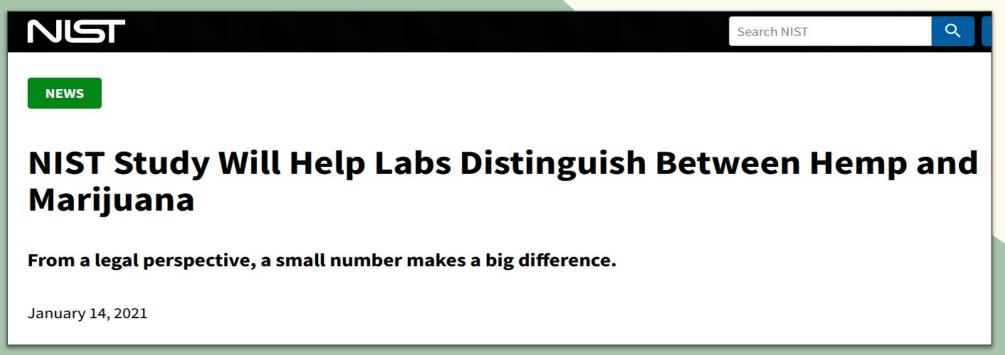


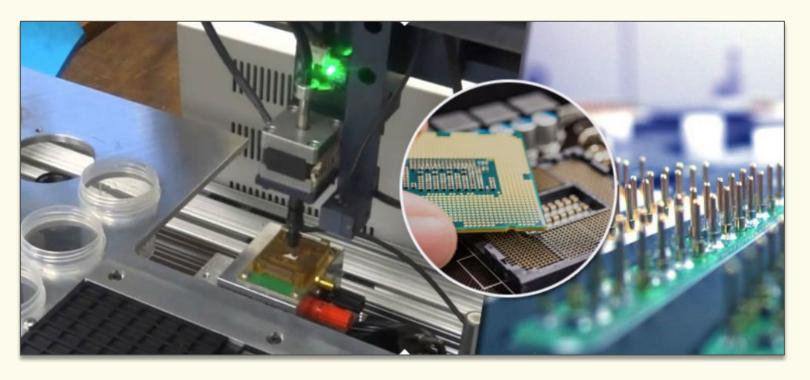


### CURRENT ADVANCES IN FORENSIC SCIENCE RELATED TO CANNABIS



...paving the way for a roadside marijuana breathalyzer







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# OJ: HISTORICAL USES OF \*CANNABIS



# WHAT IS CANNABIS?

#### Cannabis:

o Group of three plants with psychoactive properties:



#### 1. Cannabis sativa

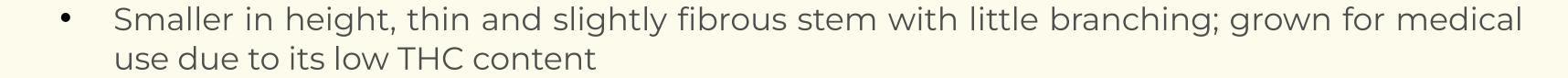
• Long stem and narrow leaves; grown for hemp fiber and psychoactive drug use



#### 2. Cannabis indica

 Shorter stem and broad, deep green leaves; grown for use in reducing nausea/pain/insomnia and increasing bodily relaxation





- O Made up of more than 120 components called "Cannabinoids"
- O Primary psychoactive compound:  $\Delta^9$ -tetrahydrocannabinol (THC)
- O Phytocannabinoid: any cannabinoid produced in the trichomes of a cannabis plant

# HISTORICAL BACKGROUND OF CANNABIS

The cannabis (hemp) plant is one of the most well-known plants across the globe. It is grown in almost every part of the world, with its principle use being a strong fibre for the manufacturing of textiles and rope. In these areas, cannabis was not used as a drug, but rather as a material. Depending on the geographical regions it can grow in, the varying temperatures and conditions modified the pharmacokinetics of cannabis, leading to its first drug-based discovery in the Himalayan region of central Asia, spreading to India, Asia Minor, North Africa, and sub-Saharan Africa.



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#### Map of Cannabis Discovery:

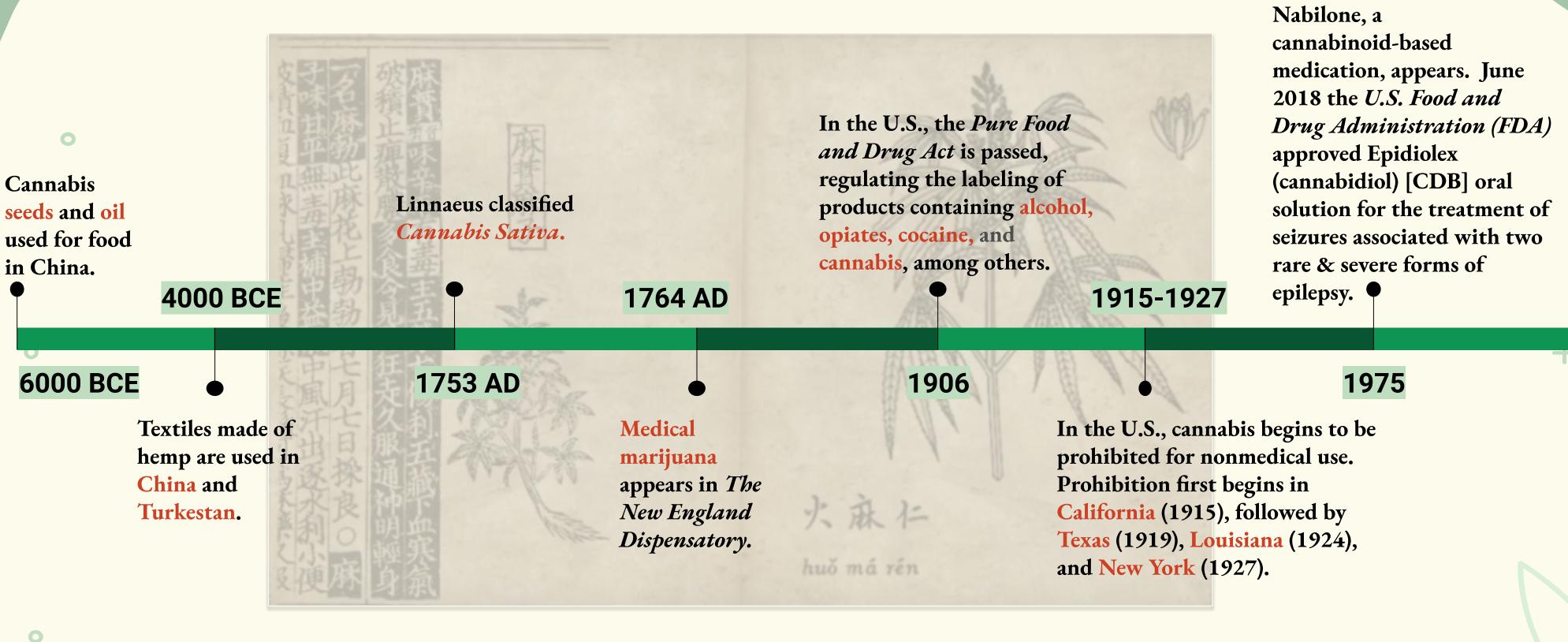
Traces of potent cannabis were identified as early as 2,500 years old in wooden artifacts found buried with people living among the Silk Road in China.

(National Geographic, 2019)

# HISTORICAL BACKGROUND OF CANNABIS

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# HISTORICAL BACKGROUND OF CANNABIS IN CANADA

**1930**s



1979

**1990**s

2018











No reliable accounts of non-medical use of cannabis predating 1930s.

for cannabis possession during the '30s in all of Canada.

#### **HIPPIE-MOVEMENT**

Cannabis introduced into the current culture, youths and adults

1962: 20 cases reported 1963: Over 2300 cases reported 1972: Nearly 12,000

cases reported

Canada: Signed UN's Convention on Psychotropic Substances (1971), which halted any plans to legalize cannabis.

#### **ONTARIO:**

Substantial increase in cannabis use:

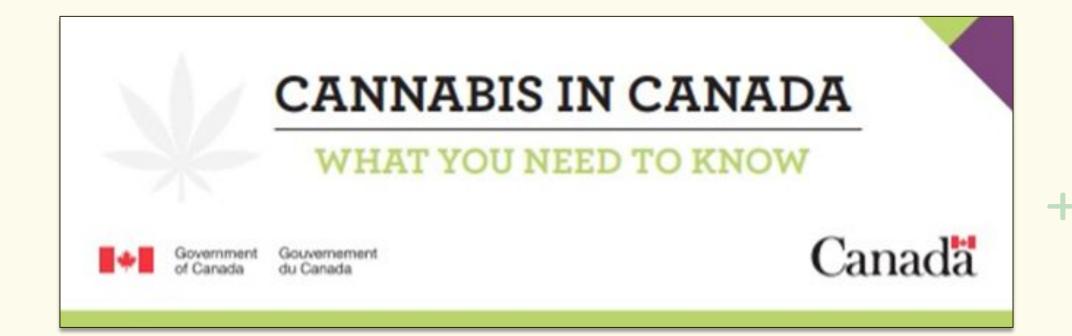
1993-1994: 4.2-7.4% increase
1996-2000: Much

higher in 18-29 age groups (18-28%)

#### LEGALIZATION:

"Cannabis Act", legalizing cannabis growth and use across the nation.

# 02: LEGAL VS. ILLEGAL ACTIVITIES



#### THE CANNABIS ACT

- Instated October 17, 2018 in Canada
- To buy or possess cannabis, **you must be of legal age** (18 or 19 or older), depending on province/territory.
- You can possess up to 30g of legal dried cannabis in public.
- Legal cannabis products have an **excise stamp** on the packaging in different colours for each province/territory.
- It's illegal to travel across the Canadian border with cannabis.
- In late 2019, edible cannabis, extracts, and topicals were legalized.
- It can take **up to two hours** to begin to feel effects of cannabis
- Don't drive high or work impaired. It is a serious criminal offence.
- Store cannabis away from children, youth, and pets.
- Under the Cannabis Act, access to cannabis for medical purposes continues to those authorized.



# LEGALITIES IN "

"

According to (2018), the following are **LEGAL** rules that are tightly regulated for cannabis use in Canada:

- 1. You must be 19 or older to buy, use, possess, and grow recreational cannabis.
- 2. You can **smoke and vape** in the following places: Private residences, many outdoor public places (ex. Sidewalks, parks), designated smoking guest rooms (hotels, motels, inns), residential vehicles and boats, research and testing facilities (for research purposes only), controlled areas such as long-term care homes/hospices/government-funded housing
- 3. You can **possess up to 30 g** (one ounce) of dried cannabis in public at any time.
- 4. You can grow up to 4 plants per residence (not per person).
- 5. Cannabis edibles were legalized on October 17, 2019.
- 6. Extracts and topicals were legalized on October 17, 2019.
- 7. Medical cannabis is legal if purchased through a federally licensed producer, by written order, or over the and delivered by secure mail.

