(1) Quarterly and annual reports of the agency.
(2) External investigations or review of agency needs and performance.
(3) Surveys, compilations, and summaries of data and information.
(4) Consumer surveys.
(5) Compliance surveys.
(6) Compliance programs, except that names of specific firms, the location of specific activities, and details about sampling numbers or sizes shall be deleted until implementation of the program is completed.
(7) Work plans prepared by Food and Drug Administration centers, field offices, and other components, except that names of specific firms, the location of specific activities, and details about sampling numbers or sizes shall be deleted until implementation of the plan is completed.
(b) The following types of reports and studies prepared by or with funds provided by the Food and Drug Administration are not available for public disclosure:
(1) Internal audits of agency needs and performance.
(2) Records relating to the internal planning and budget process.
(3) Legislative proposals or comments prior to submission to Congress.
§ 20.107 Food and Drug Administration manuals.
(a) Food and Drug Administration administrative staff manuals and instructions that affect a member of the public are available for public disclosure. An index of all such manuals is available by writing to the Division of Freedom of Information (ELEM–1029), Food and Drug Administration, 12420 Parklawn Dr., Element Bldg., Rockville, MD 20857; or by visiting the Division of Freedom of Information Public Reading Room, located in rm. 1050, at the same address. The index and all manuals created by the agency on or after November 1, 1996, will be made available through the Internet at http://www.fda.gov.
(b) Manuals relating solely to internal personnel rules and practices are not available for public disclosure except to the extent that the Commissioner determines that they should be disclosed pursuant to §20.82.
(c) All Food and Drug Administration action levels which are used to determine when the agency will take regulatory action against a violative product, limits of sensitivity and variability of analytical methods which are used in determining whether a product violates the law, and direct reference levels above which Food and Drug Administration field offices may request legal action directly to the office of the General Counsel, are available for public disclosure.
§ 20.108 Agreements between the Food and Drug Administration and other departments, agencies, and organizations.
(a) All written agreements and understandings signed by the Food and Drug Administration and other departments, agencies, and organizations are available for public disclosure.
(b) All written agreements and memoranda of understanding between FDA and any entity, including, but not limited to other departments, agencies, and organizations will be made available through the Food and Drug Administration Web site at http://www.fda.gov once finalized.
(c) Agreements and understandings signed by officials of FDA with respect to activities of the Office of Criminal Investigations are exempt from the requirements set forth in paragraph (b) of this section. Although such agreements and understandings will not be made available through the FDA Web site, these agreements will be available for disclosure in response to a request from the public after deletion of information that would disclose confidential investigative techniques or procedures, or information that would disclose guidelines for law enforcement investigations if such disclosure could reasonably be expected to risk circumvention of the law.
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