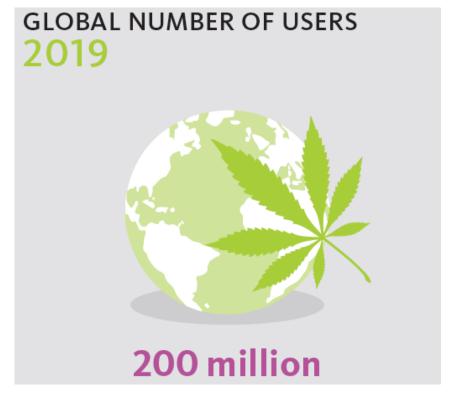
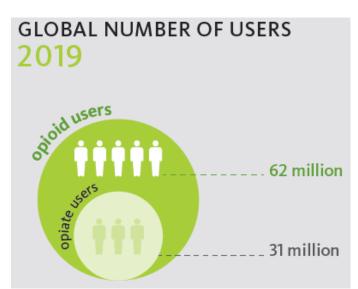


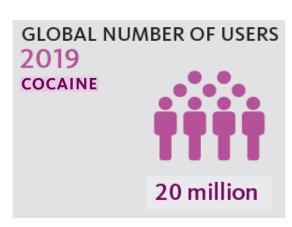


DISTURBO DA USO DI CANNABIS: IMPLICAZIONI CLINICHE E TERAPEUTICHE

Sarah Vecchio







4% OF THE GLOBAL POPULATION AGED 15–64 **18%** INCREASE BETWEEN 2010 AND 2019















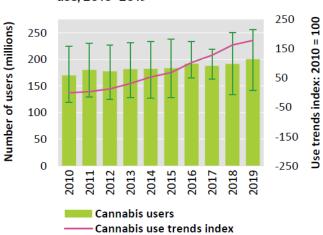


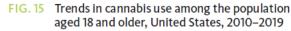


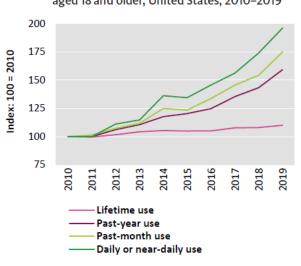


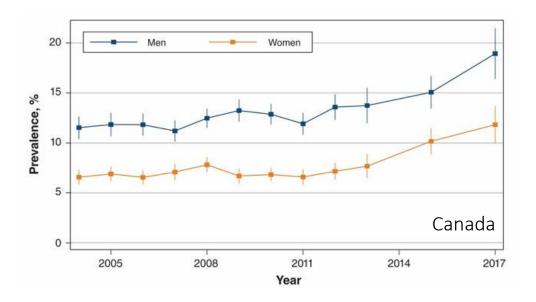


FIG. 11 Trends in the global number of people who use cannabis and reported trends in cannabis use, 2010–2019









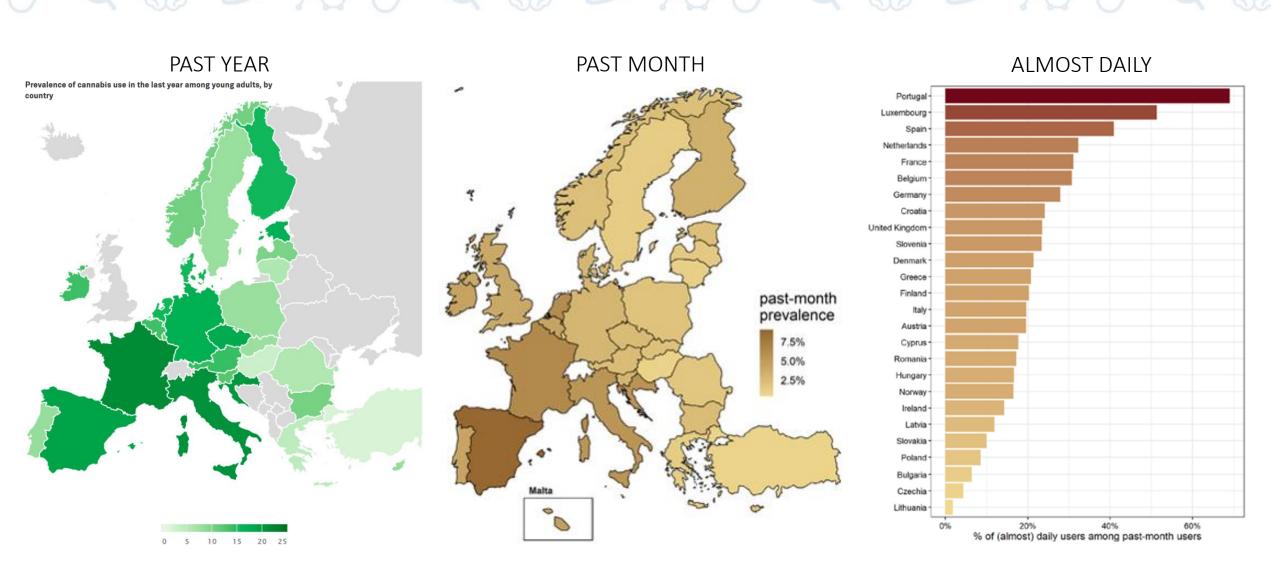
Cambiamenti negli indicatori relativi alla cannabis

Numero di paesi od ospedali che hanno segnalato un cambiamento dopo l'ultima raccolta di dati



Uso di cannabis nell'ultimo anno tra i giovani adulti (15-34 anni), 2018/19 e indagine precedente; prime richieste di trattamento correlate alla cannabis come droga primaria, 2018-19; accessi correlati alla cannabis negli ospedali Euro-DEN Plus, 2018-19.

World Drug Report 2021 (United Nations publication, Sales No. E.21.XI.8); Lowry DE, Corsi DJ. Trends and correlates of cannabis use in Canada: a repeated cross-sectional analysis of national surveys from 2004 to 2017. CMAJ Open. 2020 Jul 31;8(3):E487-E495; Osservatorio europeo delle droghe e delle tossicodipendenze (2021), Relazione europea sulla droga 2021: tendenze e sviluppi, Ufficio delle pubblicazioni dell'Unione europea, Lussemburgo



European Monitoring Centre for Drugs and Drug Addiction (2021), Cannabis: health and social responses; J. Manthey et al. Public health monitoring of cannabis use in Europe: prevalence of use, cannabis potency, and treatment rates. The Lancet Regional Health-Europe 2021

Tabella 1.1.1 - Spesa per consumi finali di droga per tipologia di sostanza stupefacente (valori in miliardi di euro)

	2016	2017	2018
Eroina	2,9	2,8	3,1
Cocaina	4,5	4,9	5,1
Cannabis	6,0	6,3	6,3
Altro	1,7	1,8	1,7
TOTALE	15,0	15,8	16,2

Fonte: ISTAT - Anni 2016 - 2018

Italy



Figura 3.1.2 - Uso di sostanze psicoattive nella vita

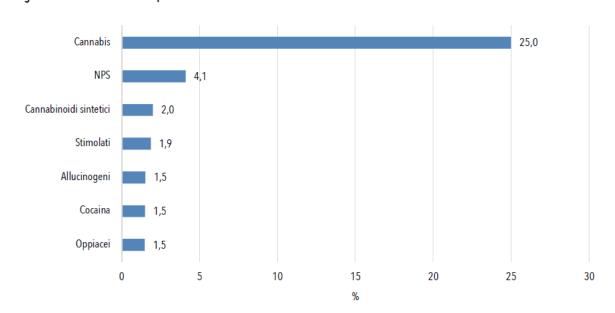
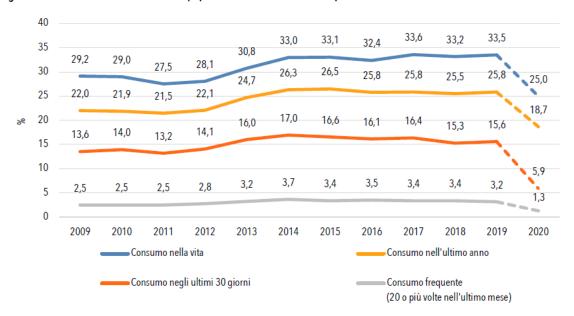


Figura 3.1.3 - Consumi di cannabis nella popolazione studentesca: trend percentuale



CANNABIS E COVID













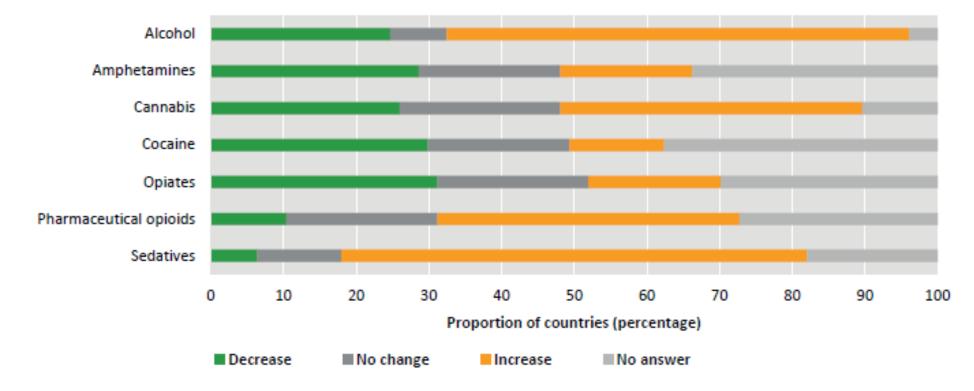












Source: Ali Farhoudian and others, "A global survey on changes in the supply, price and use of illicit drugs and alcohol, and related complications during the 2020 COVID-19 pandemic", MedRxiv (2020).

