

# MEDICINES RECALL

## CLASS 2 MEDICINES RECALL, EL(26)A/34

### Action within 48 hours

Issued 13 July 2026

Distribute to Pharmacy/Wholesaler Level

#### MARKETING AUTHORISATION HOLDER (MAH)

Beckton Dickson UK Ltd

#### MEDICINE DETAILS

##### ChloraPrep 1mL Applicator

PL: 05920/0002

Active Ingredient: Chlorhexidine gluconate 2%w/v and isopropyl alcohol 70% v/v

SNOMED code: 40818111000001106

GTIN: 20885403491761

#### AFFECTED LOT BATCH NUMBERS

| Batch No.                                                 | Expiry Date                                               | Pack Size                                 | First Distributed |
|-----------------------------------------------------------|-----------------------------------------------------------|-------------------------------------------|-------------------|
| All lots with an expiry date up to and including 05/2029* | All lots with an expiry date up to and including 05/2029* | ChloraPrep 1mL Applicator: 60 applicators | 25/06/2025        |

\*For a full list of impacted batches, see [Annex 1](#)

## Background

Becton Dickinson UK Ltd are recalling batches of ChloraPrep 2% w/v / 70% v/v cutaneous solution 1mL applicators due to a potential breach of sterility in the packaging process. The recall applies to all batches with an expiry date up to and including 05/2029. Batches with an expiry date after 05/2029 are not impacted.

This defect could increase the risk of the applicator device being contaminated with pathogens, which could lead to increased infection rates for the patients. The number of impacted units in each batch is unknown, therefore the batches included in [Annex 1](#) are being recalled as a precautionary measure.

This recall is specific to the lots included in [Annex 1](#).

### **Advice for Healthcare Professionals:**

Stop supplying the affected batches. Quarantine all remaining stock and return it to your supplier using your supplier's approved process.

### **Advice for Healthcare Professionals to Provide to Patients:**

No further action is required by patients as this product is used by healthcare professionals and they will have taken the action to remove this product from use.

The recall is a precautionary measure to mitigate any additional risk of infection.

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 email [safetyinformation@bd.com](mailto:safetyinformation@bd.com), or telephone 08000437546.

For stock control enquiries please email [info@insightbio.com](mailto:info@insightbio.com), or telephone 01707351330.

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

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