9. A summary of multilateral and U.S. bilateral assistance to the countries of sub-Saharan Africa.

The USTR requested that the Commission provide its first report by December 2000, and annually for a period of 4 years thereafter. The second report in the series was delivered to USTR on December 10, 2001. The third report shall be delivered no later than December 10, 2002. The 48 countries of sub-Saharan Africa covered in this investigation include: Angola, Benin, Botswana, Burkina Faso, Burundi, Cameroon, Cape Verde, Central African Republic, Chad, Comoros, Democratic Republic of the Congo, Côte d'Ivoire, Djibouti, Equatorial Guinea, Eritrea, Ethiopia, Gabon, The Gambia, Ghana, Guinea, Guinea-Bissau, Kenya, Lesotho, Liberia, Madagascar, Malawi, Mali, Mauritania, Mauritius, Mozambique, Namibia, Niger, Nigeria, Republic of the Congo, Rwanda, São Tomé and Principe, Senegal, Seychelles, Sierra Leone, Somalia, South Africa, Sudan, Swaziland, Tanzania, Togo, Uganda, Zambia, and Zimbabwe.

Written Submissions: The Commission does not plan to hold a public hearing in connection with this third report. However, interested persons are invited to submit written statements concerning matters to be addressed in the report. Commercial or financial information that a person desires the Commission to treat as confidential must be submitted on separate sheets of paper, each clearly marked ''Confidential Business Information'' at the top. The Commission's Rules do not authorize filing of submissions with the Secretary by facsimile or electronic means.

All written submissions must conform with the provisions of section 201.8 of the Commission's Rules of Practice and Procedure (19 C.F.R 201.8). All submissions requesting confidential treatment must conform with the requirements of section 201.6 of the Commission's Rules (19 CFR 201.6). All written statements, except for confidential business information will be made available for inspection by interested persons in the Office of the Secretary to the Commission. To be assured of consideration, written statements relating to the Commission's report should be submitted at the earliest possible date and should be received not later than July 19, 2002. All submissions should be addressed to the Secretary, United States International Trade Commission, 500 E Street, SW, Washington D.C. 20436. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000.

By order of the Commission.

Issued: May 2, 2002.

Marilyn R. Abbott,

Secretary.

[FR Doc. 02–11285 Filed 5–6–02; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated July 13, 2001, and published in the **Federal Register** on July 23, 2001, (66 FR 38323), Research Triangle Institute, Kenneth H. Davis, Jr., Hermann Building, East Institute Drive, PO Box 12194, Research Triangle Park, North Carolina 27709, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Marihuana (7360)	I
Cocaine (9041)	II

The institute will manufacture small quantities of cocaine derivatives and marihuana derivatives for use by their customers primarily in analytical kits, reagents and standards.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Research Triangle Institute to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Research Triangle Institute on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: April 18, 2002.

Laura M. Nagel,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 02–11304 Filed 5–6–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Office of Justice Programs

Agency Information Collection Activities: Proposed collection; comments requested

ACTION: 30-Day Notice of Information Collection Under Review: Reinstatement, with change, of a previously approved collection for which approval has expired, Police Public Contact Survey.

The Department of Justice (DOJ), Office of Justice Programs has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection is published to obtain comments from the public and affected agencies. This proposed information collection was previously published in the Federal Register Volume 67, Number 33, page 7379 on February 19, 2002, allowing for a 60 day comment period. The purpose of this notice is to allow for an additional 30 days for public comment until June 6, 2002. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to The Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503. Additionally, comments may be submitted to OMB via facsimile to (202)–395–7285.

Request written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agencies estimate of the burden of the