# LEGALITIES IN "THE CANNABIS ACT"

According to *The Cannabis Act* (2018), the following are **ILLEGAL** rules that are tightly regulated for cannabis use in Canada:



#### YOUTH CANNABIS DIVERSION PROGRAM

is designed to provide youth (age 12-18 years) with an online digital experience program that educates them about the risks associated with cannabis use at their stage of development. It also provides youth with information to enhance their understanding of Ontario laws related to cannabis. It promotes confidence and self-awareness for youth to assess the impacts of cannabis use to make positive choices.

- 1. An online education and prevention program for youth between the ages of 12 and 18, called **Youth Cannabis Diversion Program**, is available online. Youth who come into conflict with the law can be provided with this program by law enforcement and courts.
- 2. You **cannot smoke or vape** in the following areas: Indoor common areas (condos, apartments, university/college residences, enclosed public places / workplaces, and non-designated guest rooms (hotels, motels, inns)), schools and places where children gather, hospitals and other facilities, publicly owned spaces (sports fields), while operating a vehicle or boat.
- 3. Edible cannabis must contain zero nicotine or alcohol products.
- 4. Extracts and topicals are prohibited from being attractive to youth or making health/beauty claims.
- 5. Recreational cannabis in an enclosed workplace remains illegal.



# O3: CANNABIS EXTRACTION & PROCESSING

# HOW IS CANNABIS PROCESSED FOR USE?

In Ontario, it is legal to own and cultivate up to 4 cannabis plants in your household (aged 19+) either indoors or in your yard. To start growing a cannabis plant, there is a lot to consider:

- 1. Start with Seeds or Clones
- 2. Growing Conditions: Depends on INDOOR or OUTDOOR growth
- 3. Harvesting and Use



Apricot Kush Seeds (Regular) (OCS, 2021)

#### **Extraction of Cannabis:**

Since **THC** and **CBN** are the main psychoactive cannabinoids in cannabis, high extraction selectivity for these compounds is a key method for the purification and full use of cannabis. Conventional and domestic methods of extraction currently exist, including **distillation**, **solvent extraction**, **Soxhlet**, **maceration**, and **sonication**. Modern extraction techniques are less time-consuming and invasive, and include ultrasound-assisted extraction (**UAE**), supercritical fluid extraction (**SFE**), and pressurized hot water extraction (**PHWE**).

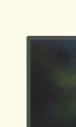


# HOW IS CANNABIS PROCESSED FOR USE?

#### **Methods:**

Soxhlet **Sonication Distillation** Maceration **Solvent Extraction** 







**Ultrasound-Assisted Extraction (UAE) Supercritical Fluid Extraction (SFE) Pressurized Hot Water Extraction (PHWE)** 

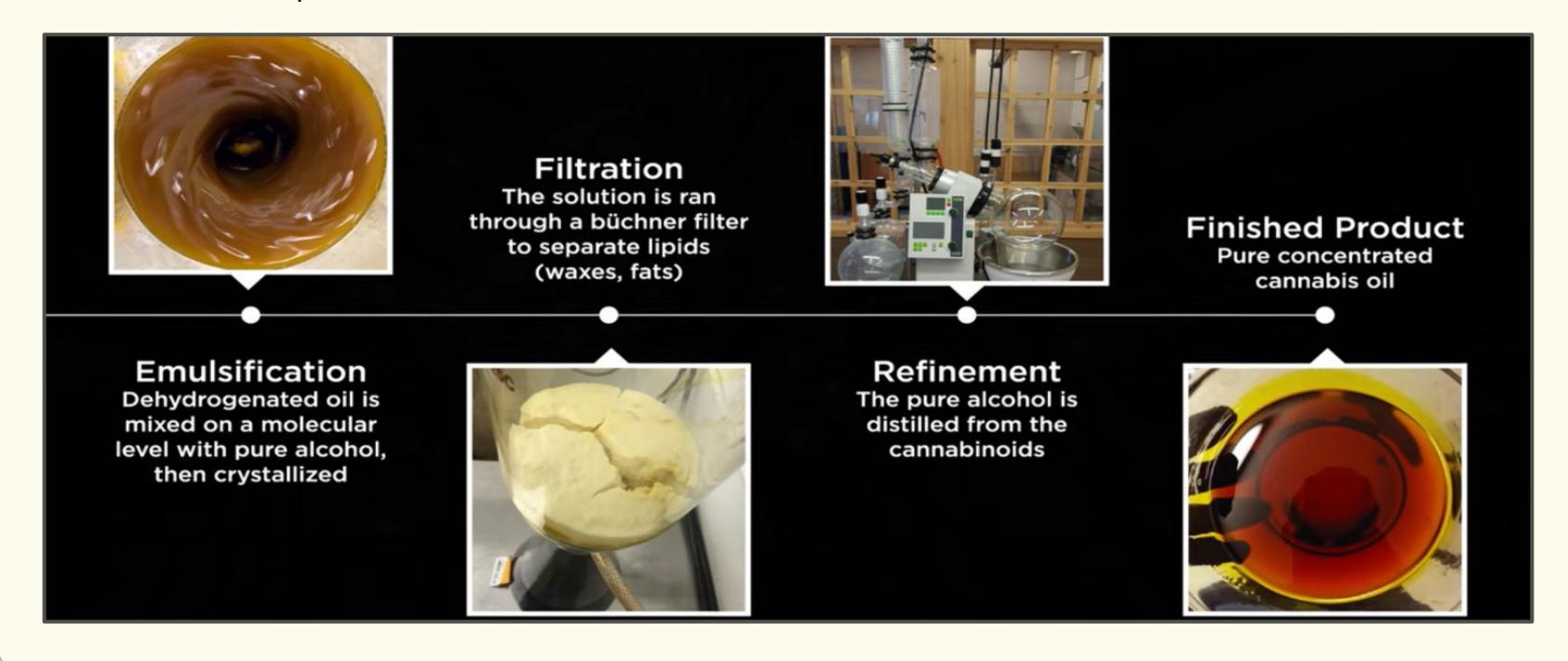
Interested in learning about

Check out this link!



# HOW IS CANNABIS PROCESSED FOR USE?

**SFE** (<u>supercritical fluid extraction</u>): supercritical fluid extraction uses carbon dioxide ( $CO_2$ ) as the solvent. The cannabis sample is placed in the extraction vessel and pressurized using  $CO_2$ , which is cooled down to ensure liquid state of the solvent.

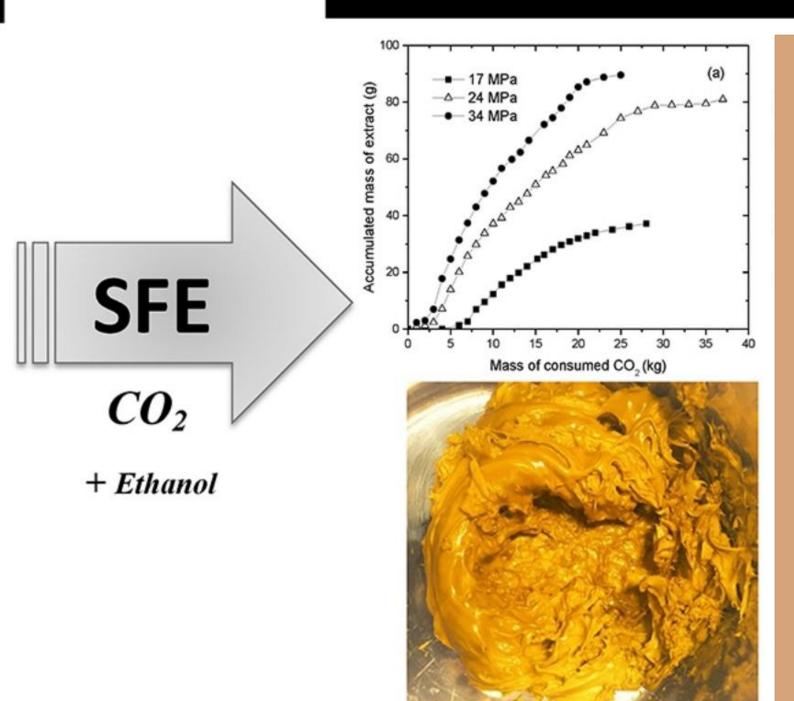


# SFE (Supercritical Fluid Extraction):

#### Cannabis Sativa L.

$$H_3C$$
 $OH$ 
 $CH$ 
 $CH$ 
 $\Delta^9$  - Tetrahydrocannabinol (THC)

#### Cannabinoids extracts



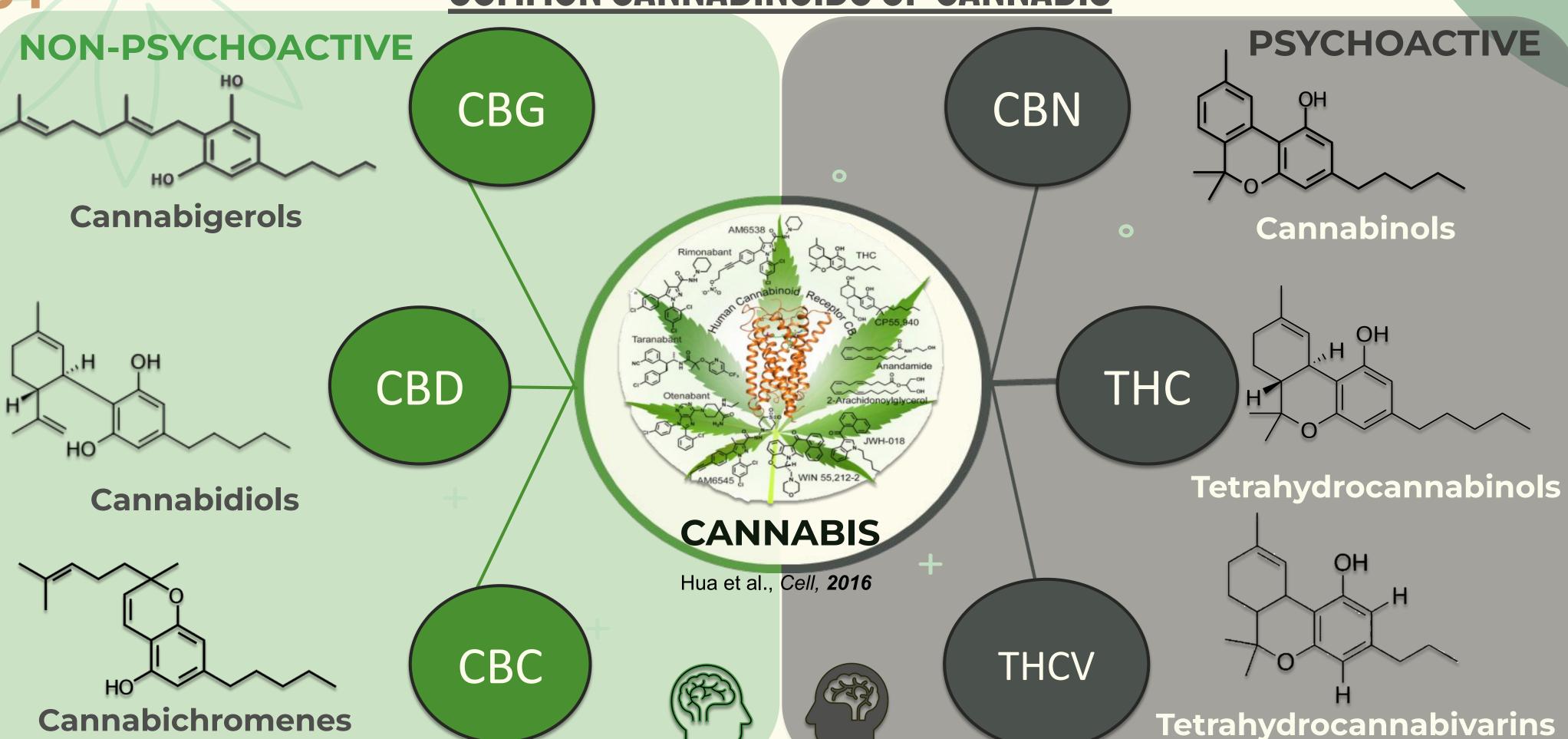
Higher amount of carbon dioxide = HIGHER amount of extract isolated

