Revised

DEFENSE MEDICAL PURCHASE DESCRIPTION | NUMBER 4 | DATE 11 June 1969

FEDERAL STOCK NO.	ITEM IDENTIFICATION	POTENCY	UNIT
6505-527-6885	PROBENECID TABLETS, USP, 0.5 Gm, 100s	60 Months	Bottle

Shall be Probenecid Tablets, U.S.P., and shall be in accordance with all applicable requirements of Federal Standard Fed. Std. No. 1L0a, dated 30 October 1966, together with the options and additions stated 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 0.50 Gm of Probenecid per tablet, within the applicable assay limits for the tablets.

S6.4.2 Color. Uncoated tablets shall be white.

S6.4.7 Scoring. Tablets shall be scored.

PREPARATION FOR DELIVERY

Shall be in accordance with all applicable requirements of Federal Specification PPP-C-186, dated 11 December 1961, together with the deletions or additions as indicated herein:

Immediate containers. Shall comply with the following classification:

GROUP A CLASS 1 TYPE e STYLE 2 GRADE 1

CLOSURE 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 identification designated as "PROBENECID TABLETS, U.S.P."

(b) the quantity of active ingredient designated as "0.5 Gm"

(See additional label information on page 2)

DPSC FORM 2087 REPLACES DMSC FORM 7-4180/1, MAR 64, WHICH WILL BE USED UNTIL DEPLETED