MODIFICATION NO. 1
DATE: 6 February 1973

MODIFICATION TO DEFENSE MEDICAL PURCHASE DESCRIPTION

This modification forms a part of Defense Medical Purchase Description No. 8, dated 20 February 1969, and covers the following item to the extent specified herein:

Federal Stock No.	Item Identification
6505-138-4225	PROPYLTHIOURACIL TABLETS, USP, 50 mg, 100s

Page 1:

Preceding the paragraph "S6.4.2 Color" - Insert the following new paragraph:

"If (i) the quality assurance representative submits samples to the laboratory at the Defense Personnel Support Center, or (ii) the bidder or offeror is required by the terms of the Schedule or otherwise to submit samples to said laboratory, a quantity of 15 grams of Propylthiouracil powder used in the manufacture of each lot of Propylthiouracil Tablets, U.S.P., accompanied by a certificate of analysis listing all test results, shall 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."

Under "PREPARATION FOR DELIVERY" - Delete the first paragraph in its entirety and substitute:

"Shall be in accordance with all applicable requirements of Interim Federal Specification PPP-C-001166a, dated 15 May 1969, and Amendment-1, dated 27 October 1969, and as specified herein:"

Pages 2 and 3:

Packaging, Packing, and Marking - Delete and substitute:

"Packaging and Packing.

"Unit of issue. One bottle containing 100 tablets, as specified, constitutes one unit of issue.

"Unit package. At the option of the contractor, each unit shall be packaged as specified in 5.2.5 of PPP-C-001166a.

"Procedure code. Procedure code No. 2 as specified in Table I of PPP-C-001166a shall apply.