

EMA/591816/2019

European Medicines Agency decision

P/0384/2019

of 4 December 2019

on the acceptance of a modification of an agreed paediatric investigation plan for ruxolitinib (phosphate) (Jakavi), (EMEA-000901-PIP04-17-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/0061/2018 issued on 16 March 2018,

Having regard to the application submitted by Novartis Europharm Limited on 8 July 2019 under Article 22 of Regulation (EC) No 1901/2006 proposing changes to the agreed paediatric investigation plan with a deferral and a waiver,

Having regard to the opinion of the Paediatric Committee of the European Medicines Agency, issued on 18 October 2019, 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 ruxolitinib (phosphate) (Jakavi), tablet, age-appropriate oral dosage form, oral use, gastric 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 Novartis Europharm Limited, Vista Building, Elm Park, Merrion Road, 4 Dublin, Ireland.



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

EMA/PDCO/410737/2019
Amsterdam, 18 October 2019

Opinion of the Paediatric Committee on the acceptance of a modification of an agreed Paediatric Investigation Plan EMA-000901-PIP04-17-M01

Scope of the application

Active substance(s):

Ruxolitinib (phosphate)

Invented name:

Jakavi

Condition(s):

Treatment of chronic Graft versus Host Disease

Authorised indication(s):

See Annex II

Pharmaceutical form(s):

Tablet

Age-appropriate oral dosage form

Route(s) of administration:

Oral use

Gastric use

Name/corporate name of the PIP applicant:

Novartis Europharm Limited

Information about the authorised medicinal product:

See Annex II



Basis for opinion

Pursuant to Article 22 of Regulation (EC) No 1901/2006 as amended, Novartis Europharm Limited submitted to the European Medicines Agency on 8 July 2019 an application for modification of the agreed paediatric investigation plan with a deferral and a waiver as set out in the European Medicines Agency's decision P/0061/2018 issued on 16 March 2018.

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

The procedure started on 20 August 2019.

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 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 chronic Graft versus Host Disease

The waiver applies to:

- newborn infants (from birth to less than 28 days);
- tablet, age-appropriate oral dosage form, oral use, gastric use;
- on the grounds that the disease or condition for which the specific medicinal product is intended does not occur in the specified paediatric subset(s).

2. Paediatric investigation plan

2.1. Condition:

Treatment of chronic Graft versus Host Disease

2.1.1. Indication(s) targeted by the PIP

Treatment of chronic Graft versus Host Disease (cGVHD) after allogeneic haematopoietic stem cell transplantation (HSCT).

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

From 28 days to less than 18 years of age

2.1.3. Pharmaceutical form(s)

Tablet

Age-appropriate oral dosage form

2.1.4. Measures

Area	Number of measures	Description
Quality-related studies	1	Study 1 Development of an age-appropriate oral dosage form <i>This study is the same as study 1 of the ruxolitinib PIP EMEA-000901-PIP03-16 and subsequent modifications thereof</i>

Non-clinical studies	1	Study 2 Definitive juvenile toxicity study in rats <i>This study is the same as study 2 of the ruxolitinib PIP EMEA-000901-PIP03-16 and subsequent modifications thereof</i>
Clinical studies	2	Study 3 Open-label, randomised, active-controlled trial to evaluate pharmacokinetics, safety and efficacy of ruxolitinib compared to best available therapy (BAT) in adults and adolescents from 12 to less than 18 years of age with corticosteroid-refractory (SR) cGvHD following allogeneic HSCT (INC424 D2301) Study 4 Open-label uncontrolled trial to evaluate pharmacokinetics, safety and activity of ruxolitinib in children from 28 days to less than 18 years of age with moderate or severe treatment-naïve or SR-cGvHD following allogeneic HSCT (INC424G12201)
Extrapolation, modelling and simulation studies	3	Study 5 Population PK (PopPK) modelling and simulation study to support the use of ruxolitinib for the treatment of cGvHD in children from 28 days to less than 18 years of age with treatment-naïve or SR-cGvHD following allogeneic HSCT Study 6 Physiologically-based PK (PBPK) modelling and simulation study to support the use of ruxolitinib for the treatment of cGvHD in children from 28 days to less than 18 years of age with treatment-naïve or SR-cGvHD following allogeneic HSCT Study 7: Extrapolation study to support the use of ruxolitinib for the treatment of cGvHD in children from 28 days to less than 18 years of age with treatment-naïve or SR-cGvHD following allogeneic HSCT
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:	Yes
Date of completion of the paediatric investigation plan:	By August 2026
Deferral for one or more measures contained in the paediatric investigation plan:	Yes

Annex II

Information about the authorised medicinal product

Condition(s) and authorised indication(s):

1. Treatment of myelofibrosis

Authorised indication(s):

Jakavi is indicated for the treatment of disease-related splenomegaly or symptoms in adult patients with primary myelofibrosis (also known as chronic idiopathic myelofibrosis), post polycythaemia vera myelofibrosis or post essential thrombocythaemia myelofibrosis.

2. Treatment of polycythaemia vera

Authorised indication(s):

Jakavi is indicated for the treatment of adult patients with polycythaemia vera who are resistant to or intolerant of hydroxyurea.

Authorised pharmaceutical form(s):

Tablet

Authorised route(s) of administration:

Oral use