Drug Enforcement Administration, Justice

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(a) Name of the prescribing practitioner.
(b) Prescribing practitioner’s Federal and State registration numbers, with the expiration dates of these registrations.
(c) Verification that the prescribing practitioner possesses the appropriate registration to prescribe this controlled substance.
(d) Patient’s name and address.
(e) Patient’s insurance provider, if available.

[70 FR 293, Jan. 4, 2005]

REPORTS

§ 1304.31 Reports from manufacturers importing narcotic raw material.

(a) Every manufacturer which imports or manufactures from narcotic raw material (opium, poppy straw, and concentrate of poppy straw) shall submit information which accounts for the importation and for all manufacturing operations performed between importation and the production in bulk or finished marketable products, standardized in accordance with the U.S. Pharmacopeia, National Formulary or other recognized medical standards. Reports shall be signed by the authorized official and submitted quarterly on company letterhead to the Drug and Chemical Evaluation Section, Drug Enforcement Administration, on or before the 15th day of the month immediately following the period for which it is submitted. See the Table of DEA Mailing Addresses in §1321.01 of this chapter for the current mailing address.
(b) The following information shall be submitted for each type of narcotic raw material (quantities are expressed as grams of anhydrous morphine alkaloid):
(1) Beginning inventory;
(2) Gains on reweighing;
(3) Imports;
(4) Other receipts;
(5) Quantity put into process;
(6) Losses on reweighing;
(7) Other dispositions and
(8) Ending inventory.
(c) The following information shall be submitted for each narcotic raw material derivative including morphine, codeine, thebaine, oxycodone,