changes in market demand abroad). Demand conditions to consider include end uses and applications; the existence and availability of substitute products; and the level of competition among the *Domestic Like Product* produced in the United States, *Subject Merchandise* produced in the *Subject Country,* and such merchandise from other countries.

(13) (OPTIONAL) A statement of whether you agree with the above definitions of the *Domestic Like Product* and *Domestic Industry;* if you disagree with either or both of these definitions, please explain why and provide alternative definitions.

Authority: This review is being conducted under authority of Title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.61 of the Commission's rules.

Issued: June 25, 2013.

By order of the Commission.

William R. Bishop,

Supervisory Hearings and Information Officer.

[FR Doc. 2013–16103 Filed 7–2–13; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

[OMB Number 1103–NEW]

Agency Information Collection Activities; Proposed New Collection; Comments Requested: Police-Led Diversion Programs: National Prevalence and Scope

ACTION: 60-Day Notice.

The Department of Justice (DOJ) Office of Community Oriented Policing Services (COPS) will be submitting 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. The purpose of this notice is to allow for 60 days for public comment until September 3, 2013. This process is conducted in accordance with 5 CFR 1320.10.

If you have comments, especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Danielle Ouellette, Department of Justice Office of Community Oriented Policing Services, 145 N Street NE., Washington, DC 20530.

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:

—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;

—Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

—Enhance the quality, utility, and clarity of the information to be collected; and

—Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of This Information Collection

(1) *Type of Information Collection:* Proposed new collection; comments requested

(2) *Title of the Form/Collection:* Police-Led Diversion Programs: National Prevalence and Scope

(3) *Agency form number, if any, and the applicable component of the Department sponsoring the collection:* None. U.S. Department of Justice Office of Community Oriented Policing Services

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Law enforcement agencies through a nationally representative sample may be asked to provide information to determine the national prevalence of police-led diversion programs and provide a portrait of their goals, target populations, and policies. Through a cooperative agreement with the COPS Office, the Center for Court Innovation (CCI, Inc.) will create a representative sample of law enforcement agencies based on data available through the FBI Uniform Crime Reporting. CCI will subcontract with a professional survey research firm to administer the survey.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond/reply:* It is estimated that 3,600 respondents annually will complete the form in approximately 1 hour.

(6) *An estimate of the total public burden (in hours) associated with the collection:* There are an estimated 3,600 total annual burden hours associated with this collection.

If additional information is required contact: Jerri Murray, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE., Room 1407B, Washington, DC 20530.

June 27, 2013.

Jerri Murray,

Department Clearance Officer, PRA, U.S. Department of Justice.

[FR Doc. 2013–15904 Filed 7–2–13; 8:45 am]

BILLING CODE 4410–AT–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–372]

Exempt Chemical Preparations Under the Controlled Substances Act

AGENCY: Drug Enforcement Administration (DEA), Department of Justice.

ACTION: Order with opportunity for comment.

SUMMARY: The applications for exempt chemical preparations received by DEA between July 1, 2012, and March 31, 2013, as listed below, were accepted for filing and have been approved or denied as indicated.

DATES: Electronic comments must be submitted and written comments must be postmarked on or before September 3, 2013. The electronic Federal Docket Management System will not accept comments after midnight Eastern Time on the last day of the comment period.

ADDRESSES: To ensure proper handling of comments, please reference ''Docket No. DEA–372'' on all electronic and written correspondence. DEA encourages that all comments be submitted electronically through *http://www.regulations.gov* using the electronic comment form provided on that site. An electronic copy of this document is also available at *http://www.regulations.gov* for easy reference. Paper comments that duplicate the electronic submission are not necessary as all comments submitted to *www.regulations.gov* will be posted for public review and are part of the official docket record. Written comments submitted via regular or express mail should be sent to the Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODL, 8701 Morrissette Drive, Springfield, Virginia 22152.

FOR FURTHER INFORMATION CONTACT: John W. Partridge, Executive Assistant, Office of Diversion Control, Drug Enforcement Administration; Mailing Address: 8701 Morrissette Drive, Springfield, Virginia 22152; Telephone: (202) 307–7165.

SUPPLEMENTARY INFORMATION:

Posting of Public Comments

Please note that all comments received are considered part of the public record and made available for public inspection online at *http://www.regulations.gov* and in the DEA's public docket. Such information includes personal identifying information (such as your name, address, etc.) voluntarily submitted by the commenter.

If you want to submit personal identifying information (such as your name, address, etc.) as part of your comment, but do not want it to be posted online or made available in the public docket, you must include the phrase ''PERSONAL IDENTIFYING INFORMATION'' in the first paragraph of your comment. You must also place all the personal identifying information you do not want posted online or made available in the public docket in the first paragraph of your comment and identify what information you want redacted.

If you want to submit confidential business information as part of your comment, but do not want it to be posted online or made available in the public docket, you must include the phrase ''CONFIDENTIAL BUSINESS INFORMATION'' in the first paragraph of your comment. You must also prominently identify confidential business information to be redacted within the comment. If a comment has so much confidential business information that it cannot be effectively redacted, all or part of that comment may not be posted online or made available in the public docket.

Personal identifying information and confidential business information identified and located as set forth above will be redacted, and the comment, in redacted form, will be posted online and placed in DEA's public docket file. Please note that the Freedom of Information Act applies to all comments received. If you wish to inspect the agency's public docket file in person by appointment, please see the **FOR FURTHER INFORMATION CONTACT** paragraph.

Legal Authority

DEA implements and enforces Titles II and III of the Comprehensive Drug Abuse Prevention and Control Act of 1970, often referred to as the Controlled Substances Act and the Controlled Substances Import and Export Act (codified at Title 21, Chapter 13 of the U.S.C.), as amended (CSA). DEA drafts and publishes the implementing regulations for these statutes in Title 21 of the Code of Federal Regulations (CFR), parts 1300 to 1321. The CSA and its implementing regulations are designed to prevent, detect, and eliminate the diversion of controlled substances and listed chemicals into the illicit market while providing for a sufficient supply of controlled substances and listed chemicals for legitimate medical, scientific, research, and industrial purposes. Controlled substances have the potential for abuse and dependence and are controlled to protect the public health and safety.

