

Administration (DEA) as importer of schedule I controlled substances.

**SUPPLEMENTARY INFORMATION:** The company listed below applied to be registered as an importer of basic class of controlled substances. Information on previously published notice is listed in the table below. No comments or objections were submitted and no requests for a hearing were submitted for this notice.

Company	FR Docket	Published
Sharp (Bethlehem), LLC.	84 FR 9837.	March 18, 2019

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrant to import the applicable basic class of schedule I 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. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing 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 DEA has granted a registration as an importer for schedule I controlled substances to the above listed company.

Dated: June 3, 2019.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2019-14023 Filed 7-1-19; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Importer of Controlled Substances Application: Bellwyck Clinical Services**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before August 1, 2019. Such persons may also file a written request for a hearing on the application on or before August 1, 2019.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement

Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on April 17, 2019, Bellwyck Clinical Services, 8946 Global Way, West Chester, Ohio 45069 applied to be registered as an importer of the following basic class of controlled substances:

Controlled substance	Drug code	Schedule
Amphetamine ...	1100	II
Methylphenidate ...	1724	II
Oxycodone .....	9143	II

The company plans to import the listed controlled substances in dosage form to conduct clinical trials. Approval of permit applications will occur only when the registrant's activity is consistent with what is authorized under 21 U.S.C. 952(a) (2). Authorization will not extend to the import of FDA approved or non-approved finished dosage forms for commercial sale.

Dated: June 18, 2019.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2019-14027 Filed 7-1-19; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Bulk Manufacturer of Controlled Substances Application: Pisgah Laboratories, Inc.**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before September 3, 2019.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement

Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on March 5, 2019, Pisgah Laboratories, Inc., 3222 Old Hendersonville Highway, Pisgah Forest, North Carolina 28768 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Difenoxin .....	9168	I
Diphenoxylate ...	9170	II
Levorphanol .....	9220	II
Meperidine intermediate-B.	9233	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

Dated: June 19, 2019.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2019-14028 Filed 7-1-19; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Importer of Controlled Substances Application: Lipomed**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before August 1, 2019. Such persons may also file a written request for a hearing on the application on or before August 1, 2019.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register

Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on March 28, 2019, Lipomed, 150 Cambridge Park Drive,

