



European Medicines Agency

Doc. Ref. EMEA/218895/2009
P/75/2009

EUROPEAN MEDICINES AGENCY DECISION

of 20 April 2009

**on the granting of a product specific waiver for pasireotide
(EMEA-000464-PIP01-08) in accordance with Regulation (EC) No 1901/2006
of the European Parliament and of the Council as amended**

(ONLY THE ENGLISH TEXT IS AUTHENTIC)

**EUROPEAN MEDICINES AGENCY DECISION
of 20 April 2009**

**on the granting of a product specific waiver for pasireotide
(EMEA-000464-PIP01-08) 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 726/2004¹,

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 Agency²,

Having regard to the application submitted by Novartis Europharm Ltd on 4 December 2008 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, issued on 6 March 2009 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 an opinion on the granting of a product specific waiver,
- (2) It is therefore appropriate to adopt a Decision granting a waiver.

¹ OJ L 378, 27.12.2006, p.1

² OJ L 136, 30.4.2004, p. 1

HAS ADOPTED THIS DECISION:

Article 1

A waiver for pasireotide, solution for injection, powder and solvent for suspension for injection, subcutaneous use, intramuscular 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 Novartis Europharm Ltd, Wimblehurst Road, RH12 5AB, Horsham West Sussex, England.

Done at London, 20 April 2009

For the European Medicines Agency
Thomas Lönngren
Executive Director

(Signature on file)



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

Doc. Ref. EMEA/PDCO/105288/2009
EMEA-000464-PIP01-08

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

Scope of the application

Active substance(s):

Pasireotide

Condition(s):

Pituitary dependant Cushing's disease
Overproduction of pituitary ACTH
Pituitary dependant hyperadrenocorticism

Pharmaceutical form(s):

Solution for injection
Powder and solvent for suspension for injection

Route(s) of administration:

Subcutaneous use
Intramuscular use

Name/corporate name of the waiver applicant:

Novartis Europharm Ltd

Basis for opinion

Pursuant to Article 13 of Regulation (EC) No 1901/2006 as amended, Novartis Europharm Ltd submitted to the EMA on 4 December 2008 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 8 January 2009.

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 all 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 do 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 Agency, together with its annex and appendix.

London, 6 March 2009

On behalf of the Paediatric Committee
Dr Daniel Brasseur, Chairman

(Signature on file)

ANNEX I
 GROUNDS FOR THE GRANTING OF THE WAIVER

GROUNDS FOR THE GRANTING OF THE WAIVER

- **Condition**

Pituitary dependant Cushing's disease

The waiver applies to all subsets of the paediatric population from birth to less than 18 years of age for solution for injection and powder and solvent for suspension for injection for subcutaneous use and for intramuscular use on the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments.

- **Condition**

Overproduction of pituitary ACTH

The waiver applies to all subsets of the paediatric population from birth to less than 18 years of age for solution for injection and powder and solvent for suspension for injection for subcutaneous use and for intramuscular use on the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments.

- **Condition**

Pituitary dependant hyperadrenocorticism

The waiver applies to all subsets of the paediatric population from birth to less than 18 years of age for solution for injection and powder and solvent for suspension for injection for subcutaneous use and for intramuscular use on the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments.