Internal pneumatic line-up clamps for welding transmission line pipe
Line-travelling coating and wrapping for pipes and tubes
Mechanical instruments, n.e.s., for measurement, transmission, or control of temperature, pressure, or other variables of liquids or gases
Metering and mixing, n.e.s.
Pipe-line cleaning

c. A new interpretation 31 is added to § 399.2, Supplement No. 1 to read as follows:

Interpretation 31: Petroleum and Natural Gas Transportation and Refining Equipment

The following is an illustrative list of petroleum and natural gas transportation and refining equipment subject to validated license control for export to the USSR, Estonia, Latvia, Lithuania, and Afghanistan. This list is illustrative only. It does not include all commodities covered by CCL entry 6198.

Reactors (Crackers)
Separators
Extraction Columns
Exchangers
Agitators
Pipes
Valves
Filtration Units
Electronic Controls
Air or Gas Compressors
Gas or Liquid Supply Meters
Gas Turbine Engines
Internal pneumatic line-up clamps for welding transmission line pipe
Line-travelling coating and wrapping equipment for pipes and tubes
Instruments for measurement, transmission, or control of temperatures, pressure, or other variables of liquids or gases
Metering and mixing equipment
Pipe-line cleaning equipment
Specialized land-based and seaborne petroleum and natural gas transportation equipment

(Sections 4, 6, 13, 15, and 21, Pub. L. 96–72, 93 Stat. 503, 50 U.S.C. app. 2401 *et seq.*; Executive Order No. 12214 (45 FR 29783, May 6, 1980); Department Organization Order 10–3 (45 FR 6141, January 25, 1980); International Trade Administration Organization and Function Orders 41–1 (45 FR 11862, February 22, 1980) and 41–4 (45 FR 65003, October 1, 1980))

Dated: December 30, 1981.

Bohdan Denysyk,

Deputy Assistant Secretary for Export Administration.

[FR Doc. 81–37468 Filed 12–30–81; 4:05 pm]

BILLING CODE 3510–25–M

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15 CFR Part 390

General Orders; Suspension of All Licensing for Exports to the U.S.S.R.

AGENCY: Office of Export Administration, International Trade Administration, Commerce.

ACTION: Interim rule.

SUMMARY: This issuance announces a General Order of the Department of Commerce suspending the processing of all licensing for exports to the U.S.S.R. to further U.S. foreign policy objectives in light of the Soviet Union's heavy and direct responsibility for the repression in Poland. This issuance provides notice that no new licenses or other authorizations for export to the U.S.S.R. will be issued by the Department of Commerce. Outstanding validated licenses and authorizations may be reviewed to determine whether suspension or revocation may be necessary to be consistent with the objectives of this action.

DATES: These rules are effective December 30, 1981. Comments must be received by the Department by March 1, 1982. However, these regulations may be revised before the close of the comment period.

ADDRESS: Written comments (six copies when possible) should be sent to: Richard J. Isadore, Director, Operations Division, Office of Export Administration, U.S. Department of Commerce, P.O. Box 273, Washington, D.C. 20044.

FOR FURTHER INFORMATION CONTACT: Mr. Archie Andrews, Director, Exporters' Service Staff, Office of Export Administration, Department of Commerce, Washington, D.C. 20230 (Telephone: (202) 377–5247 or 377–4811).

SUPPLEMENTARY INFORMATION:

Regulatory Changes

At the direction of the President, the Department of Commerce has suspended the processing of applications for validated licenses and other authorizations for export to the U.S.S.R. No changes in the General License provisions of the Export Administration Regulations are being made by this announcement.

Pursuant to section 6 of the Export Administration Act of 1979 and following consultation with the Department of State, it has been determined that this rule is necessary to further significantly the foreign policy of the United States. Appropriate persons in industry and the Congress have been consulted, and the Departments of Commerce and State have considered the criteria set forth in section 6(b) of the Act.

Pursuant to section 4(c), it has been determined that, notwithstanding foreign availability, failure to take this action would be detrimental to the foreign policy of the United States.

Pursuant to section 6(d), it has been determined that there are no feasible alternative means of achieving the purpose of this action. As provided in section 6(g), efforts are being made to obtain cooperation of countries that produce comparable items.

Accordingly, exporters are placed on notice that the processing of validated licenses and other requests for export or reexport authorization for shipments to the U.S.S.R. has been suspended.

