b. Data on the growth of *L. monocytogenes* on non-food surfaces including environmental biofilm growth.

6. Factors that influence the environmental contamination and the cross-contamination of food by *L. monocytogenes* in retail facilities, including:

a. Data and information on the potential transfer of *L. monocytogenes* to food from the retail environment, e.g., experimental studies on the transfer to food from drains, slicers, food contact surfaces, and non-food contact surfaces; and

b. Data and information on food handlers' activities, e.g., observations of food handlers' practices and monitoring of specific food safety actions in retail facilities (e.g., glove usage, hand hygiene practices, and cleaning practices).

7. Identity and effectiveness of control measures or interventions intended to reduce levels and frequency of *L. monocytogenes* in the retail environment, including:

a. Environmental sanitation procedures including the sanitizers and protocols used, frequency of application, and efficiency; and

b. Worker sanitation procedures including frequencies, protocols, and efficiency.

8. Any other data related to the occurrence, growth, and control of *L. monocytogenes* in retail facilities.

As the project progresses, additional data needs may be identified.

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments regarding this document. Submit a single copy of electronic comments or two paper copies of any mailed comments, except that individuals may submit one paper copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Please note that on January 15, 2008, the FDA Division of Dockets Management Web site transitioned to the Federal Dockets Management System (FDMS). FDMS is a Government-wide, electronic docket management system. Electronic comments or submissions will be accepted by FDA only through FDMS at *http://www.regulations.gov*.

III. References

The following references are on display in the Division of Dockets Management (see **ADDRESSES**) and may be seen by interested persons between 9 a.m. and 4 p.m., Monday through Friday. (FDA has verified the Web site addresses, but FDA is not responsible for any subsequent changes to the Web sites after this document publishes in the **Federal Register**.)

1. U.S. Department of Health and Human Services, Healthy People 2010, v. 1. Washington, DC, 2000, *http://healthypeople.gov*.

2. U.S. Department of Health and Human Services and U.S. Department of Agriculture/Food Safety and Inspection Service, ''Quantitative Assessment of Relative Risk to Public Health from Foodborne *Listeria monocytogenes* Among Selected Categories of Ready-to-Eat Foods,'' September 2003, *http://www.foodsafety.gov/~dms/lmr2-toc.html*.

3. U.S. Department of Health and Human Services, Food and Drug Administration/Centers for Disease Control and Prevention, ''Reducing the Risk of *Listeria monocytogenes* FDA/CDC 2003 Update of the Listeria Action Plan,'' November 2003, *http://www.cfsan.fda.gov/~dms/lmr2plan.html*.

4. Gombas, D.E., Chen, Y., Clavero, R.S., and Scott, V.N. (2003). Survey of *Listeria monocytogenes* in ready-to-eat foods. *Journal of Food Protection*, 66(4), 559–569.

5. Draughon, A.F. (2006). A collaborative analysis/risk assessment of *Listeria monocytogenes* in ready-to-eat processed meat and poultry collected in four FoodNet states. Symposium S–16: Contamination of ready-to-eat foods: transfer and risk: *Listeria monocytogenes* and other microorganisms. International Association for Food Protection 93rd Annual Meeting, Calgary, Alberta. August 13–16.

Dated: January 12, 2009.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E9–938 Filed 1–16–09; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Advisory Council on Migrant Health; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), notice is hereby given of the following meeting:

Name: National Advisory Council on Migrant Health.

Dates and Times: February 9, 2009, 8:30 a.m. to 5 p.m.; February 10, 2009, 8:30 a.m. to 5 p.m.

Place: The Parklawn Building, Twinbrook Room, 3rd Floor, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone: (301) 594–4303, Fax: (301) 443–0248.

Status: The meeting will be open to the public.

Purpose: The purpose of the meeting is to discuss services and issues related to the health of migrant and seasonal farmworkers and their families and to formulate recommendations for the Secretary of Health and Human Services.

Agenda: The agenda includes an overview of the Council's general business activities. The Council will also hear presentations from experts on farmworker issues, including the status of farmworker health at the local and national levels.

Agenda items are subject to change as priorities indicate.

For Further Information Contact: Gladys Cate, Office of Minority and Special Populations, Bureau of Primary Health Care, Health Resources and Services Administration, 5600 Fishers Lane, Maryland 20857; telephone (301) 594–0367.

Wendy Ponton,

Director, Office of Management.

[FR Doc. E9–1067 Filed 1–16–09; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Government-Owned Inventions; Availability for Licensing

AGENCY: National Institutes of Health, Public Health Service, HHS.

ACTION: Notice.

SUMMARY: The inventions listed below are owned by an agency of the U.S. Government and are available for licensing in the U.S. in accordance with 35 U.S.C. 207 to achieve expeditious commercialization of results of federally-funded research and development. Foreign patent applications are filed on selected inventions to extend market coverage for companies and may also be available for licensing.

ADDRESSES: Licensing information and copies of the U.S. patent applications listed below may be obtained by writing to the indicated licensing contact at the Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852–3804; telephone: 301/496–7057; fax: 301/402–0220. A signed Confidential Disclosure Agreement will be required to receive copies of the patent applications.

Mice With a Conditional LoxP-Flanked Glucosylceramide Synthase Allele Controlling Glycosphingolipid Synthesis

Description of Technology: Glycosphingolipids are organizational building blocks of plasma membranes that participate in key cellular