Section 201 of the CSA (21 U.S.C. 811) authorizes the Attorney General, by regulation, to exempt from certain provisions of the CSA certain compounds, mixtures, or preparations containing a controlled substance, if he finds that such compounds, mixtures, or preparations meet the requirements detailed in 21 U.S.C. 811(g)(3)(B).[1] DEA regulations at 21 CFR 1308.23 and 1308.24 further detail the criteria by which the DEA Deputy Assistant Administrator may exempt a chemical preparation or mixture from the application of certain provisions of the CSA. The Deputy Assistant Administrator may, pursuant to 21 CFR 1308.23(f), modify or revoke the criteria by which exemptions are granted and modify the scope of exemptions at any time.

Exempt Chemical Preparation Applications Submitted Between July 1, 2012, and March 31, 2013

The Deputy Assistant Administrator received applications between July 1, 2012, and March 31, 2013, requesting exempt chemical preparation status pursuant to 21 CFR 1308.23. Pursuant to the criteria stated in 21 U.S.C. 811(g)(3)(B) and in 21 CFR 1308.23, the Deputy Assistant Administrator has found that each of the compounds, mixtures, and preparations described in Chart I below is intended for laboratory, industrial, educational, or special research purposes and not for general administration to a human being or other animal and either: (1) Contains no narcotic controlled substance and is packaged in such a form or concentration that the packaged quantity does not present any significant potential for abuse; or (2) contains either a narcotic or non-narcotic controlled substance and one or more adulterating or denaturing agents in such a manner, combination, quantity, proportion, or concentration that the preparation or mixture does not present any potential for abuse; if the preparation or mixture contains a narcotic controlled substance, it must be formulated in such a manner that it incorporates methods of denaturing or other means so that the preparation or mixture is not liable to be abused or have ill effects if abused, and so that the narcotic substance cannot in practice be removed.

Accordingly, pursuant to 21 U.S.C. 811(g)(3)(B), 21 CFR 1308.23, and 21 CFR 1308.24, the Deputy Assistant Administrator has determined that each of the chemical preparations or mixtures generally described in Chart I below and specifically described in the application materials received by DEA, are exempt, to the extent described in 21 CFR 1308.24, from application of sections 302, 303, 305, 306, 307, 308, 309, 1002, 1003, and 1004 (21 U.S.C. 822–823, 825–829, and 952–954) of the CSA, and 21 CFR 1301.74, as of the date listed below that was provided in the approval letters to the individual requesters.

CHART I

Supplier	Product name	Form	Application date
AccuStandard, Inc	APP–9–086, a,a-Dimethylphenethylamine	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	APP–9–086–20X, a,a-Dimethylphenethylamine	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	AS–E1179, Chloral hydrate	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	CLP–HC–X1, Composite Mix #3A	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	M–551.1B, Disinfectant by-products	Amber ampule: 1 mL	1/28/2013

[1] This authority has been delegated from the Attorney General to the Administrator of the DEA by 28 CFR 0.100 and subsequently redelegated to the Deputy Assistant Administrator pursuant to the Appendix to Subpart R of 28 CFR 0.104.

CHART I—Continued

Supplier	Product name	Form	Application date
AccuStandard, Inc	M–551B, Disinfectant by-products	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	M–551B–2, Chloral hydrate	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	M–8015B/5031–21, Paraldehyde	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	M–8206B–05, Additional VOC's by Method 8206B	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	M–8270–02, Semi-Volatile by Capillary Column GC/MS Mix 2	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	M–8270–04–ASL, Method 8270B—Base/Neutrals Mix	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	M–8270–23–R1, Method 8270 Semi-Volatile Additions	Amber ampule: 1 mL	1/28/2013
AccuStandard, Inc	M–E–1179–M, Chloral hydrate	Amber ampule: 1 mL	1/28/2013
Agilent Technologies	Veterinary Drugs Checkout Mix	Amber ampule: 1 mL	8/3/2012
Agilent Technologies	Veterinary Drugs Comprehensive Mix—Submix 6	Amber ampule: 1 mL	8/3/2012
Agilent Technologies	Forensic Toxicology Comprehensive Mix—Submix 3	Amber ampule: 1 mL	8/3/2012
Agilent Technologies	Forensic Toxicology Comprehensive Mix—Submix 5	Amber ampule: 1 mL	8/3/2012
Agilent Technologies	Forensic Toxicology Comprehensive Mix—Submix 2	Amber ampule: 1 mL	8/3/2012
Agilent Technologies	Forensic Toxicology Checkout Mix	Amber ampule: 1 mL	8/3/2012
Agilent Technologies	Forensic Toxicology Comprehensive Mix—Submix 1	Amber ampule: 1 mL	12/6/2012
Alltech Associates, Inc	2C–T–2 Quik Check, 1.0 mg/mL	Glass ampule: 1 mL	9/26/2012
Alltech Associates, Inc	5-MeO–DMT Quick Check, 1.0 mg/mL	Glass ampule: 1 mL	9/26/2012
Alltech Associates, Inc	Carisoprodol Quik Check, 1.0 mg/mL	Glass ampule: 1 mL	9/26/2012
American Proficiency Institute	American Proficiency Institute SHBG/Testosterone	Amber bottle: 7 mL	12/5/2012
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Oral Fluid, OF23	Glass vials: 1 ml–200 mL	8/14/2012
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Urine, SD17	Glass vials: 1 ml–200 mL	8/14/2012
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Urine, MC176	Glass vials: 1 ml–200 mL	9/27/2012
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Urine, MC179	Glass vials: 1 ml–200 mL	10/23/2012
Biochemical Diagnostics, Inc	Salivabuse Custom Liquid Control Oral Fluid, OF25	Glass vials: 1 ml–200 mL	10/23/2012
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Urine, MC177	Glass vials: 1 ml–200 mL	11/5/2012
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Urine, MC178	Glass vials: 1 ml–200 mL	1/24/2013
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Oral Fluid, OF24	Glass vials: 1 ml–200 mL	1/24/2013
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Urine, MC181	Glass vials: 1 ml–200 mL	1/29/2013
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Urine, MC183	Glass vials: 1 ml–200 mL	3/12/2013
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Urine, MC184	Glass vials: 1 ml–200 mL	3/12/2013
Biochemical Diagnostics, Inc	Detectabuse Custom Liquid Control Urine, MC180	Glass vials: 1 ml–200 mL	3/14/2013
Bio-Rad Laboratories	Liquichek Immunoassay Plus Control Trilevel MiniPak	Box: 3 vials, 2.5 mL each	11/14/2012
Bio-Rad Laboratories	Liquichek Immunoassay Plus Control Level 1,2 and 3, Trilevel.	Box: 12 vials, 5 mL each	11/21/2012
Bio-Rad Laboratories	Liquid Assayed Multiqual Premium, Levels 1–3	Box: 6 vials, 5 mL each	11/21/2012
Bio-Rad Laboratories	Liquid Assayed Multiqual Premium, Trilevel Minipak	Box: 3 vials, 5 mL each	11/21/2012
Bio-Rad Laboratories	Lyphocheck Urine Toxicology Control MiniPak, Cat. No. 470X.	Glass vial: 25 mL	1/11/2013
Bio-Rad Laboratories	Liquichek Immunoassay Plus Control Level 1	Glass vial: 2.5 mL, Box 12 vials.	2/20/2013
Bio-Rad Laboratories	Liquichek Immunoassay Plus Control Level 2	Glass vial: 2.5 mL, Box 12 vials.	2/20/2013
Bio-Rad Laboratories	Liquichek Immunoassay Plus Control Level 3	Glass vial: 2.5 mL, Box 12 vials.	2/20/2013
Cerilliant Corporation	Benzodiazepine Mix (1 mg/mL)	Glass Ampule: 1 mL	7/2/2012
Cerilliant Corporation	Class C Mix Urinalysis Internal Standard Fortified Morphine	Glass Ampule: 1 mL	7/9/2012
Cerilliant Corporation	Spice Cannabinoid Mix 2 (0.1 mg/mL)	Amber ampule: 1 mL	9/26/2012
Cerilliant Corporation	Spice Cannabinoid Mix 3 (0.1 mg/mL)	Amber ampule: 1 mL	9/26/2012
Cerilliant Corporation	Retigabine (1 mg/mL)	Glass Ampule: 1 mL	11/7/2012
Cerilliant Corporation	±-Norpseudoephedrine-D3 HCl (0.1 mg/mL)	Glass Ampule: 1 mL	1/7/2013
Cerilliant Corporation	Amobarbital-D5 (0.1 mg/mL)	Glass Ampule: 1 mL	1/7/2013
Cerilliant Corporation	2,5-Dimethoxy-4-(n)-propylthiophenethylamine HCl (2C–T–7 HCl) (1 mg/mL).	Glass Ampule: 1 mL	1/7/2013
Cerilliant Corporation	Class B Mix Urinalysis Internal Standard-8 (0.002–0.2 mg/mL), Pregabalin-D6 (1.0 mg/mL), Codeine-6-beta-glucoronide (1.0 mg/mL), Desomorphine (1.0 mg/mL), Levorphanol-D3 (0.1 mg/mL).	Glass ampule: 1 ml	1/22/2013
Cerilliant Corporation	Zaleplon-D4 (0.1 mg/mL)	Glass Ampule: 1 mL	1/31/2013
Cerilliant Corporation	Retigabine-D4 (0.1 mg/mL)	Glass ampule: 1ml	3/22/2013
Cerilliant Corporation	Tilidine-D6 HCl (0.1 mg/mL)	Glass ampule: 1ml	3/22/2013
Cliniqa Corporation	TDM Control, Levels 1–3	Dropper bottle: 5 mL	12/4/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Controls, THC (0.5–50 ng/mL)	Glass vial: 5 ml–4 L	10/22/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Controls, Methamphetamine (2–200 ng/mL)	Glass vial: 5 ml–4 L	10/22/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Controls, Amphetamine, Benzoylecgonine, Morphine, Phencyclidine (2–200 ng/mL).	Glass vial: 5 ml–4 L	10/22/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Controls, Multidrug (THC, Benzoylecgonine, Amphetamine, Methamphetamine, Morphine, Codeine, 6-Acetylmorphine, & PCP) at 0.5–50 ng/mL.	Glass vial: 5 ml–4 L	10/22/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Calibrators, THC (0.5–50 ng/mL)	Glass vial: 5 ml–4 L	10/22/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Calibrators, Methamphetamine (2–200 ng/mL)	Glass vial: 5 ml–4 L	10/22/2012

CHART I—Continued

Supplier	Product name	Form	Application date
ElSohly Laboratories, Inc	ELI Oral Fluid Calibrators, Amphetamine, Benzoylecgonine, Morphine, Phencyclidine (2–200 ng/mL).	Glass vial: 5 ml–4 L	10/22/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Calibrators, Multidrug (THC, Benzoylecgonine, Amphetamine, Methamphetamine, Morphine, Codeine, 6-Acetylmorphine, & PCP) at 0.5–200 ng/mL.	Glass vial: 5 ml–4 L	10/22/2012
Environmental Resource Associates (ERA).	USGS–BQS LS 4434 Mix 1	Glass Ampule: 2 mL	8/9/2012
Environmental Resource Associates (ERA).	USGS–BQS LS 4434 Mix 2	Glass Ampule: 2 mL	8/9/2012
Immunalysis Corporation	Tapentadol Calibrator Level 1 (100 ng/mL)	Glass vial: 10 mL	12/11/2012
Immunalysis Corporation	Tapentadol Calibrator Level 2 (200 ng/mL)	Glass vial: 10 mL	12/11/2012
Immunalysis Corporation	Tapentadol Calibrator Level 3 (500 ng/mL)	Glass vial: 10 mL	12/11/2012
Immunalysis Corporation	Tapentadol Calibrator Level 4 (1,000 ng/mL)	Glass vial: 10 mL	12/11/2012
Immunalysis Corporation	Tapentadol Low Control (150 ng/mL)	Glass vial: 10 mL	12/11/2012
Immunalysis Corporation	Tapentadol High Control (250 ng/mL)	Glass vial: 10 mL	12/11/2012
LGC Limited	(S)-(+)-Amphetamine (Dextroamphetamine) 1.0 mg/mL in Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Amphetamine 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Amphetamine 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Amphetamine-D11 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Amphetamine-D11 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Amphetamine-D5 (D-label on ring) 0.1 mg/mL in Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	Methamphetamine ((S)-(+)-Methamphetamine) 1.0 mg/mL in Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Methamphetamine 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Methamphetamine 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Methamphetamine-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Methamphetamine-D5 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Methcathinone Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	9/24/2012
LGC Limited	Methylphenidate Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	9/24/2012
LGC Limited	Amobarbital 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Pentobarbital 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Pentobarbital-D5 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Pentobarbital-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Secobarbital 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-MDA–D5 (rac-3,4-Methylenedioxyamphetamine-D5) 0.1 mg/mL Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-MDA–D5 (rac-3,4-Methylenedioxyamphetamine-D5) 1.0 mg/mL Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-MBDB HCl (rac-N-Methyl-1-(3,4-Methylenedioxyphenyl)-2-butanamine Hydrochloride) 1.0 mg/mL Methanol (as free base).	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-MBDB–D5 (rac-N-Methyl-1-(3,4-Methylenedioxyphenyl)-2-butanamine-D5) 0.1 mg/mL Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-MDEA–D5 (rac-3,4-Methylenedioxy-N-ethylamphetamine-D5) 0.1 mg/mL Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-MDEA–D5 (rac-3,4-Methylenedioxy-N-ethylamphetamine-D5) 1.0 mg/mL Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-MDEA–D6 (rac-3,4-Methylenedioxy-N-ethylamphetamine-D6) 0.1 mg/mL Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-MDMA–D5 (rac-3,4-Methylenedioxymethamphetamine-D5) 0.1 mg/mL Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-MDMA–D5 (rac-3,4-Methylenedioxymethamphetamine-D5) 1.0 mg/mL Methanol.	Glass vial: 1 mL	9/24/2012
LGC Limited	Phencyclidine (PCP) 1.0 mg/mL Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Phencyclidine-D5 (PCP-D5) 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Phencyclidine-D5 (PCP-D5) 1.0 mg/mL Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Codeine 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Codeine 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Codeine-D3 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Codeine-D3 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Codeine-D6 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Codeine-D6 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Isocodeine 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Norcodeine 0.01 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Norcodeine 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Codeine N-Oxide 0.01 mg/mL Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Oxycodone-D3 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012

