

EMA/8373/2022

European Medicines Agency decision P/0023/2022

of 31 January 2022

on the agreement of a paediatric investigation plan and on the granting of a deferral and on the granting of a waiver for aficamten (EMEA-002958-PIP01-21) 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 Union procedures for the authorisation and supervision of medicinal products for human use and establishing a European Medicines Agency²,

Having regard to the application submitted by Cytokinetics, Inc. on 18 January 2021 under Article 16(1) of Regulation (EC) No 1901/2006 also requesting a deferral under Article 20 of said regulation and a waiver under Article 13 of said Regulation,

Having regard to the opinion of the Paediatric Committee of the European Medicines Agency, issued on 17 December 2021, in accordance with Article 17 of Regulation (EC) No 1901/2006 and Article 21 of said Regulation and Article 13 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 and on the granting of a waiver.
- (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.
- (4) It is therefore appropriate to adopt a decision granting a waiver.

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 aficamten, tablet, oral 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 aficamten, tablet, oral 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

A waiver for aficamten, tablet, oral 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 4

This decision is addressed to Cytokinetics Inc., 280 East Grand Avenue, 94080 - South San Francisco, United States.

EMA/PDCO/524958/2021
Amsterdam, 17 December 2021

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

EMA-002958-PIP01-21

Scope of the application

Active substance(s):

Aficamten

Condition(s):

Treatment of hypertrophic cardiomyopathy

Pharmaceutical form(s):

Tablet

Route(s) of administration:

Oral use

Name/corporate name of the PIP applicant:

Cytokinetics, Inc.

Basis for opinion

Pursuant to Article 16(1) of Regulation (EC) No 1901/2006 as amended, Cytokinetics, Inc. submitted for agreement to the European Medicines Agency on 18 January 2021 an application for a paediatric investigation plan for the above mentioned medicinal product and a deferral under Article 20 of said Regulation and a waiver under Article 13 of said Regulation.

The procedure started on 23 February 2021.

Supplementary information was provided by the applicant on 13 September 2021. The applicant proposed modifications to the paediatric investigation plan and the waiver.

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;
- to grant a waiver for one or more subsets of the paediatric population in accordance with Article 13 of said Regulation and concluded in accordance with Article 11(1)(c) of said Regulation, on the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments for paediatric patients.

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

2. The measures and timelines of the agreed 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 hypertrophic cardiomyopathy

The waiver applies to:

- the paediatric population from birth to less than 6 years of age;
- tablet, oral use;
- on the grounds that the specific medicinal product does not represent a significant therapeutic benefit as clinical studies are not feasible.

2. Paediatric investigation plan

2.1. Condition:

Treatment of hypertrophic cardiomyopathy

2.1.1. Indication(s) targeted by the PIP

Treatment of symptomatic sarcomeric obstructive hypertrophic cardiomyopathy (oHCM)

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

From 6 years to less than 18 years of age

2.1.3. Pharmaceutical form(s)

Tablet, oral use

2.1.4. Measures

Area	Number of measures	Description
Quality-related studies	1	Study 1 (PED-FORMDEV-1) Development of age-appropriate dosage strength(s)
Non-clinical studies	2	Study 2 (JUV-TOX-1) Dose range-finding juvenile toxicity study Study 3 (JUV-TOX-2) Definitive juvenile toxicity study

Clinical studies	1	Study 4 Randomised, double-blind, placebo-controlled, 2-arm, parallel group, 2-period study to evaluate the safety, pharmacokinetics (PK) and pharmacodynamics (PD) of aficamten compared to placebo (Period 1) followed by long-term, open-label assessment of safety and PD (Period 2) in children from 6 years to less than 18 years of age with symptomatic sarcomeric obstructive hypertrophic cardiomyopathy (oHCM).
Extrapolation, modelling and simulation studies	2	Study 5 Modelling and simulation study to support dose finding of the product in children from 6 years to less than 18 years of age with oHCM, based on data from adults and adolescents. Study 6 Modelling and simulation study to adjust the dose of the product in children from 6 years to less than 18 years of age with oHCM, on incorporation of PK/PD data from Study 4.
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 April 2028
Deferral for one or more measures contained in the paediatric investigation plan:	Yes