather expert opinion and consensus on (a) costs to environment (b) impact on personalized healthcare. 3. An extensive literature review covering current discussion on the above topics. This research will offer expert opinion and consensus of physicians and health care workers on: • (a) Costs to environment: The impact of inhaler

choice and switching inhaler delivery system on climate change, as well as short-term vs long-term solutions for reduction of impact. The research will also gather information from patients on: • (b) Personal impact: The impact of changing medication that has affected their personalized healthcare plan and inhaler use.

Study status

Finalised

Research institutions and networks

Networks

Despiratory Effectiveness Crown (DEC)
Respiratory Effectiveness Group (REG)
Belgium
☐ Denmark
France
Germany
Greece
Hungary
Italy
☐ Netherlands
Spain
Sweden
United Kingdom
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Contact details

Study institution contact

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Study contact

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Primary lead investigator

Omar Usmani

Primary lead investigator

Study timelines

Date when funding contract was signed

Planned: 01/12/2020

Actual: 11/01/2021

Study start date

Planned: 25/01/2021

Actual: 16/02/2021

Date of final study report

Planned: 31/01/2022

Actual: 17/02/2021

Sources of funding

• Pharmaceutical company and other private sector

More details on funding

AstraZeneca, Boehringer Ingelheim, Chiesi, Kindeva

Study protocol

REG inhaler choice Research Proposal_February2021.pdf(343.21 KB)

Regulatory

Was the study required by a regulatory body?

No

Is the study required by a Risk Management Plan (RMP)?

Not applicable

Methodological aspects

Study type

Study type list

Study topic:

Disease /health condition

Medical device

Study type:

Non-interventional study

Scope of the study:

Other

If 'other', further details on the scope of the study

Exploratory perspective/opinion

Data collection methods:

Primary data collection

Main study objective:

This research aims to gather patient-centric expert opinion to deliver consensus on the impact of switching inhaler type on climate change and the suggestion of green alternatives to switching inhaler types, as well as measure the impact of switching on patients and their personalised healthcare plan.

Study Design

Non-interventional study design

Other

Non-interventional study design, other

Exploratory, Survey

Study drug and medical condition

Medical condition to be studied

Asthma

Population studied

Short description of the study population

Patient inclusion criteria

- Clinically stable asthma or COPD diagnosis
- Prescribed inhaler medication
- Age >18 years
- Have switched inhaler type in the last 1 year

Exclusion criteria

- Unable to access questionnaire
- Unable to understand the electronic questionnaire process
- Using a non-MDI/DPI/SMI device

Age groups

Adults (18 to < 46 years)

Adults (46 to < 65 years)

Adults (65 to < 75 years)

Adults (75 to < 85 years)

Adults (85 years and over)

Special population of interest

Other

Special population of interest, other

Chronic obstructive pulmonary disease (COPD) or asthma patients

Estimated number of subjects

300

Study design details

Data analysis plan

Questionnaires: Descriptive statistics will be used to identify the strength of opinion/knowledge of healthcare workers and patients. Subgroup analysis will be used, as appropriate, between healthcare worker types and differences in patient age, time since inhaler switch, demographic, number of exacerbations, and whether the patient is diagnosed with asthma or COPD.

Data management

ENCePP Seal

The use of the ENCePP Seal has been discontinued since February 2025.

The ENCePP Seal fields are retained in the display mode for transparency but are no longer maintained.

Data sources

Data sources (types)

Other

Data sources (types), other

Prospective patient-based data collection

Use of a Common Data Model (CDM)

CDM mapping

No

Data quality specifications

Unknown Check completeness Unknown

Check stability

Check conformance

Unknown

Check logical consistency

Unknown

Data characterisation

Data characterisation conducted

No