

TANZANIA FOOD AND DRUGS AUTHORITY



PUBLIC NOTICE

**APPLICATION GUIDELINES FOR REGISTRATION OF HUMAN
MEDICINAL PRODUCTS**

The Tanzania Food and Drugs Authority (TFDA) would like to notify all prospective and current registration holders who intend to apply for new registration, renew or vary registration of registered human medicinal products in Tanzania, that the Authority has issued revised Guidelines for submission of applications. The guidelines and application forms are available online (free of charge) at www.tfda.or.tz or in hard copies at a nominal fee of US\$ 30 or equivalent in Tanzanian Shillings. The guidelines are:

1. Guidelines on Submission of Documentation for Registration of Human Medicinal Products, Fourth Edition, November, 2008.
2. Application Guidelines for Variation of Registered Medicinal Products, First Edition, November, 2008.

These guidelines which will come into operation from 1st January, 2009 will replace the current “Application Guidelines for Registration of Human Medicinal Products, Third Edition, June 2004”. After they have come into operation, all applications shall be submitted in accordance with the new guidelines. Applications which will not comply with the new guidelines will not be accepted.

The Authority wishes to thank you for having actively participated in the revision of the guidelines.

For more information we invite you to visit or contact us through the following address:

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