CANNABIS E COVID















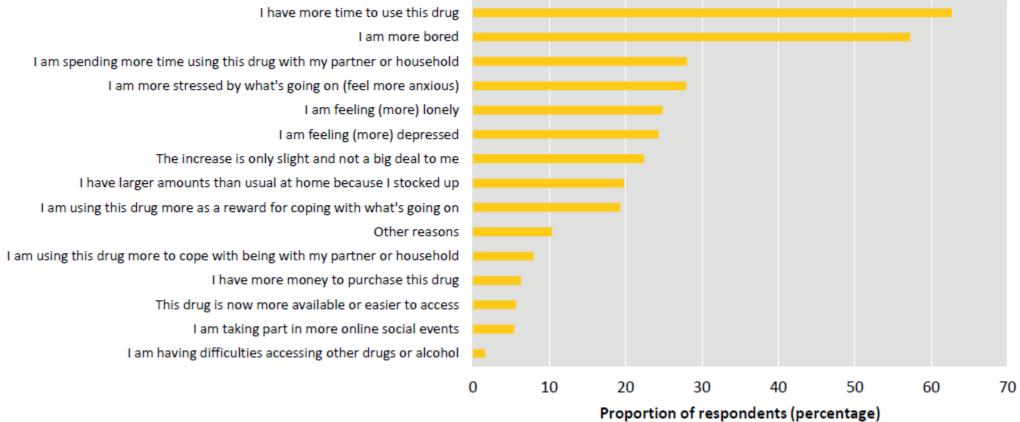








Reasons given by cannabis users for their increased use of cannabis after the onset of the COVID-19 pandemic, May-June 2020



Source: Global Drug Survey, "GDS COVID-19 special edition".













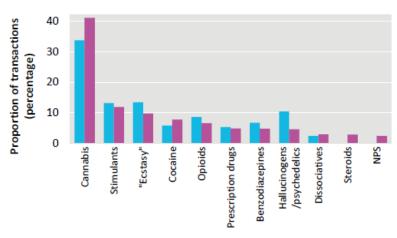






CANNABIS E DARKNET

FIG. 48 Distribution of drug transactions on 19 major darknet markets, by drug, 2011–mid-2017 and mid-2017–2020



- Former darknet markets (2011-mid-2017)
- Recent darknet markets (mid-2017-2020)



THE INTERNET: CLEAR WEB, DEEP WEB AND DARK WEB

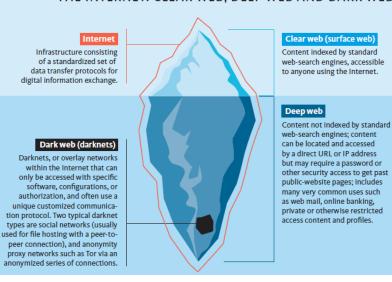


FIG. 60 Minimum drug sales on nine major global darknet markets, by drug type, 2019 and 2020

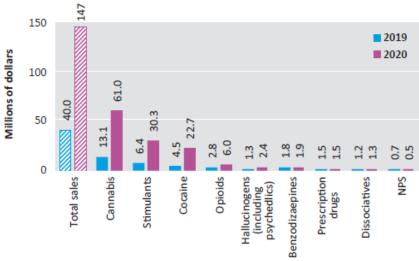
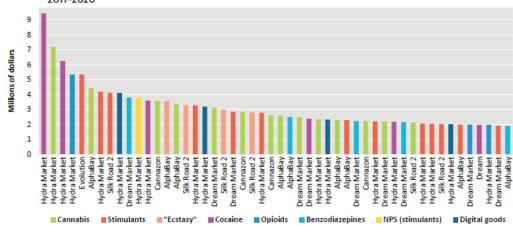


FIG. 51 Total minimum sales by the "top" 50 vendors (in terms of sales) operating on 19 major darknet markets, 2011–2020





INCREMENTO DELLA DIFFUSIONE DI CANNABIS













ELEVATA ACCETTABILITÀ SOCIALE

- Cannabis ad uso ricreazionale
- Cannabis ad uso terapeutico
- Cannabis light
- Industria della canapa (alimentare, tessile etc)

POLICIES DI DEPENALIZZAZIONE/LEGALIZZAZIONE

NUOVE MODALITÀ DI CONSUMO

RIDUZIONE DEI PREZZI







































CANNABIS: PERCEZIONE DEL RISCHIO













FIG. 26 Trends in cigarette smoking, cannabis use and risk perceptions related to smoking cigarettes and the use of cannabis among adolescents in the United States (10th grade) and Europe (aged 15-16), 1995-2019

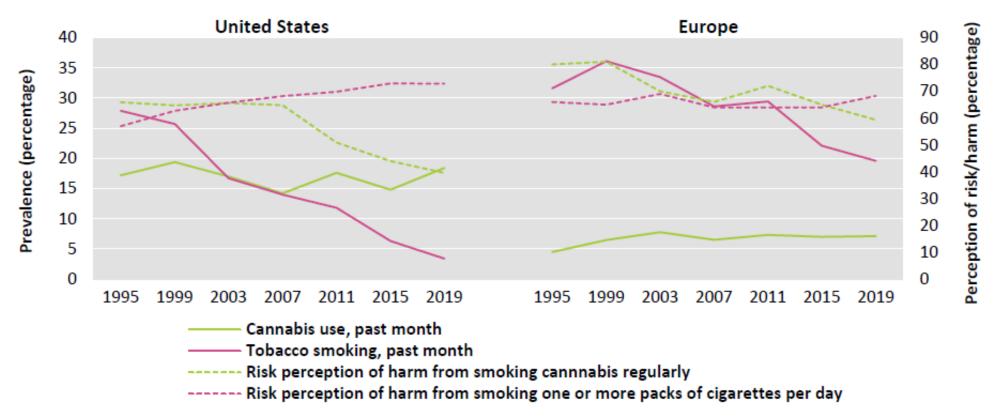














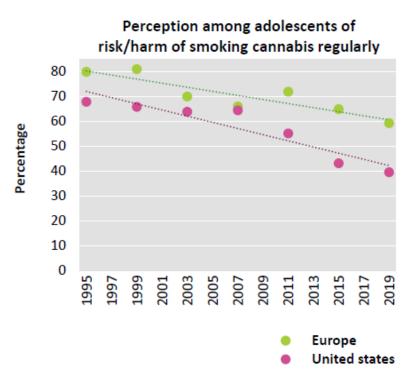


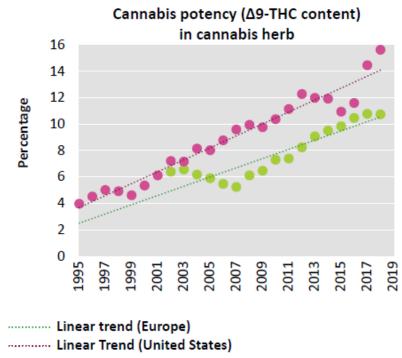


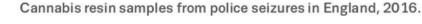


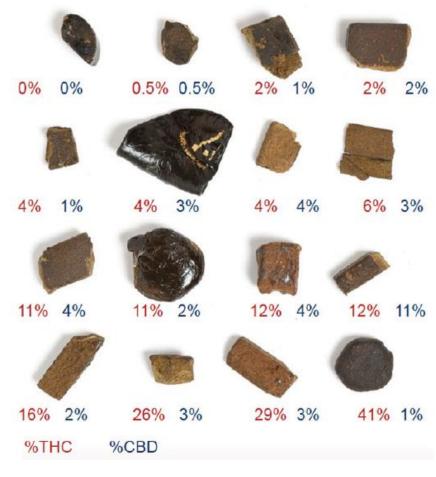


FIG. 27 Potency of cannabis and perception of risk from cannabis use among adolescents, Europe and United States, 1995–2019

