

#### **FOOD AND DRUGS AUTHORITY**

### **RECALLED PRODUCT DETAILS**

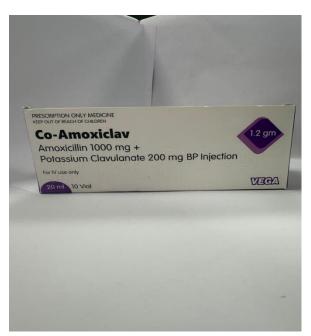
# 1. CO-AMOXICLAV (AMOXICILLIN 1000MG + POTASSIUM CLAVULANATE 200MG INJECTION)

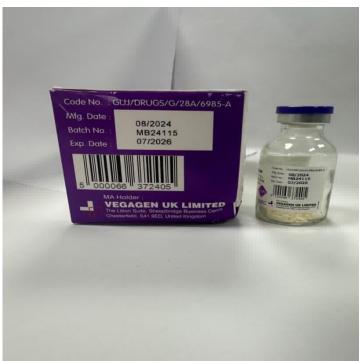
| Date Recall was issued | 22 <sup>nd</sup> July 2026                |
|------------------------|---|
| Product Name           | See list of products below                |
| Product Type           | Drug                                      |
| Manufacturer           | Vegagen UK Ltd                            |
| Recalling Firm         | Panacea Pharmaceuticals Limited           |
| Batch(es)              | See list below                            |
| Manufacturing Date     | See list below                            |
| Expiry Date            | See list below                            |
| Reason for Recall      | Substandard. The product failed assay for |
|                        | Clavulanic Acid as per the acceptance     |
|                        | criteria in the British Pharmacopoeia.    |

#### PANACEA RECALL DETAILS

| Product<br>Description   | Batch Numbers | Manufacturing<br>Dates | Expiry Dates |
|--|---------------|------------------------|--------------|
| Co- Amoxiclav (Amoxicillin 1000mg + Potassium Clavulanate 200mg Injection)   | MB24115       | 08/2024                | 07/2026      |
| Co- Amoxiclav (Amoxicillin 1000mg + Potassium Clavulanate 200mg Injection Instant Oats Morning Mills 500g, Variety | MB24034       | 03/2024                | 02/2026      |







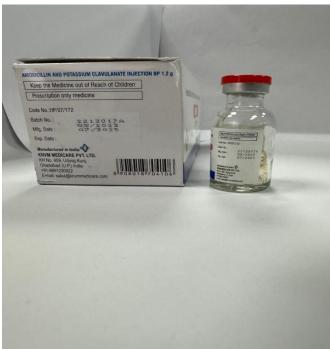
## 2. KMCLAV 1.2G INJECTION (AMOXICILLIN AND POTASSIUM CLAVULANATE INJECTION BP

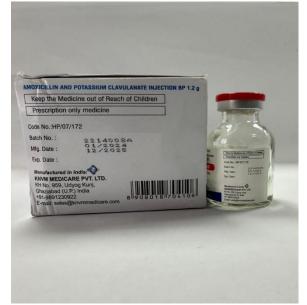
| Date Recall was issued | 22 <sup>nd</sup> July 2025                |
|------------------------|---|
| Product Name           | KMCLAV 1.2g Injection (Amoxicillin and    |
|                        | Potassium Clavulanate Injection BP)       |
| Product Type           | Drug                                      |
| Manufacturer           | KNVM Medicare Pvt Ltd, India              |
| Recalling Firm         | Med House Pharmaceuticals Ltd             |
| Batch(es)              | See list below                            |
| Manufacturing Date     | See List below                            |
| Expiry Date            | See List below                            |
| Reason for Recall      | Substandard. The product failed assay for |
|                        | Clavulanic Acid as per the acceptance     |
|                        | criteria in the British Pharmacopoeia.    |

