



MINISTRY OF HEALTH
PHARMACY AND POISONS BOARD

RAPID ALERT NOTIFICATION OF A QUALITY DEFECT / RECALL

Reference Number : **2025/005**

1. To: **World Health Organization, Market Authorization Holders, Distributors, Retailers and Consumers**

2. Product Class of Defect: **Class I**

3. **Substandard**

4. Product : **Paracetamol 1% W/V injection**

5. Marketing Authorization Number : **CTD 4994**

For use in humans

6. Brand/Trade Name : **Blink injection**

7. INN or Generic Name: **Paracetamol injection**

8. Dosage Form : **Injection**

9. Strength: **Paracetamol 1% W/V**

10. Batch Numbers: **CS4594005 and CS4594004**

11. Expiry Date: **09/2024**

12. Pack size and Presentation: **Clear, colourless solution in a 100ml bottle**

13. Date of Manufacture : **08/2026**

14. Marketing Authorization Holder : **Crown Healthcare Ltd**

15. Manufacturer : **KamlaAmrut Pharmaceutical LLP, India.**

16. Recalling Firm (if different): **Crown Healthcare, Nairobi, Kenya.**

Contact Person: **Dr. Rahab Njuguna**

15.1 Where the defect is attributed to a manufacturing site, site where defect occurred : **KamlaAmrut Pharmaceutical LLP, India.**

Telephone: **+254 721 200 400 | +254 733 200 400**

17. Recall Number Assigned (if available) : **REC/2025/014**

18. Details of Falsification :
N/A

19. Information on distribution including exports (type of customer, e.g. hospitals) :
Consumer/patients.

20. Action taken by Issuing Authority: Issue of a Mandatory Recall notice for Blink (Paracetamol 1%W/V) Injection Batch No. CS4594005 and CS4594004	
21. Proposed Action : Healthcare Professional to stop further distribution, sale, issuing, or use of the batch immediately.	
22. From (Issuing Authority): Kenya Pharmacy and Poisons Board	23. For Feedback Contact: pms@ppb.go.ke Telephone: 0795743049
24. Date : 24th April 2025	