FOI 23/140

10th March 2023

Dear

Thank you for your request under the Freedom of Information Act (FOIA).

Duodopa 20mg/ml + 5mg/ml, intestinal gel (PL 41042/0001) was authorised by a CoA on 21/12/2012. The original MA (PL 20841/0001) was authorised through an incoming MRP with Sweden as RMS (SE/H/0415/001).

There is no PAR available through the HoA MRI Product Index:
https://mri.cts-mrp.eu/portal/details?productnumber=SE/H/0415/001

However, as the product was authorised via a procedure with another EU member state as the RMS, we cannot provide any further information under S27 of the FOIA (International Relations). We advise you request this information directly from the Swedish authorities.
https://www.hma.eu/human-medicines/national-contacts.html?showctr=26&cHash=8292a649851a8f837192db30732979a4

If you disagree with how we have interpreted the Freedom of Information Act 2000 with regards to your request, you can ask for the decision to be reviewed. The review will be carried out by a senior member of the Agency who was not involved with the original decision.

If you have a query about the information provided, please reply to this email.

Please remember to quote the reference number above in any future communications.

If you were to remain dissatisfied with the outcome of the internal review, you would have the right to apply directly to the Information Commissioner for a decision. Please bear in mind that the Information Commissioner will not normally review our handling of your request unless you have first contacted us to conduct an internal review. The Information Commissioner can be contacted at:

Information Commissioner's Office
Wycliffe House
Water Lane
Wilmslow
Cheshire
SK9 5AF

Yours sincerely

MHRA Customer Experience Centre
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