Pattern of use of intravitreal drugs with antiangiogenic properties for age-related macular degeneration and other vascular retinopathies (Anti-VEGF drugs)

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Administrative details

Study description

EU PAS number	
EUPAS15749	
Study ID	
16444	
DARWIN EU® study	
No	
Study countries	
Italy	

This is a drug utilization study of antiVEGF drugs for the treatment of age related macular degeneration and other vascular retinopathies in clinical practice, in the Tuscany region of Italy, from 2011 to 2015

Study status

Finalised

Research institutions and networks

Institutions



Contact details

Study institution contact

Rosa Gini rosa.gini@ars.toscana.it

Study contact

rosa.gini@ars.toscana.it

Primary lead investigator

Rosa Gini

Primary lead investigator

Study timelines

Date when funding contract was signed

Planned: 09/03/2016 Actual: 03/03/2016

Study start date

Planned: 05/09/2016 Actual: 05/09/2016

Data analysis start date

Planned: 12/09/2016 Actual: 12/09/2016

Date of final study report

Planned: 17/10/2016 Actual: 28/11/2016

Sources of funding

Other

More details on funding

Self-funded by ARS

Study protocol

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

Human medicinal product

Study type:

Non-interventional study

Scope of the study:

Drug utilisation

Data collection methods:

Main study objective:

To describe the pattern of use of anti-VEGF drugs for the treatment of agerelated macular degeneration and other vascular retinopathies in clinical practice in Tuscany, Italy

Study Design

Non-interventional study design

Other

Non-interventional study design, other

Pharmacoepidemiological study

Study drug and medical condition

Anatomical Therapeutic Chemical (ATC) code

(L01XC07) bevacizumab

bevacizumab

(S01CB01) dexamethasone

dexamethasone

(S01LA03) pegaptanib

pegaptanib

(S01LA04) ranibizumab

ranibizumab

(S01LA05) aflibercept

aflibercept

Medical condition to be studied

Diabetic retinopathy

Diabetic retinal oedema

Age-related macular degeneration

Population studied

Short description of the study population

Patients with diabetic retinopathy, retinal oedema or age-related macular degeneration in the Tuscany region of Italy, from 2011 to 2015.

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

Patients with diabetic retinopathy, retinal oedema or age-related macular degeneration

Estimated number of subjects

15000

Study design details

Outcomes

Number of injections per year and intra-injections interval, switchingless than 5 injections in the first year

Data analysis plan

Outcomes will be compared across incident users of the drugs. Subgroup analysis will be performed in patients with a sufficient number of contacts with ophthalmic services (interval between consecutive contacts not longer than 3 months) and in patient with at least 3 injections

Documents

Study results

report antiVEGF.pdf (213.74 KB)

Study publications

Farmaco-utilizzazione di farmaci per il trattamento della degenerazione macular...

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 source(s)

ARS Toscana

Data sources (types)

Administrative healthcare records (e.g., claims)

Drug dispensing/prescription data

Other

Data sources (types), other

Disease-specific exemptions from copayment

Use of a Common Data Model (CDM)

CDM mapping

No

Data quality specifications

Check conformance

Unknown

Check completeness

Unknown

Check stability

Unknown

Check logical consistency

Unknown

Data characterisation

Data characterisation conducted

Unknown