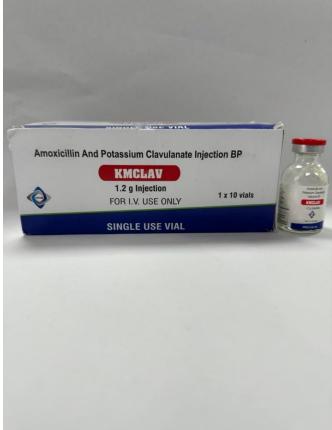
#### **Med House Pharmaceuticals Recall Details**

| Product<br>Description  | Batch Numbers | Manufacturing<br>Dates | Expiry Dates |
|---|---------------|------------------------|--------------|
| KMCLAV 1.2g Injection (Amoxicillin and Potassium Clavulanate Injection BP)) | 2213017A      | 08/2023                | 07/2025      |
| KMCLAV 1.2g Injection (Amoxicillin and Potassium Clavulanate Injection BP)) | 2214008A      | 01/2024                | 12/2025      |









## 3. CLAVU- NOVA 600G (CO- AMOXICLAV FOR INJECTION BP)

| Date Recall was issued | 22 <sup>nd</sup> July 2025                     |
|------------------------|--|
| Product Name           | Clavu- Nova 600g (Co- Amoxiclav for Injection  |
|                        | BP)  |
| Product Type           | Drug   |
| Manufacturer           | Bharat Parenterals Ltd, India                  |
| Recalling Firm         | Pharmanova Limited                             |
| Batch(es)              | P1211, P13212                                  |
| Manufacturing Date     | 11/2023  |
| Expiry Date            | 10/2025  |
| Reason for Recall      | Substandard. The product failed assay for      |
|                        | Clavulanic Acid as per the acceptance criteria |
|                        | in the British Pharmacopoeia.                  |







## 4. AMOVULIN (CO- AMOXICLAV FOR INJECTION BP 1.2GM)

| Date Recall was issued | 22 <sup>nd</sup> July 2025                |
|------------------------|---|
| Product Name           | AmoVulin (Co- Amoxiclav for Injection BP  |
|                        | 1.2gm)                                    |
| Product Type           | Drug                                      |
| Manufacturer           | Tobinco Pharmaceuticals Ltd               |
| Recalling Firm         | Tobinco Pharmaceuticals Ltd               |
| Batch(es)              | IC2478002                                 |
| Manufacturing Date     | 01/2024                                   |
| Expiry Date            | 12/2025                                   |
| Reason for Recall      | Substandard. The product failed assay for |
|                        | Clavulanic Acid as per the acceptance     |
|                        | criteria in the British Pharmacopoeia.    |



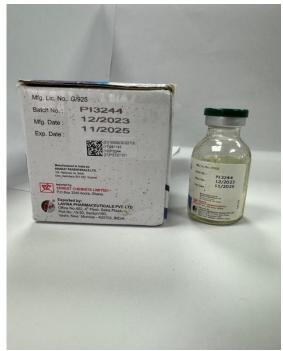




## 5. LAVI- CV 1.2G (CO- AMOXICLAV FOR INJECTION 1.2G)

| Date Recall was issued | 22 <sup>nd</sup> July 2025                        |
|------------------------|---|
| Product Name           | LAVI- CV 1.2 G (Co - Amoxiclav for Injection      |
|                        | 1.2g)   |
| Product Type           | Drug  |
| Manufacturer           | Bharat Parenterals Ltd, India                     |
| Recalling Firm         | Ernest Chemist Ltd                                |
| Batch(es)              | P13244  |
| Manufacturing Date     | 12/2023   |
| Expiry Date            | 11/2025   |
| Reason for Recall      | Substandard. The product failed assay for         |
|                        | Clavulanic Acid as per the acceptance criteria in |
|                        | the British Pharmacopoeia.                        |







## 6. OXYNIC- 600 ( AMOXICILLIN AND POTASSIUM CLAVULANATE INJECTION BP)

| Date Recall was issued | 22 <sup>nd</sup> July 2025                        |
|------------------------|---|
| Product Name           | Oxynic - 600 (Amoxicillin and Potassium           |
|                        | Clavulanate Injection BP)                         |
| Product Type           | Drug  |
| Manufacturer           | GB Pharma Ltd                                     |
| Recalling Firm         | GB Pharma (GH) Ltd                                |
| Batch(es)              | MB23162   |
| Manufacturing Date     | 12/2023   |
| Expiry Date            | 11/2025   |
| Reason for Recall      | Substandard. The product failed assay for         |
|                        | Clavulanic Acid as per the acceptance criteria in |
|                        | the British Pharmacopoeia.                        |



