rule and on the draft guidance document by January 29, 2007.

Two companies requested FDA to extend the comment period on the proposed rule by 90 days because the proposal presented complex medical and scientific issues that required the company to assemble a team of many different specialties in order to prepare their comments. Elsewhere in this issue of the **Federal Register**, FDA is reopening the comment period on the proposed rule for 30 days. Because the issues presented by the guidance document are intertwined with those presented by the proposed rule, FDA is reopening the comment period on the guidance document for the same period.

II. Electronic Access

Persons interested in obtaining a copy of the draft guidance may do so by using the Internet. To receive the draft guidance document entitled ''Class II Special Controls Document: Absorbable Hemostatic Device,'' you may either send an e-mail request to *dsmica@fda.hhs.gov* to receive an electronic copy of the document, or send a fax request to 240–276–3151 to receive a hard copy. Please use the document number 1558 to identify the guidance you are requesting.

CDRH maintains an entry on the Internet for easy access to information including text, graphics, and files that may be downloaded to a personal computer with Internet access. Updated on a regular basis, the CDRH home page includes device safety alerts, **Federal Register** reprints, information on premarket submissions (including lists of approved submissions, approved applications, and manufacturers' addresses), small manufacturer's assistance, information on video conferencing and electronic submissions, Mammography Matters, and other device-oriented information. The CDRH Web site may be accessed at *http://www.fda.gov/cdrh*. A search capability for all CDRH guidance documents are also available on the Division of Dockets Management Internet site at *http://www.fda.gov/ohrms/dockets.*

III. Request for Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments on the draft guidance document. 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. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: April 25, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–8780 Filed 5–7–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007D–0027]

Voluntary Self-Inspection of Medicated Feed Manufacturing Facilities; Draft Compliance Policy Guide; Availability; Reopening of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; reopening of comment period.

SUMMARY: The Food and Drug Administration (FDA) is reopening to June 8, 2007, the comment period for the notice of availability that appeared in the **Federal Register** of February 12, 2007 (72 FR 6572). In the notice, FDA requested comments on the draft compliance policy guide on voluntary self-inspection of medicated feed manufacturing facilities. The agency is taking this action in response to requests for an extension to allow interested persons additional time to submit comments.

DATES: Submit written and electronic comments by June 8, 2007.

ADDRESSES: Submit written comments on the draft compliance policy guide 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 documents.

FOR FURTHER INFORMATION CONTACT: Paul Bachman, Center for Veterinary Medicine (HFV–230), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9225, e-mail: *Paul.Bachman@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of February 12, 2007 (72 FR 6572), FDA published a notice of availability with a 75-day comment period to request comments on a draft compliance policy guide (CPG) on voluntary self inspection of medicated feed manufacturing facilities. The purpose of this CPG is intended to provide guidance to FDA field offices on considering, among other factors, the conduct of self-inspections when prioritizing inspections of medicated feed manufacturing facilities for compliance with Current Good Manufacturing Practices for Medicated Feeds regulations.

The agency has received a request for an extension of the comment period for the draft compliance policy guide. This request conveyed concern that the current 75-day comment period does not allow sufficient time to develop a meaningful or thoughtful response to the compliance policy guide.

FDA has considered the request and is reopening the comment period for the draft compliance policy guide until June 8, 2007. The agency believes this reopening allows adequate time for interested persons to submit comments.

II. Request for Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments on these documents. 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. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: May 1, 2007.

David Horowitz,

Assistant Commissioner for Regulatory Affairs.

[FR Doc. E7–8781 Filed 5–7–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review—Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301) 443–1129.

The following request has been submitted to OMB for review under the Paperwork Reduction Act of 1995:

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

The National Practitioner Data Bank (NPDB) was established through Title IV of Public Law (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, Department of Health and Human Services. 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.

The reporting forms and the request for information forms (query forms) are accessed, completed, and submitted to the NPDB electronically through the NPDB Web site at *http://www.npdb-hipdb.hrsa.gov.* All reporting and querying is performed through this secure Web site. Due to overlap in requirements for the Healthcare Integrity and Protection Data Bank (HIPDB), some of the NPDB's burden has been subsumed under the HIPDB.

Estimates of annualized burden are as follows:

Regulation citation	No. of respondents	Frequency of responses	Hours per response (minutes)	Total burden hours
60.6(a) Errors & Omissions	315	4	15	315
60.6(b) Revisions to Actions	109	1	30	54.5
60.7(b) Medical Malpractice Payment Reports	519	29	45	11,288.25
60.8(b) Adverse Action Reports—State Boards	0	0	0	
60.9(a)3 Adverse Action Reports—Clinical Privileges & Professional Society	480	2	45	720
Requests for Hearings by Entities	0	0	480	0
60.10(a)(1) Queries by Hospital—Practitioner Applications	0	0	0	0
60.10(a)(2) Queries by Hospitals—2 Year Cycle	5,996	213	5	106,429
60.11(a)(1) Disclosure to Hospitals	0	0	0	0
60.11(a)(2) Disclosure to Practitioners (Self-Query)	0	0	0	0
60.11(a)(3) Disclosure to Licensure Boards	87	645	5	4,676.25
60.11(a)(4) Queries by Non-Hospital Health Care Entities	7,305	322	5	196,017.5
60.11(a)(5) Queries by Plaintiffs' Attorneys	5	1	30	2.5
60.11(a)(6) Queries by Non-Hospital Health Care Entities-Peer Review	0	0	0	0
60.11(a)(7) Requests by Researchers for Aggregate Data	20	1	30	10
60.14(b) Practitioner Places a Report in Disputed Status	404	1	15	101
60.14(b) Practitioner Statement	1,415	1	45	1,061.25
60.14(b) Practitioner Requests for Secretarial Review	27	1	480	216
60.3 Entity Registration—Initial	1,447	1	60	1,447
60.3 Entity Registration—Update	13,115	1	5	1,092.92
60.11(a) Authorized Agent Designation—Initial	717	1	15	179.25
60.11(a) Authorized Agent—Update	139	1	5	11.58
60.12(c) Account Discrepancy Report	5	1	15	1.25
60.12(c) Electronic Funds Transfer Authorization	284	1	15	71
60.3 Entity Reactivation	0	1		
Total Burden Hours				323,694.25

Numbers in the table may not add up exactly due to rounding.

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: Karen Matsuoka, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: May 2, 2007.

Caroline Lewis,

Associate Administrator for Management.

[FR Doc. E7–8796 Filed 5–7–07; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office at (301) 443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995: