

MANAGING ALCOHOL USE DISORDER IN BC: NEW GUIDELINES, CHALLENGES & CLINICAL PEARLS

April 9, 2024 | 1830–2000 PT



THE UNIVERSITY OF BRITISH COLUMBIA

Continuing Professional Development

Faculty of Medicine

DISCLOSURES

Planning Team

- **Dr. Bob Bluman (UBC CPD):** No conflicts of interest
- **Nicole Esligar (UBC CPD):** No conflicts of interest
- **Kathryn Young (UBC CPD):** No conflicts of interest
- **Caldon Saunders (UBC CPD):** No conflicts of interest

Panelists

- **Dr. Simon Moore:** No conflicts of interest
- **Dr. Paxton Bach:** No conflicts of interest
- **Dr. Julius Elefante:** No conflicts of interest
- **Dr. Alana Hirsh:** No conflicts of interest
- **Dr. Sasha Langille-Rowe** No conflicts of interest

Dr. Paxton Bach –

Clinical Assistant Professor, Department of Medicine, UBC

Addiction Medicine Physician, St. Paul's Hospital

Co-Medical Director, British Columbia Centre on Substance Use



THE UNIVERSITY OF BRITISH COLUMBIA

Continuing Professional Development

Faculty of Medicine

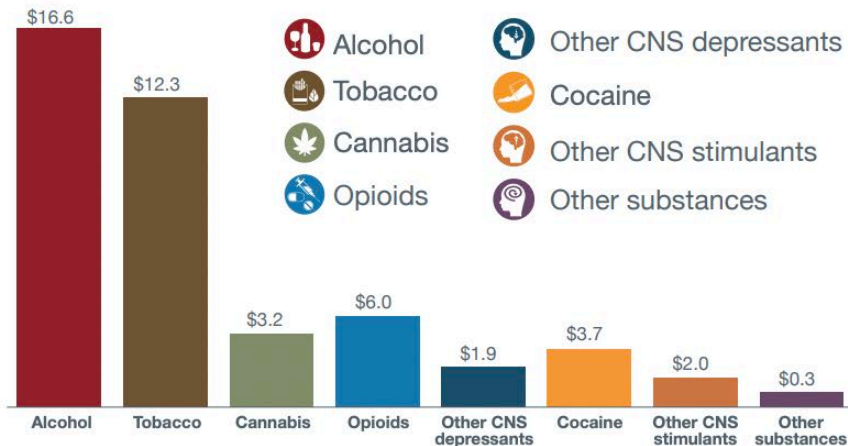
In 2017, substance use cost Canadians a total of

\$46 BILLION

Which amounts to almost \$1,260 for every Canadian regardless of age



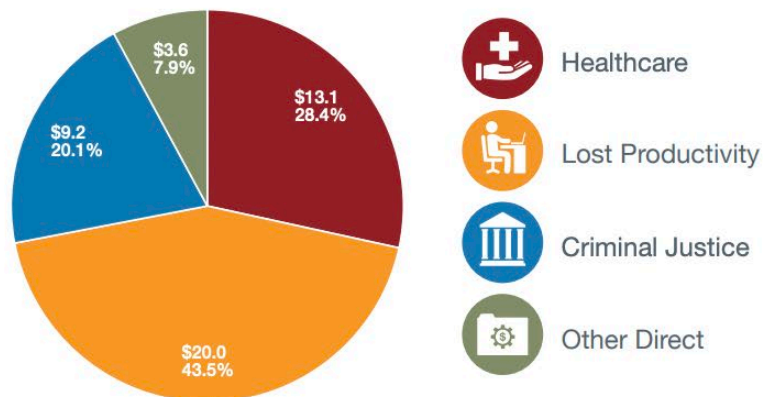
The cost of substance use (in billions)



