SECTION 3

APPLICATION FOR REGISTRATION

Under the Controlled Substances Act

INSTRUCTIONS	Save time - apply on-line at www.deadiversion.usdoj.gov	DEA OFFICIAL USE:					
	 To apply by mail complete this application. Keep a copy for your records. Mail this form to the address provided in Section 7 or use enclosed envelope. 	Do you have other DEA registration numbers?					
	 The "MAIL-TO ADDRESS" can be different than your "PLACE OF BUSINESS" address. If you have any questions call 800-882-9539 prior to submitting your application. 						
	IMPORTANT: DO NOT SEND THIS APPLICATION AND APPLY ON-LINE.						
MAIL-TO ADDRESS	Please print mailing address changes to the right of the address in this box.	FEE FOR ONE (1) YEAR - see Section 2 FEE IS NON-REFUNDABLE					
SECTION 1 APP	LICANT IDENTIFICATION	n Business Registration					
	Name of individual -OR- Business or Facility Name)						
Name 2 (First	Name and Middle Name of individual - OR- Continuation of business name)						
PLACE OF BUSIN	ESS Street Address Line 1						
PLACE OF BUSIN	ESS Address Line 2						
City		State Zip Code					
Business Phone N	umber Point of Contact						
Business Fax Num	ber Email Address						
DEBT COLLECTION INFORMATION	Tax Identification Number (<i>if registration is for business</i>)	Social Security Number (if registration is for individual					
Mandatory pursuant to Debt Collection Improvements Act	Provide TIN or SSN. See additional information note #3 on page 4.						
SECTION 2		· · · · · · · · · · · · · · · · · · ·					
BUSINESS ACTIVITY		xporterfee for one year is \$1523					
Check one business activity		nporterfee for one year is \$1523					
box only Researcher	Researcher w/Sched II - Vfee for one year is \$244	everse Distributorfee for one year is \$1523					
Researcher - See page 4 for required	Canine Handlerfee for one year is \$244	anufacturerfee for one year is \$3047					
attachments	Distributorfee for one year is \$1523	anufacturer BULKfee for one year is \$3047					

A. DRUG SCHEDULES	List 1 (L1) - manufacturers &	Schedule 2 Narcotic	Schedule 3 Narcotic	Schedule 4
Check all that apply	Schedule 1	Schedule 2 Non-Narcotic (2N)	Schedule 3 Non-Narcotic (3)	N) C Schedule 5
Enter drug codes on page 2.	Check this box if you require official	order forms - for purchase of so	chedule 2 controlled substances.	
B. MANUFACTURERS ONLY Mark each box with an 'X' to indicate which	1 2 2 NON 3 3 NON 4 5	STAGE 1 Bulk synthesis/extraction	1 2 2 NON 3 3 NON 4 5 narcotic 3 narcotic 4 5	STAGE 3 Package / Repackage Label / Relabel
drug schedule is handled	1 2 2 NON 3 3 NON 4 5	STAGE 2 Dosage form manufacture	1 2 2 NON 3 3 NON 4 5 harcotic 3 harcotic 4 5	STAGE 4 Non-human consumption NEW - Page 1

