03 April 2012

FDA MEMORANDUM CIRCULAR
No. 2012-003

TO : All Drug Manufacturers Applying for Renewal of License to Operate (LTO)

SUBJECT : Full Compliance to Good Manufacturing Practice (GMP)

In line with the implementation of RA 9711 or the “Food and Drug Administration (FDA) Act of 2009”, which aims to protect and promote the right to health of the Filipino people and to establish and maintain an effective health product regulatory system, it is hereby reiterated that all Drug Manufacturers applying for renewal of License to Operate (LTO) should ensure full compliance to Good Manufacturing Practice (GMP).

For your strict and immediate compliance.

[Signature]

SUZETTE H. LAZO, MD, FPSECP
Acting Director IV