

investigated Cambridge Isotope Lab to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, 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(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: January 18, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011-2295 Filed 2-1-11; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 3, 2010, and published in the Federal Register on September 1, 2010, (75 FR 53720), American Radiolabeled Chemicals, Inc., 101 Arc Drive, St. Louis, Missouri 63146, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

| Drug | Schedule |
|--|----------|
| Gamma Hydroxybutyric Acid (2010) | I |
| Ibogaine (7260) | I |
| Lysergic acid diethylamide (7315) | I |
| Tetrahydrocannabinols (7370) | I |
| Dimethyltryptamine (7435) | I |
| 1-[1-(2-Thienyl)cyclohexyl]piperidine (7470) | I |
| Dihydromorphine (9145) | I |
| Normorphine (9313) | I |
| Amphetamine (1100) | II |
| Methamphetamine (1105) | II |
| Amobarbital (2125) | II |
| Phencyclidine (7471) | II |
| Phenylacetone (8501) | II |
| Cocaine (9041) | II |
| Codeine (9050) | II |
| Dihydrocodeine (9120) | II |
| Oxycodone (9143) | II |
| Hydromorphone (9150) | II |
| Ecgonine (9180) | II |
| Hydrocodone (9193) | II |
| Meperidine (9230) | II |
| Metazocine (9240) | II |
| Dextropropoxyphene, bulk (non-dosage forms) (9273) | II |
| Morphine (9300) | II |

| Drug | Schedule |
|--------------------|----------|
| Oripavine (9330) | II |
| Thebaine (9333) | II |
| Oxymorphone (9652) | II |
| Phenazocine (9715) | II |
| Fentanyl (9801) | II |

The company plans to manufacture small quantities of the listed controlled substances as radiolabeled compounds for biochemical research.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of American Radiolabeled Chemicals, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated American Radiolabeled Chemicals Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, 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(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: January 18, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011-2294 Filed 2-1-11; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 2, 2010, and published in the Federal Register on September 1, 2010, 75 FR 53720, Cody Laboratories, 601 Yellowstone Avenue, Cody, Wyoming 82414, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

| Drug | Schedule |
|------------------------|----------|
| Dihydromorphine (9145) | I |
| Amphetamine (1100) | II |
| Methamphetamine (1105) | II |
| Amobarbital (2125) | II |
| Pentobarbital (2270) | II |

| Drug | Schedule |
|-----------------------|----------|
| Secobarbital (2315) | II |
| Phenylacetone (8501) | II |
| Cocaine (9041) | II |
| Codeine (9050) | II |
| Dihydrocodeine (9120) | II |
| Oxycodone (9143) | II |
| Hydromorphone (9150) | II |
| Diphenoxylate (9170) | II |
| Ecgonine (9180) | II |
| Hydrocodone (9193) | II |
| Meperidine (9230) | II |
| Methadone (9250) | II |
| Morphine (9300) | II |
| Oxymorphone (9652) | II |
| Alfentanil (9737) | II |
| Remifentanil (9739) | II |
| Sufentanil (9740) | II |
| Fentanyl (9801) | II |

The company plans on manufacturing the listed controlled substances in bulk for sale to its customers.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Cody Laboratories to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Cody Laboratories to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, 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(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: January 18, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011-2291 Filed 2-1-11; 8:45 am]

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DEPARTMENT OF JUSTICE

Federal Bureau of Investigation

[Docket No. FBI 150]

FBI Records Management Division; National Name Check Program Section; New User Fees Schedule

AGENCY: Federal Bureau of Investigation (FBI), Justice.

ACTION: Notice.

SUMMARY: Pursuant to 28 CFR 20.31(e)(3), this notice establishes a new

user fee schedule for Federal agencies requesting name-based background checks of the FBI's Central Records System through the National Name Check Program for noncriminal justice purposes. The total resource costs associated with providing these name check services have been calculated to ensure full reimbursement to the FBI.

DATES: This fee schedule is effective March 4, 2011.

FOR FURTHER INFORMATION CONTACT: FBI, RMD, National Name Check Program Section, 170 Marcel Drive, Winchester, Virginia 22602, Attention: Michael Cannon, (540) 868-4400.

SUPPLEMENTARY INFORMATION:

Pursuant to the authority in Public Law 101-515, as amended, the FBI has established user fees for Federal agencies requesting noncriminal name-based background checks of the Central Records System (CRS) through the

National Name Check Program (NNCP) of the Records Management Division (RMD). The regulations governing the revision of these user fees are set out at 28 CFR 20.31(e and f). In accordance with 28 CFR 20.31(e), the FBI is required to periodically review the amount of the fees it collects for the NNCP to determine the current cost of processing name checks for noncriminal justice purposes and publish any resulting fee adjustments in the **Federal Register**.

Accordingly, the FBI conducted a fee study to assess the proper fee amounts that should be collected by the FBI.

In accordance with 28 CFR 20.31(e)(2), the fee study employed the same Activity Based Cost (ABC) accounting method detailed in the Final Rule establishing the process for setting fees (75 FR 24796 (May 6, 2010)). The ABC methodology is consistent with widely accepted accounting principles

and complies with the provisions of 31 U.S.C. 9701 and other applicable Federal law. The fee study identified all direct and indirect costs associated with the name-based background checks incurred by the FBI in fiscal year 2009. These costs were analyzed by the ABC model to project the total reimbursable costs, by fee category, for fiscal year 2011.

The fee study recommended several adjustments to the current user fees, which have been in effect since October 1, 2007. Pursuant to the fee study, the fees imposed for electronic submissions will be increased, while the fees for manual and expedited submissions will be decreased. The following table details the fee amounts for Federal agencies requesting name-based background checks of the FBI's CRS through the NNCP for noncriminal justice purposes.

| Service | Fee currently in effect | Change in fee amount | Revised fee |
|----------------------------|-------------------------|----------------------|-------------|
| Electronic Submission: | | | |
| Batch Process Only | \$1.50 | \$0.50 | \$2.00 |
| Batch + File Review | 29.50 | 9.00 | 38.50 |
| Manual Submission | 56.00 | (5.25) | 50.75 |
| Expedited Submission | 56.00 | (5.25) | 50.75 |

This new fee schedule will become effective on March 4, 2011.

Dated: January 25, 2011.

Robert S. Mueller, III,

Director, Federal Bureau of Investigation.

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DEPARTMENT OF LABOR

Employment and Training Administration

[TA-W-74,689]

Amdocs, Inc., Global Support Services, Advertising And Media AT&T Division, New Haven, Connecticut; Notice of Affirmative Determination Regarding Application for Reconsideration

By application dated December 22, 2010, legal counsel of a member of the subject worker group requested administrative reconsideration of the negative determination regarding workers' eligibility to apply for Trade Adjustment Assistance (TAA) applicable to workers and former workers of Amdocs, Inc., Global Support Services, Advertising and Media AT&T Division, New Haven, Connecticut (subject firm). The negative determination was issued on November

9, 2010. The Notice of determination was published in the **Federal Register** on November 23, 2010 (75 FR 71461).

The negative determination was based on the findings that the worker separations are not attributable to increased imports or a shift of services by the workers' firm. Specifically, services shifted to a foreign country by Amdocs, Inc. did not contribute importantly to worker separations in Global Support Services, Advertising and Media AT&T Division.

The investigation also revealed that the firm is not a Supplier or Downstream Producer to a firm with a TAA-certified worker group.

In the request for reconsideration, the petitioner alleged that the workers of the Advertising and Media Division are eligible to apply for TAA because Section 222(a) and/or Section 222(c) of the Trade Act of 1974, as amended, has been met.

The Department has carefully reviewed the request for reconsideration and the existing record, and has determined that the Department will conduct further investigation to determine if the petitioning workers meet the eligibility requirements of the Trade Act of 1974, as amended.

Conclusion

After careful review of the application, I conclude that the claim is of sufficient weight to justify reconsideration of the U.S. Department of Labor's prior decision. The application is, therefore, granted.

Signed at Washington, DC, this 21st day of January, 2011.

Del Min Amy Chen,

Certifying Officer, Office of Trade Adjustment Assistance.

[FR Doc. 2011-2241 Filed 2-1-11; 8:45 am]

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DEPARTMENT OF LABOR

Employment and Training Administration

[TA-W-74,700]

AT&T; Reynoldsburg, OH; Notice of Affirmative Determination Regarding Application for Reconsideration

By application dated January 6, 2011, by three petitioners requested administrative reconsideration of the negative determination regarding workers' eligibility to apply for Trade Adjustment Assistance (TAA) applicable to workers and former workers of AT&T, Reynoldsburg, Ohio