



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

EMA/101962/2013

European Medicines Agency decision

P/0054/2013

of 25 March 2013

on the acceptance of a modification of an agreed paediatric investigation plan for mepolizumab (EMA-000069-PIP02-10-M01) in accordance with Regulation (EC) No 1901/2006 of the European Parliament and of the Council

Disclaimer

This Decision does not constitute entitlement to the rewards and incentives referred to in Title V of Regulation (EC) No 1901/2006.

Only the English text is authentic.



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on the acceptance of a modification of an agreed paediatric investigation plan for mepolizumab (EMEA-000069-PIP02-10-M01) 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 726/2004¹,

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 European Medicines Agency's decision P/219/2011 issued on 26 September 2011,

Having regard to the application submitted by Glaxo Group Limited on 19 November 2012 under Article 22 of Regulation (EC) No 1901/2006 proposing changes to the agreed paediatric investigation plan with a deferral and a waiver,

Having regard to the opinion of the Paediatric Committee of the European Medicines Agency, issued on 8 February 2013, in accordance with Article 22 of Regulation (EC) No 1901/2006,

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

Whereas:

- (1) The Paediatric Committee of the European Medicines Agency has given an opinion on the acceptance of changes to the agreed paediatric investigation.
- (2) It is therefore appropriate to adopt a decision on the acceptance of changes to the agreed paediatric investigation plan.

¹ OJ L 378, 27.12.2006, p.1.

² OJ L 136, 30.4.2004, p. 1.

Has adopted this decision:

Article 1

Changes to the agreed paediatric investigation plan for mepolizumab, powder for solution for infusion, intravenous use, subcutaneous use, are hereby accepted in the scope set out in the opinion of the Paediatric Committee of the European Medicines Agency annexed hereto, together with its appendices.

Article 2

This decision is addressed to Glaxo Group Limited, GlaxoWellcome House, Berkeley Avenue, UB6 0NN – Greenford, United Kingdom.

Done at London, 25 March 2013

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

EMA/PDCO/756851/2012

Opinion of the Paediatric Committee on the acceptance of a modification of an agreed Paediatric Investigation Plan EMA-000069-PIP02-10-M01

Scope of the application

Active substance(s):

Mepolizumab

Condition(s):

Treatment of asthma

Pharmaceutical form(s):

Powder for solution for infusion

Route(s) of administration:

Intravenous use

Subcutaneous use

Name / corporate name of the PIP applicant:

Glaxo Group Limited

Basis for opinion

Pursuant to Article 22 of Regulation (EC) No 1901/2006 as amended, Glaxo Group Limited submitted to the European Medicines Agency on 19 November 2012 an application for modification of the agreed paediatric investigation plan with a deferral and a waiver as set out in the European Medicines Agency's decision P/219/2011 issued on 26 September 2011.

The application for modification proposed changes to the agreed paediatric investigation plan.

The procedure started on 12 December 2012.

A meeting with the Paediatric Committee took place on 6 February 2013.

Scope of the modification

Some measures and/or timelines of the Paediatric Investigation Plan have been modified.

Opinion

1. The Paediatric Committee, having assessed the application in accordance with Article 22 of Regulation (EC) No 1901/2006 as amended, recommends as set out in the appended summary report:
 - to agree to changes to the paediatric investigation plan in the scope set out in the Annex I of this opinion.

The Norwegian Paediatric Committee member agrees with the above-mentioned recommendation of the Paediatric Committee.

2. The measures and timelines of the paediatric investigation plan are set out in the Annex I.

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

London, 8 February 2013

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

Annex I

The subset(s) of the paediatric population and condition(s) covered by the waiver and the measures and timelines of the agreed Paediatric Investigation Plan

1. Waiver

1.1. Condition: treatment of asthma

The waiver applies to:

- The paediatric population from birth to less than 6 years of age;
- for powder for solution for infusion, intravenous use and subcutaneous use;
- on the grounds that the specific medicinal product does not represent a significant therapeutic benefit as clinical studies are not feasible.

2. Paediatric Investigation Plan

2.1. Condition: treatment of asthma

2.1.1. Indication targeted by the PIP

Reduction of severe asthma exacerbations.

2.1.2. Subset(s) of the paediatric population concerned by the paediatric development

From 6 to less than 18 years of age.

2.1.3. Pharmaceutical form(s)

Powder for solution for infusion.

2.1.4. Studies

Area	Number of studies	Description
Quality	1	Study 1 Development of age-appropriate strength for intravenous or subcutaneous use.
Non-clinical	0	Not applicable.
Clinical	4	Study 2 Double-blind, randomised, multi-centre, placebo-controlled, parallel-group study to evaluate efficacy and safety of mepolizumab as add-on therapy in children from 12 to less than 18 years old (and in adults) with severe uncontrolled asthma. Study 3 Double-blind, randomised, multi-centre, placebo-controlled, parallel-group study to evaluate efficacy of mepolizumab to reduce steroid use as add-on therapy in children from 12 to less than 18 years old (and in adults) with

		severe uncontrolled asthma. Study 4 Double-blind, randomised, multi-centre, placebo-controlled, parallel-group study to evaluate efficacy of mepolizumab to reduce asthma exacerbations rate during fall and winter as add-on therapy in children from 6 to less than 12 years old with severe uncontrolled asthma. Study 5 Open-label, multi-centre extension to evaluate long-term safety and efficacy of mepolizumab in children from 6 to less than 12 years old with severe uncontrolled asthma.
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3. Follow-up, completion and deferral of PIP

Concerns on potential long term safety issues in relation to paediatric use:	Yes
Date of completion of the paediatric investigation plan:	By January 2016
Deferral for one or more measures contained in the paediatric investigation plan:	Yes