

FOI 23/593

Dear

Thank you for your request for information of 11 August 2023, where you asked:

“Please can you provide access to all iterations/revisions of the “Reg 174 Information for UK Recipients” relating to the “COVID-19 Vaccine AstraZeneca solution for injection” with confirmation of the date of each revision.”

We have noted that this information was uploaded to the <https://www.gov.uk/government/publications/regulatory-approval-of-covid-19-vaccine-astrazeneca> website, and that the [Wayback machine](#) includes an extensive index / access to the requested documentation. Therefore, we consider that the information requested is reasonably accessible to you, therefore, it is necessary to apply the below exemption.

Section 21 – Information accessible by other means

The information you have requested is already in the public domain, and can be found [Wayback machine](#). The FOIA section 21 exemption is an absolute exemption and states that there is no right of access to information via FOI if it is reasonably available to the applicant by another route.

If you are dissatisfied with the handling of your request, you have the right to ask for an internal review. Internal review requests should be submitted within two months of the date you receive this response and addressed to: info@mhra.gov.uk
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 at:

Information Commissioner’s Office
Wycliffe House
Water Lane
Wilmslow
Cheshire
SK9 5AF

Yours sincerely

MHRA Customer Experience Centre

Communications and engagement team

Medicines and Healthcare products Regulatory Agency

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