

specially denatured spirits no longer authorized by this part may—

(a) Continue to supply or use those stocks in accordance with existing permits until the stocks are exhausted;

(b) Use up those stocks in any manufacturing process approved by the Chief, Chemical Branch, pursuant to an application filed with him on ATF Form 5150.19, Formula for Articles made with Specially Denatured Alcohol and Rum;

(c) On approval of an application, filed with the regional director (compliance) and approved by him, destroy those stocks under whatever supervision the regional director (compliance) requires; or

(d) Otherwise dispose of those stocks in a manner satisfactory to the Director, pursuant to approval of an application (to be filed with the regional director (compliance) for transmittal to the Director).

#### § 21.4 Related regulations.

The procedural and substantive requirements relative to the production of denatured alcohol and specially denatured rum are prescribed in Part 19 of this chapter, and those relative to the distribution and use of denatured alcohol and specially denatured rum are prescribed in Part 20 of this chapter.

[T.D. ATF-133, 48 FR 24673, June 2, 1983, as amended by T.D. ATF-199, 50 FR 9183 Mar. 6, 1985]

#### § 21.5 Denatured spirits for export.

Spirits may be denatured in accordance with formulas prescribed by the government of a foreign country to which the denatured spirits will be exported. However, the denaturer must first apply for and obtain written permission from the Director. The application shall be submitted to the Director and shall contain the following information:

(a) A complete list of ingredients for the spirits to be denatured.

(b) The exact amount of each ingredient to be used in denaturing the spirits.

(c) A copy (accompanied by an English translation as necessary) of the law or regulations of the foreign

country to which the denatured spirits will be exported, specifying the denatured spirits formulation prescribed by that country.

#### § 21.6 Incorporations by reference.

(a) “The United States Pharmacopoeia (Twentieth Revision, Official from July, 1980) and the National Formulary (Fifteenth Edition, Official from July 1, 1980)” published together as “The USP and NF Compendia,” are incorporated by reference in this part. This incorporation by reference was approved by the Director of the Federal Register. The publication may be inspected at the Office of the Federal Register, 800 North Capitol Street, NW., suite 700, Washington, DC, and is available from the United States Pharmacopoeia Convention, Inc., 12601 Twinbrook Parkway, Rockville, Maryland 20852.

(b) Material from Parts 23, 25, and 29 of the 1980 Annual Book of ASTM Standards is incorporated by reference in this part. This incorporation by reference was approved by the Director of the Federal Register. These publications may be inspected at the Office of the Federal Register, 800 North Capitol Street, NW., suite 700, Washington, DC, and are available from the American Society for Testing and Materials, 1916 Race Street, Philadelphia, Pennsylvania 19103.

(c) Material from the “Official Methods of Analysis of the Association of Official Analytical Chemists (13th Edition 1980)” (AOAC) is incorporated by reference in this part. This incorporation by reference was approved by the Director of the Federal Register. This publication may be inspected at the Office of the Federal Register, 800 North Capitol Street, NW., suite 700, Washington, DC, and is available from the Association of Official Analytical Chemists, 11 North 19th Street, Suite 210, Arlington, Virginia 22209.

(Pub. L. 89-554, 80 Stat. 383 as amended (5 U.S.C. 552(a)))

### Subpart B—Definitions

#### § 21.11 Meaning of terms.

When used in this part and in forms prescribed under this part, unless the