



Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION



08 June 2020

TO : ALL CONCERNED DRUG IMPORTERS AND OTHER RELEVANT STAKEHOLDERS

FROM : JESUSA JOYCE N. CIRUNAY, RPh
Director IV
Center for Drug Regulation and Research (CDRR)

SUBJECT : Draft FDA Circular on Interim Guidelines for the Issuance of Foreign cGMP Clearance

Please be informed that the CDRR is soliciting for comments from the concerned stakeholders on the draft FDA Circular entitled: **Interim Guidelines for the Issuance of Current Good Manufacturing Practice (cGMP) Clearance to Foreign Drug Manufacturers**

Comments may be submitted by email to: cdrr.od@fda.gov.ph not later than **10 June 2020 (Wednesday) at 3:00PM** using the following format:

Name of Company/Organization:		
Contact information		
Telephone number:		Email address:
Section	Comments	Input/Recommendation/ Suggested text (if any)

In sending your comments, please use the subject: **[Comments] Interim Guidelines for FcGMP Clearance**

Thank you.



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