

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. 2003N-0429]

#### Prescription Drug User Fee Act III Five-Year Plan; Availability

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing the availability of an internal planning document entitled the "PDUFA III Five-Year Plan" (the plan). The plan to achieve PDUFA (Prescription Drug User Fee Act) III goals for the drug review process takes into account changes in the law under PDUFA III and projects revenue and spending in fiscal year (FY) 2003 through FY 2007.

**DATES:** Submit written or electronic comments on the plan at any time. These comments will be considered as the agency makes annual adjustments to the plan each fiscal year.

**ADDRESSES:** Submit written requests for single copies of this plan to the Office of Management and Systems, Attn: Frank Claunts (see **FOR FURTHER INFORMATION CONTACT**). Send a self-addressed adhesive label to assist that office in processing your requests. Submit written comments on the plan to the Division of Dockets Management (HFA 305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to <http://www.fda.gov/dockets/ecomments>. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the plan.

**FOR FURTHER INFORMATION CONTACT:** Frank Claunts, Office of Management and Systems (HF-20), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-4427.

#### SUPPLEMENTARY INFORMATION:

##### I. Background

FDA is announcing the availability of an internal planning document entitled "PDUFA III Five-Year Plan." PDUFA was amended and extended through the year 2007 by the Prescription Drug User Fee Amendments of 2002 (PDUFA III). PDUFA III authorizes appropriations and fees that will provide FDA with resources to sustain the drug review staff developed through FY 2002 and to achieve the even more stringent new goals through FY 2007.

The plan begins with a statement of purpose, provides background information on PDUFA along with a

summary of the new goals, and the plan documents the 10 major assumptions on which it is based. The plan summarizes individual plans of agency components with major PDUFA responsibilities and also provides a consolidated agency summary. The plan to achieve PDUFA III goals for the drug review process is based on projected revenue and spending projections through FY 2007. Appendix A of the plan is entitled the "PDUFA III Information Technology Five-Year Plan."

We (FDA) are making this plan available to interested individuals. We welcome comments, and we will consider all comments in the future as annual adjustments are made to the plan.

##### II. Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments. Submit a single copy of electronic comments or two paper copies of any mailed comments, except that individuals may submit one paper copy. Comments are to be identified with the docket number found in brackets in the heading of this document. The plan and received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

##### III. Electronic Access

Copies of this document are available on the Internet at <http://www.fda.gov/oc/pdufa3/2003plan/default.htm>.

Dated: October 7, 2003.

**Jeffrey Shuren,**

*Assistant Commissioner for Policy.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Health Resources and Services Administration

#### Agency Information Collection Activities: Proposed Collection: Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Public Law 104-13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to the Office of Management and Budget under the

Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, call the HRSA Reports Clearance Officer on (301) 443-1129.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the Agency, including whether the information shall have practical utility; (b) the accuracy of the Agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including the use of automated collection techniques or other forms of information technology.

#### Proposed Project: National Practitioner Data Bank for Adverse Information on Physicians and Other Health Care Practitioners: Regulations and Forms (OMB No. 0915-0126)—Revision

The National Practitioner Data Bank (NPDB) was established through Title IV of Pub. L. 99-660, the Health Care Quality Improvement Act of 1986, as amended. Final regulations governing the NPDB are codified at 45 CFR part 60. Responsibility for NPDB implementation and operation resides in the Bureau of Health Professions, Health Resources and Services Administration, U.S. Department of Health and Human Services (DHHS). The NPDB began operation on September 1, 1990.

The intent of Title IV of Pub. L. 99-660 is to improve the quality of health care by encouraging hospitals, State licensing boards, professional societies, and other entities providing health care services, to identify and discipline those who engage in unprofessional behavior; and to restrict the ability of incompetent physicians, dentists, and other health care practitioners to move from State to State without disclosure of the practitioner's previous damaging or incompetent performance.

The NPDB acts primarily as a flagging system; its principal purpose is to facilitate comprehensive review of practitioners' professional credentials and background. Information on medical malpractice payments, adverse licensure actions, adverse clinical privileging actions, adverse professional society actions, and Medicare/Medicaid exclusions is collected from, and disseminated to, eligible entities. It is intended that NPDB information should be considered with other relevant information in evaluating a practitioner's credentials.