Suite 705, Cambridge, Massachusetts 02140 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
3-Fluoro-N-methylcathinone (3-FMC)	1233	
Cathinone	1235	
Methcathinone	1237	
4-Fluoro-N-methylcathinone (4-FMC)	1238	
Pentedrone ( $\alpha$ -methylaminovalerophenone)	1246	
Mephedrone (4-Methyl-N-methylcathinone)	1248	
4-Methyl-N-ethylcathinone (4-MEC)	1249	
Naphyrone	1258	
N-Ethylamphetamine	1475	
N,N-Dimethylamphetamine	1480	
Fenethylamine	1503	
Aminorex	1585	
4-Methylaminorex (cis isomer)	1590	
Gamma Hydroxybutyric Acid	2010	
Methaqualone	2565	
Mecloqualone	2572	
JWH-250 (1-Pentyl-3-(2-methoxyphenylacetyl) indole)	6250	
SR-18 (Also known as RCS-8) (1-Cyclohexylethyl-3-(2-methoxyphenylacetyl) indole)	7008	
ADB-FUBINACA (N-(1-amino-3,3-dimethyl-1-oxobutan-2-yl)-1-(4-fluorobenzyl)-1H-indazole-3-carboxamide)	7010	
5-Fluoro-UR-144 and XLR11 ([1-(5-Fluoro-pentyl)1H-indol-3-yl](2,2,3,3-tetramethylcyclopropyl)methanone)	7011	
AB-FUBINACA (N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-(4-fluorobenzyl)-1H-indazole-3-carboxamide)	7012	
JWH-019 (1-Hexyl-3-(1-naphthoyl)indole)	7019	
MDMB-FUBINACA (Methyl 2-(1-(4-fluorobenzyl)-1H-indazole-3-carboxamido)-3,3-dimethylbutanoate)	7020	
FUB-AMB, MMB-FUBINACA, AMB-FUBINACA (2-(1-(4-fluorobenzyl)-1H-indazole-3-carboxamido)-3-methylbutanoate)	7021	
AB-PINACA (N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-pentyl-1H-indazole-3-carboxamide)	7023	
THJ-2201 ([1-(5-fluoropentyl)-1H-indazol-3-yl](naphthalen-1-yl)methanone)	7024	
5F-AB-PINACA (N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-(5-fluoropentyl)-1H-indazole-3-carboxamide)	7025	
AB-CHMINACA (N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-(cyclohexylmethyl)-1H-indazole-3-carboxamide)	7031	
MAB-CHMINACA (N-(1-amino-3,3-dimethyl-1-oxobutan-2-yl)-1-(cyclohexylmethyl)-1H-indazole-3-carboxamide)	7032	
5F-AMB (Methyl 2-(1-(5-fluoropentyl)-1H-indazole-3-carboxamido)-3-methylbutanoate)	7033	
5F-ADB; 5F-MDMB-PINACA (Methyl 2-(1-(5-fluoropentyl)-1H-indazole-3-carboxamido)-3,3-dimethylbutanoate)	7034	
ADB-PINACA (N-(1-amino-3,3-dimethyl-1-oxobutan-2-yl)-1-pentyl-1H-indazole-3-carboxamide)	7035	
MDMB-CHMICA, MMB-CHMINACA (Methyl 2-(1-(cyclohexylmethyl)-1H-indole-3-carboxamido)-3,3-dimethylbutanoate)	7042	
MMB-CHMICA, AMB-CHMICA (methyl 2-(1-(cyclohexylmethyl)-1H-indole-3-carboxamido)-3-methylbutanoate)	7044	
APINACA and AKB48 (N-(1-Adamantyl)-1-pentyl-1H-indazole-3-carboxamide)	7048	
5F-APINACA, 5F-AKB48 (N-(adamantan-1-yl)-1-(5-fluoropentyl)-1H-indazole-3-carboxamide)	7049	
JWH-081 (1-Pentyl-3-(1-(4-methoxynaphthoyl) indole)	7081	
5F-CUMYL-P7AICA (1-(5-fluoropentyl)-N-(2-phenylpropan-2-yl)-1H-pyrrolo[2,3-b]pyridine-3-carboxamide)	7085	
4-CN-CUMYL-BUTINACA, 4-cyano-CUMYL-BUTINACA, 4-CN-CUMYL BINACA, CUMYL-4CN-BINACA, SGT-78 (1-(4-cyanobutyl)-N-(2-phenylpropan-2-yl)-1H-indazole-3-carboximide)	7089	
SR-19 (Also known as RCS-4) (1-Pentyl-3-[(4-methoxy)-benzoyl] indole)	7104	
JWH-018 (also known as AM678) (1-Pentyl-3-(1-naphthoyl)indole)	7118	
JWH-122 (1-Pentyl-3-(4-methyl-1-naphthoyl) indole)	7122	
UR-144 (1-Pentyl-1H-indol-3-yl)(2,2,3,3-tetramethylcyclopropyl)methanone)	7144	
JWH-073 (1-Butyl-3-(1-naphthoyl)indole)	7173	
JWH-200 (1-[2-(4-Morpholinyl)ethyl]-3-(1-naphthoyl)indole)	7200	
AM2201 (1-(5-Fluoropentyl)-3-(1-naphthoyl) indole)	7201	
JWH-203 (1-Pentyl-3-(2-chlorophenylacetyl) indole)	7203	
NM2201, CBL2201 (Naphthalen-1-yl 1-(5-fluoropentyl)-1H-indole-3-carboxylate)	7221	
PB-22 (Quinolin-8-yl 1-pentyl-1H-indole-3-carboxylate)	7222	
5F-PB-22 (Quinolin-8-yl 1-(5-fluoropentyl)-1H-indole-3-carboxylate)	7225	
Alpha-ethyltryptamine	7249	
Ibogaine	7260	
CP-47,497 (5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl-phenol)	7297	
CP-47,497 C8 Homologue (5-(1,1-Dimethyloctyl)-2-[(1R,3S)-3-hydroxycyclohexyl-phenol)	7298	
Lysergic acid diethylamide	7315	
2,5-Dimethoxy-4-(n)-propylthiophenethylamine (2C-T-7)	7348	
Marihuana extract	7350	
Marihuana	7360	
Tetrahydrocannabinols	7370	
Parahexyl	7374	
Mescaline	7381	
2C-T-2, (2-(4-Ethylthio-2,5-dimethoxyphenyl) ethanamine)	7385	
3,4,5-Trimethoxyamphetamine	7390	
4-Bromo-2,5-dimethoxyamphetamine	7391	
4-Bromo-2,5-dimethoxyphenethylamine	7392	
4-Methyl-2,5-dimethoxyamphetamine	7395	

