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(54) **Eljárások transz-(-)-delta9-tetrahidrokannabinol és transz-(+)-delta9-tetrahidrokannabinol tisztítására**

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(54) METHODS FOR PURIFYING TRANS-(-)-DELTA9-TETRAHYDROCANNABINOL AND TRANS-(+)-DELTA9-TETRAHYDROCANNABINOL

VERFAHREN ZUR REINIGUNG VON TRANS-(-)-DELTA9-TETRAHYDROCANNABINOL UND TRANS-(+)-DELTA9-TETRAHYDROCANNABINOL

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- **LEVIN S ET AL: "Resolution of chiral cannabinoids on amylose tris(3,5-dimethylphenylcarbamate) chiral stationary phase: effects of structural features and mobile phase additives", JOURNAL OF CHROMATOGRAPHY A, ELSEVIER, AMSTERDAM, NL, vol. 654, 1993, pages 53-64, XP002324648, ISSN: 0021-9673**
- **FAHRENHOLTZ K E ET AL: "Total synthesis of (.+-.)-.DELTA.9-tetrahydrocannabinol and four of its isomers", JOURNAL OF THE AMERICAN CHEMICAL SOCIETY, AMERICAN CHEMICAL SOCIETY, WASHINGTON, DC; US, vol. 89, no. 23, 8 November 1967 (1967-11-08), pages 5934-5941, XP002274651, ISSN: 0002-7863, DOI: DOI: 10.1021/JA00999A034**

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(Cont. next page)

- TURK R F ET AL: "The identification, isolation, and preservation of delta-9-tetrahydrocannabinol (delta-9-THC)", JOURNAL OF PHARMACY AND PHARMACOLOGY, ROYAL PHARMACEUTICAL SOCIETY OF GREAT BRITAIN, GB, vol. 23, no. 3, 1 March 1971 (1971-03-01), pages 190-195, XP009098029, ISSN: 0022-3573

Description

1. Field of the Invention

[0001] The present invention relates to methods for purifying trans-(-)-Δ^9-tetrahydrocannabinol and/or trans-(+)-Δ^9-tetrahydrocannabinol.

2. Background of the Invention

[0002] (-)-6a,10a-Trans-Δ^9-tetrahydrocannabinol ("(-)-Δ^9-THC") is mainly responsible for the antiemetic effects associated with cannabis (S.E. Sallen et al., N. Engl. J. Med. 302:135 (1980); A.E. Chang et al., Cancer 47:1746 (1981); and D.S. Poster et al., J. Am. Med. Asso. 245:2047 (1981)). Both trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC, the trans-(-)- and (+)-enantiomers, respectively, of (±)-Δ^9-THC, are reported to be useful for treating pain, with trans-(-)-Δ^9-THC reported to be more potent than trans-(+)-Δ^9-THC *(see, e.g.,* G. Jones et al., Biochem. Pharmacol. 23:439 (1974); S.H. Roth, Can. J. Physiol. Pharmacol. 56:968 (1978); B.R. Martin et al., Life Sciences 29:565 (1981); M. Reichman et al., Mol. Pharmacol. 34:823 (1988); and M. Reichman et al., Mol. Pharmacol. 40:547 (1991)). trans-(-)-Δ^9-THC is reported to be useful as antiemetic to relieve nausea and vomiting in patients receiving cancer chemotherapy and to stimulate weight gain in patients suffering from symptomatic HIV infection (*see* U.S. Patent No. 6,703,418 B2 to Plasse). An encapsulated formulation of synthetic trans-(-)-Δ^9-THC ("dronabinol") in sesame oil is currently sold as Marinol® by Unimed Pharmaceuticals, Inc., in 2.5, 5, and 10 mg dosage strengths.

[0003] Trans-(-)-Δ^9-THC can be extracted from hashish *(see,* Y. Gaoni et al., J. Am. Chem. Soc. 93:217 (1971); and U.S. Patent No. 6,365,416 B1 to Elsohly et al.). The concentration of trans-(-)-Δ^9-THC in hashish, however, ranges from only about 1-5% depending on the source, and, even after extraction, trans-(-)-Δ^9-THC must be separated from other impurities such as cannabinoid isomers.

[0004] R.F. Turk et al., J. Pharm. Pharmac. 23:190-195 (1971) describes a method for isolating trans-(-)-Δ^9-THC from marihuana, but the product contained an undetermined amount of carboxylic precursors of THC.

[0005] The following paragraphs relate to known methods that purport making trans-(-)-Δ^9-THC or (±)-Δ^9-THC:

U.S. Patent No. 3,560,528 to Petrzilka describes the reaction of (+)-*p*-mentha-2,8-dien-1-ol with olivetol in the presence of *p*-toluenesulfonic acid monohydrate ("PTSA·H_2O") or trifluoroacetic acid in refluxing benzene to provide (-)-Δ^8-THC, which can be converted to trans-(-)-Δ^9-THC by addition of HCl followed by dehydrochlorination *(see* Y. Mcchoulam et al., J. Am. Chem. Soc. 89:4553 (1967); and R. Mechoulam et al., J. Am. Chem. Soc. 94:6159 (1972)).

U.S. Patent No. 4,025,516 to Razdan et al. describes the reaction of a mixture of *cis/trans*-(+)-*p*-mentha-2,8-dien-1-ol with olivetol in an inert organic solvent in the presence of an excess of a non-alkaline dehydrating agent and an acid catalyst to form trans-(-)-Δ^9-THC; this patent also describes the reaction of (-)-cannabidiol ("(-)-CBD") or (-)-abnormal-CBD ("(-)-abn-CBD") with a Lewis acid such as boron trifluoride diethylether ("BF_3-Et_2O") in an inert solvent under anhydrous conditions to form trans-(-)-Δ^9-THC.

R. K. Razdan et al., J. Am. Chem. Soc. 96:5860 (1974) describes the reaction of a mixture of *cis/trans*-(+)-p-mentha-2,8-dien-1-ol with olivetol in the presence of 1% BF_3·Et_2O, methylene chloride and anhydrous magnesium sulfate to form trans-(-)-Δ^9-THC.

U.S. Patent No. 4,381,399 to Olsen et al. describes a method for separating trans-(-)-Δ^9-THC from a crude synthetic mixture, the method comprising esterifying the crude mixture, isolating the resultant trans-(-)-Δ^9-THC ester, hydrolyzing the ester, and distilling trans-(-)-Δ^9-THC at reduced pressure.

K.E. Fahrenholtz et al., J. Am. Chem. Soc. 89:5934-5941 (1967) describes the hydrolysis of (±)-1-*m*-nitrobenzenesulfoanate-6a,10a-trans-Δ^9-tetrahydrocannabinol with NaOH in aqueous methanol to provide (±)-Δ^9-THC, which was subsequently crystallized from hexane.

E.G. Taylor et al., J. Am. Chem. Soc. 88:367 (1966) describes the reaction of citral with olivetol in acidified ethanol to form (±)-Δ^9-THC in about 35% yield.

S. L. Levin et al., J. Chromatogr. A 654:53-64 (1993) describe a method for resolving trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC from a composition comprising equimolar amounts of the trans-(-)- and (+)-enantiomer.

WO 2005/061480 describes a process for purifying trans-(-)-Δ^9-THC which comprises at least one chromatographic step.

[0006] Despite these described methods, there remains a need for improved methods for making trans-(-)-Δ^9-THC in pure or substantially pure form.

3. Summary of the Invention

[0007] The invention relates in a first embodiment to:

A method for removing impurities from a composition comprising trans-(-)-Δ^9-tetrahydrocannabinol and trans-(+)-Δ^9-tetrahydrocannabinol prior to crystallization of (±)-Δ^9-tetrahydrocannabinol, the method comprising:

(a) contacting the trans-(-)-Δ^9-tetrahydrocannabinol and trans-(+)-Δ^9-tetrahydrocannabinol with a first water immiscible organic solvent, a water-miscible alcohol, water, and an alkali metal hydroxide, thus forming a biphasic composition comprising (i) a first organic phase, and (ii) an alcoholic-caustic phase containing the trans-(-)-Δ^9-tetrahydrocannabinol and the trans-(+)-Δ^9-tetrahydrocannabinol;
(b) separating the alcoholic-caustic phase containing the trans-(-)-Δ^9-tetrahydrocannabinol and the trans-(+)-Δ^9-tetrahydrocannabinol from the first organic phase;
(c) contacting the alcoholic-caustic phase with an acid to provide an acid-treated alcoholic phase.

[0008] The invention relates in a second embodiment to:

A method for removing impurities from a composition comprising trans-(-)-Δ^9-tetrahydrocannabinol or trans-(+)-Δ^9-tetrahydrocannabinol prior to crystallization of (±)-Δ^9-tetrahydrocannabinol, the method comprising:

(a) contacting the trans-(-)-Δ^9-tetrahydrocannabinol or trans-(+)-Δ^9-tetrahydrocannabinol with a first water immiscible organic solvent, a water-miscible alcohol, water, and an alkali metal hydroxide, thus forming a biphasic composition comprising (i) a first organic phase, and (ii) an alcoholic-caustic phase containing the trans-(-)-Δ^9-tetrahydrocannabinol or the trans-(+)-Δ^9-tetrahydrocannabinol;
(b) separating the alcoholic-caustic phase containing the trans-(-)-Δ^9-tetrahydrocannabinol or the trans-(+)-Δ^9-tetrahydrocannabinol from the first organic phase;
(c) contacting the alcoholic-caustic phase with an acid to provide an acid-treated alcoholic phase.

[0009] The acid in both the first and the second embodiment is preferably citric acid or acetic acid. The acid is preferably added in an amount sufficient to achieve a pH of from 5 to 9.

[0010] The method of the first or second embodiment may additionally comprise the step of:

(d) contacting the acid-treated alcoholic phase with a second water immiscible organic solvent to form (i) a second organic phase comprising the Δ^9-tetrahydrocannabinol and (ii) an acid-treated alcoholic phase.

[0011] After said additional step (d) the method may comprise the additional step of:

(e) separating the second organic phase from the acid-treated alcoholic phase.

[0012] The invention relates in a third embodiment to:

A method for making crystalline (±)-Δ^9-tetrahydrocannabinol comprising:

allowing trans-(-)-Δ^9-tetrahydrocannabinol and trans-(+)-Δ^9-tetrahydrocannabinol to crystallize from a first composition comprising trans-(-)-Δ^9-tetrahydrocannabinol, trans-(+)-Δ^9-tetrahydrocannabinol, and a nonpolar organic solvent to provide crystalline (±)-Δ^9-tetrahydrocannabinol, wherein the first composition was obtained by

(a) performing the method of the first or second embodiment,
(b) separating the Δ^9-tetrahydrocannabinol from the resulting phase containing the Δ^9-tetrahydrocannabinol; and
(c) contacting the Δ^9-tetrahydrocannabinol from step (b) with a non-polar organic solvent and optionally its enantiomer to form the first composition.

[0013] The invention relates in a fourth embodiment to:

A method for preparing

- a trans-(-)-Δ^9-tetrahydrocannabinol composition, comprising: allowing a composition comprising (±)-Δ^9-tetrahy-

drocannabinol and an eluting solvent to separate on a chiral stationary phase to provide a trans-(-)-Δ^9-tetrahydrocannabinol composition, or

- a trans-(+)-Δ^9-tetrahydrocannabinol composition, comprising: allowing a composition comprising (±)-Δ^9-tetrahydrocannabinol and an eluting solvent to separate on a chiral stationary phase to provide a trans-(+)-Δ^9-tetrahydrocannabinol composition,

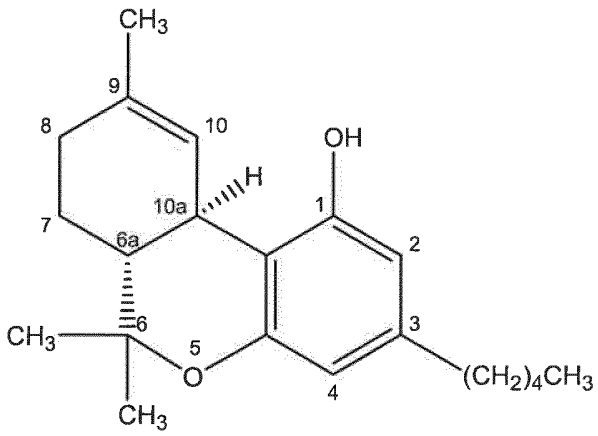
wherein the (±)-Δ^9-tetrahydrocannabinol is obtained from crystalline (±)-Δ^9-tetrahydrocannabinol, and wherein the crystalline (±)-Δ^9-tetrahydrocannabinol is obtained by the method of the third embodiment.

[0014] The present invention can be understood more fully by reference to the following detailed description and illustrative examples.

4. Detailed Description of the Invention

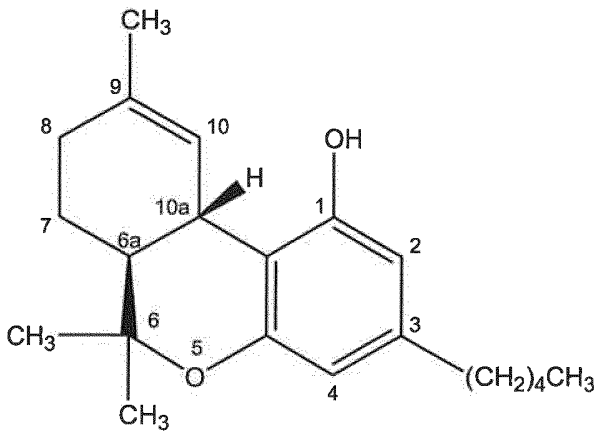
4.1. Definitions

[0015] As used herein, the generic term "Δ^9-THC" refers to trans-(-)-Δ^9-THC, trans-(+)-Δ^9-THC, (±)-Δ^9-THC, or any mixture thereof.
trans-(-)-Δ^9-THC has the structure of formula (**1a**):



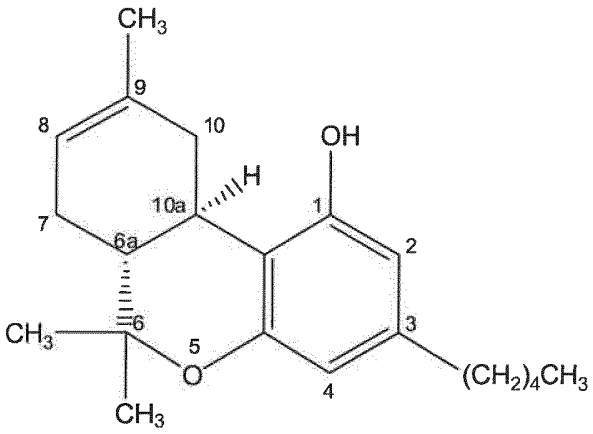
(1a)

Trans-(+)-Δ^9-THC has the structure of (**1b**):



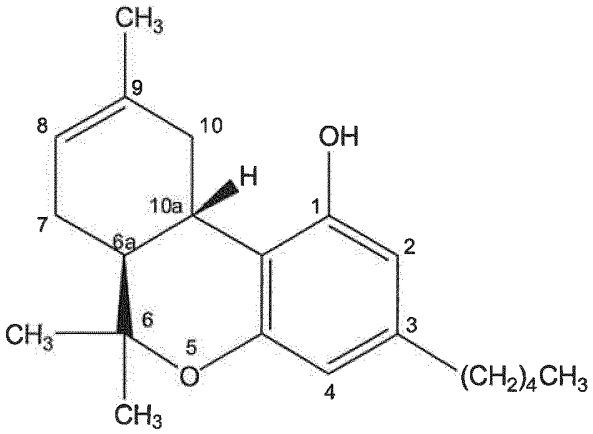
(1b)

[0016] As used herein, the generic term "Δ^8-THC" refers to (-)-Δ^8-THC, (+)-Δ^8-THC, (±)-Δ^8-THC, or any mixture thereof.
(-)-Δ^8-THC has the structure of formula (**2a**):



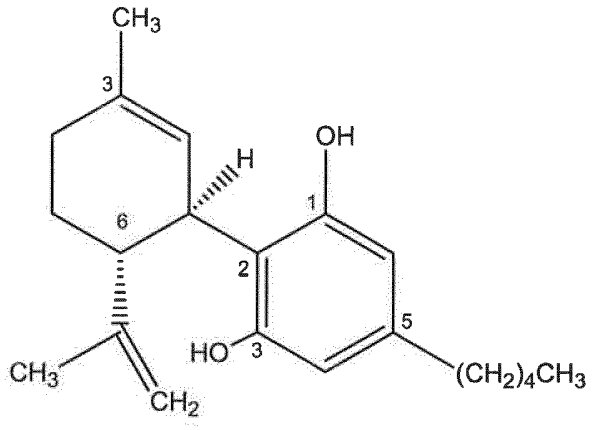
(2a)

(+)-Δ^8-THC has the structure of (**2b**):



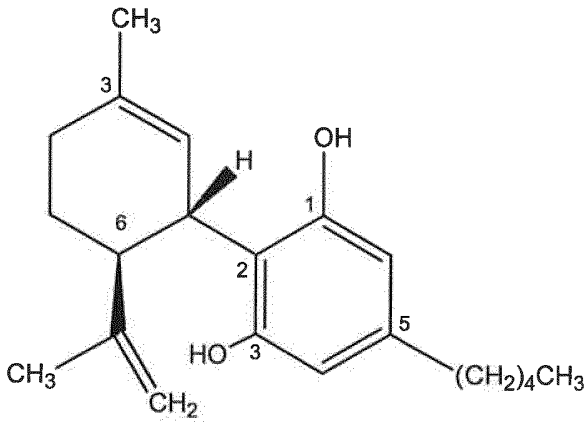
(2b)

[0017] As used herein, the generic term "CBD" refers to (-)-CBD, (+)-CBD, (±)-CBD, or any mixture thereof. (-)-CBD has the structure of formula (**3a**):



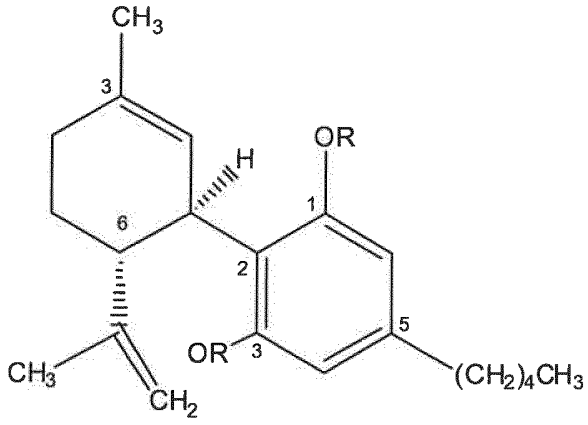
(3a)