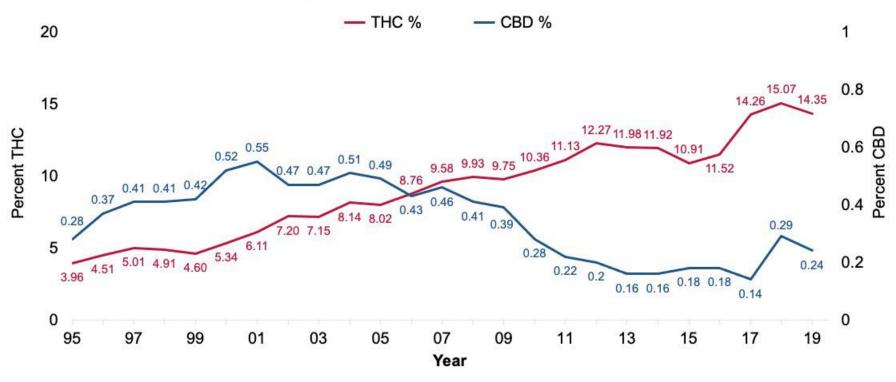










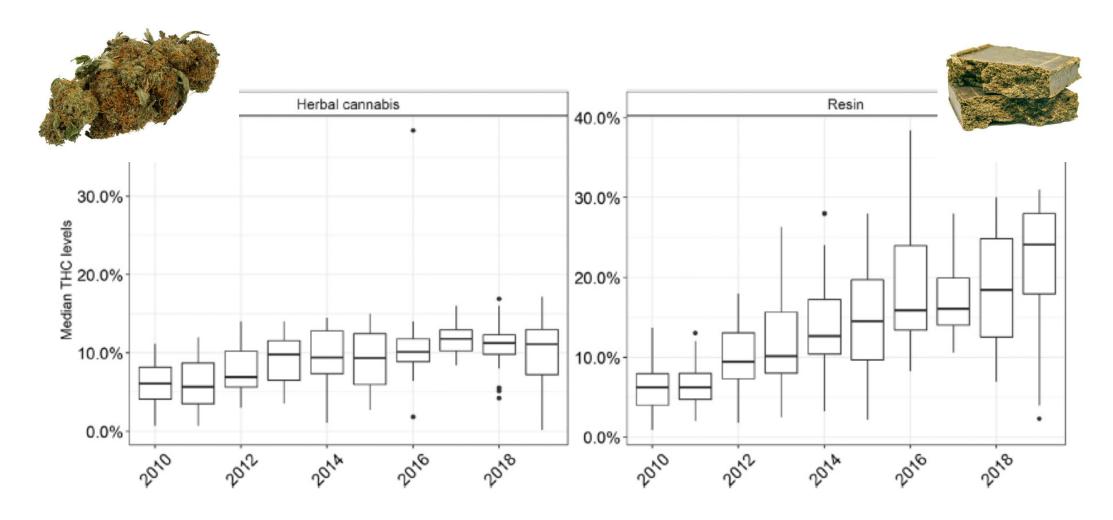












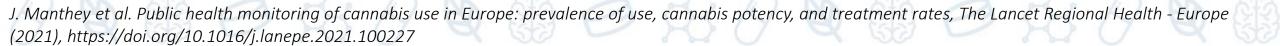
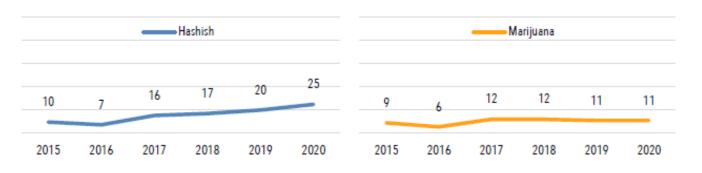
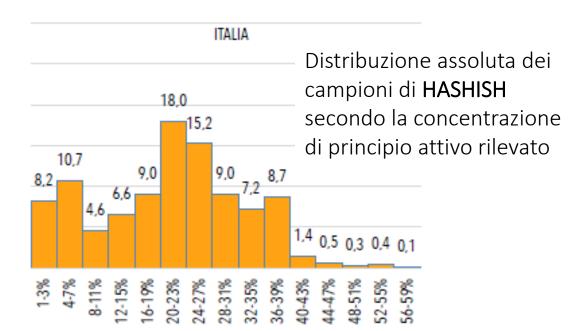
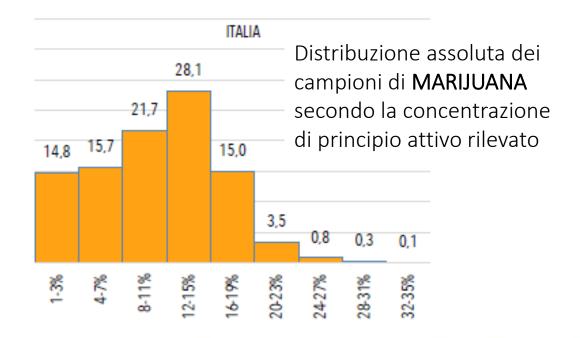


Figura 1.3.2 · Percentuali di principio attivo rilevate sulle sostanze stupefacenti analizzate

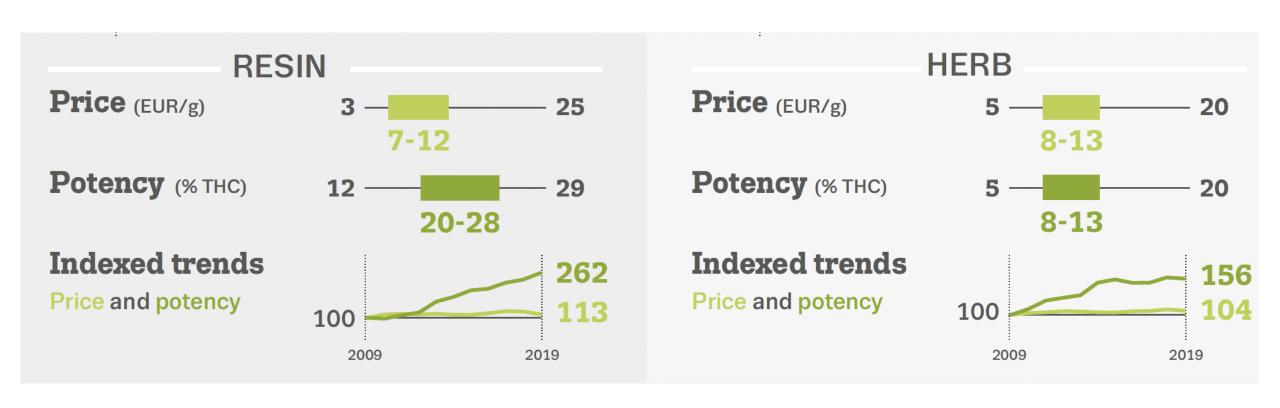








IL MERCATO DELLA CANNABIS





CANNABIS: PATTERN DI CONSUMO

















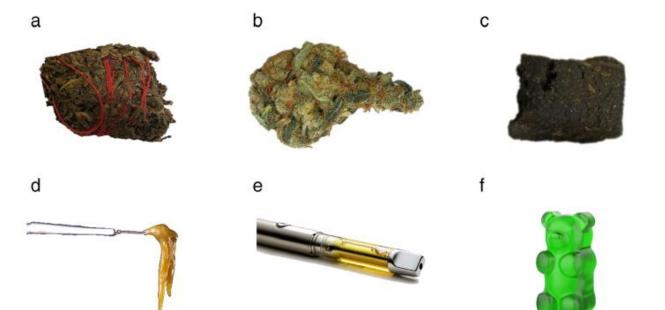


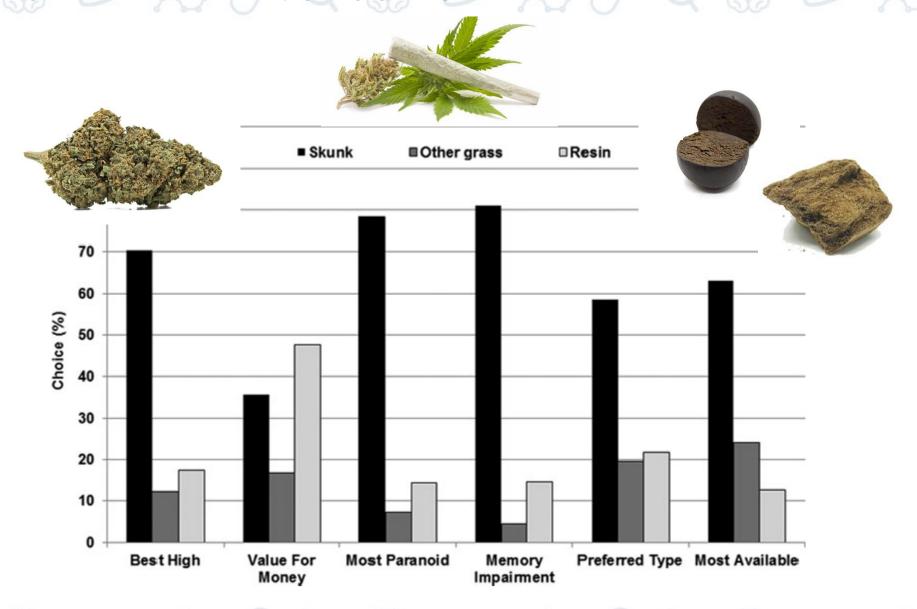
Table 2 Heterogeneity in methods of administration of cannabis products.

Method	Route	Combined with nicotine or tobacco
Joint	Inhaled, combusted	Yes/no
Pipe	Inhaled, combusted	Yes/no
Blunt	Inhaled, combusted	Yes
Bong	Inhaled, combusted	Yes/no
Dabbing	Inhaled, combusted	Yes/no
Vaporizer	Inhaled, vaporized	Yes/no
Vape pen	Inhaled, vaporized	Yes/no
Edible	Oral	No
Liquid	Oral	No

Country, year	Reference	Outdoor-grown herbal	Indoor-grown herbal	Resin	Concentrates
USA, 2017	[22]	9% THC, < 1% CBD	18% THC, $<1%$ CBD	46% THC, $<1%$ CBD	56% THC, < 1% CBD
Australia, 2010-12	[23]	15% THC, $<1%$ CBD	19% THC, $<1%$ CBD	_	_
UK, 2015-16	[24]	3% THC, $< 1%$ CBD	14% THC, $<1%$ CBD	6% THC, 2% CBD	78% THC, $<1%$ CBD
Netherlands, 2015	[25]	5% THC, $< 1%$ CBD	15% THC, $<1%$ CBD	18% THC, 8% CBD	_
France, 2016	[26]	_	_	23% THC, 4% CBD	_
Denmark, 2017	[27]	-	_	23% THC, 6% CBD	_

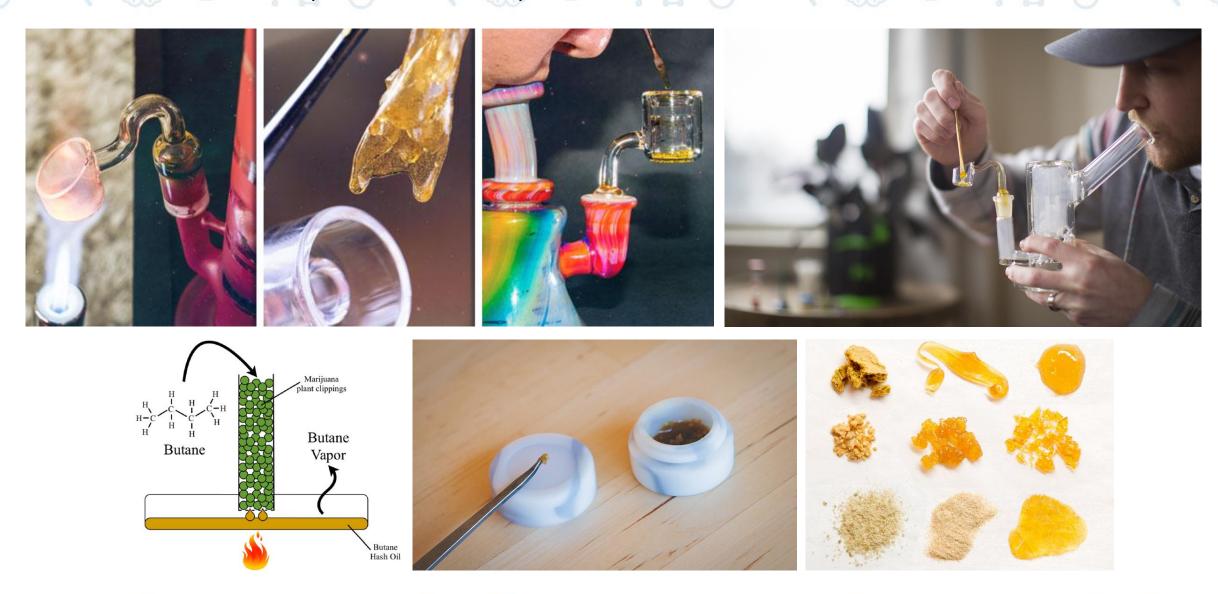


CANNABIS: PATTERN DI CONSUMO

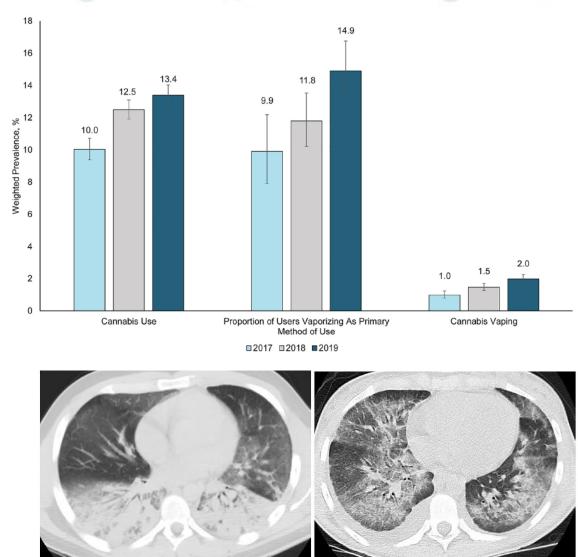




BUTANE HASH OIL (BHO, Wax...) e DABBING







Chadi N, Minato C, Stanwick R. Cannabis vaping: Understanding the health risks of a rapidly emerging trend. Paediatr Child Health. 2020 Jun;25(Suppl 1):S16-S20 Adapa S et al. Cannabis Vaping-Induced Acute Pulmonary Toxicity: Case Series and Review of Literature. J Investig Med High Impact Case Rep. 2020 Boakye E et al. Cannabis vaping among adults in the US: Prevalence, trends, and association with high-risk behaviors and adverse respiratory conditions. Prev Med. 2021

