

Viacoram® - focus on blood pressure target

Submission date 11/09/2015	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 28/10/2015	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 31/10/2019	Condition category Circulatory System	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Arterial hypertension (high blood pressure) can cause premature death. In Germany, about 30% of women and 33% of men aged between 18-79 have hypertension. In adults aged between 70-79, some 75% of people of both sexes have the condition. Based on the public importance of the disease, a timely return to a normal blood pressure should always be the aim. If the recommended initial non-pharmacological (drug) measures (for example, lifestyle changes) are not successful, a drug treatment should be started. Viacoram® is a single-pill combination of two drugs, amlodipine and perindopril, both used for treatment of arterial hypertension. It is used for the treatment of adult patients with arterial hypertension. The aim of this observational study is to gain information on the use of Viacoram® in outpatients with hypertension. This study will focus on the change of the blood pressure, time to achieve normal blood pressure, change in other treatments the patients may be taking as well as whether they keep taking the drug (therapy adherence) and an assessment of general tolerability.

Who can participate?

Adult outpatients with arterial hypertension that have been recommended treatment with Viacoram®.

What does the study involve?

All patients in the trial involved in the study are treated with Viacoram®. They are asked to come to three follow up visits after 1, 2 and 3 months. During these visits a routine practice investigation is carried out and a case report form regarding blood pressure, other diseases the patient is suffering from and what other medications they are taking is filled out by the treating physician. The patients are asked to fill out the patient questionnaire to therapy adherence at the first and the last visit.

What are the possible benefits and risks of participating?

The patients have no particular benefits or risks of participating in this observational study. The treatment given to the patient corresponds to the standard medical routine and is completely independent of the study as the study is strictly non-interventional. Patients are free to withdraw from the study at any time without giving a particular reason.

Where is the study run from?

Study will be carried out by approximately 1000 general practitioners across Germany.

When is study starting and how long is it expected to run for?
September 2015 to April 2016

Who is funding the study?
Servier Deutschland, GmbH (Germany)

Contact information

Type(s)
Public

Contact name
Dr Peter Martinka

Contact details
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Additional identifiers

Protocol serial number
IC4-05985-161-DEU

Study information

Scientific Title
Viacoram® - focus on blood pressure target. An open single-arm non-interventional observational study in hypertensive outpatients treated with Viacoram® in Germany.

Acronym
VIACORAM-BPT

Study objectives
Aim of this non-interventional study is to gain information on the use of Viacoram® in hypertensive outpatients in line with the marketing authorization under the conditions of daily practice in Germany. Special focus of the study is the change in the blood pressure (office, ABPM, central BP parameters), time to achieve the BP target, change in the concomitant treatments and therapy adherence as well as assessment of general tolerability.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Freiburg Ethics Committee International ("Freiburger Ethik Kommission International), 27/07/2015, ref: 015/1441

Study design

Prospective, open, multicentre, single-arm, observational, non-interventional study.

Primary study design

Observational

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Essential hypertension

Interventions

This is strictly a non-interventional observational study. Only interventions in line with and not exceeding the standard medical routine are allowed. All patients involved in the study will be treated with Viacoram® under the circumstance that the decision about the treatment was done prior to the study initiation. After the inclusion the patients will be asked to come to three follow up visits after 1, 2 and 3 months. During these visits a routine practice investigation will be carried out and a case report form regarding blood pressure, concomitant diseases and co-medications will be filled out by the treating physician. The patients will be asked to fill out the patient questionnaire to therapy adherence at the first and the last visit.

Intervention Type

Other

Primary outcome(s)

Office blood pressure measured within the daily medical routine at inclusion, after approx. 1 month, 2 months and at study final examination after approx. 3 months.

Key secondary outcome(s)

1. Ambulatory BP monitoring only in measured as part of the medical routine at inclusion, after approx. 1 month, 2 months and at study final examination after approx. 3 months
2. Central BP parameters only in measured as part of the medical routine at inclusion, after approx. 1 month, 2 months and at study final examination after approx. 3 months
3. Assessment of therapy adherence measured by means of the Hill-Bone-Scale

Completion date

30/04/2016

Eligibility**Key inclusion criteria**

Adult patients with essential hypertension with currently not controlled blood pressure eligible for the treatment with Viacoram® for whom the treatment decision was done prior to study inclusion.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Total final enrolment

1814

Key exclusion criteria

N/A

Date of first enrolment

21/09/2015

Date of final enrolment

31/12/2015

Locations

Countries of recruitment

Germany

Study participating centre

General practitioners

Germany

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Sponsor information

Organisation

Servier Deutschland GmbH

ROR

<https://ror.org/05wk4ae67>

Funder(s)

Funder type

Industry

Funder Name

Servier Deutschland GmbH (Germany)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not expected to be made available

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/03/2018	31/10/2019	Yes	No