

To:

Head of Paediatric Medicines
European Medicines Agency

Notification of discontinuation of a paediatric development which is covered by an agreed PIP Decision

Actives substances(s): Molidustat sodium (BAY 85-3934)

Invented name: Not available at present

Latest Decision number(s): 1) P/0273/2014

Corresponding PIP number(s): 1) EMEA-001546-PIP01-13

Date of initial marketing authorisation granted:

Date of authorisation of new indication, pharmaceutical form or route of administration:

Please note that development of the medicinal product above in the following
condition(s)/indication(s):

Anemia in chronic disorders/Treatment of symptomatic anaemia associated with chronic kidney disease

☒ has been discontinued

☐ has been suspended/put on long-term hold (with possible re-start at a later time)

for the following reason(s): (tick all that apply)

☐ (possible) lack of efficacy in adults

☐ (possible) lack of efficacy in children

☐ (possible) unsatisfactory safety profile in adults

☐ (possible) unsatisfactory safety profile in children

☐ commercial reasons (please specify:)

☐ manufacturing / quality problems

☐ other regulatory action (please specify:) (e.g. suspension, revocation of M.A.)

☒ other reason (please specify: discontinuation of development of molidustat outside of Japan)

Please add a brief description (max 2000 characters) of the reason(s) for the discontinuation / suspension:

Molidustat has been in development for patients with anemia associated with chronic kidney disease and/or end-stage renal disease. Global phase 2 studies have been completed. No subjects are currently being treated. In none of the studies, any unexpected safety findings were observed. Based

on business reasons, the Sponsor has subsequently decided to develop this compound for the Japanese market only, based on a local phase 3 program. The local phase 3 studies are currently going in Japan. There are no clinical trials planned for Europe, nor any other region or country outside Japan.

Please note that if the PIP has been submitted as part of a marketing authorisation application in order to comply with the requirements of Article 7 of the Paediatric Regulation (as a condition of the validation of the respective application) and a marketing authorisation was granted based on this application, then there is a legal obligation to complete that PIP. The same applies if there has been a successful post-authorisation application, where the PIP was included in order to comply with the requirements of Article 8 of the Paediatric Regulation.

Please confirm if any of the above applies to the PIP in question:

Yes ☐ No ☒

If yes, it means that based on the Marketing Authorisation obtained at the end of that initial procedure or the successful post-authorisation application, as applicable, you are obliged to complete that PIP. That obligation cannot be cancelled by a unilateral decision, including by withdrawing the MA. Such PIP must be completed, unless it is modified in agreement with the PDCO by removing all outstanding PIP measures or granting a full product-specific waiver instead (upon relevant circumstances in accordance with the Paediatric Regulation). Non-completion of a binding PIP establishes noncompliance with the requirements of the Paediatric Regulation, which the European Medicines Agency has an obligation to report to the European Commission.

Name and signature of the PIP contact point: Signature on file

Date: 7 August 2019

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