10 South Colonnade
Canary Wharf
London
E14 4PU
United Kingdom
gov.uk/mhra



2 May 2024

MHRA reference: FOI2024/00053

Dear ,

Thank you for your information request, which we received on 22 April. You asked for:

I would like to make an FOI request regarding the annual quantity of Androfeme (1% testosterone cream) being imported into the UK from Australia under the "specials" import licence, for the last 5 years.

We have dealt with your request under the Freedom of Information Act 2000 (FOIA).

We confirm that we hold the information you have asked for, and we are disclosing this information in full.

The MHRA is notified of the intent of a UK licensed importer to import unlicensed medicines. The MHRA is not informed of the actual physical import taking place. Importers may chose to import part of what they have notified to the MHRA or they may not import any packs at all.

The results below have been compiled manually using the search terms Androfeme and Australia as country of import.

Year of notification	Total number of packs notified
2019	9,900
2020	12,126
2021	40,200
2022	83,099
2023	42,525

Please note that these requests can be logged as a direct query to <a href="mailto:lmports@mhra.gov.uk">lmports@mhra.gov.uk</a> instead of FOIs.

We hope this information is useful for you.

This concludes our response to your request.

If you have a query about this response, please contact us at

Please remember to quote the reference number at the top of this letter in any future communications. Details of your appeal rights are below.

Yours sincerely,

Customer Experience

Medicines and Healthcare products Regulatory Agency

## **Appeal rights**

If you are dissatisfied with the handling of your request, you can ask us to conduct an internal review. Internal review requests should be submitted within two months of the date you receive this response and addressed to: <a href="mailto:foi.request@mhra.gov.uk">foi.request@mhra.gov.uk</a>

If you remain dissatisfied with the outcome of the internal review, you 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 a request unless the requester has first asked us to conduct an internal review.

The Information Commissioner can be contacted through their online webform at: <a href="https://ico.org.uk/make-a-complaints/foi-and-eir-complaints/foi-and-eir-complaints/">https://ico.org.uk/make-a-complaints/foi-and-eir-complaints/</a>

Or in writing to: Information Commissioner's Office, Wycliffe House, Water Lane, Wilmslow, Cheshire, SK9 5AF

## Re-use of our information

The MHRA information supplied in response to your request is subject to Crown copyright. Information created by the MHRA which is disclosed under the Freedom of Information Act is made available for re-use under the Open Government Licence (OGL) v3.0, except where this is otherwise stated. There are some restrictions on re-use under the OGL and these can be viewed here:



## Medicines & Healthcare products Regulatory Agency

https://www.nationalarchives.gov.uk/doc/open-government-licence/version/3/

If you re-use our information, you should include the following attribution, including a link to the OGL v3.0:

Medicines and Healthcare products Regulatory Agency, [name and date of publication], licensed under the <a href="Open Government Licence">Open Government Licence</a>.

For further information on the reproduction or re-use of MHRA information, please visit <a href="https://www.gov.uk/government/publications/reproduce-or-re-use-mhra-information/reproduce-or-re-use-mhra-information">https://www.gov.uk/government/publications/reproduce-or-re-use-mhra-information</a>.