Controlled substance	Drug code	Schedule
2,5-Dimethoxyamphetamine	7396	I
JWH-398 (1-Pentyl-3-(4-chloro-1-naphthoyl) indole)	7398	I
2,5-Dimethoxy-4-ethylamphetamine	7399	I
3,4-Methylenedioxyamphetamine	7400	I
5-Methoxy-3,4-methylenedioxyamphetamine	7401	I
N-Hydroxy-3,4-methylenedioxyamphetamine	7402	I
3,4-Methylenedioxy-N-ethylamphetamine	7404	I
3,4-Methylenedioxy-methamphetamine	7405	I
4-Methoxyamphetamine	7411	I
5-Methoxy-N-N-dimethyltryptamine	7431	I
Alpha-methyltryptamine	7432	I
Bufotenine	7433	I
Diethyltryptamine	7434	I
Dimethyltryptamine	7435	I
Psilocybin	7437	I
Psilocyn	7438	I
5-Methoxy-N,N-diisopropyltryptamine	7439	I
N-Ethyl-1-phenylcyclohexylamine	7455	I
1-(1-Phenylcyclohexyl)pyrrolidine	7458	I
1-[1-(2-Thienyl)cyclohexyl]piperidine	7470	I
1-[1-(2-Thienyl)cyclohexyl]pyrrolidine	7473	I
N-Ethyl-3-piperidyl benzilate	7482	I
N-Methyl-3-piperidyl benzilate	7484	I
N-Benzylpiperazine	7493	I
4-MePPP (4-Methyl-alpha-pyrrolidinopropiophenone)	7498	I
2C-D (2-(2,5-Dimethoxy-4-methylphenyl) ethanamine)	7508	I
2C-E (2-(2,5-Dimethoxy-4-ethylphenyl) ethanamine)	7509	I
2C-H (2-(2,5-Dimethoxyphenyl) ethanamine)	7517	I
2C-I (2-(4-iodo-2,5-dimethoxyphenyl) ethanamine)	7518	I
2C-C (2-(4-Chloro-2,5-dimethoxyphenyl) ethanamine)	7519	I
2C-N (2-(2,5-Dimethoxy-4-nitro-phenyl) ethanamine)	7521	I
2C-P (2-(2,5-Dimethoxy-4-(n)-propylphenyl) ethanamine)	7524	I
2C-T-4 (2-(4-Isopropylthio)-2,5-dimethoxyphenyl) ethanamine)	7532	I
MDPV (3,4-Methylenedioxypropylvalerone)	7535	I
25B-NBOMe (2-(4-bromo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) ethanamine)	7536	I
25C-NBOMe (2-(4-chloro-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) ethanamine)	7537	I
25I-NBOMe (2-(4-iodo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) ethanamine)	7538	I
Methylone (3,4-Methylenedioxy-N-methylcathinone)	7540	I
Butylone	7541	I
Pentylone	7542	I
N-Ethylpentylone, ephylone (1-(1,3-benzodioxol-5-yl)-2-(ethylamino)-pentan-1-one)	7543	I
alpha-PVP (alpha-pyrrolidinopentiophenone)	7545	I
alpha-PBP (alpha-pyrrolidinobutiophenone)	7546	I
AM-694 (1-(5-Fluoropentyl)-3-(2-iodobenzoyl) indole)	7694	I
Acetyldihydrocodeine	9051	I
Benzylmorphine	9052	I
Codeine-N-oxide	9053	I
Cyprenorphine	9054	I
Desomorphine	9055	I
Etorphine (except HCl)	9056	I
Codeine methylbromide	9070	I
Dihydromorphine	9145	I
Difenoxin	9168	I
Heroin	9200	I
Hydromorphinol	9301	I
Methyldesorphine	9302	I
Methyldihydromorphine	9304	I
Morphine methylbromide	9305	I
Morphine methylsulfonate	9306	I
Morphine-N-oxide	9307	I
Myrophine	9308	I
Nicocodeine	9309	I
Nicomorphine	9312	I
Normorphine	9313	I
Pholcodine	9314	I
Thebacon	9315	I
Acetorphine	9319	I
Drotebanol	9335	I
U-47700 (3,4-dichloro-N-[2-(dimethylamino)cyclohexyl]-N-methylbenzamide)	9547	I
AH-7921 (3,4-dichloro-N-[(1-dimethylamino)cyclohexylmethyl]benzamide))	9551	I
MT-45 (1-cyclohexyl-4-(1,2-diphenylethyl)piperazine))	9560	I
Acetylmethadol	9601	I
Allylprodine	9602	I
Alphacetylmethadol except levo-alpha-cetylmethadol	9603	I