Higher pressure used = HIGHER amount of extract isolated



# O4: COMMON CANNABINOIDS OF CANNABIS

### COMMON CANNABINOIDS OF CANNABIS





# COMMON CANNABINOIDS OF CANNABIS

# NON-PSYCHOACTIVE



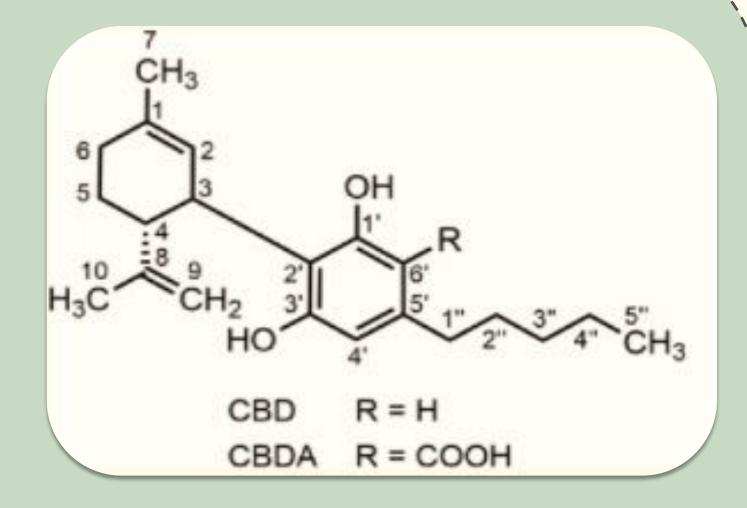
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#### **NON-PSYCHOACTIVE**

- Begins as cannabidiolic acid (CBDA); when heated, decarboxylated into CBD
- Type of phytocannabinoid that lacks detectable psychoactivity and does not bind to
   CB<sub>1</sub> or CB<sub>2</sub> receptors in the body
- Interacts with human Endocannabinoid System (ECS) to encourage release of our own endocannabinoids, rather than binding to CB1 receptors

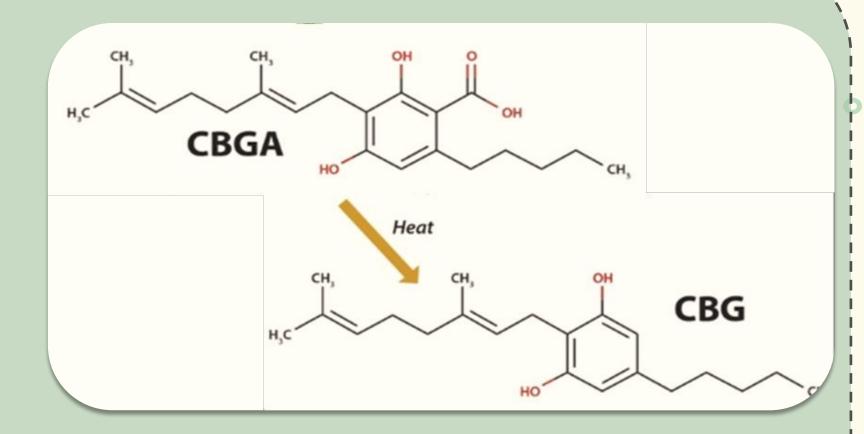


• Affects the activity of ion channels, receptors, and enzymes (e.g.) blocks activity of FAAH (increases anandamide levels), agonist of TRPV1 channel, positive allosteric modulator of glycine receptors, and acts as anti-oxidant and reactive oxygen species (ROS) scavenger

#### **NON-PSYCHOACTIVE**

 Begins as cannabigerolic acid (CBGA); when heated, decarboxylated into CBG

• Type of phytocannabinoid that lacks detectable psychoactivity and only partially binds to the CB<sub>1/2</sub> receptor (agonist)



- In vitro assays have shown at 100 µg/mL (not biologically relevant concentration), CBG can inhibit cyclooxygenase
   (COX) 1 & 2 enzymes with 30% efficiency, can block 5-HT<sub>1A</sub> receptors, and act as an α<sub>2</sub>-adrenoreceptor agonist
- Evidence further suggests partial signs of analgesia by activation of α<sub>2</sub>-adrenoreceptors

#### **NON-PSYCHOACTIVE**

- Second most abundant cannabinoid in some strains of cannabis, and <u>directly synthesized from cannabichromenic acid (CBCA)</u>
- Type of phytocannabinoid that lacks detectable psychoactivity and possesses poor affinity to the CB<sub>1</sub>
   and CB<sub>2</sub> receptors

- Recent study proved CBC acts as a selective CB<sub>2</sub> receptor agonist that signals through the G<sub>i/o</sub> type G proteins and induces CB<sub>2</sub> receptor signalling, indicating this cannabinoid may act as a potential therapeutic agent that influence the non-psychotropic CB<sub>2</sub> pathways of the ECS
- Activates the transient receptor potential ankyrin-type 1 (TRPA1) and inhibits endocannabinoid inactivation both involved in inflammatory responses



# **PSYCHOACTIVE**



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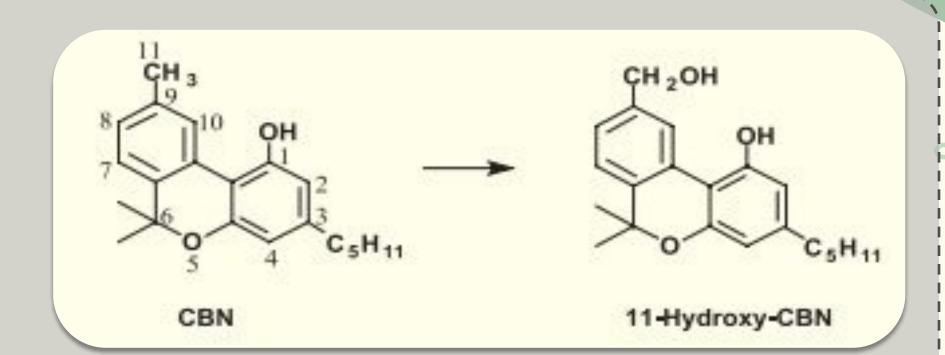
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## COMMON CANNABINOIDS & METABOLITES

#### **PSYCHOACTIVE**

#### Cannabinols (CBNs)

- CBNs are a product of the  $\Delta^9$ -THC oxidation and have 10% of the  $\Delta^9$ -THC activity at the CB $_1$  receptor
- High hydrophobicity allows CBNs to easily transport across the aqueous layer of the skin, with permeation of CBN being 10-fold higher than for  $\Delta^8$ -THC



- CBNs have a high level of distribution in blood plasma (50 L/kg), significantly higher than THC and CBD
- Polyaromatic hydrocarbons found in cannabis induce the expression of CYP1A2, with CBNs inhibiting the activity of <u>CYP1A1, 1A2, 1B1, and 2A6 enzymes</u>
- CBN is metabolized by CYP 2C9 and 3A4, but can also act as a potential substrate for CYP 2C19

## COMMON CANNABINOIDS & METABOLITES

#### **PSYCHOACTIVE**

#### Tetrahydrocannabinols (THCs)

Principal phytocannabinoid that binds to and activates the CB<sub>1</sub>
 and CB<sub>2</sub> receptors

• Derived from the inactive <u>tetrahydrocannabinolic acid (THCA)</u> and is activated once decarboxylated into THC via heating above 120°C (complete decarboxylation after 4 h)

Most commonly studied cannabinoid, as it is responsible for the majority of physical & psychotropic effects of cannabis through the
 CB₁ receptor

Acts as a partial agonist at both CB receptors, and produces CBN products

• Distribution is time-dependent & begins immediately after absorption into the bloodstream (10 L/kg)

## COMMON CANNABINOIDS & METABOLITES

#### **PSYCHOACTIVE**

#### Tetrahydrocannabivarins (THCVs)

Type of minor phytocannabinoid that acts as a CB<sub>1</sub> receptor antagonist and CB<sub>2</sub> receptor partial agonist both in vitro and in vivo, as well as a 5-HT<sub>1A</sub> receptor agonist and exhibits anti-oxidant effects

• At a dose of 10 mg/kg, has been shown to reduce acute nausea in rats, potentially through the CB<sub>1</sub> receptor-independent mechanism

• <u>Several in vitro studies</u> show that THCV binds with high affinity to CB<sub>1</sub> receptors, but it does not appear to be a potent CB<sub>1</sub> receptor agonist

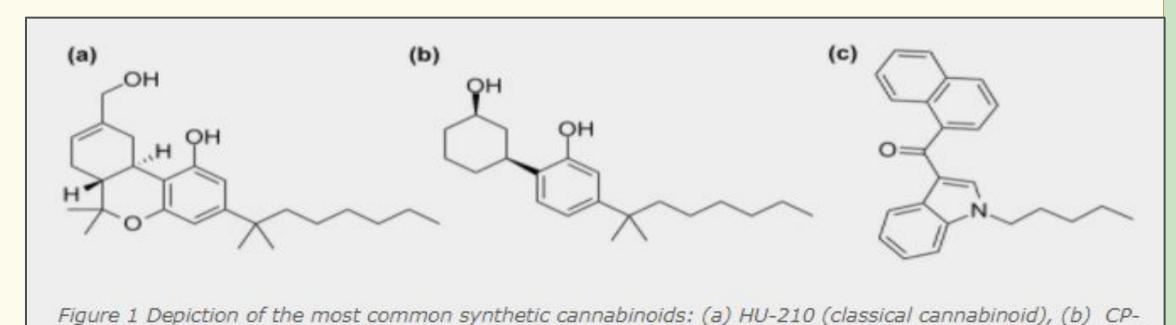
• CB<sub>1</sub> receptor antagonist and potent CB<sub>2</sub> receptor partial agonist at low-mid level concentrations

# **SYNTHETIC CANNABINOIDS**

- Synthetic Cannabinoids: class of man-made molecules that bind to the same receptors as cannabinoids; considered "designer drugs"
- Can be significantly **more toxic** than cannabis, including raising heart rate, increasing vomiting, agitation, confusion, and hallucinations

Classified as "Classical Cannabinoids" or "Non-Classical Cannabinoids"

- 1. Classical: similar behaviour to THC, much more potent than THC (ex. HU-210: 100 times more potent than THC, similar structure to THC)
- Non-Classical: include cyclohexylphenols or 3-arylcyclohexanols (CP compounds), and aminoalkylindoles (naphthoylindoles (JWH-018)) often slightly more potent than THC



47,497 (non-classical cannabinoid), (c) JWH-018 (aminoalkylindoles)

#### CASE STUDY:

On July 12, 2016, a synthetic cannabinoid caused mass intoxication of 33 persons in one New York City neighborhood, in an event described in the popular press as a "zombie" outbreak because of the appearance of the intoxicated persons.



