



Ranbaxy Pharmaceuticals (Pty) Ltd
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20 May 2026

URGENT MEDICINE RECALL – CLASS II-TYPE B

Dear Customer/Healthcare Professional

Ranbaxy Pharmaceuticals (Pty) Ltd in collaboration with the South African Health Product Regulatory Authority (SAHPRA), is recalling the below mentioned batch of RANCEPH SUSPENSION 125 mg/5 ml.

Product	Batch number(s)	Expiry date	Manufacturing date	Pack size	First release date for sale	Quantity initially released	Recall Classification (As confirmed by SAHPRA)
RANCEPH SUSPENSION 125 mg/5 ml	DFG4606A	06/2027	07/2025	100 ml	06 November 2025	15151	Class II Type B

The recall is initiated because of an Out of Specification (OOS) result observed in the Related Substance test, for an Unidentified impurity, at the 6-month stability time point (25° C/60% RH).

We request that you refrain from selling the affected batch of RANCEPH SUSPENSION 125 mg/5 ml, Batch DFG4606A. Please return all remaining stock to your supplying warehouse or distributor with immediate effect. Ranbaxy Pharmaceuticals (Pty) Ltd will collect all your stock which will be credited.

Kindly retain this letter in a prominent position for one month, in case there is stock in transit.

For further product information, please call Ranbaxy Pharmaceuticals (Pty) Ltd on 011 495 0100.

We thank you in advance for your co-operation and apologise for any inconvenience caused.

Yours faithfully

Geeta Ghela
Responsible Pharmacist

Name	Title	Designation	Contact Number
Geeta Ghela	Ms	Responsible Pharmacist/QA Head	Tel: 011 495 0100 Email: geeta.ghela@sunpharma.com