


ASAM REVIEW COURSE 2024

Cannabis Use Disorder: Science, Trends, and Clinical Implications

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Learning Objectives


- Increase knowledge about current epidemiological trends in cannabis use in the United States.
- Name the different formulations of cannabis that impact individuals today.
- Review medications that have an evidence base for treating cannabis withdrawal and cannabis use disorder.



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Presentation Outline

- Epidemiology
- Cannabis formulations
- Effects of Cannabis
- Cannabis Potency (THC concentration)
- Special Populations and Cannabis
- Cannabis Use Disorder Treatment
- Medicinal Uses of Cannabis/Cannabinoids
- Board Review Questions / Wrap Up



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Epidemiology




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Cannabis Use/Misuse

- In 2021, an estimated 65.2 million Americans- **27.1%** of the population aged 12 years or older had used cannabis in the preceding month.
- In 2015, it was 22.2 million (**8.9%**) Americans aged 12 years and older.
- Cannabis use peaks in the late teens to early 20s, then declines



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Increased Risk for Use Disorder

- 9% of users develop Cannabis Use Disorder
- The risk increases to 17% in people who start using in adolescence.
- The risk increases to 25 to 50% in people who are daily users (most of whom started using marijuana early in adolescence).


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Cannabis Basics


- The cannabis plant has 104 cannabinoids; only 2 (THC and CBD) have been extensively studied for potential therapeutic applications.
- THC is the most psychoactive component – (*inhaled, ingested*)
- CBD is postulated to have other mechanisms of action (anti-inflammatory, analgesic, etc.).

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Cannabis Plant




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Natural, Plant - Derived Cannabinoids

- Cannabis
- Sativa, Indica, or Hybrid
- Subspecies of the hemp plant

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Natural, Plant - Derived Cannabinoids

Most common preparations:

- *Marijuana
- *Hashish
- *Hash Oil

THC Concentrations vary—

For example, extraction of THC with butane ("dabs") can contain up to 90% THC.

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Synthetic Cannabinoids

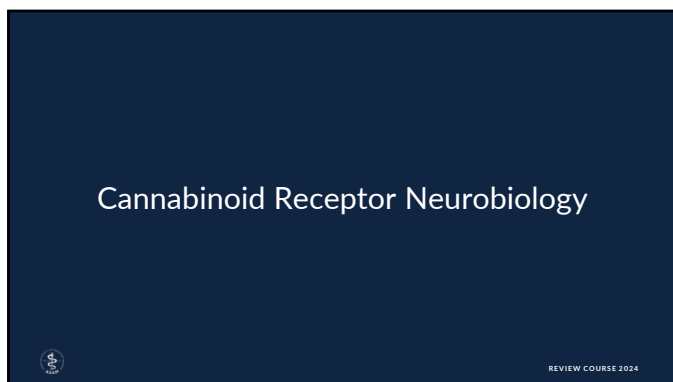
- Higher affinity for cannabinoid receptors than THC
- Have active metabolites that prolong their durations of action
- Increased potential for toxicity
- "Spice" or "K2"
- Not detected on standard UDS



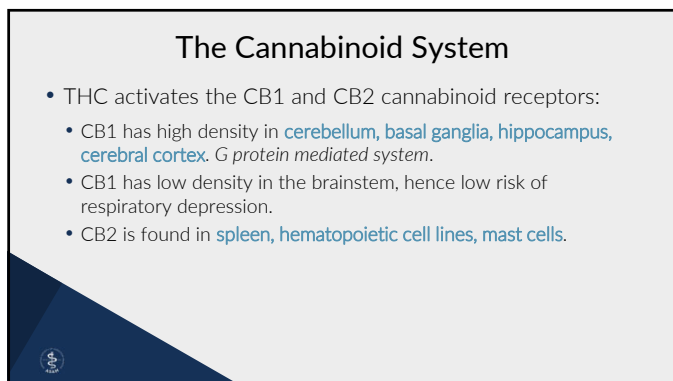
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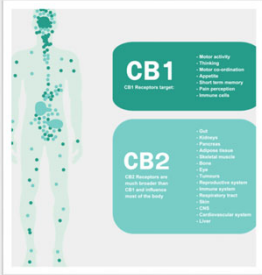
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Cannabinoid Receptors


