

EMA/152075/2021

European Medicines Agency decision P/0151/2021

of 16 April 2021

on the acceptance of a modification of an agreed paediatric investigation plan for gadopichlenol, (EMA-001949-PIP02-18-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/0145/2019 issued on 17 April 2019,

Having regard to the application submitted by Guerbet on 26 November 2020 under Article 22 of Regulation (EC) No 1901/2006 proposing changes to the agreed paediatric investigation plan with a deferral,

Having regard to the opinion of the Paediatric Committee of the European Medicines Agency, issued on 26 February 2021, 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 gadopiclesol solution for injection, 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 Guerbet, BP 57400, 95943 - Roissy CDG Cedex, France.

EMA/PDCO/654621/2020
Amsterdam, 26 February 2021

Opinion of the Paediatric Committee on the acceptance of a modification of an agreed Paediatric Investigation Plan EMA-001949-PIP02-18-M01

Scope of the application

Active substance(s):

Gadopiclenol

Condition(s):

Detection and visualisation of disorders or lesions with suspected abnormal vascularity in various body regions for diagnostic purposes

Pharmaceutical form(s):

Solution for injection

Route(s) of administration:

Intravenous use

Name/corporate name of the PIP applicant:

Guerbet

Basis for opinion

Pursuant to Article 22 of Regulation (EC) No 1901/2006 as amended, Guerbet submitted to the European Medicines Agency on 26 November 2020 an application for modification of the agreed paediatric investigation plan with a deferral as set out in the European Medicines Agency's decision P/0145/2019 issued on 17 April 2019.

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

The procedure started on 4 January 2021.

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 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:

Detection and visualisation of disorders or lesions with suspected abnormal vascularity in various body regions for diagnostic purposes

2.1.1. Indication(s) targeted by the PIP

Body MRI to detect and visualise lesions and abnormalities within the head and neck, thorax, abdomen, pelvis, and musculoskeletal system.

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)

Solution for injection

2.1.4. Measures

Area	Number of measures	Description
Quality-related studies	1	Study 1 Development of an additional volume for the solution for injection to avoid risk of overdose <i>(This study is the same as Study 1 in EMEA-001949-PIP01-16 and subsequent modifications thereof.)</i>
Non-clinical studies	2	Study 2 Dose-range finding toxicity study in rats to determine the toxicity of gadopichlenol in the neonatal and juvenile (pre-post weaning) rats in order to determine appropriate dose levels for the definitive juvenile rat study <i>(This study is the same as Study 2 in EMEA-001949-PIP01-16 and subsequent modifications thereof.)</i> Study 3 Definitive juvenile toxicity study to determine the toxicity of gadopichlenol in the neonatal and juvenile (pre-post weaning) rats <i>(This study is the same as Study 3 in EMEA-001949-PIP01-16 and subsequent modifications thereof.)</i>

Clinical studies	2	Study 4 (GDX-44-007) Non-comparative pharmacokinetics, efficacy and safety study in children from 2 to less than 18 years of age presenting central nervous system (CNS) lesions (intracranial, spine and associated tissues), who are scheduled to undergo routine contrast-enhanced MRI of CNS or body <i>(This study is the same as Study 4 in EMEA-001949-PIP01-16 and subsequent modifications thereof.)</i> Study 5 Non-comparative pharmacokinetic, efficacy and safety study in children from birth to less than 2 years of age scheduled to undergo routine contrast-enhanced MRI of CNS or body <i>(This study is the same as Study 5 in EMEA-001949-PIP01-16 and subsequent modifications thereof.)</i>
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:	No
Date of completion of the paediatric investigation plan:	By March 2024
Deferral for one or more measures contained in the paediatric investigation plan:	Yes