

COMPETITIVE PROBLEMS IN THE DRUG INDUSTRY 10203

DEFENSE MEDICAL PURCHASE DESCRIPTION		DATE
		1 November 1970
FEDERAL STOCK NO.	ITEM IDENTIFICATION	UNIT
6505-1F1-7678	CLOMID TABLETS, 50 mg, 30s	Package

Shall be The Wm. S. Merrell Company's "CLOMID TABLETS" and, in addition, shall comply with the following:

Each tablet shall contain 50 mg of Clomiphene Citrate.

Shall be supplied 30 tablets as specified herein.

Not more than 6 months shall have elapsed from the date of manufacture to the date of delivery to the Government.

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 (foil packet). Shall comply with the following:

Each tablet shall be packaged in a hermetically sealed, aluminum foil packet.

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 (packet) shall be permanently and legibly marked with the lot or control number.

Unit packages. Each unit package shall bear the following information. However, the information is not required to appear in the sequence indicated:

- (a) labeling information in accordance with commercial practice
- (b) the item name designated as "CLOMIPHENE CITRATE TABLETS"

(See additional labeling information on page 2)

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DPSC FORM 2087

REPLACES DMSC FORM T-4120/11, MAR 64, WHICH WILL BE USED UNTIL DEPLETED