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

Actives substances(s): Cotadutide

Latest Decision number(s): 1) P/0506/2021

Corresponding PIP number(s): 1) EMEA-002287-PIP01-17-M03

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:

Yes ☐ No [X]

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.

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

Treatment of type II diabetes mellitus

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: )

[X] other reason (please specify: strategic decision to stop development)

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

After careful consideration, AstraZeneca has decided to discontinue the development of cotadutide, a daily injectable GLP-1/glucagon co-agonist, to focus efforts on other assets in the pipeline. The decision to stop the development of cotadutide is exclusively a strategic decision. No new or unanticipated safety or tolerability concerns have been identified with this investigational product. The plans for a Marketing Authorisation Application for the treatment of adult patients, as well as the paediatric development of cotadutide has been discontinued.


| | |
|---|---|
| Cntact for inquiries from interested parties: | AstraZeneca AB |
| Telephone: | +46 8553 27591 |
| Email: | paediatrics@astrazeneca.com |