Controlled substance	Drug code	Schedule
Alphameprodine .....	9604	I
Alphamethadol .....	9605	I
Benzethidine .....	9606	I
Betacetylmethadol .....	9607	I
Betameprodine .....	9608	I
Betamethadol .....	9609	I
Betaprodine .....	9611	I
Clonitazene .....	9612	I
Dextromoramide .....	9613	I
Diampromide .....	9615	I
Diethylthiambutene .....	9616	I
Dimenoxadol .....	9617	I
Dimepheptanol .....	9618	I
Dimethylthiambutene .....	9619	I
Dioxaphetyl butyrate .....	9621	I
Dipipanone .....	9622	I
Ethylmethylthiambutene .....	9623	I
Etonitazene .....	9624	I
Etoxidine .....	9625	I
Furethidine .....	9626	I
Hydroxypethidine .....	9627	I
Ketobemidone .....	9628	I
Levomoramide .....	9629	I
Levophenacetylmorphan .....	9631	I
Morpheridine .....	9632	I
Noracetylmethadol .....	9633	I
Norlevorphanol .....	9634	I
Normethadone .....	9635	I
Norpipanone .....	9636	I
Phenadoxone .....	9637	I
Phenampramide .....	9638	I
Phenoperidine .....	9641	I
Piritramide .....	9642	I
Proheptazine .....	9643	I
Properidine .....	9644	I
Racemoramide .....	9645	I
Trimeperidine .....	9646	I
Phenomorphane .....	9647	I
Propiram .....	9649	I
1-Methyl-4-phenyl-4-propionoxypiperidine .....	9661	I
1-(2-Phenylethyl)-4-phenyl-4-acetoxypiperidine .....	9663	I
Tilidine .....	9750	I
Acryl fentanyl (N-(1-phenethylpiperidin-4-yl)-N-phenylacrylamide) .....	9811	I
Para-Fluorofentanyl .....	9812	I
3-Methylfentanyl .....	9813	I
Alpha-Methylfentanyl .....	9814	I
Acetyl-alpha-methylfentanyl .....	9815	I
N-(2-fluorophenyl)-N-(1-phenethylpiperidin-4-yl)propionamide .....	9816	I
Acetyl Fentanyl (N-(1-phenethylpiperidin-4-yl)-N-phenylacetamide) .....	9821	I
Butyryl Fentanyl .....	9822	I
Para-fluorobutyryl fentanyl .....	9823	I
4-Fluoroisobutyryl fentanyl (N-(4-fluorophenyl)-N-(1-phenethylpiperidin-4-yl)isobutyramide) .....	9824	I
2-methoxy-N-(1-phenethylpiperidin-4-yl)-N-phenylacetamide .....	9825	I
Para-chloroisobutyryl fentanyl .....	9826	I
Isobutyryl fentanyl .....	9827	I
Beta-hydroxyfentanyl .....	9830	I
Beta-hydroxy-3-methylfentanyl .....	9831	I
Alpha-methylthiofentanyl .....	9832	I
3-Methylthiofentanyl .....	9833	I
Furanyl fentanyl (N-(1-phenethylpiperidin-4-yl)-N-phenylfuran-2-carboxamide) .....	9834	I
Thiofentanyl .....	9835	I
Beta-hydroxythiofentanyl .....	9836	I
Para-methoxybutyryl fentanyl .....	9837	I
Ocfentanil .....	9838	I
Valeryl fentanyl .....	9840	I
N-(1-phenethylpiperidin-4-yl)-N-phenyltetrahydrofuran-2-carboxamide) .....	9843	I
Cyclopropyl Fentanyl .....	9845	I
Cyclopentyl fentanyl .....	9847	I
Fentanyl related-compounds as defined in 21 CFR 1308.11(h) .....	9850	I
Amphetamine .....	1100	II
Methamphetamine .....	1105	II
Lisdexamfetamine .....	1205	II
Phenmetrazine .....	1631	II
Methylphenidate .....	1724	II