This action affects validated export license applications, parts and components applications, special bulk license applications (to the extent they would authorize exports to the U.S.S.R.) and reexport requests that are currently pending, as well as any new applications or requests. This will serve as notice to exporters that pending applications or requests, as well as any that may be submitted in the future, will be returned without action.

The Department may review outstanding validated licenses and other authorizations for exports to the U.S.S.R. to determine if they should be suspended or revoked as a result of the President's announcement.

Rulemaking Requirements

The Office of Export Administration has determined that:

1. Under section 13(a) of the Export Administration Act of 1979 (50 U.S.C. app. 2401 *et seq.*) (Supp. III 1979) ("the Act"), this rule is exempt from the public participation in rulemaking procedures of the Administrative Procedure Act.

However, because of the importance of the issues raised by these regulations and the intent of Congress set forth in section 13(b) of the Act, these regulations are issued in interim form and comments will be considered in developing any final regulations. These regulations may be revised before the end of the comment period. Accordingly, interested persons who desire to comment are encouraged to do so at the earliest possible time to permit the fullest consideration of their views.

2. This rule does not impose a burden under the Paperwork Reduction Act of 1980, 44 U.S.C. 3501 *et seq.*

3. This rule is not subject to the requirements of the Regulatory Flexibility Act, 5 U.S.C. 3501 *et seq.*

4. This rule is exempt from the requirements of Executive Order 12291 (46 FR 13193, February 19, 1981), "Federal Regulation" because it relates

to a foreign affairs function of the United States.

PART 390—GENERAL ORDERS

The period for submission of comments will close on March 1, 1982. Comments received after the close of the comment period cannot be assured consideration in the development of the final regulations. Public comments that are accompanied by a request that part or all of the material be treated confidentially because of its business proprietary nature, or for any other reason, will not be accepted. Such comments and materials will be returned to the submitter and will not be considered in the development of the final regulations. All public comments to be considered in any revision to these regulations will be a matter of public record and will be available for public inspection and copying. In the interest of accuracy and completeness, comments in written form are preferred. If oral comments are received, they must be followed by written memoranda which will also be a matter of public record and will be available for public review. Therefore, the Export Administration Regulations (15 CFR 368, *et seq.*) are amended by adding a new § 390.8 to read:

§ 390.8 General order suspending all licensing for exports to the U.S.S.R.

Effective December 30, 1981, the processing of all applications for validated licenses, reexport authorizations, and parts and components authorizations for shipment of any commodities or transfer of any technical data to the Union of Soviet Socialist Republics (U.S.S.R.) has been suspended. Furthermore, outstanding validated licenses and authorizations to export may be reviewed to determine whether suspension or revocation is necessary.

(Sections 4, 6, 13, 15, and 21, Pub. L. 96–72, 93 Stat. 503, 50 U.S.C. app. 2401 *et seq.*; Executive Order No. 12214 (45 FR 29783, May 6, 1980); Department Organization Order 10–3 (45 FR 6141, January 25, 1980); International Trade Administration Organization and Function Orders 41–1 (45 FR 11862, February 22, 1980) and 41–4 (45 FR 65003, October 1, 1980)

Dated: December 30, 1981.

Bohdan Denysyk,

Deputy Assistant Secretary for Export Administration.

[FR Doc. 81-37346 Filed 12-30-81; 4:05 pm]

BILLING CODE 3510-25-M

DEPARTMENT OF THE TREASURY

Office of Foreign Assets Control

15 CFR Part 535

Notice to Account Parties With Standby Letters of Credit in Favor of an Iranian Entity

Cross Reference: For a notice that calls to the attention of account parties who have standby letters of credit in favor of an Iranian entity, including instruments against which no demand has been made, relevant factors concerning the purposes of certain sections of the Iranian Assets Control Regulations (15 CFR 535) and the approach of the January 19, 1982, deadline for filing claims with the Iran-United States Claims Tribunal, see FR Doc. 81-37480 appearing in the notices section of this issue.

BILLING CODE 4810-25-M

DEPARTMENT OF LABOR

Office of the Secretary

20 CFR Chs. I, V, VI

29 CFR Subtitle A and Chs. V and XVII

41 CFR Chs. 50 and 60

Display of Office of Management and Budget Control Numbers for Recordkeeping Requirements

AGENCY: Office of the Secretary, Labor.

ACTION: Technical amendments.

