

SOP-CH-002 Revision: 4 Effective Date: August 15, 2022

Drug Enforcement Administration Office of Forensic Sciences



STANDARD OPERATING PROCEDURE

for the

ANALYSIS OF SUSPECTED CANNABIS LIQUIDS AND EXTRACTS



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1.0 Introduction

SOP-CH-002 supplements the Analysis of Drugs Manual (ADM) and outlines procedures for analyzing suspected cannabis liquids and extracts, and reporting Δ^9 -tetrahydrocannabinol (THC), cannabinol (CBN), cannabidiol (CBD), and cannabichromene (CBC), if identified. Reference the ADM for evidence analysis policy.

The analytical scheme requires use of system-wide validated methods, if available, and laboratoryvalidated methods. A Decision Limit (DL) value for total THC present at 1% is established for field laboratory reporting purposes. (NOTE: Total THC = Δ^9 -tetrahydrocannabinolic acid (THCA) + Δ^9 -THC).

2.0 Scope

This procedure:

- A. Identifies THC above or below the DL.
- B. Identifies additional cannabinoids.
- C. Applies to liquid samples.
- D. Applies to suspected cannabis extracts.
- E. Does not apply to creams, edibles, and powders.
 - 1. For analysis of these matrices, follow the Analysis of Drugs Manual (ADM) 2-5 using systemwide or laboratory methods validated for the analysis of THC and other cannabinoids. See Section 4.3 for reporting.

3.0 Analytical Scheme

A. If a negative result is obtained during qualitative testing, the SOP no longer applies and analysis should proceed via the ADM or other SOP if applicable.

3.1 Qualitative Analysis

- A. Macroscopic examination: Conduct macroscopic examination on each unit for the presence or absence of plant material.
- B. Microscopic examination: Conduct microscopic examination on each unit when plant material is present.

NOTE: Effervescence of the calcium carbonate crystal (i.e., cystolith at the base of the hair) in dilute acid may be performed.

- 1. Acceptance criteria: Microscopic observation of cystolithic hairs.
- 2. If no microscopic examination was conducted, enter "N/A" in the *Microscopic Observation* finding in the *Macro/Microscopic Examination of Plant Material* test in LIMS.
- C. Gas chromatography-mass spectrometry (GC-MS):



1. Analyze one working THC positive control solution (Appendix A) prior to each sample sequence using THCSCRN_MS01.

NOTE: The data from the positive control may be used for all exhibits run during a sequence.

- a. Acceptance criteria for THC:IS ratio: THC:IS ratio > 1
- b. Use the macro associated with THCSCRN_MS01 to ensure the peak heights of THC, CBD, and IS as well as the THC:IS ratio are recorded on the data.
- 2. Analyze each selected unit using THCSCRN_MS01.
 - a. Weigh 10-20 mg (or use 1-3 drops for liquids) from each selected unit using an appropriate weighing method.
 - b. Add 5 mL of internal standard solution (ISS) to test tube; vortex/mix for 10-15 seconds.
 - c. Dilute sample solution 1:10 with ISS.
 - d. Filter each sample solution through a cotton plugged pipette or syringe filter into new auto sampler vial.
- 3. Acceptance criteria for THC:IS ratio: THC:IS ratio > *0.1*
- 4. Use the macro associated with THCSCRN_MS01 to ensure the peak heights of THC, CBD, and IS as well as the THC:IS ratio are recorded on the data.
- 5. Document ratio results in the *Remarks* finding of the *GC-MS Analysis* test in LIMS (e.g., "THC ≥1%").
- 6. Evaluate data for the presence of CBD, CBN, and CBC.
- D. Gas chromatography-flame ionization detection (GC): Analyze each selected unit using a validated method.
- E. Perform additional qualitative testing as needed.

4.0 Reporting

4.1 **Positive Results for THC in All Units Tested**

- A. Report "△9-Tetrahydrocannabinol (THC)" on the DEA-113 when:
 - 1. GC: All units are positive for THC

AND

- 2. GC-MS: <u>All</u> units are positive for THC
- B. Add statement (as applicable) in the Remarks section of the DEA-113:
 - 1. "Total delta 9-THC estimated >1%." when all units have a THC:IS ratio > 0.1



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OR

2. "*Total delta 9-THC not determined; pending further analysis upon request.*" when all units have a THC:IS ratio < 0.1

OR

3. *"Total delta 9-THC estimated >1% in X of X units tested."* when some units have a THC:IS ratio > 0.1 and others have a THC:IS ratio <0.1.

4.2 Additional Cannabinoids

A. Report CBN, CBD, and/or CBC if data is acceptable for identification.

NOTE 1: Additional testing is not required.

NOTE 2: The presence of additional cannabinoids may also be reported if data is acceptable for identification.

4.3 Creams, Edibles and Powders

- A. Report delta 9-THC and other cannabinoids if identified in accordance with ADM 2-5.
- B. Add Remark on the DEA-113:
 - 1. "Total delta 9-THC not determined."



Appendix A – THC Positive Control Preparation

- A. Prepare a 0.1 mg/mL working positive control solution of THC.
 - 1. Using Class-A volumetric glassware, prepare a 1-1.5 mg/mL THC Stock A solution by transferring 1 mL from the THC reference material (~10 mg/mL) into a tared flask or beaker.
 - 2. Evaporate to dryness and weigh the amount of residue remaining.
 - 3. Dilute using the appropriate volume of ISS. For example, if 9.6 mg of residue remain, dilute using 9 mL of ISS.
 - 4. Prepare the working THC positive control solution by performing a 1:10 dilution of the THC Stock A solution using ISS.

NOTE: Class-A glassware may be used for the preparation of the working THC solution, but it is not required.

- 5. Both stock and working solutions of the positive control can be used until they no longer meet the acceptance criteria.
- 6. Store all solutions in the refrigerator.



Effective Date/Revision History

Revision No.	Effective Date	Summary of Changes
1	10/1/2019	Original document issued.
2	03/29/2021	 Requirement of macro/microscopic examination added Requirement of identification of additional cannabinoids added Reporting for creams, edibles, and powders added Document reformatted and reorganized
3	08/01/2022	 ADM references updated Re-issued to replace SOP-CH-002 Revision 2. Major changes include: Reorganization to include only information pertaining to the analytical scheme Removal of statements when no THC is present
4	08/15/2022	• Correction of the acceptance criteria in Section 3.1.C.3 from 1 to 0.1



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