D. Where To Obtain Additional Information

For general comments or questions about this announcement, contact: Technical Information Management, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341–4146. Telephone: 770–488–2700.

For technical questions about this program, contact: Mildred Williams-Johnson, Ph.D., Scientific Program Administrator, CDC, National Center for Environmental Health, 1600 Clifton Road, NE., Mail Stop E17, Atlanta, GA 30333. Telephone: 404–498–0639. E-mail: MWilliams-Johnson@cdc.gov.


William P. Nichols,
Director, Procurement and Grants Office,
Centers for Disease Control and Prevention.

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BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Reducing Racial and Ethnic Disparities in Childhood Immunization

Announcement Type: New.
Funding Opportunity Number: RFA IP05–087.
Catalog of Federal Domestic Assistance Number: 93.185.
Application Deadline: June 27, 2005.

I. Funding Opportunity Description

Authority: Section 311 (42 U.S.C. 243) and 317(k)(1) (42 U.S.C. 247b(k)(1)) of the Public Health Service Act, as amended.

Background

Eliminating health disparities among racial and ethnic populations in the United States is a major public health goal. However, in recent years, disparities in immunization rates between black and white children have been increasing (Chu et al.). Therefore, the National Immunization Program (NIP) is seeking to support projects that may lead to reductions in these disparities.

Factors that may be related to lower immunization rates among black children include frequency and timing of well child visits, provider type (pediatrician, family practitioner, public

health clinic (PHC)), missed opportunities for immunization, socioeconomic status (SES), urban vs. rural vs. suburban settings, and parental beliefs. Missed opportunities are medical encounters during which a child fails to receive an immunization for which he/she is eligible and they have been shown to contribute to under immunization of children. The National Maternal and Infant Health Survey showed that black children were less likely than white children to receive the recommended number of well child visits and immunizations in the first seven months of life. SES has been shown to impact immunization coverage levels in many studies. Some studies have found that adjustment for SES and access to care did not completely explain racial and ethnic disparities.

Purpose

The purpose of the program is to fund a community-based demonstration project to identify, implement and evaluate interventions that will result in a statistically significant reduction in racial disparities in immunization coverage levels between black children 19–35 months of age and children of other races, particularly white children, as evidenced by a comparison of immunization coverage of black and other racial/ethnic groups before and after interventions are implemented. Throughout this announcement black refers to non-Hispanic black and white refers to non-Hispanic white. These interventions must include: (1) enhancement of healthcare utilization and (2) strategies to reduce missed opportunities for immunization. The key to the success of this program will be community-focused programs that include the full engagement of appropriate partners. These partners may include faith-communities, health care purchasers, health plans, health care providers, and many other community sectors working together. The focus of this announcement is for medium or large urban areas with populations of at least 100,000 people. This program addresses the “Healthy People 2010” focus area of Immunization and Infectious Diseases.

Measurable outcomes of the program will be in alignment with the performance goal for the Center for Disease Control and Prevention’s (CDC) National Immunization Program (NIP) to reduce the number of indigenous vaccine-preventable diseases and will be evidenced by a significant increase in immunization coverage levels among black children in the study communities before and after implementation of study interventions. A significant increase is defined as 90 percent confidence in having achieved an increase in coverage among black children of at least five percentage points with no increase in disparities.

Research Objectives

1. Identify factors related to disparities in childhood immunization rates between black children and children of other racial/ethnic groups within an urban area. These factors must include community and practice level factors related to utilization of health services and practice level factors related to missed opportunities for immunization.

2. Develop and implement interventions to address factors related to disparities in immunization rates between black children and children of other racial/ethnic groups. The applicant must address community and practice level factors related to enhancing utilization of health services and practice level factors related to missed opportunities for immunization.

3. Evaluate the effectiveness of these interventions in decreasing racial disparity in immunization rates between blacks and all other children within the urban area.

Activities

Awardee activities for this program are as follows:

1. Select a medium or large urban area with a total population of at least 100,000 people, with documented significant racial/ethnic disparities in childhood immunization rates. At least 25 percent of this urban area should be black.

