



Dear Healthcare Professional

URGENT MEDICINE RECALL (CLASS II TYPE A)

Class II recall is intended for medicines that possibly could cause temporary or medically reversible adverse health problem or mistreatment.

Type A recall is designed to reach all suppliers of medicines (all distribution points) i.e., wholesalers, hospitals, retail pharmacy outlets, doctors, nurses, pharmacists, authorised prescribers and dispensers, and individual customers or patients

YAZ PLUS

Registration No.: 45/21.8.2/0534

Bayer (Pty) Ltd, in consultation with the South African Health Products Regulatory Authority (SAHPRA), is conducting a Class II Type A recall of the following batch of YAZ PLUS Tablets:

Product Name	Dosage Form	Pack Size	Batch No	Expiry Date
YAZ PLUS	Tablet	28's	WEW96J	03/2026

Reason for Recall:

This recall is initiated due to a limited number of packs found in retail Pharmacy, with mix-up of the sequence of hormone-containing and hormone-free tablets, wherein some packs had

- 24 light orange hormone-free, and
- 4 pink film-coated hormone tablets

Instead of

- 24 pink film-coated hormone tablets and
- 4 light orange hormone-free tablets.

Mix-up of arrangement



Incorrect blister



Correct blister

The defective packs contain only 4 hormone tablets instead of 24 hormone tablets and will not provide the expected contraceptive efficacy. A return to supplier of Batch WEW96J is recommended.

Action required:

- Please return all stocks of this batch to DSV Healthcare or the wholesaler/distributor that you purchased it from, for credit.
- Kindly retain this letter in a prominent position for one month, in case stock is still in transit.

For further information about this Recall, please contact Bayer at 011 921 5900

Yours sincerely
Bayer (Pty) Ltd

Eric Chauke

Regulatory Affairs Head/Responsible Pharmacist



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Regulatory Affairs Department

Bayer (Pty) Ltd

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