A-A-51936A
30 May 1989
SUPERSEDING
A-A-0051936(DM)
10 April 1987

COMMERCIAL ITEM DESCRIPTION

ALLOPURINOL TABLETS, USP

The General Services Administration has authorized the use of this Commercial Item Description as a replacement for Military Specification MIL-A-37487 which is cancelled.

Salient characteristics:

Shall be Allopurinol Tablets, USP.

Shall be in accordance with the requirements of the USP.

Material used in manufacture of the tablets shall be of pharmaceutical grade.

For Strength I, shall contain 100 mg of Allopurinol per tablet, within the applicable assay limits.

For Strength II, shall contain 300 mg of Allopurinol per tablet, within the applicable assay limits.

Each tablet shall be marked/embossed with a product identification code to identify the active ingredient(s), strength, and manufacturer. The materials used in marking/embossing the dosage forms shall be acceptable to the FDA. The actual code used for the product shall be left to the discretion of the supplier.

Workmanship. The tablets shall be free from defects which detract from their appearance or impair their serviceability.

AMSC/NA FSC 6505

Unit.

For Size 1 - Bottle (BT). One bottle containing 30 tablets, as specified, constitutes one unit. Bottle shall be supplied with a child-resistant safety closure and shall have a minimum height of 51 mm base to shoulder.

For Size 2 - Bottle (BT). One bottle containing 100 tablets, as specified, constitutes one unit.

For Size 3 - Bottle (BT). One bottle containing 500 tablets, as specified, constitutes one unit.

For Size 4 - Bottle (BT). One bottle containing 1000 tablets, as specified, constitutes one unit.

For Size 5 - Package (PG). One package containing 100 tablets individually sealed in commercially available blister or strip packaging, as specified, constitutes one unit.

Contractor certification. The contractor shall certify that the product offered meets the salient characteristics of this description and conforms to the producers' own drawings, specifications, standards, and quality assurance practices. The Government reserves the right to require proof of such conformance prior to first delivery and thereafter as may be otherwise provided for under the provisions of the contract.

Regulatory requirements.

Federal Food, Drug and Cosmetic Act. If the product covered by this document has been determined by the U.S. Food and Drug Administration to be under its jurisdiction, the offeror/contractor shall comply, and be responsible for compliance by its subcontractors/suppliers, with the requirements of the Federal Food, Drug and Cosmetic Act, as amended, and regulations promulgated thereunder. In addition, the offeror/contractor shall comply, and be responsible for compliance by its subcontractors/suppliers, with the requirements of all other applicable Federal, State, and local statutes, ordinances, and regulations.

Recovered materials. The offeror/contractor is encouraged to use recovered material in accordance with Public Law 94-580, as amended, and as implemented by Federal Acquisition Regulation Subpart 23.4 to the maximum extent practical.

Preservation, packaging, packing, labeling, and marking. Unless otherwise specified, preservation, packaging, and packing shall be to a degree of protection to preclude damage to containers and/or contents thereof under normal shipping conditions, handling, etc., involving shipment from the supply source to the receiving activity, plus reshipment from receiving activity, and shall conform to applicable carrier's rules and regulations. Intermediate and exterior package quantities and labeling and marking shall be as specified in the contract and/or order.

NOTE: The following National Stock Numbers are covered by this document:

NSN	ITEM IDENTIFICATION	STRENGTH	SIZE
6505-00-998-4381	ALLOPURINOL TABLETS, USP 100 mg, 100s.	I	2
6505-01-004-3952	ALLOPURINOL TABLETS, USP 300 mg, 100s.	II	2
6505-01-006-5974	ALLOPURINOL TABLETS, USP 300 mg, 30s.	II	1
6505-01-044-2579	ALLOPURINOL TABLETS, USP 100 mg, 1000s.	I	4
6505-01-044-9395	ALLOPURINOL TABLETS, USP 300 mg, Individually Sealed, 100s.	II	5
6505-01-090-9251	ALLOPURINOL TABLETS, USP 300 mg, 500s.	II	3

Ordering data. Expiration dating required, intermediate/exterior package quantities, labeling, and marking must be specified in the contract and/or order.

For Size 4 - Package (PG) - One package containing 10[illegible]

MILITARY INTERESTS:

Custodians:

Army - MD
Navy - MS
Air Force - 03

PREPARING ACTIVITY:

DoD - MB

Agent:

DLA-DM

CIVIL AGENCY COORDINATING ACTIVITIES:

VA-OSS
PHS
FDA-MPQAS

PGC 05758

Project No. 6505-2808