

London, 21 February 2018
EMA/CMDh/92835/2018
Co-ordination group for Human Use
EMA/H/N/PSR/J/0008

Position of the Co-ordination Group for Mutual Recognition and Decentralised Procedures for human use on non-interventional imposed Post-Authorisation Safety Study (PASS) final report for

Medicinal product(s)

Invented name(s): See PRAC assessment report for non-interventional imposed PASS final report appendix
Active substance(s): Thiocolchicoside

Basis for Position

Pursuant to Article 107p of Directive 2001/83/EC, the Marketing Authorisation Holder(s) submitted to the European Medicines Agency the final study report for non-interventional imposed post-authorisation safety study (PASS) for the nationally authorised medicinal product(s) mentioned above containing Thiocolchicoside (for details see PRAC assessment report appendix).

The evaluation procedure started on 29 August 2017.

The steps taken for the assessment of the non-interventional imposed PASS final report are detailed in the appended Pharmacovigilance Risk Assessment Committee (PRAC) assessment report.

The recommendation was adopted by the PRAC on 08 February 2018.

Position

1. The CMDh, having considered in accordance with Article 107q(2) of Directive 2001/83/EC the results of the study on the basis of the PRAC recommendation and the PRAC assessment report as appended, reaches its position by consensus on the variation to the terms of the Marketing Authorisation(s) for the medicinal products mentioned above concerning the following change(s):

The following changes to the conditions of the marketing authorisation(s) of medicinal products containing the active substance thiocolchicoside concerned by the PASS final report are recommended:

The MAH(s) should provide within the risk management plan submission, a protocol for results of the drug utilisation study to characterise prescribing practices for the medicinal products during typical clinical use in representative groups of prescribers and to	November 2017 2019
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assess main reasons for prescription. Final study report by:	
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The Icelandic and the Norwegian CMDh member(s) agree(s) with the above-mentioned position of the CMDh.

2. The scientific conclusions are set out in Annex I.
3. The changes to the conditions to the marketing authorisation(s) of the medicinal products mentioned above are set out in Annex II.
4. The timetable for the implementation of this CMDh position is set out in Annex III.

To the extent that other medicinal products containing thiocolchicoside not listed in the PRAC assessment report are currently authorised in the EU, or are subject to future authorisation procedures by the Member States, the CMDh recommends that the concerned Member States and Marketing Authorisation Holders take due consideration of this CMDh position.

This position is forwarded, to Member States, to Iceland and Norway, to the MAHs together with its annexes and appendix(ces).