VAPING

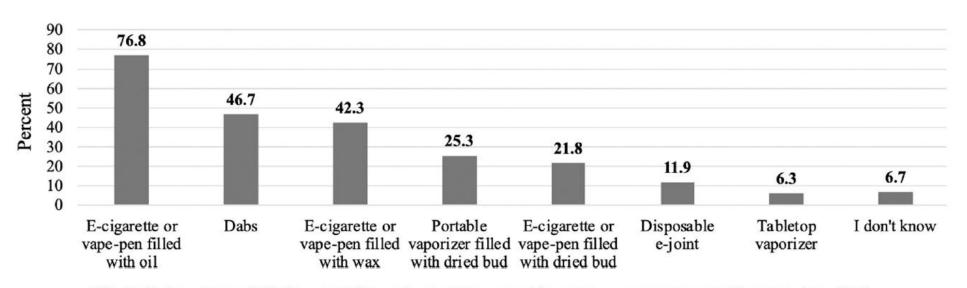
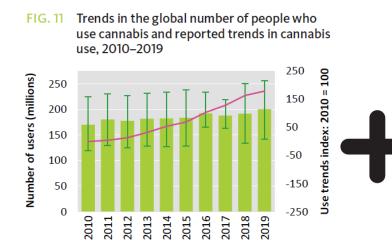


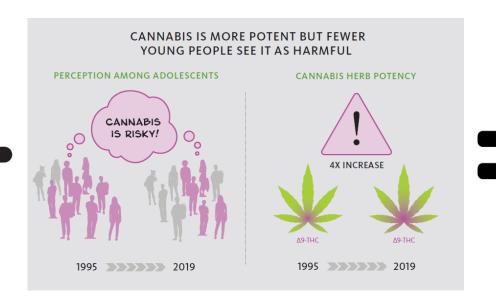
Fig. 2. Endorsement of lifetime modalities of vaporizing cannabis among past-month cannabis vapers (n = 524).



IMPATTO DELLE NUOVE TENDENZE?









INCREMENTO DELLA DIFFUSIONE

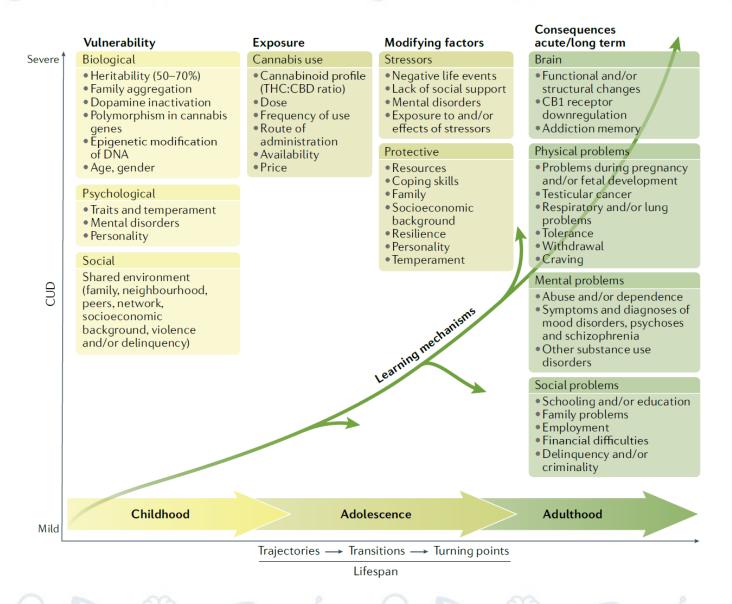
Cannabis use trends index

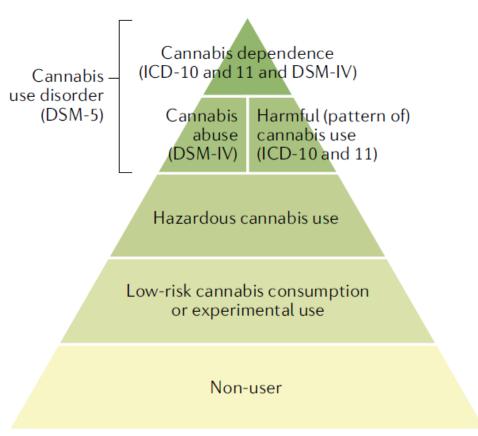
Cannabis users

RIDUZIONE PERCEZIONE DEL RISCHIO

INCREMENTO DELLA POTENZA INCREMENTO
DISTURBO DA USO
DI CANNABIS (CUD)

CUD: MODELLO MULTIFATTORIALE E GERARCHIA DIAGNOSTICA





DISTURBO DA USO DI CANNABIS (CUD)

Broad domain	DSM-5 CUD 'diagnostic criteria'4	ICD-11 Cannabis dependence 'description'5		
Impaired control	$\mbox{\bf 1}^a \mbox{Cannabis} \mbox{is taken} \mbox{in larger amounts or over} \\ \mbox{longer periods than intended}$	"Cannabis dependence is a disorder of regulation of cannabis use arising from repeated or continuous		
	2ª There is a persistent desire or unsuccessful attempts to cut down or control cannabis use	use of cannabis. The characteristic feature is a strong internal drive to use cannabis, which is manifested by impaired ability to control use"		
	3ª A great deal of time spent in activities necessary to obtain cannabis, use cannabis or recover from its effects	"increasing priority given to use over other activities"		
	4 Craving, or a strong desire or urge to use cannabis	"These experiences are often accompanied by a subjective sensation of urge or craving to use cannabis."		
Increasing priority resulting in	5 Recurrent cannabis use resulting in a failure to fulfil major role obligations at work, school or home	"and persistence of use despite harm or negative consequences."		
social and physical risk	6 Continued cannabis use despite having persistent or recurrent social or interpersonal problems caused or exacerbated by the effects of cannabis	"and persistence of use despite harm or negative consequences."		
	7 ^a Important social, occupational, or recreational activities are given up or reduced because of cannabis use	"increasing priority given to use over other activities"		
	8 Recurrent cannabis use in situations in which it is physically hazardous	"and persistence of use despite harm or negative consequences."		
	9ª Cannabis use is continued despite knowledge of having a persistent or recurrent physical or psychological problem that is likely to have been caused or exacerbated by cannabis	"and persistence of use despite harm or negative consequences."		
Physiological dependence	10 ^a Tolerance, as evidenced by a markedly diminished effect	"Physiological features of dependence may also be present, including tolerance to the effects of		
	11ª Withdrawal syndrome, or drinking to prevent withdrawal	cannabis, withdrawal symptoms following cessation or reduction in use of cannabis, or repeated use of cannabis or pharmacologically similar substances to prevent or alleviate withdrawal symptoms."		

Un pattern problematico di uso di cannabis che porta a disagio o compromissione clinicamente significativi, come manifestato da almeno due delle seguenti condizioni, che si verificano entro un periodo di 12 mesi

Gravità CUD

- CUD lieve: 2–3 condizioni
- CUD moderato: 4–5 condizioni
- CUD grave: ≥6 condizioni

DSM-5 specificatori

- Remissione precoce: 3–12 mesi
- Remissione protratta: >12 mesi

CUD: ASTINENZA

Criteri diagnostici

- A. Cessazione dell'uso di cannabis che è stato pesante e prolungato.
- B. Tre (o più) dei seguenti segni e sintomi, che si sviluppano approssimativamente entro 1 settimana dopo il Criterio A:
 - 1. Irritabilità, rabbia, aggressività
 - 2. Nervosismo, ansia
 - 3. Difficoltà del sonno (per es., insonnia, sogni inquietanti)
 - 4. Diminuzione dell'appetito o perdita di peso
 - 5. Irrequietezza
 - 6. Umore depresso
 - 7. Almeno uno dei seguenti sintomi fisici causa malessere significativo: dolori addominali, instabilità/tremori, sudorazione, febbre, brividi, cefalea
- C. I segni o sintomi causano disagio clinicamente significativo o compromissione del funzionamento in ambito sociale, lavorativo o in altre aree importanti.
- D. I segni o sintomi non sono attribuibili a un'altra condizione medica e non sono meglio spiegati da un altro disturbo mentale.

Criteri diagnostici

- A. Recente uso di cannabis.
- B. Comportamento problematico clinicamente significativo o cambiamenti psicologici (per es., coordinazione motoria compromessa, euforia, ansia, sensazione di rallentamento del tempo, capacità critica compromessa, isolamento sociale) che si sviluppano durante, o subito dopo, l'uso della cannabis.
- C. Due (o più) dei seguenti segni o sintomi, che si sviluppano entro 2 ore dall'uso della cannabis:
 - 1. Iperemia congiuntivale.
 - 2. Aumento dell'appetito.
 - 3. Secchezza delle fauci.
 - 4. Tachicardia.
- D. I segni o sintomi non sono attribuibili a un'altra condizione medica e non sono meglio spiegati da un altro disturbo mentale, compresa intossicazione da altra sostanza.

Specificare se:

Con alterazioni percettive: Allucinazioni con esame di realtà integro o illusioni uditive, visive o tattili che si verificano in assenza di un delirium.



