Additional written submissions to the Commission, including requests pursuant to section 201.12 of the Commission's rules, shall not be accepted unless good cause is shown for accepting such submissions, or unless the submission is pursuant to a specific request by a Commissioner or Commission staff.

In accordance with sections 201.16(c) and 207.3 of the Commission's rules, each document filed by a party to the investigations must be served on all other parties to the investigations (as identified by either the public or BPI service list), and a certificate of service must be timely filed. The Secretary will not accept a document for filing without a certificate of service.

Authority: These investigations are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.21 of the Commission's rules.

By order of the Commission.

Issued: August 15, 2012.

Lisa R. Barton,

Acting Secretary to the Commission.

[FR Doc. 2012–20372 Filed 8–17–12; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; SA INTL GMBH C/O., Sigma Aldrich Co., LLC

Correction

In notice document 2012–19191 appearing on pages 47106–47108 in the issue of Tuesday, August 7, 2012, make the following corrections:

1. On page 47106, in the third column, the document heading should appear as set forth above.

2. On page 4707, in the sixth paragraph following the table, in the eighth line of text, ''September 6, 2012'' should read ''September 19, 2012''.

[FR Doc. C1–2012–19191 Filed 8–17–12; 8:45 am]

BILLING CODE 1505–01–D

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; Clinical Supplies Management, Inc.

Correction

In notice document 2012–19197 appearing on pages 47109–47110 in the issue of Tuesday, August 7, 2012, make the following corrections:

1. On page 47109, in the third column, the document heading should appear as set forth above.

2. On page 47110, in the second column, in the first full paragraph, in the eighth line of text, ''September 6, 2012'' should read ''September 19, 2012''.

[FR Doc. C1–2012–19197 Filed 8–17–12; 8:45 am]

BILLING CODE 1505–01–D

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; Lipomed

Correction

In notice document 2012–19196 appearing on page 47108 in the issue of Tuesday, August 7, 2012, make the following corrections:

1. On page 47108, in the first column, the document heading should appear as set forth above.

2. On page 47108, in the second column, in the second full paragraph, in the eighth line of text, ''September 6, 2012'' should read ''September 19, 2012''.

[FR Doc. C1–2012–19196 Filed 8–17–12; 8:45 am]

BILLING CODE 1505–01–D

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; R & D Systems, Inc.

Correction

In notice document 2012–19193 appearing on pages 47110–47111 in the issue of Tuesday, August 7, 2012, make the following corrections:

1. On page 47110, in the third column, the document heading should appear as set forth above.

2. On page 47110, in the fifth paragraph following the table, in the eighth line of text, ''September 6, 2012'' should read ''September 19, 2012''.

[FR Doc. C1–2012–19193 Filed 8–17–12; 8:45 am]

BILLING CODE 1505–01–D

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; Cerilliant Corporation

Correction

In notice document 2012–19199 appearing on pages 47108–47109 in the issue of Tuesday, August 7, 2012, make the following corrections:

1. On page 47108, in the third column, the document heading should appear as set forth above.

2. On page 47109, in the sixth paragraph following the table, in the eighth line of text, ''September 6, 2012'' should read ''September 19, 2012''.

[FR Doc. C1–2012–19199 Filed 8–17–12; 8:45 am]

BILLING CODE 1505–01–D

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; Almac Clinical Services, Inc.

By Notice dated April 17, 2012, and published in the **Federal Register** on April 26, 2012, 77 FR 24985, Almac Clinical Services, Inc., (ACSI), 25 Fretz Road, Souderton, Pennsylvania 18964, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

Drug	Schedule
Oxycodone (9143)	II
Hydromorphone (9150)	II
Tapentadol (9780)	II
Fentanyl (9801)	II

The company plans to import small quantities of the listed controlled substances in dosage form to conduct clinical trials.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of Almac Clinical Services, Inc. (ACSI) to import the basic classes of controlled substances is consistent with the public interest, and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. DEA has investigated Almac Clinical Services, Inc. (ACSI) 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic classes of controlled substances listed.