SUMMARY: This document amends certain Department of Labor regulations to include OMB control numbers approved prior to December 31, 1981, at the places in the regulations where current information collection requirements are described. It includes items listed in the December 29, 1981, **Federal Register** (46 FR 62844) and thereby updates that list. It also removes obsolete references to previous GAO and OMB approvals of information collection requirements. OMB control numbers for other DOL recordkeeping requirements will be published in a subsequent issue of the **Federal Register**.

EFFECTIVE DATE: January 5, 1982.

FOR FURTHER INFORMATION CONTACT: Paul Larson, Director, Office of Management Reports and Analysis, Directorate of Management Policy and Systems, Room S–5526, Frances Perkins Building, U.S. Department of Labor, 200 Constitution Avenue, NW, Washington, D.C. 20210, Telephone 202–523–6331.

SUPPLEMENTARY INFORMATION:

Paperwork Reduction Act

The information collection requirements contained in the regulatory sections listed below have been approved by the Office of Management and Budget under the provisions of the Paperwork Reduction Act of 1980 (Pub. L. 96–511) and assigned the control numbers contained in the listing.

Guide to Agency Abbreviation Code

BLS— Bureau of Labor Statistics
ESA—Employment Standards Administration
ETA—Employment and Training Administration

Text of Amendments

Following the text of each paragraph cited in the second column of the table, add parenthetically the corresponding OMB number listed in the third column:

Agency	Regulatory citation	OMB No.
BLS	29 CFR 1904.2, 1904.4, and 1904.5.	1220-0029
	29 CFR 1904.5	1220-0045
ESA	20 CFR 702.111	1215-0117
	29 CFR 575.3	1215-0120
	20 CFR 10.145	1215-0115
	20 CFR 702.148	1215-0118
	29 CFR 549.1	1215-0122
	20 CFR 725.531	1215-0124
	29 CFR 516 Subpart A and B	1215-0017
	29 CFR 3.4(b), 4.6(g), 5.5(a)(3), and 5.5(e).	1215-0017
	41 CFR 50-201.501	1215-0017
	41 CFR Parts 60-1, 60-2, 60-4, 60-20, 60-30, 60-40, 60-50, 60-60, 60-250, and 60-741.	1215-0072
	41 CFR Part 60-3	3046-0017
	29 CFR 547.1	1215-0019
	29 CFR 570.35(b)(3)(vi)	1215-0121
	29 CFR 4.6(b)(2)(1)	1215-0126
	29 CFR 4.6(1)(i)	1215-0127
	29 CFR 40.51, 40.52, 40.53	1215-0128
	20 CFR 10.410(a)	1215-0133
ETA	20 CFR 653.103, 653.106, 653.107, 653.108, 653.109, 653.111, 653.112, 653.501, 653.503.	1205-0039
	20 CFR 658.410 and 658.413	1205-0039
	29 CFR 56.63	1205-0175

Signed at Washington, D.C. this 31st day of December 1981.

Raymond J. Donovan,

Secretary of Labor.

[FR Doc. 81-37481 Filed 12-31-81; 3:04 pm]

BILLING CODE 4510-23-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 173

[Docket No. 80F-0482]

Secondary Direct Food Additives Permitted in Food for Human Consumption; Ethyl Acetate

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) amends the food additive regulations to provide for the safe use of ethyl acetate as a solvent in the decaffeination of coffee. This action is based on data contained in a petition filed by the General Foods Corp.

DATES: Effective January 5, 1982; objections by February 4, 1982.

ADDRESS: Written objections to the Dockets Management Branch (HFA–305), Food and Drug Administration, Rm. 4–62, 5600 Fishers Lane, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: James B. Lamb, Bureau of Foods (HFF–334), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–472–5690.

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of January 16, 1981 (46 FR 3983), FDA announced that a petition (FAP 8A3353) had been filed by General Foods Corp., White Plains, NY 10625, proposing to amend Part 173—Secondary Direct Food Additives Permitted in Food for Human Consumption (21 CFR Part 173) to provide for the safe use of ethyl acetate as a solvent for the decaffeination of coffee. In response to the notice of filing, FDA received one comment. The comment and FDA's response are summarized below.

The petition asked that FDA approve ethyl acetate for use as the extraction solvent for moisturized green coffee beans. The comment expressed the belief that the petition is too narrow because it limits the extraction of caffeine to moisturized green coffee beans. According to the comment, ethyl acetate can be used to remove caffeine from whole roasted coffee beans, ground roasted coffee, or aqueous extracts of coffee beans as well as from moisturized green coffee beans. The comment further stated that the good manufacturing practices involved in producing the finished product would limit solvent residues. Thus, there is no reason to restrict the forms of coffee in which ethyl acetate may safely be used, so that manufacturers may have flexibility in their choice of decaffeination processes.