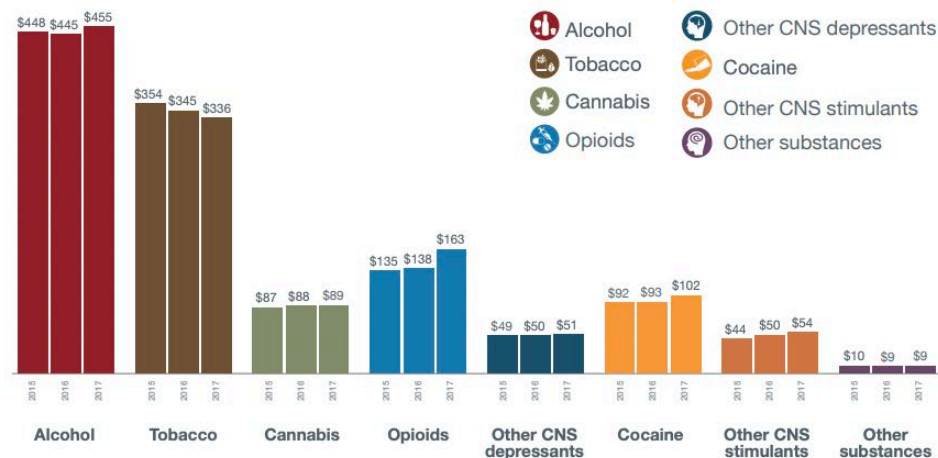
63% of total costs are due to alcohol and tobacco



Overall costs attributable to substance use in billions



Per person costs attributed to substance use



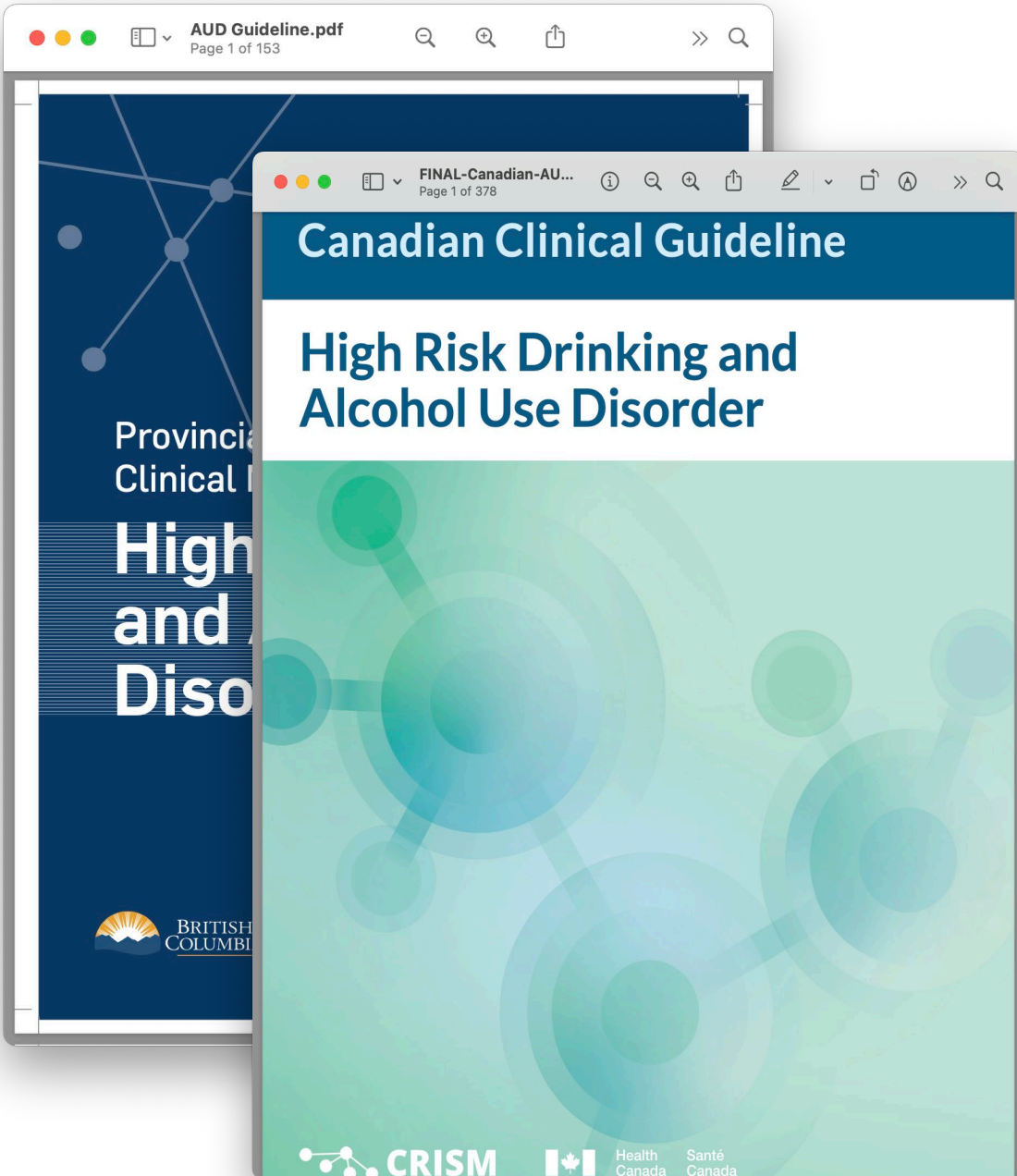


- **There are more hospitalizations for alcohol than for heart attacks.** In 2015–2016, there were about 77,000 hospitalizations entirely caused by alcohol compared with about 75,000 for heart attacks.

