COMPETITIVE PROBLEMS IN THE DRUG INDUSTRY

DEFENSE	MEDICAL PURCHASE DESCRIPTION	NUMBER 3 DATE	June 1971
FEDERAL STOCK NO.	ITEM IDENTIFICATION		UNIT
6505-890-2010 and 6505-926-9026	PROMETHAZINE HYDROCHLORIDE, CHLOROFORI EXTRACT, AND POTASSIUM GUAIACOLSUL	M, IPECAC FLUID- FONATE SYRUP	Bottle

1. SCOPE

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1.1 This specification ndicated for the appropr	covers the following items in the quantity per souther iate Federal Stock No. (FSN) and Item Identification:
FSN	Item Identification
6505-890-2010	PROMETHAZINE HYDROCHLORIDE, CHLOROFORM, IPECAC FLUIDEXTRACT, AND POTASSIUM GUAIACOLSULFONATE SYRUP, 1 gal (3.78 liters)
6505-926-9026	PROMETHAZINE HYDROCHLORIDE, CHLOROFORM, IPECAC FLUIDEXTRACT, AND POTASSIUM QUAIACOLSULFONATE SYRUP, 4 fl oz (118 cc)

2. APPLICABLE DOCUMENTS .

2.1 Specifications and standards. Unless otherwise indicated, the issue in effect on date of invitation for bids or request for proposals of the specifications and standards, referenced in the body of this specification shall apply to the extent specified herein. These documents may be obtained as circcted by the contraction. by the contracting officer.

3. REQUIREMENTS

- 3.1 Material. Shall be a clear syrupy liquid. Each 5 oc of the preparation shall contain 5 mg promethazine hydrochloride, 0.17 minims ipecac fluidextract, 44 mg potassium guaiacolsulfonate, 0.25 minims chloroform, 60 mg citric acid (anhydrous), 197 mg sodium citrate, in a syrup base, and 7 percent alcohol.
- 3.2 Description. The finished preparation shall be green in color and have a pleasant taste and pleasing odor. Shall be suitable as an antihistamini expectorant for symptomatic cough control in adults, infants, and children.
- 3.3 Additives. The finished preparation shall contain switable preparatives. The coloring agent(s) or other ingredients, if such are using and the preservatives in the product shall be in amounts as approved by the Federal Food and Drug Administration (F.D.A.) which will make the preparation attractive and palmable, and will properly preserve the product.

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