HOUSE HEALTH AND HUMAN SERVICES COMMITTEE SUBSTITUTE FOR HOUSE BILL 139

53rd Legislature - STATE OF NEW MEXICO - second session, 2018

.210165.2

AN ACT

RELATING TO PHARMACEUTICALS; AMENDING A SECTION OF THE

CONTROLLED SUBSTANCES ACT TO REMOVE FROM LIABILITY UNDER THE

CONTROLLED SUBSTANCES ACT CERTAIN ACTIVITIES RELATING TO

PRESCRIPTION DRUGS CONTAINING A MARIJUANA DERIVATIVE; PROVIDING

FOR A CONTINGENT EFFECTIVE DATE; AUTHORIZING RECONCILIATION OF

MULTIPLE AMENDMENTS TO THE SAME SECTION OF LAW.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:

SECTION 1. Section 30-31-6 NMSA 1978 (being Laws 1972, Chapter 84, Section 6, as amended) is amended to read:

"30-31-6. SCHEDULE I.--The following controlled substances are included in Schedule I:

A. any of the following opiates, including their isomers, esters, ethers, salts, and salts of isomers, esters and ethers, unless specifically exempted, whenever the

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1	existence of these isomers, esters, ethers and salts is				
2	possible within the specific chemical designation:				
3	(1) acetylmethadol;				
4	(2) allylprodine;				
5	(3) alphacetylmethadol;				
6	(4) alphameprodine;				
7	(5) alphamethadol;				
8	(6) benzethidine;				
9	(7) betacetylmethadol;				
10	(8) betameprodine;				
11	(9) betamethadol;				
12	(10) betaprodine;				
13	(11) clonitazene;				
14	(12) dextromoramide;				
15	(13) dextrorphan;				
16	(14) diampromide;				
17	(15) diethylthiambutene;				
18	(16) dimenoxadol;				
19	(17) dimepheptanol;				
20	(18) dimethylthiambutene;				
21	(19) dioxaphetyl butyrate;				
22	(20) dipipanone;				
23	(21) ethylmethylthiambutene;				
24	(22) etonitazene;				
25	(23) etoxeridine;				

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underscored material = new
[bracketed material] = delete

1	(24) furethidine;		
2	(25) hydroxypethidine;		
3	(26) ketobemidone;		
4	(27) levomoramide;		
5	(28) levophenacylmorphan;		
6	(29) morpheridine;		
7	(30) noracymethadol;		
8	(31) norlevorphanol;		
9	(32) normethadone;		
10	(33) norpipanone;		
11	(34) phenadoxone;		
12	(35) phenampromide;		
13	(36) phenomorphan;		
14	(37) phenoperidine;		
15	(38) piritramide;		
16	(39) proheptazine;		
17	(40) properidine;		
18	(41) racemoramide; and		
19	(42) trimeperidine;		
20	B. any of the following opium derivatives, their		
21	salts, isomers and salts of isomers, unless specifically		
22	exempted, whenever the existence of these salts, isomers and		
23	salts of isomers is possible within the specific chemical		
24	designation:		
25	(1) acetorphine;		

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1	(2)	acetyldihydrocodeine;	
2	(3)	benzylmorphine;	
3	(4)	codeine methylbromide;	
4	(5)	codeine-N-oxide;	
5	(6)	cyprenorphine;	
6	(7)	desomorphine;	
7	(8)	dihydromorphine;	
8	(9)	etorphine;	
9	(10)	heroin;	
10	(11)	hydromorphinol;	
11	(12)	methyldesorphine;	
12	(13)	methyldihydromorphine;	
13	(14)	morphine methylbromide;	
14	(15)	morphine methylsulfonate;	
15	(16)	morphine-N-oxide;	
16	(17)	myrophine;	
17	(18)	nicocodeine;	
18	(19)	nicomorphine;	
19	(20)	normorphine;	
20	(21)	pholcodine; and	
21	(22)	thebacon;	
22	C. any ma	terial, compound, mixture or preparation	
23	that contains any quantity of the following hallucinogenic		
24	substances, their sal	ts, isomers and salts of isomers, unless	
25	specifically exempted	, whenever the existence of these salts,	

