

NATIONAL



AUTHORITY

8th February 2024

Ref: 0232G/DR/NDA/02/2024

To: All Licence Holders
Manufacturers
Local Technical Representatives

CIRCULAR NO: 01/DPAR/2024

SUBMISSION OF VETERINARY PHARMACEUTICAL PRODUCT DOSSIERS IN THE COMMON TECHNICAL DOCUMENT (CTD) FORMAT.

National Drug Authority is mandated to ensure the availability, at all times, of essential, efficacious and cost-effective drugs to the entire population of Uganda as a means of providing satisfactory healthcare and safeguarding the appropriate use of drugs. This is accomplished through the regulation and control the production, importation, distribution and use of human and veterinary medicines.

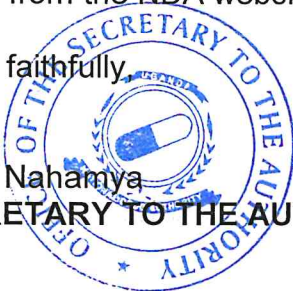
This is to notify all Marketing Authorization Holders, Manufacturers, and Local Technical Representatives for veterinary pharmaceutical products that National Drug Authority is no longer receiving applications for marketing authorization of veterinary pharmaceutical products compiled in non-CTD format.

For the registered products and those under assessment, a grace period ending December 2025 has been granted to convert the dossiers to CTD format.

The new guidelines and the corresponding registration application form can be accessed online from the NDA website; <https://www.nda.or.ug>.

Yours faithfully,

David Nahamya
SECRETARY TO THE AUTHORITY



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OUR MISSION

To protect and promote human and animal health through the effective regulation of drugs and healthcare products.

REGIONAL OFFICES

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