2. Develop and implement plans to identify factors which are related to the disparity differences in immunization coverage between black children and children of other racial/ethnic groups in this urban area. These factors must include community and practice level factors related to utilization of health services and practice level factors related to missed opportunities for immunization. Examples include number and timing of well child visits, pattern of missed opportunities, SES status, provider type (family practitioner, pediatrician, PHC), and availability of social services and transportation within the urban area.

3. Design interventions for addressing the factors related to disparities in immunization coverage in this urban area. These interventions must address community and practice level factors related to utilization of health services and practice level factors related to missed opportunities for...
immunization. Interventions also need to involve collaboration between the community and practice-based activities, as well as, a plan for sustainability of these activities. Programs are expected to employ multiple strategies, including innovative strategies as well as evidence-based public health strategies based at least partially on the existing and emerging research base and careful scientific review such as the Guide to Community Preventive Services (http://www.thecommunityguide.org/).

Effective public health strategies may include changes to the social and physical environments; health promotion, public education, and information; media and other communication strategies; technological advances; economic incentives and disincentives; system improvements; provider education and medical office-based improvement strategies. While they may be included, mass media campaigns should not constitute the sole intervention aimed at the community. While project activities should reach all persons in an identified intervention area, special efforts should be taken to ensure focus on black populations experiencing disparities in access to and use of preventive services.

Because sustainability is important, the program must include a plan for sustaining interventions past the funding period.

Programs must be culturally competent, and meet the health literacy and linguistic needs of target populations in the intervention area.

Programs could optimize resources by coordinating and partnering with existing programs and resources in the community, surrounding areas, and the state.

Collaborative partnerships with, for example, professional organizations; health care providers, employers, purchasers, and health plans; faith-based organizations; schools; child care, early childhood programs, Women, Infants, and Children (WIC) program, and other organizations that serve children; and many others are key to reaching affected populations and delivering and sustaining effective programs. Strong, cooperative linkages between clinical preventive care and community public health should be established and maintained.

4. Implement interventions within multiple immunization provider practices. At a minimum, a representative sample of at least 30 practices in the urban area should participate in the intervention and be evaluated. This sample should be representative of where whites receive care and of where whites receive care in a geographically defined area (city or region) where blacks account for at least 25 percent of the population. Each sample must include a sufficient number of clinics for meaningful comparisons to be made. Because disparities persist across socioeconomic categories, it is important that clinics that serve patients of higher SES be represented as well as clinics serving patients of lower SES.

5. Validate or document degree of implementation of interventions, including number of persons reached by, and use of intervention strategies; tracking the accomplishment of activities and the achievement of short-term and intermediate outcomes; monitoring changes in health outcomes; and using program evaluation findings to adjust plans and strengthen the program. This would involve identification and collection of appropriate process measures through multiple means and would also involve direct observation of practices.

6. Determine effectiveness of interventions by comparing immunization rates between black children and children of racial/ethnic groups within and between practice sites. The evaluation must include a comparison of immunization coverage of black and other racial/ethnic groups before and after interventions are implemented. In addition, if available, population-based measures (cluster surveys or random digit dial telephone surveys) can also be used to monitor coverage rates.

7. Identify the most effective, feasible, and sustainable interventions in reducing disparities in immunization rates in this urban area.

8. Collaboratively disseminate research findings in peer reviewed publications and for use in determining national policy.

Because sustainability is important and the program included a plan for sustaining interventions, we encourage measures of progress past the project period. In a cooperative agreement, CDC staff is substantially involved in the program activities, above and beyond routine grant monitoring.

CDC Activities for this program are as follows:

1. Provide CDC investigator(s) to monitor the cooperative agreement as project officer(s).

2. Participate as active project team members in the development, implementation and conduct of the research project and as coauthors of all scientific publications that result from the project.

3. Provide technical assistance on the selection and evaluation of data collection and data collection instruments.

4. Assist in the development of research protocols for Institutional Review Boards (IRB) review. The CDC IRB will review and approve the project protocol or will defer to outside IRB, and will do so on at least an annual basis until the research project is completed.

5. Contribute subject matter expertise in the areas of epidemiologic methods and statistical analysis, and survey research consultation.

6. Participate in the analysis and dissemination of information, data and findings from the project, facilitating dissemination of results.

