



EUROPEAN MEDICINES AGENCY  
SCIENCE MEDICINES HEALTH

EMA/169085/2014

## European Medicines Agency decision

P/0100/2014

of 11 April 2014

on the granting of a product specific waiver for alirocumab (EMEA-001169-PIP02-13) in accordance with Regulation (EC) No 1901/2006 of the European Parliament and of the Council

**Only the English text is authentic.**



# European Medicines Agency decision

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on the granting of a product specific waiver for alirocumab (EMA-001169-PIP02-13) in accordance with Regulation (EC) No 1901/2006 of the European Parliament and of the Council

The European Medicines Agency,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 1901/2006 of the European Parliament and of the Council of 12 December 2006 on medicinal products for paediatric use and amending Regulation (EEC) No. 1768/92, Directive 2001/20/EC, Directive 2001/83/EC and Regulation (EC) No 726/2004<sup>1</sup>,

Having regard to Regulation (EC) No 726/2004 of the European Parliament and of the Council of 31 March 2004 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency<sup>2</sup>,

Having regard to the application submitted by sanofi-aventis recherche & développement on 13 December 2013 under Article 13 of Regulation (EC) No 1901/2006,

Having regard to the opinion of the Paediatric Committee of the European Medicines Agency, issued on 21 March 2014 in accordance with Article 13 of Regulation (EC) No 1901/2006,

Having regard to Article 25 of Regulation (EC) No 1901/2006,

Whereas:

- (1) The Paediatric Committee has given an opinion on the granting of a product specific waiver.
- (2) It is therefore appropriate to adopt a decision granting a waiver.

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<sup>1</sup> OJ L 378, 27.12.2006, p.1.

<sup>2</sup> OJ L 136, 30.4.2004, p. 1.

Has adopted this decision:

**Article 1**

A waiver for alirocumab, solution for injection, subcutaneous use, the details of which are set out in the opinion of the Paediatric Committee of the European Medicines Agency annexed hereto, together with its appendices, is hereby granted.

**Article 2**

This decision is addressed to sanofi-aventis recherche & développement, 1 avenue pierre brossollette, 91385 - Chilly-Mazarin Cedex, France.

Done at London, 11 April 2014

For the European Medicines Agency  
Guido Rasi  
Executive Director  
(Signature on file)



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EMA/PDCO/10431/2014

## Opinion of the Paediatric Committee on the granting of a product-specific waiver

EMA-001169-PIP02-13

### Scope of the application

**Active substance(s):**

Alirocumab

**Condition(s):**

Treatment of mixed dyslipidaemia

**Pharmaceutical form(s):**

Solution for injection

**Route(s) of administration:**

Subcutaneous use

**Name/corporate name of the PIP applicant:**

Sanofi-aventis recherche & développement

### Basis for opinion

Pursuant to Article 13 of Regulation (EC) No 1901/2006 as amended, sanofi-aventis recherche & développement submitted to the European Medicines Agency on 13 December 2013 an application for a product-specific waiver on the grounds set out in Article 11 of said Regulation for the above mentioned medicinal product.

The procedure started on 21 January 2014.

A meeting with the Paediatric Committee took place on 20 March 2014.



## Opinion

1. The Paediatric Committee, having assessed the waiver application in accordance with Article 13 of Regulation (EC) No 1901/2006 as amended, recommends as set out in the appended summary report:

- to grant a product-specific waiver for all subsets of the paediatric population and the above mentioned condition(s) in accordance with Article 11(1)(c) of said Regulation, on the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments for paediatric patients.

The Icelandic and the Norwegian Paediatric Committee members agree with the above-mentioned recommendation of the Paediatric Committee.

2. The grounds for the granting of the waiver are set out in Annex I.

This opinion is forwarded to the applicant and the Executive Director of the European Medicines Agency, together with its annex and appendix.

London, 21 March 2014

On behalf of the Paediatric Committee  
Dr Dirk Mentzer, Chairman  
(Signature on file)

## **Annex I**

### **Grounds for the granting of the waiver**

# 1. Waiver

## ***1.1. Condition: treatment of mixed dyslipidaemia***

The waiver applies to:

- the paediatric population from birth to less than 18 years;
- for solution for injection, subcutaneous use;
- on the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments.