

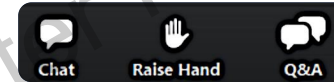
Welcome!



Ensure we can identify you for attendance.



Use the Zoom Q&A function to ask questions.



Have your camera on, if possible.



We'll send you a copy of the slides.



Remain muted when not speaking.



Complete post-training eval within 2 weeks.



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Massachusetts recording law stipulates that it is a two-party consent state. In Massachusetts, it is a criminal offense to use any device to record and/or disseminate communications, whether they're wire, oral or electronic, without the consent of all contributing parties. Mass. Ann. Laws ch. 272, § 99(C). This means that in Massachusetts you are not legally allowed to record a conversation you are taking part in unless all parties are in agreement.



Cannabis 301: Understanding & Responding to Psychoactive Effects

Megan (Meg) Hudson, MSN, PMHNP-BC
April 17, 2026

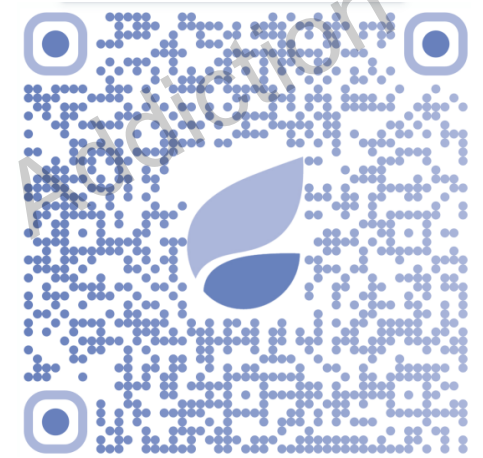


Grayken Center for Addiction
Training & Technical Assistance
Boston Medical Center



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CANNABIS AND NICOTINE EDUCATIONAL TRAINING SERIES FOR YOUTH AND ADULT CARE



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- Thu, 1/15, 1-2pm – Nicotine 101
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Funders



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The faculty and planning committee have no relevant financial relationships to disclose.

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In support of improving patient care, Boston University Chobanian & Avedisian School of Medicine is jointly accredited by the Accreditation Council for Continuing Medical Education (ACCME), the Accreditation Council for Pharmacy Education (ACPE), and the American Nurses Credentialing Center (ANCC), to provide continuing education for the healthcare team.

CME: Boston University Chobanian & Avedisian School of Medicine designates this live activity for a maximum of **1.00 AMA PRA Category 1 Credit(s)**[™]. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

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Accreditation Information



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APA CE Designation Statement: Continuing Education (CE) credits for psychologists are provided through the co-sponsorship of the American Psychological Association (APA) Office of Continuing Education in Psychology (CEP). The APA CEP Office maintains responsibility for the content of the programs.

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Boston University Chobanian & Avedisian School of Medicine Barry M. Manuel Center for Continuing Education is recognized by the New York State Education Department's State Board for Psychology as an approved provider of continuing education for licensed psychologists #PSY-0181. Note: NYS psychologists must attend all sessions for credit. Partial credit is not allowed.

The Department's approval of a provider of continuing education does not constitute the Department's endorsement of the content, positions or practices that may be addressed in any specific continuing education course offered by the approved provider

Accreditation Information



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LADC/CADC & Recovery Coach: Boston Medical Center Grayken Center for Addiction Training and Technical Assistance is approved by the Massachusetts Department of Public Health, Bureau of Substance Addiction Services (DPH/BSAS) to award LADC/CADC and Recovery Coaches who complete this training **1.00** continuing education credits.

Objectives

Following this training, participants will have the knowledge to:

- Identify risk factors for psychosis and psychotic-like experiences associated with cannabis use.
- Recognize symptoms of cannabis-induced psychosis.
- Describe at least two interventions for the management of acute psychosis associated with cannabinoid use.
- Evaluate interventions to reduce risk for harm when providing long-term care for individuals with a history of cannabis-induced psychosis or psychotic-like experiences.

Cannabis & Psychosis Overview

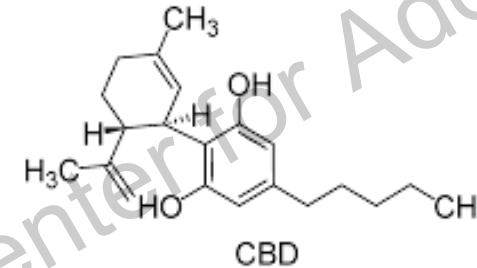
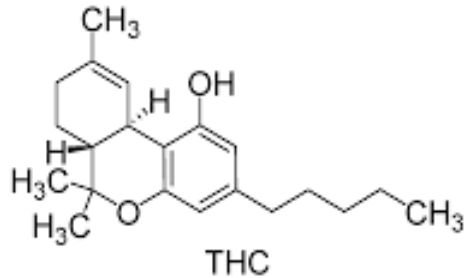
Terminology

Cannabis – general term referring to psychoactive substances produced by the *Cannabis sativa* plant

Cannabinoids – compounds structurally similar to THC or share many of the pharmacological properties of THC (includes synthetic THC)

weed (weed)
marijuana (mar-uh-wah-nuh)
pot (pot)
chronic (kron-ik)
cannabis (kan-uh-bis)

Roles of THC and CBD



Tetrahydrocannabinol (THC)

- Psychoactive, appetite stimulant
- Dose-response relationship between THC and adverse (e.g., neurocognitive) effects
- Delta-9 THC isomer = most common
- *Delta-8 THC naturally exists in small quantities, ~ 50-75% as psychoactive; Growing quasi-legal market.

Cannabidiol (CBD)

- No psychotomimetic properties
- Neuroprotective, anti-inflammatory, and anticonvulsant effects.
- Mixed evidence about its ability to improve/reduce psychosis with THC
- Antagonist at CB1 receptors (opposite of THC)

Acute Intoxication

Transient, linked to cannabis intoxication, self-limiting

Anxiety, paranoia, perceptual disturbances

Cannabis-Induced Psychosis (CIP)

Typically develops during or soon after use, persists beyond expected intoxication period (days to weeks / ~1 month)

Hallucinations and delusions that are prominent and impair functioning

Primary Psychotic Disorder

Symptoms exist independent of substance use, may be worsened with active use or withdrawal

Hallucinations, delusions, cognitive disorganization, negative symptoms, may include mood symptoms

Withdrawal

Symptoms exist in the setting of acute withdrawal

Paranoia most common, hallucinations & delusions possible

Prevalence & Risk Factors

Prevalence of Psychosis with Cannabis Use

Acute Psychotic Symptoms with Intoxication:

- 5-50% of adults report transient psychotic symptoms during intoxication
 - Symptoms are typically dose-dependent and last a few hours
- Data is limited in adolescents

Cannabis-Induced Psychosis

(CIP): Limited data on prevalence, recent estimates 3-6/100,000

Conversion to Chronic Psychotic Disorders:

- 20-50% of patients with CIP develop long-term psychotic disorders (i.e., schizophrenia)
- Approximately 50% of those who transition do so within 2 years
- Cannabis-induced psychosis has the highest conversion rate to schizophrenia across substances

(Gorelick, 2023)
(Hua et al., 2021)
(Kansagara et al., 2026)
(Osborne et al., 2025)
(Javed et al., 2026)

Risk Factors

Age of Initiation:

- Adolescent cannabis use is associated with an 11-fold increased risk of psychotic disorders
- Greatest risk when started in adolescence and continued into adulthood

Frequency and Potency:

- Daily use: 3.2x greater chance of psychotic disorder vs never using
- High potency THC: 1.6x greater chance of psychotic disorder

(Gorelick, 2023)
(Hua et al., 2021)
(Jin et al., 2026)
(Kansagara et al., 2026)
(Volkow et al., 2016)

Risk Factors Continued

Genetic Predisposition:

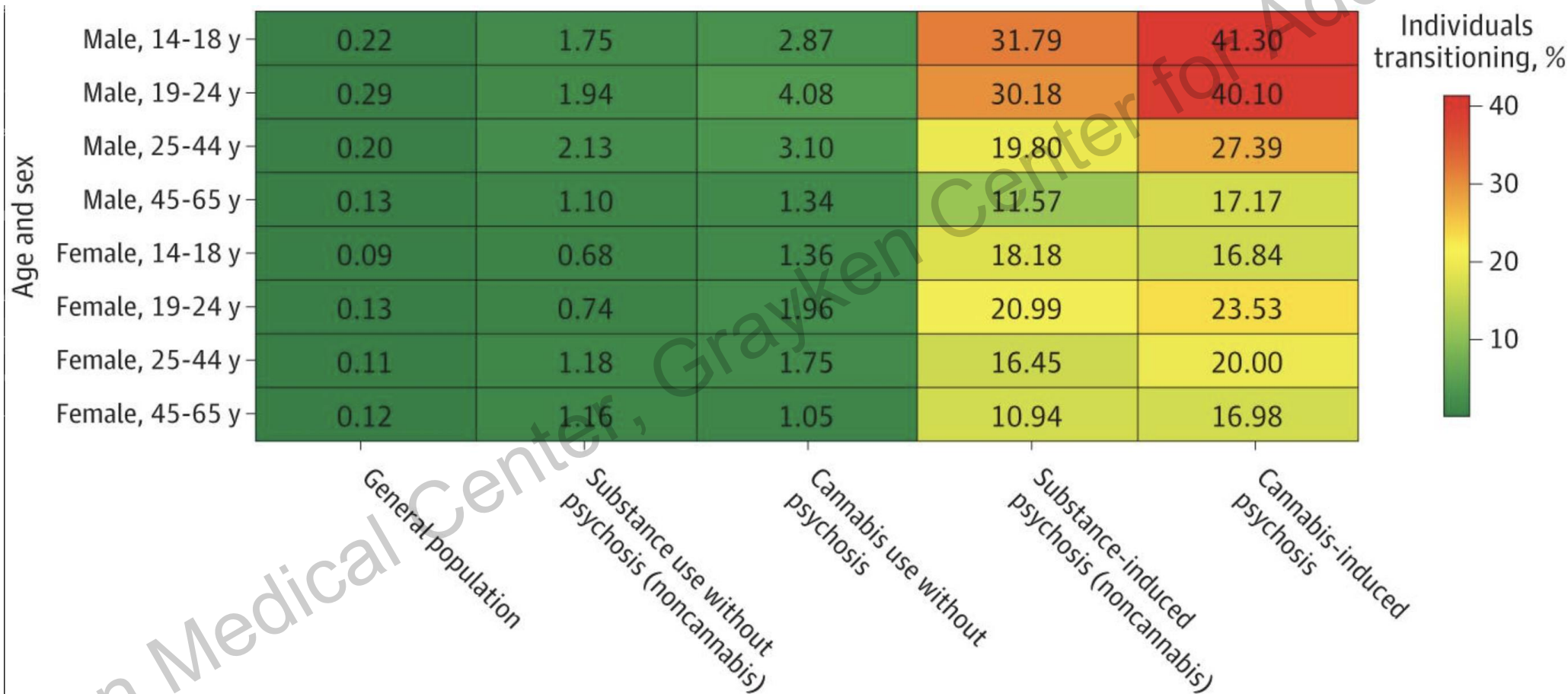
- Family or personal history of psychotic disorders
- Catechol-O-Methyltransferase (COMT) gene variations associated with increased risk of CIP
 - Impacts how the brain processes dopamine

Impact on Existing Psychotic Disorders:

- Cannabis use worsens illness course
- Ongoing use after first psychotic episode associated with poorer prognosis

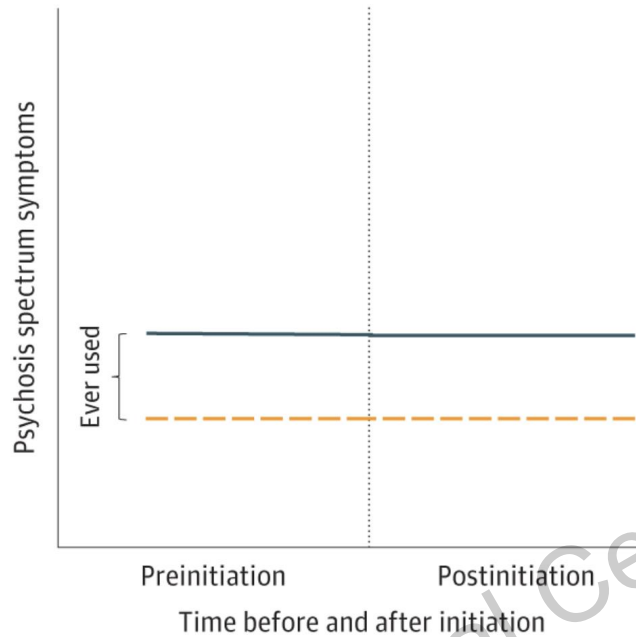
(Gorelick, 2023)
(Hua et al., 2021)
(Jin et al., 2026)
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Risk of Transition to Schizophrenia

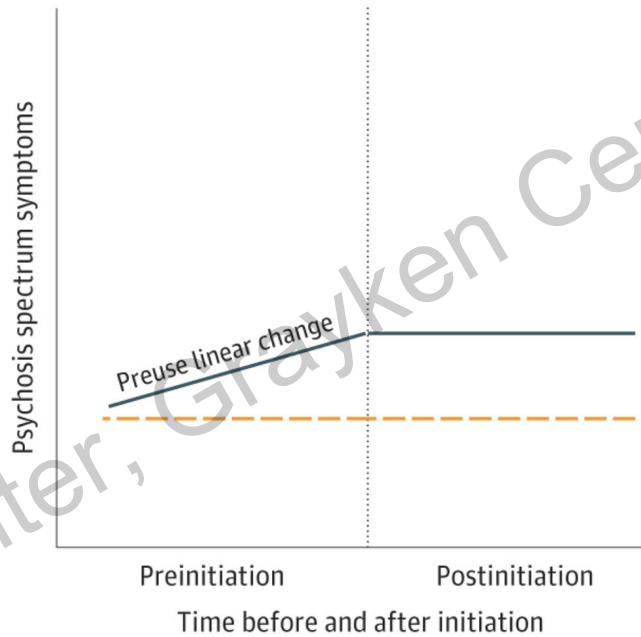


Potential Conceptual Patterns

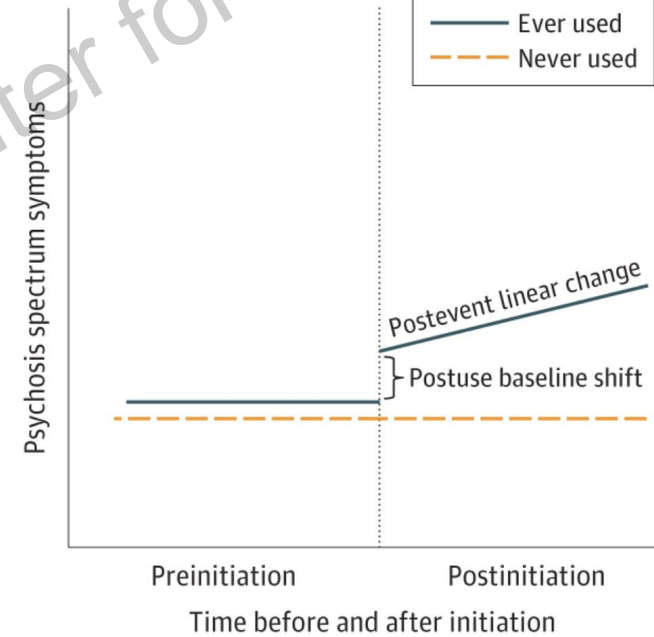
A Shared vulnerability hypothesis



B Self-medication hypothesis



C Contributing risk hypothesis



The figure shows possible patterns of how psychosis symptoms may change over time under three different explanations: a shared vulnerability, self-medication, or contributing risk model.

Cannabis & the Brain

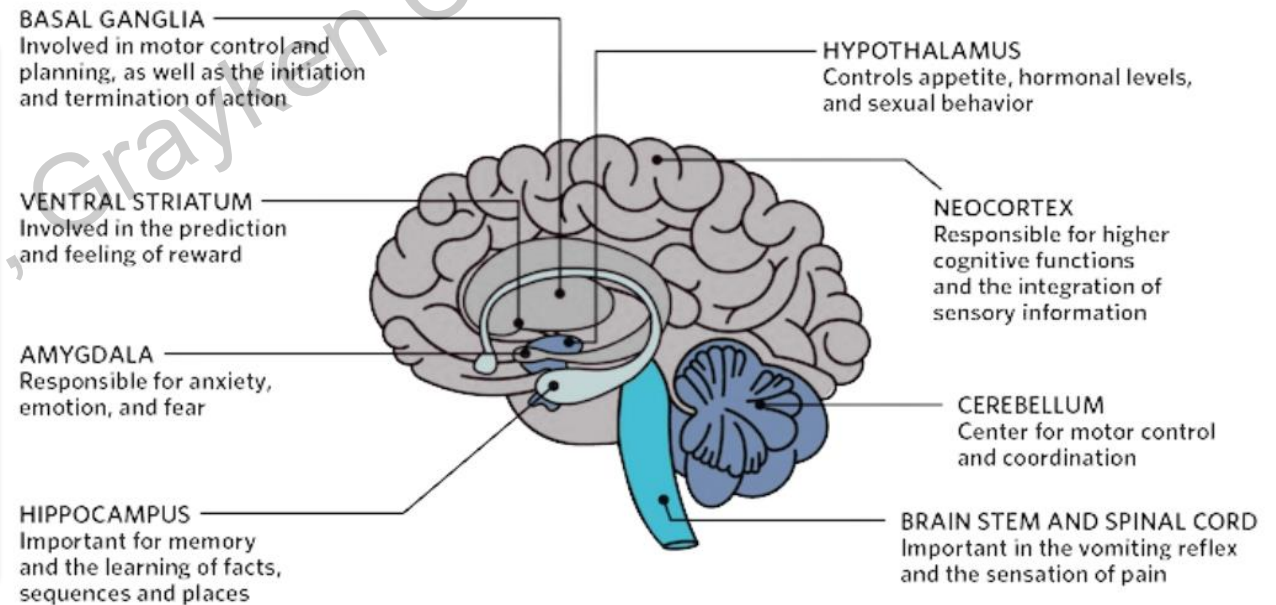
Cannabis & the Brain

- Tetrahydrocannabinol (THC) activates CB1 receptors in the brain
- CB1 part of the endocannabinoid system that regulates mood, memory, reward, and pain
 - Highly concentrated in the prefrontal cortex, hippocampus, basal ganglia, and cerebellum
 - Impacts attention, memory, decision-making, coordination, and perception

THC increases dopamine release in reward pathways, contributing to its reinforcing effects and potential for dependence.

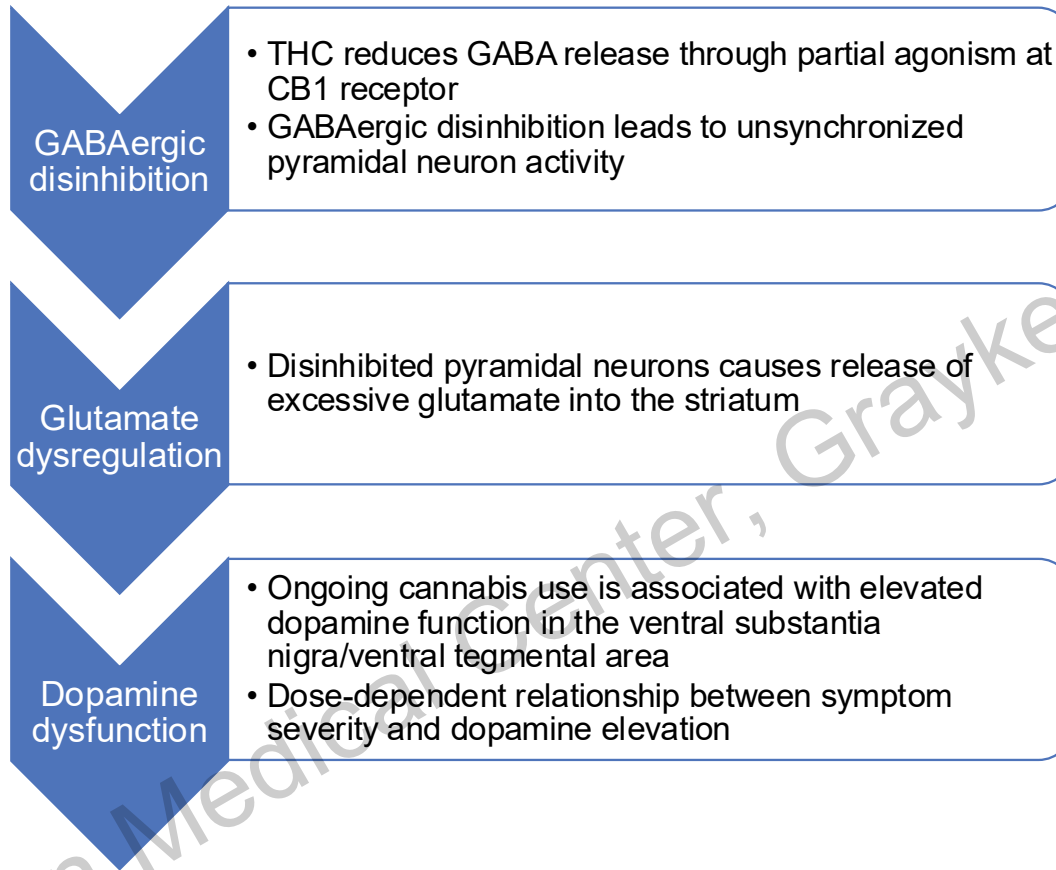
Acute cannabis use may produce euphoria, relaxation, altered perception, impaired short-term memory, and slowed reaction time.

Heavy and long-term use can result in going structural changes/dysregulation



(Simons, 1998)
(Simons, Correia & Carey, 2000)
Image Source: Adapted from APA

How Does Cannabis Cause Psychosis?



Final Common Pathway: Disrupted Salience Processing

- Breakdown of coordination between prefrontal cortex and striatum causes the brain to assign importance to irrelevant stimuli
 - Hallucinations, delusions, paranoia
- Aberrant salience is also seen in other psychotic disorders (i.e. schizophrenia)

Ongoing Psychiatric Consequences of Cannabis

Acute Cognitive Impairment

- Dose-dependent deficits in verbal learning, working memory, executive functioning, processing speed

Chronic Cognitive Effects

- Small decline in IQ seen in adolescents with regular use
- Chronic use in adults associated with ongoing cognitive deficits, likely to improve with prolonged abstinence (more persistent with adolescent onset)

Mood/Anxiety Disorders

- Increased risk for bipolar affective disorder (more likely to have psychotic features present), MDD, and anxiety disorders

Dementia

- Emerging evidence shows individuals with ED visits for cannabis use had 1.7 fold increased risk of dementia
- Long-term use shows smaller hippocampal volume

Acute Intoxication & Withdrawal

Symptoms & Management

Acute Cannabis Intoxication

Psychological

- Euphoria
- Relaxation
- Sedation
- Increased appetite
- Anxiety
- Psychotic symptoms
 - Hallucinations, delusions, paranoia

Physical

- Impaired motor coordination
- Slurred speech
- Conjunctival injection
- Tachycardia
- Orthostatic hypotension