underscored material = new
[bracketed material] = delete

1	isomers and salts of isomers is possible within the specific				
2	chemical designation:				
3	(1) 3,4-methylenedioxy amphetamine;				
4	(2) 5-methoxy-3,4-methylenedioxy amphetamine;				
5	(3) 3,4,5-trimethoxy amphetamine;				
6	(4) bufotenine;				
7	(5) diethyltryptamine;				
8	(6) dimethyltryptamine;				
9	(7) 4-methyl-2,5-dimethoxy amphetamine;				
10	(8) ibogaine;				
11	(9) lysergic acid diethylamide;				
12	(10) marijuana;				
13	(11) mescaline;				
14	(12) peyote, except as otherwise provided in				
15	the Controlled Substances Act;				
16	(13) N-ethyl-3-piperidyl benzilate;				
17	(14) N-methyl-3-piperidyl benzilate;				
18	(15) psilocybin;				
19	(16) psilocyn;				
20	(17) tetrahydrocannabinols;				
21	(18) hashish;				
22	(19) synthetic cannabinoids, including:				
23	(a) l-[2-(4-(morpholinyl)ethyl]				
24	-3-(1-naphthoy1)indole;				
25	(b) 1-butyl-3-(1-napthoy1)indole;				
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1
                                   1-hexy1-3-(1-naphthoy1)indole;
                             (c)
 2
                             (d)
                                   1-penty1-3-(1-naphthoy1)indole;
 3
                             (e)
                                   1-penty1-3-(2-methoxyphenylacety1)
 4
      indole;
 5
                             (f)
                                  cannabicyclohexanol (CP 47, 497 and
      homologues: 5-(1,1-dimethylheptyl)-2-[(1R,3S)
 6
 7
       -3-hydroxycyclohexyl]-phenol (CP-47,497); and 5-(1,
       1-dimethyloctyl)-2-[(lR,3S)-3-hydroxycyclohexyl]-phenol;
 8
 9
                             (g) 6aR, 10aR) - 9 - (hydroxymethy1)
      -6,6-dimethyl-3-(2-methyloctan-2-yl)-6a,7,10,
10
       10a-tetrahydrobenzo[c]chromen-1-o1);
11
12
                             (h)
                                  dexanabinol, (6aS, 10aS)
       -9-(hydroxymethyl)-6,6-dimethyl-3-(2-methyloctan-2-yl)
13
      -6a,7,10,10a-tetrahydrobenzo[c]chromen-1-ol;
14
                             (i)
                                   1-penty1-3-(4-chloro naphthoy1)
15
      indole;
16
                                   (2-methyl-l-propyl-lH-indol-3-yl)
                             (j)
17
       -1-naphthalenyl-methanone; and
18
                                  5-(1,1-dimethylheptyl)-2-(3-hydroxy
19
      cyclohexyl)-phenol;
20
                              3,4-methylenedioxymethcathinone;
                        (20)
21
                              3,4-methylenedioxypyrovalerone;
                        (21)
22
                        (22)
                              4-methylmethcathinone;
23
                        (23)
                              4-methoxymethcathinone;
24
                        (24)
                              3-fluoromethcathinone; and
25
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(25) 4-fluoromethcathinone;

- D. the enumeration of peyote as a controlled substance does not apply to the use of peyote in bona fide religious ceremonies by a bona fide religious organization, and members of the organization so using peyote are exempt from registration. Any person who manufactures peyote for or distributes peyote to the organization or its members shall comply with the federal Comprehensive Drug Abuse Prevention and Control Act of 1970 and all other requirements of law;
- E. the enumeration of marijuana, tetrahydrocannabinols or chemical derivatives of tetrahydrocannabinol as Schedule I controlled substances does not apply to:
- (1) the use of marijuana, tetrahydrocannabinols or chemical derivatives of tetrahydrocannabinol by certified patients pursuant to the Controlled Substances Therapeutic Research Act or by qualified patients pursuant to the provisions of the Lynn and Erin Compassionate Use Act; [and] or
- (2) the use, dispensing, possession, prescribing, storage or transport of a prescription drug that the United States food and drug administration has approved and that contains marijuana, a tetrahydrocannabinol derivative or a chemical derivative of tetrahydrocannabinol; and
- F. controlled substances added to Schedule I by .210165.2

rule adopted by the board pursuant to Section 30-31-3 NMSA 1978."

SECTION 2. TEMPORARY PROVISION--COMPILATION INSTRUCTION--RECONCILIATION.--If acts making amendments to Section 30-31-6 NMSA 1978 are enacted by the first and second sessions of the fifty-third legislature, the provisions of those acts shall be reconciled and compiled in accordance with the provisions of Section 12-1-8 NMSA 1978, notwithstanding that the amendments were not made in the same session of the legislature.

SECTION 3. CONTINGENT EFFECTIVE DATE--NOTIFICATION.--The effective date of the provisions of this act is thirty days following the date that the board of pharmacy certifies to the New Mexico compilation commission and the director of the legislative council service that the United States food and drug administration has approved one or more drugs containing a marijuana derivative. The board of pharmacy shall notify the New Mexico compilation commission and the director of the legislative council service immediately upon the board's knowledge that the United States food and drug administration has approved for the first time a drug containing a marijuana derivative.

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