

usefulness and cost-effectiveness of this teen asthma intervention program. Sample participants will come from students, parents, program facilitators, and school personnel (school nurses and teachers) in the selected two school districts. Self-administered questionnaires will be given to students at baseline (pre-intervention program), immediately post-program, and at 6-months post-program, while parents

receive baseline and 6-month post-program surveys. The student survey will focus on: knowledge, attitudes, and behaviors regarding their asthma; perception of their health status and quality of life; assessment of the program; and impact of the program on their asthma management skills. Parents will be asked about their child's asthma condition, assessment of the program, and cost-related issues for their child's

asthma. Individual, one-time interviews will be conducted with program facilitators and school personnel regarding their perceptions of the intervention program and its impact on the students. Two focus groups will be conducted with students post-program to obtain additional, in-depth information about their perceptions of the program.

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hrs.)	Total burden (in hrs.)
Students:				
Baseline	524	1	30/60	262
Post-program	524	1	15/60	131
6-month follow-up	524	1	30/60	262
Focus group	16	1	1	16
Parents:				
Baseline	524	1	10/60	87
6-month follow-up	524	1	15/60	131
Program facilitators:				
Interview	6	1	40/60	4
Program sessions	6	12	30/60	36
School nurses:				
School profile	6	1	10/60	1
Record abstraction	6	87	10/60	87
Interview	6	1	40/60	4
Teachers Interview	12	1	40/60	8
Total				1029

Dated: May 13, 2003.

Thomas Bartenfeld,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Endocrinologic and Metabolic Drugs Advisory Committee; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

The Food and Drug Administration (FDA) is announcing an amendment to the notice of the meeting of the Endocrinologic and Metabolic Drugs Advisory Committee. This meeting was announced in the **Federal Register** of May 6, 2003 (68 FR 24003). The amendment is being made to reflect a change in the *Agenda* portion of the document. There are no other changes.

FOR FURTHER INFORMATION CONTACT:

Dornette Spell-LeSane, Center for Drug Evaluation and Research (HFD-21),

Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301-827-7001, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 12536. Please call the Information Line for up-to-date information on this meeting.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of May 6, 2003 (68 FR 24003), FDA announced that a meeting of the Endocrinologic and Metabolic Drugs Advisory Committee would be held on June 10, 2003. On page 24003, in the third column, the *Agenda* portion of the meeting is amended to read as follows:

Agenda: The committee will discuss supplemental new drug application (sNDA) 19-604/S-033 HUMATROPE (somatropin recombinant deoxyribonucleic acid (rDNA) origin) for injection, Eli Lilly and Co., for the proposed indication of treatment of nongrowth hormone deficiency short stature.

The notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to advisory committees.

Dated: May 13, 2003.

Peter J. Pitts,

Associate Commissioner for External Relations.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of the Committee: Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee:

To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on June 9, 2003, from 12:30 p.m.