



Medicines & Healthcare products
Regulatory Agency

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

foi.request@mhra.gov.uk.

[MHRA Website](#)

Our Ref: **FOI2026/00441**

21 May 2026

Dea [REDACTED]

Thank you for your Freedom of Information (Fol) request received on 22 April. You wrote:

Good day,

Based on the MHRA Inspections Listing we received, we kindly request that you please provide the inspection reports for the following:

DR REDDY'S LABORATORIES (UK) LIMITED, located at 410 CAMBRIDGE SCIENCE PARK, MILTON ROAD, CAMBRIDGE, CB4 0PE, UNITED KINGDOM, Inspection End Date: January 8, 2026, INSP GDP 8553/19889136-0008.

PHIL INTER PHARMA COMPANY LIMITED, located at NUMBER. 20, HUU NGHI BOULEVARD, VIETNAM-SINGAPORE INDUSTRIAL PARK (VSIP) BINH HOA WARD, HO CHI MINH CITY, CITY VN-590000, VIET NAM, Inspection End Date: November 21, 2025, INSP GMP 46387/14673770-0004.

AMANAH PHARMACEUTICALS LIMITED, located at UNIT 13, FIRST QUARTER, LONGMEAD BUSINESS PARK, BLENHEIM ROAD, EPSOM, KT19 9QN, UNITED KINGDOM, Inspection End Date: October 28, 2025, INSP GDP 61385/37432789-0001.

NORDIC PHARMA LIMITED, located at BUILDING 1410, ARLINGTON BUSINESS PARK, THEALE, READING, RG7 4SA, UNITED KINGDOM, Inspection End Date: October 7, 2025, INSP GDP 5827/34897770-0002.

RIA GENERICS LIMITED, located at 36 INGLEBY WAY, WALLINGTON, SM6 9LR, UNITED KINGDOM, Inspection End Date: September 24, 2025, INSP GMP/GDP 36282/8148588-0012.

TILLOMED LABORATORIES LIMITED, located at 220 BUTTERFIELD, GREAT MARLINGS, LUTON, LU2 8DL, UNITED KINGDOM, Inspection End Date: September 10, 2025, INSP GMP/GDP 11311/16282866-0008.

NORBROOK LABORATORIES LIMITED, located at 105 ARMAGH ROAD, NEWRY, BT35 6PU, UNITED KINGDOM, Inspection End Date: July 24, 2025, INSP GLP 2000/12919-0023.

Kindly let us know if any questions arise from the processing of this request.

Thank you!

MHRA Response

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

Please find the five inspection reports that we hold attached.

An Inspection report is not held for the Dr Reddy's Laboratories inspection case as this was a 'Named Person' assessment.

An inspection report is not held for Amanah Pharmaceuticals Limited as the application was withdrawn by the site before a report was issued.

Please note that some of the information in the inspection reports we have provided cannot be disclosed and is being exempt from release for the reasons below.

We consider that the information is exempt under Section 40(2) and Section 43(2) of the FoI Act.

Section 40(2)

We are unable to provide you with some of the information requested as it constitutes personal data of someone other than yourself and as such, it is being withheld in accordance with section 40(2) of the Freedom of Information Act.

Section 40(2) exempts information in response to a request if it is personal data belonging to an individual other than the requester and it satisfies one of the conditions listed in the legislation. In this case the condition contained in section 40(3A)(a) applies - that disclosure would breach one of the data protection principles, specifically that "Personal data shall be processed lawfully, fairly and in a transparent manner...".

We do not consider that disclosing this information is necessary or justified in order to satisfy your information request and the requirements of the FoI Act. In relation to this request, we consider that there is no strong legitimate interest that would override the prejudice to the rights and freedoms of the data subject.

Personal data are subject to UK General Data Protection Regulation (UK GDPR) and the Data Protection Act 2018.

MHRA maintains the following policy on disclosure of personal information:

"All personal information held in MHRA records is regarded as confidential. Information will not normally be disclosed to third parties without the consent of the person concerned. Information may normally be disclosed without consent to meet statutory requirements; to comply with a court order; to prevent duplication of payments from public funds; or where there is a compelling public interest in making the disclosure."

All disclosures made by MHRA, in order to be considered authorised must be able to demonstrate that at least one of the above criteria apply.

Section 43(2)

Release of all, or part of, the information 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).

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

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