Dated: August 7, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–20370 Filed 8–17–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; Research Triangle Institute

Correction

In notice document 2012–19208 appearing on pages 47111–47114 in the issue of Tuesday, August 7, 2012, make the following correction:

On page 47111, in the second column, the document heading should appear as set forth above.

[FR Doc. C1–2012–19208 Filed 8–17–12; 8:45 am]

BILLING CODE 1505–01–D

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; Catalent Pharma Solutions, Inc.

Correction

In notice document 2012–19202 appearing on page 47114 in the issue of Tuesday, August 7, 2012, make the following correction:

On page 47114, in the first column, the document heading should appear as set forth above.

[FR Doc. C1–2012–19202 Filed 8–17–12; 8:45 am]

BILLING CODE 1505–01–D

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances, Notice of Registration, Cody Laboratories, Inc.

By Notice dated March 8, 2012, and published in the **Federal Register** on March 20, 2012, 77 FR 16263, Cody Laboratories, Inc., 601 Yellowstone Avenue, Cody, Wyoming 82414, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
4-Anilino-N-phenethyl-4-piperidine (8333).	II
Thebaine (9333)	II

The company plans on manufacturing the listed controlled substances as bulk intermediates for distribution 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, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Cody Laboratories, 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, 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: August 7, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–20369 Filed 8–17–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Mine Safety and Health Administration

Affirmative Decisions on Petitions for Modification Granted in Whole or in Part

AGENCY: Mine Safety and Health Administration (MSHA), Labor.

ACTION: Notice.

SUMMARY: Section 101(c) of the Federal Mine Safety and Health Act of 1977 and 30 CFR part 44 govern the application, processing, and disposition of petitions for modification. This **Federal Register** Notice notifies the public that MSHA has investigated and issued a final decision on certain mine operator petitions to modify a safety standard.

ADDRESSES: Copies of the final decisions are posted on MSHA's Web Site at *http://www.msha.gov/indexes/petition.htm.* The public may inspect the petitions and final decisions during normal business hours in MSHA's Office of Standards, Regulations and Variances, 1100 Wilson Boulevard, Room 2349, Arlington, Virginia 22209. All visitors must first stop at the receptionist desk on the 21st Floor to sign-in.

FOR FURTHER INFORMATION CONTACT: Roslyn B. Fontaine, Office of Standards, Regulations and Variances at 202–693–9475 (Voice), *fontaine.roslyn@dol.gov* (Email), or 202–693–9441 (Telefax), or Barbara Barron at 202–693–9447 (Voice), *barron.barbara@dol.gov* (Email), or 202–693–9441 (Telefax). [These are not toll-free numbers].

SUPPLEMENTARY INFORMATION:

I. Introduction

Under section 101 of the Federal Mine Safety and Health Act of 1977, a mine operator may petition and the Secretary of Labor (Secretary) may modify the application of a mandatory safety standard to that mine if the Secretary determines that: (1) an alternative method exists that will guarantee no less protection for the miners affected than that provided by the standard; or (2) that the application of the standard will result in a diminution of safety to the affected miners.

MSHA bases the final decision on the petitioner's statements, any comments and information submitted by interested persons, and a field investigation of the conditions at the mine. In some instances, MSHA may approve a petition for modification on the condition that the mine operator complies with other requirements noted in the decision.

II. Granted Petitions for Modification

On the basis of the findings of MSHA's investigation, and as designee of the Secretary, MSHA has granted or partially granted the following petitions for modification:

- *Docket Number:* M–2009–050–C.
  FR Notice: 75 FR 3256 (1/20/2010).
  Petitioner: Wolf Run Mining Company, 300 Corporate Centre Drive, Scott Depot, West Virginia 25560.
  Mine: Sentinel Mine, MSHA I.D. No. 46–04168, located in Barbour County, West Virginia.
  Regulation Affected: 30 CFR 75.1700 (Oil and gas wells).
- *Docket Number:* M–2009–052–C.
  FR Notice: 75 FR 3257 (1/20/2010).