CHART I—Continued

Supplier	Product name	Form	Application date
LGC Limited	Oxycodone-D6 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Oxycodone-D6 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Hydromorphone 1.0 mg/mL Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Norhydromorphone Hydrochloride 1.0 mg/mL Methanol (as free base).	Glass vial: 1 mL	9/24/2012
LGC Limited	Pethidine (Meperidine) 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Pethidine-D4 (Meperidine-D4) 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Pethidine-D4 (Meperidine-D4) 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Morphine 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Morphine 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Morphine 3-beta-D-Glucuronide 0.1 mg/mL in Methanol/Water (1/1).	Glass vial: 1 mL	9/24/2012
LGC Limited	Morphine 3-beta-D-Glucuronide 1.0 mg/mL in Methanol with 0.05 percent Sodium Hydroxide (w/v).	Glass vial: 1 mL	9/24/2012
LGC Limited	Morphine 6-beta-D-Glucuronide 0.1 mg/mL in Methanol/Water (1/1).	Glass vial: 1 mL	9/24/2012
LGC Limited	Morphine 6-beta-D-Glucuronide 1.0 mg/mL Water/Methanol (80/20).	Glass vial: 1 mL	9/24/2012
LGC Limited	Morphine-D3 0.1 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Morphine-D3 1.0 mg/mL in Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Morphine N-Oxide 0.01 mg/mL Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	Normorphine 1.0 mg/mL Methanol	Glass vial: 1 mL	9/24/2012
LGC Limited	rac-Amphetamine-D10 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Amphetamine-D5 (D-label on ring) 0.1 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Amphetamine-D5 (D-label on side chain) 0.1 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Amphetamine-D5 (D-label on side chain) 1.0 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Amphetamine-D6 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Amphetamine-D6 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Amphetamine-D8 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Amphetamine-D8 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methamphetamine-D11 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methamphetamine-D11 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methamphetamine-D14 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methamphetamine-D14 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methamphetamine-D8 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methamphetamine-D8 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methamphetamine-D9 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methamphetamine-D9 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	(+)-Norpseudoephedrine Hydrochloride (Cathine Hydrochloride) 0.1 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	(R)-(+)-Cathinone Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	(S)-(-)-Cathinone Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	(S)-(-)-Methcathinone Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Phentermine 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Fenfluramine Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Methylphenidate-D9 Hydrochloride 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Pipradrol Hydrochloride 0.1 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Pipradrol Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	GHB Sodium Salt (Sodium Gammahydroxybutyrate) 1.0 mg/mL in Methanol (as free acid).	Glass vial: 1 mL	11/30/2012
LGC Limited	GHB–D6 Sodium Salt (Sodium Gammahydroxybutyrate-D6) 0.1 mg/mL in Methanol (as free acid).	Glass vial: 1 mL	11/30/2012
LGC Limited	GHB–D6 Sodium Salt (Sodium Gammahydroxybutyrate-D6) 1.0 mg/mL in Methanol (as free acid).	Glass vial: 1 mL	11/30/2012
LGC Limited	Hexobarbital 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Methohexital 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Methohexital-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Phenobarbital 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Phenobarbital-D5 (D-label on ring) 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Phenobarbital-D5 (D-label on ring) 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Phenobarbital-D5 (D-label on side chain) 0.1 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012

CHART I—Continued

Supplier	Product name	Form	Application date
LGC Limited	Phenobarbital-D5 (D-label on side chain) 1.0 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	Methaqualone 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Methaqualone-D7 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Clonazepam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Clonazepam-D4 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Chlordiazepoxide 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Chlordiazepoxide-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Bromazepam 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Bromazepam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Clobazam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Delorazepam 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Estazolam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Estazolam-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Flunitrazepam 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Flunitrazepam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Flunitrazepam-D7 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Prazepam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Prazepam-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Diazepam 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Diazepam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Diazepam-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Diazepam-D5 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Flurazepam 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Flurazepam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Lormetazepam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Zaleplon 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Pregabalin 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Pregabalin-D6 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Zolpidem Tartrate 1.0 mg/mL in Methanol (as free base)	Glass vial: 1 mL	11/30/2012
LGC Limited	Zolpidem-D6 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Zopiclone 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Zopiclone Oxide 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Zopiclone-D4 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Meprobamate 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Nitrazepam 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Nitrazepam 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Nitrazepam-D5 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Oxazepam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Oxazepam-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Oxazepam-D5 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Nordazepam (Nordiazepam) 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Nordazepam (Nordiazepam) 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Nordazepam-D5 (Nordiazepam-D5) 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Nordazepam-D5 (Nordiazepam-D5) 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Alprazolam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Alprazolam-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Alprazolam-D5 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Methyltestosterone (17alpha-Methyltestosterone) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Midazolam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Midazolam-D4 Maleate 0.1 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Lorazepam 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Lorazepam 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Lorazepam Acetate ((3RS)-7-Chloro-5-(2-chlorophenyl)-2-oxo-2,3-dihydro-1H-1,4-benzodiazepin-3-yl Acetate) 0.1 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Lorazepam-D4 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Lorazepam-D4 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Tetrazepam 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Triazolam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Triazolam-D4 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Temazepam 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Temazepam 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Temazepam-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Temazepam-D5 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Androstanolone (5alpha-Dihydrotestosterone, Stanolone) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Androstenedione (Androst-4-ene-3, 17-dione) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012

