



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

EMA/417834/2011

European Medicines Agency decision P/161/2011

of 4 July 2011

on the acceptance of a modification of an agreed paediatric investigation plan for ciclosporin (EMA-000575-PIP01-09-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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on the acceptance of a modification of an agreed paediatric investigation plan for ciclosporin (EMA-000575-PIP01-09-M01) in accordance with Regulation (EC) No 1901/2006 of the European Parliament and of the Council

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/49/2010 issued on 6 April 2010,

Having regard to the application submitted by Novagali Pharma S.A. on 7 February 2011 under Article 22 of Regulation (EC) No 1901/2006 proposing changes to the agreed paediatric investigation plan with a waiver,

Having regard to the opinion of the Paediatric Committee of the European Medicines Agency, issued on 20 May 2011, 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.
- (2) It is therefore appropriate to adopt a decision on the acceptance of changes to the agreed paediatric investigation plan.

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 ciclosporin, eye drops, emulsion, ocular use 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 Novagali Pharma S.A., 1 rue Pierre Fontaine - Bâtiment Genavir IV, 91058 Evry cedex, France.

Done at London, 4 July 2011

For the European Medicines Agency
Andreas Pott
Acting Executive Director
(Signature on file)



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

EMA/PDCO/193860/2011

Opinion of the Paediatric Committee on the acceptance of a modification of an agreed Paediatric Investigation Plan EMA-000575-PIP01-09-M01

Scope of the application

Active substance(s):

Ciclosporin

Condition(s):

Keratoconjunctivitis Sicca

Vernal Keratoconjunctivitis

Pharmaceutical form(s):

Eye drops, emulsion

Route(s) of administration:

Ocular use

Name/corporate name of the PIP applicant:

Novagali Pharma S.A.

Basis for opinion

Pursuant to Article 22 of Regulation (EC) No 1901/2006 as amended, Novagali Pharma S.A. submitted to the European Medicines Agency on 7 February 2011 an application for modification of the agreed paediatric investigation plan with a waiver as set out in the European Medicines Agency's decision P/49/2010 issued on 6 April 2010.

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

The procedure started on 23 February 2011.

A meeting with the Paediatric Committee took place on 19 May 2011.



Scope of the modification

Some measures 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 in the scope set out in the Annex I of this opinion.

The Icelandic and the Norwegian Paediatric Committee members agree 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 annex(es) and appendix.

London, 20 May 2011

On behalf of the Paediatric Committee
Dr Daniel Brasseur, Chairman
(Signature on file)

Annex I

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

1. Condition(s)

Keratoconjunctivitis Sicca

Vernal Keratoconjunctivitis

2. Waiver

2.1. Condition

Keratoconjunctivitis Sicca

The waiver applies to:

- All subsets of the paediatric population from birth to less than 18 years of age;
- for eye drops, emulsion for ocular use;
- on the grounds that the specific medicinal product does not represent a significant therapeutic benefit as clinical studies(s) are not feasible.

2.2. Condition

Vernal Keratoconjunctivitis

The waiver applies to:

- Children from birth to less than 4 years of age;
- for eye drops, emulsion for ocular 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).

3. Paediatric Investigation Plan

3.1. Condition to be investigated

Vernal Keratoconjunctivitis

3.1.1. Indication targeted by the PIP

Treatment of active severe vernal keratoconjunctivitis with severe keratitis.

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

From 4 to less than 18 years of age.

3.1.3. Pharmaceutical form(s)

Eye drops, emulsion

3.1.4. Studies

Area	Number of studies	Description
Quality	0	Not applicable.
Non-clinical	0	Not applicable.
Clinical	1	Double masked randomized multicenter dose regimen placebo controlled study to assess the efficacy and safety of ciclosporin (NOVA22007) in paediatric patients from 4 to less than 18 years of age with active severe vernal keratoconjunctivitis with severe keratitis

4. Follow-up, completion and deferral of PIP

Measures to address long term follow-up of potential safety issues or efficacy in relation to paediatric use:	Yes
Date of completion of the paediatric investigation plan:	By December 2013
Deferral for one or more studies contained in the paediatric investigation plan:	No