

**FOI 23/069**

10<sup>th</sup> February 2023

Dear

Thank you for your request under the Freedom of Information Act (FOIA).

The main efficacy and safety results for the Phase I, II and III trials for all authorised vaccines have been submitted to MHRA, sufficient that these vaccines can be authorised for use in the patient populations stated in the Summary of Product Characteristics for each vaccine. Some studies are currently ongoing to follow-up vaccine recipients to collect additional safety data, in the same way that all clinical trials for new medicines follow up their study subjects after the main results of the study have been reported. Other studies that are currently in progress are either for cohorts/subpopulations of recipients or to investigate different regimens (such as giving different brands of vaccine for the first and second doses).

Information on ongoing clinical trials is publicly available in various registries, including [ISRCTN Registry](#) and [Home - ClinicalTrials.gov](#). Please use the advanced search functions to identify trials of interest based on product name, condition under investigation.

If you disagree with how we have interpreted the Freedom of Information Act 2000 with regards to your request, you can ask for the decision to be reviewed. The review will be carried out by a senior member of the Agency who was not involved with the original decision.

If you have a query about the information provided, please reply to this email.

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  
10 South Colonnade, Canary Wharf, London E14 4PU  
Telephone 020 3080 6000