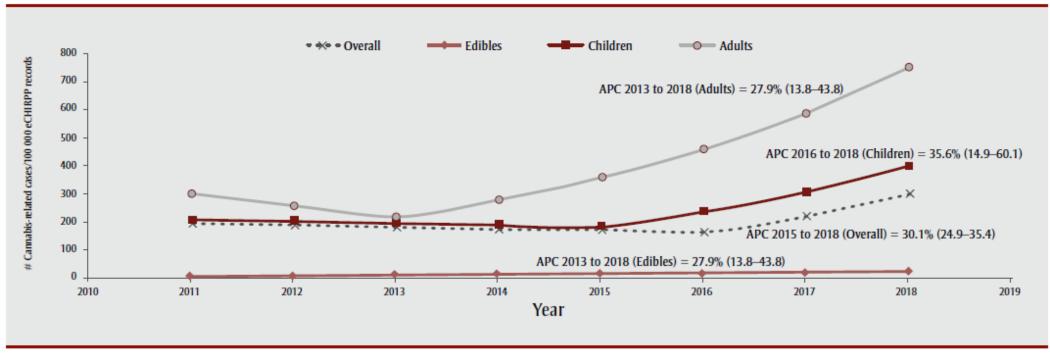








Time trend of cannabis-related cases presenting to emergency departments, children, adults and overall cases, eCHIRPP, 2011 to 2018^a



Abbreviations: APC, annual percent change; eCHIRPP, electronic Canadian Hospitals Injury Reporting and Prevention Program.



^a Records for 2019 were suppressed due to varying entry times in eCHIRPP.





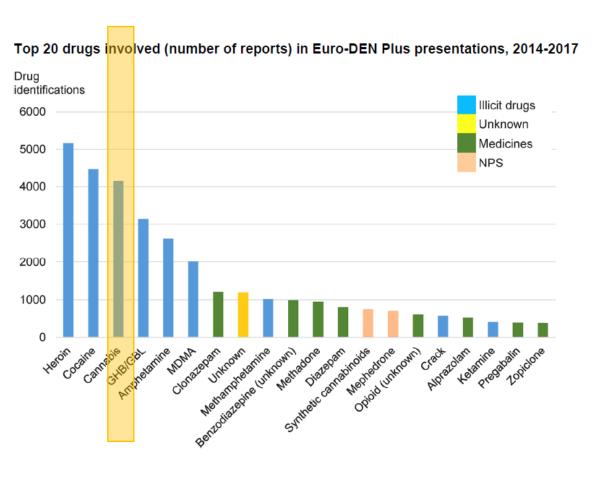


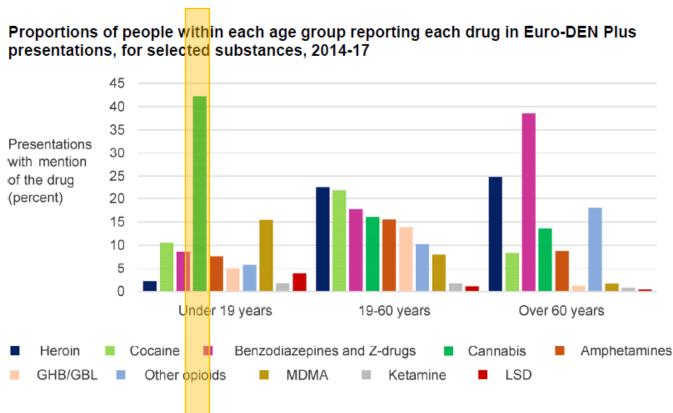






























Most common clinical features in the cases with cannabis alone (N = 186).

Clinical feature	Occurrence as the only clinical feature	Occurrence with other clinical features	Total occurrence
Cardiovascular symptoms			
Tachycardia (HR ≥ 100 bpm) on arrival	6	60	66
Palpitations	3	44	47
Chest pain	2	25	27
Nausea / vomiting	5	43	48
Anxiety	1	41	42
Dizziness	0	27	27
Impaired consciousness (GCS 8-14/"drowsy") on arrival	5	16	21
Agitation / aggression	1	18	19
Respiratory symptoms			
Dyspnoea	2	17	19
Hyperventilation	0	17	17
Panic attack	0	14	14
Psychosis	1	11	12
Mydriasis	0	11	11
Seizures	2	7	9
Hallucinations	0	7	7

Number of patients presenting with the clinical features in descending order of frequency. HR: heart rate; bpm: beats per minute; GCS: Glasgow Coma Score.

Substances reported or analytically detected in combination with cannabis (n).

alcohol 238 219 cocaine 121 77 amphetamine/methamphetamine 45 39 benzodiazepines/sedatives 40 72 MDMA/ecstasy 30 72 heroin 23 21 2 LSD 21 2 2 opioids other than heroin or methadone 12 12 12 methadone 9 22 2 2 opiates 31 1 31 31 31 31 32 32 32 32 32 32 32 32 33 33 33 33 33 33 33 33 33 33 33 33 33 33 34 33 34 <td< th=""><th>Substances</th><th>Reported</th><th>Analytically detected</th></td<>	Substances	Reported	Analytically detected
amphetamine/methamphetamine 45 39 benzodiazepines/sedatives 40 72 MDMA/ecstasy 30 heroin 23 LSD 21 2 opioids other than heroin or methadone 12 methadone 9 22 opiates 31 methylphenidate 7 1 psychedelic mushrooms 5 neuroleptics 4 3 ketamine 3 1 poppers 2 antidepressants 2 6 antihistamines 1 laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil 1 bupropion 1 tizanidine clomethiazole	alcohol	238	219
benzodiazepines/sedatives 40 72 MDMA/ecstasy 30 heroin 23 LSD 21 2 opioids other than heroin or methadone 12 methadone 9 22 opiates 31 methylphenidate 7 1 psychedelic mushrooms 5 neuroleptics 4 3 ketamine 3 1 poppers 2 antidepressants 2 6 antihistamines 1 laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil bupropion 1 tizanidine clomethiazole 1	cocaine	121	77
MDMA/ecstasy 30 heroin 23 LSD 21 2 opioids other than heroin or methadone 12 methadone 9 22 opiates 31 methylphenidate 7 1 psychedelic mushrooms 5 neuroleptics 4 3 ketamine 3 1 poppers 2 6 antidepressants 2 6 antihistamines 1 1 laughing gas 1 4 dextromethorphan 1 1 testosterone 1 1 sildenafil 1 1 bupropion 1 1 tizanidine 1 1 clomethiazole 1 1	amphetamine/methamphetamine	45	39
heroin 23 LSD 21 2 opioids other than heroin or methadone 12 methadone 9 22 opiates 31 methylphenidate 7 1 psychedelic mushrooms 5 neuroleptics 4 3 ketamine 3 1 poppers 2 6 antidepressants 2 6 antihistamines 1 1 laughing gas 1 4 dextromethorphan 1 1 testosterone 1 1 sildenafil 1 1 bupropion 1 1 tizanidine 1 1 clomethiazole 1 1	benzodiazepines/sedatives	40	72
LSD 21 2 opioids other than heroin or methadone 12 methadone 9 22 opiates 31 methylphenidate 7 1 psychedelic mushrooms 5 1 neuroleptics 4 3 ketamine 3 1 poppers 2 6 antidepressants 2 6 antihistamines 1 1 laughing gas 1 4 dextromethorphan 1 1 testosterone 1 1 sildenafil 1 1 bupropion 1 1 tizanidine 1 1 clomethiazole 1 1	MDMA/ecstasy	30	
opioids other than heroin or methadone methadone 9 22 opiates 31 methylphenidate 7 1 psychedelic mushrooms 5 neuroleptics 4 3 ketamine 3 1 poppers 2 antidepressants 2 6 antihistamines 1 laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil bupropion 1 tizanidine clomethiazole 1 laughing clomethiazole	heroin	23	
methadone 9 22 opiates 31 methylphenidate 7 1 psychedelic mushrooms 5 neuroleptics 4 3 ketamine 3 1 poppers 2 antidepressants 2 6 antihistamines 1 laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	LSD	21	2
opiates 31 methylphenidate 7 1 psychedelic mushrooms 5 neuroleptics 4 3 ketamine 3 1 poppers 2 antidepressants 2 6 antihistamines 1 laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	opioids other than heroin or methadone	12	
methylphenidate 7 1 psychedelic mushrooms 5 neuroleptics 4 3 ketamine 3 1 poppers 2 antidepressants 2 6 antihistamines 1 laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	methadone	9	22
psychedelic mushrooms neuroleptics 4 3 ketamine 3 1 poppers 2 antidepressants 2 6 antihistamines 1 laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	opiates		31
neuroleptics 4 3 ketamine 3 1 poppers 2 6 antidepressants 2 6 antihistamines 1 1 laughing gas 1 1 dextromethorphan 1 1 testosterone 1 1 sildenafil 1 1 bupropion 1 1 tizanidine 1 1 clomethiazole 1 1	methylphenidate	7	1
ketamine 3 1 poppers 2 6 antidepressants 2 6 antihistamines 1 1 laughing gas 1 1 dextromethorphan 1 1 testosterone 1 1 sildenafil 1 1 bupropion 1 1 tizanidine 1 1 clomethiazole 1 1	psychedelic mushrooms	5	
2	neuroleptics	4	3
antidepressants 2 6 antihistamines 1 laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	ketamine	3	1
antihistamines 1 laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	poppers	2	
laughing gas 1 dextromethorphan 1 testosterone 1 sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	antidepressants	2	6
dextromethorphan 1 testosterone 1 sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	antihistamines	1	
testosterone 1 sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	laughing gas	1	
sildenafil 1 bupropion 1 tizanidine 1 clomethiazole 1	dextromethorphan	1	
bupropion 1 tizanidine 1 clomethiazole 1	testosterone	1	
tizanidine 1 clomethiazole 1	sildenafil	1	
clomethiazole 1	bupropion	1	
	tizanidine	1	
	clomethiazole	1	
caffeine 1	caffeine	1	
melatonin (self-reported as "melanin") 1	melatonin (self-reported as "melanin")	1	
self-reported "smileys" 1	self-reported "smileys"	1	
self-reported "synthetic drugs" 1	self-reported "synthetic drugs"	1	
unknown substance 10	unknown substance	10	







CANNABINOID HYPEREMESIS SYNDROME (CSH)



Frequent vomiting from use of cannabis (pot/marijuana)

CHS - Cannabinoid Hyperemesis Syndrome



If you are vomiting more than 5 times per day, for a day or longer, immediately contact your health care provider or go to the nearest emergency department.



