Medicines & Healthcare products Regulatory Agency

MHRA Central Freedom of Information Team
10 South Colonnade
Canary Wharf
London
E14 4PU

foi.request@mhra.gov.uk.

MHRA Website

[redacted]

Our Ref: **FOI2024/00638**

13 November 2024

Dear [redacted]

Thank you for your Freedom of Information (FoI) request received on 17 October. You wrote:

> *Please provide a copy of the MHRA GMDP Inspection Report arising from the 21 Feb 2022 inspection of the following site:*
>
> *PHARMARON MANUFACTURING SERVICES (UK) LTD, WINDMILL INDUSTRIAL ESTATE, SHOTTON LANE, CRAMLINGTON, NE23 3JL, UNITED KINGDOM*
>
> *The corresponding GMP Certificate is: UK API 22857 Insp GMP 22857/36790-0008<https://cms.mhra.gov.uk/mhra/gmp/uk-api-22857-insp-gmp-2285736790-0008>*
>
> *(The site was previously owned by Rosemont)*

**MHRA Response**

The Agency has completed its search for the information you have requested and we are able to confirm that we do hold the information you have requested.

Please note that some of the information within the inspection report cannot be disclosed and is being exempt from release for the reasons below.

We consider that the information is exempt under section 40 and section 43.

**Section 40:**
This information contains elements of personal data, the disclosure of which would be unfair in that it would breach the first principle of the Data Protection Act which says that information must be processed fairly and lawfully.

**Section 43:**
Release of all, or part of, the information would, or would be likely to, cause harm to the third party's commercial interests.

We have considered the balance of the public interest when applying this exemption.
The exemption is to safeguard the commercially sensitive information / industrial secrets of a third party / commercial enterprise (which can include a Government Department). This exemption is conditional on the public interest in releasing it not outweighing the

company's/commercial enterprise's right to confidentiality and the probable damage that the company/commercial enterprise could suffer as a result of the information being released. In this case we have not identified any issues which would benefit the public as a whole by being brought to their attention (examples of issues would be a major public health risk or a major procedural failure or irregularity).

This concludes our response to your request.

If you have any queries about this letter, please contact us quoting the reference number above.

Yours sincerely,

MHRA Central Freedom of Information Team
Medicines & Healthcare products Regulatory Agency

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**Your right to complain under the Freedom of Information Act**

If you are not happy with this response you may request an internal review by e-mailing foi.request@mhra.gov.uk or by writing to: MHRA Central Freedom of Information Team, 10 South, Colonnade, Canary Wharf, London, E14 4PU

Any request for an internal review must be received by us within 40 working days of the date of this letter. Please note we are not obliged to provide a review if it is requested after more than 40 working days.

If you are not content with the outcome of the internal review you may apply directly to the Information Commissioner's Office for a decision. Generally, the Commissioner cannot make a decision unless you have exhausted our own complaints procedure. The Information Commissioner can be contacted at: The Information Commissioner's Office, Wycliffe House, Water Lane, Wilmslow, Cheshire SK9 5AF.
Website: ICO FOI and EIR complaints or telephone 0303 123 1113.

**Re-use of our information**