



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

EMA/83575/2020

European Medicines Agency decision P/0074/2020

of 21 March 2020

on the acceptance of a modification of an agreed paediatric investigation plan for Vadadustat (EMA-001944-PIP01-16-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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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/0035/2017 issued on 31 January 2017,

Having regard to the application submitted by Akebia Therapeutics, Inc on 28 October 2019 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 31 January 2020, 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 Vadadustat, tablet, age-appropriate oral formulation, oral 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 Akebia Therapeutics, Inc., 245 First Street, Suite 1100, 02142 - Cambridge, MA, USA.



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

EMA/PDCO/610168/2019
Amsterdam, 31 January 2020

Opinion of the Paediatric Committee on the acceptance of a modification of an agreed Paediatric Investigation Plan EMA-001944-PIP01-16-M01

Scope of the application

Active substance(s):

Vadadustat

Condition(s):

Treatment of anaemia due to chronic disorders

Pharmaceutical form(s):

Tablet

Age-appropriate oral formulation

Route(s) of administration:

Oral use

Name/corporate name of the PIP applicant:

Akebia Therapeutics, Inc

Basis for opinion

Pursuant to Article 22 of Regulation (EC) No 1901/2006 as amended, Akebia Therapeutics, Inc submitted to the European Medicines Agency on 28 October 2019 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/0035/2017 issued on 31 January 2017.

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

The procedure started on 3 December 2019.



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 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 anaemia due to chronic disorders

The waiver applies to:

- the paediatric population from birth to less than 4 months of age;
- tablet and age appropriate oral formulation, oral use;
- on the grounds that the specific medicinal product does not represent a significant therapeutic benefit as clinical studies(s) are not feasible.

2. Paediatric investigation plan

2.1. Condition

Treatment of anaemia due to chronic disorders

2.1.1. Indication(s) targeted by the PIP

Treatment of anaemia secondary to chronic kidney disease

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

From 4 months to less than 18 years of age

2.1.3. Pharmaceutical form(s)

Tablet

Age appropriate oral formulation

2.1.4. Measures

Area	Number of measures	Description
Quality-related studies	1	Study 1 Development of an age-appropriate oral dosage form.
Non-clinical studies	1	Study 2 Pilot oral gavage study in juvenile rats Study 3 <i>This study was deleted as a result of procedure EMEA-001944-PIP01-16-M01.</i>

Clinical studies	4	Study 4 Open-label, single-arm, externally controlled trial to evaluate the activity, safety, tolerability, PK and PD of oral vadadustat for the correction of anaemia in children from 4 months to less than 18 years of age with anaemia secondary to chronic kidney disease (CKD). Study 5 Medical record review study to assess the activity and safety of ESA treatment to correct and maintain haemoglobin levels in children from 4 months to less than 18 years of age with anaemia secondary to CKD and naïve to ESA treatment. Study 6 Open-label, single-arm, externally controlled trial to evaluate the activity, safety, tolerability, PK and PD of oral vadadustat for the maintenance treatment of anaemia in children from 4 months to less than 18 years of age with anaemia secondary to CKD receiving ESA treatment. Study 7 Medical record review study to assess the activity and safety of ESA treatment to maintain haemoglobin levels in children from 4 months to less than 18 years of age with anaemia secondary to CKD receiving ESA treatment.
Extrapolation, modelling and simulation studies	1	Study 8 Modelling and simulation study to develop PK/PD simulations for the prediction of exposure and selection of doses to be used in children from 4 months to less than 18 years of age with anaemia secondary to CKD naïve to ESA treatment or receiving ESA treatment.
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 2027
Deferral for one or more measures contained in the paediatric investigation plan:	Yes