If you continue to experience CHS symptoms and are not improving as expected, talk with your health care provider.

What is CHS?

People with cannabinoid hyperemesis syndrome (CHS) experience persistent nausea and stomach pain that can lead to frequent and repetitive vomiting and weight loss. Frequent use of cannabis (at least once a week for more than a year) can increase the risk.

Treatment

- People with CHS may take a hot shower or bath to help temporarily relieve their symptoms. Caution should be taken as these can cause dehydration or scald/burn the skin.
- Capsaicin cream (brand name Zostrix) may be prescribed to help reduce the symptoms of CHS. The cream is to be applied to your stomach, back, or arms.
- · Other medications may be prescribed to relieve nausea and vomiting.
- If you go to the emergency department, let your health care provider know if you are using cannabis to manage a medical condition. Your health care provider may do some blood tests. You may be given fluids intravenously to keep you hydrated and help you feel better.

Prevention

Choosing not to use cannabis is the only way to completely avoid CHS. If CHS does occur, the symptoms will usually resolve within 2 weeks after you stop using cannabis. However, if you've had CHS once, even a small amount of cannabis can cause the symptoms to come back. If this happens, talk to your health care provider about treatment options or programs for stopping cannabis use.

Reduce your risk of CHS by following "Cannabis & Your Health: 10 WAYS to Reduce Risks When Using". https://bit.ly/Ircugphac

For more information visit: www.ccsa.ca/cannabis

To access a PDF of this handout visit: safemedicationuse.ca/tools_resources/tips.html

<u>i/m</u>p

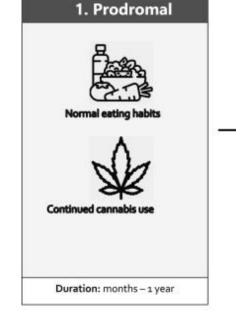


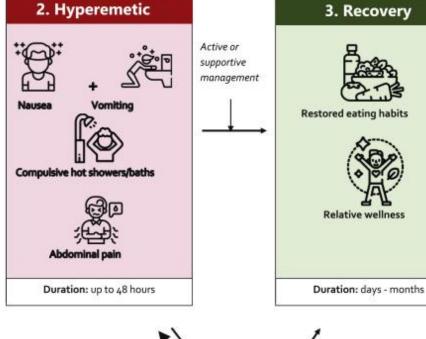














https://www.ccsa.ca/public-education

CANNABINOID HYPEREMESIS SYNDROME (CSH)











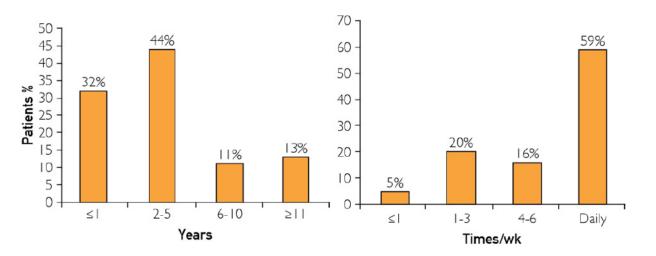


Table 1 Rome IV criteria for CHS diagnosis

Category	Features
Essential	Stereotypical episodic vomiting resembling CVS in terms of onset, duration, and frequency Presentation after prolonged, excessive cannabis use Relief of vomiting episodes by sustained cessation of cannabis use
Supportive remarks	May be associated with pathologic bathing behavior (prolonged hot baths or showers)

CHS = cannabinoid hyperemesis syndrome; CVS = cyclic vomiting syndrome.

Mechanism	GRADE rating
The emetogenic and anti-emetic effects of Δ9-THC and its analogs are mediated through CB-1 receptors (CB1r) and thus underlie the syndrome of CHS [106, 135]	Very low
Cannabinoids may bind to CB-1 receptors in the gastrointestinal tract and decrease GI motility and gastric emptying, which may override brainstem-mediated antiemetic effects and precipitate hyperemesis [9, 92, 95, 132]	Very low
Chronic cannabis use may lead to paradoxical and plastic changes in expression and downstream effects of cannabinoid receptors [133]	Very low
Chronic cannabis use leads to desensitization and downregulation of CB1 receptors that ordinarily have peripheral antiemetic effects, causing rebound vomiting and spasmodic pain that abates with abstinence and corresponding recovery of CB-1 receptor activity [98, 136, 185]	Very low
In chronic cannabis users, cannabinoid metabolites may accumulate in the brain and fatty tissues inducing a toxic effect [90, 94]	Very low
CHS may be caused by a non-THC, cannabinoid-like structure within <i>Cannabis sativa</i> , such as cannabidiol [96, 186]	Very low
Patients susceptible to developing CHS may have genetic variation in their metabolic enzymes resulting in toxic levels of cannabinoid metabolites [131]	Very low
Δ9-THC may act as a partial agonist on CB1 receptors and thus relatively antagonize the effects of full endogenous agonists on these receptors, thus precipitating sudden withdrawal and hyperemesis in sensitive patients [97, 105]	Very low
THC causes dilation of splanchnic vasculature, resulting in CHS. Hot bathing leads to peripheral venodilation and shunts blood away from the splanchnic bed, resulting in symptom improvement [102, 137]	Very low

Simonetto DA et al. Cannabinoid hyperemesis: a case series of 98 patients. Mayo Clin Proc. 2012 Feb;87(2):114-9 Zhu JW et al. Diagnosis and Acute Management of Adolescent Cannabinoid Hyperemesis Syndrome: A Systematic Review. J Adolesc Health. 2021 Feb;68(2):246-254

Sorensen CJ et al. Cannabinoid Hyperemesis Syndrome: Diagnosis, Pathophysiology, and Treatment-a Systematic Review. J Med Toxicol. 2017 Mar;13(1):71-87

CUD: IMPATTO A BREVE E LUNGO TERMINE











Table 1 Common clinical adverse effects associated with cannabinoids use

Psychiatric conditions	An increased risk of psychotic disorders following acute and repeated consumption of cannabis in vulnerable individuals and naïve users. 27,28,41–44 Anxiety and panic attacks following intoxication especially in naïve users. 38 Chronic use is associated with mood disturbances, mania, and depression. 36,37,39,40 Cannabis addiction and dependency. 9,13		
Cognitive and CNS alterations	Impairment of a wide range of cognitive functions following cannabis intoxication in a dose-relation manner. 38–42,44,45		
	Impaired cognitive function following cannabis consumption was associated with an increased risk of having a road accident. ^{31–35} Chronic use is associated with long-term brain functional and structural alterations. ^{31,32,45–50}		
Effects on recoiretery system	Acute cannabis consumption decreases airway resistance. 12		
Effects on respiratory system	Chronic cannabis use is associated with an increased risk for developing airway diseases and lung cancer. 59-66		
Effect on cardiovascular system	An increase of cardiovascular activity, increase of heart rate, and decrease of blood pressure. ⁶⁴ Several reports have described a temporal relationship between cannabis use and acute myocardial infarction, cardiomyopathy, and sudden cardiac death. ^{68–72}		

CUD E PATOLOGIE PSICHIATRICHE















Figure 2. Forest Plot Showing Adjusted Odds Ratio (OR) and 95% CIs for Depression and Anxiety in Young Adulthood According to Cannabis Use in Individual Studies

Figure 3. Forest Plot Showing Adjusted Odds Ratio (OR) and 95% CIs for Suicidal Ideations and Attempts According to Cannabis Use in Individual Studies

	OR	Favors Controls	Favors		OR	Favors Controls ! Favors
Study	(95% CI)	(Non-Cannabis	Cannabis	Study	(95% CI)	(Non-Cannabis Cannabis
Depression in young adulthood		Users)	Users	Suicide Ideations		Users) Users
Brook et al, ³⁴ 2002, United States	1.44 (1.08 to 1.91)		-	Fergusson et al, ⁴¹ 1996, New Zealand	1.40 (0.70 to 2.80)	
Brook et al, ¹⁶ 2011, United States and Puerto Rico	1.50 (0.90 to 3.20)		-	McGee et al, ⁴⁷ 2005, New Zealand	1.10 (0.58 to 2.07)	
Degenhardt et al, ³⁸ 2013, Australia	1.10 (0.60 to 1.90)	_		Weeks and Colman, ⁵⁷ 2016, Canada	1.74 (1.16 to 2.60)	
Gage et al, 44 2015, United Kingdom	1.30 (0.98 to 1.72)		-			≈
Georgiades and Boyle, 45 2007, Canada	1.48 (0.65 to 3.40)	_	 -	Pooled OR for all studies: Q = 1.49, df = 2 (P = .48); I = 0%	1.50 (1.11 to 2.03)	
Marmorstein and Iacono, 46 2011, USA	2.62 (1.22 to 5.65)		<u> </u>	Suicide attempts		
Silins et al, ¹⁰ 2014, Australia and New Zealand	1.02 (0.52 to 2.01)			Roberts et al, ⁵⁴ 2010, United States	4.81 (1.82 to 12.66)	
Pooled OR for all studies: Q=3.26, df=6 (P=.62); I ² =0%	1.37 (1.16 to 1.62)			Silins et al, ¹⁰ 2014, Australia and New Zealand	6.83 (2.04 to 22.90)	
Anxiety in young adulthood				Weeks and Colman, 57 2016, Canada	1.87 (1.09 to 3.22)	
Brook et al, ¹⁶ 2011, United States and Puerto Rico	1.60 (0.90 to 2.90)		-	Pooled OR for all studies: Q = 5.38, df = 2 (P = .07); I^2 = 61.3%	3.46 (1.53 to 7.84)	
Degenhardt et al, 38 2013, Australia	1.40 (0.84 to 2.50)	-	-		0.1	
Gage et al, ⁴⁴ 2015, United Kingdom	0.96 (0.75 to 1.24)	-	-		U.	OR (95% CI)
Pooled OR for all studies: Q = 3.26, df = 2 (P = .20); I2 = 42%	1.18 (0.84 to 1.67)	-	<u> </u>			OR (33% CI)
	0.1		1 10			

Gobbi G. Association of Cannabis Use in Adolescence and Risk of Depression, Anxiety, and Suicidality in Young Adulthood: A Systematic Review and Meta-analysis. JAMA Psychiatry. 2019 Apr 1;76(4):426-434

OR (95% CI)