- CB1 – CNS site of CB binding
 - Memory, learning, problem solving, coordination
 - Activated by anandamide, other CBs
 - Modulates neurotransmitters
- CB2 – immune cells outside CNS
 - Anti-inflammatory effects



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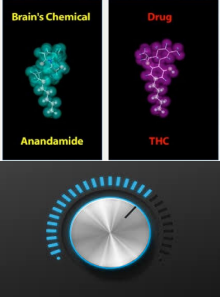
Cannabinoids (CBs)

- > 400 chemicals, ↓ neurotransmitter release
- **Natural CBs**
 - **Endogenous** – Anandamide, 2-Arachidonoylglycerol (AEA, 2-AG)
 - **Exogenous** – Sativa or Indica plant (marijuana)
 - Tetrahydrocannabinol (THC) – psychoactive
 - Cannabidiol (CBD) – no effect in brain



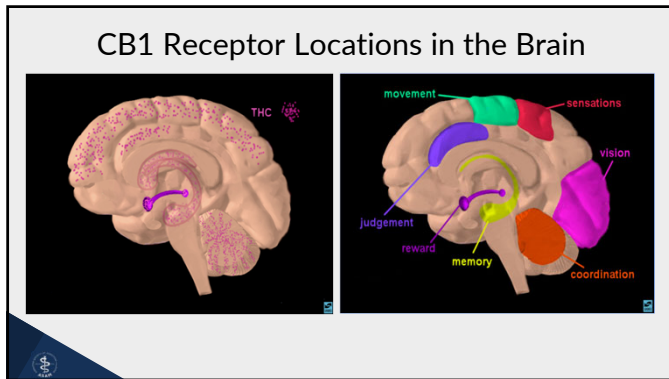
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9-tetrahydrocannabinol (THC)



- Primary psychoactive constituent
- Endocannabinoid system
 - Brain development
- Mimics anandamide
 - Dial down neuron activity

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Neurotransmitter modulation

- Dopamine – euphoria, reward, pleasure
- GABA- muscle relaxation and sleepiness
- ↓ Glutamate- relaxation, ↓ memory

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Cannabis Intoxication

- *Desired effects:* relaxation, euphoria, slowed time perception, altered sensory perception, increased appetite.
- *Undesired effects:* impaired concentration, anterograde amnesia, anxiety, panic attacks, paranoia, derealization/depersonalization, psychosis (visual – not auditory hallucinations).

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Synthetic Cannabinoid Toxicity

Central Nervous System	Seizures	Cardiovascular	Tachycardia
	Agitation		Hypertension
	Irritation		Chest pain
	Loss of consciousness		Cardiac Ischemia
	Anxiety	Gastrointestinal	Nausea
	Confusion		Vomiting
Paranoia			
Metabolic	Hypokalemia	Autonomic	Fever
	Hyperglycemia		Mydriasis
		Other	Conjunctivitis

Seely et al. Marijuana-based Drugs: Innovative Therapeutics or Designer Drugs of Abuse? *Med Interv*. 2011;11(1):36-51

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Routes of Administration

- Smoked:
 - Reaches the brain in minutes
 - Effects last 1 - 3 hours
 - Delivers significant amount of THC into the bloodstream

Smoked	Vaporized	Eaten/Drunk
Smoked in a pipe, bowl, cigarette	Inhaled through machine that converts active compounds into inhalable form	Consumed as ingredient in baked goods, candies, sodas
Rapid effects	Rapid effects	Takes time to reach brain, so effects are delayed