7. Serve as liaisons between the recipients of the project award and other administrative units within the CDC.

8. Facilitate an annual meeting between awardee and CDC to coordinate planned efforts and review progress.

II. Award Information

Type of Award: Cooperative Agreement. CDC involvement in this program is listed in the Activities Section above.

Mechanism of Support: U01.

Fiscal Year Funds: 2005.

Approximate Total Funding: $300,000 (Includes direct and indirect costs. This amount is an estimate, and is subject to availability of funds.)

Approximate Number of Awards: One.

Appro biased Average Award: $300,000 (Includes direct and indirect costs. This amount is for the first 12-month budget period.)

Floor of Award Range: None.

Ceiling of Award Range: $300,000 (Includes direct and indirect costs. This ceiling is for the first 12-month budget period.)

Anticipated Award Date: August 31, 2005.

Budget Period Length: 12 months.

Project Period Length: Three (3) years. Throughout the project period, CDC's commitment to continuation of awards will be conditioned on the availability of funds, evidence of satisfactory progress by the recipient (as documented in required reports), and the determination that continued funding is in the best interest of the Federal Government.

III. Eligibility Information

III.1. Eligible applicants

Applications are limited to public and private nonprofit organizations and by governments and their agencies, such
as: (For profit organizations are not eligible under Section 317(k)(1) [42 U.S.C. 247(b)(1)] of the Public Health Service Act, as amended.)

- Public nonprofit organizations.
- Private nonprofit organizations.
- Small, minority, women-owned businesses.
- Universities.
- Colleges.
- Research institutions.
- Hospitals.
- Community-based organizations.
- Faith-based organizations.
- Federally recognized Indian tribal governments.
- Indian tribes.
- Indian tribal organizations.
- State and local governments or their Bona Fide Agents (this includes the District of Columbia, the Commonwealth of Puerto Rico, the Virgin Islands, the Commonwealth of the Northern Mariana Islands, American Samoa, Guam, the Federated States of Micronesia, the Republic of the Marshall Islands, and the Republic of Palau).
- Political subdivisions of States (in consultation with States).
- A Bona Fide Agent is an agency/or organization identified by the state as eligible to submit an application under the state eligibility in lieu of a state application. If you are applying as a bona fide agent of a State or local government, you must provide a letter from the State or local government as documentation of your status. Place this documentation behind the first page of your application form.

III.2. Cost Sharing or Matching

Matching funds are not required for this program.

III.3. Other

If you request a funding amount greater than the ceiling of the award range, your application will be considered non-responsive, and will not be entered into the review process. You will be notified that your application did not meet the submission requirements.

Special Requirements: If your application is incomplete or non-responsive to the requirements listed in this section, it will not be entered into the review process. You will be notified that your application did not meet submission requirements.

Late applications will be considered non-responsive. See section "IV.3. Submission Dates and Times" for more information on deadlines.

Note: Title 2 of the United States Code Section 1611 states that an organization described in Section 501(c)(4) of the Internal Revenue Code that engages in lobbying activities is not eligible to receive Federal funds constituting an award, grant, or loan.

Individuals Eligible to Become Principal Investigators: Any individual with the skills, knowledge, and resources necessary to carry out the proposed research is invited to work with their institution to develop an application for support. Individuals from underrepresented racial and ethnic groups as well as individuals with disabilities are always encouraged to apply for CDC programs.

IV. Application and Submission Information

IV.1. Address To Request Application Package

To apply for this funding opportunity, use application form PHS 398 (OMB number 0920–0001 rev. 9/2004). Forms and instructions are available in an interactive format on the CDC Web site, at the following Internet address: http://www.cdc.gov/od/pgo/forminfo.htm.

IV.2. Content and Form of Application Submission

Letter of Intent (LOI): Your LOI must be written in the following format:

- Maximum number of pages: 2.
- Font size: 12-point unreduced.
- Double spaced.
- Paper size: 8.5 by 11 inches.
- Page margin size: One inch.
- Printed only on one side of page.
- Written in plain language, avoid jargon.

Your LOI must contain the following information:

- Descriptive title of the proposed research.
- Name, address, E-mail address, telephone number, and FAX number of the Principal Investigator.
- Names of other key personnel.
- Participating institutions.
- Number and title of this Announcement.