CUD E PATOLOGIE PSICHIATRICHE

Cannabis Exposure

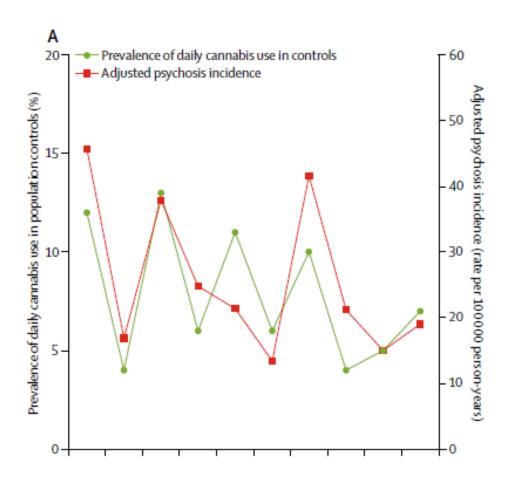
Psychosis risk distribution ES (95% CI) ∞ Psychotic symptoms Tien 1990 Tien 1990 1.85 (1.33, 2.57) Degenhardt 2001 Degenhart 2001 7.45 (3.99, 13.90) 7 Zammit 2002 Henquet 2005 5.16 (3.13, 8.50) Arseneault 2002 **Wiles 2006** 3.04 (1.41, 6.59) Henguet 2005 Miettunen 2008 4.67 (3.66, 5.96) 9 Wiles 2006 McGrath 2010 1.89 (1.32, 2.69) Miettunen 2008 McGrath 2010 Zammit 2011 4.36 (2.38, 7.99) 2 Zammit 2011 Arseneault 2002 4.29 (1.45, 12.70) R GAP data 2012 Subtotal 3.59 (2.43, 5.32) 4 Diagnosis of psychosis Zammit 2002 6.20 (4.19, 9.17) 3 GAP data 2012 4.38 (3.30, 5.81) **Subtotal** 5.07 (3.62, 7.09) 2 Overall effect 3.90 (2.84, 5.34) 16 0.0 0.2 0.6 8.0 1.0 0.4

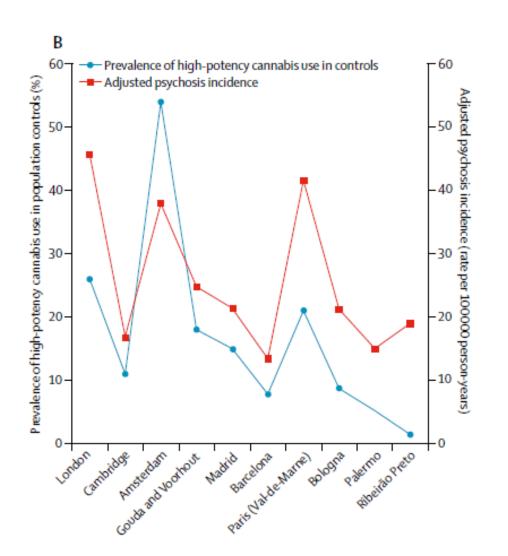
PATTERN DI CONSUMO E PATOLOGIE PSICHIATRICHE

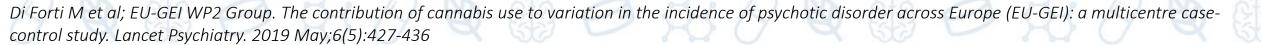






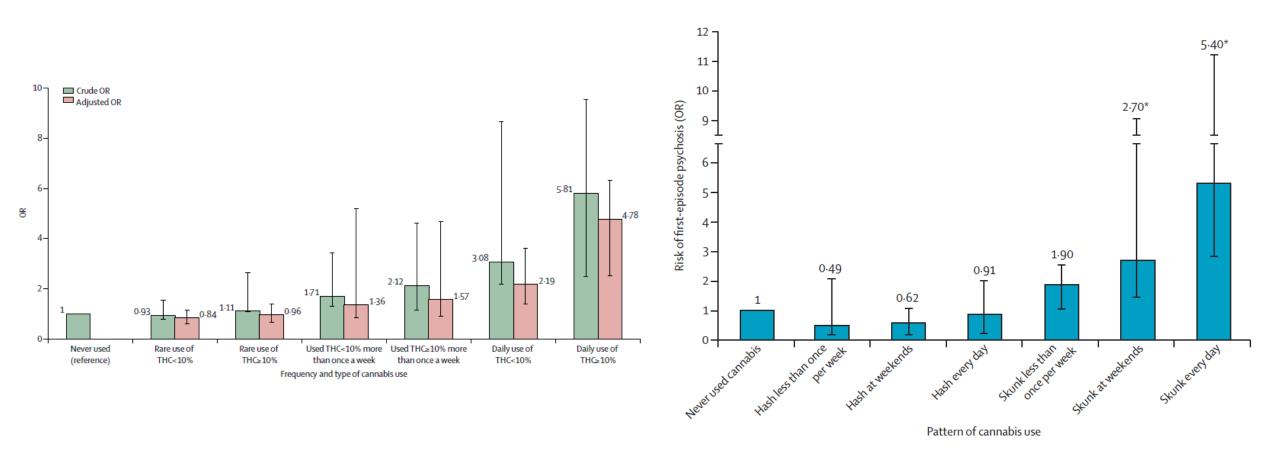






PATTERN DI CONSUMO E PATOLOGIE PSICHIATRICHE





Di Forti M et al; EU-GEI WP2 Group. The contribution of cannabis use to variation in the incidence of psychotic disorder across Europe (EU-GEI): a multicentre case-control study. Lancet Psychiatry. 2019 May;6(5):427-436; Di Forti M et al. Proportion of patients in south London with first-episode psychosis attributable to use of high potency cannabis: a case-control study. Lancet Psychiatry. 2015 Mar;2(3):233-8

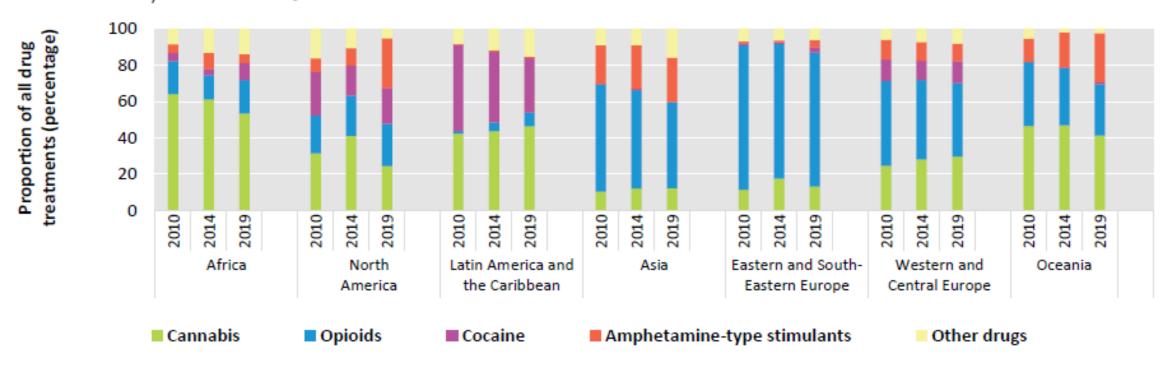
DISTURBO DA USO DI CANNABIS: RICHIESTA DI TRATTAMENTO







FIG. 20 Trends in the primary drug of concern in people in treatment for drug use disorders, by region, 2010, 2014 and 2019











