What is the MHRA?

The **Medicines and Healthcare products Regulatory Agency (MHRA)** is the government agency responsible for ensuring that medicines and medical devices work and are acceptably safe. When any possible problem is found, the MHRA takes prompt action to protect the public and reduce risk.

Medicines & Healthcare products Regulatory Agency

For more information about the MHRA:

Visit: **gov.uk/mhra**

Email: **info@mhra.gsi.gov.uk**

Tel: **020 3080 6000**

Reporting Adverse Drug Reactions to the Yellow Card Scheme

Healthcare professionals are asked to help improve medicines safety by reporting all suspected Adverse Drug Reactions (ADRs) to the Yellow Card Scheme that are:

- associated with **newer drugs and vaccines** - identified by the black triangle ▼ symbol: **mhra.gov.uk/blacktriangle**
- **serious, medically significant or result in harm** from established vaccines and medicines, including unlicensed medicines, herbal remedies, and medicines used off-label. Serious events are fatal, life-threatening, disabling or incapacitating, or result in or prolong hospitalisation.
- **If you are unsure, please report anyway.**

Help make medicines safer for everyone

mhra.gov.uk/yellowcard

MHRA

How do I report a suspected side effect?

There are three main ways to report to the Yellow Card Scheme

- the easiest way to report is **online** at **mhra.gov.uk/yellowcard**
- the free **Yellow Card mobile app** from iTunes Yellow Card or PlayStore Yellow Card for iOS or Android devices
- **complete a paper Yellow Card form** which you can post to FREEPOST YELLOW CARD

Yellow Cards can be found in the **BNF**, **MIMS**, **ABPI Compendium** or ordered by calling the **Yellow Card Information Service freephone on 0800 731 6789**

What happens to Yellow Cards?

The MHRA (see overleaf) collects Yellow Card reports from all healthcare professionals and patients. These reports are used to identify ADRs and other problems which might not have been known about before about the medicine.

If a new ADR is found, the MHRA will review the way that the medicine can be used, and the warnings that are given to prescribers and patients.

The information you provide will be kept safe, secure and confidential. No details that could identify you or your patient(s) will be passed on without your permission.

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