Controlled substance	Drug code	Schedule
Amobarbital .....	2125	II
Pentobarbital .....	2270	II
Secobarbital .....	2315	II
Glutethimide .....	2550	II
Dronabinol in an oral solution in a drug product approved for marketing by the U.S. Food and Drug Administration	7365	II
Nabilone .....	7379	II
1-Phenylcyclohexylamine .....	7460	II
Phencyclidine .....	7471	II
ANPP (4-Anilino-N-phenethyl-4-piperidine) .....	8333	II
Phenylacetone .....	8501	II
1-Piperidinocyclohexanecarbonitrile .....	8603	II
Alphaprodine .....	9010	II
Anileridine .....	9020	II
Cocaine .....	9041	II
Codeine .....	9050	II
Etorphine HCl .....	9059	II
Dihydrocodeine .....	9120	II
Oxycodone .....	9143	II
Hydromorphone .....	9150	II
Diphenoxylate .....	9170	II
Ecgonine .....	9180	II
Ethylmorphine .....	9190	II
Hydrocodone .....	9193	II
Levomethorphan .....	9210	II
Levorphanol .....	9220	II
Isomethadone .....	9226	II
Meperidine .....	9230	II
Meperidine-intermediate-A .....	9232	II
Meperidine intermediate-B .....	9233	II
Meperidine intermediate-C .....	9234	II
Metazocine .....	9240	II
Methadone .....	9250	II
Methadone intermediate .....	9254	II
Metopon .....	9260	II
Dextropropoxyphene, bulk (non-dosage forms) .....	9273	II
Morphine .....	9300	II
Oripavine .....	9330	II
Thebaine .....	9333	II
Dihydroetorphine .....	9334	II
Levo-alphaacetylmethadol .....	9648	II
Oxymorphone .....	9652	II
Noroxymorphone .....	9668	II
Phenazocine .....	9715	II
Thiafentanil .....	9729	II
Piminodine .....	9730	II
Racemethorphan .....	9732	II
Racemorphan .....	9733	II
Alfentanil .....	9737	II
Remifentanil .....	9739	II
Sufentanil .....	9740	II
Carfentanil .....	9743	II
Tapentadol .....	9780	II
Bezitramide .....	9800	II
Fentanyl .....	9801	II
Moramide-intermediate .....	9802	II

The company plans to import analytical reference standards for distribution to its customers for research and analytical purposes. Placement of these drug codes onto the company's registration does not translate into automatic approval of subsequent permit applications to import controlled substances. Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized in 21 U.S.C. 952(a)(2). Authorization will not extend to the import of FDA approved or non-approved finished dosage forms for commercial sale.