(+)-CBD has the structure of formula (**3b**):



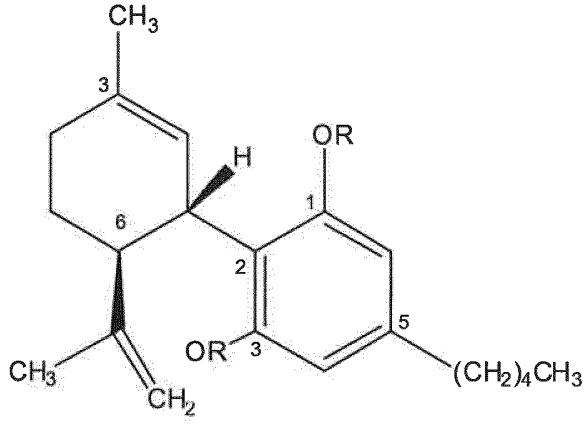
(3b)

[0018] As used herein, the generic term "CBD-bis-1,3-(3,5-dinitrobenzoate)" refers to (-)-CBD-bis(3,5-dinitrobenzoate), (+)-CBD-bis(3,5-dinitrobenzoate), (±)-CBD-bis(3,5-dinitrobenzoate), or any mixture thereof.
(-)-CBD-bis(3,5-dinitrobenzoate) has the structure of formula (**4a**):



(4a)

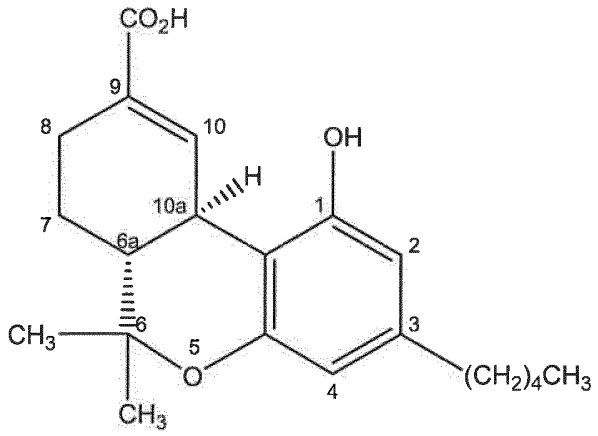
where R is -C(O)(3,5-$C_6H_3(NO_2)_2$).
(+)-CBD-bis(3,5-dinitrobenzoate) has the structure of formula (**4b**):



(4b)

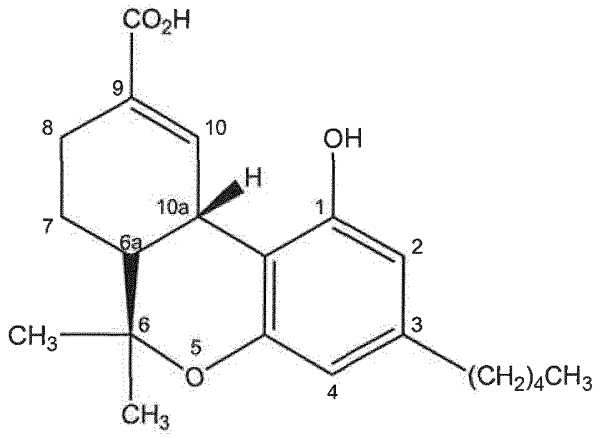
where R is -C(O)(3,5-$C_6H_3(NO_2)_2$).

[0019] As used herein, the generic term "trans-Δ^9-THC carboxylic acid" refers to trans-(-)-Δ^9-THC carboxylic acid, trans-(+)-Δ^9-THC carboxylic acid, trans-(±)-Δ^9-THC carboxylic acid, or any mixture thereof
trans-(-)-Δ^9-THC carboxylic acid has the structure of formula (**5a**):



5a

trans-(+)-Δ^9-THC carboxylic acid has the structure of formula (**5b**):



5b

[0020] The term "-halo" means -F, -Cl, -Br or -I.

[0021] The term "-(C_1-C_4)alkyl" means a saturated straight-chain or branched hydrocarbon having from 1 to 4 carbon atoms. Representative saturated straight chain (C_1-C_4)alkyls are -methyl, -ethyl, -n-propyl, and -n-butyl. Representative saturated branched -(C_1-C_4)alkyls are -isopropyl, -sec-butyl, - isobutyl, and -tert butyl.

[0022] The phrase "anhydrous organic solvent," unless otherwise defined herein, means an organic solvent having an amount of water that is less than 0.01 % by weight of the total amount of water and organic solvent.

[0023] The term "cannabinoids" refers to Δ^9-THC including trans-Δ^9-THC and cis-Δ^9-THC; structural isomers of Δ^9-THC having a molecular formula $C_{21}H_{30}O_2$, including Δ^8-THC, (-)-Δ^8-iso-THC and (+)-Δ^8-iso-THC; cannabinol and structural isomers of cannabinol having a molecular formula of $C_{21}H_{28}O_2$; Δ^9-THC-carboxylic acid; Δ^9-THC precursors including CBD, abn-CBD, (+)-abn-CBD, olivetol, (+)-p-mentha-2,8-dien-1-ol and (-)-p-mentha-2,8-dien-1-ol; salts thereof; and derivatives thereof including acids, ethers, esters, amines.

[0024] Unless otherwise specified herein, the phrase "cannabinoid impurities" means cannabinoids other than trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC.

[0025] Unless otherwise specified herein, the generic term "Δ^9-THC-carboxylic acid" means (-)-Δ^9-THC-carboxylic acid, (+)-Δ^9-THC-carboxylic acid, or (±)-THC-carboxylic acid.

[0026] As used herein, the phrase "crystalline (±)-Δ^9-THC" means a solid form of Δ^9-THC comprising equimolar amounts of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC and having an amount of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC that

is at least 95% by weight based on the total weight of cannabinoids. As used herein, the term "patient" means an animal such as cow, horse, sheep, pig, chicken, turkey, quail, cat, dog, mouse, rat, rabbit, guinea pig, and is more preferably a mammal, and most preferably a human.

4.2. Methods for Purifying: trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC

[0027] The present invention relates to methods for making compositions comprising at least 98% by weight of trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC based on the total amount of cannabinoids.

[0028] In one embodiment, the invention relates to a method comprising allowing a composition comprising (±)-Δ^9-THC and an eluting solvent to separate on a chiral stationary phase to provide a trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC composition, wherein the (±)-Δ^9-THC is obtained from crystalline (±)-Δ^9-THC. Without being limited by theory, Applicants believe that cannabinoid impurities typically present in Δ^9-THC compositions are substantially, if not completely, removed when trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC are allowed to form crystalline (±)-Δ^9-THC. Subsequent resolution of (±)-Δ^9-THC obtained from crystalline (±)-Δ^9-THC with an eluting solvent on a chiral stationary phase provides a composition comprising at least 98% by weight of trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC based on the total amount of cannabinoids.

[0029] In one embodiment, the invention relates to a method for preparing a composition comprising trans-(-)-Δ^9-THC, comprising:

> allowing a composition comprising (±)-Δ^9-THC and an eluting solvent to separate on a chiral stationary phase to provide a trans-(-)-Δ^9-THC composition, wherein the (±)-Δ^9-THC is obtained from crystalline (±)-Δ^9-THC made by the method of the third embodiment as listed above.

[0030] In another embodiment, the invention relates to a method for preparing a composition comprising trans-(+)-Δ^9-THC, comprising:

> allowing a composition comprising (±)-Δ^9-THC and an eluting solvent to separate on a chiral stationary phase to provide a trans-(+)-Δ^9-THC composition, wherein the (±)-Δ^9-THC is obtained from crystalline (±)-Δ^9-THC made by the method of the third embodiment as listed above.

[0031] Crystalline (±)-Δ^9-THC used in these embodiments of the present invention is obtained by a method including crystallization from a first composition comprising trans-(-)-Δ^9-THC, trans-(+)-Δ^9-THC, and a non-polar organic solvent to provide crystalline (±)-Δ^9-THC as described below in Section 4.3.

[0032] In another embodiment, the invention relates to a method for preparing a trans-(-)-Δ^9-THC composition, comprising:

> allowing a composition comprising (±)-Δ^9-THC and an eluting solvent to separate on a chiral stationary phase to provide a trans-(-)-Δ^9-THC composition comprising at least 98% by weight of trans-(-)-Δ^9-THC based on the total amount of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC;
> wherein the (±)-Δ^9-THC was obtained by allowing trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC to crystallize from a first composition comprising trans-(-)-Δ^9-THC, trans-(+)-Δ^9-THC, and a non-polar organic solvent to provide crystalline (±)-Δ^9-THC and a liquid phase according to the method of the third embodiment as listed above.

[0033] In another embodiment, the invention relates to a method for preparing a trans-(+)-Δ^9-THC composition, comprising:

> allowing a composition comprising (±)-Δ^9-THC and an eluting solvent to separate on a chiral stationary phase to provide a trans-(+)-Δ^9-THC composition comprising at least 98% by weight of trans-(+)-Δ^9-THC based on the total amount of trans-(+)-Δ^9-THC and trans-(-)-Δ^9-THC;
> wherein the (±)-Δ^9-THC was obtained by allowing trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC to crystallize from a first composition comprising trans-(-)-Δ^9-THC, trans-(+)-Δ^9-THC, and a non-polar organic solvent to provide crystalline (±)-Δ^9-THC and a liquid phase according to the method of the third embodiment as listed above.

[0034] Compositions comprising trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC useful for obtaining crystalline (±)-Δ^9-THC are obtained by methods described in Section 4.3.

4.3. The Crystallizing Step

[0035] Crystalline (±)-Δ^9-THC is, in one embodiment, obtained by allowing trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC to

crystallize from a composition comprising trans-(-)-Δ^9-THC, trans-(+)-Δ^9-THC and a non-polar organic solvent (the "Crystallizing Step") to provide crystalline (±)-Δ^9-THC and a liquid phase. Compositions comprising trans-(-)-Δ^9-THC, trans-(+)-Δ^9-THC and a non-polar organic solvent useful for the Crystallizing Step are obtained by performing the method of the first or second embodiment.

[0036] In one embodiment, crystalline (±)-Δ^9-THC is obtained by contacting a suitable amount of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC with a non-polar organic solvent. The order and rate of addition of the trans-(-)-Δ^9-THC, the trans-(+)-Δ^9-THC and the non-polar organic solvent is not critical and can be carried out sequentially or substantially simultaneously. As an example, trans-(-)-Δ^9-THC, optionally in the presence of a non-polar organic solvent, and trans-(+)-Δ^9-THC, optionally in the presence of a non-polar organic solvent, can be added to a non-polar organic solvent. Likewise, trans-(+)-Δ^9-THC in the presence of a non-polar organic solvent and trans-(-)-Δ^9-THC in the presence of a non-polar organic solvent can be admixed.

[0037] Trans-(-)-Δ^9-THC can be obtained from natural products or by synthetic methods. Trans-(-)-Δ^9-THC may be obtained from a natural product such as, *e.g.,* hashish or marijuana *(see* Y. Gaoni et al., J. Am. Chem. Soc. 93:217 (1971); and U.S. Patent No. 6,365,416 B1 to Elsohly et al.).

[0038] Trans-(-)-Δ^9-THC can also be obtained by known synthetic methods including reaction of *cis/trans* mixture of (+)-*p*-mentha-2,8-dien-1-ol with olivetol in the presence of an acid catalyst such a para-toluenesulfonic acid and a dehydrating agent (*see* U.S. Patent No. 3,560,528 to Petrizilka and U.S. Patent No. 4,025,516 to Razdan et al.); reaction of (-)-CBD with a Lewis acid such as $BF_3{\cdot}Et_2O$ in an inert solvent under anhydrous conditions *(see* U.S. Patent No. 4,025,516 to Razdan et al.; and International publication no. WO 03/070506); or reaction of (-)-Δ^8-THC with HCl followed by dehydrochlorination *(see* Y. Mechoulam et al., J. Am. Chem. Soc. 89:4553 (1967); and R. Mechoulam et al., J. Am. Chem. Soc. 94:6159 (1972)). Alternatively, trans-(-)-Δ^9-THC can be obtained by methods described in Section 5.

[0039] Trans-(+)-Δ^9-THC, which is not known to occur in nature, can be made by known synthetic methods including reaction of (+)-Δ^8-THC with HCl followed by dehydrochlorination (*see* R. Mechoulam et al., J. Am. Chem. Soc. 94:6159 (1972). Alternatively, trans-(+)-Δ^9-THC can be obtained by methods described in Section 5. In one embodiment, trans-(+)-Δ^9-THC used in the Crystallizing Step is "recycled" from a previous resolution of (±)-Δ^9-THC on a chiral stationary phase as described in Section 4.4.

[0040] In another embodiment, the trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC used in the Crystallizing Step can be obtained as a mixture of enantiomers by a direct synthetic method. When a synthetic method is used, the ratio of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC can vary depending on the optical purity of the reagents and the synthetic process. In one embodiment, trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC are obtained in equimolar amounts by a synthetic route using racemic reagents. Methods for preparing trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC by a direct synthetic route include reaction of citral and olivetol in the presence of a Lewis acid *(see* R. Mechoulam et al., J. Am. Chem. Soc. 94:6159 (1972)) or hydrolysis of (±)-1-*m*-nitrobenzenesulfoanate-6a,10a-trans-Δ^9-THC with NaOH in aqueous methanol (K.E. Fahrenholtz et al., J. Am. Chem. Soc. 89:5934-5941 (1967)). Alternatively, (±)-Δ^9-THC can be obtained by methods described in Section 5.

[0041] The ratio of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC used in the Crystallizing Step can vary. In one embodiment, the trans-(-)-Δ^9-THC is present in an amount from 0.75 to 1.25 molar equivalents per molar equivalent of trans-(+)-Δ^9-THC. In another embodiment, the trans-(-)-Δ^9-THC is present in an amount from 0.9 to 1.1 molar equivalents per molar equivalent of trans-(+)-Δ^9-THC. In another embodiment, the trans-(-)-Δ^9-THC is present in an amount from 0.95 to 1.05 molar equivalents per molar equivalent of trans-(+)-Δ^9-THC. And in another embodiment, the trans-(-)-Δ^9-THC is present in an amount of 1 molar equivalent per molar equivalent of trans-(+)-Δ^9-THC.

[0042] Examples of non-polar organic solvents that are useful in the Crystallizing Step include aliphatic (C_4-C_{10})hydrocarbons such as butane, pentane, hexane, heptane, octane, nonane, decane, including straight-chained aliphatic hydrocarbons, branched aliphatic hydrocarbons and cyclic aliphatic hydrocarbons, or any mixture thereof.

[0043] In one embodiment, the non-polar organic solvent used in the Crystallizing Step is a straight-chain or branch-chain heptane. In another embodiment, the non-polar organic solvent used in the Crystallizing Step is pentane, hexane, heptane, octane or isooctane. In another embodiment, the non-polar organic solvent used in the Crystallizing Step is n-heptane.

[0044] The amount of the non-polar organic solvent used in the Crystallizing Step can vary and will depend, in part, on the amount and type of cannabinoid impurities and temperature. Typically, the non-polar organic solvent is present in an amount sufficient to provide a mixture having a Δ^9-THC concentration from 1 % to 95%, preferably from 20% to 75%, more preferably from 40% to 60% by weight based on the total amount of Δ^9-THC and the non-polar organic solvent.

[0045] The Crystallizing Step is carried out for a time and at a temperature sufficient to provide (±)-Δ^9-THC crystals. A time sufficient to crystallize (±)-Δ^9-THC is from 1 hour to 200 hours; in another embodiment, from 5 hours to 150 hours; in another embodiment, from 25 hours to 100 hours; and in another embodiment, from 30 hours to 75 hours.

[0046] Typically, a temperature sufficient to provide crystalline (±)-Δ^9-THC is from -78°C to 100°C; in another embodiment, from -50°C to 25°C; in another embodiment, from -30°C to 0°C; and in another embodiment, from -25°C to -15°C.

[0047] In certain embodiments, the Crystallizing Step is carried out at two or more temperatures. In one embodiment,

the composition comprising trans-(-)-Δ^9-THC, trans-(+)-Δ^9-THC and a non-polar organic solvent is prepared at a first temperature, e.g., 20°C or higher. Without being limited by theory, Applicants believe that forming the composition at a temperature of 20°C or higher increases the solubility of the trans-(-)-Δ^9-THC and the trans-(+)-Δ^9-THC in the non-polar organic solvent. The temperature of the admixture can then be decreased to a second temperature, e.g., 0°C or lower. Without being limited by theory, Applicants believe that holding the admixture at a temperature of 0°C or lower decreases the solubility of (±)-Δ^9-THC and promotes crystallization. Optionally, the temperature of the admixture can be further decreased to a third temperature, e.g., -20 to -15°C. As noted above, such a decrease in temperature is believed to enhance the (±)-Δ^9-THC crystallization process.

[0048] In one embodiment, trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC are dissolved in a non-polar organic solvent; the resultant solution is cooled to 0°C; the resultant mixture is further cooled to -15°C; and the resultant crystalline (±)-Δ^9-THC is separated from the liquid phase.

[0049] In another embodiment, the Crystallizing Step is carried out in the presence of a seed crystal. Typically, the seed crystal, when used, is added to the cold (*e.g.*, 0°C or lower) admixture comprising trans-(-)-Δ^9-THC, trans-(+)-Δ^9-THC and the non-polar organic solvent. In one embodiment, the seed crystal is (±)-Δ^9-THC.

[0050] The progress of the Crystallizing Step can be monitored visually or using conventional analytical techniques, including thin-layer chromatography ("TLC"), high-performance liquid chromatography ("HPLC"), gas chromatography ("GC"), gas-liquid chromatography ("GLC"), infrared spectroscopy ("IR"), Raman spectroscopy ("Raman") and nuclear magnetic resonance spectroscopy ("NMR") such as ^{1}H or ^{13}C NMR.

[0051] The Crystallizing Step can be carried out at reduced pressure, atmospheric pressure or elevated pressure. In one embodiment, the Crystallizing Step is carried out at atmospheric pressure.

[0052] In the present invention, impurities are removed from the trans-(-)-Δ^9-THC and/or trans-(+)-Δ^9-THC compositions prior to carrying out the Crystallizing Step by extraction under basic conditions as described below.

[0053] In the present invention, (+)-Δ^9-THC, (-)-Δ^9-THC, or (±)-Δ^9-THC is contacted with base prior to carrying out the Crystallizing Step.

[0054] The invention also relates to a method for purifying trans-(+)-Δ^9-THC, trans-(-)-Δ^9-THC, or trans-(±)-Δ^9-THC (the "Δ^9-THC Purification Method") comprising:

> contacting trans-(+)-Δ^9-THC, trans-(-)-Δ^9-THC, or trans-(±)-Δ^9-THC with a first water-immiscible organic solvent, a water-miscible alcohol, water, and an alkali metal hydroxide (the "Caustic Contacting Step") to form a biphasic mixture comprising (i) a first organic phase and (ii) an alcoholic-caustic phase comprising trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC.

[0055] Without being limited by theory, it is believed that the Caustic Contacting Step extracts impurities from the Δ^9-THC-containing alcoholic-caustic phase into the first organic phase, which impurities may impede or prevent (±)-Δ^9-THC from crystallizing from the composition comprising the trans-(-)-Δ^9-THC, the trans-(+)-Δ^9-THC, and the non-polar organic solvent.

[0056] The amount of alkali metal hydroxide used in the Caustic Contacting Step typically ranges from 1 to 1000 molar equivalents per molar equivalent of Δ^9-THC; in another embodiment, the amount of alkali metal hydroxide ranges from 10 to 100 molar equivalents per molar equivalent of trans-Δ^9-THC; and in another embodiment, amount of alkali metal hydroxide ranges from 25 to 55 molar equivalents per molar equivalent of trans-Δ^9-THC.

[0057] Examples of water-miscible alcohols useful in the Caustic Contacting Step include methanol, ethanol, isopropanol, or any combination thereof. In one embodiment, the water-miscible alcohol is methanol.

[0058] The amount of water-miscible alcohol used in the Caustic Contacting Step typically is from 1 part to 100 parts by weight based on the weight of the alkali metal hydroxide; in another embodiment, the amount of water-miscible alcohol is from 1 part to 25 parts by weight based on the weight of the alkali metal hydroxide; and in another embodiment, the amount of water-miscible alcohol is from 5 parts to 10 parts by weight based on the weight of the alkali metal hydroxide.

[0059] Examples of first water-immiscible organic solvents useful in the Caustic Contacting Step include the non-polar organic solvents described above for the Crystallizing Step. In one embodiment, the first water-immiscible solvent is heptane.

[0060] The amount of first water-immiscible organic solvent used in the Caustic Contacting Step typically is from 1 part to 1000 parts by weight based on the weight of the Δ^9-THC; in another embodiment, the amount of water-immiscible organic solvent is from 5 parts to 100 parts by weight based on the weight of the Δ^9-THC; and in another embodiment, the amount of water-immiscible organic solvent is from 5 parts to 20 parts by weight based on the weight of the Δ^9-THC.

[0061] The Caustic Contacting Step can be carried out by methods known in the art such as stirring, shaking, countcrcurrcnt cascade, and ultrasound admixing, pumping. The Caustic Contacting Step can also be carried out by methods useful for liquid-liquid extraction (*see*, *e.g.*, Lo et al., Extraction, in 7 Kirk-Othmer Encyc. of Chem. Technol. 349-381 (4th ed. 1993)).

[0062] The Caustic Contacting Step typically is carried out from 0.25 hours to 50 hours; in another embodiment, from

0.25 hours to 10 hours; and in another embodiment, from 0.25 hours to 2 hours.

[0063] The Caustic Contacting Step is typically carried at a temperature of from 0°C to 100 °C; in another embodiment, from 20°C to 50°C; and in another embodiment, from 20°C to 30°C.

[0064] The Caustic Contacting Step can be carried out at reduced pressure, atmospheric pressure (*i.e.*, 1 atmosphere), or elevated pressure. In one embodiment, the Caustic Contacting Step is carried out at atmospheric pressure.

[0065] The progress of the Caustic Contacting Step can be monitored using conventional techniques as described above for the Crystallizing Step.

[0066] The trans-Δ^9-THC Purification Method of the present invention further comprises contacting the alcoholic-caustic phase with an acid to provide an acid-treated alcoholic phase. Without being limited by theory, it is believed that trans-Δ^9-THC is immiscible in the acidified alcoholic phase. Examples of useful acids include citric acid, acetic acid. In one embodiment, the acid is citric acid.

[0067] Typically, the acid, is added in an amount sufficient to achieve a pH of from 5 to 9. In another embodiment, the acid is added in an amount sufficient to achieve a pH from 6 to 8; in another embodiment, the acid is added in an amount sufficient to achieve a pH of from 7 to 8.

[0068] The Δ^9-THC Purification Method of the present invention may further comprise contacting the acid-treated alcoholic phase with a second water-immiscible organic solvent to form (i) a second organic phase comprising trans-(-)Δ^9-THC and (ii) an acid-treated alcoholic phase.

[0069] Examples of second water-immiscible organic solvents useful for contacting the acid-treated alcoholic phase to form a second organic phase comprising trans-Δ^9-THC include the non-polar organic solvents described above for the Crystallizing Step. In one embodiment, the second water-immiscible organic solvent is heptane. The amount of first water-immiscible organic solvent used is typically is from 1 part to 1000 parts by weight based on the weight of the trans-Δ^9-THC; in another embodiment, the amount of water-immiscible organic solvent is from 1 part to 50 parts by weight based on the weight of the trans-Δ^9-THC; and in another embodiment, the amount of water-immiscible organic solvent is from 1 part to 10 parts by weight based on the weight of the trans-Δ^9-THC. Methods useful for contacting the acid-treated alcoholic phase with a second water-immiscible organic solvent include those described above for the Caustic Contacting Step.

[0070] The Δ^9-THC Purification Method of the present invention may further comprise separating the second organic phase from the acid-treated alcoholic phase. Methods useful for separating the second organic phase from the acid-treated alcoholic phase include those described above for separating the first organic phase from the alcoholic-caustic phase. After separation from the acid-treated alcoholic phase, the second organic phase is typically dried by, *e.g.*, azeotropic distillation and/or contacting the second organic phase with a drying agent (*e.g.*, Na_2SO_4 or $MgSO_4$).

[0071] The Δ^9-THC Purification Method of the present invention may further comprise concentrating the second organic phase to form a concentrated second organic phase comprising trans-Δ^9-THC. A method useful for concentrating the second organic phase is distillation. When the second organic phase is concentrated by distillation, the distillation can be carried out at elevated pressure, atmospheric pressure, or at reduced pressure. In one embodiment, the distillation is carried out at atmospheric pressure. In another embodiment, the distillation is carried out at reduced pressure.

[0072] The Δ^9-THC Purification Method of the present invention may further comprise contacting the concentrated second organic phase with a non-polar organic solvent to form a first organic composition comprising trans-Δ^9-THC. The amount and type of non-polar organic solvent are those described above in the Crystallizing Step for the non-polar organic solvent.

[0073] In one embodiment, the trans-Δ^9-THC used in the Δ^9-THC Purification Method comprises trans-(-)-Δ^9-THC. In another embodiment, the trans-Δ^9-THC used in the Δ^9-THC Purification Method comprises trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC. In another embodiment, the trans-Δ^9-THC used in the Δ^9-THC Purification Method comprises trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC, wherein the trans-(-)-Δ^9-THC is present in an amount from 0.75 to 1.25 molar equivalents per molar equivalent of trans-(+)-Δ^9-THC.

[0074] In another embodiment, the Δ^9-THC Purification Method of the present invention further comprises allowing the trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC to crystallize from the first organic composition to provide crystalline (±)-Δ^9-THC as described above for the Crystallizing Step, wherein (a) the first organic composition comprises (-)-Δ^9-THC and trans-(+)-Δ^9-THC, and (b) the trans-(-)-Δ^9-THC is present in the first organic composition in an amount from 0.75 to 1.25 molar equivalents per molar equivalent of trans-(+)-Δ^9-THC.

[0075] The invention also relates to a method for making crystalline (±)-Δ^9-THC comprising:

allowing trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC to crystallize from a first composition comprising trans-(-)-Δ^9-THC, trans-(+)-Δ^9-THC, and a non-polar organic solvent to provide crystalline (±)-Δ^9-THC,
wherein the trans-(-)-Δ^9-THC and the trans-(+)-Δ^9-THC were obtained by:

(a) forming a biphasic composition comprising (i) a first organic phase comprising a first water-immiscible organic solvent, and (ii) an alcoholic-caustic phase containing trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC;

(b) separating the trans-(-)-Δ^9-THC and the trans-(+)-Δ^9-THC from the alcoholic-caustic phase according to the method of the first or second embodiment described above; and
(c) forming the first composition comprising (i) the trans-(-)-Δ^9-THC and the trans-(+)-Δ^9-THC of step (b), and (ii) the non-polar organic solvent.

[0076] Methods for forming the biphasic composition as well as the amounts and type of first water-immiscible organic solvent, water-miscible alcohol, water, and alkali metal hydroxide include those described above for the Caustic Contacting Step. Similarly, methods for separating the trans-(-)-Δ^9-THC and the trans-(+)-Δ^9-THC from the alcoholic-caustic phase, and methods for forming the first composition comprising (i) the trans-(-)-Δ^9-THC and the trans-(+)-Δ^9-THC of step (b), and (ii) the non-polar organic solvent include those described above for the (±)-Δ^9-THC Purification Method.

[0077] Once obtained, crystalline (±)-Δ^9-THC formed in the Crystallizing Step can be separated from the liquid phase by methods known in the art. Methods for separating the crystalline (±)-Δ^9-THC from the liquid phase include, *e.g.*, filtration, centrifugation and decantation. In one embodiment, crystalline (±)-Δ^9-THC is separated from the liquid phase by filtration.

[0078] Crystalline (±)-Δ^9-THC formed in the Crystallizing Step can, optionally, be washed with an organic wash solvent, and separated from the liquid phase as described above. When crystalline (±)-Δ^9-THC is washed, the temperature of the organic wash solvent can vary. Typically, the washing, when done, is carried out with an organic wash solvent at a temperature from -78°C to 50°C; in another embodiment, from -30°C to 30°C; and in another embodiment, from -20°C to 25°C.

[0079] Examples of useful organic wash solvents include the non-polar organic solvents described above. In one embodiment, the organic wash solvent, when used, is n-heptane.

[0080] The separated (±)-Δ^9-THC can, optionally, be dried. The drying can be carried out at atmospheric pressure, optionally with the aid of a sweep gas such as dry air, nitrogen, helium or argon. Alternatively, the (±)-Δ^9-THC can be dried at reduced pressure.

[0081] When the separated (±)-Δ^9-THC is dried, the drying temperature can vary. Typically, the drying, when done, can be carried out at a temperature from -25° to 65°C; in another embodiment, from 0° to 60°C; and in another embodiment, from 25° to 50°C.

[0082] Typically, the (±)-Δ^9-THC obtained in the Crystallizing Step comprises at least 95% by weight of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC based on the total amount of cannabinoids. In another embodiment, the (±)-Δ^9-THC obtained in the Crystallizing Step comprises at least 98% by weight of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC based on the total amount of cannabinoids. In another embodiment, the (±)-Δ^9-THC obtained in the Crystallizing Step comprises at least 99% by weight of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC based on the total amount of cannabinoids.

[0083] The separated (±)-Δ^9-THC can then be resolved on a chiral stationary phase as described below in Section 4.4.

4.4. The Resolving Step

[0084] In the present invention, (±)-Δ^9-THC obtained from crystalline (±)-Δ^9-THC and an eluting solvent are contacted with a chiral stationary phase to resolve the trans-(-)- and (+)-enantiomers (the "Resolving Step"). This provides a composition comprising at least 98% by weight of trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC based on the total amount of cannabinoids. Without being limited by theory, Applicants believe that resolving (±)-Δ^9-THC obtained from crystalline (±)-Δ^9-THC provides a trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC composition having low levels, if any, of the cannabinoid impurities found in trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC obtained by known methods.

[0085] The composition comprising (±)-Δ^9-THC used in the Resolving Step can contain an amount of trans-(-)-Δ^9-THC that is less than, equal to or greater than the amount of trans-(+)-Δ^9-THC. For example, the composition comprising (±)-Δ^9-THC may be obtained by admixing crystalline (±)-Δ^9-THC with a trans-(-)-Δ^9-THC composition and/or a trans-(+)-Δ^9-THC prior to carrying out the Resolving Step. Typically, the composition comprising (±)-Δ^9-THC contains an equimolar amount of the trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC.

[0086] Any known or later-developed chiral stationary phase that resolves trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC can be used. For example, a method for resolving trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC enantiomers on a chiral stationary phase is described in S. L. Levin et al., J. Chromatogr. A 654:53-64 (1993). Typically, the chiral stationary phase contains a chiral group or derivative immobilized on a support such as, *e.g.*, a polymer or inorganic oxide. An example of a useful polymer support is polystyrene in bead form. Examples of useful inorganic oxide supports include silica, magnesium silicate, magnesia, alumina and molecular sieves. In one embodiment, the inorganic oxide support is silica.

[0087] The chiral derivative comprises at least one chiral center. Examples of useful chiral derivatives include tris(arylcarbamate) derivatives of saccharides such as, e.g., amylose, cellulose, chitosin, xylan, curdlan, dextran, and inulan. In one embodiment, the saccharide is amylose.

[0088] In one embodiment, the tris(arylcarbamate) is tris(3,5-dimethylphenylcarbamate), tris(4-chlorophenylcarbamate), tris(4-methylcarbamate), tris(4-methylbenzoate) or tris[(S)-phenylethylcarbamate]. In another embodiment, the

tris(arylcarbamate) is tris(3,5-dimethylphenylcarbamate). In another embodiment, the chiral stationary phase is amylose tris(3,5-dimethylcarbonate) immobilized on silica, available as Chiralpak® AD™ from Daicel Chemical Industries, Tokyo, Japan.

[0089] Other examples of useful chiral stationary phases include cellulose triacetate; cellulose tribenzoate; poly[(S)-N-acrylolyphenylalanine ethyl ester]; 3,5-dinitrobenzoylphenylglycine; crosslinked di-(3,5-dimethylbenzoyl)-L diallyltartramide; crosslinked di-(4-*tert*-butylbenzoyl)-L diallyltartramide; and tetrahydro-aminophenanthrene 3,5-dinitrobenzamide *(see* E.R. Francotte, J. Chromatogr. A 906:379-397 (2001)).

[0090] Typically, a concentrated solution of (±)-Δ^9-THC and an eluting solvent is added to the top (or front) of a column containing a chiral stationary phase. The (±)-Δ^9-THC is then eluted with the eluting solvent (*i.e.*, the mobile phase) to provide eluents containing trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC.

[0091] The Resolving Step can be carried out using batch chromatography, continuous chromatography or simulated moving bed chromatography (see, *e.g.*, E.R. Francotte, J. Chromatogr. A 906:379-397 (2001)). In one embodiment, the Resolving Step is carried out using continuous chromatography.

[0092] The Resolving Step can be carried out at 1 atmosphere of pressure or, optionally, at reduced pressure or elevated pressure. In one embodiment, the Resolving Step is carried out at 1 atmosphere of pressure. In another embodiment, the Resolving Step is carried out at elevated pressure. In one embodiment, the Resolving Step is carried out at using flash chromatography at moderately elevated pressure, *e.g.*, from 1.1 to 10 atmospheres; from 1.1 to 5 atmospheres; or from 1.1 to 1.3 atmospheres. In another embodiment, the Resolving Step is carried out at using flash chromatography at highly elevated pressure, *e.g.*, from 10 to 175 atmospheres; from 100 to 175 atmospheres; from 125 to 175 atmospheres; or at 150 atmospheres.

[0093] Examples of eluting solvents useful in the Resolving Step include straight-chain or branch-chain (C_1-C_4)alkyls substituted with one or more -OH, -OR_1, -OC(O)R_1, -C(O)OR_1, -halo, or -CN; straight-chain or branch-chain (C_4-C_{10})aliphatic hydrocarbons; (C_5-C_7)cycloaliphatic hydrocarbon optionally substituted with one or more -R_1; (C_4-C_7)cyclic ethers optionally substituted with one or more -R_1; aromatic hydrocarbons optionally substituted with one or more -R_1, -halo, -CH_2(halo), -CH(halo)$_2$, -C(halo)$_3$, -O(C_1-C_6)alkyl; or any mixture thereof, where R_1 is (C_1-C_4)alkyl.

[0094] Examples of straight-chain or branch-chain (C_1-C_4)alkyls substituted with one or more -OH, -OR_1, -OC(O)R_1, -C(O)OR_1, -halo, or -CN include methanol, ethanol, n-propanol, isopropanol, n-butanol, isobutanol, tert-butanol, chloromethane, methylene chloride, chloroform, carbon tetrachloride chloride, diethyl ether, diisopropyl ether, tert-butyl methyl ether, acetonitrile, methyl formate, ethyl formate, methyl acetate, ethyl acetate, isopropyl acetate, butyl acetate, or any mixture thereof.

[0095] Examples of straight-chain or branch-chain (C_4-C_{10})aliphatic hydrocarbons include butane, pentane, hexane, heptane, isooctane, nonane, decane, or any mixture thereof.

[0096] Examples of (C_5-C_7)cycloaliphatic hydrocarbons optionally substituted with one or more -R_1 include cyclopentane, cyclohexane, methylcyclohexane, cycloheptane or any mixture thereof.

[0097] Examples of (C_4-C_7)cyclic ethers optionally substituted with one or more -R_1 include tetrahydrofuran, methyltetrahydrofuran, 1,4-dioxane, 1,3-dioxolane, or any mixture thereof.

[0098] Examples of aromatic hydrocarbons optionally substituted with one or more -R_1, -halo, -CH_2(halo), -CH(halo)$_2$, -C(halo)$_3$, -O(C_1-C_6)alkyl include toluene, xylene, chlorobenzene, benzotrifluoride, or any mixture thereof.