Alcohol Use Disorder in Canada

	Lifetime (%)	12-month (%)
Substance use disorder	21.6	4.4
Alcohol abuse or dependence	18.1	3.2
Cannabis abuse or dependence	6.8	1.3
Other drug abuse or dependence (excluding cannabis)	4.0	0.7

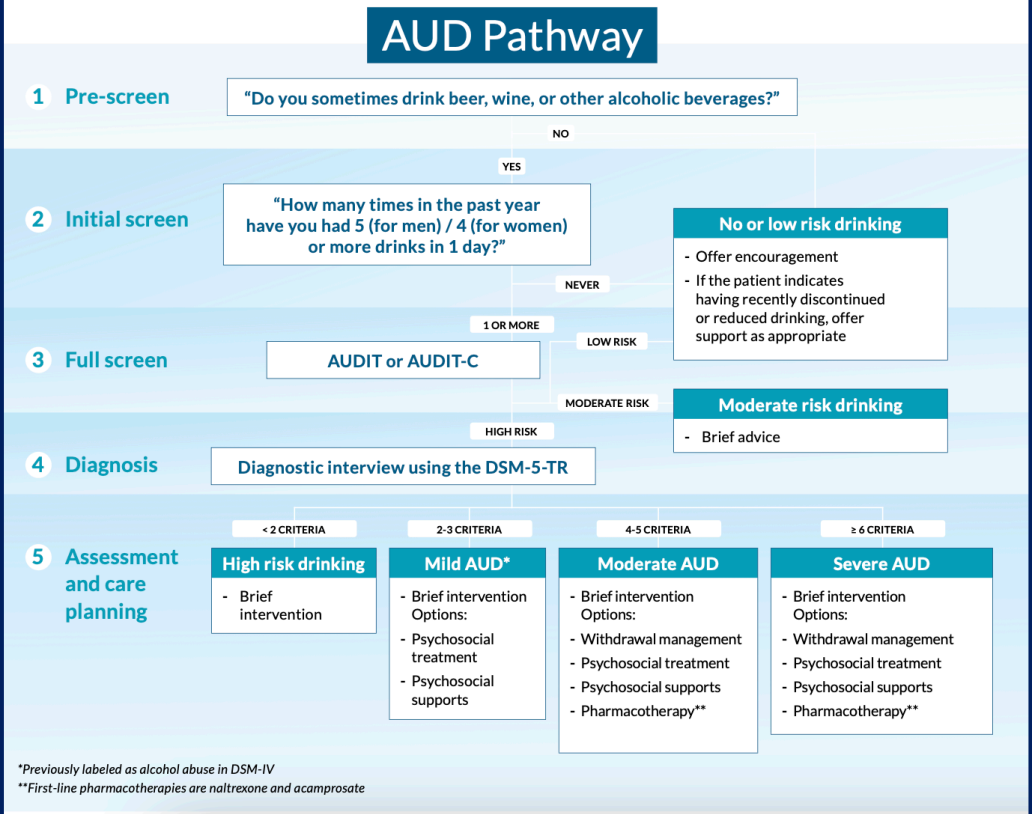
*DSM-IV diagnoses



Appendix 2: Screening and Diagnosis


Universal alcohol use screening of adult and youth patients has a significant role in health promotion, as the identification of high-risk alcohol use facilitates the prevention of the wide range of alcohol-related conditions as well as AUD. This appendix provides an instructive overview of the screening and diagnosis process as depicted in Figure 1.

Figure 1. Screening, Diagnosis, and Treatment Pathway



To reduce the risk of harm from alcohol, it is recommended that people living in Canada consider reducing their alcohol use.

