

NATIONAL



AUTHORITY

047/NDA/DPS/01/2019

22nd January 2019

Attention: Health Care Providers

CIRCULAR NO. 028

FUROSEMIDE INJECTION MANUFACTURED BY GRAND PHARMACEUTICAL (CHINA) Co., Ltd; BATCH NUMBER 180120

National Drug Authority in its routine Post Marketing Surveillance discovered that the product with details below has mismatching information on the Patient Information Leaflet (PIL) and both the primary and secondary packages. The product strength indicated on both the primary and secondary packages is 10mg/2ml whereas the PIL shows product to be of strength 10mg/ml.

Product Name	Batch	Mfg. Date	Expiry Date	Manufacturer
Furosemide Injection BP	180120	01/2018	01/2021	Grand Pharmaceutical (China) Co., Ltd.

The above mismatching information is confusing, misleading and can lead to wrong/irrational use of the drug.

The purpose of this communication is to advise you to;

- i. Stop the purchase, supply and use of the above product with immediate effect. The license holder of this product has been instructed to recall the product and cease further distribution.
- ii. Quarantine any available stock in your possession and return to your supplier.
- iii. Be vigilant and report to NDA any other product defects and any adverse drug reactions associated with the use of above product or other products. Reports can be sent to NDA offices or by email to druginfo@nda.or.ug or calling **0414-255665** or by sending a message on WhatsApp to **0791415555**.

Thank you for your continued cooperation.

Helen Byomire Ndagije
DIRECTOR PRODUCT SAFETY

HEAD OFFICE

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OUR MISSION

Promoting and protecting public health through the effective regulation of human and animal medicines and healthcare products

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