[0099] In one embodiment, the eluting solvent comprises an aliphatic hydrocarbon and an alcohol. In another embodiment, the eluting solvent comprises n-hcptanc and iso-propanol. In another embodiment, the organic solvent comprises a 95:5 (v/v) mixture of n-heptane:2-propanol.

[0100] The progress of the Resolving Step can be monitored using analytical methods described above in Section 4.3.

[0101] The eluents containing trans-(-)-Δ^9-THC and being substantially free of other cannabinoids can be combined. In one embodiment, the eluents comprise at least 98% by weight of trans-(-)-Δ^9-THC; in another embodiment, at least 99% by weight of trans-(-)-Δ^9-THC; in another embodiment, at least 99.5% by weight of trans-(-)-Δ^9-THC; and in another embodiment, at least 99.9% by weight of trans-(-)-Δ^9-THC based on the total amount of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC.

[0102] Similarly, eluents containing trans-(+)-Δ^9-THC and being substantially free of other cannabinoids can be combined. In one embodiment, the eluents comprise at least 98% by weight of trans-(+)-Δ^9-THC; in another embodiment, at least 99% by weight of trans-(+)-Δ^9-THC; in another embodiment, at least 99.5% by weight of trans-(+)-Δ^9-THC; and in another embodiment, at least 99.9% by weight of trans-(+)-Δ^9-THC based on the total amount of trans-(+)-Δ^9-THC and trans-(-)-Δ^9-THC.

[0103] The eluents comprising a first solvent and trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC can, optionally, be separated from the volatiles to provide each enantiomer as an oil. Methods for separating the trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC from volatile components include, *e.g.*, distillation at atmospheric pressure or reduced pressure. For example, the trans-(-)-Δ^9-THC or trans-(+)-Δ^9THC can, if desired, be distilled by fractional distillation to provide a trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC distillate (*see* U.S. Patent No. 4,381,399 to Olsen et al.).

4.5. Compositions Comprising: trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC

[0104] The present invention relates to methods for preparing compositions comprising trans-(-)-Δ^9-THC or trans-(+)-Δ^9-THC.

[0105] In one embodiment, a method according to the invention provides a composition comprising at least 99.0% by weight of trans-(-)-Δ^9-THC; in another embodiment, at least 99.5% by weight of trans-(-)-Δ^9-THC; and in another embodiment, at least 99.9% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids.

[0106] In one embodiment, a method according to the invention provides a composition comprising at least 99.0% up to 99.95% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids. In another embodiment, a methods according to the invention provides a composition comprising at least 99.0% up to 99.98% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids.

[0107] The compositions comprising at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids may be formulated as a pharmaceutical composition as described in Section 4.6.

[0108] In one embodiment, a method according to the invention provides a composition comprising at least 99.0% by weight of trans-(+)-Δ^9-THC; in another embodiment, at least 99.0.0% by weight of trans-(+)-Δ^9-THC; in another embodiment, at least 99.5.0% by weight of trans-(+)-Δ^9-THC; and in another embodiment, at least 99.9% by weight of trans-(+)-Δ^9-THC based on the total amount of cannabinoids.

[0109] The trans-(-)-Δ^9-THC compositions prepared according to the present invention typically contain no Δ^9-THC carboxylic acid, which can be found in trans-(-)-Δ^9-THC compositions derived from natural sources (see R.F. Turk et al., J. Pharm. Pharmac. 23:190-195 (1971)). In one embodiment, the trans-(-)-Δ^9-THC compositions prepared according to the present invention contain less than 0.05% Δ^9-THC carboxylic acid; in another embodiment, less than 0.01% Δ^9-THC carboxylic acid; in another embodiment, less than 0.005% Δ^9-THC carboxylic acid; and in another embodiment, less than 0.001% Δ^9-THC carboxylic acid based on the based on the total amount of cannabinoids. In another embodiment, the trans-(-)-Δ^9-THC compositions contain no Δ^9-THC carboxylic acid.

[0110] In another embodiment, a method according to the invention provides a composition comprising at least 99.0% by weight of trans-(-)-Δ^9-THC and less than of 0.05% of Δ^9-THC carboxylic acid; in another embodiment, at least 99.5% by weight of trans-(-)-Δ^9-THC and less than 0.05% of Δ^9-THC carboxylic acid; and in another embodiment, at least 99.9% by weight of trans-(-)-Δ^9-THC and less than 0.05% of Δ^9-THC carboxylic acid based on the total amount of cannabinoids.

[0111] In another embodiment, a method according to the invention provides a composition comprising at least 99.0% by weight of trans-(+)-Δ^9-THC and less than of 0.05% of Δ^9-THC carboxylic acid; in another embodiment, at least 99.5% by weight of trans-(+)-Δ^9-THC and less than 0.05% of Δ^9-THC carboxylic acid; and in another embodiment, at least 99.9% by weight of trans-(+)-Δ^9-THC and less than 0.05% of Δ^9-THC carboxylic acid based on the total amount of cannabinoids.

4.6. Therapeutic/Prophylactic Administration of Compositions Comprising trans-(-)-Δ^9-THC

[0112] Compositions comprising at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids are useful for treating the same diseases, ailments, or disorders ("Conditions") for which trans-(-)-Δ^9-THC is known to be useful, or for any Condition for which trans-(-)-Δ^9-THC is later found to be useful for treating or preventing. For example, trans-(-)-Δ^9-THC compositions comprising at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids can be used for treating or preventing emesis, loss of weight loss of appetite, multiple sclerosis, Tourette's syndrome, Parkinson's disease, or palsies such as cerebral palsy. Accordingly, methods for treating or preventing a Condition comprise administering to a patient in need thereof an effective amount of a trans-(-)-Δ^9 composition, wherein the trans-(-)-Δ^9-THC composition comprises at least 99.0% by weight of trans-(-)-Δ^9-THC; or at least 99.5% by weight of trans-(-)-Δ^9-THC; or at least 99.9% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids.

[0113] Other methods for treating or preventing a Condition comprise administering to a patient in need thereof an effective amount of a trans-(-)-Δ^9 composition, wherein the trans-(-)-Δ^9-THC composition comprises at least 99.0% by weight of trans-(-)-Δ^9-THC and less than 0.05% of Δ^9-THC carboxylic acid; or at least 99.5% by weight of trans-(-)-Δ^9-THC and less than 0.05% of Δ^9-THC carboxylic acid; or at least 99.9% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids and less than 0.05% of Δ^9-THC carboxylic acid.

[0114] In particular, the Condition may be pain.

[0115] In particular, the Condition may be emesis, e.g., as the result of cancer chemotherapy.

[0116] In particular, the Condition may be loss of appetite.

[0117] In particular, the Condition may be weight loss, *e.g.*, as the result of symptomative HIV infection including acquired immunodeficiency syndrome (AIDS) or AIDS related complex (ARC).

[0118] When administered to a patient, the trans-(-)-Δ^9-THC compositions containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids comprise a suitable amount of a pharmaceutically acceptable carrier so as to provide the form for proper administration to the patient.

[0119] In particular, the term "pharmaceutically acceptable" means approved by a regulatory agency of the Federal or a state government or listed in the U.S. Pharmacopeia or other generally recognized pharmacopeia for use in animals, and more particularly in humans. The term "carrier" refers to a diluent, adjuvant, excipient, or vehicle with which the trans-(-)-Δ^9-THC, containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids, is administered. Such pharmaceutical carriers can be liquids, such as water and oils, including those of petroleum, animal, vegetable or synthetic origin, such as peanut oil, soybean oil, mineral oil, sesame oil. The pharmaceutical carriers can be saline, gum acacia, gelatin, starch paste, talc, keratin, colloidal silica, urea. In addition, auxiliary, stabilizing, thickening, lubricating and coloring agents may be used. The present compositions, if desired, can also contain minor amounts of wetting or emulsifying agents, and/or pH buffering agents. When administered to a patient, the pharmaceutically acceptable carriers are preferably sterile.

[0120] The present compositions can take the form of solutions, suspensions, emulsion, tablets, pills, pellets, capsules, capsules containing liquids, powders, sustained-release formulations, suppositories, emulsions, aerosols, sprays, suspensions, or any other form suitable for use.

[0121] In one example, the trans-(-)-Δ^9-THC composition containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids further comprises sesame oil. In another example, the trans-(-)-Δ^9-THC compositions containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids further comprises sesame oil, and the resultant admixture is encapsulated (*see, e.g.*, U.S. Patent No. 6,703,418 B2).

[0122] In another example, the trans-(-)-Δ^9-THC composition containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids is formed as a tablet.

[0123] The trans-(-)-Δ^9-THC compositions containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total weight of cannabinoids can be administered by any convenient route, for example by infusion or bolus injection, by absorption through epithelial or mucocutaneous linings (e.g., oral mucosa, rectal and intestinal mucosa) and may be administered together with another biologically active agent. Administration can be systemic or local. Various delivery systems are known, e.g., encapsulation in liposomes, microparticles, microcapsules, capsules, and can be used to administer the pharmaceutical compositions. Methods of administration include intradermal, intramuscular, intraperitoneal, intravenous, subcutaneous, intranasal, epidural, oral, sublingual, intranasal, intracerebral, intravaginal, transdermal, rectally, by inhalation, or topically to the ears, nose, eyes, or skin. The preferred mode of administration is oral, but other modes of administration can be left to the discretion of the practitioner.

[0124] When used for oral delivery, the trans-(-)-Δ^9-THC composition containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids can be in the form of tablets, lozenges, aqueous or oily suspensions, granules, powders, emulsions, capsules, syrups, or elixirs, for example. Orally administered compositions can contain one or more optional agents, for example, sweetening agents such as fructose, aspartame or saccharin; flavoring agents such as peppermint, oil of wintergreen, or cherry; coloring agents; and preserving agents, to provide a pharmaceutically palatable preparation. Moreover, where in tablet or pill form, the compositions can be coated to delay disintegration and absorption in the gastrointestinal tract thereby providing a sustained action over an extended period of time. Selectively permeable membranes surrounding an osmotically active driving compound are also suitable for orally administered pharmaceutical compositions. In these later platforms, fluid from the environment surrounding the capsule is imbibed by the driving compound, which swells to displace the agent or agent composition through an aperture. These delivery platforms can provide an essentially zero-order delivery profile as opposed to the spiked profiles of immediate release formulations. A time delay material such as glycerol monostearate or glycerol stearate may also be used. Oral compositions can include standard carriers such as mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate. Such carriers are preferably of pharmaceutical grade.

[0125] When used for intravenous delivery, the trans-(-)-Δ^9-THC composition containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids is formulated in accordance with routine procedures for intravenous administration to human beings. Preferably, the pharmaceutical compositions for intravenous administration are solutions in sterile isotonic aqueous buffer, optionally with a solublizing agent. Compositions for intravenous administration may optionally include a local anesthetic such as lignocaine to ease pain at the site of the injection. Generally, the ingredients are supplied either separately or mixed together in unit dosage form, for example, as a dry lyophilized powder or water free concentrate in a hermetically sealed container such as an ampoule or sachette indicating the quantity of active agent. Where the pharmaceutical compositions are to be administered by infusion, they can be dispensed, for example, with an infusion bottle containing sterile pharmaceutical grade water or saline, optionally with a solublizing agent. Where the pharmaceutical compositions are administered by injection, an ampoule of sterile water for injection or saline can be provided so that the ingredients can be mixed prior to administration.

[0126] The amount of the trans-(-)-Δ^9-THC composition containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids that is effective in the treatment or prevention of a Condition can be determined by standard clinical techniques. In addition, *in vitro* or *in vivo* assays can optionally be employed to help identify optimal dosage amounts. The precise dose to be employed will also depend on the route of administration, and the seriousness of the Condition and can be decided according to the judgment of a practitioner and/or each animal's circumstances.

When trans-(-)-Δ^9-THC compositions containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids are administered orally, the effective dosage amount is from 0.005 mg/kg of body weight to 0.4 mg/kg of body weight every 4 hours, although it is typically 0.1 mg/kg of body weight or less. In one example, the effective dosage amount is from 0.005 mg/kg of body weight to 0.4 mg/kg of body weight; in another example, the effective dosage amount is from 0.01 mg/kg of body weight to 0.1 mg/kg of body weight; and in another example, the effective dosage amount is from 0.01 mg/kg of body weight to 0.075 mg/kg of body weight.

[0127] The oral dosage forms typically comprise an amount of trans-(-)-Δ^9-THC from 0.1 mg to 20 mg; in another example, from 2.5 mg to 10 mg; in another example, 2.5 mg; in another example, 5 mg; and in another example, 10 mg.

[0128] In one example, an effective dosage amount is administered every 24 hours until the Condition is abated. In another example, an effective dosage amount is administered every 12 hours until the Condition is abated. In another example, an effective dosage amount is administered every 8 hours until the Condition is abated. In another example, an effective dosage amount is administered every 6 hours until the Condition is abated. And in another example, an effective dosage amount is administered every 4 hours until the Condition is abated.

[0129] In certain cases, it may be desirable to introduce the pharmaceutical compositions into the central nervous system by any suitable route, including intraventricular and intrathecal injection. Intraventricular injection may be facilitated by an intraventricular catheter, for example, attached to a reservoir, such as an Ommaya reservoir.

[0130] Pulmonary administration can also be employed, e.g., by use of an inhaler or nebulizer, and formulation with an aerosolizing agent, or via perfusion in a fluorocarbon or synthetic pulmonary surfactant. In certain cases, the pharmaceutical compositions can be formulated as a suppository, with traditional binders and carriers such as triglycerides.

[0131] The trans-(-)-Δ^9-THC composition containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids can be delivered in a vesicle, in particular a liposome *(see* Langer, Science 249:1527-1533 (1990); Treat et al., in Liposomes in the Therapy of Infectious Disease and Cancer, Lopez-Berestein and Fidler (eds.), Liss, New York, pp. 353-365 (1989); Lopez-Berestein, ibid., pp. 317-327).

[0132] The trans-(-)-Δ^9-THC composition containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids can be delivered in a controlled-release system. In one example, a pump can be used *(see* Langer, *supra*; Sefton, CRC Crit. Ref. Biomed. Eng. 14:201 (1987); Buchwald et al., Surgery 88:507 (1980); Saudek et al., N. Engl. J. Med. 321:574 (1989)). In another example, polymeric materials can be used (*see* Medical Applications of Controlled Release, Langer and Wise (eds.), CRC Pres., Boca Raton, Fla. (1974); Controlled Drug Bioavailability, Drug Product Design and Performance, Smolen and Ball (eds.), Wiley, New York (1984); Ranger and Peppas, J. Macromol. Sci. Rev. Macromol. Chem. 23:61 (1983); *see also* Levy et al., Science 228:190 (1985); During et al., Ann. Neurol. 25:351 (1989); Howard et al., J. Neurosurg. 71:105 (1989)). In another example, a controlled-release system can be placed in proximity of the target of the pharmaceutical compositions, thus requiring only a fraction of the systemic dose *(see, e.g.,* Goodson, in Medical Applications of Controlled Release, supra, vol. 2, pp. 115-138 (1984)). Other controlled-release systems discussed in the review by Langer (Science 249:1527-1533 (1990)) can be used.

[0133] Pharmaceutical packs or kits comprising one or more containers filled with the trans-(-)-Δ^9-THC composition containing at least 99.0% by weight of trans-(-)-Δ^9-THC based on the total amount of cannabinoids are also considered. Optionally associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration.

[0134] The following examples are set forth to assist in understanding the invention and do not limit the invention described and claimed herein.

5. Examples

[0135] Unless otherwise stated, all reactions were carried out under argon or nitrogen atmosphere.

[0136] Unless otherwise stated, the phrase "cold water, " "cold hexane," or "cold heptane" means water, hexane, or heptane, respectively, at a temperature of from about 0° to about 5°C.

[0137] *Reagents and Solvents:* Unless otherwise stated, all reagents and solvents were purchased from Aldrich Chemical Company and used without further purification.

[0138] *High Performance Liquid chromatography:* High performance liquid chromatography (HPLC) was carried out under the following conditions, and the purity of the samples eluents was calculated from the resultant area percentages:

Standard HPLC was carried out using a 3 μm C_{18}-stationary phase column (150 x 4.6 mm); a mobile phase of the following composition: THF (71%), MeOH (24%) and water (5%) for 25 min, gradient to THF (71%), MeOH (5%) and water (24%) in 10 min, and THF (71%), MeOH (24%) and water (5%) for 10 min; a flow rate of 1 mL/min; and a UV detector at 228 nm.

Chiral HPLC method 1 was carried out using a 20 μm Chiralpak AD 250 x 4.6 mm column; a mobile phase of heptane:isopropanol (95:5 (v:v)); a flow rate of 1 mL/min; and a UV detector at 228 nm. The concentration of the

sample was about 1 mg per 1 mL of heptane.

Chiral HPLC method 2 was carried out using a 5 μm Chiralpak AD-H 250 x 4.6 mm (Diacel) column; a mobile phase of hexane:ethanol (95:5 (v:v)) for CBD and hexane:isopropanol (90:10 (v:v)) for Δ^9-THC; a flow rate of 1 mL/min; and a UV detector at 228 nm. The concentration of the sample was about 1 mg per 1 mL of hexane.

Gas chromatography: Gas chromatography (GC) was carried out under the following conditions, and the purity of the eluents was calculated from the resultant area percentages:

Standard GC was carried using a HP-5 capillary column (length - 30 m, ID - 0.25 mm); a stationary phase of 5% diphenyl/95% dimethyl)polysiloxane (0.25 μm film); an injection temperature of 230°C; a detector/temperature (FID) of 270°C; and an oven temperature program using a hold at 100°C for 3 min, increasing to 240°C at 10°C per minute, holding at 240°C for 10 min, increasing to 270°C per min, and holding at 270°C for 10 min. The concentration of the GC sample was about 1 mg per 1 mL of EtOH.

Chiral GC was carried out in a manner similar to that described above for standard GC, except that an Alpha-DEX-120, 30 m x 0.25 mm column was used; the injection temperature was 250°C; and the oven temperature was 90°C (isothermal).

Powder x-ray diffraction patterns: Powder x-ray diffraction analysis was carried out by known methods using a PANALYTICAL (Philips) X'Pert Pro MPD x-ray powder diffraction system (CuK_α radiation, PW3050/60 goniometer, PW3011/20 proportional detector). The Bragg-Brentano scheme was used for beam focusing.

