



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

EMA/528359/2023

European Medicines Agency decision P/0505/2023

of 29 December 2023

on the agreement of a paediatric investigation plan and on the granting of a deferral and on the granting of a waiver for broadly neutralizing anti-HIV human monoclonal antibody (VH3810109) (EMA-003392-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 726/2004¹,

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 Viiv Healthcare UK Limited on 20 January 2023 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 10 November 2023, 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 broadly neutralizing anti-HIV human monoclonal antibody (VH3810109), solution for injection/infusion, subcutaneous use, intravenous 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 broadly neutralizing anti-HIV human monoclonal antibody (VH3810109), solution for injection/infusion, subcutaneous use, intravenous 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 broadly neutralizing anti-HIV human monoclonal antibody (VH3810109), solution for injection/infusion, subcutaneous use, intravenous 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 ViiV Healthcare UK Limited, 980 Great West Road, TW8 9GS – Brentford, United Kingdom.



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

EMA/PDCO/363365/2023 Corr.¹
Amsterdam, 10 November 2023

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

EMA-003392-PIP01-23

Scope of the application

Active substance(s):

Broadly neutralizing anti-HIV human monoclonal antibody (VH3810109)

Invented name and authorisation status:

See Annex II

Condition(s):

Treatment of human immunodeficiency virus (HIV-1) infection

Pharmaceutical form(s):

Solution for injection/infusion

Route(s) of administration:

Subcutaneous use

Intravenous use

Name/corporate name of the PIP applicant:

ViiV Healthcare UK Limited

Basis for opinion

Pursuant to Article 16(1) of Regulation (EC) No 1901/2006 as amended, ViiV Healthcare UK Limited submitted for agreement to the European Medicines Agency on 20 January 2023 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 27 February 2023.

¹ 11 December 2023.



Supplementary information was provided by the applicant on 4 August 2023. The applicant proposed modifications to the paediatric investigation plan.

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 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 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 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

1.1. Condition:

Treatment of human immunodeficiency virus (HIV-1) infection

The waiver applies to:

- the paediatric population from birth to less than 2 years of age;
- solution for injection/infusion; subcutaneous use, intravenous use;
- on the grounds that the specific medicinal product does not represent a significant therapeutic benefit over existing treatments.

2. Paediatric investigation plan

2.1. Condition:

Treatment of human immunodeficiency virus (HIV-1) infection.

2.1.1. Indication(s) targeted by the PIP

Treatment of HIV-1-infected children and adolescents from 2 years to less than 18 years of age to replace the current antiretroviral (ARV) regimen in those who are virologically suppressed (HIV-1 RNA <50 copies/mL) on a stable ARV regimen, given with cabotegravir (CAB) (as a separate injection given at the same visit).

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

From 2 to less than 18 years of age.

2.1.3. Pharmaceutical form(s)

Solution for injection/infusion.

2.1.4. Measures

Area	Description
Quality-related studies	Not applicable
Non-clinical studies	Not applicable
Clinical studies	Study 1 Open-label, multiple-cohort study to evaluate VH3810109, administered IV or SC, in combination with cabotegravir (CAB) (as a separate injection given at the same visit) in HIV-1-infected (HIV-1 RNA <50 copies/mL) paediatric patients from 12 years to less than 18 years of age (and adults) on an existing suppressive antiretroviral therapy (ART). Study 2

	Open-label, single arm study to evaluate pharmacokinetics and safety of VH3810109 in HIV-1-infected (HIV-1 RNA <50 copies/mL) in combination with cabotegravir (CAB) (as a separate injection given at the same visit) in paediatric patients from 2 years to less than 12 years of age on an existing suppressive antiretroviral therapy (ART).
Modelling and simulation studies	Study 3 Population PK model to identify doses providing exposures of VH3810109 in paediatric patients from 2 years to less than 18 years of age that match the safe and efficacious exposures that have been confirmed to be safe and efficacious in the adult programme and to support extrapolation of efficacy.
Other studies	Not applicable.
Extrapolation plan	Study 3 is part of an extrapolation of efficacy plan agreed by PDCO.

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 December 2030
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.