

To:

Head of Paediatric Medicines  
European Medicines Agency

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

Actives substance(s): AZD8233 sodium, PCSK9-targeted, N-acetylgalactosamine (GalNAc)conjugated antisense oligonucleotide (ASO)

Invented name: Not available at present

Latest Decision number(s): 1) P/0465/2021

Corresponding PIP number(s): 1) EMEA-002962-PIP01-21

Date of initial marketing authorisation granted: N/A

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

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

treatment of mixed dyslipidaemia; treatment of elevated cholesterol / treatment of heterozygous and of homozygous familial hypercholesterolemia

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: not sufficiently differentiated to be competitive)

manufacturing / quality problems

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

other reason (please specify: )

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

For commercial reasons and following the Phase 2b clinical study (SOLANO, D7990C0004) results, AstraZeneca has decided not to advance AZD8233 into Phase 3 development for hypercholesterolemia. AZD8233 was generally safe and well tolerated.

The agreed PIP included a deferral for the initiation of the paediatric study in relation to the planned initial MAA for the reduction of low-density lipoprotein cholesterol (LDL-C) in adults. Since it has been decided that AstraZeneca will not further develop AZD8233 in adults, AstraZeneca would like to withdraw the agreed PIP.

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: 25 Oct 2022

Contact for inquiries from interested parties: AstraZeneca

Telephone: +46 8 553 244 00

Email: paediatrics@astrazeneca.com