


FDA MEMORANDUM CIRCULAR
No. 2013-019

DATE : 10 May 2013

TO : ALL CONCERNED PHARMACEUTICAL COMPANIES

FROM : 
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Acting Director IV

SUBJECT : DECKING SCHEDULE OF DRUG PRODUCT REGISTRATION APPLICATIONS

In consonance with FDA Memorandum Circular No. 2013-04 Subject : Follow-Up of Applications, the Center for Drug Regulation and Research (CDRR) will no longer accept Follow-Up Forms for Product Registration Applications.

Please be advised that effective 14 May 2013, the CDRR will regularly post at its bulletin boards the list of initial and renewal registration applications filed **beginning February 2013** with corresponding decking schedule following the First-In First-Out Policy and CDRR Memo No. 0003, s, 2013 Subject: Facilitation of Application for Product Registration. The same list will be uploaded in the FDA website.

Renewal registration applications filed in January 2013 are currently being processed while there are some that were already granted Certificates of Product Registration (CPR). Also, all pending 2012 applications are already being acted upon.

For your information and compliance.

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