Nuclear Magnetic Resonance Spectroscopy: Nuclear magnetic resonance (NMR) spectra were recorded on a Bruker AM-200 (^1H at 200 MHz, ^{13}C at 50 MHz) or a Bruker AM-400 (^1H at 400 MHz) instruments using $CDCl_3$ (unless otherwise stated) as a solvent. Chemical shifts are in δ (ppm) relative to internal TMS.

Melting points: Melting point determinations were carried out in open capillary tubes using a Buchi B-545 capillary melting point apparatus or with a Mettler-Toledo FP-81 Melting point accessory with FP-900 processor. The melting points are uncorrected.

5.1. Example 1: Synthesis of (-)-*cis*-*p*-Menth-2,8-dien-1-ol

[0139] *Preparation of(-)-(1R,2R,S5)-2-phenylthio-8-p-menthen-1-ol:* A mixture of (-)-limonene oxide (152.2 g, 1.00 mol (about 1:1 cis:trans diastereomeric mixture) (Aldrich Chemical), thiophenol (60.6 g, 0.55 mol) (Fluka Chemical, Buchs, Switzerland), potassium carbonate (82.9 g, 0.60 mol), N,N-dimethylformamide (18.9 g, 0.26 mol) and toluene (400 mL) was stirred at 117°C for 19 hours under an Ar atmosphere. The mixture was cooled to 25°C and water (300 mL) was added. The resultant organic phase was collected, and the water layer was extracted with toluene (3 x 200 mL). The combined organic phases were washed with water (1 x 400 mL) and a 15% solution of brine (1 x 410 mL). The organic phase was then dried over Na_2SO_4 (30 g), filtered, and the resultant filtrate concentrated under pressure at 65°C. The resultant brown oil (200.5 g) was fractionally distilled under reduced pressure to provide (-)-*cis*-limonene oxide (33.7 g) (28.1° to 32.1 °C @ 1.1 mbar) and (-)-(1R, 2R,4S)-2-phenylthio-8-*p*-menthen-1-ol (147.4 g) (128.1° to 138.2°C @ 1.2 mbar) having a purity (GC) of 90.2%. An analytical sample of (-)-(1*R*,2*R*,4*S*)-2-phenylthio-8-*p*-menthen-1-ol had mp of 50-51 °C (hexane) and a purity (GC) of 99.0%.

Optical rotation: $[\alpha]_D^{20}$ -110° (c = 1.55, $CHCl_3$).

^1H NMR agreed with the structure.

[0140] *Preparation of (1R,2R,4S)-1-Hydroxy-8-p-menthen-2-phenylsulfoxide*: (-)-(1R,2R,4S)-2-phenylthio-8-*p*-menthen-1-ol (147 g; 0.56 mol) was dissolved in methyl alcohol (1.35 L) with stirring at 25°C under an Ar atmosphere, and the resultant solution was cooled to -10° to -5°C. A solution of OXONE® (potassium peroxymonosulfate) (279.1 g, 0.448 mol) (Aldrich Chemical) in water (1.35 L) was added dropwise to the methyl alcohol solution over 2 hours at - 10° to -5°C, and the resultant mixture was stirred for an additional 30 min at -10° to - 5°C. The mixture was warmed to 20° to 25°C, water (2.1 L) was added, and the resultant biphasic mixture was extracted with dichloromethane (3 x 910 mL). The combined organic phases were dried over sodium sulfate and filtered, and the resultant filtrate was concentrated under reduced pressure at 60° C to provide 150.9 g of a residue. The residue was then purified by chromatography on a silica gel column (eluent: n-heptane/ethyl acetate 9:1 then 8:2). The fractions containing mainly (1R,2R,4S)-1-hydroxy-8-*p*-menthen-2-phenyl sulfoxide were combined and concentrated under vacuum for 10 hours at 40° to 50°C to provide (1R,2R,4S)-1-hydroxy-8-*p*-menthen-2-phenyl sulfoxide as a mixture of two diastereomers. Yield: 86.1 g; 55.2%. The product was stored in freezer.

[0141] *(-)-cis-p-Mentha-2,8-dien-1-ol*: A mixture of (1R,2R,4S)-1-hydroxy-8-*p*-menthen-2-phenylsulfoxide (86 g, 0.31 mol) and piperidine (71.0 g, 0.83 mol) in dimethylsulfoxide (910 mL) was heated to 163°C under a flowing Ar atmosphere, and the resultant mixture was stirred at 163°C for 3 hours. The mixture was cooled to 20° to 25°C, treated with water (800 ml), and extracted with diethyl ether (2 x 400 mL). The combined organic phases were washed with 1N HCl (160 mL), a 7% solution of sodium hydrogen carbonate (150 mL), brine (150 mL), and dried over sodium sulfate. The organic phase was then concentrated under reduced pressure. The resultant residue (93.3 g) was purified by silica gel column

chromatography (eluent: n-heptane followed by n-heptane:ethyl acetate (1:9 (v:v)), the fractions containing mainly (-)-*cis*-*p*-mentha-2,8-dien-1-ol were combined and concentrated under reduced pressure at 40° to 50°C over 10 hours to provide (-)-*cis*-*p*-mentha-2,8-dien-1-ol. Yield: 26.1 g; 55%. Analysis (GC) of the product indicated that it was 90.9 % pure.
Optical rotation: $[\alpha]_D^{25}$ -69° (neat).
^{1}H NMR agreed with the structure.

5.2. Example 2: Synthesis of (+)-*cis*-*p*-Menth-2,8-dien-1-ol

[0142] (+)-p-mentha-2,8-dien-1-ol was prepared as described in Example 1, except that (+)-limonene oxide (1:1 cis/trans diastereomeric mixture) was used instead of (-)-limonene oxide. Analysis (GC) of the resultant product indicated that it had a purity of 91.0%.
Optical rotation: $[\alpha]_D^{25}$ +78° (neat).

5.3. Example 3: Synthesis of (+)-*cis*-*p*-Menth-2,8-dien-1-ol

[0143] (±)-*p*-Mentha-2,8-dien-1-ol was prepared by mixing equivalent quantities of (-)-*p*-mentha-2,8-dien-1-ol of Example 2 with (+)-*p*-mentha-2,8-dien-1-ol of Example 1.

5.4. Example 4: Synthesis of (+)-CBD

[0144] *Synthesis of crude (+)-CBD (**3b**)*: A mixture of olivetol (3.6 g, 20 mmol), zinc chloride (3.5 g, 26 mmol), water (3.5 mL, 19 mmol) and dichloromethane (35 mL) was refluxed for 1 hour. A solution of (-)-p-mentha-2,8-dien-1-ol (3.0 g, 20 mmol) in dichloromethane (10 mL) was added drop-wise over 0.75 hour to the refluxing mixture, and the resultant reaction mixture was mixed for 0.5 hours at reflux. The mixture was cooled to 25°C, ice water (50 mL) was added, and the resultant biphasic mixture stirred for 20 minutes at 0°C. The resultant organic phase was collected, washed with water (2 x 20 mL) and 5% $NaHCO_3$ (20 mL). The organic phase was dried over Na_2SO_4, filtered, and concentrated under reduced pressure to provide 6.0 g of first crude (+)-CBD residue. Analysis (GC) of the first crude (+)-CBD residue indicated that it contained (+)-CBD (46.9%) and abn-(-)-CBD (19.7%). The first crude (+)-CBD residue was purified by column chromatography on silica gel (eluent MTBE/hexane) to provide 2.4 g of a second crude (+)-CBD residue.
[0145] *Synthesis of (+)-CBD-bis(3,5-dinitrobenzoate)(**4b**)*: A solution of 3,5-dinitrobenzoyl chloride (3.4 g, 14.7 mmol) in dichloromethane (10 mL) was added dropwise to a stirred mixture of the second crude (+)-CBD residue (2.4 g), 4, N,N-dimethylaminopyridine (0.05 g), pyridine (6 mL) and dichloromethane (15 mL) at 0° to 5°C. The mixture was allowed to warm to 25°C and stirred for 2 hour at 25°C. The mixture was then poured into a mixture of 37% HCl (6 mL), ice (75 g) and dichloromethane (50 mL). The resultant organic phase was collected, washed with brine (15 mL), 5% $NaHCO_3$ (15 mL), dried over Na_2SO4, and filtered. The resultant filtrate was concentrated under reduced pressure to provide 5.2 g of crude (+)-CBD-bis(3,5-dinitrobenzoate) (**4b**). A solution of the crude (+)-CBD-bis(3,5-dinitrobenzoate) (5.2 g) in a 10:1 (vol:vol) mixture of isopropanol and ethylacetate (70 mL) was stirred overnight at 25°C and filtered. The resultant precipitate was washed with 10:1 (vol:vol) mixture of isopropanol and ethylacetate (3 x 10 mL) and dried under reduced pressure to provide crystalline (+)-CBD-bis(3,5-dinitrobenzoate) (**4b**). Yield: 3.7 g, 26.5%.
Melting point: 90-92°C (dec.).
Optical rotation: $[\alpha]_D^{20}$ +80° (c = 0.4, $CHCl_3$).
[0146] *Synthesis of (+)-CBD (**3b**)*: A mixture of the crystalline (+)-CBD bis(3,5-dinitrobenzoate) (**4b**) (3.5 g, 5.0 mmol), butylamine (3.7 g, 50 mmol) and toluene (20 mL) was stirred at room temperature for 12 hours and concentrated under reduced pressure. The resultant residue was purified by column chromatography on silica gel (eluent hexane:MTBE (70:1 (v:v)) to provide 1.3 g of (+)-CBD as an oil. A solution of the (+)-CBD (1.3 g) in hexane (1 mL) was kept overnight at -15°C. The resultant mixture was then filtered, and the resultant solids were dried under reduced pressure to provide (+)-CBD (**3b**) as white crystals. Yield: 1.2 g, 64%. Analysis (GC) of the product indicated that it was 98.6% pure.
Melting point: 64-66°C
Optical rotation: $[\alpha]_D^{20}$: +126° (c = 0.12, 95% EtOH).

5.5. Example 5: Preparation of (±)-Δ^8-THC

[0147] A solution of methanesulfonic acid (1.1 g, 11 mmol) in dichloromethane (6 mL) was added to solution of olivetol (10.0 g, 55.5 mmol) and (±)-*p*-mentha-2,8-dien-1-ol (8.5 g, 55.5 mmol) in dichloromethane (130 mL). The resultant mixture was refluxed for 4 hours with removal of water using a Dean-Stark separator. The mixture was then cooled to 25°C and treated with aqueous $NaHCO_3$. The resultant organic phase was collected and concentrated under reduced pressure. The resultant residue was dissolved in heptane (110 mL) and washed with 10% NaOH (130 mL), and the resultant organic phases were concentrated under reduced pressure to provide 15.6 g of crude (±)-Δ^8-THC. Analysis

(GC) of the crude product indicated that it had a purity of 61.7%.

5.6. Example 6: Preparation of (-)-Δ^8-THC

[0148] Crude (-)-Δ^8-THC (**2a**) was prepared in a manner similar to that described in Example 5 for the preparation of crude (±)-Δ^8-THC, except that (+)-*p*-mentha-2,8-dien-1-ol was used instead of (±)-*p*-mentha-2,8-dien-1-ol.

5.7. Example 7: Reparation of (+)-Δ^8-THC

[0149] Crude (+)-Δ^8-THC (**2b**) was prepared in a manner similar to that described in Example 5 for the preparation of crude (+)-Δ^8-THC, except that (-)-*p*-mentha-2,8-dien-1-ol was used instead of (±)-*p*-mentha-2,8-dien-1-ol.

5.8. Example 8: Two-part Synthesis of trans-(-)-Δ^9-THC

[0150] *Synthesis of (-)-CBD (**3a**)*: A solution of (+)-p-mentha-2,8-dien-1-ol (84.5 g, 0.56 mol) in dichloromethane (325 mL) was added drop-wise over 1 hour to a stirred mixture of olivetol (100.0 g, 0.56 mol), zinc chloride (100.3 g, 0.72 mol), water (10.0 mL, 0.56 mol) and dichloromethane (1 L) at 40°C. The mixture was stirred for an additional 30 minutes at 40°C. The mixture was cooled to 25°C, poured into ice water (500 g), and the resultant biphasic mixture stirred for 20 minutes at 0°C. The resultant organic phase was collected and washed with cold water (2 x 250 mL). The organic phase was collected and concentrated under reduced pressure to provide a first residue (185.5 g). Analysis (GC) of the first residue indicated that it contained (-)-CBD (51.8%), *abn*-CBD (13.2%), olivetol (8.0%) and dialkylated olivetol (13.4%).

[0151] The first residue (185.5 g) was dissolved in n-heptane (1.1 L), and the resultant solution was admixed with a solution of 10% sodium hydroxide (1.3 L). The resultant organic phase was collected, washed with water (250 mL), and concentrated under reduced pressure to provide an oily-brown second residue (124.3 g). Analysis (GC) of the second residue indicated that it contained (-)-CBD (66.0%), *abn*-CBD (0.0%), olivetol (0.0%) and dialkylated olivetol (16.8%).

[0152] The second residue (124.3 g) was fractionally distilled (171°-178°C; 0.1 mm Hg) to provide a 87.0 g of a distillate. Analysis (GC) of the distillated indicated that it contained 74.3% of (-)-CBD.

[0153] The distillate (87.0 g) was dissolved in heptane (425 mL) at 57°C and filtered. The resultant filtrate was cooled to 0° to 5°C and seeded with ~0.02 mg of powdered crystalline (-)-CBD (**3a**). The seeded solution was stirred at 0° to 5°C for 5 hours then at -15° to -20°C for 48 hours. The resultant mixture was filtered, and the resultant solids were washed with cold heptane. The solids were then dried under reduced pressure at 40°C to provide (-)-CBD (**3a**). Yield: 39.2 g; 22%. Analysis (GC) of the product indicated that it contained (-)-CBD (**3a**) (97.1 %) and trans-(-)-Δ^9-THC (**1a**) (1.44%). The structure of **3a** was confirmed by ^{1}H NMR spectroscopy. An analytical sample was prepared by recrystallizing a portion of the crude 3a from heptane as described above.

Melting point: 64°-65°C.

Optical rotation: $[\Delta]_D^{20}$ -132° (c = 0.12, 95% EtOH).

[0154] *Synthesis of trans-(-)-Δ^9-THC (**1a**)*: A solution of 15.0 g (47.8 mmol) of the crystallized (-)-CBD (**3a**) in anhydrous dichloromethane (45 mL) was added drop-wise over 1 hour to a stirred solution of $BF_3{\cdot}Et_2O$ (8.4 g, 59.2 mmol) in anhydrous dichloromethane (180 mL) at -10°C under an Ar atmosphere. The mixture was stirred for 2 hours at -10°C and poured into ice water (100 g). The resultant biphasic mixture was further stirred for 20 minutes at 0°C. The resultant organic phase was collected, washed with cold water (50 mL), 7 % aqueous sodium bicarbonate (50 mL), and water (50 mL). The organic phase was dried with Na_2SO_4 and filtered. The resultant filtrate was concentrated under reduced pressure at 40°C to provide trans-(-)-Δ^9-THC (**1a**) as a yellow oil. Yield: 14.9 g, 99%. Analysis (CG) of the product indicated that it contained 81.9% of trans-(-)-Δ^9-THC (**1a**).

5.9. Example 9: One-Pot Synthesis of trans-(-)-Δ^9-THC

[0155] A mixture of olivetol (50.0 g, 0.28 mol), zinc chloride (50.0 g, 0.36 mol) and anhydrous dichloromethane (510 mL) was stirred at 40°C for 1 hour under an Ar atmosphere. A solution of (+)-*p*-mentha-2,8-dien-1-ol (42.2 g, 0.28 mol) and dichloromethane (155 mL) was added drop-wise over 1 hour to the stirred olivetol-containing mixture, and the resultant mixture was stirred for an additional 40 minutes at 40°C. The mixture was cooled to -10°C, and a solution of BF_3Et_2O (23.6 g, 166 mmol) in anhydrous dichloromethane (37 mL) was added dropwise over one hour. The resultant mixture was stirred for 1.5 hours at -10°C. Cold water (250 mL) was added, and the resultant organic phase was collected and washed with cold water (120 mL), 7% aqueous sodium bicarbonate (120 mL), and water (120 mL). The organic phase was dried with Na_2SO_4 (30 g) and filtered. The resultant filtrate was concentrated under reduced pressure to provide trans-(-)-Δ^9-THC (**1a**) as a brown oil. Yield: 89.14 g, 46% based on the trans-(-)-Δ^9-THC content in the oil. Analysis (GC) of the product indicated that it contained trans-(-)-Δ^9-THC (**1a**) (45.1%), (-)-Δ^8-THC (5.06%) (**2a**), (-)-Δ^8-iso-THC (17.6%), CBD (**3a**) (0.71%), olivetol (7.95%) and dialkylated olivetol (10.8 wt.%); no trans-(+)-Δ^9-THC (**1b**)

was detected.

[0156] A solution of the trans-(-)-Δ^9-THC oil (20.0 g) in heptane (120 mL) was thoroughly washed with 10% NaOH (150 mL) and water (50 mL), dried over Na_2SO_4, and filtered. The resultant filtrate was then concentrated under reduced pressure to provide a first crude residue (16.6 g) containing 38.5 wt. % of trans-(-)-Δ^9-THC (**1a**) using HPLC; and trans-(-)-Δ^9-THC (**1a**) (47.4%), Δ^8-THC (**2a**) (8.6%), Δ^8-iso-THC (19.6%), CBD (0.5%), olivetol (0.0%) and dialkylated olivetol (10.9%) using GC.

[0157] A solution of the first crude residue (16.5 g) in heptane (240 mL) was extracted with an aliquot of 9% NaOH in 80% methanol (3 x 180 mL). The combined basic methanolic extracts were acidified to approximately pH 7 with 20% citric acid and extracted with heptane (3 x 90 mL). The combined organic fractions were washed with water (50 mL), dried over Na_2SO_4, and filtered. The resultant filtrate was then concentrated under reduced pressure to provide 13.7 g of crude residue which contained 44.0 wt. % of trans-(-)-Δ^9-THC using HPLC; and trans-(-)-Δ^9-THC (**1a**) (51.8%), Δ^8-THC (**2a**) (10.0%), Δ^8-iso-THC (22.3%), CBD (0.0%), olivetol (0.0%) and dialkylated olivetol (1.3%) using GC.

