2301, will continue to be counted as part of OMB control number 0910-0284.

The reporting and recordkeeping burden estimates, including the total number of annual responses, are based

on the submission of reports to the Division of Surveillance, Center for Veterinary Medicine. The annual frequency of responses was calculated as the total annual responses divided by the number of respondents.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN¹

21 CFR Section or Section of the Act	FDA Form No.	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
514.80(b)(1), (b)(2)(i), (b)(2)(ii), and (b)(3)	1932 ²	404	44.26	17,882.5	1	17,882.5
Voluntary reporting FDA Form 1932a for the public	1932a²	81.5	1	81.5	1 ³	81.5
514.80(b)(4)	2301	84	17.0	1,428	16	22,848
514.80(b)(5)(i)	2301	84	0.31	26	2	52
514.80(b)(5)(ii)	2301	84	33.92	2,849	2	5,698
514.80(b)(5)(iii)	2301	646	0.08	49	2	98
Total Hours	46,660					

¹There are no capital costs or operating and maintenance costs associated with this collection of information. ²Burden hours were determined as explained above.

TABLE 2.—ESTIMATED ANNUAL RECORDKEEPING BURDEN¹

21 CFR Section	No. of Recordkeepers	Annual Frequency per Recordkeeping	Total Annual Records	Hours per Record	Total Hours
514.80(e) ²	646	7.20	4651	14	65,116.8
Total					1,541

¹There are no capital costs or operating and maintenance costs associated with this collection of information.

Dated: October 7, 2009.

David Horowitz.

Assistant Commissioner for Policy. [FR Doc. E9-24734 Filed 10-14-09; 8:45 am] BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Health Disparities Subcommittee, **Advisory Committee to the Director** (ACD), Centers for Disease Control (CDC); Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC) announces the following meeting of the aforementioned subcommittee.

Time and Date: 2 p.m.-4:30 p.m., October 28, 2009.

Place: The meeting will be convened at the CDC, 1600 Clifton Road, NE., Atlanta, GA

30333, Building 19, Auditorium B1, Global Communications Center. Please see Supplementary Information for details on accessing the meeting location.

Status: Open to the public, limited only by the availability of space. The meeting room accommodates approximately 90 people.

Purpose: The Subcommittee will provide advice to the CDC Director through the Advisory Committee to the Director on strategic and other broad issues facing CDC.

Matters To Be Discussed: ACD Health Disparities Subcommittee 2009 Action Agenda; CDC Director's Health Disparity Indicator Project Update, Director's Priorities and Reorganization/Structure.

Agenda items are subject to change as priorities dictate.

Supplementary Information: To participate in the meeting, please plan to register with CDC Security Officials at the Visitor's Center at least one hour prior to the meeting. A government-issued picture ID will be required. All persons who do not have a CDC/Health and Human Services identification will have to be escorted to the

Contact Person for More Information: Walter W. Williams, M.D., M.P.H., Designated Federal Officer, Health

Disparities Subcommittee, ACD, CDC, 1600 Clifton Road, NE., M/S E-67, Atlanta, Georgia 30333. Telephone 404/498-2310, Email: http://www1@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: October 8, 2009.

Andre Tyler,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. E9-24859 Filed 10-14-09; 8:45 am] BILLING CODE 4163-18-P

³The hours per response for paper versions of Forms FDA 1932 and 1932a are assumed to be 1 hour. The hours per response for the electronic version of Form FDA 1932 is assumed to be 1 hour, while the electronic version of Form FDA 1932a is assumed to take .6 hours to complete the form and gather the required information as part of the MedWatch^{Plus} Portal information collection (see 74 FR 23721 at 23727, May 20, 2009).

² Section 514.80(e) covers all recordkeeping hours for all adverse event reporting.