European Medicines Agency

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EUROPEAN MEDICINES AGENCY DECISION

of 11 December 2007

on the application for a product specific waiver for Everolimus (EMEA-000019-PIP01-07) in accordance with Regulation (EC) No 1901/2006 of the European Parliament and of the Council as amended

(ONLY THE ENGLISH TEXT IS AUTHENTIC)

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on the application for a product specific waiver for Everolimus (EMEA-000019-PIP01-07) in accordance with Regulation (EC) No 1901/2006 of the European Parliament and of the Council as amended

THE EUROPEAN MEDICINES AGENCY,

Having regard to the Treaty establishing the European Community,

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 as amended and amending Regulation (EEC) No. 1768/92, Directive 2001/20/EC, Directive 2001/83/EC and Regulation (EC) No 726/2004[1],

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[2],

Having regard to the application submitted by Novartis Europharm Ltd. United Kingdom on 23 July 2007 under Article 13 of Regulation (EC) No 1901/2006 as amended,

Having regard to the opinion of the Paediatric Committee of the European Medicines Agency, formulated on 26 October 2007 in accordance with Article 13 of Regulation (EC) No 1901/2006 as amended,

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

WHEREAS:

(1) The Paediatric Committee has given a positive opinion,

(2) It is therefore appropriate to adopt a Decision granting a waiver.

HAS ADOPTED THIS DECISION:

[1] OJ L 378, 27.12.2006, p. 1
[2] OJ L 136, 30.4.2004, p. 1

Article 1

A waiver for Everolimus, tablet, oral use, in renal cell carcinoma and pancreatic neuroendocrine tumour, the details of which are set out in the Opinion of the Paediatric Committee the European Medicines Agency annexed hereto, together with its appendices, is hereby granted.

Article 2

This decision is addressed to Novartis Europharm Limited, Wimblehurst Road, Horsham, West Sussex RH12 5AB, United Kingdom.

Done at London, 11 December 2007

For the European Medicines Agency
Thomas Lönngren
Executive Director

Signature: (On file)

European Medicines Agency
Pre-authorisation Evaluation of Medicines for Human Use

EMEA/575610/2007/EN
EMEA-000019-PIP01-07

POSITIVE OPINION OF THE PAEDIATRIC COMMITTEE ON A PRODUCT-SPECIFIC WAIVER

Scope of the application

Active substance:
Everolimus

Condition(s):
Renal cell carcinoma and pancreatic neuroendocrine tumour

Pharmaceutical form(s):
Tablet

Route(s) of administration:
Oral use

Name/corporate name of the waiver applicant:
Novartis Europharm Ltd. United Kingdom

Basis for opinion

Pursuant to Article 13(1) of Regulation (EC) No 1901/2006 of 12 December 2006, as amended, Novartis Europharm Ltd. submitted for agreement to the EMEA on 23 July 2007 an application for a waiver pursuant to Article 11(1) of Regulation (EC) No 1901/2006, as amended for the above mentioned medicinal product.

The procedure started on 30 August 2007.

Opinion

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 waiver for all subsets of the paediatric population and all above mentioned conditions in accordance with Article 11(1)(b) of Regulation (EC) No 1901/2006, as amended, on the grounds that the disease or condition for which the specific medicinal product is intended occurs only in adult populations.

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

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

London, 26 October 2007

On behalf of the Paediatric Committee
Dr Daniel Brasseur, Chairman

Signature: (On file)