5.10. Example 10: Synthesis of trans-(+)-Δ^9-THC

[0158] A solution of $BF_3{\cdot}Et_2O$ (0.34 g, 2.4 mmol) in anhydrous dichloromethane (8 mL) was added dropwise with stirring over 1 hour to a solution of the crystalline (+)-CBD (**3a**) from Example 4 (1.1 g, 3.6 mmol) in anhydrous dichloromethane (50 mL) at -5°C. The resultant mixture was stirred for 1.5 hours at -5°C. The mixture was added to a mixture of ice (100 g) and 7% $NaHCO_3$ (100 mL). The resultant organic phase was collected and the aqueous phase extracted with dichloromethane (2 x 20 mL). The combined organic phases were washed with water (20 mL), dried with Na_2SO_4, and filtered. The resultant filtrate was concentrated under reduced pressure at 40°C. The resultant residue was purified by column chromatography on silica gel (stationary phase) using MTBE:hexane (1:100 to 3:100 (v:v)) as eluent to provide crude trans-(+)-Δ^9-THC (**1b**) as a yellow oil: Yield: 0.7 g. Analysis (GC) of the crude trans-(+)-Δ^9-THC indicated that it had a purity of 92.6%.

5.11. Example 11: One-pot Synthesis of trans-(+)-Δ^9-THC

[0159] A mixture of olivetol (14.21 g, 79.6 mmol), zinc chloride (14.25 g, 102.6 mmol) and anhydrous dichloromethane (145 mL) was stirred at 40°C for 1 hour. A solution of (-)-*p*-mentha-2,8-dien-1-ol (12.00 g, 76.6 mol) and anhydrous dichloromethane (45 mL) was added drop-wise over 1 hour at 40°C to the stirred olivetol-containing mixture, and the resultant mixture was stirred for an additional 40 minutes at 40°C. The mixture was cooled to -10°C, and a solution of $BF_3.Et_2O$ (6.7 g, 47 mmol) in anhydrous dichloromethane (12 mL) was added drop-wise over 1 hour at -10°C. The mixture was stirred for 30 minutes at -10°C. Cold water (50 mL) was added, and the resultant biphasic mixture was stirred for an additional 20 minutes at 0°C. The resultant organic phase was collected, washed with cold water (2 x 50 mL), 5% aqueous sodium bicarbonate (50 mL), and water (50 mL). The organic phase was then concentrated under reduced pressure at 40°C, and the resultant residue (24.8 g) was dissolved in n-heptane (140 mL) at 25°C. The resultant solution was washed with 10% aqueous KOH (124 mL), water (2 x 50 mL), dried with $MgSO_4$ (10 g), and filtered. The resultant filtrate was concentrated under reduced pressure at 40°C. The resultant residue (20.7 g) was then fractionally distilled at reduced pressure (0.1 mbar) to provide trans-(+)-Δ^9-THC (**1b**). Yield: 17.16 g, 69 %. Analysis (GC) of the product indicated that it contained trans-(+)-Δ^9-THC (**1b**) (49.2%), Δ^8-iso-THC (25.31 %) and dialkylolivetol (1.29 %); no trans-(-)-Δ^9-THC (**1a**) was detected.

5.12. Example 12: Synthesis of (±)-Δ^9-THC

[0160] A solution of $BF_3{\cdot}Et_2O$ (0.3 g, 2.1 mmol) in anhydrous dichloromethane (8 mL) was added dropwise with stirring over 1 hour to a solution of (±)-CBD (1.0 g, 3.2 mmol) in anhydrous dichloromethane (45 mL) at -5°C. The resultant mixture was stirred for 1.5 hours at -5°C. The mixture was then added to 7% $NaHCO_3$ (50 mL). The resultant organic phase was collected and the aqueous phase extracted with dichloromethane (3 x 30 mL). The combined organic phases were washed with brine (20 mL), dried with Na_2SO_4 and filtered. The resultant filtrate was concentrated under reduced pressure. The resultant residue was purified by column chromatography on silica gel (stationary phase) and MTBE:hexane (1:100 to 2:100 (v:v)) as eluent to provide crude (±)-Δ^9-THC as a yellow oil. Yield: 0.6 g, 56%. Analysis (GC) of the (±)-Δ^9-THC oil indicated that it had a purity of 92.6% purity. The oily (±)-Δ^9-THC (0.6 g) was dissolved in hexane (0.5 mL), and the resultant mixture was maintained at -15 °C for 24 hours. The resultant mixture was filtered, washed with cold hexane (3 x 1 mL) and dried under reduced pressure to provide (±)-Δ^9-THC as slightly rose crystals. Yield: 0.4 g. Melting point: 65-66 °C.

5.13. Example 13: One-Pot Synthesis of (±)-Δ^9-THC

[0161] A mixture of olivetol (11.84 g, 65.7 mmol), zinc chloride (11.87 g, 85.4 mmol) and anhydrous dichloromethane (120 mL) was stirred at 40°C for 1 hour. A solution of (+) *p*-mentha-2,8-dien-1-ol (5.00 g, 32.84 mol), (-)-*p*-mentha-2,8-dien-1-ol from Example 1 (5.00 g, 32.84 mol) and anhydrous dichloromethane (37 mL) was added drop-wise over 1 hour at 40° to the stirred olivetol-containing mixture, and the resultant mixture was stirred for an additional 40 minutes at 40°C. The mixture was cooled to -10°C, and a solution of $BF_3{\cdot}Et_2O$ (5.6 g, 39.4 mmol) in anhydrous dichloromethane (10 mL) was added drop-wise over 1 hour at -10°C. The mixture was stirred for 30 minutes at -10°C, and 50 mL of cold water were added. The resultant biphasic mixture was stirred for an additional 20 minutes at 0°C. The resultant organic phase was collected and washed with cold water (2 x 50 mL), 8% aqueous sodium bicarbonate (50 mL), and water (50 mL). The organic phase was concentrated under reduced pressure at 40°C. The resultant residue (20.5 g) was dissolved in n-heptane (115 mL) at 25°C and washed with 10% aqueous KOH (100 mL) for 40 min at 25°C and water (50 mL). The organic phase was then concentrated under reduced pressure at 50°C to provide 17.1 g of crude (±)-Δ^9-THC as a brown oil.

[0162] A portion of the crude (±)-Δ^9-THC oil (2.4 g) was dissolved in a minimal amount of heptane and purified by chromatography in a single pass using a Merck-Knauer PP K-1800 preparative chromatograph with one cylinder (50 mm x 210 mm of LUNA CM 10 μm; loading capacities 600 mg; eluent: *n*-heptane). Fractions containing (±)-Δ^9-THC were combined and concentrated under reduced pressure at 40°C to provide (±)-Δ^9-THC (1). Yield: 1.1 g. Analysis (GC) of the product indicated that it contained (±)-Δ^9-THC (**1**) (91.27 %), iso-Δ^8-THC (1.87 %) and Δ^8-THC (1.08 %).

5.14. Example 14: Preparation of (±)-Δ^9-THC

[0163] A mixture of olivetol (15.0 g, 83.2 mmol), zinc chloride (15.0 g, 108 mmol) and anhydrous dichloromethane (150 mL) was stirred at 40°C for 1 hour. A solution of (±)-*p*-mentha-2,8-dien-1-ol (12.7 g, 83.2 mmol) and anhydrous dichloromethane (45 mL) was added drop-wise over 1 hour at 40° to the stirred olivetol-containing mixture, and the resultant mixture was stirred for an additional 0.50 hours at 40°C. The mixture was cooled to -10°C, and a solution of $BF_3{\cdot}Et_2O$ (7.1 g, 49.4 mmol) in anhydrous dichloromethane (11 mL) was added drop-wise to the mixture over 1 hour at -10°C. The mixture was stirred for 0.50 hours at -10°C, and 80 mL of cold water was added with stirring to form a biphasic mixture. The organic phase was collected and washed with cold water (80 mL), 5% aqueous sodium bicarbonate (80 mL), and water (80 mL). The organic phase was dried over Na_2SO4 and filtered, and the resultant filtrate was concentrated under reduced pressure to provide 28.5 g of a first crude (±)-Δ^9-THC residue. Analysis of the residue indicated that it contained (±)-Δ^9-THC (30.3 %) using HPLC; and (±)-Δ^9-THC (45.2 %), Δ^8-THC (3.2 %), (±)-Δ^8-iso-THC (17.3 %). CBD (4.0 %), olivetol (8.3 %), and dialkylated olivetol (11.7 %) using GC.

[0164] A portion of the first crude (±)-Δ^9-THC residue (28.5 g) was dissolved in heptane (165 mL), and the resultant solution was washed with 10% NaOH (200 mL) and water (80 mL). The organic solution was then dried by azeotropic distillation and concentrated under reduced pressure to provide a second crude (±)-Δ^9-THC residue. Yield: 23.5 g, 37.6 %. Analysis of the second crude (±)-Δ^9-THC residue indicated that it contained (±)-Δ^9 -THC (37.6 %) using HPLC; and (±)-Δ^9-THC (50.7 %), Δ^8-THC (3.8%), (+)-Δ^8-iso-THC (19.6 %), CBD (4.4 %), olivetol (0.0 %), and dialkylated olivetol (12.8 %) using GC.

5.15. Example 15: Preparation of (±)-Δ^9-THC from a Mixture of Crude trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC

[0165] Trans-(-)-Δ^9-THC was prepared as described in Example 14 for 9 preparing the crude second (+)-Δ^9 -THC residue, except that (+)-p-mentha-2,8-dien-1-ol was used instead of (±)-*p*-mentha-2,8-dien-1-ol. Analysis (HPLC) of the resultant crude trans-(-)-Δ^9-THC indicated that it contained 41.4% by weight of trans-(-)-Δ^9-THC.

[0166] Trans-(+)-Δ^9-THC was prepared as described in Example 14 for preparing the crude second (±)-Δ^9-THC residue, except that (-)-p-mentha-2,8-dien-1-ol was used instead of (±)-p-mentha-2,8-dien-1-ol. Analysis (HPLC) of the resultant crude trans-(+)-Δ^9-THC indicated that it contained 37.5% by weight of trans-(-)-Δ^9-THC.

[0167] The crude trans-(-)-Δ^9-THC (24.3g; 10.0 g of trans-(-)-Δ^9-THC) and trans-(+)-Δ^9-THC (26.7g; 10.0 g of trans-(+)-Δ^9-THC) were dissolved in heptane (425 mL) at 25°C. The resultant solution was admixed with 2 x 180 mL of a solution of 9% aqueous NaOH:methanol (20:80 (v:v)). The methanolic phases were combined and treated with 10% citric acid at 0°C to about 5°C until the pH was about 7. Heptane (290 mL) was added, and the resultant organic phase was washed with water. The organic phase was then dried over Na_2SO_4 and filtered, and the resultant filtrate was concentrated under reduced pressure to provide 41.8 g of crude (±)-Δ^9-THC as a brown oil. Analysis (HPLC) of the crude (±)-Δ^9-THC indicated that it had a purity of 48%.

[0168] The crude (±)-Δ^9-THC (41.8 g) was dissolved in heptane (85 mL), and the resultant solution was cooled to 0°C and seeded with crystalline (±)-Δ^9-THC (100 mg). The resultant mixture was further cooled to -15°C for 12 hour and filtered. The resultant solids were washed with cold heptane (3 x 10 mL) and dried under reduced pressure to provide

(±)-Δ^9-THC as a white crystalline solid. Yield: 8.7g, 43%. Analysis (HPLC) of the product indicated that it had a purity of 96.5%. The crystalline (+)-Δ^9-THC remained white after at least three days at 25°C.

5.16. Example 16: Preparation of (±)-Δ^9-THC from (±)-Δ^8-THC

[0169] *Preparation of (±)-9-chloro-trans-hexahydrocannabinol*: A mixture of crude (±)-Δ^8-THC from Example 5 (15.6 g; 9.63 g of (±)-Δ^8-THC), zinc chloride (4.66g, 34.23 mmol) and anhydrous dichloromethane (310 mL) was stirred for 0.5 hours at 25°C under an Ar atmosphere. The mixture was cooled to 0°C, and gaseous hydrogen chloride was bubbled through the mixture for 1.5 hours. The mixture was poured into an ice bath (150 g), and the resultant biphasic mixture was stirred for 1 hour at 0 to 5°C. The organic phase was collected and washed with cold water (2 x 100 mL), 8% sodium bicarbonate solution (100 mL), and water (100 mL). The organic phase was dried over anhydrous Na_2SO_4 (15 g), and filtered. The resultant filtrate was then concentrated under reduced pressure at 30°C. The resultant residue (16.3 g) was dissolved in n-heptane (33 mL), cooled to 0°C, and seeded with (±)-9-chloro-trans-hexahydrocannabinol (0.01 g). The resultant mixture was then stirred at 0°C for 5 hours, cooled to -15°C, and stirred at -15°C for 60 hours. The mixture was filtered and the resultant solids washed with cold n-heptane (14 mL). The solids were then dried under reduced pressure at 50°C to provide (±)-9-chloro-trans-hexahydrocannabinol. Yield: 5.7 g; 32.7%. Analysis (HPLC) of the (±)-9-chloro-trans-hexahydrocannabinol indicated that it had a purity of 95.2%. An analytical sample, recrystallized from heptane, had a melting point of 89-90 °C. The purity (HPLC) of the analytical same was 99.6%.
Optical rotation: $[\alpha]_D^{20}$ 0.0° (c = 0.53, $CHCl_3$).
^1H NMR agreed with the structure.
^{13}C NMR ($CDCl_3$) δ 13.9, 19.1, 22.4, 24.2, 27.6, 30.4, 31.3, 31.5, 34.1, 35.3, 42.0, 44.8, 48.7, 72.6, 76.7, 107.7, 108.9, 110.0, 142.8, 154.5, 155.0.

[0170] The x-ray powder diffraction pattern of the crystalline (±)-9β-Cl-HHC had characteristic peaks expressed in degrees 2θ at approximately 7.5, 11.2, 13.3, 14.9, 15.4, 15.9, 19.4, 19.7, 20.0 and 22.5.

[0171] *Preparation of (±)-Δ^9-THC*: A mixture of potassium-tert-amylate (6.6 g), (±)-9-chloro-trans-hexahydrocannabinol (5.7 g, 16.2 mmol) and anhydrous toluene (280 mL) was stirred for 75 minutes at 65°C. The mixture was cooled to 25°C and poured into ice water (100 g). The resultant organic phase was collected and washed with cold water (2 x 100 mL), 7% sodium bicarbonate, and water (2 x 100 mL). The organic phase was then dried over anhydrous Na_2SO_4 and concentrated under reduced pressure. The resultant residue (5.35 g) was dissolved in 9 n-heptane (3.4 mL), cooled to 0°C, and seeded with (±)-Δ^9-THC (0.01 g). The resultant mixture was stirred at 0°C for 5 hours, cooled to -15°C, and stirred at -15°C for 60 hours. The mixture was filtered and the resultant solids washed with cold n-heptane (4 mL). The solids were then dried under reduced pressure at 50°C to provide (±)-Δ^9-THC. Yield: 3.3g, 64.7%. Analysis (HPLC) of the product indicated that it had a purity of 97.23%.

5.17. Example 17: Purification of (±)-Δ^9-THC

[0172] *Preparation of (±)-Δ^9-THC m-nitrobenzenesulfonate:* A mixture of the second crude (±)-Δ^9-THC residue of Example 14 (20.0 g; 7.52 g of (±)-Δ^9-THC), 3-nitrobenzenesulfonyl chloride (14.5 g, 65.4 mmol), triethylamine (9.7 g) and dichloromethane (300 mL) was stirred at 25°C for 1 hour. The resultant admixture was then treated with cold water (200 mL). The resultant organic phase was collected and washed sequentially with 10% HCl (80 mL), water (100 mL), 5% $NaHCO_3$ (100 mL) and water (100 mL). The organic phase was then dried over Na_2SO_4 and filtered. The resultant filtrate was concentrated under reduced pressure to provide 25.8 g of a first crude (±)-Δ^9-THC *m*-nitrobenzenesulfonate residue. Analysis (HPLC) of the first crude (±)-Δ^9-THC *m*-nitrobenzenesulfonate residue indicated that it had a purity of 42.9 wt.%.

[0173] The first crude (±)-Δ^9-THC *m*-nitrobenzenesulfonate residue was dissolved in isopropanol (95 mL) at 50°C. The resultant solution was cooled to room temperature, seeded with powdered crystalline (±)-Δ^9-THC *m*-nitrobenzenesulfonate, cooled to 0°C, and stirred for 12 hour at 0°C. The resultant mixture was filtered, and the resultant solids were washed with cold heptane (65 mL). The solids were then dried under reduced pressure to provide 10.3 g of second crude (±)-Δ^9-THC *m*-nitrobenzenesulfonate residue as a yellow solid. Analysis (HPLC) of the second crude (±)-Δ^9-THC *m*-nitrobenzenesulfonate residue indicated that it had a purity of 79.1 %.

[0174] The second crude (±)-Δ^9-THC *m*-nitrobenzenesulfonate (10.0 g) was dissolved in dichloromethane (13 mL), and the resultant solution was added to a distillation pot equipped with a 10 cm Vigreux column and an addition port. The contents of the distillation pot were then distilled while isopropanol (40 mL) was continuously added dropwise to the mixture through the addition port. The distillation was stopped when the temperature of vapors in the head of the column reached 82.4 °C. The contents of the distillation pot where cooled to 0°C to 5°C, and the resultant suspension was stirred for 12 hours at 0°C to about 5°C. The suspension was filtered, and the resultant solids were washed with cold heptane (22 mL). The solids were then dried under reduced pressure to provide crystalline (±)-Δ^9-THC *m*-nitrobenzenesulfonate. Yield: 7.0 g, 59%. Analysis (HPLC) of the product indicated that it had a purity of 99.0%.

Melting point: 105-107°C.

X-ray powder diffraction pattern: Characteristic peaks expressed in degrees 2θ where observed at approximately 9.3, 10.6, 12.5, 15.2, 18.7, 19.3, 21.2 and 22.9.