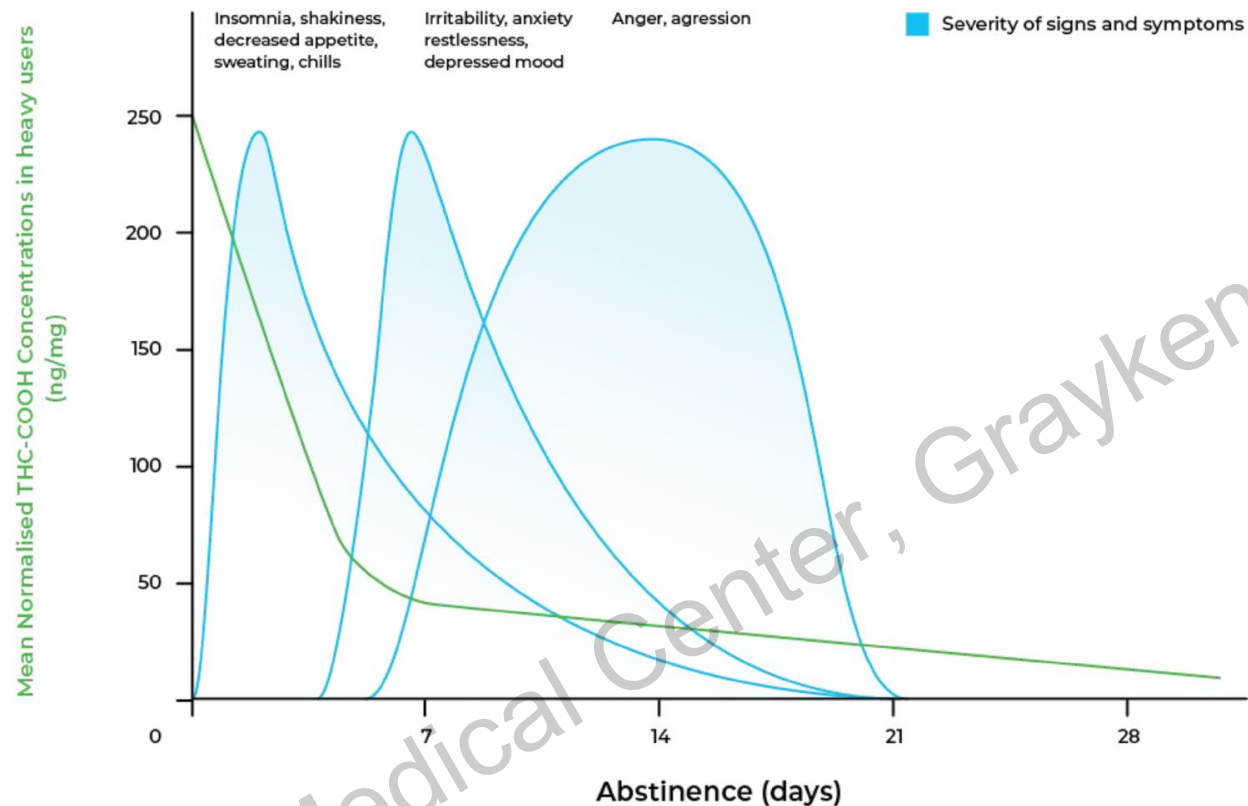
Acute Management

There are currently no FDA-approved medications for acute management of cannabis intoxication/acute cannabis-induced psychosis.

- Supportive care is first line: reduction of stimulation, supportive reassurance.
- Benzodiazepines can be used for acute agitation or anxiety
- As-needed second-generation antipsychotics for symptoms of psychosis
 - Olanzapine (2nd gen) & haloperidol (1st gen) have been directly compared in their use, equally effective but lower risk of side effects with olanzapine

Always connect to long-term care when possible! Assess for ongoing psychosis in the absence of cannabis use

Cannabis Withdrawal & Psychosis



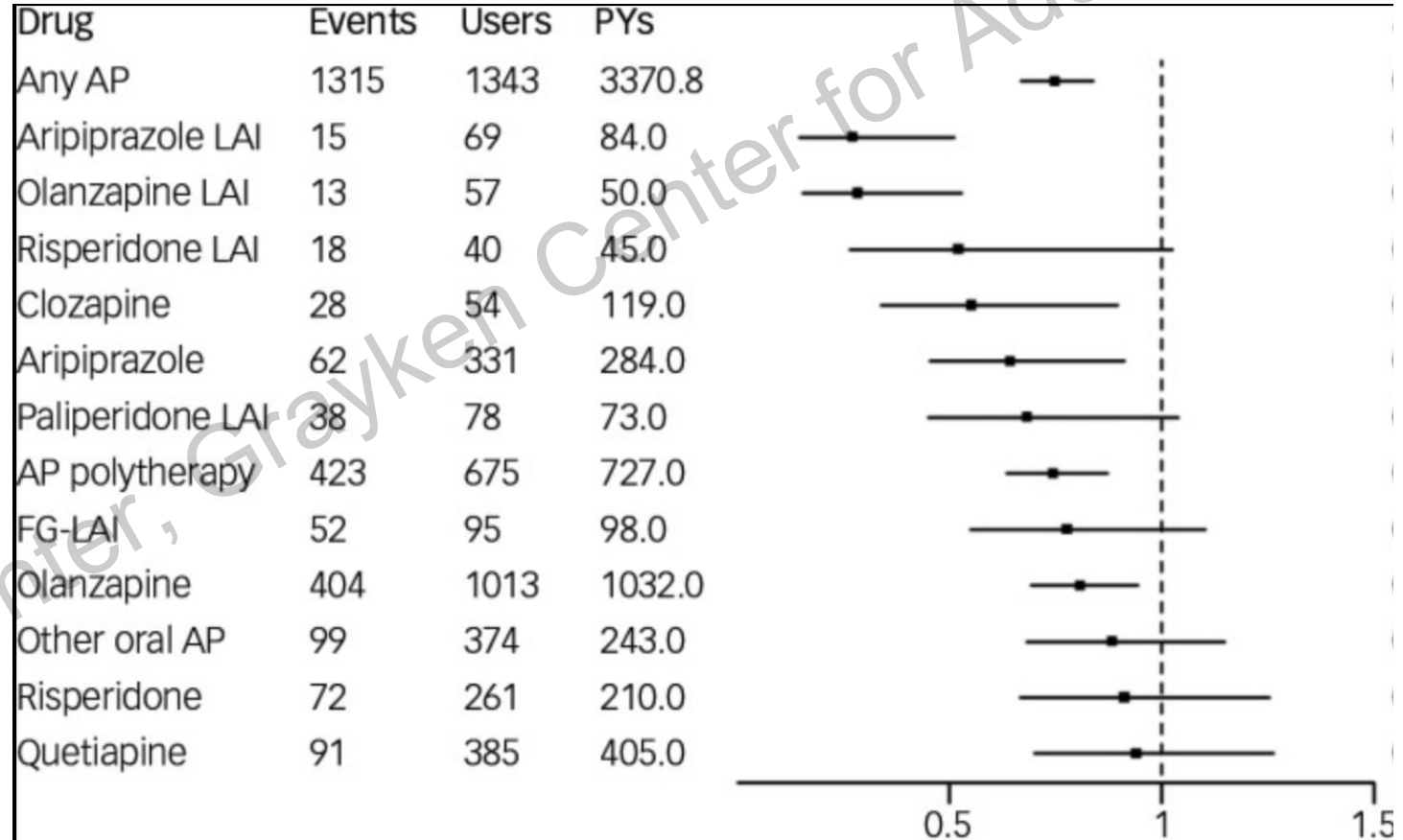
- Most common withdrawal symptoms include sleep disruption, decreased appetite, irritability
- Onset of frank psychotic symptoms often within 1 week of abrupt cessation, highest incidence around day 4
- Most common psychotic symptoms include paranoia, but can include hallucinations and delusions