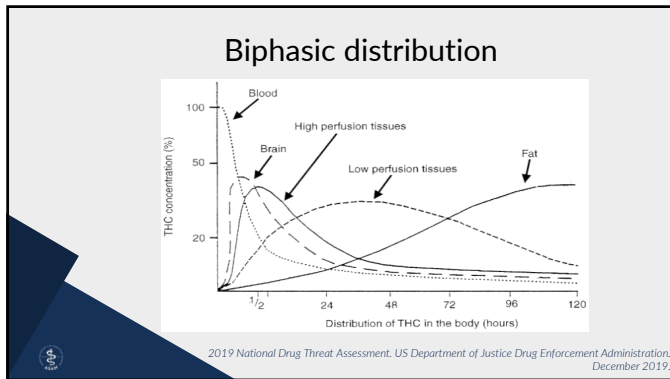
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Routes of Administration

- Eating or drinking marijuana:
 - Takes ½ - 1 hour to have an effect
 - Effects last up to 4 hours
 - THC is metabolized by the liver into 11-hydroxy-THC
 - 11-Hydroxy-THC is more lipophilic, potent and has a longer half-life.

Smoked	Vaporized	Eaten/Drunk
Smoked in a pipe, bowl, cigarette	Inhaled through machine that converts active compounds into inhalable form	Consumed as ingredient in baked goods, candies, sodas
Rapid effects	Rapid effects	Takes time to reach brain, so effects are delayed

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Toxicology Testing

- Casual use:
 - Up to 10 days in urine
 - 50% positive in hair samples
- Heavy use:
 - Up to 30 days in urine
 - 85% positive in hair samples
- Measures THC
- Weight loss gives serial UTox spike
- Dronabinol gives positive test
- Passive inhalation gives negative test

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Cannabis Withdrawal

- Reported by up to 1/3 of persons who use cannabis frequently.
- Cannabis withdrawal is recognized by the DSM 5.
- Clinical trials – show reduction of withdrawal symptoms with synthetic THC (dronabinol), nabilone, nabiximol, and gabapentin.

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Cannabis Withdrawal

Causing distress & ≥ 3 of the following:

- Irritability
- Anxiety
- Sleep problems
- \downarrow Appetite/weight loss
- Depressed Mood
- Restlessness

AND ≥ 1 of the following:

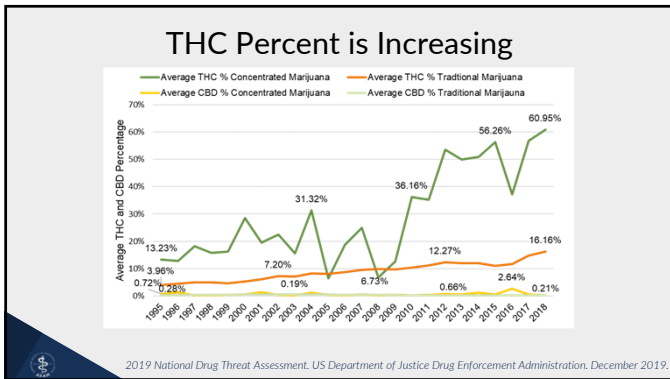
- Abdominal pain
- Sweating
- Shakiness/tremors
- Fever/chills
- Headache

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THC Potency is Increasing

- Up to 31 % in products
- Widespread availability of THC edibles (food and beverage products) and butane-extracted hash oil products (“dabs”, “budder”, “shatter”, “wax”)
- Rate of ED visits per 100,000 for cannabis-related adverse reactions has dramatically risen: **96.2 to 146.2** (2004 in 2011).


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Special Populations and Cannabis


- Adolescents
- Pregnant persons



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Decreased Harm Perception: Adolescents


- 36% of teens think cannabis is harmless
 - 43% favor legalization
 - 80s: 15%
 - 90s-00s: 30%
- Harm perception lowest in 40 yrs
 - Often precedes ↑ prevalence



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Rates ↑ Across Adolescence


<ul style="list-style-type: none">• Ever tried<ul style="list-style-type: none">• ~17% 8th graders• ~50% 12th graders• Past year use<ul style="list-style-type: none">• 12% 8th graders• 35% 12th graders	<ul style="list-style-type: none">• Current use (past month)<ul style="list-style-type: none">• 7% 8th graders• 21% 12th graders• Surpasses current alcohol and tobacco use
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Pregnancy

- Children exposed to THC:
 - Problem-solving skills, memory, attention deficit
- THC-specific vs. associated environmental factors hard to sort out; ongoing debate and research.



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Effects of Use




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Physiological Effects


- Adrenergic look-alike:
 - Tachycardia
 - Hypertension (but orthostatic hypotension)
 - Tachypnea
 - Dry mouth
- Conjunctival injection
- Appetite increase



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Impaired Cognition


- ↓ Ability to learn
- ↓ Attention, concentration
- ↓ Abstract reasoning and decision-making
- ↓ Memory



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Neurocognitive Effects

- Short-term memory impairment
- Judgment impairment
- Motor coordination impairment (increased risk of MVA)



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Impaired Driving


- Acute THC
 - → ↓ Peripheral vision
 - → ↓ Motor coordination
 - → ↑ reaction time
 - → ↓ time/distance judgment
- #1 reported illicit drug in accidents/fatalities
 - 2x accident risk
 - 3-7x risk of causing accident



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Physical Health


- Respiratory
 - ↓ Function
 - ↑ Infections
- ↑ Stroke/Temporary brain blood constriction



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Psychiatric


- Anxiety
 - Acute THC → ↓ anxiety
 - Long-term THC → ↑ anxiety
- ↑ Depression
- ↑ Psychosis



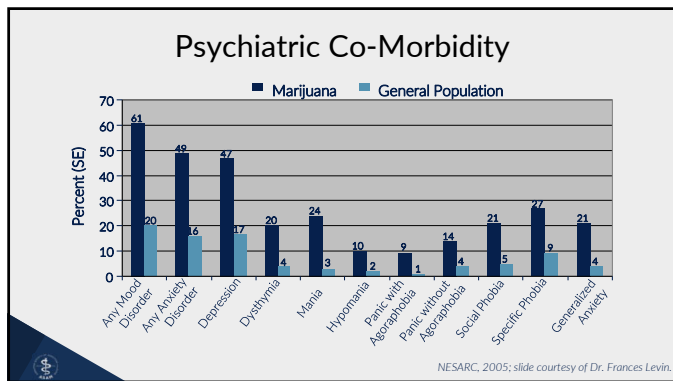
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Amotivational Syndrome

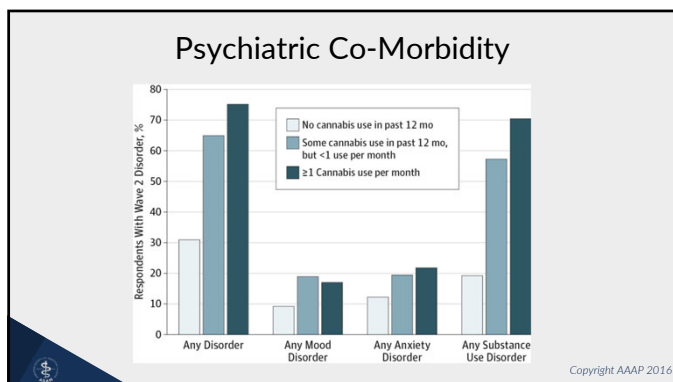
- Mental slowing
- ↓ Planning ability
- ↓ Judgment, concentration, memory
- Apathy, ↓ pursuit of goals



