Re: FOI 24/265 - Missing PAR Prednisolone 5mg Tablets

FOILicensing <FOILicensing@mhra.gov.uk>
Wed 27/03/2024 19:03

[redacted]

Dear [redacted]

Regarding your request for the Public Assessment Report (PAR) for Prednisolone 5mg Tablets (PL 00142/0842), this marketing authorisation was granted by a Change of Authorisation Holder (CoA) on 02 June 2016. The original marketing authorisation for this product (PL 24668/0297) was granted by a decentralised procedure with the UK as Reference Member State (RMS).

A link to this PAR is provided below:
https://mhraproducts4853.blob.core.windows.net/docs/16dbd2f4f0518966d13f9208006fe831f7a83eac

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

If you are dissatisfied with the handling of your request, you have the right to ask for an internal review. Internal review requests should be submitted within two months of the date you receive this response and addressed to: info@mhra.gov.uk

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,

FOI Team, MHRA

From: [redacted]
Sent: Thursday, March 14, 2024 1:28 PM
To: MHRA Customer Services <MHRACustomerServices@mhra.gov.uk>
Subject: Missing PAR

Good Afternoon,

I can find the SPC and PIL for the following on your website but not the Public Assessment Report. Is the PAR available for this product?

Prednisolone 5mg Tablets
Accord-UK Ltd
PL00142/0842

Kind regards,

[redacted]