## Vol.6 Issue 5: November 2025



**MEDwatch** is the e-bulletin for all NHS Grampian Staff who are involved with patients and medicine management.

Its aim is to improve the safety of medicines by sharing learning, and encouraging adverse event reporting from all staff groups.



3rd - 9th November sees the annual #MedSafetyWeek campaign, during which the MHRA joins forces with other regulatory bodies around the world to promote reporting adverse drug reactions (ADRs).

This year's theme is "we can all help make medicines safer".

A side effect is an unexpected effect that happens in addition to the main intended effect of the medicine. An ADR is when the resulting effect causes harm.



Anyone; healthcare professionals, patients, caregivers and the public can report ADRs to the MHRA for any medicine (this includes blood products, vaccines and herbal/homeopathic medicines).

Report electronically using the <u>Yellow Card Scheme</u> and provide as much information as possible including the following if known:

- medical history
- any concomitant medication
- onset timing
- treatment dates.

If you're unsure whether someone has already submitted a report it's better to report than not, the MHRA are able to identify duplicate reports for the same incident.



Each time a report is submitted we all benefit; medicines become safer for patients to use and more information is available to healthcare professionals. For example:

- warnings may be added to the product information
- advice on how to take the medicine may be changed
- warnings may be given to groups of patients or healthcare professionals.

## **Turas Modules**

Six ADR e-learning modules created by Yellow Card Centre Scotland and NES are available on Turas:

- 1. Basic principles of ADR
- 2. Categorisation
- 3. Drug allergy classification
- 4. Diagnosis, interpretation and management
- 5. Avoiding adverse drug reactions
- 6. Pharmacovigilance

You can find them by searching 'adverse drug reactions' in Turas.

## **Contact**

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