

10320 COMPETITIVE PROBLEMS IN THE DRUG INDUSTRY

DEFENSE MEDICAL PURCHASE DESCRIPTION		NUMBER	DATE
		1	5 April 1973
FEDERAL STOCK NO.	ITEM IDENTIFICATION	POTENCY	UNIT
6505-655-8355	TETRACYCLINE HYDROCHLORIDE CAPSULES, USP, 0.25 Gram, 100s	Not less than 48 Months	Bottle

Shall be Tetracycline Hydrochloride Capsules, U.S.P., as produced by Lederle Laboratories as "Achromycin V Capsules."

Each capsule shall contain 250 mg of Tetracycline Hydrochloride.

Shall conform to the pertinent provisions of the General Regulations for the Certification of Antibiotics and Antibiotic-Containing Drugs as promulgated by the Food and Drug Administration (F.D.A.), U.S. Department of Health, Education, and Welfare.

Each lot shall be certified by the F.D.A.

Not more than 4 months of the expiration dating period (potency period) shall have elapsed at the time of delivery to the Government.

Shall be supplied 100 capsules in a commercially available bottle.

Labeling shall comply with the requirements of the Federal Food, Drug, and Cosmetic Act and, in addition, shall include labeling in accordance with commercial practice.

PREPARATION FOR DELIVERY

Packaging and packing. Level C. The subject commodity shall be packaged and packed in substantial commercial containers of the type, size, and kind commonly used for the purpose, so constructed as to insure acceptance and safe delivery by common or other carriers, at the lowest rate, to point of delivery called for in the contract or purchase order.

Marking.

Exterior containers. Exterior containers shall be marked with the Federal Stock No., item identification, quantity, expiration date, name of consignee, name of consignor, and contract or purchase order number. Marking shall include the legend:

"STORE AT CONTROLLED ROOM TEMPERATURE (59° - 86° F.)."

NOTE: The item identification shall be shown on exterior containers as follows:

TETRACYCLINE HYDROCHLORIDE CAPSULES, U.S.P.,
0.25 Gram, 100s

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