final rule do not impose an additional burden since the members will be complying with the current organizations' standards which are comparable to the requirements in the final rule. To account for persons or establishments that may not be a member of an industry organization and, for whom therefore, the extent of compliance with the requirements of the final rule is unknown, FDA will be using 1 percent as an estimation of the information collection burden on the tissue industry.

Industry estimates that in 1994 there were 350,000 bone transplants, 42,000 corneal transplants, 5,000 patellar tendon transplants, and the transplantation of 5,000 square feet of skin. There are approximately 300 persons and 170 tissue banks currently operating in the Untied States affected by the regulations.

The total annual estimated burden imposed by this collection of information is 32,260 hours annually.

TABLE 1—ESTIMATED ANNUAL RECORDKEEPING BURDEN

21 CFR Section	No. of Recordkeepers	Annual Frequency of Recordkeeping	Total Annual Records	Hours per Recordkeeper	Total Hours
1270.31 (b)–(d)	11	4	44	28	308
1270.35 (a)–(b)	11	420	4,620	290	3,190
1270.35 (c)	11	2,893	31,823	4,782	52,602
1270.35 (d)	11	17	187	17	187
Total	.....	.....	.....	.....	56,287

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

Dated: December 5, 1997.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 97–32459 Filed 12–10–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that published in the **Federal Register** of November 25, 1997 (62 FR 62777). The notice announced a meeting of the Orthopaedics and Rehabilitation Devices Panel of the Medical Devices Advisory Committee that is scheduled for December 11 and 12, 1997. The notice published with an error. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: LaJuana D. Caldwell, Office of Policy (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–2994.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of November 25, 1997 (62 FR 62777), in FR Doc. 97–30914, FDA announced that a meeting of the Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee would be held on December 11 and 12, 1997. The notice published with an error in the first sentence in the *Agenda* portion of the meeting.

Beginning on page 62777, in the 2d column, under the *Agenda* portion of the meeting, the first sentence should be corrected to read ''On December 11, 1997, the committee will discuss, make recommendations, and vote on one premarket approval application (PMA) for a spinal intervertebral fusion device and a second PMA for a spinal intervertebral fusion system.''

Dated: December 9, 1997.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 97–32584 Filed 12–9–97; 3:03 pm]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Council; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of February 1998:

Name: National Advisory Committee on Rural Health.

Dates and Time: February 1–4, 1998.

Place: J.W. Marriott Hotel, 1331 Pennsylvania Avenue, N.W., Washington, D.C. 20004, Phone: (202) 393–2000, FAX: (202) 626–6915.

The meeting is open to the public.

Agenda: The plenary session on Monday morning, February 2, will include a national legislative update, discussion of quality issues, universal service provisions of the Telecommunications Act, definition of rural, and telemedicine payment issues.

Presentations on graduate medical education, the rural minority health project, and long-term care; assisted living housing will be the bases of discussion for the Committee of the Whole on Tuesday.

The final plenary session will be convened on Wednesday, February 4, at 8:00 a.m. During this session there will be an update of the Office of Rural Health Policy activities, a report of the Committee of the Whole regarding the discussions that took place on Tuesday, and information regarding the next agenda and future meeting dates and places will be discussed. The meeting will be adjourned at approximately 11:30 a.m.

Anyone requiring information regarding the subject Committee should contact Dena S. Puskin, Sc.D., Executive Secretary, National Advisory Committee on Rural Health, Health Resources and Services Administration, Room 9–05, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857, telephone (301) 443–0835, FAX (301) 443–2803.

Persons interested in attending any portion of the meeting should contact Arlene A. Granderson or Lilly Smetana, Office of Rural Health Policy, (301) 443–0835.

Agenda items change as priorities dictate.

Dated: December 4, 1997.

Jane M. Harrison,

Committee Management Office, HRSA.

[FR Doc. 97–32409 Filed 12–10–97; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Human Genome Research Institute; Notice of Meeting

Pursuant to Section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meeting: