**FOOD AND DRUGS AUTHORITY**

## RECALLED PRODUCT DETAILS

### 1. DORIFA VIGO PLUS CAPSULES

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|---|---|
| Date Recall was issued | 22nd May 2025 |
| Product Name | Dorifa Vigo Plus Capsules |
| Product Type | Herbal |
| Manufacturer | Dorifa Manufacturing Limited |
| Recalling Firm | Dorifa Manufacturing Limited |
| Batch(es) | All batches |
| Manufacturing Date | N/A |
| Expiry Date | N/A |
| Reason for Recall | **Unapproved packaging material** |



## 2. 4FR SINGLE LUMEN POWDER PICC

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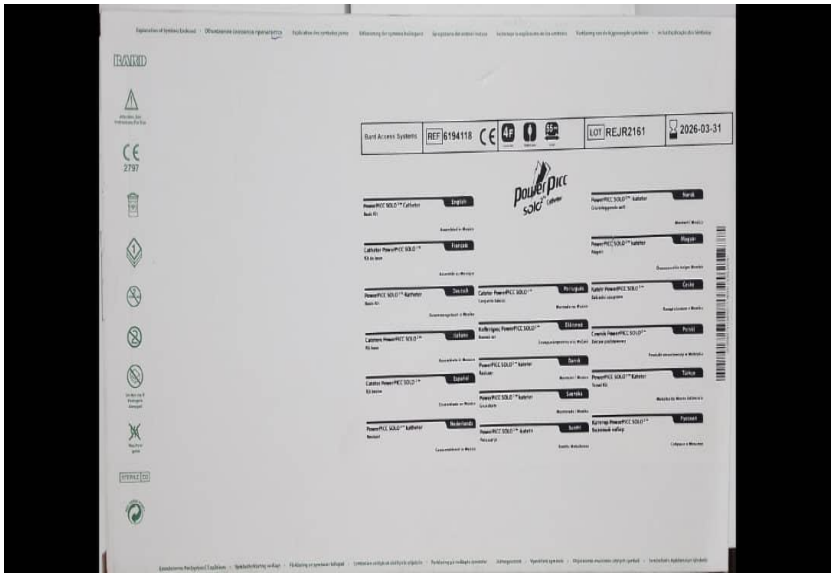
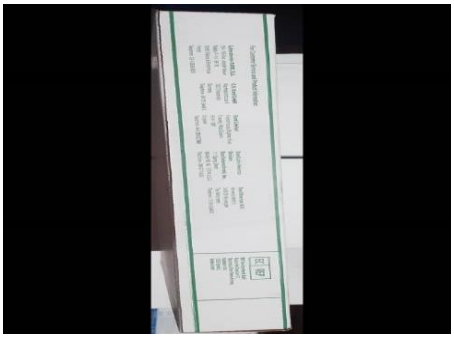
|---|---|
| Date Recall was issued | 27th May 2025 |
| Product Name | 4Fr Single Lumen Powder PICC |
| Product Type | Medical Device |
| Manufacturer | Becton Dickinson (PTY) Limited: |
| Recalling Firm | East Cantonments Pharmacy Limited |
| Batch(es) | REJS3341, REHY3938, REJR2161 |
| Manufacturing Date | N/A |
| Expiry Date | N/A |
| Reason for Recall | This recall is necessitated because of a global product removal of specific lots of 4Fr single -lumen power PICC catheters due to an increase of material fatigue leaks in specific geographies. |



## 3. COMPOUND SODIUM LACTATE INFUSION BP (RINGER LACTATE SOLUTION FOR INJECTION

| | |
|---|---|
| Date Recall was issued | 2nd May 2025 |
| Product Name | Compound Sodium Lactate Infusion BP (Ringer Lactate Solution for Injection) |
| Product Type | Drug |
| Manufacturer | Atlantic Life Science Ltd |
| Recalling Firm | Atlantic Life Science Ltd |
| Updated Batch(es) | 02C221, 02D24, 02D25, 02D26, 02D206, 02D208, 02E19, 02E20, 02E107w, 02E108w |
| Manufacturing Date | 02/2025 |
| Expiry Date | 01/2028 |
| Reason for Recall | **Substandard.** This recall is necessitated as a result of laboratory analysis conducted on samples of the products by the FDA Centre for Laboratory Services and Research, which indicated that the samples did not meet the requirements of NMT 0.25iu/ml for bacterial endotoxin as per the BP specifications. |

**PICTURES**