DEFENSE MEDICAL PURCHASE DESCRIPTION	NUMBER 8	DATE 20 January 1972

FEDERAL STOCK NO.	ITEM IDENTIFICATION	UNIT
6505-116-7750	DIGOXIN TABLETS, USP, 0.25 mg, 100s	Bottle

Shall be Digoxin Tablets, U.S.P., and shall be in accordance with all applicable requirements of Federal Standard Fed. Std. No. 1l0a, dated 30 October 1966, and Amendment-1, dated 25 March 1970, and as specified herein:

S2. Classification. Shall be type I, class 1.

S5.2 The following additional requirements and tests are added to this paragraph:

Shall be tablets containing 0.25 mg of Digoxin per tablet, within the applicable assay limits for the tablets.

S6.4.2 Color. Uncoated tablets shall be white.

PREPARATION FOR DELIVERY

Shall be in accordance with all applicable requirements of Interim Federal Specification PPP-C-00186a, dated 15 May 1969, and Amendment-1, dated 27 October 1969, and as specified herein:

Immediate containers. Shall comply with the following classification:

GROUP A	CLASS 1	TYPE e	STYLE 1	GRADE 1
CLOSURE A, B, or F				SEAL A or B

Labeling. Labeling shall be in accordance with the requirements of the Federal Food, Drug, and Cosmetic Act, and shall include the information required below:

Immediate containers. Each immediate container label shall bear the following information. However, the information is not required to appear in the sequence indicated:

(a) the item name designated as "DIGOXIN TABLETS, U.S.P."

(b) the quantity of active ingredient per tablet designated as "0.25 mg"

(c) the phrase "100 tablets" or a similar phrase

(See additional label information on page 2)