Application: Follow the PHS 398 application instructions for content and formatting of your application. For further assistance with the PHS 398 application form, contact PGO–TIM staff at 770–488–2700, or contact GrantsInfo, Telephone (301) 435–0714, E-mail: GrantsInfo@nih.gov.

Your research plan should address activities to be conducted over the entire project period.

You are required to have a Dun and Bradstreet Data Universal Numbering System (DUNS) number to apply for a grant or cooperative agreement from the Federal Government. Your DUNS number must be entered on line 11 of the face page of the PHS 398 application form. The DUNS number is a nine-digit identification number, which uniquely identifies business entities. Obtaining a DUNS number is easy and there is no charge. To obtain a DUNS number, access http://www.dunandbroadstreet.com or call 1–866–705–5711.

For more information, see the CDC Web site at: http://www.cdc.gov/od/pgo/funding/pubcomm1.htm.

This announcement uses the non-modular budgeting format.

Additional requirements that may require you to submit additional documentation with your application are listed in section “VI.2. Administrative and National Policy Requirements.”

IV.3. Submission Dates and Times

LOI Deadline Date: June 10, 2005.

CDC requests that you send a LOI if you intend to apply for this program. Although the LOI is not required, not binding, and does not enter into the review of your subsequent application, the LOI will be used to gauge the level of interest in this program, and to allow CDC to plan the application review.

Application Deadline Date: June 27, 2005.

Explanation of Deadlines: LOIs must be received in the CDC Office of Public Health (OPHR) and applications must be received in the CDC Procurement and Grants Office by 4 p.m. eastern time on the deadline date. If you submit your LOI and Application by the United States Postal Service or commercial delivery service, you must ensure that the carrier will be able to guarantee delivery by the closing date and time. If CDC receives your submission after closing due to: (1) Carrier error, when the carrier accepted the package with a guarantee for delivery by the closing date and time, or (2) significant weather delays or natural disasters, you will be given the opportunity to submit documentation of the carriers guarantee. If the documentation verifies a carrier problem, CDC will consider the
submission as having been received by the deadline.

This announcement is the definitive guide on LOI and application content, submission address, and deadline. It supersedes information provided in the application instructions. If your application does not meet the deadline above, it will not be eligible for review, and will be discarded. You will be notified that you did not meet the submission requirements.

CDC will not notify you upon receipt of your submission. If you have a question about the receipt of your LOI or application, first contact your courier. If you still have a question concerning your LOI, contact the OPHR staff at 404–371–5277. If you still have a question concerning your application, contact the PGO–TIM staff at: 770–488–2700. Before calling, please wait two to three days after the submission deadline. This will allow time for submissions to be processed and logged.

IV.4. Intergovernmental Review of Applications

Your application is subject to Intergovernmental Review of Federal Programs, as governed by Executive Order (EO) 12372. This order sets up a system for State and local governmental review of proposed federal assistance applications. You should contact your state single point of contact (SPOC) as early as possible to alert the SPOC to your LOI by express mail, delivery service, fax, or E-mail to: Mary Lerchen, DrPH, Scientific Review Administrator, CDC/Office of Public Health Research, One West Court Square, Suite 7000, MS D–72, Telephone: 404–371–5277, Fax: 404–371–5215; E-mail: M Lerchen@cdc.gov.

Application Submission Address: Submit the original and one hard copy of your application by mail or express delivery service to: Technical Information Management—RFA IP05–087, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341.

At the time of submission, four additional copies of the application, and all appendices must be sent to: Mary Lerchen, DrPH, Scientific Review Administrator, CDC/Office of Public Health Research, One West Court Square, Suite 7000, MS D–72, Telephone: 404–371–5277, Fax: 404–371–5215, E-mail: M Lerchen@cdc.gov.

Applications may not be submitted electronically at this time.

IV.5. Funding Restrictions

Restrictions, which must be taken into account while writing your budget, are as follows:

• Funds relating to the conduct of research will not be released until the appropriate assurances and Institutional Review Board approvals are in place.
• Reimbursement of pre-award costs is not allowed.

