EMA/554964/2023

European Medicines Agency decision

P/0518/2023

of 29 December 2023

on the agreement of a paediatric investigation plan and on the granting of a deferral for allogeneic cultured postnatal thymus-derived tissue (EMEA-003496-PIP01-23) 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 Union procedures for the authorisation and supervision of medicinal products for human use and establishing a European Medicines Agency[2],

Having regard to the application submitted by Enzyvant Therapeutics Ireland Limited on 4 August 2023 under Article 16(1) of Regulation (EC) No 1901/2006 also requesting a deferral under Article 20 of said Regulation,

Having regard to the opinion of the Paediatric Committee of the European Medicines Agency, issued on 10 November 2023, in accordance with Article 17 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 agreement of a paediatric investigation plan and on the granting of a deferral.

(2) It is therefore appropriate to adopt a decision agreeing a paediatric investigation plan.

(3) It is therefore appropriate to adopt a decision granting a deferral.

[1] OJ L 378, 27.12.2006, p.1, as amended.
[2] OJ L 136, 30.4.2004, p. 1, as amended.

Has adopted this decision:

Article 1

A paediatric investigation plan for allogeneic cultured postnatal thymus-derived tissue, living tissue equivalent, implant use, the details of which are set out in the opinion of the Paediatric Committee of the European Medicines Agency annexed hereto, together with its appendices, is hereby agreed.

Article 2

A deferral for allogeneic cultured postnatal thymus-derived tissue, living tissue equivalent, implant use, the details of which are set out in the opinion of the Paediatric Committee of the European Medicines Agency annexed hereto, together with its appendices, is hereby granted.

Article 3

This decision is addressed to Enzyvant Therapeutics Ireland Limited, 88 Harcourt Street, D02 DK18 – Dublin, Ireland.

EMA/PDCO/475810/2023
Amsterdam, 10 November 2023

Opinion of the Paediatric Committee on the agreement of a Paediatric Investigation Plan and a deferral

EMEA-003496-PIP01-23

Scope of the application

Active substance(s):

Allogeneic cultured postnatal thymus-derived tissue

Invented name and authorisation status:

See Annex II

Condition(s):

Treatment of congenital athymia

Pharmaceutical form(s):

Living tissue equivalent

Route(s) of administration:

Implant use

Name/corporate name of the PIP applicant:

Enzyvant Therapeutics Ireland Limited

Basis for opinion

Pursuant to Article 16(1) of Regulation (EC) No 1901/2006 as amended, Enzyvant Therapeutics Ireland Limited submitted for agreement to the European Medicines Agency on 4 August 2023 an application for a paediatric investigation plan for the above mentioned medicinal product and a deferral under Article 20 of said Regulation.

The procedure started on 11 September 2023.

Opinion

1. The Paediatric Committee, having assessed the proposed paediatric investigation plan in accordance with Article 17 of Regulation (EC) No 1901/2006 as amended, recommends as set out in the appended summary report:

   - to agree the paediatric investigation plan in accordance with Article 17(1) of said Regulation;
   - to grant a deferral in accordance with Article 21 of said Regulation.

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

2. The measures and timelines of the agreed 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 annexes 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 congenital athymia

2.1.1. Indication(s) targeted by the PIP

Immune reconstitution in paediatric patients with congenital athymia

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)

Living tissue equivalent

2.1.4. Measures

Area	Description
Quality-related studies	Not applicable
Non-clinical studies	Not applicable
Clinical studies	**Study 1 (668-2)** Open-label single dose trial to evaluate activity and safety of allogeneic cultured postnatal thymus-derived tissue in children from birth to less than 18 years of age with congenital athymia. **Study 2 (884)** Open-label single dose trial to evaluate activity, safety, and tolerability of allogeneic cultured postnatal thymus-derived tissue combined with immunosuppression in children from birth to less than 18 years of age with congenital athymia. **Study 3 (25966)** Open-label single dose trial to evaluate activity and safety of allogeneic cultured postnatal thymus-derived tissue combined with immunosuppression adjusted to the subject's immune status in children from birth to less than 18 years of age with congenital athymia. **Study 4 (RVT-802-4001)** Open-label observational trial to evaluate long-term activity and safety

	of allogeneic cultured postnatal thymus-derived tissue in children from birth to less than 18 years of age with congenital athymia.
Modelling and simulation studies	Not applicable
Other studies	Not applicable
Extrapolation plan	Not applicable

3. Follow-up, completion and deferral of PIP

Concerns on potential long term safety/efficacy issues in relation to paediatric use:	No
Date of completion of the paediatric investigation plan:	By January 2028
Deferral for one or more measures contained in the paediatric investigation plan:	Yes

Annex II

Information about the authorised medicinal product

Information provided by the applicant:

The product is not authorised anywhere in the European Community.