Alcohol Consumption Per Week

<p>0 drinks per week Not drinking has benefits, such as better health and better sleep.</p>	No risk	<p>0 </p>
<p>1 to 2 standard drinks per week You will likely avoid alcohol-related consequences for yourself and others.</p>	Low risk	<p>1 </p> <p>2 </p>
<p>3 to 6 standard drinks per week Your risk of developing several different types of cancer, including breast and colon cancer, increases.</p>	Moderate risk	<p>3 </p> <p>4 </p> <p>5 </p> <p>6 </p>
<p>7 or more standard drinks per week Your risk of heart disease or stroke increases.</p> <p>Each additional standard drink Radically increases the risk of these alcohol-related consequences.</p>	Increasingly high risk	<p>7 </p> <p>8 </p> <p>+  ++</p>



24/7 ADDICTION
MEDICINE 
CLINICIAN SUPPORT LINE

Telephone consultation for physicians, nurse practitioners, nurses, mid-wives, and pharmacists providing addiction and substance use care.

Available 24/7, 365 days a year. More info at www.bccsu.ca/24-7.

CALL 778-945-7619

**Dr. Sasha Langille-Rowe, CCFP –
UBC Rural North West Residency Program Site Director
and Terrace Addiction Medicine Lead**

A Rural Family Medicine Perspective



THE UNIVERSITY OF BRITISH COLUMBIA

Continuing Professional Development

Faculty of Medicine

1. Screening can be quick

2. The relationship you have with your patients can make a difference

3. There are several effective interventions for the outpatient setting



Dr. Alana Hirsh–

Clinical Assistant Professor, UBC;

Physician at Kilala Lelum Urban Indigenous Health and Healing Cooperative

AUD Pharmacotherapy



THE UNIVERSITY OF BRITISH COLUMBIA

Continuing Professional Development

Faculty of Medicine

Recommendations 10 & 11

- 1st Line:
 - Naltrexone
 - Acamprosate
- 2nd Line:
 - Gabapentin
 - Topiramate

To prevent one individual from returning to any drinking, the number needed to treat (NNT) is:



Acamprosate

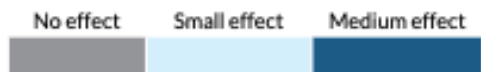
12 people must be treated
to prevent 1 relapse



Naltrexone

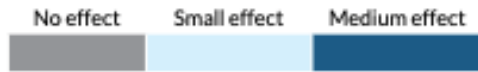
20 people must be treated
to prevent 1 relapse

Table 24. Comparison of AUD Pharmacotherapies



	Naltrexone	Acamprosate	Gabapentin	Topiramate
Efficacy				
Abstinence	Small effect	Medium effect	No effect	Medium effect
Heavy Drinking	Small effect	No effect	Medium effect	Medium effect
Craving	Small effect	No effect	No effect	No effect
Contraindications (▲) and Cautions (●)				
Opioid Use	▲		●	
Liver Failure / Hepatitis	▲			
Severe Kidney Impairment	●	▲	●	
Kidney Stones				▲
Narrow angle glaucoma				▲
Current alcohol use				
Safe to use while drinking?	✓	✓	✗	✓
Pre-treatment abstinence is beneficial	✓	✓	✓	

Table 24. Comparison of AUD Pharmacotherapies

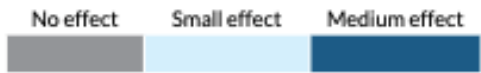


	Naltrexone	Acamprosate	Gabapentin	Topiramate
Efficacy				
Abstinence	Small effect	No effect	No effect	No effect
Heavy Drinking	Small effect	No effect	Small effect	No effect
Craving	Small effect	No effect	No effect	No effect
Contraindications (▲) and Cautions (●)				
Opioid Use	Medium effect (▲)		No effect (●)	
Liver Failure / Hepatitis	Medium effect (▲)			
Severe Kidney Impairment	Medium effect (●)	Small effect (▲)	No effect (●)	
Kidney Stones				Small effect (▲)
Narrow angle glaucoma				Small effect (▲)
Current alcohol use				
Safe to use while drinking?	Yes (✓)	Yes (✓)	No (X)	Yes (✓)
Pre-treatment abstinence is beneficial	Yes (✓)	Yes (✓)	Yes (✓)	

NALTREXONE

- Mu-opioid receptor antagonist, blocks euphoria associated with alcohol consumption.
- Treatment goal abstinence/Etoh reduction
- Start at 25mg OD for 3-4 days, titrate: to 50mg OD. Can go up to 150mg if needed.
- Some evidence for using PRN
- Monitor LFTs at 0,1,3,6 mo

