DEFENSE MEDICAL PURCHASE DESCRIPTION	NUMBER 5	DATE 22 June 1972

FEDERAL STOCK NO.	ITEM IDENTIFICATION	UNIT
6505-116-9660	DIMENHYDRINATE TABLETS, USP, 50 mg, 100s	Bottle

Shall be Dimenhydrinate Tablets, U.S.P., and shall be in accordance with all applicable requirements of Federal Standard No. 1l0a, dated 30 October 1966, and Amendment-1, dated 2[illegible] March 1970, and as specified herein:

3.2. Classification. [illegible] shall be type I, class 1.

3.2.2 The following additional requirements are added to this paragraph:

[illegible] shall be tablets containing 50 mg of Dimenhydrinate, within the applicable assay limits for the [illegible].

3.6.3 Uncoated tablets. Tablets shall be yellow in color.

3.6.7 Scoring. Tablets shall be scored.

PREPARATION FOR DELIVERY

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

Immediate containers. Shall comply with the following classification:

[illegible] UP A	[illegible] 1	TYPE [illegible]	STYLE 2 GRADE 1 or 2
CLOSURE A, [illegible], or [illegible]			SEAL A or B

Labeling. Labels 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 [illegible] required to appear in the sequence indicated:

(a) The item name designated as "DIMENHYDRINATE TABLETS, U.S.P."

(b) The quantity of active ingredient designated as "50 mg"

(See additional label information on page 2)

DPSC FORM 2087 OCT 65

REPLACES DMAC FORM [illegible]/20/11 MAR 64, WHICH WILL BE USED UNTIL DEPLETED.