CHART I—Continued

Supplier	Product name	Form	Application date
LGC Limited	Boldenone 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Fluoxymesterone 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Mesterolone 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Methandienone 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Nandrolone 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Norandrostenedione (19-Norandrost-4-ene-3,17-dione) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Norethandrolone 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Stanozolol 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Stanozolol-D3 0.1 mg/mL in 1,2-Dimethoxyethane	Glass vial: 1 mL	11/30/2012
LGC Limited	Testosterone 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Testosterone Benzoate 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Testosterone Isocaproate 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Testosterone Propionate 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Trenbolone 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	RCS–4 (DD–001, (4-Methoxyphenyl)(1-pentylindol-3-yl)methanone) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	JWH–081 ((4-Methoxynaphthalen-1-yl)(1-pentylindol-3-yl)methanone) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	JWH–122 ((4-Methylnaphthalen-1-yl)(1-pentylindol-3-yl)methanone) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	JWH–203 (2-(2-Chlorophenyl)-1-(1-pentylindol-3-yl)ethanone) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Ketamine Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Ketamine-D4 Hydrochloride 0.1 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Iso-LSD (Lysergic Acid Diethylamide) 0.1 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	LAMPA (Lysergic Acid N-Methyl-N-propylamide) 0.025 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	LSD (Lysergic Acid Diethylamide) 0.025 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	LSD–D3 (Lysergic Acid Diethylamide-D3) 0.025 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	PMA HCl (p-Methoxyamphetamine) 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-MDA (rac-3,4-Methylenedioxyamphetamine) 1.0 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-MDEA (rac-3,4-Methylenedioxy-N-ethylamphetamine) 1.0 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	1-Methylamino-1-(3,4-methylenedioxyphenyl)propane Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-BDB HCl (rac-3,4-Methylenedioxyphenyl-2-butanamine Hydrochloride) 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-MDMA (rac-3,4-Methylenedioxymethamphetamine) 1.0 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	Bufotenine 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Methylone Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Methylone-D3 Hydrochloride 0.1 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	AM–694 (1-(5-Fluoropentyl)-3-(2-iodobenzoyl)indole) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Cocaine 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Cocaine-D3 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Cocaine-D3 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Norcocaine Hydrochloride 1.0 mg/mL in Acetonitrile (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Norcocaine-D3 Hydrochloride 0.1 mg/mL in Acetonitrile (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	O-Methylcodeine 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Buprenorphine 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Buprenorphine 3-beta-D-Glucuronide 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Buprenorphine-D4 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Dihydrocodeine Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Dihydrocodeine-D6 Hydrochloride 0.1 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Noroxycodone Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012

CHART I—Continued

Supplier	Product name	Form	Application date
LGC Limited	Noroxycodone-D3 Hydrochloride 0.1 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Oxycodone 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Oxycodone-D3 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Dihydromorphine 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Hydromorphone-D3 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Anhydroecgonine Hydrochloride 1.0 mg/mL in Acetonitrile (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Anhydroecgonine Methyl Ester 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Benzoylecgonine 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Benzoylecgonine-D3 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Benzoylecgonine-D3 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Benzoylecgonine-D8 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Benzoylecgonine-D8 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Cocaethylene (Benzoylethylecgonine) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Cocaethylene-D3 (Benzoylethylecgonine-D3) 0.1 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Cocaethylene-D8 (Benzoylethylecgonine-D8) 0.1 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Ecgonine 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Ecgonine Ethyl Ester 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Ecgonine Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Ecgonine Methyl Ester 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Ecgonine Methyl Ester 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Ecgonine-D3 Hydrochloride 0.1 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Ecgonine-D3 Methyl Ester 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Ethylmorphine 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Hydrocodone 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Hydrocodone-D3 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Hydrocodone-D6 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Norhydrocodone Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Heroin (Diacetylmorphine) 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Heroin-D9 (Diacetylmorphine-D9) 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Heroin-D9 (Diacetylmorphine-D9) 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methadone 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methadone 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methadone-D3 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methadone-D3 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methadone-D9 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Methadone-D9 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Dextropropoxyphene 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Dextropropoxyphene 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Propoxyphene-D11 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Propoxyphene-D11 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Propoxyphene-D5 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	rac-Propoxyphene-D5 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	6-Acetylmorphine 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	6-Acetylmorphine 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	6-Acetylmorphine-D3 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	6-Acetylmorphine-D3 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	6-Acetylmorphine-D6 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	6-Acetylmorphine-D6 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Acetylcodeine 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Acetylcodeine 1.0 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
LGC Limited	Nordihydromorphine Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Oxymorphone 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Oxymorphone-D3 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Oxymorphone-D3 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Noroxymorphone Hydrochloride 0.1 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Noroxymorphone Hydrochloride 1.0 mg/mL in Methanol (as free base).	Glass vial: 1 mL	11/30/2012
LGC Limited	Butorphanol Tartrate 1.0 mg/mL in Methanol (as free base)	Glass vial: 1 mL	11/30/2012
LGC Limited	Sufentanil Citrate 0.1 mg/mL in Methanol (as free base)	Glass vial: 1 mL	11/30/2012

