

REPUBLIC OF THE PHILIPPINES DEPARTMENT OF FINANCE BUREAU OF CUSTOMS MANILA

CUSTOMS MEMORANDUM ORDER

AUG 0 5 1993

All Collectors of Customs
Division/Service/Unit Chiefs
(POM, MICP and NAIA)
All Importers/Customs Brokers
All Others Concerned

Subject: Chemical/Pharmaceutical Shipments Requiring Mandatory Referral for Laboratory Analysis

OBJECTIVE:

To ensure collection of proper duties and taxes and strictly enforce all Customs laws, rules and regulations governing chemical and pharmaceutical shipments.

GENERAL PROVISIONS:

It shall be a mandatory requirement that all shipments of pharmaceuticals and chemicals falling under Section II to VII of the TCCP be subject to Laboratory Analysis in any of the following cases:

- Shipments with invoice value (F.O.B) below \$500 and therefore not covered by a SGS Clean Report of Findings. Exempted from this provision are:
 - Chemicals and pharmaceutical products registered with the BOC Laboratory as provided for in III-4 below;
 - (2) Shipments of regulated/controlled drugs, pharmaceuticals and other substances covered by Dangerous Drugs Board import permits, the handling and treatment of which shall be covered in a separate memorandum order.
 - (3) Finished products of pharmaceuticals ready for use.
 - The shipment is covered by alert/derogatory information related to valuation, classification/description and issued pursuant to existing regulations.

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III OPERATIONAL PROCEDURES:

- The analysis required herein shall be performed by the Phil Customs Laboratory (PCL) unless such cannot be performed thereat for absence of the required facilities/expertise.
- 2. It shall be the duty of the COO III to insure that the shipment covered by this order are referred to the PCL.
- 3. The assigned COO III shall obtain samples of the shipment following the procedures defined in the Customs Laboratory Operations Manual. The sample shall be submitted to the laboratory accompanied by a photocopy of the import entry and brochure/literature showing the chemical name as described in III-1 of CMC No. 210-93 dated July 20,1993.
- 4. To register chemicals and pharmaceuticals, the importing firm must submit the following to the PCL (Attn.: The Chief, PCL):
 - (1) Sample of the chemical/pharmaceutical;
 - (2) A statement under oath describing its chemical composition, its commercial name (if any), its physical properties, usage and handling/packaging and manufacturer; and
 - (3) Brochures or technical materials supporting 2 above.
- 5. Application for registration shall be acted upon within five (5) working days. A record of such registration will be made containing the following:
 - a. Registration No.
 - b. Date
 - c. Firm
 - d. Chemical Name
 - e. Commercial Name
 - f. Manufacturer
 - g. IR trace
 - h. Sample (to be deposited in sample room)
 - i. Location in sample room
- 6. A Registration Certificate shall be issued to the firm to be valid for a period of one year from date of registration.

IV EFFECTIVITY:

This CMO shall take effect fifteen (15) calendar days after signing of this order.

Commissioner