information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, Division of Information Resource Management Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

The Department specifically requests comments 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) 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 through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: July 15, 1997.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 97–19075 Filed 7–18–97; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Agency Recordkeeping/Reporting Requirements Under Emergency Review by the Office of Management and Budget (OMB)

Title: Emergency Temporary Assistance for Needy Families Data Report.

OMB No.: New Request.

Description: This information is being collected to meet the statutory requirements of section 411 of the Social Security Act and section 116 of the Personal Responsibility and Work Opportunity Reconciliation Act of 1996. It consists of disaggregated demographic and program information that will be used to determine participation rates and other statutorily required indicators for the Temporary Assistance for Needy Families (TANF) program.

Respondents: States, Puerto Rico, Virgin Islands, Guam and the District of Columbia.

Annual Burden Estimates:

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
TANF Data Report	54	4	451	97,416

Estimated Total Annual Burden Hours: 97,416.

Additional Information: ACF is requesting that OMB grant a 180 day approval for this information collection under procedures for emergency processing by September 1, 1997. A copy of this information collection, with applicable supporting documentation, may be obtained by calling the Administration for Children and Families, Reports Clearance Officer, Robert Driscoll at (202) 401–9313.

Comments and questions about the information collection described above should be directed to the Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for ACF, Office of Management and Budget, Paperwork Reduction Project, 725 17th Street NW., Washington, DC 20503, (202) 395–7316.

Dated: July 18, 1997.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 97–19068 Filed 7–18–97; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97N–0215]

Babineaux's Veterinary Products, Inc., et al.; Withdrawal of Approval of NADA's

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is withdrawing approval of one new animal drug application (NADA) held by Babineaux's Veterinary Products, Inc., and two NADA's held by Schein Pharmaceutical, Inc. / Steris Laboratories, Inc. The sponsors requested voluntary withdrawal of approval of the NADA's because the products are no longer being marketed. In a final rule published elsewhere in this issue of the **Federal Register**, FDA is amending the regulations by removing those portions which reflect approval of these NADA's.

EFFECTIVE DATE: July 31, 1997

FOR FURTHER INFORMATION CONTACT: Mohammad I. Sharar, Center for Veterinary Medicine (HFV–216), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1722.

SUPPLEMENTARY INFORMATION:

Babineaux's Veterinary Products, Inc., 6425 Airline Hwy., Metairie, LA 70003, is the sponsor of NADA 46–147 Dirocide (diethylcarbamazine citrate) Syrup. Schein Pharmaceutical, Inc. / Steris Laboratories, Inc., 620 North 51st Ave., Phoenix, AZ 85043–4705, is the sponsor of NADA 48–391 phenylbutazone injection and NADA 49–183 oxytocin injection. The sponsors requested withdrawal of approval of the NADA's under 21 CFR 514.115(d) because the products are no longer being marketed.

Therefore, under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.10) and redelegated to the Center for Veterinary Medicine (21 CFR 5.84), and in accordance with § 514.115 *Withdrawal of approval of applications* (21 CFR 514.115), notice is given that approval of NADA's 46–147, 48–391, and 49–183 and all supplements and amendments thereto is hereby withdrawn, effective July 31, 1997.

In a final rule published elsewhere in this issue of the **Federal Register**, FDA is amending 21 CFR 510.600, 520.622b, 522.1680, and 522.1720 to reflect