

COMPETITIVE PROBLEMS IN THE DRUG INDUSTRY 10393

6505-926-9055 (P. D. No. 3)

5. PREPARATION FOR DELIVERY

5.1 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:

5.1.1 Immediate containers. Shall comply with the following classification:

GROUP A	CLASS 1	TYPE e	STYLE 1	GRADE 1
CLOSURE A				SEAL A

5.2 Labeling. Labeling shall be in accordance with the requirements of the Federal Food, Drug, and Cosmetic Act, and shall include the information required below:

5.2.1 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
"ACETAMINOPHEN ELIXIR, N.F."
- (b) the quantity of active ingredient per 5 cc (1 teaspoonful)
designated as "120 mg"
- (c) the quantity of contents designated as
"1 gal (3.78 liters)"
- (d) the Federal Stock No.
- (e) the lot or control number
- (f) the name and address of the manufacturer. When
the manufacturer is not the contractor, the name
and address of the contractor shall also appear.
When both names are placed on the label, the
following designations shall precede the names:
"MFR" for the manufacturer and
"CONTR" for the contractor.
- (g) the date of manufacture
- (h) the statement "KEEP FROM FREEZING."