If you are requesting indirect costs in your budget, you must include a copy of your indirect cost rate agreement. If your indirect cost rate is a provisional rate, the agreement should be less than 12 months of age.

IV.6. Other Submission Requirements

LOI Submission Address: Submit your LOI by express mail, delivery service, fax, or E-mail to: Mary Lerchen, DrPH, Scientific Review Administrator, CDC/Office of Public Health Research, One West Court Square, Suite 7000, MS D–72, Telephone: 404–371–5277, Fax:

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Applications may not be submitted electronically at this time.

V. Application Review Information

V.1. Criteria

Applicants are required to provide measures of effectiveness that will demonstrate the accomplishment of the various identified objectives of the cooperative agreement. Measures of effectiveness must relate to the performance goals stated in the "Purpose" section of this announcement. Measures must be objective and quantitative, and must measure the intended outcome. These measures of effectiveness must be submitted with the application and will be an element of evaluation.

The goals of CDC-supported research are to advance the understanding of biological systems, improve the control and prevention of disease and injury, and enhance health. In the written comments, reviewers will be asked to evaluate the application in order to judge the likelihood that the proposed research will have a substantial impact on the pursuit of these goals.

The scientific review group will address and consider each of the following criteria equally in assigning the application’s overall score, weighting them as appropriate for each application. The application does not need to be strong in all categories to be judged likely to have major scientific impact and thus deserve a high priority score. For example, an investigator may propose to carry out important work that by its nature is not innovative, but is essential to move a field forward.

The review criteria are as follows:

Significance: Does this study address an important problem? If the aims of the application are achieved, how will scientific knowledge be advanced? What will be the effect of these studies on the concepts or methods that drive this field?

Approach: Are the conceptual framework, design, methods, and analyses adequately developed, well-integrated, and appropriate to the aims of the project? Does the applicant acknowledge potential problem areas and consider alternative tactics? Are disparities in immunization rates documented and significant?

Applicants must document the targeted community has statistically significant disparities in immunization rates between black and children of other racial/ethnic groups for children 19–35 months of age. Documentation of population should be placed behind the application face page.

Innovation: Does the project employ novel concepts, approaches or methods? Are the aims original and innovative? Does the project challenge existing paradigms or develop new methodologies or technologies?

Investigator: Is the investigator appropriately trained and well suited to carry out this work? Is the work proposed appropriate to the experience level of the principal investigator and other researchers (if any)?

Environment: Does the scientific environment in which the work will be done contribute to the probability of success? Do the proposed experiments take advantage of unique features of the scientific environment or employ useful collaborative arrangements? Is there evidence of institutional support? Are letters of support included, if appropriate?

Additional Review Criteria: In addition to the above criteria, the following items will be considered in the determination of scientific merit and priority score:

Preference will be given to communities with greater disparities in immunization rates as evidenced by National Immunization Survey data or other indicators. These communities are frequently located in the Northeastern United States.

Protection of Human Subjects from Research Risks: Does the application adequately address the requirements of Title 45 Part 46 for the protection of human subjects? The involvement of human subjects and protections from research risk relating to their participation in the proposed research will be assessed.

Inclusion of Women and Minorities in Research: Does the application adequately address the CDC Policy requirements regarding the inclusion of women, ethnic, and racial groups in the proposed research? This includes: (1) The proposed plan for the inclusion of both sexes and racial and ethnic minority populations for appropriate...
representation; (2) The proposed justification when representation is limited or absent; (3) A statement as to whether the design of the study is adequate to measure differences when warranted; and (4) A statement as to whether the plans for recruitment and outreach for study participants include the process of establishing partnerships with community(ies) and recognition of mutual benefits.

**Budget:** The reasonableness of the proposed budget and the requested period of support in relation to the proposed research. The priority score should not be affected by the evaluation of the budget.

**V.2. Review and Selection Process**

Applications will be reviewed for completeness by the Procurement and Grants Office (PGO) and for responsiveness by the OPHR. Incomplete applications and applications that are non-responsive to the eligibility criteria will not advance through the review process. Applicants will be notified that their application did not meet submission requirements.

