

**List of Subjects in 19 CFR Part 12**

Cultural property, Customs duties and inspection, Imports, Prohibited merchandise.

**Amendment to CBP Regulations**

For the reasons set forth above, part 12 of Title 19 of the Code of Federal Regulations (19 CFR part 12), is amended as set forth below:

**PART 12—SPECIAL CLASSES OF MERCHANDISE**

■ 1. The general authority citation for part 12 and the specific authority citation for § 12.104g continue to read as follows:

**Authority:** 5 U.S.C. 301; 19 U.S.C. 66, 1202 (General Note 3(i), Harmonized Tariff Schedule of the United States (HTSUS)), 1624;

\* \* \* \* \*

Sections 12.104 through 12.104i also issued under 19 U.S.C. 2612;

\* \* \* \* \*

**§ 12.104g [Amended]**

■ 2. In § 12.104g, paragraph (a), the table is amended in the entry for Colombia by adding, after the reference to “CBP Dec. 06–09”, the words “extended by CBP Dec. 11–06”.

**Alan Bersin,**

*Commissioner, U.S. Customs and Border Protection.*

Approved: March 9, 2011.

**Timothy E. Skud,**

*Deputy Assistant Secretary of the Treasury.*

[FR Doc. 2011–5879 Filed 3–14–11; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES****Food and Drug Administration****21 CFR Parts 312 and 314**

[Docket No. FDA–2011–N–0130]

**Investigational New Drug Applications and Abbreviated New Drug Applications; Technical Amendment**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule; technical amendment.

**SUMMARY:** The Food and Drug Administration (FDA) is amending its investigational new drug application (IND) regulations and abbreviated new drug application regulations to correct inaccurate cross-references to the IND regulations and the Federal Food, Drug, and Cosmetic Act (the FD&C Act). This

action is being taken to ensure accuracy and clarity in the Agency’s regulations.

**DATES:** This rule is effective March 15, 2011.

**FOR FURTHER INFORMATION CONTACT:**

Olivia A. Pritzlaff, Center for Drug Evaluation and Research, Food and Drug Administration, Bldg. 51, Rm. 6308, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002, 301–796–3506.

**SUPPLEMENTARY INFORMATION:** FDA is amending its regulation in 21 CFR 312.83 to correct an inaccurate cross-reference to other sections of the IND regulations. FDA is amending its regulation in 21 CFR 314.94 to correct an inaccurate cross-reference to a section of the FD&C Act.

Publication of this document constitutes final action on these changes under the Administrative Procedure Act (5 U.S.C. 553). FDA has determined that notice and public comment are unnecessary because this amendment to the regulations provides only technical changes to correct inaccurate cross-references to the IND regulations and the FD&C Act.

**List of Subjects***21 CFR Part 312*

Drugs, Exports, Imports, Investigations, Labeling, Medical research, Reporting and recordkeeping requirements, Safety.

*21 CFR Part 314*

Administrative practice and procedure, Confidential business information, Drugs, Reporting and recordkeeping requirements.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR parts 312 and 314 are amended as follows:

**PART 312—INVESTIGATIONAL NEW DRUG APPLICATION**

■ 1. The authority citation for 21 CFR part 312 continues to read as follows:

**Authority:** 21 U.S.C. 321, 331, 351, 352, 353, 355, 360bbb, 371; 42 U.S.C. 262.

**§ 312.83 [Amended]**

■ 2. Section 312.83 is amended by removing “312.34 and 312.35” and by adding in its place “312.305 and 312.320”.

**PART 314—APPLICATIONS FOR FDA APPROVAL TO MARKET A NEW DRUG**

■ 3. The authority citation for 21 CFR part 314 continues to read as follows:

**Authority:** 21 U.S.C. 321, 331, 351, 352, 353, 355, 356, 356a, 356b, 356c, 371, 374, 379e.

**§ 314.94 [Amended]**

■ 4. Section 314.94 is amended in paragraph (a)(8)(iv) by removing “505(j)(4)(D)” and by adding in its place “505(j)(5)(F)”.

Dated: March 9, 2011.

**Leslie Kux,**

*Acting Assistant Commissioner for Policy.*

[FR Doc. 2011–5946 Filed 3–14–11; 8:45 am]

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**DEPARTMENT OF THE TREASURY****Internal Revenue Service****26 CFR Part 301**

[TD 9516]

**RIN 1545–BG73**

**Disclosure of Return Information in Connection With Written Contracts Among the IRS, Whistleblowers, and Legal Representatives of Whistleblowers**

**AGENCY:** Internal Revenue Service (IRS), Treasury.

**ACTION:** Final regulations and removal of temporary regulations.

**SUMMARY:** This document contains final regulations relating to the disclosure of return information by an officer or employee of the Treasury Department, to a whistleblower and, if applicable, the legal representative of the whistleblower, to the extent necessary in connection with a written contract among the IRS, the whistleblower and, if applicable, the legal representative of the whistleblower, for services relating to the detection of violations of the internal revenue laws or related statutes. The final regulations will affect officers and employees of the Treasury Department who disclose return information to whistleblowers or their legal representatives in connection with written contracts among the IRS, whistleblowers and, if applicable, their legal representatives, for services relating to the detection of violations of the internal revenue laws or related statutes. The final regulations will also affect any whistleblower or legal representative of a whistleblower who receives return information in connection with a written contract among the IRS, the whistleblower and, if applicable, the legal representative of the whistleblower, for services relating to the detection of violations of the internal revenue laws or related statutes.