# Batch Recall

## Warfant 3mg Tablets, PA0073/139/002

| Batch number | Expiry date |
|---|---|
| 25002 | 03/2030 |

17th of August 2026

Dear Pharmacist/Healthcare Professional,

We wish to advise you that batch no. 25002 of Warfant 3mg Tablets, PA 0073/139/002 is being recalled with immediate effect.

This recall is going to **patient level**. This action has been agreed with the Health Products Regulatory Authority (HPRA).

The impacted batch is being recalled due to a potential product mix up issue, whereby a report was received in which one sealed unit labelled as Warfant 3mg Tablets was found to contain Warfant 1 mg Tablets; all tablets within the container were the 1 mg presentation. One product mix-up report has been received to date, and it is considered that the rate of this defect is low within the batch.

**You are requested to perform the following actions**:

1. For stock within your pharmacy or facility, immediately identify and quarantine any units (including partial units) of Warfant 3mg Tablets from batch no. 25002. For hospital pharmacies, clinics, nursing homes and other such facilities, this includes units at wards, clinics and any other relevant locations within your facility.

2. For Warfant 3mg Tablets which you have dispensed since 02/02/2026:
    - Check your dispensing records to identify any patients to whom Warfant 3mg Tablets were dispensed since 02/02/2026.
    - Contact those patients or their carers via telephone to request them to check that Warfant 3mg Tablets within their possession are the correct strength. The correct Warfant 3mg Tablets are blue in colour and are marked with 'W3' on one side. The incorrect Warfant 1 mg Tablets are brown in colour and are marked with the 'W1' on one side.
    - If any 1mg tablets present within a container labelled as 3mg are identified at patient level, advise the patient or their carer not to use the tablets, and to return the container to the pharmacy at their earliest opportunity, where they will obtain a replacement. Quarantine any units returned by your customers and notify Advanz Pharma of any patient returns via the contact details provided below.
    - Advise the patient to contact their prescriber at their earliest opportunity if they have consumed incorrect 1mg strength tablet(s).

3. Please provide the prescribers of patients who have been dispensed Warfant 3mg Tablets since 02/02/2026 with a copy of the attached Dear Doctor Letter.

4. If you have supplied any units of batch no. 25002 of Warfant 3mg Tablets to another pharmacy or facility, please forward a copy of this recall letter to them so that they can perform the requested actions.

5. Please contact your wholesaler to arrange uplift of the quarantined units within the next 14 days. Credit will be arranged at that time.

Replacement unaffected stock is available to order at this time.

Please report any suspect adverse reactions to Advanz Pharma at medicalinformation@advanzpharma.com and to the HPRA at medsafety@hpra.ie.

We sincerely apologise for any inconvenience this may cause. If you have any queries regarding this recall please contact Advanz Pharma at medicalinformation@advanzpharma.com or telephone +353 1800 851 119.

Yours sincerely,

Signed by:
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