

[REDACTED]

From: [REDACTED]
Sent: 31 March 2021 12:32
To: Pharmacovigilanceservice
Subject: RE: FOI 21/276 - Freedom of Information Act Request of adverse drug reaction (ADR) data regarding all Covid-19 vaccines in use in the UK which is not available on gov.uk website

Yep standard idap lines is fine- thanks

From: Pharmacovigilanceservice <Pharmacovigilanceservice@mhra.gov.uk>
Sent: 28 March 2021 14:58
To: [REDACTED]
Subject: FW: FOI 21/276 - Freedom of Information Act Request of adverse drug reaction (ADR) data regarding all Covid-19 vaccines in use in the UK which is not available on gov.uk website
Importance: High

Hi [REDACTED]

Just to confirm, all of this will be covered by iDAPs?

Thanks,
[REDACTED]

From: MHRA Customer Services <MHRACustomerServices@mhra.gov.uk>
Sent: 22 March 2021 15:51
To: Pharmacovigilanceservice <Pharmacovigilanceservice@mhra.gov.uk>
Cc: [REDACTED]
Subject: FOI 21/276 - Freedom of Information Act Request of adverse drug reaction (ADR) data regarding all Covid-19 vaccines in use in the UK which is not available on gov.uk website
Importance: High

Dear All,

This is a Freedom of Information request FOI 21/276.

The deadline for reply is 20/04/2021 but this is not a target. Please aim to send it out as soon as possible to avoid backlogs.

- Please review the case immediately and establish:
 - If it's not for you or
 - If it's sensitive and might need wider input on handling. If it does, please alert wider corporate leads and local management in advance and ensure that they have the opportunity to input before the reply is sent.
- **The [FOI Policy team](#) and [News Centre](#) need to see draft responses to sensitive cases with 5 days' notice – notice – please can you move these ones to the front of the queue.** This is to provide enough time to review and comment and arrange high level clearance, if needed. Sensitive cases are flagged in the twice-weekly reminder emails. The [Customer Services](#) team will also send these out. **Each division is responsible for ensuring a full and final draft is sent to customer service.**
- If the case concerns a wider ongoing topic, please ensure consistency with other communications (eg providing additional context in the reply).
- Click [here](#) for FOI guidance and [here](#) for reply templates.
- Divisional FOI leads, click [here](#) for the shared FOI case tracker.

- Finally, please copy [FOI Policy mailbox](#) and [Customer Services](#) when you send the final cleared reply.

The [FOI Policy team](#) are happy to help if you have any questions.

Many thanks

MHRA Customer Service Centre

Medicines and Healthcare products Regulatory Agency
10 South Colonnade, Canary Wharf, London E14 4PU
Telephone 020 3080 6000”

From: Marco Tullio Suadoni <marco.suadoni@pm.me>

Sent: 19 March 2021 10:46

To: MHRA Customer Services <MHRACustomerServices@mhra.gov.uk>

Subject: Freedom of Information Act Request of adverse drug reaction (ADR) data regarding all Covid-19 vaccines in use in the UK which is not available on gov.uk website

CHM

To whom it might concern:

Dear Sir or Madam,

Re: Freedom of Information Act Request of adverse drug reaction (ADR) data regarding all Covid-19 vaccines in use in the UK which is not available on gov.uk website

First of all, I am aware of the information available

here: <https://www.gov.uk/government/publications/coronavirus-covid-19-vaccine-adverse-reactions>

However, the information linked as above does no report all ADRs data, but only summary data.

I request in spreadsheet or database format, e.g., comma-separated-values (CSV) (not PDF format), the full body of all anonymised raw data with the level of details as close as possible to that one available for Interactive Drug Analysis Profile (iDAP) and related CSV files, for all Covid-19 vaccines currently in use in the UK. Especially to include for EACH event, but not limited to:

SEX
AGE
DATE
REPORTER
REPORT SUBMISSION
ROUTE OF ADMINISTRATION
SERIOUSNESS
SYSTEM ORGAN CLASS

Yours faithfully,

Marco Tullio Suadoni

Sent with [ProtonMail](#) Secure Email.