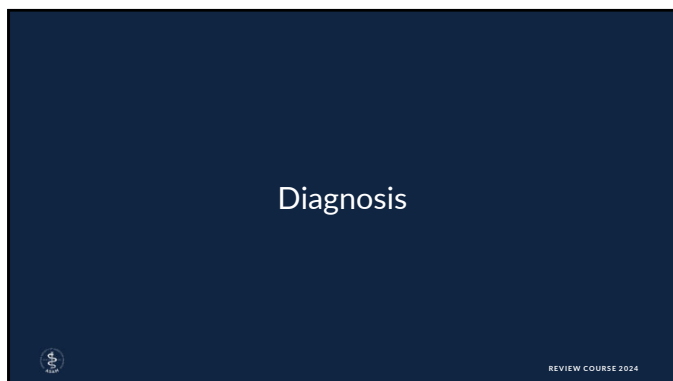
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Substance Use Disorder


In Same Year, ≥ 2 of:

- Tolerance
- Withdrawal
- Use more/longer
- Unable to \downarrow use
- Use despite problems
- Craving
- Failed roles
- Hazardous use
- Social problems
- \downarrow Activities
- Lots time use



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Cognitive Effects




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Residual Cognitive Effects


- Memory
 - Learning & retaining new information
- Attention and concentration
 - Response speed & variability
- Executive functioning
 - Working memory
 - Verbal fluency



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Likely Reversible with Abstinence

- Biological markers normalize ~4wks
 - CB receptor density in brain
 - Cortical blood volumes
- Especially in cognitive areas



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Treatment




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Treatment for CUD is Challenging

- Few evidence-based supported approaches
- ~ 50% achieve remission
- ~ 70% return to use
- No FDA-approved medications



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Psychosocial Treatments


- Motivational Enhancement Therapy
- Cognitive Behavior Therapy
- Contingency Management
- Family-Based Programs



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Pharmacologic Treatment Options


Medication	Mechanism	Comments	Literature in Adolescents?
Atomoxetine	Norepinephrine reuptake inhibitor	<ul style="list-style-type: none"> • No change in cannabis use • Worsened irritability and GI side effects 	• Thurstone et al., 2010 ⁷
Bupropion	Norepinephrine reuptake inhibitor	<ul style="list-style-type: none"> • Exacerbated withdrawal (irritability, insomnia) 	• Riggs, et al., 2013 ⁸
Buspirone	Serotonin partial agonist	<ul style="list-style-type: none"> • Conflicting evidence on cravings and irritability 	..
Dronabinol	CB1 receptor agonist	<ul style="list-style-type: none"> • Reduced symptoms of withdrawal • Contains THC 	..
Gabapentin	GABA modulation	<ul style="list-style-type: none"> • Decrease self-reported cannabis use • Reduced withdrawal symptoms 	..
N-acetylcysteine	Correct glutamate dysregulation	<ul style="list-style-type: none"> • Decreased use in adolescents • Did not show same benefit in adults 	• Gray et al., 2012 ⁹
Naltrexone	Mu-opioid receptor antagonist	<ul style="list-style-type: none"> • Enhanced subjective effects of cannabis • No change in frequency of cannabis use 	..



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Medication for CUD

- N-acetylcysteine (NAC)
 - Amino acid derivative, OTC supplement
 - Restores normal glutamate activity
 - Pros: ↓ use in Non-Treatment Seeking adolescents, *not in adults*
 - Cons: did not ↓ craving



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N-acetylcysteine (NAC)

Risks	<ul style="list-style-type: none"> Nausea/vomiting Drowsiness/insomnia Vivid reams Anaphylactoid reactions seen with IV admin, not PO
Pharmacokinetics	<ul style="list-style-type: none"> Bioavailability for oral: 9% Metabolized to cysteine and glutathione Half-life: ~ 18 hours

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Gabapentin

Mechanism of Action	<ul style="list-style-type: none"> Blocks alpha-2d subunit of the voltage-gated calcium channel which modulates GABA in the amygdala
Notes	<ul style="list-style-type: none"> FDA approved for multiple indications, including partial seizures in ages 3-12
Doses	<ul style="list-style-type: none"> Goal of ~1200mg/day Mason (2012) ¹⁸: 50 cannabis-dependent adults (18-65 years old) Gabapentin 1200mg vs placebo for 12 weeks <ul style="list-style-type: none"> • Titrated up to 300mg / 300mg / 600mg over the course of 4 days
Clinical benefit	<ul style="list-style-type: none"> Increase in negative UDS Decrease self-reported cannabis use Reduction in withdrawal symptoms (mood disturbance, craving, and sleep disturbances)

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Gabapentin

Risks	<ul style="list-style-type: none"> Well tolerated @ Headache, nausea, insomnia and depression
Pharmacokinetics	<ul style="list-style-type: none"> Bioavailability: Inversely proportional due to saturable absorption <ul style="list-style-type: none"> • Immediate release <ul style="list-style-type: none"> • 900mg/day: 60% • 1200mg/day: 47% • 3600mg/day: 33% • 4800mg/day: 27% • Extended release: increased with higher fat content Half-life: <ul style="list-style-type: none"> • ≤ 12 years old: 5hr • > 12 years: 5-7hr • Longer in patients with decreased renal function

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CB1 Receptor Agonists

Cannabidiol (CBD) Epidiolex®	Dronabinol (THC) Marinol® Syndros® Nabilone (THC) Cesamet®	Nabiximols (THC + CBD) Stavivex® <i>not FDA approved</i>
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Medicinal Uses of Cannabis/Cannabinoids

- Dronabinol: FDA approved for treatment of anorexia associated with weight loss in patients with AIDS, chemotherapy-induced nausea/vomiting.
- Nabilone: FDA approved for treatment of chemotherapy-induced nausea/vomiting.
- Studies also ongoing re: effects on other disease states (epilepsy, MS).

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Therapeutic Potential

- Pain (cancer, multiple sclerosis)
- Nausea (cancer)
- Loss of appetite and wasting (HIV/AIDS)
- Increased ocular pressure (glaucoma)
- Inflammation (rheumatoid arthritis, Crohn's disease, ulcerative colitis)
- Epilepsy

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In Summary


Cannabis includes plants and synthetic cannabinoids.	Cannabis use is common, risk of a use disorder increases with earlier onset of use.	Cannabis contains more THC now than in the past.
Cannabis can affect cognition, but this is reversible in adults, impacts on adolescents less clear.	Most treatment is psychosocial, but several drug targets are being investigated.	



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Which of the following trends in youth from the Monitoring the Future study about marijuana use and perception of harm is true?


- A. Since the early 1990's, the percentage with perceived risk of harm from marijuana has been higher than past year use of marijuana.
- B. Since about 2009, there has been a growing gap between decreased perception of harm and increased past year use of cannabis.
- C. The lowest past year cannabis use was in the late 1970's.
- D. The perceived risk of harm for cannabis fell throughout the 1980's.



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Which of the following medications has a trial supporting efficacy in cannabis use disorder in adolescents?

- A. N-acetylcysteine
- B. Baclofen
- C. Quetiapine
- D. Mirtazapine




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Cannabis use is reported in greater than 10% of pregnancies. Which correctly lists the reasons cannabis users who are planning to become pregnant should be cautioned against cannabis use:

- A. THC easily passes into breast milk and crosses membranes and is transferred to the developing fetus, and therefore impacts pregnancy success in females only.
- B. While THC does not pass into breast milk, studies show that it does easily cross membranes and is transferred to the developing fetus.
- C. While human studies on the effect of prenatal THC exposure on the developing brain are preliminary, they correlate with studies carried out in animals and show that THC easily passes into breast milk and crosses membranes and is transferred to the developing fetus.
- D. While no human studies have been done on the effect of prenatal THC exposure, animal studies show that it does easily pass into breast milk, crosses membranes, and is transferred to the developing fetus.

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