Reducing Risk

Antipsychotic Effectiveness after CIP

Recent study of nearly 2,000 patients with CIP showed that ANY antipsychotic use reduced hospitalizations for psychosis

- Long-acting injectables and clozapine showed the best benefit
- Quetiapine had no benefit



Reducing Risk with Ongoing Cannabis Use

Interventions to reduce harm:

- Lower THC content
- Reduce frequency of use
- Avoid products with inconsistent potency
- Ongoing therapeutic alliance/open communication about cannabis use and psychosis
- Consider off-label medications for CUD

Digital Harm Reduction: (Cannabis Harm-Reducing App to Manage Practices Safely, 'CHAMPS' app)

- Pilot RCT in first episode psychosis
- 82% retention; improved motivation to change and implement harm reduction strategy use
- Not yet available to the public

(Kapler et al., 2023)
(Gonzalez-Ortega et al., 2022)
(Coronado et al., 2025)
(Walker et al., 2025)

THC Potency Reduction

Gradually reducing THC potency instead of reducing frequency of cannabis intake can decrease dependency and withdrawal symptoms in a manageable way

- Reduces risks of higher THC-potency products
- Supports moving towards reduction of overall use or abstinence based on patient's goal

Example Reduction Plan

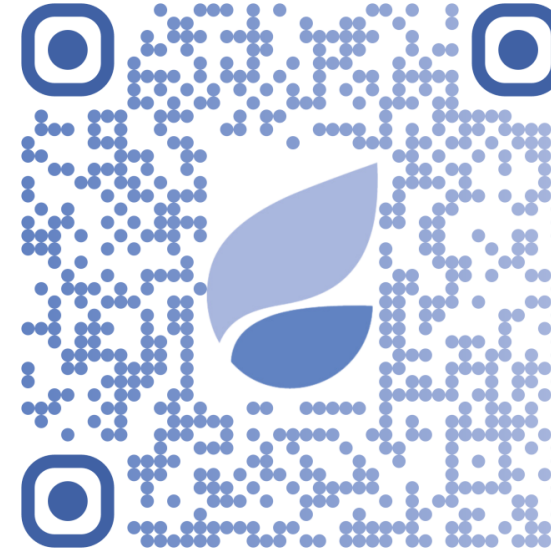
- Step 1: Know your potency/product
- Step 2: Choose a lower potency option over the next several weeks
 - Week 1: 40-50% THC
 - Week 2: 30-40%
 - Week 3: 20-30%
- Step 3: Track how you feel

Cannabis Risk Reduction



Resource:

Watch Grayken TTA's short animated educational video, "Harm reduction strategies for cannabis use". [Click here](#) or scan the QR code to access:



