The overarching purpose of the Hospital Data Abstraction Form, formerly entitled Evaluation of Emergency Department Crisis Center Follow-up, is to examine the impact of crisis center follow-up with patients admitted to emergency departments or inpatient behavioral health units following a suicide attempt or serious suicidal ideation on subsequent readmissions for suicidal behavior. This effort assesses the capacity of follow-up to save both lives and critical hospital resources. This evaluation effort includes one data collection activity. Clearance is being requested for the continuation and expansion of the already-approved abstraction form of hospital data on patients admitted to emergency departments or inpatient behavioral health units following a suicide attempt or serious ideation. This effort will continue to examine the impact of crisis center follow-up on readmissions for suicidal behavior. The data collected through this project will ultimately help SAMHSA to understand and direct crisis center follow-up lifesaving initiatives. The data collection activity is described below.

Hospitals collaborating with two cohorts (cohorts IV and V) of Lifeline crisis centers will participate in this expanded initiative. Fifteen hospitals per cohort will participate for a total of 30. Patient data will be collected for patients admitted for a suicide attempt in the two years prior to collaboration between the hospital and crisis center and for patients admitted for a suicide attempt for the two-year period after collaboration.

The Hospital Data Abstraction Form will be utilized to collect systematic patient data for patients seen in the 30 participating hospitals' emergency departments or inpatient behavioral health units. Information to be abstracted from patient data include: Demographic data, historical data, and subsequent suicidal behavioral and admission data. Data will be de-identified. Hospital staff will review patient data for qualifying (*i.e.,* admission to the emergency department for suicide attempt) records. Records to be reviewed will include emergency department or inpatient behavioral health unit admissions for the two years prior to crisis center and hospital collaboration and for two years following collaboration. It is expected that a total of 30,000 records will be abstracted by hospital staff and provided to the evaluation team.

This revision involves an increase in the number of participating hospital respondents and burden associated with the continuation/expansion of the already-approved Hospital Data Abstraction Form (OMB No. 0930–0337; Expiration 09/30/2016), as well as the discontinuation of data collection and burden associated with the Crisis Center Data Abstraction Form.

The estimated response burden to collect this information is as follows annualized over the requested three-year clearance period is presented below:

TOTAL AND ANNUALIZED AVERAGES: RESPONDENTS, RESPONSES, AND HOURS

Instrument	Number of respondents	Responses per respondent *	Total number of responses	Burden per response	Annual burden *
Hospital Data Abstraction Form	30	334	10,020	.04	401

* Rounded to the nearest whole number

Written comments and recommendations concerning the proposed information collection should be sent by October 26, 2015 to the SAMHSA Desk Officer at the Office of Information and Regulatory Affairs, Office of Management and Budget (OMB). To ensure timely receipt of comments, and to avoid potential delays in OMB's receipt and processing of mail sent through the U.S. Postal Service, commenters are encouraged to submit their comments to OMB via email to: *OIRA_Submission@omb.eop.gov*. Although commenters are encouraged to send their comments via email, commenters may also fax their comments to: 202–395–7285. Commenters may also mail them to: Office of Management and Budget, Office of Information and Regulatory Affairs, New Executive Office Building, Room 10102, Washington, DC 20503.

Summer King,

Statistician.

[FR Doc. 2015–24290 Filed 9–23–15; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Accreditation and Approval of Amspec Services, LLC, as a Commercial Gauger and Laboratory

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security.

ACTION: Notice of accreditation and approval of AmSpec Services, LLC, as a commercial gauger and laboratory.

SUMMARY: Notice is hereby given, pursuant to CBP regulations, that AmSpec Services, LLC, has been approved to gauge petroleum and certain petroleum products and accredited to test petroleum and certain petroleum products for customs purposes for the next three years as of April 29, 2015.

DATES: *Effective Dates:* The accreditation and approval of AmSpec Services, LLC, as commercial gauger and laboratory became effective on April 29, 2015. The next triennial inspection date will be scheduled for April 2018.

FOR FURTHER INFORMATION CONTACT: Approved Gauger and Accredited Laboratories Manager, Laboratories and Scientific Services Directorate, U.S. Customs and Border Protection, 1300 Pennsylvania Avenue NW., Suite 1500N, Washington, DC 20229, tel. 202–344–1060.

SUPPLEMENTARY INFORMATION: Notice is hereby given pursuant to 19 CFR 151.12 and 19 CFR 151.13, that AmSpec Services, LLC, 100 Wheeler St., Unit G, New Haven, CT 06512, has been approved to gauge petroleum and certain petroleum products and accredited to test petroleum and certain petroleum products for customs purposes, in accordance with the provisions of 19 CFR 151.12 and 19 CFR 151.13. AmSpec Services, LLC is approved for the following gauging procedures for petroleum and certain petroleum products from the American Petroleum Institute (API):

API chapters	Title
1	Vocabulary.
3	Tank Gauging.
7	Temperature Determination.
8	Sampling.
12	Calculations.

API chapters	Title
17	Maritime Measurement.

