10360 COMPETITIVE PROBLEMS IN THE DRUG INDUSTRY

FEDERAL STO	CK NO.		ITEM IDENT	IFICATION .	THE REST PORT	UNIT
1			No. 1. Sept.	5"		
6505-126	-9375	MEPERIDINE	HYDROCHLORIDE 1	TABLETS, USP,	50 mg, 100s	Bottle
	· · · · · · · · · · · · · · · · · ·					
		10 to 10 miles				

Shall be Meperidine Hydrochloride Tablets, U.S.P., and shall be in accordance with all applicable requirements of Federal Standard Fed. Std. No. 140a, dated 30 October 1966, and Amendment-1, dated 25 March 1970, and as specified herein:

S2. Classification. Shall be type I, class 1.

Shall be suitable for use as an oral narcotic analgesic.

S5.2 The following additional requirements and tests are added to the paragraph:

Shall be tablets containing 50 mg of Meperidine Hydrochloride per tablet, within the applicable assay limits for the tablets.

S6.4.2 Color. Uncoated tablets shall be white.

PREPARATION FOR DELIVERY

Shall be in accordance with all applicable requirements of Interim Federal Specification PPP-C-00186s, 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

CLOSURF A, 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 name designated as "MEPERIDINE HYDROCHLORIDE TABLETS, U.S.P."
- (b) the quantity per tablet designated as "50 mg"
- "(c) the phrase "100 tablets" or a similar phrase (See additional label information on page 2)

SSC-1

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DPSC FORM 2087

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

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