European Medicines Agency decision
P/0244/2012

of 22 October 2012


Only the English text is authentic.
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P/0244/2012

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The European Medicines Agency,

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


Having regard to the application submitted by Menarini Ricerche SpA on 5 June 2012 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 7 September 2012 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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Has adopted this decision:

**Article 1**

A waiver for dexketoprofen (trometamol) / tramadol (hydrochloride), film-coated tablet, oral 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 Menarini Ricerche SpA, Via Tito Speri 10, 00040 Pomezia, Roma, Italy.

Done at London, 22 October 2012

For the European Medicines Agency
Guido Rasi
Executive Director
(Signature on file)
Opinion of the Paediatric Committee on the granting of a product-specific waiver
EMEA-001302-PIP01-12

Scope of the application

Active substance(s):
Dexketoprofen (trometamol) / tramadol (hydrochloride)

Condition(s):
Treatment of acute pain

Pharmaceutical form(s):
Film-coated tablet

Route(s) of administration:
Oral use

Name/corporate name of the PIP applicant:
Menarini Ricerche SpA

Basis for opinion

Pursuant to Article 13 of Regulation (EC) No 1901/2006 as amended, Menarini Ricerche SpA submitted to the European Medicines Agency on 5 June 2012 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 11 July 2011.
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, 7 September 2012

On behalf of the Paediatric Committee
Dr Daniel Brasseur, Chairman
(Signature on file)
Annex I

Grounds for the granting of the waiver
1. Waiver

1.1. Condition: Treatment of moderate to severe acute pain

The waiver applies to:

- All subsets of the paediatric population from birth to less than 18 years of age;
- For film-coated tablet, oral use;
- On the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments.