



MEDICINES NOTIFICATION

CLASS 4 MEDICINES DEFECT INFORMATION, EL(25)A/41

Caution In Use

Issued: 21 August 2025

Distribute to Pharmacy/Wholesaler Level

MARKETING AUTHORISATION HOLDER (MAH)

Chanelle Medical Unlimited Company

MEDICINE DETAILS

Fexofenadine Hydrochloride 120mg film-coated tablets

PL: 13931/0039

Active Ingredient: Fexofenadine Hydrochloride

GTIN: 5099299199583

SNOMED: 14161911000001109

AFFECTED LOT BATCH NUMBERS

Batch No.	Expiry Date	Pack Size	First Distributed
HM11123A	28/02/2026	30's	20/09/2023
HM10523	28/02/2026	30's	29/08/2024
HM10623	28/02/2026	30's	19/09/2024
HM10723A	28/02/2026	30's	19/09/2024
HM10823	28/02/2026	30's	31/10/2024
HM10923	28/02/2026	30's	15/01/2025
HM11023	28/02/2026	30's	31/01/2025
HM11223	28/02/2026	30's	11/02/2025

HM11323	28/02/2026	30's	17/02/2025
HM11423	28/02/2026	30's	25/02/2025
HM11623	28/02/2026	30's	25/02/2025
HM11523A	31/03/2026	30's	25/02/2025
HM11723	31/03/2026	30's	11/03/2025
HM11823	31/03/2026	30's	31/03/2025
HM11923	31/03/2026	30's	02/04/2025
HM12023	31/03/2026	30's	02/04/2025
HM12123	31/03/2026	30's	02/04/2025
HM12223	31/03/2026	30's	02/04/2025
HM12323A	31/03/2026	30's	02/04/2025
HM12423	31/03/2026	30's	08/04/2025
HM12523	31/03/2026	30's	24/04/2025
HM12623	31/03/2026	30's	30/04/2025
HM12723	30/04/2026	30's	16/04/2025
HM12823	30/04/2026	30's	20/05/2025
HM12923	30/04/2026	30's	11/06/2025
HM13023	30/04/2026	30's	13/06/2025

Background

Chanelle Medical Unlimited Company has informed the MHRA of an error with the European Article Number (EAN) / Global Trade Item Number (GTIN) barcode on the cartons of the above batch of Fexofenadine Hydrochloride 120mg film-coated tablets, distributed by Healthcare Pharma Limited. When scanned, the EAN/GTIN barcode identifies the product as Naratriptan 2.5mg Tablets.

The other product details on the carton, including the name, strength and pharmaceutical form of the medicine are correct. The quality of the medicine is not impacted by the labelling defect.

Advice for Healthcare Professionals:

Healthcare professionals are advised to use caution and consider extra safeguards for these batches in robotic or automated dispensing system or stocking systems and should carry out manual dispensing and stocking, as appropriate.

The product quality of the Fexofenadine Hydrochloride 120mg film-coated tablets is not impacted by this issue, therefore the affected batches are not being recalled.

All remaining stock of Naratriptan 2.5mg Tablets which has not been distributed from UK wholesaler will be placed on hold until April 2026 (Last exp of impacted Fexofenadine stock) to provide further mitigation that the wrong product may be selected.

Advice for Healthcare Professionals to Provide to Patients:

No action is needed from patients, continue to take medication from these batches as prescribed by your healthcare professional. The issue is related to the wrong barcode on the cartons of the listed batch of Fexofenadine Hydrochloride 120mg film-coated tablets and will be controlled by the healthcare professionals prescribing/dispensing the medication. The quality of the medication is not affected. Patients who experience adverse reactions or have any questions about their medication should seek medical attention. Any suspected adverse reactions should also be reported via the [MHRA Yellow Card scheme](#).

Additional information:

For all medical information enquiries and information on this product, please contact Medical Information Department chanelle@medinformation.co.uk or telephone +44 808 304 2429 (Toll free-UK). For stock control enquiries please email David Hammond: david.hammond@healthcarepharma.co.uk.

Recipients of this Medicines Recall should bring it to the attention of relevant contacts by copy of this notice. NHS regional teams are asked to forward this to community pharmacists and dispensing general practitioners for information.

Yours faithfully

Defective Medicines Report Centre

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