SECTION 4 STATE LICENSE(S)	You MUST be currently author in the schedules for which yo	bu are applying under the law	vs of the state	or juriso	diction in	which you	are operating	g or prop	ose to op	berate.
Be sure to include both state license numbers if applicable	State License Number						Expiration Date (REQUIRED)		/	
		What state issued this licen:	se ?					MM -	DD- YY	ΥY
	State Controlled Substance License Number (if required)						Expiration Date (if required)	/	/	2.0.4
	(- 1)	What state issued this licen:	se ?				(MM -	DD-YY	ΥY
SECTION 5									YES	NO
0	as the applicant ever been con r been excluded or directed to b ction pending?	victed of a crime in connect be excluded from participation	ction with cont on in a medica	rolled su re or sta	bstance te health	(s) under s care prog	tate or federa ram, or is any	al law, y such		
D IMPORTANT	ate(s) of incident MM-DD-YYY	Y:							YES	NO
2. H All questions in re this section must	as the applicant ever surrender estricted, or denied, or is any su	uch action pending?	eral controlled	substar	ice regist	ration revo	oked, suspen	ded,		
be answered.	ate(s) of incident MM-DD-YYY								YES	NO
	as the applicant ever surrender evoked, suspended, denied, res		e professiona on, or is any s	l license such acti	or contro on pendi	olled subst ng?	ance registra	tion		
	ate(s) of incident MM-DD-YYY								YES	NO
re	the applicant is a corporation artnership, or pharmacy, has ar ontrolled substance(s) under st. ggistration revoked, suspended ggistration revoked, suspended	, restricted, denied, or ever r	nad a state pr	otession	ai iicense	e or contro	lied substand	ition, ion with stance se		
D	ate(s) of incident MM-DD-YYY	Y:	No	te: If que	estion 4 d	does not a cessing of	pply to you, b your applicat	e sure to	mark 'No	O'. blank
EXPLANATION OF "YES" ANSWERS										
Applicants who have	Liability question #	Location(s) of incide	ent:						_	
answered "YES" to any of the four questions	Nature of incident:									
above must provide a statement to explain each "YES" answer.										
each i Lo answei.										
Use this space or attach a separate sheet and return with application	Disposition of incident:									
Use this space or attach a separate sheet and return with application		IFEE								
Use this space or attach a separate sheet and return with application	Disposition of incident:		nent official or	institutio	n. Does	not apply	to contractor-	operated	l institutio	ons.
Use this space or attach a separate sheet and return with application SECTION 6 EXEL Check	Disposition of incident:	deral, state, or local governm				,		•	l institutio	ons.
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Print or type name and title of applicant

WARNING: 21 USC 843(d), states that any person who knowingly or intentionally furnishes false or fraudulent information in the application is subject to a term of imprisonment of not more than 4 years, and a fine under Title 18 of not more than \$250,000, or both.

Listed below are examples of schedules 1-5 a For more information, see our website at www					
Canine Handler must mark schedule 1		stributor	must mark all schedule 1, drug code 2012		
Exporter must mark all schedule 1-5		tributor must mark all schedule 1, drug code 2012			
Importer must mark all schedule 1-5 & List 1 co			w/Sched 1 must mark schedule 1		ļ
Manufacturer must mark all schedule 1, 2 & List 1 cod	des Re	searcher	w/Sched 2-5 must mark schedule 2 to be manufactured or imported as part of research	1	
If you bulk manufacture a s	ubstance, cl	neck the	'BULK?' column after the applicable class code.		
CHEDULE 1 NARCOTIC & NON-NARCOTIC		BULK?	SCHEDULE 2 NARCOTIC & NON-NARCOTIC	CODE	BULK
3,4-Methylenedioxyamphetamine (MDA)	7400		Amobarbital (Amytal, Tuinal)	2125	
3,4-Methylenedioxymethamphetamine (MDMA)	7405		Amphetamine (Dexedrine, Adderall)	1100	
4-Methyl - 2,5 - Dimethoxyamphetamine (DOM, STP)	7395		Cocaine (Methyl benzoylecgonine)	9041	
4-Methylaminorex (cis isomer) (U4Euh, McN-422)	1590		Codeine (Morphine methyl ester)	9050	
Alphacetylmethadol (except LAAM)	9603		Dextropropoxyphene (bulk)	9273	
Bufotenine (Mappine)	7433		Diphenoxylate	9170	
Marihuana / Cannabidiol	7360/7372		Fentanyl (Duragesic)	9801	
Diethyltryptamine (DET) (7434		Hydrocodone (Dihydrocodeinone)	9193	
Difenoxin 1MG/25UG AtSO4 /DU (Motofen)	9167		Hydromorphone (Diaudid)	9150	
Dimethyltryptamine (DMT)	7435		Levo-Alphacetylmethadol (LAAM)	9648	
Etorphine (except HCL)	9056		Levorphanol (Levo-Dromoran)	9220	
Gamma Hydroxybutyric Acid (GHB)	2010		Meperidine (Demerol, Mepergan)	9230	
Heroin (Diamorphine)	9200		Methadone (Dolophine, Methadose)	9250	
Ibogaine	7260		Methamphetamine (Desoxyn)	1105	
Lysergic acid diethylamide (LSD)	7315		Methylphenidate (Concerta, Ritalin)	1724	
Mescaline	7381		Morphine (MS Contin, Roxanol)	9300	
Marihuana	7360		Opium, powdered	9639	
Methaqualone (Quaalude)	2565		Oxycodone (Oxycontin, Percocet)	9143	
Normorphine	9313		Oxymorphone (Numorphan)	9652	
Peyote	7415		Pentobarbital (bulk) (Nembutal)	2270	
Psilocybin	7437		Phencyclidine (PCP)	7471	
Tetrahydrocannabinols (THC)	7370		Secobarbital (Seconal, Tuinal)	2315	
CHEDULE 3 NARCOTIC & NON-NARCOTIC	CODE	BULK?	SCHEDULE 4 NARCOTIC & NON-NARCOTIC	CODE	BUL
Anabolic Steroids	4000		Alprazolam (Xanax	2882	
Barbituric acid derivative	2100		Barbital (Veronal, Plexonal)	2145	
Benzphetamine (Didrex, Inapetyl)	1228		Chloral Hydrate (Noctec)	2465	
Buprenorphine (Buprenex, Temgesic)	9064		Chlordiazepoxide (Librium)	2744	
Butabarbital	2100/2175		Clonazepam (Klonopin)	2737	
Butalbital	2100/2165		Clorazepate (Tranxene)	2768	
Codeine combo product (Empirin)	9804		Diazepam (Valium)	2765	
Dihydrocodeine combo product (Compal)	9807		Flurazepam (Dalmane)	2767	
Dronabinol in sesame oil soft cap (Marinol)	7369		Lorazepam (Ativan)	2885	
Gamma-Hydroxybutyric Acid preparations (Zyrem)	2012		Meprobamate (Milltown, Equanil)	2820	
Hydrocodone combo products (Lorcet, Vicodin)	9806		Midazolam (Versed)	2884	
Ketamine (Ketaset, Ketalar)	7285		Oxazepam (Serax, Serenid-D)	2835	
Morphine combo product	9810		Phenobarbital (Fastin, Zantryl)	2285	
Nalorphine (Nalline)	9400		Phentermine	1640	
Opium combo product (Paregoric)	9809		Temazepam (Restoril)	2925	
Pentobarbital suppository dosage (FP3)	2270		Zolpidem (Ambien, Stilnox)	2783	
Phendimetrazine (Plegine, Bontril)	1615		LIST 1 REGULATED CHEMICALS	CODE	BUL
Thiopental	2100/2329		** ONLY manufacturers & importers may select List 1		
HEDULE 5 NARCOTIC & NON-NARCOTIC	CODE	BULK?	Ephedrine	8113	
Codeine preparations (Robitussin A-C, Pediacof)	9050		Phenylpropanolamine	1225	
Pyrovalerone (Centroton, Thymergix)	1485		Pseudoephedrine	8112	