CHART I—Continued

Supplier	Product name	Form	Application date
LGC Limited	cis-Tilidine (Ethyl (1R,2RS)-2-(Dimethylamino)-1-phenylcyclohex-3-enecarboxylate) 0.1 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	Tilidine 0.1 mg/mL in Acetonitrile	Glass vial: 1 mL	11/30/2012
Microgenics Corporation	Abbott TDM Multiconstituent Calibrator (CAL 1–6) Catalog Number: 5P04–01.	Box: 7 vials; 5 mL each	7/2/2012
Microgenics Corporation	Thermo Scientific DRI Opiate Assay Catalog Number: 100014602.	Vial: 18 mL, Box: 3 vials	1/30/2013
Microgenics Corporation	Thermo Scientific DRI Cannabinoid Assay Catalog Number: 10014665.	Vial: 18 mL, Box: 3 vials	1/30/2013
Microgenics Corporation	Thermo Scientific Phencyclidine Assay Catalog Number: 10014673.	Vial: 18 mL, Box: 3 vials	1/30/2013
Microgenics Corporation	Thermo Scientific Ecstacy Assay Catalog Number: 10014681	Vial: 18 mL, Box: 3 vials	1/30/2013
Microgenics Corporation	Thermo Scientific CEDIA Methamphetamine OFT Calibrator 1: Catalog Number 10016362.	Vial: 5 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Methamphetamine OFT Calibrator 2: Catalog Number 10063.	Vial: 5 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Methamphetamine OFT Calibrator 3: Catalog Number 10016364.	Vial: 5 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Methamphetamine OFT Control 1: Catalog Number 10017686.	Vial: 10 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Methamphetamine OFT Control 2: Catalog Number 10017687.	Vial: 10 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Methamphetamine OFT Control 3: Catalog Number 1007688.	Vial: 10 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Multi-Drug OFT Calibrator 1: Catalog Number 10016882.	Vial: 10 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Multi-Drug OFT Calibrator 2: Catalog Number 10016883.	Vial: 10 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Multi-Drug OFT Calibrator 3: Catalog Number 10016884.	Vial: 10 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Multi-Drug OFT Control 1: Catalog Number 10017711.	Vial: 15 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Multi-Drug OFT Control 2: Catalog Number 10017712.	Vial: 15 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA Multi-Drug OFT Control 3: Catalog Number 10017713.	Vial: 15 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA THC OFT Calibrator 1: Catalog Number 10016700.	Vial: 5 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA THC OFT Calibrator 2: Catalog Number 10016701.	Vial: 5 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA THC OFT Calibrator 3: Catalog Number 10016702.	Vial: 5 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA THC OFT Control 1: Catalog Number 10017702.	Vial: 10 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA THC OFT Control 2: Catalog Number 10017703.	Vial: 10 mL; Box: 1 vial	2/11/2013
Microgenics Corporation	Thermo Scientific CEDIA THC OFT Control 3: Catalog Number 10017704.	Vial: 10 mL; Box: 1 vial	2/11/2013
Ortho Clinical Diagnostics, Inc	VITROS Immunodiagnostics Products Testosterone Reagent Pack.	Plastic chamber: 8.4 mL	9/28/2012
Ortho Clinical Diagnostics, Inc	VITROS Immunodiagnostics Products Estradiol Reagent Pack.	Plastic chamber: 13.3 mL	9/28/2012
Ortho Clinical Diagnostics, Inc	VITROS Immunodiagnostics Products Testosterone Calibrators (1–3).	Box: 3 vials; 2 mL each	10/19/2012
Ortho Clinical Diagnostics, Inc	VITROS Immunodiagnostics Products Testosterone Controls (1–3).	Box: 9 vials; 1 mL each	10/19/2012
Ortho Clinical Diagnostics, Inc	VITROS Immunodiagnostics Products Testosterone Range Verifiers (Low and High).	Box: 4 vials; 1–1.5 mL each	10/19/2012
Restek	Custom Revised Appendix IX Mix #1	Glass vial: 1.3 mL	2/14/2013
Restek Corporation	Product and Reactor Samples Standard	Ampule: 2 mL	10/12/2012
Restek Corporation	8270 Supplemental Standard #1	Ampule: 2 mL	12/13/2012
Siemens Healthcare Diagnostics Inc.	TTST Stock X (Bulk)	Stock container: 50 mL–1 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	VS TTST Cal Bulk Soln Level 1	Bulk container: 4 mL–100 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	VS TTST Cal Bulk Soln Level 2	Bulk container: 4 mL–100 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	VS TTST Cal Bulk Soln Level 3	Bulk container: 4 mL–100 L	7/26/2012

CHART I—Continued

Supplier	Product name	Form	Application date
Siemens Healthcare Diagnostics Inc.	VS TTST Cal Bulk Soln Level 4	Bulk container: 4 mL–100 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	VS TTST Cal Bulk Soln Level 5	Bulk container: 4 mL–100 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	VS TTST Cal Bulk Soln Level 6	Bulk container: 4 mL–100 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	Dimension Vista TTST CAL A	Vial: 1 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	Dimension Vista TTST CAL B	Vial: 1 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	Dimension Vista TTST CAL C	Vial: 1 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	Dimension Vista TTST CAL D	Vial: 1 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	Dimension Vista TTST CAL E	Vial: 1 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	Dimension Vista TTST CAL F	Vial: 1 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	Dimension Vista System TTST CAL	Box: 12 vials; 1 ml each	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST Lvl 1 Bulk	Bulk container: 4 mL–20 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST Lvl 2 Bulk	Bulk container: 4 mL–20 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST Lvl 3 Bulk	Bulk container: 4 mL–20 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST Lvl 4 Bulk	Bulk container: 4 mL–20 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST Lvl 5 Bulk	Bulk container: 4 mL–20 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST Lvl 6 Bulk	Bulk container: 4 mL–20 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST CAL Lvl 1 FC	Vial: 1–5 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST CAL Lvl 2 FC	Vial: 1–5 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST CAL Lvl 3 FC	Vial: 1–5 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST CAL Lvl 4 FC	Vial: 1–5 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST CAL Lvl 5 FC	Vial: 1–5 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	MP TTST CAL Lvl 6 FC	Vial: 1–5 mL	7/26/2012
Siemens Healthcare Diagnostics Inc.	TTST Stock S (Bulk)	Stock container: 50 mL–1 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	TTST Stock T (Bulk)	Stock container: 50 mL–1 L	7/26/2012
Siemens Healthcare Diagnostics Inc.	Estradiol Diluent	Plastic container: 0.5L–1.0L	11/14/2012
Siemens Healthcare Diagnostics Inc.	Dimension Vista Drug 1 CAL Pilot, Level B	Plastic test tube: 15 mL	12/3/2012
Supelco, Inc	Titan C18 Batch Test 1 Mix	Glass ampule: 1 mL	2/28/2013

The Deputy Assistant Administrator has found that each of the compounds, mixtures, and preparations described in Chart II below is not consistent with the criteria stated in 21 U.S.C. 811(g)(3)(B) and in 21 CFR 1308.23. Accordingly, the Deputy Assistant Administrator has determined that the chemical preparations or mixtures generally described in Chart II below and specifically described in the application materials received by DEA, are not exempt from application of any part of the CSA or from application of any part of the CFR, with regard to the requested exemption pursuant to 21 CFR 1308.23, as of the date listed below that was provided in the determination letters to the individual requesters.