Applications that are complete and responsive to the announcement will be evaluated for scientific and technical merit by an appropriate peer review group or charter study section, a Special Emphasis Panel (SEP), convened by the OPHR in accordance with the review criteria listed above. As part of the initial merit review, all applications will:

- Undergo a process in which only those applications deemed to have the highest scientific merit by the review group, generally the top half of the applications under review, will be discussed and assigned a priority score.
- Receive a written critique.
- Receive a second programmatic level review by the Office of Science, National Immunization Program.
- Undergo a peer review by a Special Emphasis Panel (SEP). The SEP will be selected from the National Institutes of Health (NIH) pool of scientists or recommendations from the NIP to serve as reviewers on SEPs. Applications will be ranked for the secondary review according to scores submitted by the SEP. Only those applications deemed to have the highest scientific merit by the review group, generally the top half of the applications under review, will be discussed and assigned a priority score.

**Award Criteria:** Criteria that will be used to make award decisions during the programmatic review include:

- Scientific merit (as determined by peer review).
- Availability of funds.
- Programmatic priorities.
- Disparities in immunization rates.

**V.3. Anticipated Announcement and Award Dates**

**Award Date:** August 31, 2005.

**VI. Award Administration Information**

**VI.1. Award Notices**

Successful applicants will receive a Notice of Award (NoA) from the CDC Procurement and Grants Office. The NoA shall be the only binding, authorizing document between the recipient and CDC. The NoA will be signed by an authorized Grants Management Officer, and mailed to the recipient fiscal officer identified in the application.

Unsuccessful applicants will receive notification of the results of the application review by mail from the Scientific Review Administrator.

**VI.2. Administrative and National Policy Requirements**

45 CFR Part 74 and Part 92

For more information on the Code of Federal Regulations, see the National Archives and Records Administration at the following Internet address: http://www.access.gpo.gov/nara/cfr/cfr-table-search.html.

The following additional requirements apply to this project:

- AR–1 Human Subjects Requirements.
- AR–2 Requirements for Inclusion of Women and Racial and Ethnic Minorities in Research.
- AR–7 Executive Order 12372.
- AR–10 Smoke-Free Workplace Requirements.
- AR–11 Healthy People 2010.
- AR–12 Lobbying Restrictions.
- AR–15 Proof of Non-Profit Status.
- AR–22 Research Integrity.

Additional information on these requirements can be found on the CDC Web site at the following Internet address: http://www.cdc.gov/od/pgo/funding/ARIs.htm.

**VI.3. Reporting**

You must provide CDC with an original, plus two hard copies of the following reports:

1. Interim progress report, (use form PHS 2390, OMB Number 0925–0001, rev. 9/2004 as posted on the CDC website) no less than 90 days before the end of the budget period. The progress report will serve as your non-competing continuation application, and must contain the following additional elements:
   - b. Additional Information Requested by Program.
   - 2. Financial status report, no more than 90 days after the end of the budget period.
   - 3. Final financial and performance reports, no more than 90 days after the end of the project period.

These reports must be mailed to the Grants Management Specialist listed in the “Agency Contacts” section of this announcement.

**VII. Agency Contacts**

We encourage inquiries concerning this announcement.

For general questions, contact: Technical Information Management Section, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341; Telephone: (770) 488–2700.

For scientific/research issues, contact: Susan Chu, PhD, MSPH, Extramural Program Official, Centers for Disease Control and Prevention, National Immunization Program, MS E–05, 1600 Clifton Road NE., Atlanta, GA 30333, Telephone: (404) 639–8727; E-mail: SChu@cdc.gov.

For questions about peer review, contact: Mary Lerchen, DrPH, Scientific Review Administrator, CDC/Office of Public Health Research, One West Court Square, Suite 7000, MS D–72, Telephone: 404–371–5277, Fax: 404–371–5215; E-mail: Mlerchen@cdc.gov.

For financial, grants management, or budget assistance, contact: Peaches Brown, Grants Management Specialist, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341, Telephone: (770) 488–2738; E-mail: POBrown@cdc.gov.

**VIII. Other Information**

This and other CDC funding opportunity announcements can be found on the CDC Web site, Internet address: http://www.cdc.gov. Click on “Funding” then “Grants and Cooperative Agreements.”


**William P. Nichols,**

Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 05–9364 Filed 5–10–05; 8:45 am]