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# Batch Recall

**MYNZEPLI® 40mg/mL solution for injection in pre-filled syringe, EU/1/25/1964/002**

| Batch number | Expiry date |
|---|---|
| 2325718 | May-2027 |
| 2461576 | Sep-2027 |

28th of October 2026

Dear Pharmacist,

We wish to advise you that batches **2325718** and **2461576** of MYNZEPLI® 40mg/mL solution for injection in pre-filled syringe are being recalled with immediate effect.

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

The recall is due to out-of-specification dimensions of the finger flange on the pre-filled syringe, which may impair the handling of the syringe during administration.

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

1. Immediately identify and quarantine any units of MYNZEPLI® 40mg/mL solution for injection from **batches 2325718** and **2461576** within your facility. For hospital pharmacies and clinics, this includes units at wards, clinics and other relevant locations within your facility.
2. If you have supplied any units of **batches 2325718** and **2461576** to another pharmacy or clinic, please forward a copy of this letter to them so that they can perform the requested actions.
3. Please contact Allphar to arrange return 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.

If you have any queries regarding this recall please contact Advanz Pharma at medicalinformation@advanzpharma.com or telephone +353 1800 851 119.

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

We apologise for any inconvenience this may have caused you.

Yours sincerely,

Signed by:
Rebecca Burgess October 29, 2026 | 14:29 GMT
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**Rebecca Burgess**
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