CHART II

Supplier	Product name	Form	Application date
Agilent Technologies	Forensic Toxicology Comprehensive Mix–Submix 9A	Ampule: 1 mL	12/6/2012

CHART II—Continued

Supplier	Product name	Form	Application date
Biochemical Diagnostics, Inc ..	Detectabuse Custom Liquid Control Urine, MC177	Glass vials: 1 ml–200 mL	10/11/2012
Biochemical Diagnostics, Inc ..	Detectabuse Custom Liquid Control Urine, MC178	Glass vials: 1 ml–200 mL	10/23/2012
Biochemical Diagnostics, Inc ..	Salivabuse Custom Liquid Control Oral Fluid, OF24	Glass vials: 1 ml–200 mL	10/23/2012
Biochemical Diagnostics, Inc ..	Detectabuse Custom Liquid Control Urine, MC180	Glass vials: 1 ml–200 mL	12/14/2012
Biochemical Diagnostics, Inc ..	Detectabuse Custom Liquid Control Urine, MC181	Glass vials: 1 ml–200 mL	12/14/2012
Biochemical Diagnostics, Inc ..	Detectabuse Custom Liquid Control Urine, MC182	Glass vials: 1 ml–2 L	1/22/2013
Biochemical Diagnostics, Inc ..	Detectabuse Custom Liquid Control Urine, MC180	Glass vials: 1 ml–200 mL	1/29/2013
Biochemical Diagnostics, Inc ..	Detectabuse Custom Liquid Control Urine, MC182	Glass vials: 1 ml–2 L	2/14/2013
Cerilliant Corporation	(-)-11-nor-9-Carboxy-delta9-THC (1 mg/mL)	Glass ampule: 5 mL	7/6/2012
Cerilliant Corporation	Cocaine HCl (10 mg/mL)	Glass Ampule: 1 mL	7/6/2012
Cerilliant Corporation	Lacosamide (10 mg/mL)	Glass Ampule: 1 mL	9/12/2012
Cliniqa Corporation	TDM Control, Levels 1–3 Bulk	Plastic Container: 500 L	12/4/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Controls, THC (0.5–50 ng/mL)	Glass vial: 5 ml–4 L	8/9/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Controls, Methamphetamine (2–200 ng/mL)	Glass vial: 5 ml–4 L	8/9/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Controls, Amphetamine, Benzoylecgonine, Morphine, Phencyclidine (2–200 ng/mL).	Glass vial: 5 ml–4 L	8/9/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Controls, Multidrug (selected from THC, Cocaine/Benzoylecgonine, Amphetamines, Opiates, & PCP at 0.5–50 ng/mL).	Glass vial: 5 ml–4 L	8/9/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Calibrators, THC (0.5–50 ng/mL)	Glass vial: 5 ml–4 L	8/9/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Calibrators, Methamphetamine (2–200 ng/mL)	Glass vial: 5 ml–4 L	8/9/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Calibrators, Amphetamine, Benzoylecgonine, Morphine, Phencyclidine (2–200 ng/mL).	Glass vial: 5 ml–4 L	8/9/2012
ElSohly Laboratories, Inc	ELI Oral Fluid Calibrators, Multidrug (selected from THC, Cocaine/Benzoylecgonine, Amphetamines, Opiates, & PCP at 0.5–200 ng/mL).	Glass vial: 5 ml–4 L	8/9/2012
LGC Limited	LAMPA (Lysergic Acid N-Methyl-N-propylamide) 1.0 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	LSD (Lysergic Acid Diethylamide) 1.0 mg/mL in Acetonitrile ..	Glass vial: 1 mL	11/30/2012
LGC Limited	LSD–D3 (Lysergic Acid Diethylamide-D3) 0.1 mg/mL in Acetonitrile.	Glass vial: 1 mL	11/30/2012
LGC Limited	(-)-Cannabidiol 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Cannabinol 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	(-)-delta9-THC (Dronabinol) 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	(-)-delta9-THC-D3 (Dronabinol-D3) 0.1 mg/mL in Methanol ...	Glass vial: 1 mL	11/30/2012
LGC Limited	trans-11-Hydroxy-delta9-THC ((6aRS,10aRS)-11-Hydroxy-delta9-tetrahydrocannabinol) 0.1 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	trans-11-Hydroxy-delta9-THC ((6aRS,10aRS)-11-Hydroxy-delta9-tetrahydrocannabinol) 1.0 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	trans-11-Hydroxy-delta9-THC-D3 (trans-11-Hydroxy-delta9-tetrahydrocannabinol-D3) 0.1 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	trans-11-Nor-9-carboxy-delta9-THC 0.1 mg/mL in Methanol ..	Glass vial: 1 mL	11/30/2012
LGC Limited	trans-11-Nor-9-carboxy-delta9-THC-D3 0.1 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	trans-11-Nor-9-carboxy-delta9-THC-D3 1.0 mg/mL in Methanol.	Glass vial: 1 mL	11/30/2012
LGC Limited	Fentanyl 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Fentanyl 1.0 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
LGC Limited	Fentanyl-D5 0.1 mg/mL in Methanol	Glass vial: 1 mL	11/30/2012
Siemens Healthcare Diagnostics Inc.	Dimension Vista Drug 1 CAL Bulk, Level B	Plastic container: 4–100 L	12/3/2012

Scope of Approval

The exemptions are applicable only to the precise preparation or mixture described in the application submitted to DEA in the form(s) listed in this order and only for those sections of the CSA and the CFR that are specifically identified. Pursuant to 21 CFR 1308.24(h), any change in the quantitative or qualitative composition of the preparation or mixture or change in the trade name or other designation of the preparation or mixture after the date of application requires a new application. Pursuant to 21 CFR 1308.24(g), DEA may prescribe requirements other than those set forth in 1308.24(b)–(e) on a case-by-case basis for materials exempted in bulk quantities. Accordingly, in order to limit opportunity for diversion from the larger bulk quantities, DEA has determined that each of the exempted bulk products listed in this order may only be used by the manufacturer and may not be distributed by the manufacturer for any purpose or transported to other facilities.