Table 24. Comparison of AUD Pharmacotherapies

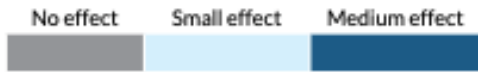


	Naltrexone	Acamprosate	Gabapentin	Topiramate
		Efficacy		
Abstinence		Medium effect	Small effect	Small effect
Heavy Drinking		No effect	Small effect	Small effect
Craving		No effect	Small effect	Small effect
	Contraindications (▲) and Cautions (●)			
Opioid Use	▲		●	
Liver Failure / Hepatitis	▲			
Severe Kidney Impairment	●	▲	●	
Kidney Stones				▲
Narrow angle glaucoma				▲
	Current alcohol use			
Safe to use while drinking?	✓	✓	✗	✓
Pre-treatment abstinence is beneficial	✓	✓	✓	

ACAMPROSATE

- Mechanism of action not well understood.
- Treatment goal abstinence (not reduced drinking)
- Start at maintenance dosage: 2 x 333mg tablets (666mg) TID
- Avoid in severe renal dysfunction

Table 24. Comparison of AUD Pharmacotherapies

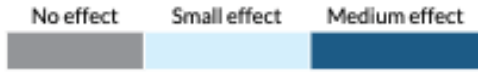


	Naltrexone	Acamprosate	Gabapentin	Topiramate
	Efficacy			
Abstinence	Small effect	Small effect	No effect	Small effect
Heavy Drinking	Small effect	Small effect	Medium effect	Small effect
Craving	Small effect	Small effect	No effect	Small effect
	Contraindications (▲) and Cautions (●)			
Opioid Use	▲		●	
Liver Failure / Hepatitis	▲			
Severe Kidney Impairment	●	▲	●	
Kidney Stones				▲
Narrow angle glaucoma				▲
	Current alcohol use			
Safe to use while drinking?	✓	✓	✗	✓
Pre-treatment abstinence is beneficial	✓	✓	✓	

GABAPENTIN

- Anticonvulsant, used for Etoh withdrawal, and off-label for AUD
- Dosing: Start at 100–300mg TID, titrate up to 1800mg daily
- Monitor for renal function, CNS side effects

Table 24. Comparison of AUD Pharmacotherapies



	Naltrexone	Acamprosate	Gabapentin	Topiramate
Efficacy				
Abstinence	Small effect	Medium effect	No effect	Medium effect
Heavy Drinking	Small effect	No effect	Small effect	Medium effect
Craving	Small effect	No effect	No effect	No effect
Contraindications (▲) and Cautions (●)				
Opioid Use	▲		●	
Liver Failure / Hepatitis	▲			
Severe Kidney Impairment	●	▲	●	
Kidney Stones				▲
Narrow angle glaucoma				▲
Current alcohol use				
Safe to use while drinking?	✓	✓	✗	✓
Pre-treatment abstinence is beneficial	✓	✓	✓	

TOPIRAMATE

- Anticonvulsant, off label for AUD
- Dosing: start at 25mg qhs, increase by 25mg weekly (divide bid) up to target of 50mg BID
- Monitor for CNS related side-effects
- Avoid in nephrolithiasis and narrow angle glaucoma



Dr. Julius Elefante–

Clinical Assistant Professor, UBC Faculty of Medicine

Addiction Medicine Consultation Liaison, Department of Psychiatry, St. Paul's Hospital



THE UNIVERSITY OF BRITISH COLUMBIA

Continuing Professional Development

Faculty of Medicine

WHAT THE 2023 GUIDELINES SAY

Table 2: Summary of recommendations

Recommendation		Strength of recommendation*	Certainty of evidence ¹⁵
Screening†			
12	Adult and youth patients should not be prescribed antipsychotics or SSRI antidepressants for the treatment of AUD.	Strong	Moderate
13	Prescribing SSRI antidepressants is not recommended for adult and youth patients with AUD and a concurrent anxiety or depressive disorder.	Strong	Moderate
14	Benzodiazepines should not be prescribed as ongoing treatment for AUD.	Strong	High

WHAT OTHER GUIDELINES HAVE SAID

The Canadian Network for Mood and Anxiety Treatments (CANMAT) task force recommendations for the management of patients with mood disorders and comorbid substance use disorders

SSRIs		
Escitalopram	Alcohol	Level 2 ^{46,47}
Fluoxetine	Alcohol	Level 2 ⁴⁸⁻⁵¹ ^a Level 2: negative ⁵²⁻⁵⁵
	Cannabis	^a Level 2: negative ^{52,56}
	Cocaine	Level 3: negative ^{57,58}
	Opiate	Add-on to methadone: level 2: negative ^{50,59}
Nefazodone	Alcohol	Level 1 ⁶⁰⁻⁶²
	Cocaine	Level 2: negative ⁶³
Sertraline	Alcohol	Level 2: negative ^{50,64-68} Level 2: naltrexone plus sertraline ⁶⁹
	Opiate	Add-on to methadone: level 2: negative ^{50,70}
OTHER		
Mirtazapine	Alcohol	Level 2 ^{36,71}

Substance	MDD
Alcohol	<i>First choice:</i> Mirtazapine Add-on naltrexone or alone Add-on naltrexone to sertraline ^a
	<i>Second choice:</i> Add-on disulfiram
	<i>Third choice:</i> Valproic acid Amitriptyline Desipramine Imipramine Escitalopram Memantine
	<i>Not recommended:</i> Fluoxetine ^b Lithium Sertraline Nefazodone (withdrawn from the market)

WHAT OTHER GUIDELINES HAVE SAID



—
Psychiatry

- 9 **Don't routinely prescribe antidepressants as first-line treatment for depression comorbid with an active alcohol use disorder without first considering the possibility of a period of sobriety and subsequent reassessment for the persistence of depressive symptoms.**



The concurrent management of psychiatric illness and alcohol use disorders requires evaluation of the role alcohol plays as a causative factor for depressive symptoms. Studies have found that response rates to antidepressants are higher when antidepressants are reserved for persistence of symptoms after a period of sobriety lasting from two to four weeks. Additionally, studies have demonstrated remission from depressive symptoms with sobriety in the absence of antidepressant treatment in a significant percentage of cases. Management of comorbid psychiatric illness and substance use disorders including alcohol dependence involves assessment and treatment delivered in a concurrent manner.



by
Canadian Academy of Child and Adolescent Psychiatry
Canadian Academy of Geriatric Psychiatry
Canadian Psychiatric Association
Last updated: September 2023

⌘ Navigating the nuances of the Canadian guideline's stance on selective serotonin reuptake inhibitors in concurrent alcohol use disorder and mood or anxiety disorders

Raymond Julius O. Elefante, Clara Lu and Paxton J. Bach

CMAJ March 18, 2024 196 (10) E348; DOI: <https://doi.org/10.1503/cmaj.150034-l>

- The complete AUD guideline provides an expanded rationale for recommendation 13:
 - “lack of high-quality evidence supporting the effectiveness of SSRIs for those with concurrent AUD and depression, a potentially higher risk of adverse events including worsening drinking outcomes, and research demonstrating a rapid reduction of depressive symptoms following a period of abstinence from alcohol use”



⌘ Navigating the nuances of the Canadian guideline's stance on selective serotonin reuptake inhibitors in concurrent alcohol use disorder and mood or anxiety disorders

Raymond Julius O. Elefante, Clara Lu and Paxton J. Bach

CMAJ March 18, 2024 196 (10) E348; DOI: <https://doi.org/10.1503/cmaj.150034-l>

- The AUD guidelines appropriately caution that their recommendation “does not address severe psychiatric conditions” and that, among patients who “demonstrated benefit from SSRI therapy, continued use of the medication could be considered with close monitoring of clinical response as well as unintended effects”



⌘ Navigating the nuances of the Canadian guideline's stance on selective serotonin reuptake inhibitors in concurrent alcohol use disorder and mood or anxiety disorders

Raymond Julius O. Elefante, Clara Lu and Paxton J. Bach

CMAJ March 18, 2024 196 (10) E348; DOI: <https://doi.org/10.1503/cmaj.150034-l>

- We agree that prescribers should pause before starting SSRIs in the context of AUD with comorbid anxiety or mood disorders
- However, in reviewing the evidence cited by the guidelines, we found that the risk of worsening drinking outcomes is inconsistent, and the cited randomized controlled trials have substantial limitations that prevent definitive conclusions



⌘ Navigating the nuances of the Canadian guideline's stance on selective serotonin reuptake inhibitors in concurrent alcohol use disorder and mood or anxiety disorders

Raymond Julius O. Elefante, Clara Lu and Paxton J. Bach

CMAJ March 18, 2024 196 (10) E348; DOI: <https://doi.org/10.1503/cmaj.150034-l>

- Treating people with concurrent alcohol use and mood/anxiety disorders is complex
 - Heterogeneous population
 - Difficulty attaining abstinence in the face of ongoing psychiatric symptoms
 - Refractory symptoms despite abstinence
 - Past success with pharmacotherapy



THANK YOU

JULIUS.ELEFANTE@UBC.CA



@RJELEFANTE

Comments,
questions and
suggestions