Dated: June 18, 2019.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2019-14026 Filed 7-1-19; 8:45 am]

**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

[OMB Number 1125-0001]

### Agency Information Collection Activities; Proposed Collection; Comments Requested; Application for Cancellation of Removal (42A) for Certain Permanent Residents; and Application for Cancellation of Removal and Adjustment of Status (42B) for Certain Nonpermanent Residents

**AGENCY:** Executive Office for Immigration Review, Department of Justice.

**ACTION:** 30-Day notice.

**SUMMARY:** The Department of Justice (DOJ), Executive Office for Immigration Review, 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.

**DATES:** Comments are encouraged and will be accepted for an additional 30 days until August 1, 2019.

**FOR FURTHER INFORMATION CONTACT:** If you have additional 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 Lauren Alder Reid, Assistant Director, Office of Policy, Executive Office for Immigration Review, 5107 Leesburg Pike, Suite 2500, Falls Church, VA 22041, telephone: (703) 305-0289. Written comments and/or suggestions can also be directed to the Office of Management and Budget, Office of

Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20530 or sent to [OIRA\\_submissions@omb.eop.gov](mailto:OIRA_submissions@omb.eop.gov).

**SUPPLEMENTARY INFORMATION:** 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/or
- 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:* Extension with changes to a currently approved collection.

2. *The Title of the Form/Collection:* Application for Cancellation of Removal for Certain Permanent Residents; and Application for Cancellation of Removal and Adjustment of Status for Certain Nonpermanent Residents.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* The form numbers are EOIR-42A and EOIR-42B, Executive Office for Immigration Review, United States Department of Justice.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: Individual aliens determined to be removable from the United States. Other: None. Abstract: This information collection is necessary to determine the statutory eligibility of individual aliens who have been determined to be removable from the United States for cancellation of their removal, as well as to provide information relevant to a favorable exercise of discretion.

5. *An estimate of the total number of respondents and the amount of time*

*estimated for an average respondent to respond:* It is estimated that 27,999 respondents will complete the form annually with an average of 5 hours and 50 minutes per response.

6. *An estimate of the total public burden (in hours) associated with the collection:* The estimated public burden associated with this collection is 162,394 hours.

*If additional information is required contact:* Melody Braswell, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE, 3E.405B, Washington, DC 20530.

Dated: June 27, 2019.

**Melody D. Braswell,**

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

[FR Doc. 2019-14064 Filed 7-1-19; 8:45 am]

**BILLING CODE 4410-30-P**

## NUCLEAR REGULATORY COMMISSION

[Docket Nos. 52-025 and 52-026; NRC-2008-0252]

### Southern Nuclear Operating Company, Inc.; Vogtle Electric Generating Plant, Units 3 and 4; Passive Residual Heat Removal Instrumentation Minimum Inventory Displays

**AGENCY:** Nuclear Regulatory Commission.

**ACTION:** Exemption and combined license amendment; issuance.

**SUMMARY:** The U.S. Nuclear Regulatory Commission (NRC) is granting an exemption to allow a departure from the certification information of Tier 1 of the generic design control document (DCD) and is issuing License Amendment Nos. 162 and 160 to Combined Licenses (COLs), NPF-91 and NPF-92. The COLs were issued to Southern Nuclear Operating Company, Inc., and Georgia Power Company, Oglethorpe Power Corporation, MEAG Power SPVM, LLC, MEAG Power SPVJ, LLC, MEAG Power SPVP, LLC, and the City of Dalton, Georgia (collectively SNC); for construction and operation of the Vogtle Electric Generating Plant (VEGP) Units 3 and 4, located in Burke County, Georgia.

The granting of the exemption allows the changes to Tier 1 information asked for in the amendment. Because the acceptability of the exemption was determined in part by the acceptability of the amendment, the exemption and