State Public Assistance Agencies (SPAAs).

C.F.D.A. Number: 93.647

Statutory Authority: Privacy Act of 1974, as amended by Public Law 100–503.

SUMMARY: In compliance with the Privacy Act of 1974, as amended by Public Law 100–503, the Computer Matching and Privacy Protection Act of 1988, ACF is publishing a notice of a computer matching program. The purpose of this computer match is to identify specific individuals who receive benefits from the Department of Veterans Affairs (VA) and also receive payments pursuant to various benefit programs administered by both the Department of Health and Human Services (HHS) and the Department of Agriculture. ACF will facilitate this program on behalf of SPAAs that participate in PARIS for verification of continued eligibility for public assistance. The match will utilize VA and SPAA records.

DATES: Submit written comments on or before May 14, 2012

ACF will file a report of the subject matching program with the Committee on Homeland Security and Governmental Affairs of the Senate and the Committee on Oversight and Government Reform of the House of Representatives and the Office of Information and Regulatory Affairs within the Office of Management and Budget (OMB). The dates for the matching program will be effective as indicated in ''*E. Inclusive Dates of the Matching Program''* in this notice.

ADDRESSES: Interested parties may comment on this notice by writing to the Director, Office of Financial Services, Office of Administration, 370 L'Enfant Promenade SW., Washington, DC 20047. All comments received will be available for public inspection at this address.

FOR FURTHER INFORMATION CONTACT: Director, Office of Financial Services, Office of Administration, 370 L'Enfant Promenade SW., Washington, DC 20047. Telephone: (202) 401–7237.

SUPPLEMENTARY INFORMATION: The Privacy Act of 1974, as amended by Public Law 100–503, the Computer Matching and Privacy Protection Act of 1988, (5 U.S.C. 552a), adds certain protections for individuals applying for and receiving Federal benefits. The law regulates the use of computer matching by Federal agencies when records in a system of records are matched with other Federal, State, and local government records.

Federal agencies that provide or receive records in computer matching programs must:

1. Negotiate written agreements with source agencies;

2. Provide notification to applicants and beneficiaries that their records are subject to matching;

3. Verify match findings before reducing, suspending, or terminating an individual's benefits or payments;

4. Furnish detailed reports to Congress and OMB; and

5. Establish a Data Integrity Board that must approve matching agreements.

This computer matching program meets the requirements of Public Law 100–503.

Jason Donaldson,

Deputy Assistant Secretary for Administration, ACF.

Notice of Computer Matching Program

A. Participating Agencies

VA and SPAAs.

B. Purpose of the Match

To identify specific individuals who receive benefits from the VA and also receive payments pursuant to HHS and Department of Agriculture benefit programs, and to verify their continued eligibility for such benefits. SPAAs will contact affected individuals and seek to verify the information resulting from the match before initiating any adverse actions based on the match results.

C. Authority for Conducting the Match

The authority for conducting the matching program is contained in section 402(a)(6) of the Social Security Act [42 U.S.C. 602(a)(6)].

D. Records To Be Matched

VA will disclose information from the system of records identified as Compensation, Pension, Education, and Vocational Rehabilitation and Employment Records—VA (58VA21/22/28) published at 74 FR 29275, (June 19, 2009), last amended at 75 FR 22187, (April 27, 2010). VA's disclosure of information for use in this computer match is listed as a routine use in this system of records.

VA, as the source agency, will prepare electronic files containing the names and other personal identifying data of eligible veterans receiving benefits. These records are matched electronically against SPAA files consisting of data regarding monthly Medicaid, Temporary Assistance for Needy Families, general assistance, and Supplemental Nutrition Assistance Program beneficiaries.

1. The electronic files provided by the SPAAs will contain client names and Social Security numbers (SSNs).

2. The resulting output returned to SPAAs will contain personal identifiers, including names, SSNs, employers, current work or home addresses, etc.

E. Inclusive Dates of the Matching Program

The effective date of the matching agreement and date when matching may actually begin shall be at the expiration of the 40-day review period for OMB and Congress, or 30 days after publication of the matching notice in the **Federal Register**, whichever date is later. The matching program will be in effect for 18 months from the effective date, with an option to renew for 12 additional months, unless one of the parties to the agreement advises the others by written request to terminate or modify the agreement.