APPLICATION FOR REGISTRATION

SECTION 1. APPLICANT IDENTIFICATION - Information must be typed or printed in the blocks provided to help reduce data entry errors. A physical address is required in address line 1; a post office box or continuation of address may be entered in address line 2. Fee exempt applicant must list the address of the fee exempt institution. Applicant must enter a valid social security number (SSN), or a tax identification number (TIN) if applying as a business entity.

Debt collection information is mandatory pursuant to the Debt Collection Improvement Act of 1996.

SECTION 2. BUSINESS ACTIVITY - Indicate only one. Each type of business activity requires a separate application. You are required to register as a "manufacturer" if you manufacture a controlled substance or list 1 chemical and then distribute it.

SECTION 3A. SCHEDULES - Applicant should check all schedules to be handled. However, applicant must still comply with state requirements; federal registration does not overrule state restrictions. Check the order form box only if you intend to purchase or to transfer schedule 1 and 2 controlled substances. Order forms will be mailed to the registered address following issuance of a Certificate of Registration.

3B. MANUFACTURER ONLY - Mark the chemical/controlled substance schedule(s) handled in each manufacturing stage listed.

3C. SCHEDULE CODES - Report all chemical/drug codes as required for your business activity. Controlled substances manufacturers and importers must obtain a separate chemical registration if they handle chemicals other than an FDA-approved drug product containing 1225, 8112, or 8113.

SECTION 4. STATE LICENSE(S) - Federal registration by DEA is based upon the applicant's compliance with applicable state and local laws. Applicant should contact the local state licensing authority prior to completing this application. If your state requires a license, provide that number on this application.

SECTION 5. LIABILITY - Applicant must answer all four questions for the application to be accepted for processing. If you answer "Yes" to a question, provide an explanation in the space provided. If you answer "Yes" to several questions, then you must provide a separate explanation describing the date, location, nature, and result of each incident. If additional space is required, you may attach a separate page.

