

Product Recall**Palmeux prolonged-release suspensions for injection in prefilled syringe****All in date batches**

Product Name	License Number
Palmeux 50 mg prolonged-release suspension for injection in prefilled syringe	PA1142/040/002
Palmeux 75 mg prolonged-release suspension for injection in prefilled syringe	PA1142/040/003
Palmeux 100 mg prolonged-release suspension for injection in prefilled syringe	PA1142/040/004
Palmeux 150 mg prolonged-release suspension for injection in prefilled syringe	PA1142/040/005

Date: 24th December 2025

Dear Pharmacist,

We wish to advise you that all batches of the above-listed products are being recalled with immediate effect.

The recall is to pharmacy level.

This action has been agreed with the Health Products Regulatory Authority (HPRA).

This recall is a precautionary measure due to GMP deficiencies identified at the manufacturing site.

You are requested to perform the following actions:

1. Please immediately quarantine any units of this product which you have in your possession. For hospital pharmacies, this includes units at wards, clinics and any other relevant locations within the hospital.
2. If you have supplied these products to any other pharmacy, clinic or other such healthcare establishment, please forward a copy of this recall letter to them, requesting that they immediately quarantine and return any unused units back to you.
3. Return all quarantined units to your wholesaler. The final date for recalled stock to be received for credit is 19th January 2026.

Unaffected stock is not available to order.

Suspected adverse reactions should be reported to medicalinformation@advanzpharma.com and the Health Products Regulatory Authority (medsafety@hpra.ie).

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

Best regards,



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