



Medicines & Healthcare products
Regulatory Agency



MHRA
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11th July 2023

Dear [REDACTED]

FOI 23/164

Thank you for your email dated 22nd February 2023, where you requested information held by the MHRA relating to the CPRD study “Near real time vaccine safety monitoring for COVID-19 vaccines”. Apologies for the delay in responding to you on this request.

Specifically you asked, would you be able to provide all versions of the protocol for this study.

The protocol and reports from this study will be published in the future and thus provision of this information is exempt under Section 22 of the FOI Act. Section 22 of the Act allows public authorities to refuse requests where the authority intends to publish the information at a future date. This study will be published in a peer reviewed journal.

Yours sincerely,

FOI Team,
Safety and Surveillance Group

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Medicines & Healthcare products
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