SECTION 6. EXEMPTION - Exemption from payment of application fee is limited to federal, state or local government official or institution. The applicant's superior or agency officer must certify exempt status. The signature, authority title, and telephone number of the certifying official (other than the applicant) must be provided. The address of the fee exempt institution must appear in Section 1.

SECTION 7. METHOD OF PAYMENT - Indicate the desired method of payment. Make checks payable to "Drug Enforcement Administration". Third-party checks or checks drawn on foreign banks will not be accepted. FEES ARE NON-REFUNDABLE.

SECTION 8. APPLICANT'S SIGNATURE - Applicant MUST sign in this section or application will be returned. Card holder signature in section 7 does not fulfill this requirement.

ATTACHMENTS: Researcher or canine handler must attach 3 copies of protocol, including curriculum vitae, to conduct research with schedule 1 controlled substances. For clinical investigations, researcher must first submit to FDA a "Notice of Claimed Investigational Exemption for New Drug (IND)". See DEA web site or CFR 1301.18 for details.

NOTICE TO REGISTRANTS MAKING PAYMENT BY CHECK

Authorization to Convert Your Check: If you send us a check to make your payment, your check will be converted into an electronic fund transfer. "Electronic fund transfer" is the term used to refer to the process in which we electronically instruct your financial institution to transfer funds from your account to our account, rather than processing your check. By sending your completed, signed check to us, you authorize us to copy your check and to use the account information from your check to make an electronic fund transfer from your account for the same amount as the check. If the electronic fund transfer cannot be processed for technical reasons, you authorize us to process the copy of your check.

Insufficient Funds: The electronic funds transfer from your account will usually occur within 24 hours, which is faster than a check is normally processed. Therefore, make sure there are sufficient funds available in your checking account when you send us your check. If the electronic funds transfer cannot be completed because of insufficient funds, we may try to make the transfer up to two more times.

Transaction Information: The electronic fund transfer from your account will be on the account statement you receive from your financial institution. However, the transfer may be in a different place on your statement than the place where your checks normally appear. For example, it may appear under "other withdrawals" or "other transactions". You will not receive your original check back from your financial institution. For security reasons, we will destroy your original check, but we will keep a copy of the check for record-keeping purposes.

Your Rights: You should contact your financial institution immediately if you believe that the electronic fund transfer reported on your account statement was not properly authorized or is otherwise incorrect. Consumers have protections under Federal law called the Electronic Fund Transfer Act for an unauthorized or incorrect electronic fund transfer.

ADDITIONAL INFORMATION

No registration will be issued unless a completed application form has been received (21 CFR 1301.13).

In accordance with the Paperwork Reduction Act of 1995, no person is required to respond to a collection of information unless it displays a valid OMB control number. The OMB number for this collection is 1117-0012. Public reporting burden for this collection of information is estimated to average 12 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the information.

The Debt Collection Improvements Act of 1996 (31 U.S.C. §7701) requires that you furnish your Taxpayer Identification Number (TIN) or Social Security Number (SSN) on this application. This number is required for debt collection procedures if your fee is not collectible.

PRIVACY ACT NOTICE: Providing information other than your SSN or TIN is voluntary; however, failure to furnish it will preclude processing of the application. The authorities for collection of this information are §§302 and 303 of the Controlled Substances Act (CSA) (21 U.S.C. §§822 and 823). The principle purpose for which the information will be used is to register applicants pursuant to the CSA. The information may be disclosed to other Federal law enforcement and regulatory agencies for law enforcement and regulatory purposes, State and local law enforcement and regulatory agencies for law enforcement and regulatory purposes, and person registered under the CSA for the purpose of verifying registration. For further guidance regarding how your information may be used or disclosed, and a complete list of the routine uses of this collection, please see the DEA System of Records Notice "Controlled Substances Act Registration Records" (DEA-005), 52 FR 47208, December 11, 1987, as modified.

Your Local DEA Office CONTACT INFORMATION

All offices are listed on web site (800, 877, and 888 are toll-free

INTERNET: www.deadiversion.usdoj.gov

TELEPHONE: HQ Call Center (800) 882-9539

WRITTEN INQUIRIES: DEA Attn: Registration Section/ODR

P.O. Box 2639 Springfield, VA 22152-2639