



CENTER FOR DRUG REGULATION AND RESEARCH

**CERTIFICATE OF LISTING OF IDENTICAL DRUG PRODUCT (CLIDP)
APPLICATION**

- Who May Avail** : All Manufacturers, Distributors, Importers, Exporters, Wholesalers, and Traders of Pharmaceutical Products
- Fees to be Paid** : **AO No. 50 s. 2001 and AO No. 2005-0031**
Branded: PHP 3,000.00/year + PHP 500.00 (per proposed brand name, for brand name clearance) + 1% LRF
Unbranded: PHP 2,000.00/year + 1% LRF
*per year – depending on the remaining validity of the Principal Certificate of Product Registration (PCPR)

CHECKLIST OF REQUIREMENTS

1. Duly Accomplished and Notarized Integrated Application Form
2. Proof of payment (based on AO No. 50 s. 2001 and AO No. 2005-0031)
3. Copy of the current and valid LTO of the PCPR and Identical Drug Applicant
4. Copy of current and valid PCPR and copies of previously approved certificates and/or notifications during the validity of the CPR
5. Authenticated copy of the duly notarized Distributorship Agreement, license Agreement, or other written contract between the principal CPR holder and the identical Drug Applicant
6. For Imported Products:
Copy of valid Certificate of GMP Clearance (and/or renewal application, whichever is applicable)
7. Facsimile of Labeling Materials

END

