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24th February 2026

[redacted]

Dear Sir/Madam

Freedom of Information Act (FOI) 2000 - Request for Information Reference: MS treatment

I am writing in response to your request for information under the FOI 2000.

I can confirm in accordance with Section 1 (1) of the Freedom of Information Act 2000 that we do hold some of the information you have requested. A response to each part of your request is provided below.

FOI Request / Question	Question Response	Is there an exemption?	Exemption	Exemption Details
1. How many patients who have a current diagnosis of Multiple Sclerosis received treatment in the last six months with any of the following treatment regimens (July 2025 – December 2025)? • Ocrevus (ocrelizumab) IV (300mg/30ml) • Ocrevus (ocrelizumab) Subcutaneous (920mg/23ml) • Kesimpta (ofatumumab) • Briumvi (ublituximab) • Mavenclad (cladribine) • Dimethyl Fumarate (Tecfidera + Dimethyl Fumarate generic) • Natalizumab (Tysabri and Tyruko) 1a. Of the patients identified in question one, how many patients received the same treatment in the previous six-month period? (i.e. How many of the same patients received treatment with each treatment regimen in January 2025 to June 2025 AND July 2025 – December 2025?) • Ocrevus (ocrelizumab) IV (300mg/30ml)	The Trust has not used any of these treatments within the time frames requested.			

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• Ocrevus (ocrelizumab) Subcutaneous (920mg/23ml) • Kesimpta (ofatumumab) • Briumvi (ublituximab) • Mavenclad (cladribine) • Dimethyl Fumarate (Tecfidera + Dimethyl Fumarate generic) • Natalizumab (Tysabri and Tyruko)				
2. How many patients have received a new diagnosis for Multiple Sclerosis in the latest six-month period (July 2025 – December 2025)? 2a. Of the patients identified in question two, how many of these patients have received any of the following treatment regimens as their first treatment following their Multiple Sclerosis diagnosis? • Ocrevus (ocrelizumab) IV (300mg/30ml) • Ocrevus (ocrelizumab) Subcutaneous (920mg/23ml) • Kesimpta (ofatumumab) • Briumvi (ublituximab) • Mavenclad (cladribine) • Dimethyl Fumarate (Tecfidera +	N/A			

Dimethyl Fumarate generic) • Natalizumab (Tysabri and Tyruko)				

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I trust this information answers your request. Should you have any further enquiries or queries about this response please do not hesitate to contact me. However, if you are unhappy with the way in which your request has been handled, you have the right to ask for an internal review. Internal review requests should be submitted within two months of the date of receipt of the response to your original letter and should be addressed to: Sally Brook Shanahan, Director of Corporate Affairs, King's Mill Hospital, Mansfield Road, Sutton in Ashfield, Nottinghamshire, NG17 4JL or email sally.brookshanahan@nhs.net.

If you are dissatisfied with the outcome of the internal review, you can apply to the Information Commissioner's Office, who will consider whether we have complied with our obligations under the Act and can require us to remedy any problems. Generally, the Information Commissioner's Office cannot decide unless you have exhausted the internal review procedure. You can find out more about how to do this, and about the Act in general, on the Information Commissioner's Office website at: https://ico.org.uk/your-data-matters/official-information/.

Complaints to the Information Commissioner's Office should be sent to FOI/EIR Complaints Resolution, Information Commissioner's Office, Wycliffe House, Water Lane, Wilmslow, Cheshire, SK9 5AF. Telephone 0303 1231113, email casework@ico.org.uk.

If you would like this letter or information in an alternative format, for example large print or easy read, or if you need help with communicating with us, for example because you use British Sign Language, please let us know. You can call us on 01623 672232 or email sfh-tr.foi.requests@nhs.net.

Yours faithfully

Information Governance Team

All information we have provided is subject to the provisions of the Re-use of Public Sector Information Regulations 2015. Accordingly, if the information has been made available for re-use under the Open Government Licence (OGL) a request to re-use is not required, but the licence conditions must be met. You must not re-use any previously unreleased information without having the consent from Sherwood Forest Hospitals NHS Foundation Trust. Should you wish to re-use previously unreleased information then you must make your request in writing. All requests for re-use will be responded to within 20 working days of receipt.