Additional exempt chemical preparation requests received between July 1, 2012, and March 31, 2013, and not otherwise referenced in this order may remain under consideration until DEA receives additional information required, pursuant to 21 CFR 1308.23(d), as detailed in separate correspondence to individual requesters. DEA's order on such requests will be communicated to the public in a future **Federal Register** publication.

DEA also notes that these exemptions are limited to exemption from only those sections of the CSA and the CFR that are specifically identified in 21 CFR

1308.24(a). All other requirements of the CSA and the CFR apply, including, but not limited to, registration as an importer as required by 21 U.S.C. 957.

Chemical Preparations Containing Newly Controlled Substances

The statutory authority for exempt chemical preparations is based on the control status of substances contained within a preparation, the intended administration of a preparation, and the packaged form of a preparation. DEA conducts a case-by-case analysis of each application for exemption to determine whether exemption of a preparation from certain provisions of the CSA is appropriate pursuant to the specified statutory and regulatory requirements.

Most exempt chemical preparations have remained effective until the holder of a specific exempt chemical preparation specifically requested that the exemption be terminated. The CSA allows for modifications to the controlled substances schedules to add, remove, or change the schedule of substances thus resulting in periodic modifications to the control status of various substances. 21 U.S.C. 811(a). Since the CSA was enacted in 1970, DEA has on several occasions added to, removed from, or modified the schedules of controlled substances in accordance with the CSA. Such changes may result in the non-compliance of exempt chemical preparations with current statutes or regulations if chemical preparations that have already obtained exempt status contain newly controlled substances. For example, although an exempt chemical preparation may continue to be packaged in the same manner as when it was approved, non-controlled substances in the preparation may become controlled, thus prompting the need for a new application for exemption of the chemical preparation to ensure continued compliance. Other preparations that previously contained no controlled substances may contain newly controlled substances and thus would require an application for exemption.

DEA reviews applications for chemical preparation exemptions based on the statutes and regulations that are in place at the time of the application, including the control status of substances included in the preparation. DEA must remain vigilant to ensure that exempt chemical preparations remain consistent with the standards set forth in the CSA and its implementing regulations. As such, DEA reminds the public that any chemical preparation, regardless of whether it was previously exempt, that contains a newly controlled substance will require a new application for exemption pursuant to 21 U.S.C. 811(g)(3)(B) and 21 CFR 1308.23–1308.24.

Review of Exemptions Pursuant to 21 U.S.C. 811(g)(3)

Based on inquiries received from industry, DEA is conducting a comprehensive review of the exempt chemical preparation regulations. DEA's regulations at 21 CFR 1308.24(a) state that approved chemical preparations are exempt from certain provisions of both Subchapter I and Subchapter II of the CSA: ''The chemical preparations and mixtures approved pursuant to 1308.23 are exempt from application of sections 302, 303, 305, 306, 307, 308, 309, 1002, 1003 and 1004 of the Act (21 U.S.C. 822–823, 825–829, 952–954) and 1301.74 of this chapter, to the extent described in paragraphs (b) to (h) of this section.'' Pursuant to its regulations, DEA has provided exemptions from the application of section 302, 303, 305, 306, 307, 308, 309, 1002, 1003, and 1004 of the Act (21 U.S.C. 822–823, 825–829, 952–954) and 21 CFR 1301.74 since the implementation of the regulations in the early 1970s. Until DEA's analysis of the exemption regulations is complete, DEA will continue to review and provide exemptions to chemical preparations consistent with the implementing regulations, when warranted. DEA will publish a future notice regarding the outcome of DEA's review of its regulations with respect to the exemption of chemical preparations.

Opportunity for Comment

Pursuant to 21 CFR 1308.23, any interested person may submit written comments on or objections to any chemical preparation in this order that has been approved or denied as exempt. If any comments or objections raise significant issues regarding any finding of fact or conclusion of law upon which this order is based, the Deputy Assistant Administrator will immediately suspend the effectiveness of any applicable part of this order until he may reconsider the application in light of the comments and objections filed.

Approved Exempt Chemical Preparations Are Posted on DEA's Web Site

A list of all current exemptions, including those listed in this order, is available on DEA's Web site at *http://www.DEAdiversion.usdoj.gov/schedules/exempt/exempt_chemlist.pdf.* The dates of applications of all current exemptions are posted for easy reference.

Dated: June 17, 2013.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

[FR Doc. 2013–16010 Filed 7–2–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–378]

Proposed Aggregate Production Quotas for Schedule I and II Controlled Substances and Proposed Assessment of Annual Needs for the List I Chemicals Ephedrine, Pseudoephedrine, and Phenylpropanolamine for 2014

AGENCY: Drug Enforcement Administration (DEA), Department of Justice.

ACTION: Notice with request for comments.

SUMMARY: This notice proposes initial year 2014 aggregate production quotas for controlled substances in Schedules I and II of the Controlled Substances Act (CSA) and assessment of annual needs for the List I chemicals ephedrine, pseudoephedrine, and phenylpropanolamine.

DATES: Electronic comments must be submitted and written comments must be postmarked on or before August 2, 2013. The electronic Federal Docket Management System will not accept comments after midnight Eastern Time on the last day of the comment period.

ADDRESSES: To ensure proper handling of comments, please reference ''Docket No. DEA–378'' on all electronic and written correspondence. DEA encourages that all comments be submitted electronically through *http://www.regulations.gov* using the electronic comment form provided on that site. An electronic copy of this document is also available at *http://www.regulations.gov* for easy reference. Paper comments that duplicate the electronic submission are not necessary as all comments submitted to *www.regulations.gov* will be posted for public review and are part of the official docket record. Written comments submitted via regular or express mail should be sent to the Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODL, 8701 Morrissette Drive, Springfield, VA 22152.

FOR FURTHER INFORMATION CONTACT: John W. Partridge, Executive Assistant, Office of Diversion Control, Drug Enforcement Administration, 8701 Morrissette Drive,