FOI 23/790

Dear

Many thanks for your information request dated 18 October 2023, where you asked the details of *which version of the Pfizer Covid-19 vaccine ("Comirnaty Original/Omicron BA.1, tozinameran/riltozinameran", "Comirnaty Original/Omicron BA.4-5, tozinameran/famtozinameran", or "Comirnaty Omicron XBB.1.5, raxtozinameran") the vaccine batch number "GM8341" relates to as this information is not provided online.*

Our response:

We can confirm that Comirnaty batch GM8341 was the Original / Omicron BA.4/5 presentation.

We trust that you will find this information of use. However, if you disagree with how we have interpreted the Freedom of Information Act 2000 in answering your request, you have the right to ask for an internal review. Please reply to this email, within two months of this reply, specifying that you would like an Internal Review to be carried out.

Please remember to quote the reference number above in any future communications.

If you were to remain dissatisfied with the outcome of the internal review, you would 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 your request unless you have first contacted us to conduct an internal review.

The Information Commissioner can be contacted online via an electronic form: https://ico.org.uk/make-a-complaint/foi-and-eir-complaints/foi-and-eir-complaints/
Or
by writing to:
Information Commissioner's Office
Wycliffe House
Water Lane
Wilmslow
Cheshire
SK9 5AF

Yours sincerely,
HQA FOI Team

Medicines and Healthcare products Regulatory Agency
10 South Colonnade, Canary Wharf, London E14 4PU