the administrative record shall submit them with a new petition to modify the decision. FDA uses the information provided in the request to determine whether to grant the petition for reconsideration. Respondents to this collection of information are individuals of households, State or local governments, not-for-profit institutions, and businesses or other for-profit instructions who are requesting from the Commissioner a reconsideration of a matter.

Section 10.35 (21 CFR 10.35) issued under section 701(a) of the act, sets forth the format and procedures by which an interested person may request, in accordance with § 10.20 (submission of documents to the Dockets Management Branch), the Commissioner to stay the effective date of any administrative action.

Such a petition must: (1) Identify the decision involved, (2) state the action requested—including the length of time for which a stay is requested, and (3) include a statement of the factual and legal grounds on which the interested person relies in seeking the stay. FDA uses the information provided in the request to determine whether to grant the petition for stay of action.

Respondents to this information collection are interested persons who choose to file a petition for an administrative stay of action.

Section 10.85 (21 CFR 10.85), issued under section 701(a) of the the act, sets forth the format and procedures by which an interested person may request, in accordance with § 10.20 (submission of documents to the Dockets Management Branch), an advisory opinion from the Commissioner on a matter of general applicability. An advisory opinion represents the formal position of FDA on a matter of general applicability. When making a request, the petitioner must provide a concise statement of the issues and questions on which an opinion is requested, and, a full statement of the facts and legal points relevant to the request. Respondents to this collection of information are interested persons seeking an advisory opinion from the Commissioner on the agency's formal position for matters of general applicability.

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

TABLE 1.— ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
10.30	150	3	450	12	5,400
10.33	10	1	10	10	100
10.35	13	1	13	10	130
10.85	3	1	3	16	48
Total					5,678

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

The burden estimates for this collection of information is based on agency records and experience over the past 3 years. Agency personnel handling the petitions for § 10.30 estimate 150 (citizen petitions) received by the agency annually, each requiring an average of 12 hours preparation time. Agency personnel handling the petitions for § 10.33 (administrative reconsideration of an action) estimate 10 requests are received by the agency annually, each requiring an average of 10 hours preparation time. Agency personnel handling the petitions for § 10.35 (administrative stay of an action) estimate 13 requests are received by the agency annually, each requiring an average of 10 hours preparation time. Agency personnel handling the petitions for § 10.85 (advisory opinions) estimate 3 requests are received by the agency annually, each requiring an average of 16 hours preparation time.

Dated: June 27, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–17077 Filed 7–8–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Statement of Delegation of Authority

Notice is hereby given that I have delegated to the Director, National Institutes of Health (NIH), the authorities under Public Law 107–84, Muscular Dystrophy Community Assistance, Research and Education Amendments (MD–CARE) Act, Part A, Title IV, Section 404E(d) of the Public Health Service Act, as amended, to establish the Muscular Dystrophy Coordinating Committee.

I reserve to myself the authority to appoint members of the Coordinating Committee, including the Chair of the Coordinating Committee.

This delegation shall be exercised in accordance with the Department's applicable policies, procedures, guidelines and regulations. In addition, I ratify and affirm any actions taken by the NIH Director or his subordinates which involve the exercise of the authorities delegated herein prior to the effective date of this delegation.

The delegation is effective upon date of signature.

Dated: June 26, 2002.

Tommy G. Thompson,

Secretary.

[FR Doc. 02–17129 Filed 7–8–02; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR–4739–N–27]

Notice of Proposed Information Collection: Comment Request; Capital Advance Program Requirements for Section 202 Housing for the Elderly, and Section 811 Housing for Persons With Disabilities

AGENCY: Office of the Assistant Secretary for Housing-Federal Housing Commissioner, HUD.

ACTION: Notice.

SUMMARY: The proposed information collection requirement described below will be submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

DATES: *Comments Due Date:* September 9, 2002.