

EMA/568463/2021

European Medicines Agency decision

P/0414/2021

of 29 October 2021

on the acceptance of a modification of an agreed paediatric investigation plan for recombinant human A disintegrin and metalloprotease with thrombospondin type-1 motifs 13 (EMEA-001160-PIP01-11-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/0048/2012 issued on 29 February 2012 and the decision P/0324/2017 issued on 31 October 2017,

Having regard to the application submitted by Baxalta Innovations GmbH on 4 June 2021 under Article 22 of Regulation (EC) No 1901/2006 proposing changes to the agreed paediatric investigation plan and proposing a deferral,

Having regard to the opinion of the Paediatric Committee of the European Medicines Agency, issued on 10 September 2021, in accordance with Article 22 of Regulation (EC) No 1901/2006 and Article 21 of said Regulation,

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 on the refusal of a deferral.
- (2) It is therefore appropriate to adopt a decision on the acceptance of changes to the agreed paediatric investigation plan.
- (3) It is therefore appropriate to adopt a decision refusing a 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 recombinant human A disintegrin and metalloprotease with thrombospondin type-1 motifs 13, powder and solvent for solution for injection, intravenous 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

A deferral for recombinant human A disintegrin and metalloprotease with thrombospondin type-1 motifs 13, powder and solvent for solution for injection, intravenous use, the details of which are set out in the opinion of the Paediatric Committee the European Medicines Agency annexed hereto, together with its appendices, is hereby refused.

Article 3

This decision is addressed to Baxalta Innovations GmbH, Industriestrasse 67, 1221 – Vienna, Austria.

EMA/PDCO/325791/2021 Corr
Amsterdam, 10 September 2021

Opinion of the Paediatric Committee on the acceptance of a modification of an agreed Paediatric Investigation Plan EMA-001160-PIP01-11-M02

Scope of the application

Active substance(s):

Recombinant human A disintegrin and metalloprotease with thrombospondin type-1 motifs 13

Condition(s):

Treatment of thrombotic thrombocytopenic purpura (TTP)

Pharmaceutical form(s):

Powder and solvent for solution for injection

Route(s) of administration:

Intravenous use

Name/corporate name of the PIP applicant:

Baxalta Innovations GmbH

Basis for opinion

Pursuant to Article 22 of Regulation (EC) No 1901/2006 as amended, Baxalta Innovations GmbH submitted to the European Medicines Agency on 4 June 2021 an application for modification of the agreed paediatric investigation plan as set out in the European Medicines Agency's decision P/0048/2012 issued on 29 February 2012 and the decision P/0324/2017 issued on 31 October 2017.

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

The procedure started on 13 July 2021.

Scope of the modification

Some timelines of the Paediatric Investigation Plan have been modified.

Opinion

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;
- to refuse a deferral as it does not meet the grounds detailed in Article 20(1) of said Regulation.

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

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 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

Not applicable

2. Paediatric Investigation Plan

2.1. Condition: Treatment of thrombotic thrombocytopenic purpura (TTP)

2.1.1. Indication targeted by the PIP

Treatment of hereditary thrombotic thrombocytopenic purpura (TTP)

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

From birth to less than 18 years of age

2.1.3. Pharmaceutical form(s)

Powder and solvent for solution for infusion

2.1.4. Measures

Area	Number of studies	Description
Quality-related studies	1	Study 1 Development of a paediatric vial
Non-clinical studies	0	Not applicable
Clinical studies	2	Study 2 A Phase 1, prospective, uncontrolled, open-label, multicentre, dose-escalation study evaluating the safety, including immunogenicity, and pharmacokinetics of BAX 930 (rADAMTS13) in congenital thrombotic thrombocytopenic purpura (cTTP) in children from 12 years of age to less than 18 years of age (and in adults). Study 3 Prospective, randomised, active-controlled, open-label, multicenter, 2-period crossover study (rADAMTS13 versus standard of care [SoC], 1:1 randomisation, Periods 1 and 2), followed by a single arm continuation period (rADAMTS13, Period 3) to evaluate the safety and efficacy of rADAMTS13 in the prophylactic and on-demand treatment of patients with severe congenital thrombotic thrombocytopenic purpura cTTP in children from birth to less than 18 years of age (and in adults).

3. Follow-up, completion and deferral of PIP

Concerns on potential long term efficacy and safety issues in relation to paediatric use:	No
Date of completion of the paediatric investigation plan:	By December 2023
Deferral for one or more studies contained in the paediatric investigation plan:	No