[0175] *Preparation of (±)-Δ9-THC*: A mixture of the crystalline (±)-Δ9-THC m-nitrobenzenesulfonate (4.5 g, 7.5 mmol), 50% NaOH (5.3 g), and methanol (110 mL) was stirred at 50°C for about 1-2 hours then cooled to room temperature. The cooled mixture was then treated with cold water (1 x 150 mL) followed by treatment with 10% HCl until the pH was about 7. The resultant mixture was extracted with heptane (3 x 75 mL), and the combined organic extracts were washed with 7% $NaHCO_3$ (100 mL) and water (100 mL). The organic phase was dried over Na_2SO_4 and filtered. The resultant filtrate was then concentrated under pressure to provide 2.5 g of crude (±)-Δ9-THC. Analysis (HPLC) of the crude product indicated that it contained 92.6 % by weight of (±)-Δ9-THC.

[0176] The crude (±)-Δ9-THC was dissolved in heptane (5 mL) at 40°C. The resultant solution was cooled to 0°C, seeded with powdered crystalline (±)-Δ9-THC, and stirred for 12 hours at -15°C. The resultant mixture was filtered and the resultant solids washed with cold heptane (3.5 mL). The solids were then dried under reduced pressure to provide (±)-Δ9-THC as off-white crystals. Yield: 2.1 g, 74%. The crystalline (±)-Δ9-THC was stable at 25°C in the presence of air and laboratory lighting. Analysis (HPLC) of the product indicated that it had a purity of 99.0%. An analytical sample that was recrystallized from hexane had a melting point of 65°-66°C.

Optical rotation: $[\alpha]_D^{20}$ 0.0° (c = 0.53, $CHCl_3$).

^1H NMR: The spectra of the product agreed with the structure.

5.18. Example 18: Reparation of (+)-Δ9-THC from trans-(-)-Δ9-THC and trans-(+)-Δ9-THC

[0177] A solution of trans-(-)-Δ9-THC (**1a**) (10 g; 9.35 g of trans-(-)-Δ9-THC based on a purity of 93.5%), trans-(+)-Δ9-THC (**1b**) from Example 11 (17.0 g, 8.36 g based on a purity of 49.2%) and heptane (28 mL) was cooled to 0°C, seeded with (±)-Δ9-THC (0.02 g), and stirred for 5 hours at 0°C. The resultant mixture was cooled to -15°C and stirred for an additional 48 hours at -15°C. The mixture was filtered and the resultant solids washed with cold n-heptane (4 mL). The solids were then dried under reduced pressure at 35°C to provide crude (±)-Δ9-THC. Yield: 11.4 g, 68%. Analysis (HPLC) of the crude (+)-Δ9-THC indicated that it had a purity of 93.6%.

[0178] The crude (±)-Δ9-THC (11.2 g) was dissolved in heptane (15 g) at 50°C, and the mixture was cooled with stirring to 0°C. The resultant mixture was stirred at 0°C for 2 hours, cooled to -15°C, and stirred for an additional 48 hours at - 15°C. The mixture was filtered and the resultant crystalline solids washed with cold n-heptane (4 mL). The solids were then dried under reduced pressure at 35°C to provide crystalline (±)-Δ9-THC. Yield: 9.2 g, 82%. Analysis (HPLC) of the crystalline (±)-Δ9-THC indicated that it had a purity of 97.7 %.

5.19. Example 19: Preparation of Crystalline (+)-Δ9-THC

[0179] (+)-Δ9-THC (2.70g, 2.55 g of trans-(+)-Δ9-THC based on a purity of 94.3%) (obtained from enantioselective chromatography of crystalline (±)-Δ9-THC) as described in Example 21) and trans-(-)-Δ9-THC from Example 9 (3.36 g, 2.76 g of trans-(-)-Δ9-THC based on purity of 82.2%) were dissolved in heptane (9.5 mL). The resultant solution was cooled to 0°C and seeded with crystalline (±)-Δ9-THC (0.01 g). The resultant admixture was stirred for 5 hours at 0°C and for 72 hour at - 15°C. The resultant mixture was filtered and the resultant solids washed with cold heptane (8 mL). The solids were then dried under reduced pressure at 35°C to provide crystalline (±)-Δ9-THC. Yield: 4.4 g, 79.7%. Analysis (HPLC) of the product indicated that it had a purity of 98.7%.

5.20. Example 20: Preparation of Crystalline (+)-Δ9-THC

[0180] Crude trans(-)-Δ9-THC and crude trans-(+)-Δ9-THC were prepared by processes as described in Examples 9 and 11, respectively. Crude trans-(-)-Δ9-THC (27.7 g; containing 10.0 g of trans-(-)-Δ9-THC) in 65 mL of heptane and crude trans-(+)-Δ9-THC (24.3 g; containing 10.0 g of trans-(+)-Δ9-THC), 50% and heptane (315 mL) was admixed with a methanolic caustic solution containing 50% caustic (33 g), water (16.5 mL) and methanol (190 mL) for 20 minutes at 25°C. The resultant purple methanolic caustic (lower) phase was collected, and the organic phase was admixed again with a methanolic caustic solution containing 50% caustic (33 g), water (16.5 mL) and methanol (190 mL) for 20 minutes at 25°C. The resultant methanolic caustic phase was collected, and the combined methanolic caustic phases were treated slowly with a 10% solution of citric acid in water (545 g). The resultant yellow admixture was then extracted with heptane (200 g). The resultant organic phase collected and washed with water (150 mL), dried over Na_2S_{O4}, and filtered. The resultant filtrate was dried by azeotropic distillation and concentrated under reduced pressure. The resultant red oil (41.76 g) was dissolved in heptane (57 g), cooled to 0°C, and seeded with 100 mg of crystalline (±)-Δ9-THC. The resultant admixture was cooled to -15°C and stirred at -15°C for 12 hours. The resultant mixture was suction-filtered, and the solids were washed with cold heptane (3 x 10 mL). The resultant yellow solids were allowed to dry under suction

to provide 12.45 g of crude (±)-Δ^9-THC.

[0181] The crude (+)-Δ^9-THC (12.45 g) was dissolved in heptane (25 mL) at 50°C, and the resultant solution was cooled to -10°C for 2-3 hours. The resultant mixture was suction-filtered and the solids washed 3 times with cold heptane (10, 10, and 20 mL). The solids were then allowed to dry under suction to provide (±)-Δ^9-THC as white crystals. Yield: 8.70 g; 14% yield (based on olivetol); 44% yield based on (-)-Δ^9-THC and (+)-Δ^9-THC. Analysis (HPLC) of the crystalline (±)-Δ^9-THC indicated that it had a purity of 96.45%.

5.21. Example 21: Resolution of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC from (±)-Δ^9-THC

[0182] (±)-Δ^9-THC (2.00 g, 97.7% pure) was eluted by flash chromatography on a Merck column (210 x 50 mm) using Chiralpak® AD™ 20 μm chiral (Daicel, Tokyo, Japan) as the stationary phase (loading capacity 500 mg per injection, UV at 228 nm) and n-heptane:2-propanol (95:5 (v:v)) as the mobile phase at a flow rate of 200 mL/min at 20° to 25°C. The fractions in which only trans-(-)-Δ^9-THC was observed were combined, and the volatiles removed using a rotary evaporator at 35° to 40°C to provide trans-(-)-Δ^9-THC (**1a**). Yield: 0.89 g; 89%. Analysis of the product (HPLC) indicated that it was at least 99.9% pure, *i.e.*, no other cannabinoids were detected.

5.22. Example 22: Resolution of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC from (±)-Δ^9-THC

[0183] Crystalline (±)-Δ^9-THC from Example 15 (3.8 g) was dissolved in 8 ml of heptane:2-propanol (95:5 (v:v)) mixture. The resultant solution was injected into a 2 inch stainless steel "Load and Lock" column (Varian) packed with Chiralpak® AD chiral derivatized silica (Chiral Technologies, Inc. Exton, PA). Elution was carried out under isocratic conditions with a solution of heptane:isopropanol (95:5 (v:v)) at a temperature of about 25°C and at a flow rate of 250 mL of eluent/min. Detection of compounds in the eluent was carried out by UV absorption at 235 nm.

[0184] Trans-(+)-Δ^9-THC eluted first, and the combined trans-(+)-Δ^9-THC eluents were concentrated under reduced pressure to provide 1.5 g of trans-(+)-Δ^9-THC (**1b**) as a reddish-yellow oil.

[0185] Trans-(-)-Δ^9-THC eluted after the trans-(+)-Δ^9-THC, and the combined trans-(-)-Δ^9-THC eluents were concentrated under reduced pressure to provide trans-(-)-Δ^9-THC (**1a**) as a thick viscous reddish-yellow oil. Yield: 1.4 g. Analysis (HPLC) of the trans-(-)-Δ^9-THC product indicated that it had a purity of 99.4%.

5.23. Example 23: Resolution of trans-(-)-Δ^9-THC and trans-(+)-Δ^9-THC from (±)-Δ^9-THC

[0186] Crystalline (±)-Δ^9-THC from Example 13 (about 2.0 g) was dissolved in a about 26 ml of 95:5 heptane:IPA (v:v) mixture to provide a 10 wt.% solution. A portion of the 10% solution (about 5g) was injected into 220 x 50 mm stainless steel column (Merck) packed with Chiralpak® AD 20 μm chiral derivatized silica (Daicel, Tokyo, Japan). Elution was carried out under isocratic conditions with a solution of heptane:2-propanol (95:5 (v:v)) solvent at about 25°C and at a flow rate of 200 mL of eluent/min. Detection of products in the eluent was carried out by UV absorption at 228 nm. The elution of the remaining portions of the 10% solution was carried on about 3 x 5 g samples as described above.

[0187] The fractions containing (+)-Δ^9-THC were combined and concentrated under reduced pressure to provide (+)-Δ^9-THC as reddish-yellow oil. Yield: 1.0 g. Analysis (HPLC) of the oil indicated that it had a purity of 97.0%.

[0188] The fractions containing trans-(-)-Δ^9-THC were combined and concentrated under reduced pressure to provide trans-(-)-Δ^9-THC (**1a**) as a thick viscous reddish-yellow oil. Yield: 1.0 g. Analysis (HPLC) of the product indicated that it had a purity of 99.9%.

[0189] The product was stored in a freezer and protected from light and oxygen.

Claims

1. A method for removing impurities from a composition comprising trans-(-)-Δ^9-tetrahydrocannabinol and trans-(+)-Δ^9-tetrahydrocannabinol prior to crystallization of (±)-Δ^9-tetrahydrocannabinol, the method comprising:

   (a) contacting the trans-(-)-Δ^9-tetrahydrocannabinol and trans-(+)-Δ^9-tetrahydrocannabinol with a first water immiscible organic solvent, a water-miscible alcohol, water, and an alkali metal hydroxide, thus forming a biphasic composition comprising (i) a first organic phase, and (ii) an alcoholic-caustic phase containing the trans-(-)-Δ^9-tetrahydrocannabinol and the trans-(+)-Δ^9-tetrahydrocannabinol;
   (b) separating the alcoholic-caustic phase containing the trans-(-)-Δ^9-tetrahydrocannabinol and the trans-(+)-Δ^9-tetrahydrocannabinol from the first organic phase;
   (c) contacting the alcoholic-caustic phase with an acid to provide an acid-treated alcoholic phase.

2. A method for removing impurities from a composition comprising trans-(-)-Δ^9-tetrahydrocarmabinol or trans-(+)-Δ^9-tetrahydrocannabinol prior to crystallization of (±)-Δ^9-tetrahydrocannabinol, the method comprising:

(a) contacting the trans-(-)-Δ^9-tetrahydrocannabinol or trans-(+)-Δ^9-tetrahydrocannabinol with a first water immiscible organic solvent, a water-miscible alcohol, water, and an alkali metal hydroxide, thus forming a biphasic composition comprising (i) a first organic phase, and (ii) an alcoholic-caustic phase containing the trans-(-)-Δ^9-tetrahydrocannabinol or the trans-(+)-Δ^9-tetrahydrocannabinol;
(b) separating the alcoholic-caustic phase containing the trans-(-)-Δ^9-tetrahydrocannabinol or the trans-(+)-Δ^9-tetrahydrocannabinol from the first organic phase;
(c) contacting the alcoholic-caustic phase with an acid to provide an acid-treated alcoholic phase.

3. The method of claim 1 or 2, wherein the acid is citric acid or acetic acid.

4. The method of any one of claims 1 to 3, wherein the acid is added in an amount sufficient to achieve a pH of from 5 to 9.

5. The method of any one of claims 1 to 4, comprising the additional step of:

(d) contacting the acid-treated alcoholic phase with a second water immiscible organic solvent to form (i) a second organic phase comprising the Δ^9-tetrahydrocannabinol and (ii) an acid-treated alcoholic phase.

6. The method of claim 5, comprising the additional step of:

(e) separating the second organic phase from the acid-treated alcoholic phase.

7. A method for making crystalline (+)-Δ^9-tetrahydrocannabinol comprising:

allowing trans-(-)-Δ^9-tetrahydrocannabinol and trans-(+)-Δ^9-tetrahydrocannabinol to crystallize from a first composition comprising trans-(-)-Δ^9-tetrahydrocannabinol, trans-(+)-Δ^9-tetrahydrocannabinol, and a non-polar organic solvent to provide crystalline (±)-Δ^9-tetrahydrocannabinol,
wherein the first composition was obtained by

(a) performing the method of any one of claims 1 to 6,
(b) separating the Δ^9-tetrahydrocannabinol from the resulting phase containing the Δ^9-tetrahydrocannabinol; and
(c) contacting the Δ^9-tetrahydrocannabinol from step (b) with a non-polar organic solvent and optionally its enantiomer to form the first composition.

8. The method of claim 7, wherein the non-polar organic solvent is a straight-chain or branch-chain (C_4-C_{10})aliphatic hydrocarbon, a (C_4-C_{10})cycloaliphatic hydrocarbon, or any mixture thereof, wherein the straight-chain or branch-chain hydrocarbon is preferably pentane, hexane, heptane, isooctane, or any mixture thereof and is more preferably n-heptane.

9. A method for preparing
a trans-(-)-Δ^9-tetrahydrocannabinol composition, comprising:

allowing a composition comprising (±)-Δ^9-tetrahydrocannabinol and an eluting solvent to separate on a chiral stationary phase to provide a trans-(-)-Δ^9-tetrahydrocannabinol composition, or

a trans-(+)- Δ^9-tetrahydrocannabinol composition, comprising:

allowing a composition comprising (±)-Δ^9-tetrahydrocannabinol and an eluting solvent to separate on a chiral stationary phase to provide a trans-(+)-Δ^9-tetrahydrocannabinol composition,

wherein the (±)-Δ^9-tetrahydrocannabinol is obtained from crystalline (±)-Δ^9-tetrahydrocannabinol, and wherein the crystalline (+)-Δ^9-tetrahydrocannabinol is obtained by the method of claim 7 or 8.

10. The method of any one of claims 7 to 9, wherein the trans-(-)-Δ^9-tetrahydrocannabinol is present in the first composition in an amount from 0.75 to 1.25 molar equivalents, preferably from 0.9 to 1.1 molar equivalents, more

preferably from 0.95 to 1.05 molar equivalents and even more preferably 1 molar equivalent per molar equivalent of trans-(+)-Δ^9-tetrahydrocannabinol.

11. The method of any one of claims 7 to 10, wherein the (±)-Δ^9-tetrahydrocannabinol comprises at least 95% by weight, preferably at least 98% by weight and more preferably at least 99% by weight of trans-(-)-Δ^9-tetrahydrocannabinol and trans-(+)-Δ^9-tetrahydrocannabinol based on the total amount of cannabinoids.

Patentansprüche

1. Ein Verfahren zur Entfernung von Verunreinigungen aus einer Zusammensetzung enthaltend trans-(-)-Δ^9-Tetrahydrocannabinol und trans-(+)-Δ^9-Tetrahydrocannabinol bevor kristallines (±)-Δ^9-Tetrahydrocannabinol hergestellt wird, das Verfahren umfassend:

(a) In-Kontakt-Bringen des trans-(-)-Δ^9-Tetrahydrocannabinols und trans-(+)-Δ^9-Tetrahydrocannabinols mit einem ersten nicht mit Wasser mischbaren organischen Lösungsmittel, einem mit Wasser mischbaren Alkohol, Wasser, und einem Alkalimetallhydroxid, um eine zweiphasige Zusammensetzung herzustellen, welche umfasst (i) eine erste organische Phase und (ii) eine alkoholisch-kaustische Phase enthaltend das trans-(-)-Δ^9-Tetrahydrocannabinol und das trans-(+)-Δ^9-Tetrahydrocannabinol;
(b) Abtrennen der alkoholisch-kaustischen Phase enthaltend das trans-(-)-Δ^9-Tetrahydrocannabinol und das trans-(+)-Δ^9-Tetrahydrocannabinol aus der ersten organischen Phase;
(c) In-Kontakt-Bringen der alkoholisch-kaustischen Phase mit einer Säure, um eine säurebehandelte alkoholische Phase herzustellen.

2. Ein Verfahren zur Entfernung von Verunreinigungen aus einer Zusammensetzung enthaltend trans-(-)-Δ^9-Tetrahydrocannabinol oder trans-(+)-Δ^9-Tetrahydrocannabinol bevor kristallines (±)-Δ^9-Tetrahydrocannabinol hergestellt wird, das Verfahren umfassend:

(a) In-Kontakt-Bringen des trans-(-)-Δ^9-Tetrahydrocannabinols oder trans-(+)-Δ^9-Tetrahydrocannabinols mit einem ersten nicht mit Wasser mischbaren organischen Lösungsmittel, einem mit Wasser mischbaren Alkohol, Wasser, und einem Alkalimetallhydroxid, um eine zweiphasige Zusammensetzung herzustellen, welche umfasst (i) eine erste organische Phase und (ii) eine alkoholisch-kaustische Phase enthaltend das trans-(-)-Δ^9-Tetrahydrocannabinol oder das trans-(+)-Δ^9-Tetrahydrocannabinol;
(b) Abtrennen der alkoholisch-kaustischen Phase enthaltend das trans-(-)-Δ^9-Tetrahydrocannabinol oder das trans-(+)-Δ^9-Tetrahydrocannabinol aus der ersten organischen Phase;
(c) In-Kontakt-Bringen der alkoholisch-kaustischen Phase mit einer Säure, um eine säurebehandelte alkoholische Phase herzustellen.

