6505-104-8672 (P. D. No. 1)

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

(a) the item name designated as "MEPROBAMATE TABLETS, U.S.P."

(b) the quantity of active ingredient designated as "0.4 Gram" or "400 mg"

Note: The official abbreviation "g." may be used in lieu of the word "gram."

(c) the Federal Stock No.

(d) the lot or control number

(e) the date of manufacture

(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 statement "Caution: Federal law prohibits dispensing without prescription."

(h) the following statements or similar statements:

1. Multiple dispensing package.
2. This package not for household use.

(i) the usual dosage

(j) all labeling information and the controlled substance schedule symbol as required by the Bureau of Narcotics and Dangerous Drugs regulations

(k) the unit of issue designated as

"1 BOX
(1 roll of 25 tablets)"

The parenthetical phrase shall appear in smaller characters than the unit of issue designation.