EMA/190114/2019

European Medicines Agency decision

P/0130/2019

of 17 April 2019

on the acceptance of a modification of an agreed paediatric investigation plan for luspatercept (EMEA-001521-PIP01-13-M03) 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 726/2004[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 European Medicines Agency's decision P/0245/2014 issued on 29 September 2014, the decision P/0219/2017 issued on 9 August 2017 and the decision P/0122/2018 issued on 11 April 2018,

Having regard to the application submitted by Celgene Europe B.V. on 26 November 2018 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 1 March 2019, 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.

(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 luspatercept, powder for solution for injection, 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 Celgene Europe B.V., Winthontlaan 6 N, 3526 KV - Utrecht, The Netherlands.

EMA/PDCO/836819/2018
London, 1 March 2019

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

EMEA-001521-PIP01-13-M03

Scope of the application

Active substance(s):

Luspatercept

Condition(s):

Treatment of myelodysplastic syndromes

Treatment of beta-thalassaemia

Pharmaceutical form(s):

Powder for solution for injection

Route(s) of administration:

Subcutaneous use

Name/corporate name of the PIP applicant:

Celgene Europe B.V.

Basis for opinion

Pursuant to Article 22 of Regulation (EC) No 1901/2006 as amended, Celgene Europe B.V. submitted to the European Medicines Agency on 26 November 2018 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/0245/2014 issued on 29 September 2014, the decision P/0219/2017 issued on 9 August 2017 and the decision P/0122/2018 issued on 11 April 2018.

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

The procedure started on 3 January 2019.

Scope of the modification

Some measures 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 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 annex and appendix.

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 myelodysplastic syndromes

The waiver applies to:

- all subsets of the paediatric population from birth to less than 18 years of age;
- powder for solution for injection, subcutaneous use;
- on the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments.

1.2. Condition

Treatment of beta-thalassaemia

The waiver applies to:

- the paediatric population from birth to less than 6 months;
- powder for solution for injection, subcutaneous use;
- on the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments.

2. Paediatric investigation plan

2.1. Condition

Treatment of beta-thalassaemia

2.1.1. Indication(s) targeted by the PIP

Treatment of anaemia in patients with beta-thalassaemia intermedia and major

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

From 6 months to less than 18 years of age

2.1.3. Pharmaceutical form(s)

Powder for solution for injection

2.1.4. Measures

Area	Number of measures	Description
Quality-related studies	1	**Study 1** Development of an age-appropriate self-administration device for subcutaneous use, allowing sufficient accuracy of dose measurement and delivery in all age groups to be treated.
Non-clinical studies	2	**Study 2** (1383657025333) Dose range-finding juvenile toxicity study **Study 3** (1381154982795) Definitive juvenile toxicity study
Clinical studies	3	**Study 4** Study to evaluate safety and pharmacokinetics of luspatercept in paediatric patients from 6 months to less than 18 years of age with transfusion-dependent beta-thalassaemia. **Study 5** (1406216840211) Two-part, double-blind, randomised, placebo controlled trial to evaluate pharmacokinetics, safety and efficacy of luspatercept in children from 6 months to less than 18 years of age with non-transfusion-dependent beta-thalassaemia. **Study 6** Double-blind, randomised, placebo controlled trial to evaluate safety and efficacy of luspatercept in paediatric patients from 6 months to less than 18 years of age with transfusion-dependent beta-thalassaemia.
Extrapolation, modelling and simulation studies	0	Not applicable
Other studies	0	Not applicable
Other measures	0	Not applicable

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 May 2024
Deferral for one or more measures contained in the paediatric investigation plan:	Yes