Providing Ongoing Care

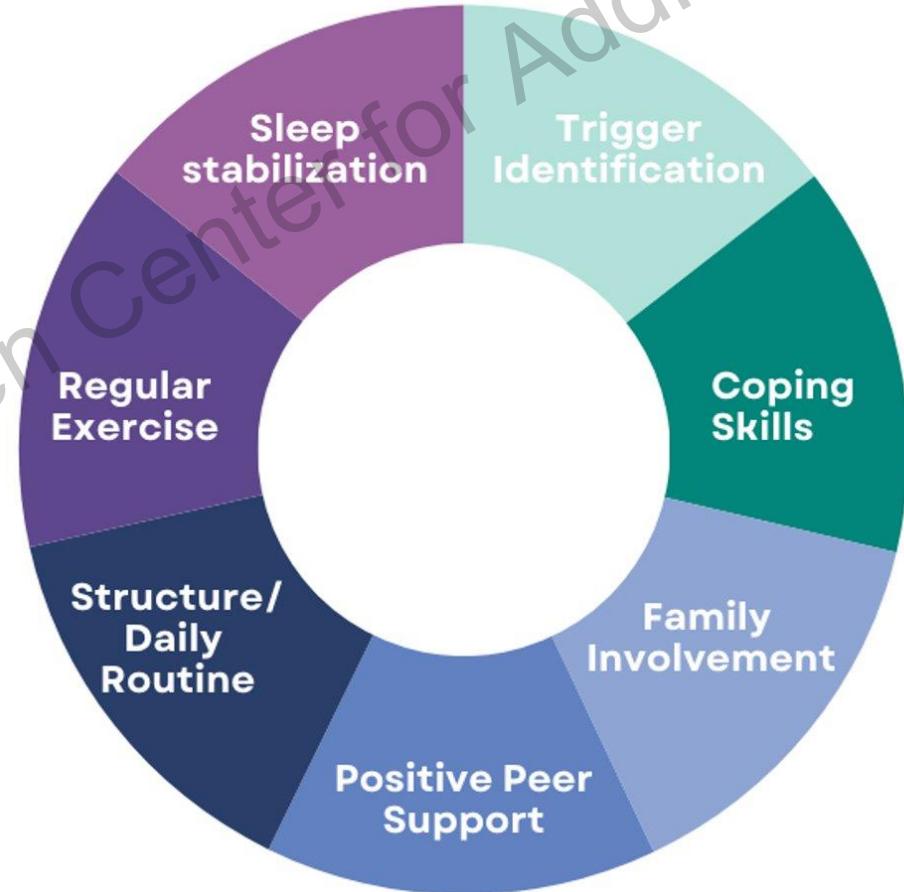
Psychosocial Interventions for CIP

Cannabis cessation: the most important modifiable factor

- Decreased risk of hospitalization, psychotic relapse
- First Episode Psychosis (FEP) with cannabis use show similar outcomes 10 years after stopping as those who never used

Evidence-based interventions:

- Cognitive Behavioral Therapy
- Motivational Enhancement Therapy (Cannabis Check-Up for Psychosis)



(Kapler et al., 2023)
(Gonzalez-Ortega et al., 2022)
(Coronado et al., 2025)
(Walker et al., 2025)

Recovery Capital Domains in CIP

Recovery Capital: Tangible & intangible resources individuals draw upon to initiate and sustain recovery from substance use and psychosis

Human/personal capital

- Self-esteem
- Insight & awareness
- Coping skills

Social capital (Strongest predictor of recovery in FEP)

- Social support network/peer support
- Family involvement
- Therapeutic alliance

Physical Capital

- Housing stability (substances in the home?)
- Access to healthcare

Cultural/Community

- Stigma management
- Meaningful activities
- Recovery identity development

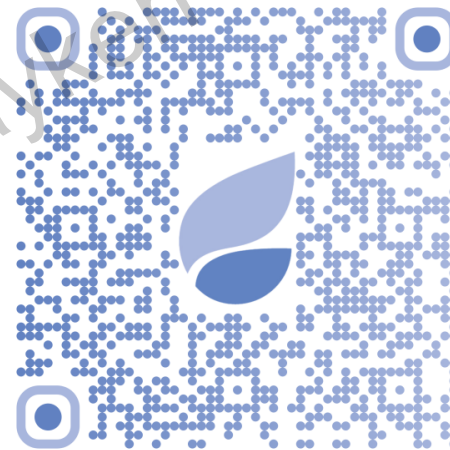
Barriers & Assessment Tools for Recovery Capital

Barriers

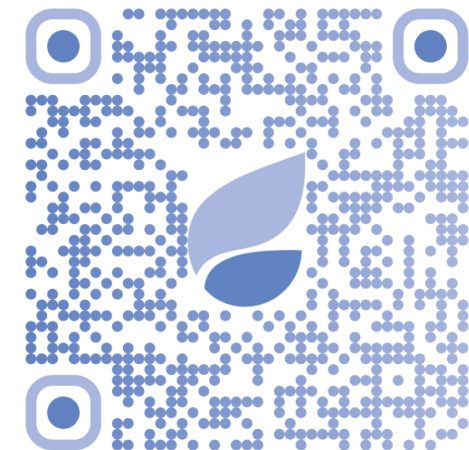
- Continued cannabis use
- Longer duration of untreated psychosis
- Prominent negative symptoms
- Poor premorbid functioning
- Internalized stigma, discrimination

Assessment Tools

Assessment of
Recovery Capital, ARC:



Brief Assessment of
Recovery Capital,
BARC-10:



Using the CGI to Track Progress and Care

Clinical Global Impression (CGI):

- Brief clinician-rated scale, validated for first episode psychosis
- Provides a quick clinical snapshot of overall clinical status (current severity and improvement)

Clinical Global Impression (CGI)

1. Severity of illness

Considering your total clinical experience with this particular population, how mentally ill is the patient at this time?

- | | |
|-----------------------------|---|
| 0 = Not assessed | 4 = Moderately ill |
| 1 = Normal, not at all ill | 5 = Markedly ill |
| 2 = Borderline mentally ill | 6 = Severely ill |
| 3 = Mildly ill | 7 = Among the most extremely ill patients |

2. Global improvement: Rate total improvement whether or not, in your judgement, it is due entirely to drug treatment.

Compared to his condition at admission to the project, how much has he changed?

- | | |
|------------------------|---------------------|
| 0 = Not assessed | 4 = No change |
| 1 = Very much improved | 5 = Minimally worse |
| 2 = Much improved | 6 = Much worse |
| 3 = Minimally improved | 7 = Very much worse |

3. Efficacy index: Rate this item on the basis of drug effect only.

Select the terms which best describe the degrees of therapeutic effect and side effects and record the number in the box where the two items intersect.

EXAMPLE: Therapeutic effect is rated as 'Moderate' and side effects are judged 'Do not significantly interfere with patient's functioning'.

	Therapeutic effect	Side effects			
		None	Do not significantly interfere with patient's functioning	Significantly interferes with patient's functioning	Outweighs therapeutic effect
Marked	Vast improvement. Complete or nearly complete remission of all symptoms	01	02	03	04
Moderate	Decided improvement. Partial remission of symptoms	05	06	07	08
Minimal	Slight improvement which doesn't alter status of care of patient	09	10	11	12
Unchanged or worse		13	14	15	16
	Not assessed = 00				

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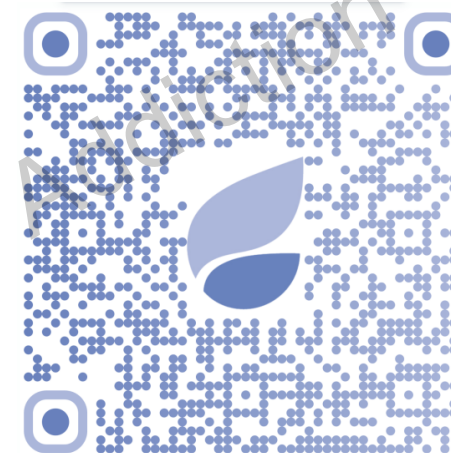
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Resources

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Virtual Drop-in Stimulant Office Hours

Monthly opportunities to ask your addiction-related questions



To learn more and join an upcoming session, [click here](#) or scan QR code!

Stimulant-Focused Office Hours:

3rd Thursday of each month from 5 – 6pm ET

- Hosted by BMC Grayken TTA Clinical Educators
- Open to all clinical providers and staff supporting those with substance use



FREE Pre-Recorded Trainings



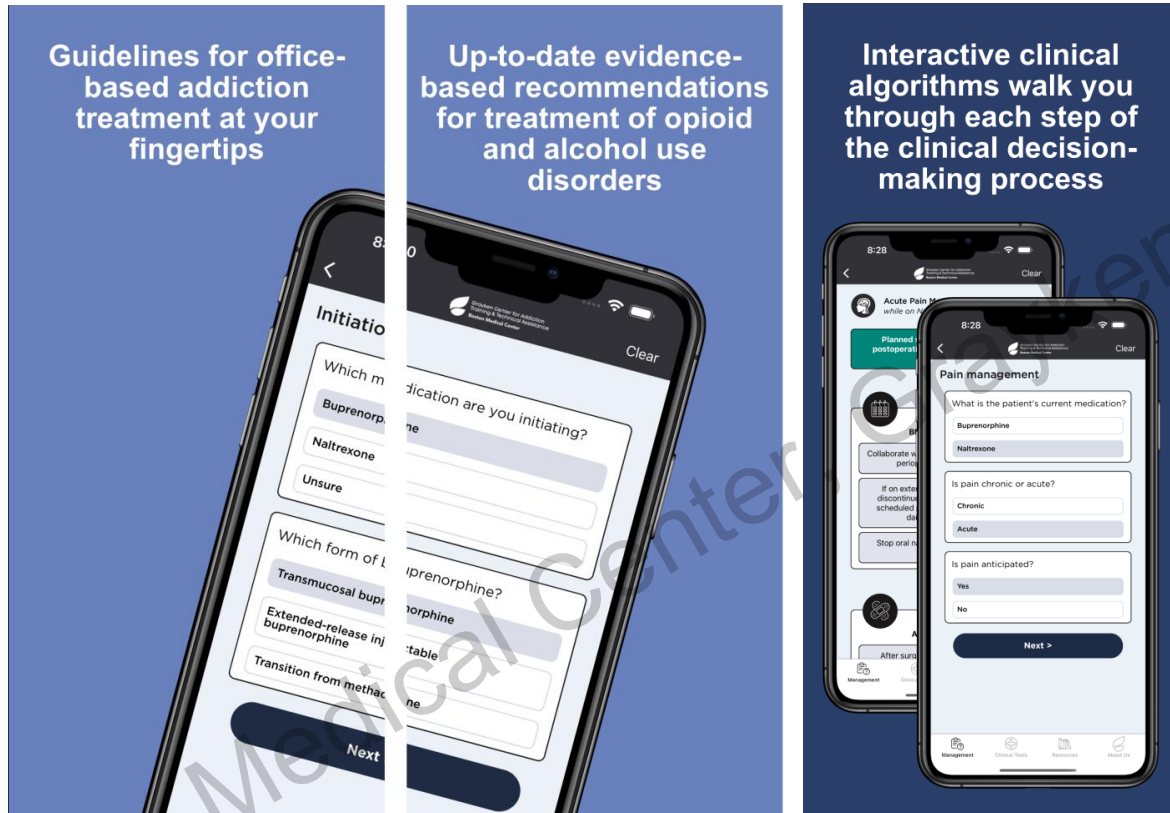
The screenshot shows the website header with the logo for Grayken Center for Addiction Training & Technical Assistance, Boston Medical Center. It includes a search bar with 'ENHANCED BY Google' and navigation links for LOGIN and CREATE ACCOUNT. The main navigation menu includes Home, DEA MATE Act, MAT Mobile App, Training (selected), Request TTA, Project ECHO®, Resources, News, and About Us. A sub-menu under Training shows Pre-recorded Trainings and Live Trainings. The main content area features the heading 'Pre-recorded Trainings' and a paragraph: 'Our free pre-recorded trainings are available on demand when it's convenient for you. Browse options by training topic and register online to get started.' Below this is a photo of a person in a blue lab coat working at a computer. At the bottom of the screenshot, there is a call to action: 'To access our free recorded trainings, [click here](#) or scan the QR code!'.

- ✓ Access trainings on various **specialty topics**
- ✓ Count towards **DEA MATE Act** requirement
- ✓ **FREE** CME/CE & completion certificates
- ✓ **On-demand 24/7**



BMC MAT Quick Start App

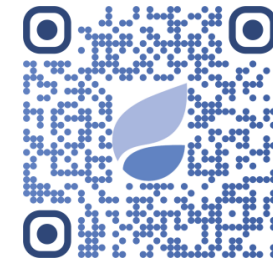
Free interactive clinical tools, decision trees, treatment protocols & resources



Provides real-time access to:

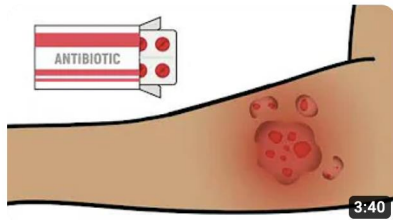
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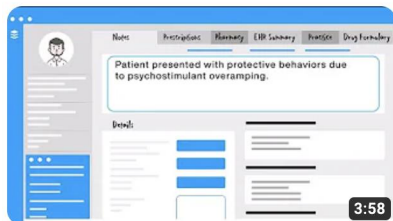
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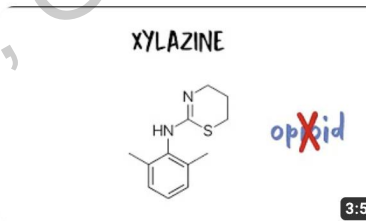
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