[FR Doc. 2012–8901 Filed 4–12–12; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–D–0889]

Draft Guidance for Industry on New Animal Drugs and New Animal Drug Combination Products Administered in or on Medicated Feed or Drinking Water of Food-Producing Animals: Recommendations for Drug Sponsors for Voluntarily Aligning Product Use Conditions With GFI #209; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry (draft GFI #213) entitled ''New Animal Drugs and New Animal Drug Combination Products Administered in or on Medicated Feed or Drinking Water of Food-Producing Animals: Recommendations for Drug Sponsors for Voluntarily Aligning Product Use Conditions With GFI #209.'' The purpose of this document is to provide information to sponsors of certain new animal drug products who are interested in developing revised conditions of use for those products consistent with FDA's GFI #209, ''The Judicious Use of Medically Important Antimicrobial Drugs in Food-Producing Animals'' and to set timelines for stakeholders wishing to comply voluntarily with this guidance.

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance, submit either electronic or written comments on the draft guidance by July 12, 2012.

ADDRESSES: Submit written requests for single copies of the guidance to the Communications Staff (HFV–12), Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855. Send one self-addressed adhesive label to assist that office in processing your request. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

Submit electronic comments on the draft guidance to *http://www.regulations.gov*. Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: William T. Flynn, Center for Veterinary Medicine (HVF–1), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9084, *william.flynn@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

This document is related to two documents published elsewhere in this issue of the **Federal Register**, wherein FDA is announcing: (1) The availability of a final guidance entitled ''The Judicious Use of Medically Important Antimicrobial Drugs in Food-Producing Animals'' (GFI #209) and (2) the availability of a draft proposed regulation for veterinary feed directives.

FDA is announcing the availability of a draft guidance for industry entitled ''New Animal Drugs and New Animal Drug Combination Products Administered in or on Medicated Feed or Drinking Water of Food-Producing Animals: Recommendations for Drug Sponsors for Voluntarily Aligning Product Use Conditions With GFI #209'' (draft GFI #213). The audience for this draft guidance is sponsors of approved applications for new animal drug products containing medically important antimicrobial new animal drugs for use in or on medicated feed or in drinking water of food-producing animals. The purpose of this draft guidance is to provide sponsors of the affected new animal drug products with more specific information on how to supplement their approved new animal drug applications to align with FDA's recommendations in GFI #209.

Final GFI #209, published elsewhere in this edition of the **Federal Register,** discusses FDA's concerns regarding the development of antimicrobial resistance in human and animal bacterial pathogens when medically important antimicrobial drugs are used in food-producing animals in an injudicious manner. GFI #209 recommends that the use of medically important antimicrobial drugs be limited to uses in animals that are considered necessary for assuring animal health and include veterinary oversight or consultation (namely through the use of prescription or veterinary feed directive products).

FDA encourages all sponsors of new animal drug products covered by draft GFI #213 to participate in the voluntary program outlined in the draft guidance. FDA believes a voluntary approach, conducted in a cooperative and timely manner, will be a far faster and less burdensome route to achieving the common goal of more judicious use of medically important antimicrobials in animal agriculture. However, FDA also believes it is critical to see meaningful progress toward reaching this goal. Therefore, in order to ensure an orderly, equitable, and timely transition, draft GFI #213 also includes clear timelines for sponsors of affected products wishing to revise their approved applications.

II. Significance of Guidance

This level 1 draft guidance is being issued consistent with FDA's good guidance practices regulation (21 CFR 10.115). The draft guidance, when finalized, will represent the Agency's current thinking on this topic. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirements of the applicable statutes and regulations.

III. Paperwork Reduction Act of 1995

This draft guidance refers to previously approved collections of information found in FDA regulations. These collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in 21 CFR part 514 have been approved under OMB control numbers 0910–0032 and 0910–0669.

IV. Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) either electronic or written comments regarding this document. It is only necessary to send one set of comments. Identify comments 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.

V. Electronic Access

Persons with access to the Internet may obtain the draft guidance at either *http://www.fda.gov/AnimalVeterinary/GuidanceComplianceEnforcement/GuidanceforIndustry/default.htm* or *http://www.regulations.gov*.

Dated: April 5, 2012.

David Dorsey,

Acting Associate Commissioner for Policy and Planning.

[FR Doc. 2012–8845 Filed 4–11–12; 11:15 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–D–0094]

Guidance for Industry on the Judicious Use of Medically Important Antimicrobial Drugs in Food-Producing Animals; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a guidance for industry (GFI #209) entitled ''The Judicious Use of Medically Important Antimicrobial Drugs in Food-Producing Animals.'' This guidance is intended to inform the public of FDA's current thinking on the use of medically important antimicrobial drugs in animal agriculture.

DATES: Submit either electronic or written comments on Agency guidances at any time.

ADDRESSES: Submit written requests for single copies of the guidance to the Communications Staff (HFV–12), Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855. Send one self-addressed adhesive label to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.

Submit electronic comments on the guidance to *http://www.regulations.gov*. Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration,