practical utility; (ii) the accuracy of the agency's estimate of the burden of the proposed collection of information; (iii) ways to enhance the quality, utility, and clarity of the information to be collected; and (iv) ways to minimize the burden of the collection of information on respondents, including the use of automated collection techniques for other forms of information technology. Send comments to Anne O'Connor, Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, Georgia 30333. Written comments should be received within 60 days of this notice.

Proposed Project

Survey of User Satisfaction with National Health Care Survey Data—New—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC). This Survey of User Satisfaction with National Health Care Survey Data is needed to provide current information on the use and usefulness of the variety of data products describing health care delivery systems in the United States. The National Health Care Survey comprises several component surveys: National Hospital Discharge Survey, National Nursing Home Survey, National Home and Hospice Care Survey, National Ambulatory Medical Care Survey, National Hospital Ambulatory Medical Care Survey and occasional other similar surveys when funded, such as the National Health Provider Inventory. Unlike other national surveys conducted by CDC National Center for Health Statistics, the National Health Care surveys address the health care delivery systems rather than the vital statistics, health status, health-related behavior, and access to care experienced by individuals and households who are consumers of the health care delivery systems. Between the years of 1968 and 1984, a number of surveys were conducted to learn more about National Center for Health Statistics (NCHS) data users and to assess the quality of data dissemination activities conducted by NCHS. Studies focusing solely on user satisfaction with National Health Care Survey data products have not been conducted since 1984. We need current specific information on how well our users' needs are being met, how to improve our data products, and how to serve current non-users of our data who are, nonetheless, potential users. Our data products consist mainly of published reports and web-published data sets including Data Highlights and E-Stats. Our published reports include Advance Data Reports, a newsletter-like summary of more detailed analyses to be published later, and Series Reports, which are in-depth analyses of specific topics addressed by our collected data. As the contractor for this project, CHPS Consulting will conduct a multi-mode survey using a web-based survey for those in the sample for whom an email address is available and a mail survey for those without an email address. Current users will be asked questions about what publications they use, how they use them, and their opinion of the timeliness, accessibility, format, and quality of the data publications. Non-users will be asked why they do not use our publications, their current sources of health care provider data, and how we improve data products to meet their needs. Our target population will include the following groups of persons: researchers, educators, health facility administrators, practitioners, and policymakers. Our goal for this survey is to obtain 600 returned surveys with an approximately equal number of returned surveys from users and non-users. There is no cost to respondents other than their time in responding.The total annualized burden is 75 hours.

Respondents	Number of respondents	Number of responses per respondent	Avg. burden per response (in hrs.)	Total response per burden (in hrs.)
Users	300	1	10/60	50
Non-Users	300	1	5/60	25
Total	600			75

Dated: January 16, 2001.

Nancy Cheal,

Acting Associate Director for Policy, Planning, and Evaluation Centers for Disease Control and Prevention (CDC).

[FR Doc. 01–1995 Filed 1–22–01; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60 Day–01–16]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 639–7090.

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 through the use of automated collection techniques for other forms of information technology. Send comments to Seleda Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project

Foreign Quarantine Regulations—Extension—OMB No.0920–0134 National Center for Infectious Diseases (NCID), Centers for Disease Control and Prevention (CDC) Section 361 of the Public Health Service (PHS) Act (42 USC 264) authorizes the Secretary of Health and Human Services to make and enforce regulations necessary to prevent the introduction, transmission, or spread of communicable diseases from foreign countries into the United States. Legislation and the existing regulations governing quarantine activities (42 CFR part 71) authorize quarantine officers and other personnel to inspect and undertake necessary control measures with respect to conveyances, persons, and shipments of

animals and etiologic agents in order to protect the public health. Currently, with the exception of rodent inspections and the cruise ship sanitation program, inspections are performed only on those vessels and aircraft which report illness prior to arrival or when illness is discovered upon arrival. Other inspection agencies assist quarantine officers in public health screening of persons, pets, and other importations of public health importance and make referrals to PHS when indicated. These practices and procedures assure protection against the introduction and spread of communicable diseases into the United States with a minimum of recordkeeping and reporting as well as a minimum of interference with trade and travel. Respondents would include airplane pilots, ships' captains, importers, and travelers. The nature of the quarantine response would dictate which forms are completed by whom. Thus, the *respondents* portion of the information below is replaced by the requisite form title. The estimated cost to the public is $22,225.

Respondents	Number of respondents	Number of responses/ respondent	Avg. burden per respondent (in hrs.)	Total burden (in hrs.)
Radio reporting of death/illness:				
(1) Aircraft	130	1	2/60	4.00
(2) Cruise ships	90	23	1/60	34.00
(3) Other ships	22	1	1/60	0.04
Report by persons held in isolation/surveillance	11	1	30/60	5.50
Report of death or illness on carrier during stay in port	5	1	3/60	0.25
Requirements for admission of dogs and cats:				
(1)	5	1	3/60	0.25
(2)	2,650	1	15/60	662.50
Application for permits to import turtles	10	1	30/60	5.00
Requirements for registered importers of nonhuman primates:				
(1)	40	1	10/60	6.70
(2)	50	1	30/60	25.00
Total				743.60

Dated: January 12, 2001.

Nancy Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 01–1996 Filed 1–22–01; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01N–0006]

Agency Information Collection Activities; Proposed Collection; Comment Request; New Animal Drug Application, Form FDA 356 V, 21 CFR Part 514

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed reinstatement of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on requirements for submission of a new animal drug application (NADA).

DATES: Submit written or electronic comments on the collection of information by March 26, 2001.

ADDRESSES: Submit electronic comments on the collection of information via the Internet at: http://www.accessdata.fda.gov/scripts/oc/dockets/edockethome.cfm. Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information listed below.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.