3. Das Verfahren gemäß Anspruch 1 oder 2, wobei die Säure Zitronensäure oder Essigsäure ist.

4. Das Verfahren gemäß irgendeinem der Ansprüche 1 bis 3, wobei die Säure in einer Menge zugegeben wird, die ausreicht, um einen pH von 5 bis 9 zu erreichen.

5. Das Verfahren gemäß irgendeinem der Ansprüche 1 bis 4, umfassend den zusätzlichen Schritt:

(d) In-Kontakt-Bringen der säurebehandelten alkoholischen Phase mit einem zweiten nicht mit Wasser mischbaren organischen Lösungsmittel, um (i) eine zweite organische Phase, enthaltend das Δ^9-Tetrahydrocannabinol und (ii) eine säurebehandelte alkoholische Phase herzustellen.

6. Das Verfahren gemäß Anspruch 5, umfassend den zusätzlichen Schritt:

(e) Abtrennen der zweiten organischen Phase von der säurebehandelten alkoholischen Phase.

7. Ein Verfahren zur Herstellung von kristallinem (±)-Δ^9-Tetrahydrocannabinol, umfassend:

Kristallisieren von trans-(-)-Δ^9-Tetrahydrocannabinol und trans-(+)-Δ^9-Tetrahydrocannabinol aus einer ersten Zusammensetzung, enthaltend trans-(-)-Δ^9-Tetrahydrocannabinol, trans-(+)-Δ^9-Tetrahydrocannabinol und ein unpolares organisches Lösungsmittel, zur Herstellung von kristallinem (±)-Δ^9-Tetrahydrocannabinol,

wobei die erste Zusammensetzung erhalten wurde durch:

(a) Durchführung des Verfahrens gemäß irgendeinem der Ansprüche 1 bis 6,
(b) Abtrennen des Δ^9-Tetrahydrocannabinols aus der resultierenden Phase, welche das Δ^9-Tetrahydrocannabinol enthält; und
(c) In-Kontakt-Bringen des Δ^9-Tetrahydrocannabinols aus Schritt (b) mit einem unpolaren organischen Lösungsmittel und optional seinem Enantiomer, um die erste Zusammensetzung herzustellen.

8. Das Verfahren gemäß Anspruch 7, wobei das unpolare organische Lösungsmittel ein unverzweigter oder verzweigter aliphatischer (C_4-C_{10})-Kohlenwasserstoff, ein cycloaliphatischer (C_4-C_{10})-Kohlenwasserstoff, oder eine Mischung daraus ist, wobei der unverzweigte oder verzweigte Kohlenwasserstoff vorzugsweise Pentan, Hexan, Heptan, Isooktan, oder eine Mischung daraus ist, und noch bevorzugter n-Heptan ist.

9. Ein Verfahren zur Herstellung
einer trans-(-)-Δ^9-Tetrahydrocannabinol-Zusammensetzung, umfassend:

Auftrennen einer Zusammensetzung enthaltend (±)-Δ^9-Tetrahydrocannabinol und ein Elutionsmittel auf einer chiralen stationären Phase zur Herstellung einer trans-(-)-Δ^9-Tetrahydrocannabinol-Zusammensetzung,
oder

einer trans-(+)-Δ^9-Tetrahydrocannabinol-Zusammensetzung, umfassend:

Auftrennen einer Zusammensetzung enthaltend (±)-Δ^9-Tetrahydrocannabinol und ein Elutionsmittel auf einer chiralen stationären Phase zur Herstellung einer trans-(+)-Δ^9-Tetrahydrocannabinol-Zusammensetzung,

wobei das (±)-Δ^9-Tetrahydrocannabinol aus kristallinem (±)-Δ^9-Tetrahydrocannabinol erhalten wird, und wobei das kristalline (±)-Δ^9-Tetrahydrocannabinol mit dem Verfahren gemäß Anspruch 7 oder 8 hergestellt wird.

10. Das Verfahren gemäß irgendeinem der Ansprüche 7 bis 9, wobei das trans-(-)-Δ^9-Tetrahydrocannabinol in der ersten Zusammensetzung in einer Menge von 0,75 bis 1,25 Mol-Äquivalenten, vorzugsweise von 0,9 bis 1,1 Mol-Äquivalenten, noch bevorzugter von 0,95 bis 1,05 Mol-Äquivalenten und sogar noch bevorzugter von 1 Mol-Äquivalent pro Mol-Äquivalent trans-(+)-Δ^9-Tetrahydrocannabinol vorhanden ist.

11. Das Verfahren gemäß irgendeinem der Ansprüche 7 bis 10, wobei das (±)-Δ^9-Tetrahydrocannabinol mindestens 95 Gew.-%, vorzugsweise mindestens 98 Gew.-%, und noch bevorzugter mindestens 99 Gew.-% trans-(-)-Δ^9-Tetrahydrocannabinol und trans-(+)-Δ^9-Tetrahydrocannabinol, bezogen auf die Gesamtmenge der Cannabinoide, enthält.

Revendications

1. Procédé pour éliminer les impuretés d'une composition comprenant du trans-(-)-Δ^9-tétrahydrocannabinol et du trans-(+)-Δ^9-tétrahydrocannabinol avant cristallisation du (±)-Δ^9-tétrahydrocannabinol, le procédé comprenant :

(a) la mise en contact du trans-(-)-Δ^9-tétrahydrocannabinol et du trans-(+)-Δ^9-tétrahydrocannabinol avec un premier solvant organique non miscible à l'eau, un alcool miscible à l'eau, de l'eau et un hydroxyde de métal alcalin, en formant ainsi une composition biphasique comprenant (i) une première phase organique et (ii) une phase alcooliquecaustique contenant le trans-(-)-Δ^9-tétrahydrocannabinol et le trans-(+)-Δ^9-tétrahydrocannabinol ;
(b) la séparation de la phase alcoolique-caustique contenant le trans-(-)-Δ^9-tétrahydrocannabinol et le trans-(+)-Δ^9-tétrahydrocannabinol de la première phase organique ;
(c) la mise en contact de la phase alcoolique-caustique avec un acide pour fournir une phase alcoolique traitée avec un acide.

2. Procédé pour éliminer les impuretés d'une composition comprenant du trans-(-)-Δ^9-tétrahydrocannabinol et du trans-(+)-Δ^9-tétrahydrocannabinol avant cristallisation du (±)-Δ^9-tétrahydrocannabinol, le procédé comprenant :

(a) la mise en contact du trans-(-)-Δ^9-tétrahydrocannabinol ou du trans-(+)-Δ^9-tétrahydrocannabinol avec un

premier solvant organique non miscible à l'eau, un alcool miscible à l'eau, de l'eau et un hydroxyde de métal alcalin, en formant ainsi une composition biphasique comprenant (i) une première phase organique et (ii) une phase alcooliquecaustique contenant le trans-(-)-Δ^9-tétrahydrocannabinol ou le trans-(+)-Δ^9-tétrahydrocannabinol ;

(b) la séparation de la phase alcoolique-caustique contenant le trans-(-)-Δ^9-tétrahydrocannabinol ou le trans-(+)-Δ^9-tétrahydrocannabinol de la première phase organique ;

(c) la mise en contact de la phase alcoolique-caustique avec un acide pour fournir une phase alcoolique traitée avec un acide.

3. Procédé suivant la revendication 1 ou 2, dans lequel l'acide est l'acide citrique ou l'acide acétique.

4. Procédé suivant l'une quelconque des revendications 1 à 3, dans lequel l'acide est ajouté en une quantité suffisante pour atteindre un pH de 5 à 9.

5. Procédé suivant l'une quelconque des revendications 1 à 4, comprenant l'étape supplémentaire de :

(d) mise en contact de la phase alcoolique traitée avec un acide avec un second solvant organique non miscible à l'eau pour former pour (i) une seconde phase organique comprenant le trans-(-)-Δ^9-tétrahydrocannabinol et (ii) une phase alcoolique traitée avec un acide.

6. Procédé suivant la revendication 1, comprenant l'étape supplémentaire de :

(e) séparation de la seconde phase organique de la phase alcoolique traitée avec un acide.

7. Procédé pour préparer du (±)-Δ^9-tétrahydrocannabinol cristallin, comprenant :

l'étape consistant à laisser le trans-(-)-Δ^9-tétrahydrocannabinol et le trans-(+)-Δ^9-tétrahydrocannabinol cristalliser à partir d'une première composition comprenant du trans-(-)-Δ^9-tétrahydrocannabinol, du trans-(+)-Δ^9-tétrahydrocannabinol et un solvant organique non polaire pour fournir du (±)-Δ^9-tétrahydrocannabinol cristallin, dans lequel la première composition a été obtenu

(a) en mettant en oeuvre le procédé de l'une quelconque des revendications 1 à 6,
(b) en séparant le Δ^9-tétrahydrocannabinol de la phase résultante contenant le Δ^9-tétrahydrocannabinol ; et
(c) en mettant en contact le Δ^9-tétrahydrocannabinol de l'étape (b) avec un solvant organique non polaire et, éventuellement, son énantiomère pour former la première composition.

8. Procédé suivant la revendication 7, dans lequel le solvant organique non polaire est un hydrocarbure aliphatique en C_4 à C_{10} à chaîne droite ou à chaîne ramifiée, un hydrocarbure cycloaliphatique en C_4 à C_{10}, ou n'importe lequel de leurs mélanges, l'hydrocarbure à chaîne droite ou à chaîne ramifiée étant avantageusement le pentane, l'hexane, l'heptane, l'isooctane ou n'importe lequel de leurs mélanges et étant plus avantageusement le n-heptane.

9. Procédé pour préparer :

une composition de trans-(-)-Δ^9-tétrahydrocannabinol, comprenant :

l'étape consistant à laisser une composition comprenant du (±)-Δ^9-tétrahydrocannabinol et un solvant d'élution se séparer sur une phase stationnaire chirale, pour fournir une composition de trans-(-)-Δ^9-tétrahydrocannabinol, ou

une composition de trans-(+)-Δ^9-tétrahydrocannabinol, comprenant :

l'étape consistant à laisser une composition comprenant du (±)-Δ^9-tétrahydrocannabinol et un solvant d'élution se séparer sur une phase stationnaire chirale, pour fournir une composition de trans-(+)-Δ^9-tétrahydrocannabinol,

dans lequel le (±)-Δ^9-tétrahydrocannabinol est obtenu à partir de (±)-Δ^9-tétrahydrocannabinol cristallin, et dans lequel le (±)-Δ^9-tétrahydrocannabinol est obtenu par le procédé de la revendication 7 ou 8.

10. Procédé suivant l'une quelconque des revendications 7 à 9, dans lequel le trans-(-)-Δ^9-tétrahydrocannabinol est présent dans la première composition en une quantité de 0,75 à 1,25 équivalent molaire, avantageusement de 0,9 à 1,1 équivalent molaire, plus avantageusement de 0,95 à 1,05 équivalent molaire et encore plus avantageusement de 1 équivalent molaire, par équivalent molaire de trans-(+)-Δ^9-tétrahydrocannabinol.

11. Procédé suivant l'une quelconque des revendications 7 à 10, dans lequel le ($\pm$)-Δ^9-tétrahydrocannabinol comprend au moins 95 % en poids, avantageusement au moins 98 % en poids, et plus avantageusement au moins 99 % en poids de trans-(-)-Δ^9-tétrahydrocannabinol et de trans-(+)-Δ^9-tétrahydrocannabinol sur la base de la quantité totale de cannabinoïdes.

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ELJÁRÁSOK TRANSZ-(-)-DELTA9-TETRAHIDROKANNABINOL ÉS TRANSZ-(+)-DELTA9-TETRAHIDROKANNABINOL TISZTÍTÁSÁRA

Szabadalmi igénypontok:

1. Eljárás szennyezések eltávolítására transz-(-)-Δ^9-tetrahidrokannabinolt és transz-(+)-Δ^9-tetrahidrokannabinolt tartalmazó kompozícióból (±)-Δ^9-tetrahidrokannabinol kristályosítása előtt, az eljárás során:

(a) a transz-(-)-Δ^9-tetrahidrokannabinolt és transz-(+)-Δ^9-tetrahidrokannabinolt érintkezésbe hozzuk egy első, vízzel nem-elegyedő szerves oldószerrel, vízzel elegyedő alkohollal, vízzel és alkálifém-hidroxiddal, így kétfázisú kompozíciót kapunk, amely tartalmaz (i) egy első szerves fázist, és (ii) egy alkoholos-lúgos fázist, amely tartalmazza a transz-(-)-Δ^9-tetrahidrokannabinolt és a transz-(+)-Δ^9-tetrahidrokannabinolt;

(b) elválasztjuk a transz-(-)-Δ^9-tetrahidrokannabinolt és a transz-(+)-Δ^9-tetrahidrokannabinolt tartalmazó alkoholos-lúgos fázist az első szerves fázistól;

(c) érintkezésbe hozzuk savval az alkoholos-lúgos fázist savval kezelt alkoholos fázis előállítására.

2. Eljárás szennyezések eltávolítására transz-(-)-Δ^9-tetrahidrokannabinolt vagy transz-(+)-Δ^9-tetrahidrokannabinolt tartalmazó kompozícióból (±)-Δ^9-tetrahidrokannabinol kristályosítása előtt, az eljárás során:

(a) a transz-(-)-Δ^9-tetrahidrokannabinolt vagy transz-(+)-Δ^9-tetrahidrokannabinolt érintkezésbe hozzuk egy első, vízzel nem-elegyedő szerves oldószerrel, vízzel elegyedő alkohollal, vízzel és alkálifém-hidroxiddal, így kétfázisú kompozíciót kapunk, amely tartalmaz (i) egy első szerves fázist, és (ii) egy alkoholos-lúgos fázist, amely tartalmazza a transz-(-)-Δ^9-tetrahidrokannabinolt vagy a transz-(+)-Δ^9-tetrahidrokannabinolt;

(b) elválasztjuk a transz-(-)-Δ^9-tetrahidrokannabinolt vagy a transz-(+)-Δ^9-tetrahidrokannabinolt tartalmazó alkoholos-lúgos fázist az első szerves fázistól;

(c) érintkezésbe hozzuk savval az alkohlos-lúgos fázist savval kezelt alkoholos fázis előállítására.

3. Az 1. vagy 2. igénypont szerinti eljárás, amelyben a sav citromsav vagy ecetsav.

4. Az 1-3. igénypontok bármelyike szerinti eljárás, amelyben a savat pH 5 – 9 eléréséhez elegendő mennyiségben adagoljuk.

5. Az 1-4. igénypontok bármelyike szerinti eljárás, amely tartalmazza a következő további lépést:

(d) a savval kezelt alkoholos fázist érintkezésbe hozzuk egy második, vízzel nem-elegyedő szerves oldószerel (i) egy második, a Δ^9-tetrahidrokannabinolt tartalmazó szerves fázis és (ii) és egy savval kezelt alkoholos fázis előállítására.

6. Az 5. igénypont szerinti eljárás, amely tartalmazza a következő további lépést:

(e) elválasztjuk a második szerves fázist a saval kezelt alkoholos fázistól.

7. Eljárás kristályos (±)-Δ^9-tetrahidrokannabinol előállítására, amely során:

hagyunk transz-(-)-Δ^9-tetrahidrokannabinolt és transz-(+)-Δ^9-tetrahidrokannabinolt kristályosodni egy első kompozícióból, amely tartalmaz transz-(-)-Δ^9-tetrahidrokannabinolt, transz-(+)-Δ^9-tetrahidrokannabinolt és nem-poláros szerves oldószert, kristályos (±)-Δ^9-tetrahidrokannabinol előállítására,

ahol az első kompozíciót úgy kapjuk, hogy

(a) elvégezzük az 1-6. igénypontok bármelyike szerinti eljárást,

(b) elválasztjuk a Δ^9-tetrahidrokannabinolt a képződött, a Δ^9-tetrahidrokannabinolt tartalmazó fázistól; és

(c) érintkezésbe hozzuk a (b) lépésben kapott Δ^9-tetrahidrokannabinolt nem-poláros szerves oldószerrel és adott esetben enantiomerjével egy első kompozíció előállítására.

8. A 7. igénypont szerinti eljárás, ahol a nem-poláros szerves oldószer egyenes láncú vagy elágazó láncú (C_4-C_{10})alifás szénhidrogén, (C_4-C_{10})cikloalifás szénhidrogén vagy ezek bármilyen elegye, ahol az egyenes láncú vagy elágazó láncú szénhidrogén előnyösen pentán, hexán, heptán, izooktán vagy ezek bármilyen elegye, és előnyösebben n-heptán.

9. Eljárás

transz-(-)-Δ^9-tetrahidrokannabinol kompozíció előállítására, amely során:

hagyunk egy (±)-Δ^9-tetrahidrokannabinolt és eluáló oldószert tartalmazó kompozíciót szétválni egy királis álló fázison transz-(-)-Δ^9-tetrahidrokannabinol kompozíció előállítására, vagy

transz-(+)-Δ^9-tetrahidrokannabinol-kompozíció előállítására, amely során:

hagyunk egy (±)-Δ^9-tetrahidrokannabinolt és eluáló oldószert tartalmazó kompozíciót szétválni egy királis álló fázison transz-(+)-Δ^9-tetrahidrokannabinol kompozíció előállítására,

ahol a (±)-Δ^9-tetrahidrokannabinolt kristályos (±)-Δ^9-tetrahidrokannabinolból kapjuk, és ahol a kristályos (±)-Δ^9-tetrahidrokannabinolt a 7. vagy 8. igénypont szerinti eljárással kapjuk.

10. A 7- 9. igénypontok bármelyike szerinti eljárás, amelyben a transz-(-)-Δ^9-tetrahidrokannabinol az első kompozícióban 0,75 – 1,25 mólekvivalens, előnyösen 0,9 – 1,1 mólekvivalens, előnyösebben 0,95 – 1,05 mólekvivalens, és még előnyösebben 1 mólekvivalens mennyiségben van jelen a transz-(+)-Δ^9-tetrahidrokannabinol mólekvivalensére vonatkoztatva.

11. A 7-10. igénypontok bármelyike szerinti eljárás, amelyben a (±)-Δ^9-tetrahidrokannabinol legalább 95 tömeg%, előnyösen legalább 98 tömeg%, és előnyösebben legalább 99 tömeg% transz-(-)-Δ^9-tetrahidrokannabinolt és transz-(+)-Δ^9-tetrahidrokannabinolt tartalmaz a kannabinoidok összes mennyiségére vonatkoztatva.