

Tuesday, 21 July 2026

URGENT MEDICINE RECALL – CLASS II, TYPE B

Dear Customer,

We hereby inform you that we are recalling the following batches of **Nintedanib 100 mg Cipla** and **Nintedanib 150 mg Cipla**

(SAHPRA allocated registration numbers: 56/26/0704.702 & 56/26/0705.703, respectively).

Product	Batch number	Manufacturing date	First release date for sale	Expiry Date	Pack Size	Quantity initially released	Recall Classification (As confirmed by SAHPRA)
Nintedanib 100mg Cipla	6KA0781	02.2026	14.04.2026	01.2028	60's	1689	CLASS II, TYPE B
Nintedanib 150mg Cipla	6KA0782	02.2026	14.04.2026	01.2028	60's	1675	

This recall is initiated as a result of product quality complaints received due to soft-gel capsules clumping/sticking together.

We request that you refrain from selling or dispensing any pack of the affected batches of Nintedanib 100 mg and 150 mg Cipla and return the products to your supplying warehouse or distributor with immediate effect for a full credit/replacement

For further product information, please call Cipla Medpro (Pty) Ltd. Customer Care line (080 222 6662).

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

Yours faithfully,



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Mrs Nicole Carter

Responsible Pharmacist and Director Quality Operations
Cipla Medpro (Pty) Ltd.

GLF-RC-INSP-05B_v2-Recall-Letter-to-customer - Nintedanib 100mg and 150mg

Final Audit Report

2026-07-21

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