FDA has evaluated the comment, data in the petition, and other relevant material and has concluded that ethyl acetate has a wide margin of safety. Further, the data submitted in support of the petition demonstrate that ethyl acetate effectively decaffeinates coffee. In sum, the agency agrees with the comment that it is not necessary to limit the use of ethyl acetate in the decaffeination of coffee to moisturized green coffee beans, and that ethyl acetate may safely be used in accordance with good manufacturing practice as a solvent in the decaffeination of coffee. Accordingly, Part 173 is amended to provide for the safe use of ethyl acetate as set forth below.

PART 173—SECONDARY DIRECT FOOD ADDITIVES PERMITTED IN FOOD FOR HUMAN CONSUMPTION

Therefore, under the Federal Food, Drug, and Cosmetic Act (Secs. 201(s), 409, 72 Stat. 1784–1788 as amended (21 U.S.C. 321(s), 348)) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.10 (formerly 5.1; see 46 FR 26052; May 11, 1981)), Part 173 is amended in Subpart C by adding new § 173.228, to read as follows:

§ 173.228 Ethyl acetate.

Ethyl acetate (CAS Reg. No. 141–78–6) may be safely used in food in accordance with the following conditions:

(a) The additive meets the specifications of the Food Chemicals Codex,[1] (Ethyl Acetate; p. 372, 3d Ed., 1981), which are incorporated by reference.

(b) The additive is used in accordance with good manufacturing practice as a solvent in the decaffeination of coffee.

Any person who will be adversely affected by the foregoing regulation may at any time on or before February 4, 1982, submit to the Dockets Management Branch (address above), written objections thereto and may make a written request for a public hearing on the stated objections. Each objection shall be separately numbered and each numbered objection shall specify with particularity the provision of the regulation to which objection is made. Each numbered objection on which a hearing is requested shall specifically so state; failure to request a hearing for any particular objection shall constitute a waiver of the right to a hearing on that objection. Each numbered objection for which a hearing is requested shall include a detailed description and analysis of the specific factual information intended to be presented in support of the objection in the event that a hearing is held; failure to include such a description and analysis for any particular objection shall constitute a waiver of the right to a hearing on the objection. Three copies of all documents shall be submitted and shall be identified with the docket number found in brackets in the heading of this regulation. Received objections may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday.

Effective date. This regulation shall become effective January 5, 1982.

(Secs. 201(s), 409, 72 Stat. 1784–1788 as amended (21 U.S.C. 321(s), 348).)

Note.—Incorporation by reference was approved by the Director, Office of the Federal Register, September 23, 1981.

Dated: December 21, 1981.

William F. Randolph,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 82–23 Filed 1–4–82; 8:45 am]

BILLING CODE 4160–01–M

[1] Copies may be obtained from: National Academy Press, 2101 Constitution Ave. NW., Washington, D.C. 20418 or examined at the Office of the Federal Register, 1100 L St. NW., Washington, D.C. 20408.

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21 CFR Parts 510 and 522

New Animal Drugs; Change of Sponsor

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor for a canine, narcotic antagonist (naloxone hydrochloride injection) from Endo Laboratories, Inc., to Endo Pharmaceuticals, Inc. Endo Pharmaceuticals, Inc., filed a supplemental new animal drug application (NADA) providing for the change.

EFFECTIVE DATE: January 5, 1982.

FOR FURTHER INFORMATION CONTACT: John R. Markus, Bureau of Veterinary Medicine (HFV–104), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–4313.

SUPPLEMENTARY INFORMATION: Endo Pharmaceuticals, Inc., P.O. Box 363, Manati, PR 00701, filed a supplemental NADA (035–825) providing for its sponsorship of a naloxone hydrochloride injection. By letter of October 6, 1981, Endo Laboratories, Inc., the former sponsor, confirmed the transfer of sponsorship.

The supplemental NADA for the change of sponsor is approved and the regulations are amended to reflect the approval.

Under the Bureau of Veterinary Medicine's supplemental approval policy (42 FR 64367; December 23, 1977), the intracorporate transfer of an NADA is a Category I change that does not require reevaluation of the safety and effectiveness data in the parent application.