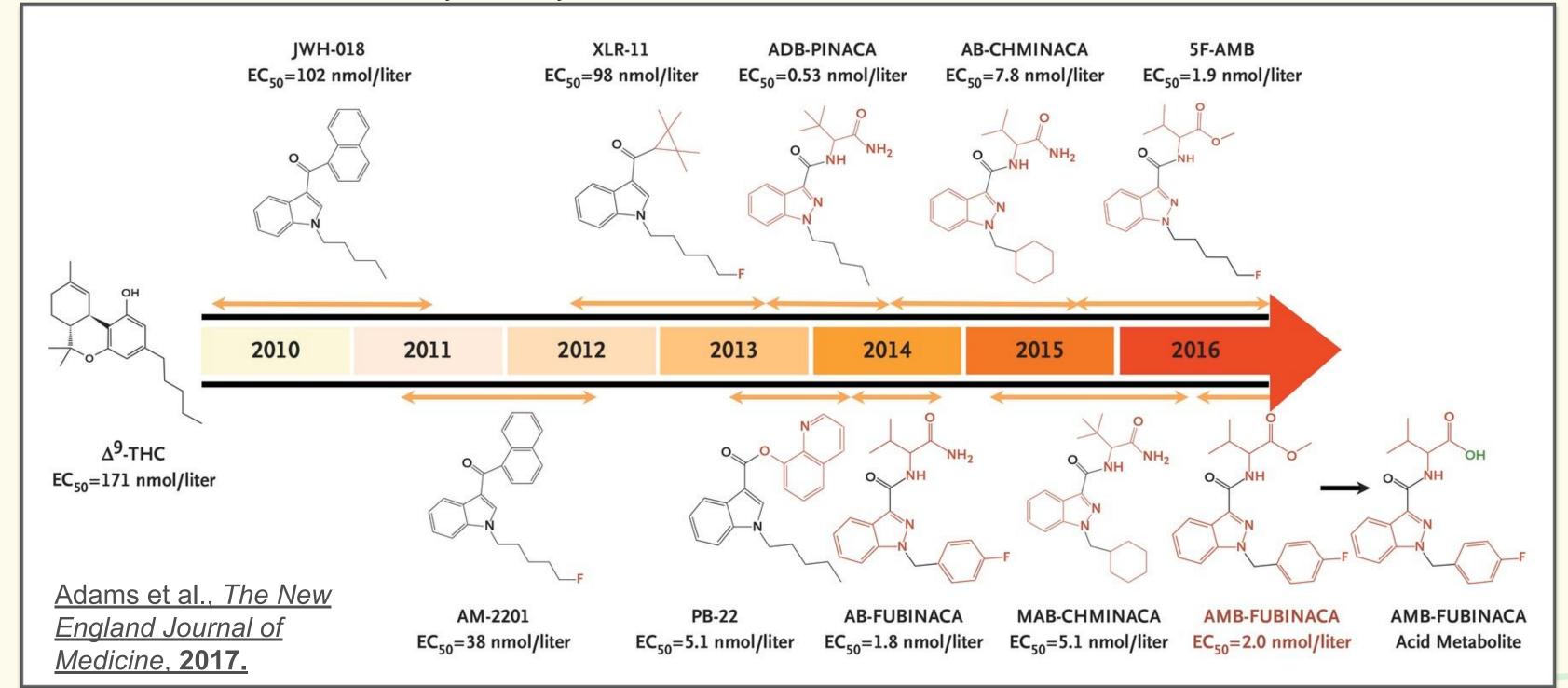
N Engl J Med 2017; 376:235-242 DOI: 10.1056/NEJMoa1610300 AK-47 24 Karat Gold Foil Wrapper Containing Herbal Products Recovered from a Patient Involved in the Outbreak

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# SYNTHETIC CANNABINOIDS

#### **Evolution of Synthetic Cannabinoids:**

"Zombie" outbreak caused by the synthetic cannabinoid AMB-FUBINACA in New York



# SYNTHETIC CANNABINOIDS

- Canada's Addiction Services warning about the uses of Synthetic Cannabinoids
- Age distribution of synthetic cannabinoid and marijuana-related emergency department (ED) visits in
   2010

# Canada's Lower-Risk Cannabis Use Guidelines

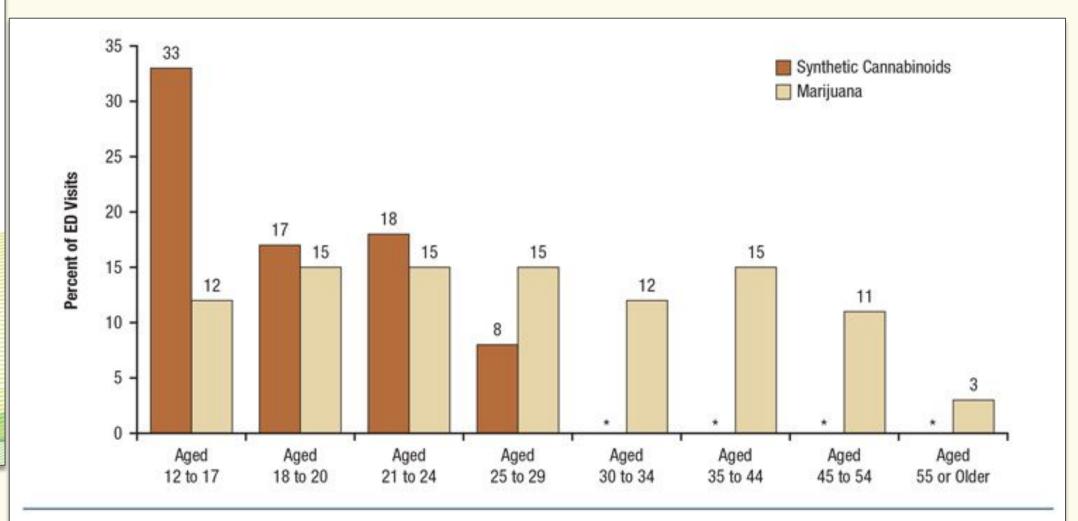


#### 4. Don't use synthetic cannabinoids

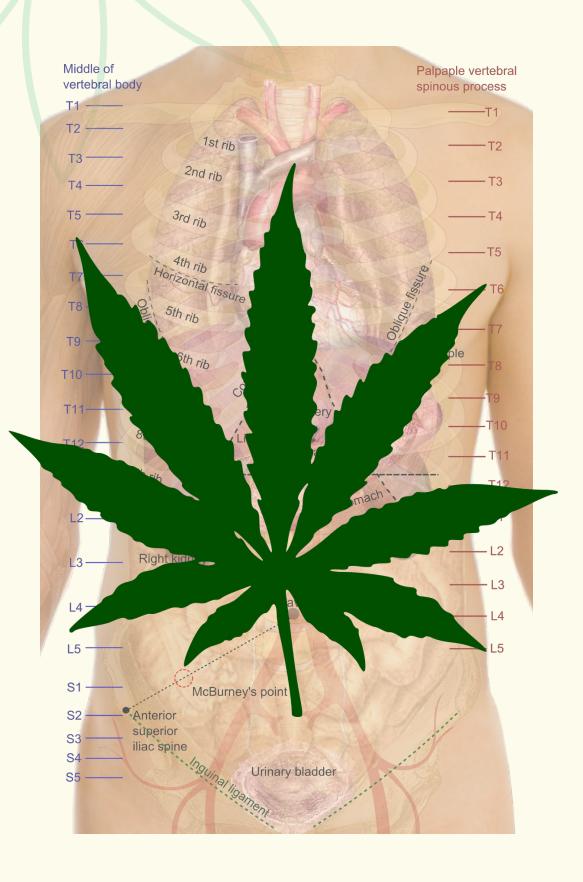
Synthetic cannabinoids (K2, Spice) are relatively new. Synthetics generally have more severe psychoactive impacts and health risks, including cases of death.

Recent reviews on synthetic cannabinoids indicate markedly more acute and severe adverse health effects from the use of these products. The use of these products should be avoided.





<sup>\*</sup> Estimates for ED visits involving synthetic cannabinoids for patients aged 30 or older were suppressed due to low statistical precision. Note: ED visits in which the patient age was unknown are excluded. Source: 2010 SAMHSA Drug Abuse Warning Network (DAWN).



# O5: COMMON ROUTES OF CANNABIS EXPOSURE

# **ROUTES OF ADMINISTRATION**

- Smoking
- Joints, spliffs (cannabis rolled in cigarette paper, can be mixed with tobacco), pipes and bongs, blunts (cigar wrappers filled with cannabis)

Drinking / Eating

Teas, sodas, cannabis oil, baked goods (cookies, brownies, loaves, etc.)

Vaporizing & Vaping

Breathing in dried cannabis or liquid cannabis vapours through a vaporizer or vaping device

Dabbing

Breathing in very hot vapours from heating cannabis concentrates

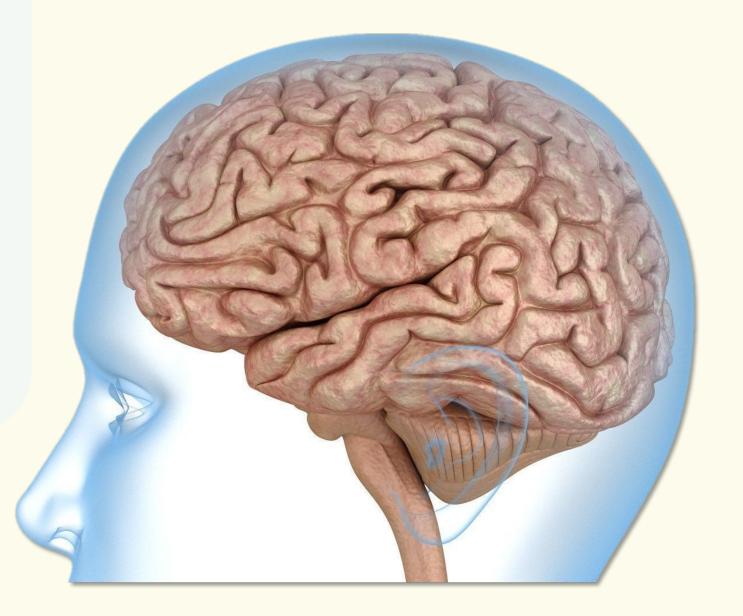
No.	Form	Source	Methods of Use
1.		Dried leaves, stalks, flower, and seeds	Smoked as joint
2.		Fresh leaves and stalk	Mixed with food items (consumed orally)
3.		Leaves, seeds, stem and flower soaked in oil/solvent	Smoked as joint, consumed orally





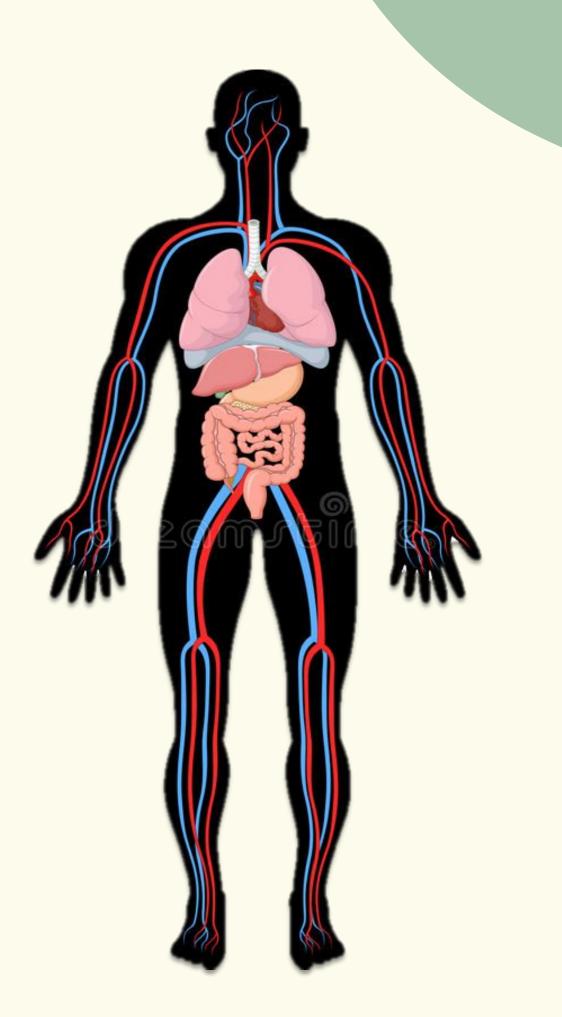
# CANNABIS EFFECTS ON THE BODY

- 1. Impairs ability to drive safely or operate equipment: Causes drowsiness, slower reaction times, lowers ability to pay attention and impairs coordination
- 1. Impairs thinking, concentration, memory and decision-making
- 1. Can cause **anxiety and panic**, as well as possible triggering of psychotic episodes (hallucinating, experiencing paranoia, etc.)



# CANNABIS EFFECTS ON THE BODY

- 1. Hurt your lungs permanently, make it harder to breathe (if smoked)
- 2. Frequent use of THC over a long period increases the risk of **cannabis dependence**, also called: addiction, cannabis use disorder, problematic cannabis use.
- 3. Associated with risk of developing / worsening disorders relating to **depression and** anxiety
- 4. THC products with levels higher than 20% [200 mg/g] or more additionally increases the **risk of mental health** problems over time
- 5. Cannabis poisoning □ consuming too much can cause effects to last up to 12 hr



# CANNABINOID DRUG PROPERTIES

fit like a lock and key in the receptors of the human ECS. These receptors are , and responsible for modulating pain mechanisms, memory, appetite, anti-inflammatory responses.





CB<sub>1</sub> receptors are primarily found in the brain and central nervous system (CNS).

- Brain
- Immune System
- Lungs
- Vascular System
- Muscles

- GI Tract
- Liver
- Pancreas
  - Reproductive Organs
  - Bone Marrow



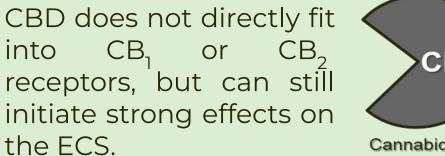
Tetrahydrocannabinol

CB<sub>2</sub> receptors are mostly in the peripheral organs, especially immune-associated cells.

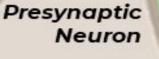


- Immune System
- Skin
- Liver Bone Marrow
- Pancreas
- Spleen
- Bones









Cannabinoid Receptor



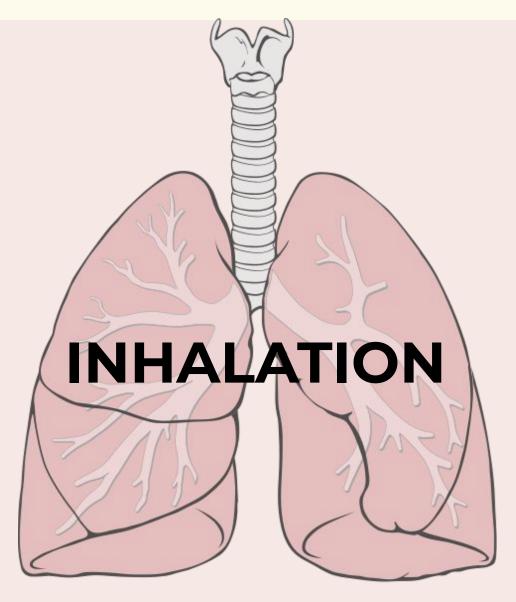
**Veurotransmitters** 

Hua et al., Cell, 2016

# O6: METABOLISM OF CANNABIS



## METABOLISM: ROUTE OF ADMINISTRATION



- Rapidly absorbed into the lungs
- 30% THC destroyed by pyrolysis
- 90 % in blood plasma
- In plasma within seconds of inhalation, peak
   [plasma] = 3-10 min
- [THC] in plasma = 1-4 ng/mL after 3-4 hrs

2



- Systemic absorption **slow**
- Extensive **liver** metabolism
- Rapidly penetrates into fat tissues
- Low [plasma] levels

# METABOLITES OF THC

 $\Delta^9$ -THC \_\_\_\_\_\_ II-OH-THC \_\_\_\_\_ THC-COOH \_\_\_\_\_ Glucuronide

**THC-COOH-Gluc** 

## **DID YOU KNOW?**

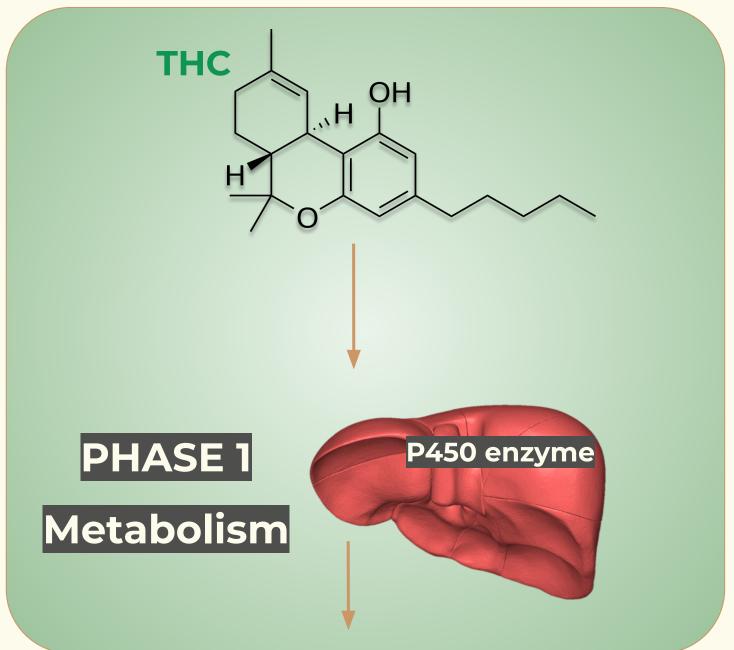
- THC is a **lipophilic** compound
- THC rapidly spreads from blood to fat deposits and organs
- THC lasting up to 30 days after use in chronic users

1.3

# METABOLISM OF THC: PHASE I

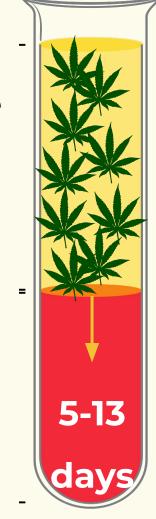
## **INFREQUENT USER**

- Half-life = 1.3 days in
   plasma serum
- Average plasma clearancerate = 36 L/hr



## **FREQUENT USER**

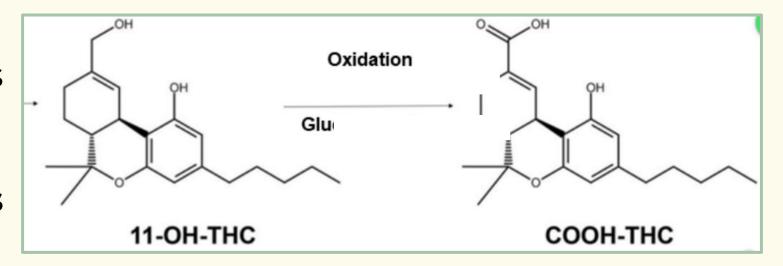
- Half-life = 5-13 days in
   plasma serum
- Average plasma clearancerate = 60 L/hr



Excreted cannabis

(80-90%)

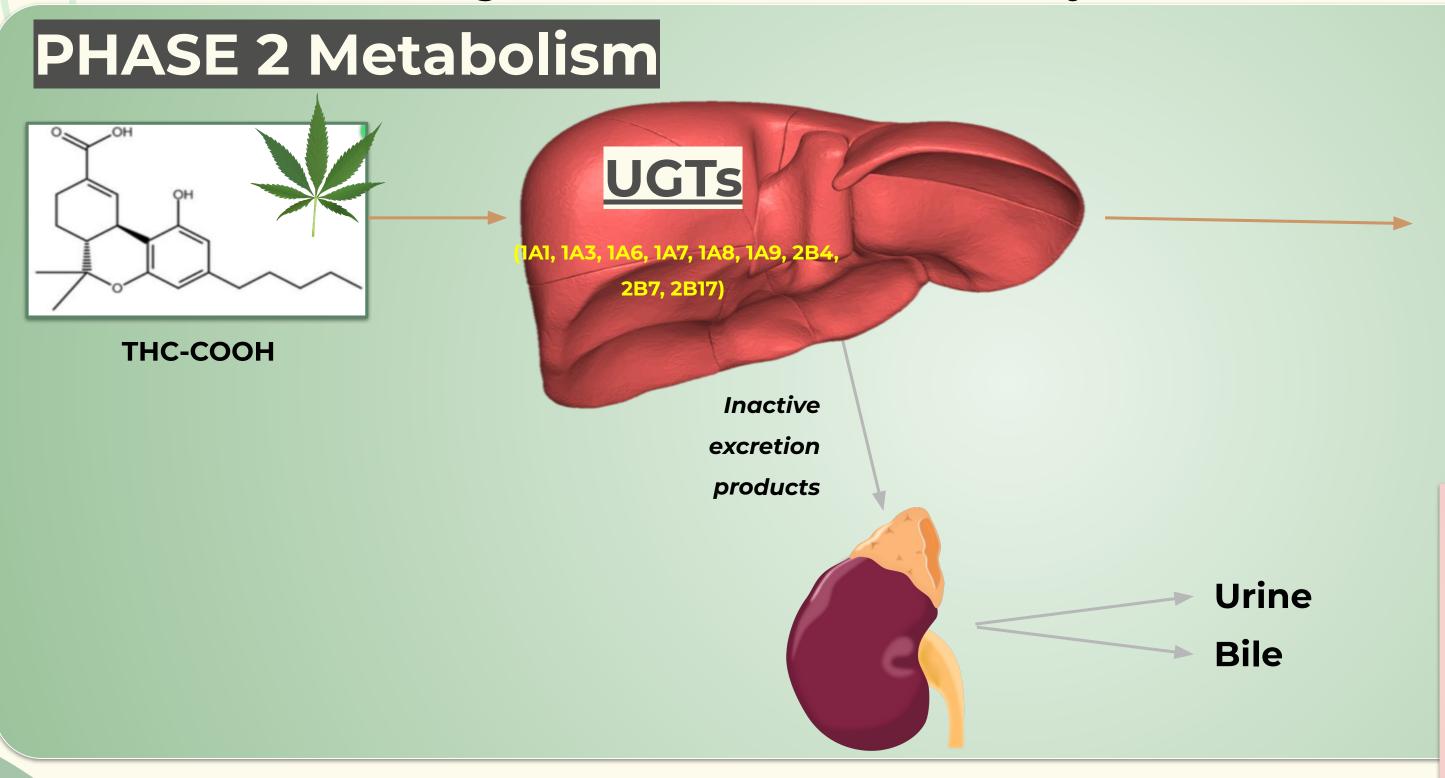
= excreted in **5 days** 

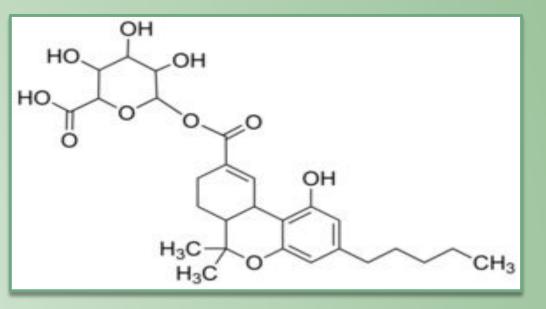


Hydroxylated & carboxylated metabolites

# METABOLISM OF THC: PHASE 2

Glucuronidation of drug enhances its water solubility - increase in elimination rate





THC-COOH-Gluc

Conjugation,
detoxification, & excretion
occurs with catalytic
enzymes during Phase II
Metabolism.

## Did you know?

18 gluc-metabolites exist in urine

06

# METABOLISM OF CBD

## Large portion of CBD is

excreted intact

Ujváry I, Hanuš L. *Cannabis Cannabinoid Res.* 2016;1(1)

## HIGH

СООН

OH CH<sub>2</sub> H<sub>3</sub>C

7-COOH

Hydroxylation products (Phase 1)

# Relative Amounts LOW COOH OH OH OH Hack CHa

Glucuronide product (Phase 2)

CBD-glucuronide

- Oral Administration: 1.09 and 1.97 hr
- Oromucosal Spray Administration: 1.44 10.86 hr
- Inhalation: 1.1 and 2.4 hr
- **I.V. Infusion**: 24 hr
- Chronic Oral Intake: 2-5 days
- **Smoking:** 30 hr

## **Volume of Distribution (CBD)**

4"-OH-7-COOH

- I.V. = 3,000 L
- Oromucosal spray = 30,000 L

## **Bioavailability (CBD)**

- Oral: ~20%
- Inhaled: ~30%

# O7: CANNABIS IN BIOLOGICAL FLUIDS



# BIOLOGICAL FLUIDS CANNABIS IS PRESENT IN: BLOOD



Cannabinoids do **not** distribute well into **erythrocytes**, thus increasing cannabinoid plasma concentration to twice the blood cannabinoid concentration. Most THC and CBD cannabinoids have been detected in the blood plasma!



When THC and alcohol were both consumed, alcohol concentrations **could not** be accurately measured.

## **DID YOU KNOW?**

THC-glucuronide, CBD, and CBN may be present up to 4 hr!



**Blood Alcohol** 

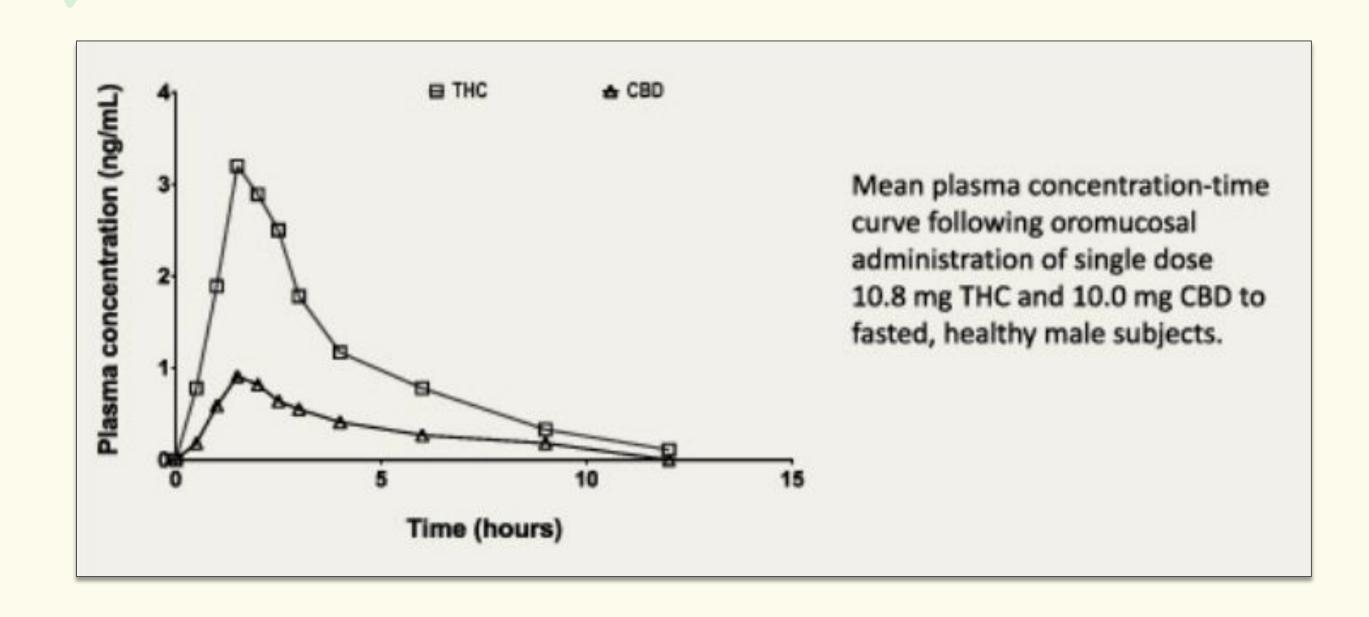
0.08%

## = Same Impairment Level

Alcohol is absorbed mostly from the small intestine, resulting in the highest level of ethanol concentration in the blood which was found to be **delayed** due to the cannabis.



## CANNABIS FOUND IN PLASMA: CASE STUDY



THC and CBD are found in blood plasma

Greater concentration of THC than CBD

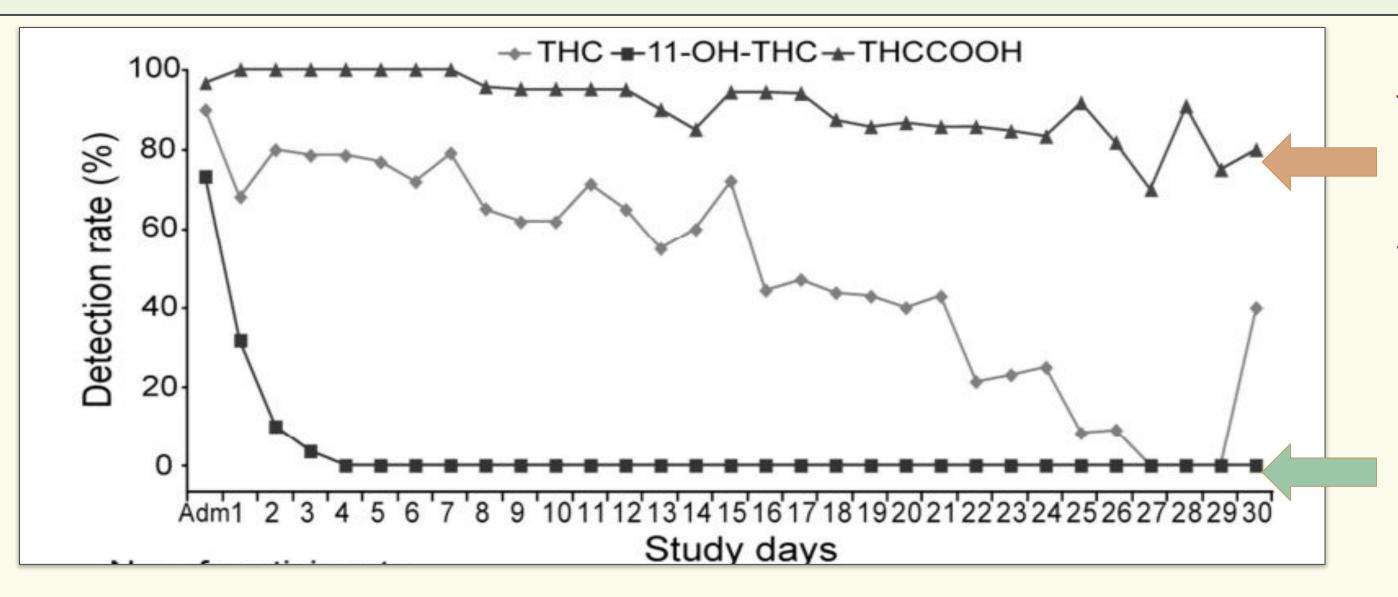
Both compounds peak at ~ 2 min

Both compounds eliminated by 12 hours

# CANNABIS FOUND IN BLOOD: CASE STUDY

Study of frequent users of cannabis smokers.

Whole blood detection rates for THC and its metabolites (**Phase 1**) were analyzed over 30 days of abstinence.



THC-COOH metabolite persists over time

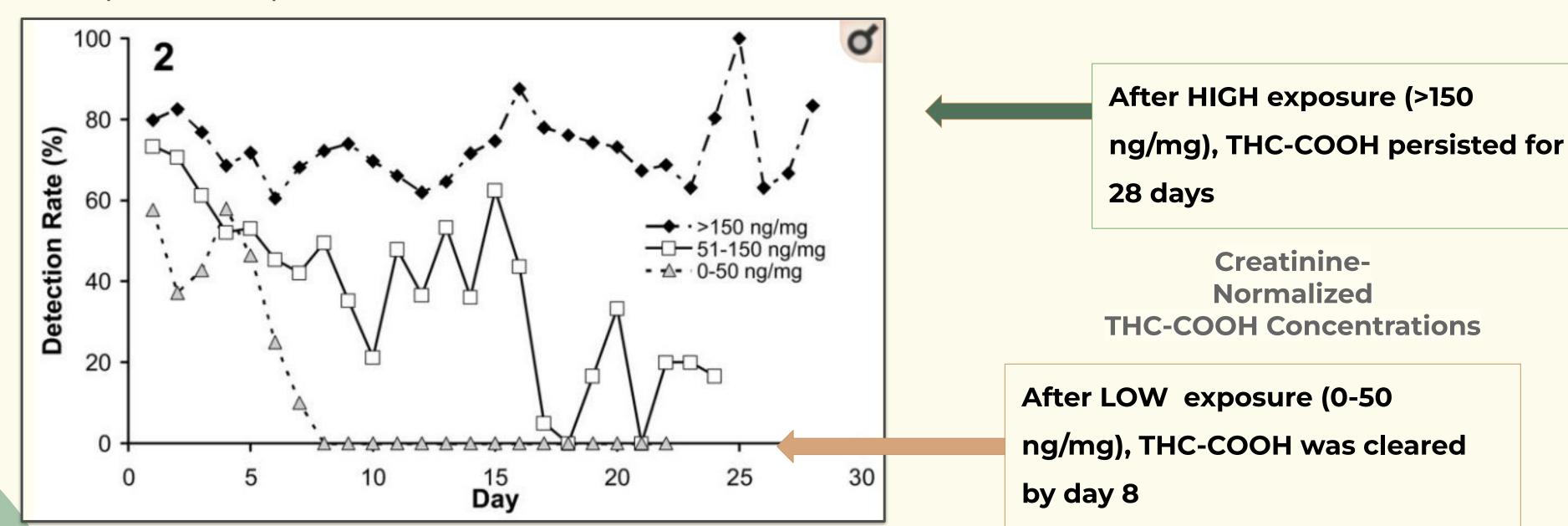
110H-THC metabolite eliminated within 4 days

# BIOLOGICAL FLUIDS CANNABIS IS PRESENT IN: URINE

<u>Urine Markers</u>

## THC-COOH-glucuronide

- Enzymatic hydrolysis step to liberate THC-COOH is analyzed *via* immunoassays, GC-MS, LC-MS/MS, HR-MS
- Characterization of the time course of THC-COOH elimination in urine following variable cannabis exposure to provide data for improved interpretation of urine cannabinoid tests



07

## OTHER BIOLOGICAL FLUIDS WITH CANNABIS

### Oral Fluid (OF):

- Non-invasive, applicable for workplace and drug treatment plans
- 2 µg/L THC in OF considered to be a marker of cannabis intake with the last 24 hr
- [CBD] positive in OF at 0.3  $\mu$ g/L for up to 20 hr in both frequent and infrequent users

#### **Hair Markers:**

- Non-invasive, long detection times for drug intake (basic drugs transfer into hair and remain for years)
- Cannabinoids are acidic, so do not transfer to hair well □ still quantifiable in pg/mg (THC, 11-OH-THC, THC-COOH, CBN, and CBD)

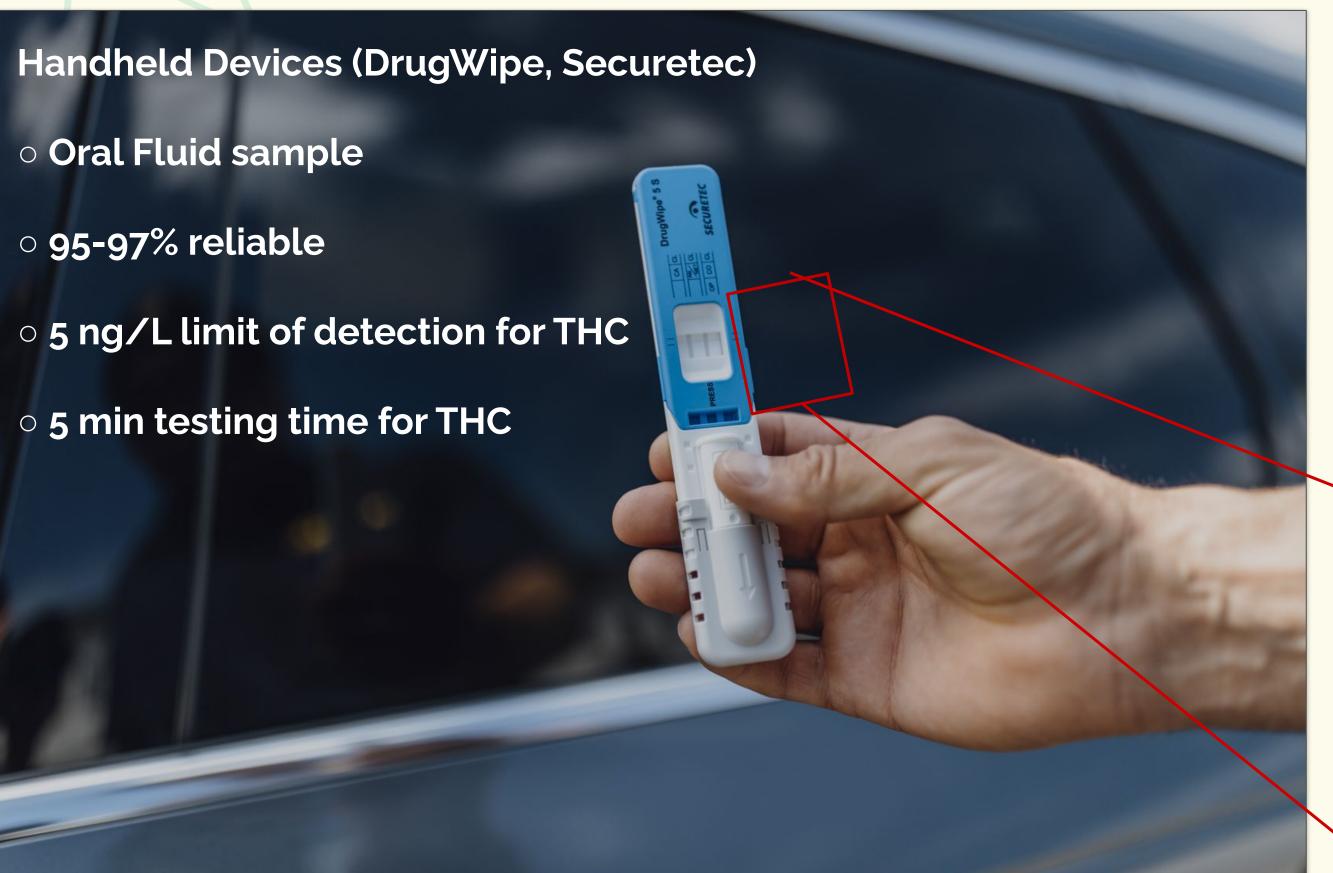
#### **Sweat Markers:**

- Sweat patches worn for one week, monitor drug intake from 3 days before application until removal
- Good method for evaluating THC levels within a week

#### **Breath Markers:**

- THC can be collected via roadside test (breathalyzer) and transported to lab for LC-MS/MS from a filter
- THC ≥50 pg/filter was detected up to 4 hr after smoking (shorter for infrequent users)
- No THC-COOH has been detected in breath □ need to develop selective on-site instruments for metabolites

# ON-SITE CANNABIS TESTING

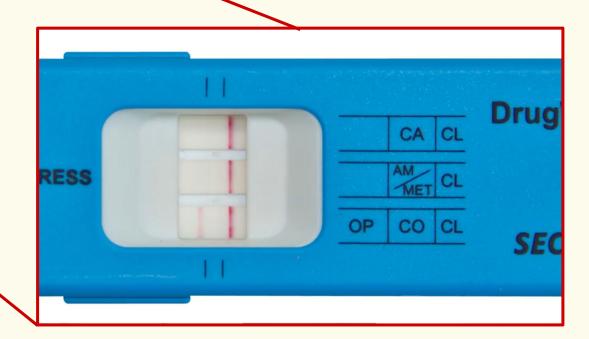


## Saliva drug screener

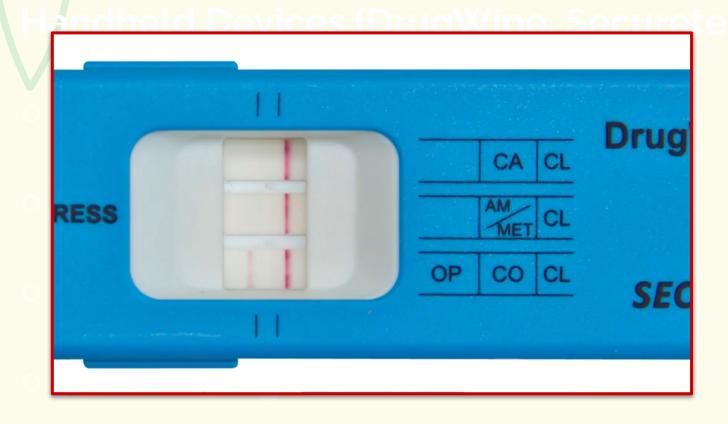
A LESS invasive method of sample collecting compared to blood and urine tests.

The LOWEST sampling volume required.

Potential application and uses by law enforcement, border control, rehabilitation clinics and prison systems worldwide.

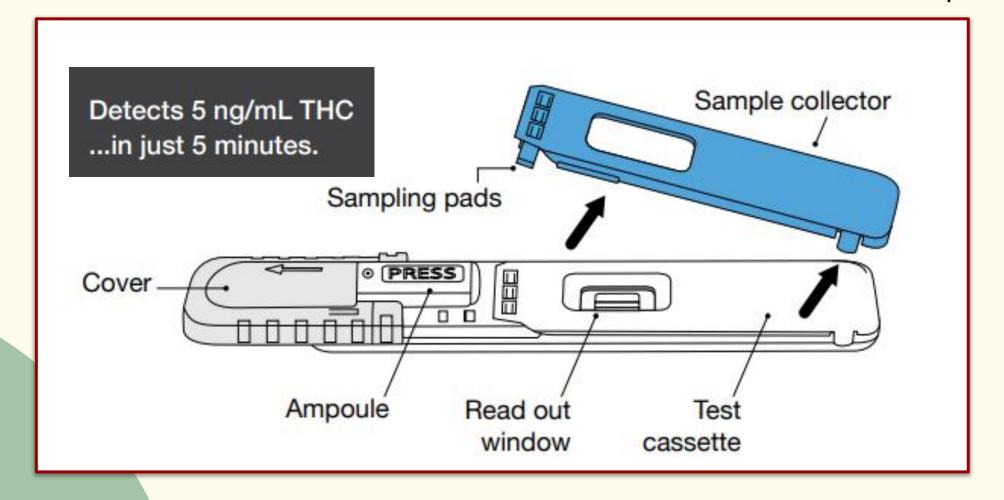


# ON-SITE CANNABIS TESTING



### How does it work?

- Collect sample
- Break the ampoule and release the liquid
- Liquid moves drug particles that are bound to the antibodies on the test strip migrate to the test line
- A sequence of red lines appears on the readout window, depending on the results.



## What drugs does it detect?

**Cannabis**, amphetamine, methamphetamines, cocaine, opiates, ketamine and benzodiazepines.



Let's consider a forensic case based on cannabis!

08: FROM THE CRIME SCENE TO THE LAB: CANNABIS CASE FILES



# **CAN-LII CASE TIMELINE FROM ARREST TO DRE:**

**R. v. D.A.H.** 

June 12, 2015

11:30 AM

12:03 PM

12:05 PM

2:00 PM

4:15 PM

D.A.H. struck a tree

Hit the tree on the left side of the vehicle.
Airbags deployed, no one else hurt.

Officer dispatched

Constable Thomas
Mondoux
dispatched, arrests
D.A.H. for driving
under the influence.

Officer makes arrest at scene of the crime

Constable Thomas
Mondoux issued a
Drug Recognition
Evaluation (DRE) on
grounds that the
defendant's ability
to operate a motor
vehicle was
impaired by drug.

Constable Lisa Grison conducts DRE

DRE completed at 2:00pm. Notably, physiological effects observed (cannabis), matching the observation from Constable Mondoux. Biological Sample Collection

Urine sample was collected at the hospital from the accused. Sample was sent to the CFS for toxicological analysis.

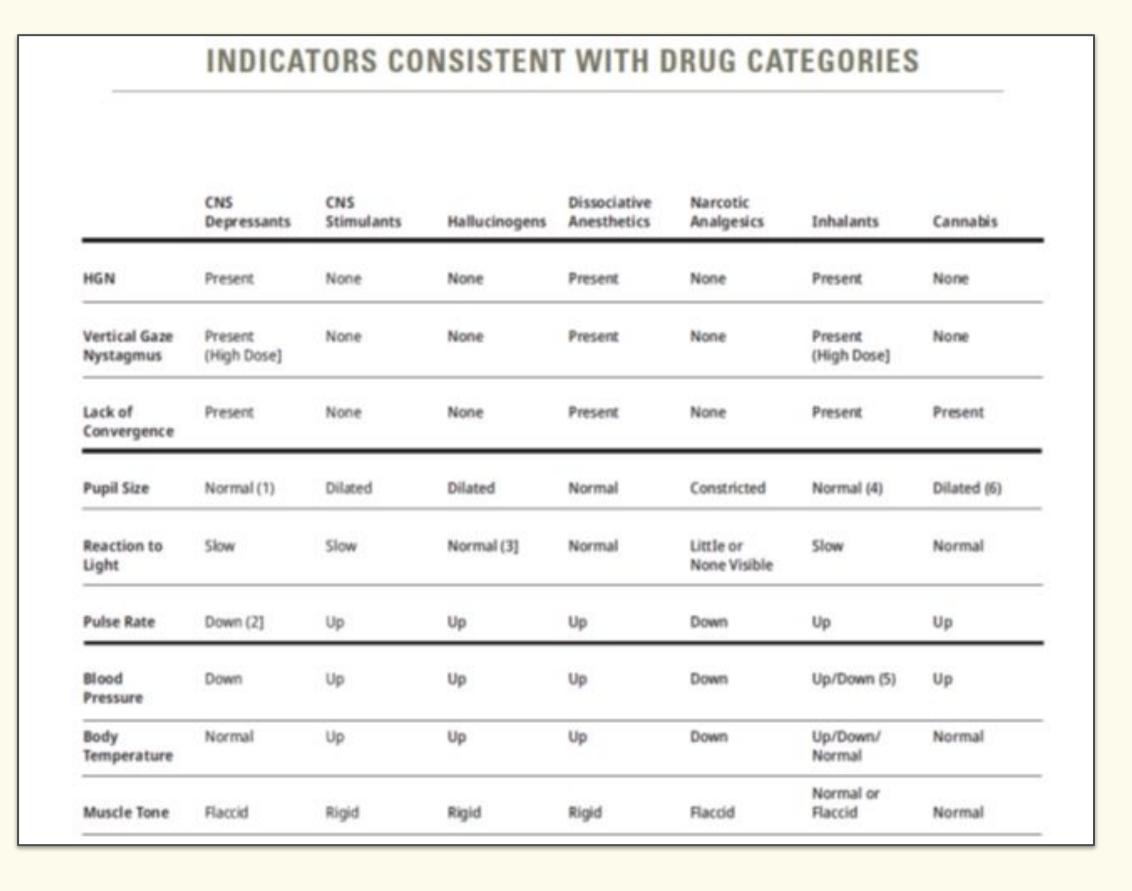
# WHAT ARE DRE ROLLING LOGS?

- A \_\_is a police officer who is trained and certified by the International Association of Chiefs of Police (IACP) to correctly identify drug-impaired persons.
- In Canada, all DRE officers are trained by the Royal Canadian Mounted Police (RCMP).
- DREs are trained on a 12-step protocol called the Drug Influence Evaluation to determine:
  - [1] If the driver is actually impaired and,
  - [2] The classification of the drug causing the incident.



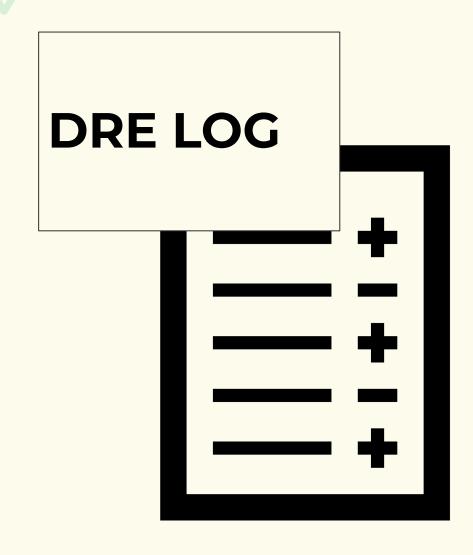
# DRE ROLLING LOGS

- 1. Breath Alcohol Test
- 2. Interview of the Arresting Officer
- 3. Preliminary Examination and First Pulse
- 4. Eye Examinations
- 5. Divided Attention Psychophysical Tests
- 6. Vital Signs and Second Pulse
- 7. Dark Room Examinations
- 8. Examination for Muscle Tone
- 9. Check for Injection Sites and Third Pulse
- 10. Subject's Statements and OtherObservations
- 11. Analysis and Opinions of the Evaluator
- 12. After completing the evaluation, the DRE normally requests a urine, blood marker, and/or OF sample from the subject for a toxicology lab analysis



## CLICK LINK FOR DRE EXAMPLES

## **CAN-LII CASE TOXICOLOGICAL ANALYSIS:**



2:00 pm DRE log completed

4:15 pm Urine sample collected



Urine sample sent to



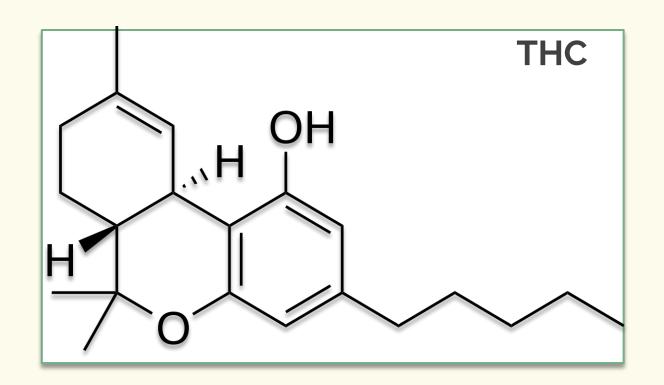
Centre of Forensic Sciences

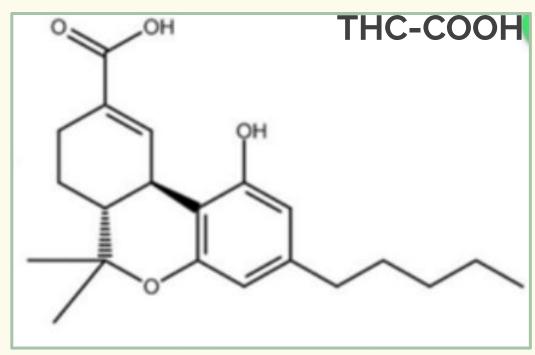
## **CAN-LII CASE TOXICOLOGICAL ANALYSIS:**

## Dr. Daryl Mayers:

- -testified as an expert witness for this cannabis-specific case.
- -Is an experienced forensic toxicologist employed by the Centre of Forensic Sciences (CFS) since 1992.
- -holds a doctorate in forensic toxicology, as well as other degrees

- At 4:15 PM, urine sample was collected from suspect by Cst.
   Grison
- 2. Urine sample sent to CFS
- 3. Dr. Mayers analyzed urine and found **THC** and **Carboxy-THC** (inactive metabolite, principal psychoactive ingredient in cannabis).





# **CAN-LII CASE TOXICOLOGICAL ANALYSIS:**

## Dr. Daryl Mayers testimony:

- 1. Since the metabolite may be found in the urine weeks after ingestion of cannabis, and THC was found mixed in, it was noted that the suspect probably ingested cannabis within the **last several hours**.
- 2. While THC acts immediately on the body, the **metabolite** demonstrates a much longer time course due to **slow release** and **metabolism** in tissues.
- 3. Dr. Mayers stated THC can affect one's mental ability, including the ability to operate a vehicle, 3-4 hours after ingestion.
- 4. He also stated it is difficult to be definitive about its effects without THC concentrations from blood.
- 5. His final remark: it is **unlikely** that a person would be impaired by cannabis **after the 4-hour window,** even though urine detection is still possible.

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# LIST OF REFERENCES

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