GOVERNMENT OF THE PEOPLE'S REPUBLIC OF BANGLADESH
MINISTRY OF HEALTH AND FAMILY WELFARE
DIRECTORATE GENERAL OF DRUG ADMINISTRATION
AUSHADH BHABAN
MOHAKHALI, DHAKA, BANGLADESH
www.dgda.gov.bd

Ref. WHO TRS 902, 2002 ANNEX-9 Date: 01/04/2020

Guidelines on Packaging for Pharmaceutical Products

DGDA, as a competent medicine regulatory authority gives marketing authorization of a new pharmaceutical product for the purpose of marketing or distribution after evaluation of its safety, efficacy and quality.

During product registration MA holder should submit the relevant packaging materials as per this guideline

To approve packaging materials properly and effectively guidelines are needed. In this purpose WHO Guidelines on packaging for pharmaceuticals products is very extensive guidelines which also covered all the aspect and issues of packaging for marketing authorization.

So here this is to certify that WHO Guidelines on Packaging for pharmaceuticals products WHO TRS 902, 2002, Annex-9 is adopted for the purpose of variation of marketing authorization in DGDA from the date given below and valid up to next office order.

01/04/2020

Major General Md Mahbubur Rahman
Director General
Directorate General of Drug Administration
&
Licensing Authority (Drugs)