AmSpec Services, LLC is accredited for the following laboratory analysis procedures and methods for petroleum and certain petroleum products set forth by the U.S. Customs and Border Protection Laboratory Methods (CBPL) and American Society for Testing and Materials (ASTM):

CBPL No.	ASTM	Title
27–01	D287	Standard Test Method for API Gravity of crude Petroleum and Petroleum Products.
27–02	D1298	Standard Practice for Density, Relative Density (Specific Gravity), or API Gravity of Crude Petroleum and Liquid Petroleum Products by Hydrometer Meter.
27–04	D95	Standard Test Method for Water in Petroleum Products and Bituminous Materials by Distillation.
27–05	D4928	Standard Test Method for Water in Crude Oils by Coulometric Karl Fischer Titration.
27–06	D473	Standard Test Method for Sediment in Crude Oils and Fuel Oils by the Extraction Method.
27–08	D86	Standard Test Method for Distillation of Petroleum Products.
27–11	D445	Standard Test Method for Kinematic Viscosity of Transparent and Opaque Liquids.
27–13	D4294	Standard Test Method for Sulfur in Petroleum and Petroleum Products by Energy-Dispersive X-ray Fluorescence Spectrometry
27–20	D4057	Standard Practice for Manual Sampling of Petroleum and Petroleum Products.
27–48	D4052	Standard Test Method for Density and Relative Density of Liquids by Digital Density Meter.
27–50	D93	Standard Test Methods for Flash-Point by Pensky-Martens Closed Cup Tester.
27–53	D2709	Standard Test Method for Water and Sediment in Middle Distillate Fuels by Centrifuge.
27–54	D1796	Standard Test Method for Water and Sediment in Fuel Oils by the Centrifuge Method.
27–58	D5191	Standard Test Method For Vapor Pressure of Petroleum Products.
Pending	D97	Standard Test Method for Pour Point of Petroleum Products.
Pending	D2500	Standard Test Method for Cloud Point of Petroleum Products.

Anyone wishing to employ this entity to conduct laboratory analyses and gauger services should request and receive written assurances from the entity that it is accredited or approved by the U.S. Customs and Border Protection to conduct the specific test or gauger service requested. Alternatively, inquiries regarding the specific test or gauger service this entity is accredited or approved to perform may be directed to the U.S. Customs and Border Protection by calling (202) 344–1060. The inquiry may also be sent to *CBPGaugersLabs@cbp.dhs.gov*. Please reference the Web site listed below for a complete listing of CBP approved gaugers and accredited laboratories.

http://www.cbp.gov/about/labs-scientific/commercial-gaugers-and-laboratories

Dated: September 10, 2015.

Ira S. Reese,

Executive Director, Laboratories and Scientific Services Directorate.

[FR Doc. 2015–24229 Filed 9–23–15; 8:45 am]

BILLING CODE 9111–14–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Notice of Issuance of Final Determination Concerning Certain Analytical-Grade Acetonitrile

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security.

ACTION: Notice of final determination.

SUMMARY: This document provides notice that U.S. Customs and Border Protection (''CBP'') has issued a final determination concerning the country of origin of certain analytical-grade acetonitrile. Based upon the facts presented, CBP has concluded that the country of origin of the analytical-grade acetonitrile is the country of origin of the crude acetonitrile for purposes of U.S. Government procurement.

DATES: The final determination was issued on September 18, 2015. A copy of the final determination is attached. Any party-at-interest, as defined in 19 CFR 177.22(d), may seek judicial review of this final determination within October 26, 2015.

FOR FURTHER INFORMATION CONTACT: Ross Cunningham, Valuation and Special Programs Branch, Regulations and Rulings, Office of International Trade (202) 325–0034.

SUPPLEMENTARY INFORMATION: Notice is hereby given that on September 18, 2015 pursuant to subpart B of Part 177, U.S. Customs and Border Protection Regulations (19 CFR part 177, subpart B), CBP issued a final determination concerning the country of origin of certain analytical-grade acetonitrile, which may be offered to the U.S. Government under an undesignated government procurement contract. This final determination, HQ H265712, was issued under procedures set forth at 19 CFR part 177, subpart B, which implements Title III of the Trade Agreements Act of 1979, as amended (19 U.S.C. 2511–18). In the final determination, CBP concluded that the processing in the United States does not result in a substantial transformation. Therefore, the country of origin of the analytical-grade acetonitrile is the country of origin of the crude acetonitrile for purposes of U.S. Government procurement.

Section 177.29, CBP Regulations (19 CFR 177.29), provides that a notice of final determination shall be published in the **Federal Register** within 60 days of the date the final determination is issued. Section 177.30, CBP Regulations (19 CFR 177.30), provides that any party-at-interest, as defined in 19 CFR 177.22(d), may seek judicial review of a final determination within 30 days of publication of such determination in the **Federal Register**.

Dated: September 18, 2015.

Harold Singer,

Acting Executive Director, Regulations and Rulings, Office of International Trade.

HQ H265712

September 18, 2015

OT:RR:CTF:VS H265712 RMC

CATEGORY: Country of Origin

David R. Stepp

Bryan Cave LLP

120 Broadway, Suite 300, Santa Monica, CA 90401–2386

Re: U.S. Government Procurement; Country of Origin of Acetonitrile; Substantial Transformation

Dear Mr. Stepp: This is in response to your letter dated April 1, 2015, requesting a country-of-origin determination on behalf of the Sigma-Aldrich Corporation (''Sigma-Aldrich''). You state that Sigma-Aldrich wishes to sell ''analytical-grade acetonitrile'' to the U.S. Government and thus seeks a determination that the country of origin of its product will be the United States.