6505-290-0022 (P. D. No. 7)

All other ingredients shall comply with S5.1.

S6.4.2 Color. Uncoated tablets shall be white.

S6.4.7 Scoring. Tablets shall be scored.

If samples of the tablets are submitted to Defense Personnel Support Center, a quantity of not less than 5 grams of each lot of Reserpine Powder, U.S.P., shall also be submitted with the finished product, to the Technical Operations Division, Directorate of Medical Materiel, Defense Personnel Support Center, 2800 South 20th Street, Philadelphia, Pa. 19101, Attention: Quality Assurance Branch.

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 2	GRADE 1
CLOSURE A, B, or F			SEAL A or B for Closures A and B only.	

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 "RESERPINE TABLETS, U.S.P."

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

(c) the phrase "1000 tablets" or a similar phrase

(d) the Federal Stock No.

(e) the lot or control number

(See additional label information on page 4)