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: **FOI2025/00037**

07 February 2025

Dear [redacted],

Thank you for your Freedom of Information (FoI) request received on 10 January. You wrote:

> ***Could you please share full latest WDA & GDP inspection report:***
>
> ***1- AHP Medical Supplies Limited - WDA(H) 19142***
> ***2- Primius Lab Limited - WDA(H) 40556***
> ***3- Simply Meds Pro Limited - WDA(H) 50648***
> ***4- VERTICAL PHARMA RESOURCES LIMITED, 41 CENTRAL AVENUE, WEST MOLESEY, KT8 2QZ, UNITED KINGDOM***
> ***5- CLINIGEN HEALTHCARE LIMITED, PITCAIRN HOUSE, CROWN SQUARE, FIRST AVENUE, BURTON-ON-TRENT, DE14 2WW, UNITED KINGDOM***
> ***6- CLINIGEN HEALTHCARE LIMITED, UNIT 3, CANADA ROAD, BYFLEET, WEST BYFLEET, KT14 7JL, UNITED KINGDOM***
> ***7- CLINIGEN HEALTHCARE LIMITED, IDIS HOUSE, CHURCHFIELD ROAD, WEYBRIDGE, KT13 8DB, UNITED KINGDOM***
> ***8- ALLIANCE HEALTHCARE (DISTRIBUTION) LIMITED, AVENUE 1, LETCHWORTH BUSINESS PARK, LETCHWORTH, SG6 2HB, UNITED KINGDOM***

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 majority of the information you have requested.

Wholesale Distribution Authorisations (WDAs) can be found by searching the MHRA-GMDP database at the following link:
https://cms.mhra.gov.uk/mhra/wda

The authorisations for AHP Medical Supplies and Simply Meds Pro Limited were revoked and so finalised GDP reports are not available for the latest inspections that took place at these sites. We have instead included the post-inspection letters that were sent out to the sites following the inspections. These letters describe the deficiencies that were identified.

The authorisation for Primius Labs Limited was also revoked in 2022. This was due to the apparent ceasing of operations at the site. The MHRA Inspection Action Group (IAG) proposed revocation and when no representations were received from the company the WDA(H), API and MIA were revoked. The most recent inspection took place in 2017 so we have provided the associated inspection report.

A list of revoked licences can be found at the link below:
Revoked manufacturing and wholesale distribution authorisations - GOV.UK

Please find the requested reports for the other inspections attached.

Please note that some of the information within the reports and post-inspection letters 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

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