

EMA/719224/2014

European Medicines Agency decision

P/0317/201

of 12 December 2014

on the acceptance of a modification of an agreed paediatric investigation plan for vedolizumab (Entyvio) (EMA-000645-PIP01-09-M02) 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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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/145/2010 issued on 30 July 2010 and the decision P/0053/2013 issued on 20 March 2013,

Having regard to the application submitted by Takeda Pharma A/S on 26 August 2014 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 14 November 2014, 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 plan and to the deferral.
- (2) It is therefore appropriate to adopt a decision on the acceptance of changes to the agreed paediatric investigation plan, including changes to the deferral.

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 vedolizumab (Entyvio), powder for concentrate for solution for infusion, intravenous use, including changes to the deferral, 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 Takeda Pharma A/S, Dybendal Alle 10, 2630 – Taastrup, Denmark.

Done at London, 12 December 2014

For the European Medicines Agency
Zaide Frias
Head of Division (ad interim)
Human Medicines Research and Development Support
(Signature on file)



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

EMA/PDCO/529183/2014 **Corr**

Opinion of the Paediatric Committee on the acceptance of a modification of an agreed Paediatric Investigation Plan

EMA-000645-PIP01-09-M02

Scope of the application

Active substance(s):

Vedolizumab

Invented name:

Entyvio

Condition(s):

Treatment of Crohn's disease

Treatment of ulcerative colitis

Authorised indication(s):

See Annex II

Pharmaceutical form(s):

Powder for concentrate for solution for infusion

Route(s) of administration:

Intravenous use

Name / corporate name of the PIP applicant:

Takeda Pharma A/S

Information about the authorised medicinal product:

See Annex II



Basis for opinion

Pursuant to Article 22 of Regulation (EC) No 1901/2006 as amended, Takeda Pharma A/S submitted to the European Medicines Agency on 26 August 2014 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/145/2010 issued on 30 July 2010 and the decision P/0053/2013 issued on 20 March 2013.

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

The procedure started on 16 September 2014.

Scope of the modification

Some measures and 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 and to the deferral in the scope set out in the Annex I of this opinion.

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

2. The measures and timelines of the paediatric investigation plan and the subset(s) of the paediatric population and condition(s) covered by the waiver 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 annexes and appendix.

London, 14 November 2014

On behalf of the Paediatric Committee
Dr Dirk Mentzer, 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 (PIP)

1. Waiver

1.1. Condition: treatment of Crohn's disease

The waiver applies to:

- children from birth to less than 4 years,
- for powder for concentrate for solution for infusion, intravenous use,
- on the grounds that the specific medicinal product is likely to be unsafe.

1.2. Condition: treatment of ulcerative colitis

The waiver applies to:

- children from birth to less than 4 years,
- for powder for concentrate for solution for infusion, intravenous use,
- on the grounds that the specific medicinal product is likely to be unsafe.

2. Paediatric Investigation Plan

2.1. Condition: treatment of Crohn's disease

2.1.1. Indication(s) targeted by the PIP

Treatment of moderately to severely active Crohn's disease.

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

- From 4 to less than 18 years of age.

2.1.3. Pharmaceutical form(s)

Powder for concentrate for solution for infusion.

2.1.4. Measures

Area	Number of measures	Description
Quality	0	Not applicable.
Non-clinical	0	Not applicable.
Clinical	2	Study 1) MLN0002-2003 Open label clinical pharmacology study to determine the pharmacokinetics, pharmacodynamics, safety and tolerability of vedolizumab in paediatric patients with inflammatory bowel disease (IBD)

Area	Number of measures	Description
		Study 2) MLN0002-3025 Randomised, double-blind, placebo-controlled two-dose, three-arm, multicentre study of the induction and maintenance of clinical response and remission by vedolizumab in paediatric patients with moderate to severe Crohn's disease.

2.2. Condition: treatment of ulcerative colitis

2.2.1. Indication(s) targeted by the PIP

Treatment of moderately to severely active ulcerative colitis.

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

From 4 to less than 18 years of age.

2.2.3. Pharmaceutical form(s)

Powder for concentrate for solution for infusion.

2.2.4. Measures

Area	Number of measures	Description
Quality	0	Not applicable.
Non-clinical	0	Not applicable.
Clinical	2	Study 1) MLN0002-2003 Open label clinical pharmacology study to determine the pharmacokinetics, pharmacodynamics, safety and tolerability of vedolizumab in paediatric patients with inflammatory bowel disease (IBD) Study 3) MLN0002-3024 Randomised, double-blind, placebo-controlled two-dose, three-arm, multicentre study of the induction and maintenance of clinical response and remission by vedolizumab in paediatric patients with moderate to severe ulcerative colitis.

3. Follow-up, completion and deferral of PIP

Concerns on potential long term safety/efficacy issues in relation to paediatric use:	Yes
Date of completion of the paediatric investigation plan:	By June 2021
Deferral for one or more measures contained in the paediatric investigation plan:	Yes