FOI 23-530

MHRA RESPONSE

Thank you for your FOI request dated 24th July.

> *I would like to request the following information: The number of reports made on the Yellow Card Scheme in the last 5 years, (if possible) relating to; (*Hormonal* long acting reversible contraceptives). Specifically, of side effects experienced. (not relating to pain on insertion or migration of devices as already published). If possible - can the side effects be stated, or categorised into groups. If possible - can the reports be separated into who reported them (eg professionals/ patients) i) Nexplanon ii) Hormonal intrauterine devices; Mirena, Jaydess, Kyleena, Levosert (please do not include the non-hormonal IUD) iii) Injectable methods; Depo-Provera, Sayana-Press*

Please note that the MHRA regularly publishes Yellow Card data on our website and this can be accessed using the link below:
https://yellowcard.mhra.gov.uk/idaps

You may look up the information by active substance and filter by different criteria using the interactive profiles. Should you require further assistance, please do not hesitate to contact us.
Kind regards,

FOI Team
Safety & Surveillance Group
Medicines and Healthcare Products Regulatory Agency
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