



# Macugen

pegaptanib

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WITHDRAWN

This medicine is now withdrawn from use in the European Union.

#### Overview

The marketing authorisation for Macugen has been withdrawn at the request of the marketingauthorisation holder.



Macugen: EPAR - Summary for the public (PDF/520.99 KB)

First published: 31/05/2007 Last updated: 27/06/2019

Available languages (21) >



More detail is available in the summary of product characteristics

This EPAR was last updated on 27/06/2019

## Authorisation details

Name

Macugen



Agency product number

EMEA/H/C/000620

Active substance

pegaptanib

International non-proprietary name (INN) or common name

pegaptanib

Therapeutic area (MeSH)

Wet Macular Degeneration

Anatomical therapeutic chemical (ATC) code

S01LA03

Marketing-authorisation holder

PharmaSwiss Ceska Republika s.r.o

Revision

15

Date of issue of marketing authorisation valid throughout the European Union

31/01/2006

Contact address

Jankovcova 1569/2c Lighthouse 17000 Prague 7 Czech Republic

## **Product information**

20/07/2016 Macugen - EMEA/H/C/000620 - IB/0065



Macugen: EPAR - Product Information (PDF/832.53 KB)

First published: 03/08/2009 Last updated: 27/06/2019

Available languages (24) >





#### **Contents**

- Annex I Summary of product characteristics
- Annex IIA Manufacturing-authorisation holder responsible for batch release
- Annex IIB Conditions of the marketing authorisation
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Please note that the size of the above document can exceed 50 pages.

You are therefore advised to be selective about which sections or pages you wish to print.



Macugen: EPAR - All Authorised presentations (PDF/457.96 KB)

First published: 05/02/2008 Last updated: 27/06/2019

Available languages (21) V





Macugen : EPAR - Conditions imposed on member states for safe and effective use - Annex IV ( $PDF/477.37\ KB$ )

First published: 13/07/2007 Last updated: 27/06/2019

Available languages (21) 🗸



## Pharmacotherapeutic group

Ophthalmologicals

## Therapeutic indication

Macugen is indicated for the treatment of neovascular (wet) age-related macular degeneration (AMD).

## **Assessment history**

## Changes since initial authorisation of medicine



Macugen: EPAR - Procedural steps taken and scientific information after authorisation (PDF/688.92 KB)



First published: 03/08/2009 Last updated: 27/06/2019

## Initial marketing-authorisation documents



Macugen: EPAR - Procedural steps taken before authorisation (PDF/482.92 KB)

First published: 31/05/2007 Last updated: 27/06/2019



Macugen: EPAR - Scientific Discussion (PDF/984.11 KB)

First published: 31/05/2007 Last updated: 27/06/2019

# More information on Macugen



• Macugen: Withdrawn application

• Macugen: Paediatric investigation plan

#### **CONTACT**

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