

# Republic of the Philippines Department of Health



#### FOOD AND DRUG ADMINISTRATION

# CENTER FOR DRUG REGULATION AND RESEARCH

## INITIAL REGISTRATION OF VETERINARY DRUGS

Who May Avail : All Manufacturers, Distributors, Importers, Exporters, Wholesalers,

and Traders of Veterinary Drug Products

Fees to be Paid : Initial:

Branded: PHP 3,000.00/year + PHP 500.00 (Brand Name Clearance)

+ 1% LRF

Unbranded: PHP 2,000.00/year + 1% LRF

The applicant may apply for 2/5-year CPR validity.

2-year validity:

Branded: PHP 6,000.00 + PHP 500.00 (for Brand Name Clearance) =

PHP 6,500.00 + 1% LRF

Unbranded: PHP 4,000.00 + 1% LRF

5-year validity:

Branded: PHP 15,000.00 + PHP 500.00 (for Brand Name Clearance)

= PHP 15,500.00 + 1% LRF

Unbranded: PHP 10,000.00 + 1% LRF

### **CHECKLIST OF REQUIREMENTS**

- 1. Duly accomplished and notarized Integrated Application Form (with proof of payment)
- 2. Valid agreements between the manufacturer, trader, importer, distributor, where applicable
- 3. Unit Dose and Batch Formulation
- 4. Technical Specifications of all Raw Materials
- 5. Certificate of Analysis of Active Raw Material(s)
  - a. From supplier of API
  - b. From manufacturer of finished product
- 6. Technical Specifications of Finished Product
- 7. Certificate of Analysis (CA) of Finished Product (from the same batch of representative sample)
- 8. Manufacturing Procedure, Production, Equipment, Sampling, In-process controls, and Master Packaging Procedure (including specification for container closure system)
- 9. Assay and Other Test Procedures including Identity, Purity Tests, with Data Analysis, where applicable
- 10. Stability Studies
- 11. Labeling Materials (facsimile labels)
- 12. Representative Sample (upon request of the evaluator)

#### **Additional Requirements:**

- 1. For products in plastic container: Certificate of Analysis for Test of Migratable Substances/ Leachability
- 2. For imported products:
  - a. Certificate of Pharmaceutical Product (CPP)
  - b. Foreign Good Manufacturing Practice (GMP) Clearance



- 3. For new veterinary drugs: a. Pre-clinical studies

  - b. Protocol for monitored release
- 4. For fixed-dose combination: Rationale of the Combination
- 5. Valid LTO (Importer/Manufacturer/Distributor/Trader)

**END**