



Medicines & Healthcare products
Regulatory Agency

10 South Colonnade
Canary Wharf
London
E14 4PU
United Kingdom
[gov.uk/mhra](https://www.gov.uk/mhra)

[REDACTED]

23 May 2024

MHRA reference **FOI2024/00079**

Dear [REDACTED],

Thank you for your information request, which we received on **24th April 2024**. You asked for:

Please see attached the electrodes used on ECG forwarded by the manager at Lonsdale Medical Centre. I would rather wait for your reply in regards to the ingredients or if you could send me a link to the exact brand with ingredient list on your website.

As for the symptoms I have had to the allergy they include Allergic Contact Dermatitis causing blister type rash, Facial Myclonus, Increased blood pressure, full body muscle spasms & sleep issues.

Hope this helps with your search for anyone else experiencing this type of allergic reaction to ECG.

We have dealt with your request under the Freedom of Information Act 2000 (FOIA). After a search of our paper and electronic records in relevant and appropriate locations we have not been able to locate the information to address your request.

Therefore, under Section 1(1) (a) of the FOIA we confirm that the information is not held and having exhausted all the usual avenues in our search for this information, we have concluded that it is no longer on our systems in a retrievable form.

This concludes our response to your request.

If you have a query about this response, please contact us at foi.request@mhra.gov.uk



Medicines & Healthcare products Regulatory Agency

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,

Safety and Surveillance Group

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: foi.request@mhra.gov.uk

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: <https://ico.org.uk/make-a-complaint/foi-and-eir-complaints/foi-and-eir-complaints/>

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 FOIA requests 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:

<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 [Open Government Licence](#).

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



Medicines & Healthcare products
Regulatory Agency