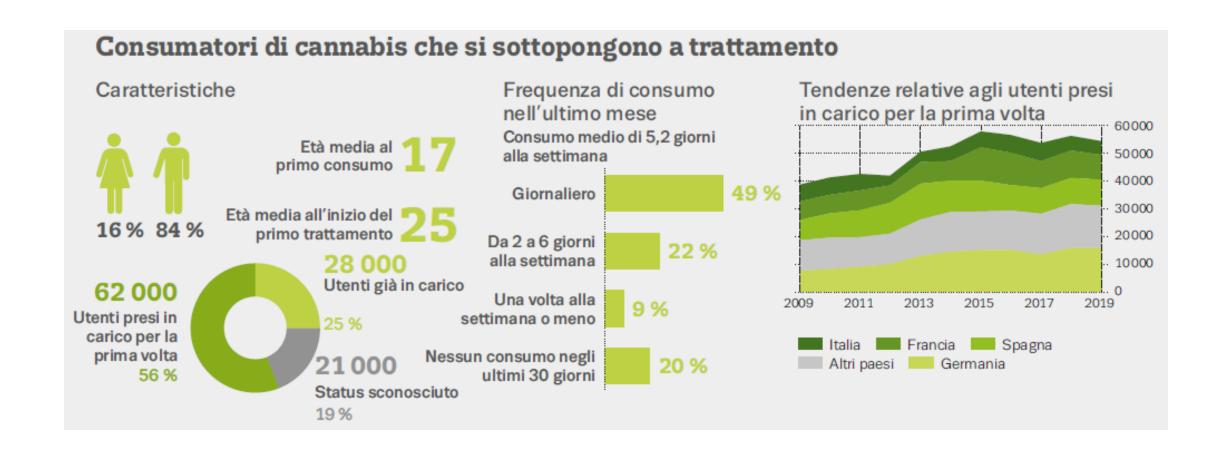


DISTURBO DA USO DI CANNABIS: RICHIESTA DI TRATTAMENTO







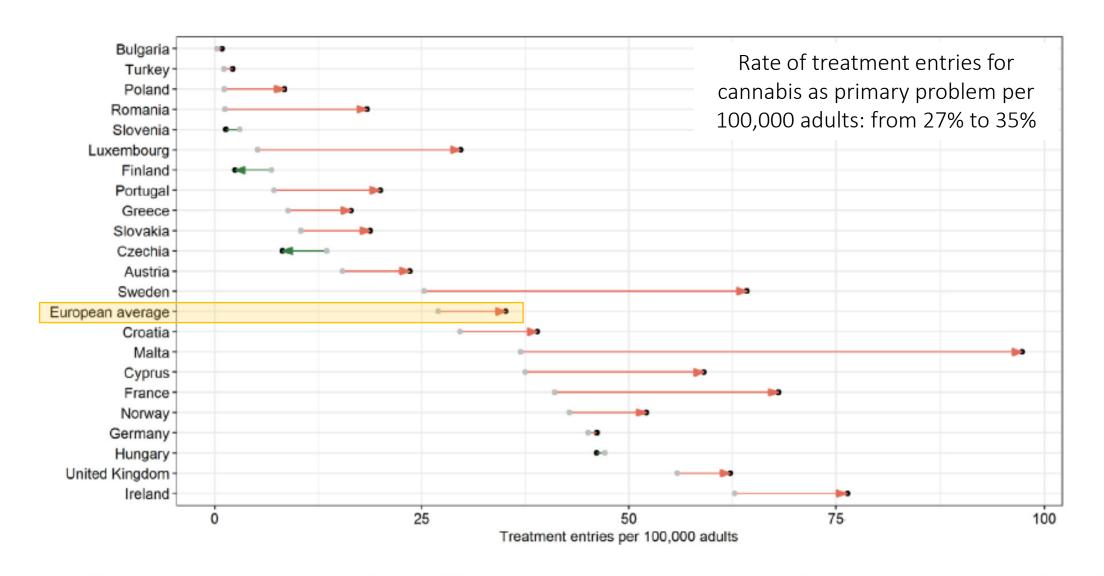


DISTURBO DA USO DI CANNABIS: RICHIESTA DI TRATTAMENTO









DISTURBO DA USO DI CANNABIS: RICHIESTA DI TRATTAMENTO



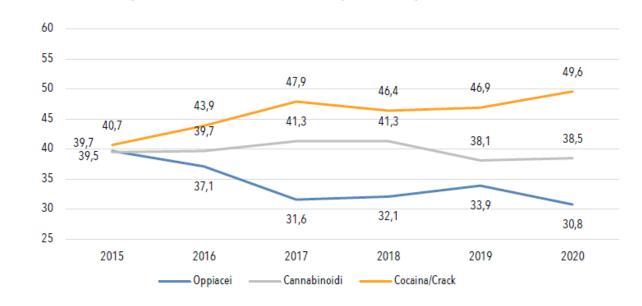
Figura 6.1.5 - Distribuzione degli utenti trattati nei SerD per sostanza primaria 100 86,5 63,8 22,7 20 11,9

Cannabinoidi

2020

2009

Figura 6.1.7 - Andamento temporale dei nuovi utenti trattati nei SerD per sostanza (primaria o secondaria)



Cannabinoidi

1999

Oppiacei

SOSTANZA PRIMARIA 11,9 14.968

Cocaina/Crack

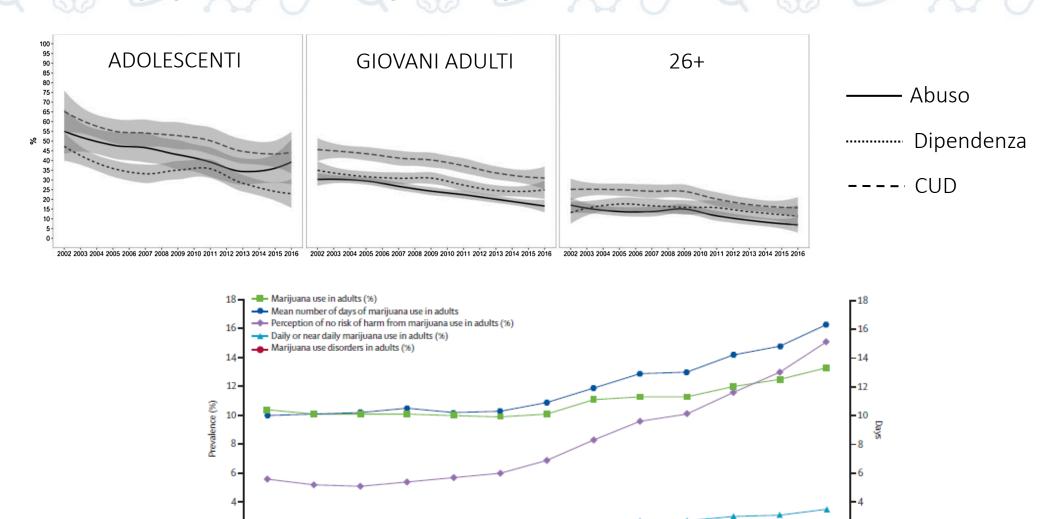
SOSTANZA SECONDARIA 19.565 15,6

TOTALE

34.533

27,5

CUD: DIMENSIONI DEL FENOMENO



Santaella-Tenorio J et al. Cannabis use disorder among people using cannabis daily/almost daily in the United States, 2002-2016. Drug Alcohol Depend. 2019 Dec 1;205:107621; Compton WM et al. Marijuana use and use disorders in adults in the USA, 2002-14: analysis of annual cross-sectional surveys. Lancet Psychiatry. 2016 Oct;3(10):954-964

CUD: DIMENSIONI DEL FENOMENO

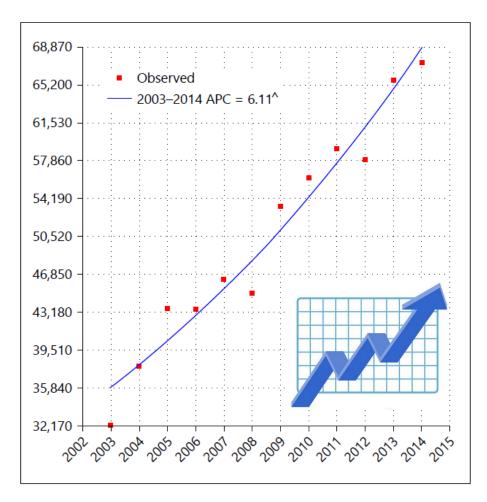
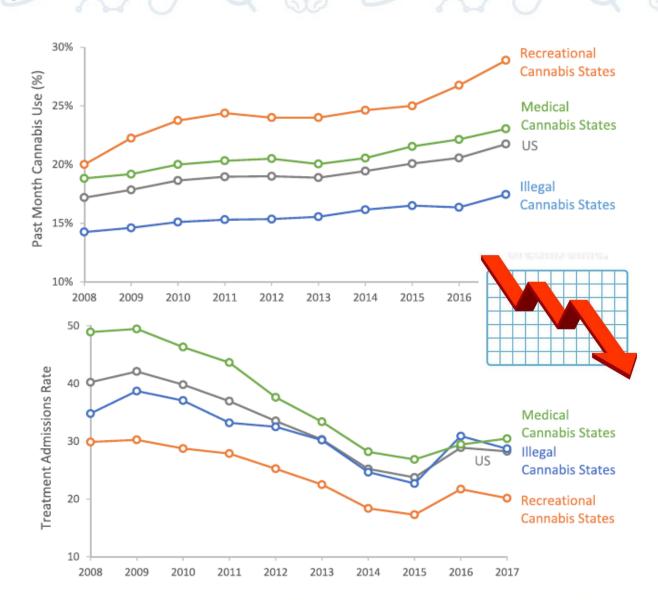


Fig. 1. Trends in the number of first-time cannabis treatment admission in Eu-22 – joinpoint regression outcome, 2003–2014. APC, annual percent change – observed.



Montanari L et al. Cannabis Use among People Entering Drug Treatment in Europe: A Growing Phenomenon? Eur Addict Res. 2017;23(3):113-121

Mennis J et al. Young adult cannabis use disorder treatment admissions declined as past month cannabis use increased in the U.S.: An analysis of states by year, 2008-2017. Addict Behav. 2021 Dec;123

CUD: BARRIERE AL TRATTAMENTO

Factor	Sought Treatment in Lifetime (n = 126) $\%$ (SE)	Never sought Treatment in Lifetime (n = 225) $\%$ (SE)	Total (n = 351) % (SE)
PR EDISP OSING			
Fear of Stigma/Social Consequences			
Too embarrassed to discuss it	32.15 (5.91)	24.32 (2.79 ¹)	27.23 (2.80 ¹)
Afraid of what others would think	27.96 (5.82)	17.63 (3.21)	21.47 (2.99)
Hated answering personal questions	12.91 (3.56)	13.38 (2.62)	13.20 (2.06)
Afraid would lose job	13.15 (3.82)	8.66 (2.30)	10.33 (2.22)
Was afraid children would be taken away	4.36 (1.63)	6.14 (1.34)	5.48 (1.01)
Family member objected	0.95 (0.94)	0.98 (0.53)	0.97 (0.49)
Fear of Treatment			
Afraid would be put into the hospital	17.92 (4.60)	15.14 (3.01)	16.17 (2.51)
Afraid of the treatment	15.62 (4.25)	10.93 (3.28)	12.67 (2.46)
Treatment Pessimism			
Did not think anyone could help	27.85 (5.06)	11.62 (2.41)	17.66 (2.73)
Tried to get help before and it didn't work	17.83 (3.84)	7.48 (2.17)	11.33 (2.08)
ENABLING			
Financial			
Could not afford to pay	19.07 (4.63)	18.19 (2.87)	18.52 (2.76)
Health insurance didn't pay for it	20.01 (4.48)	13.99 (3.47)	16.23 (2.75)
Could not arrange child care	0.66 (0.49)	1.61 (1.12)	1.26 (0.73)
Logistical/Structural			
Did not know any place to go for help	10.89 (3.23)	13.86 (2.70)	12.75 (2.10)
Did not have time	15.50 (4.29)	6.81 (2.06)	10.04 (2.21)
Did not have a way to get there	10.46 (3.77)	4.91 (1.33)	6.97 (1.65)
Had to wait too long	8.10 (3.22)	2.45 (0.93)	4.55 (1.40)
Inconvenient hours	4.70 (2.17)	1.91 (1.17)	2.95 (1.10)
Could not arrange child care	0.66 (0.49)	1.61 (1.12)	1.26 (0.73)
Cannot speak English well	2.51 (2.48)	0.29 (0.29)	1.11 (0.94)
PER CEIVED NEED			
Self-Reliance/Minimizing Problems			
Thought should be strong enough to handle it alone	47.96 (6.27)	44.38 (4.56)	45.71 (3.94)
Thought the problem would get better by itself	41.34 (6.43)	39.36 (3.84)	40.10 (3.41)
Wanted to keep using	37.85 (5.86)	31.14 (3.33)	33.64 (3.05)
Didn't want to go	36.38 (4.93)	31.59 (3.52)	33.38 (3.19)
Thought problem was not serious enough	23.17 (5.11)	27.30 (3.44)	25.77 (2.97)
Stopped using on my own	16.43 (4.24)	26.64 (3.58)	22.85 (2.59)
Did not think it was necessary (despite family requests)	23.78 (4.72)	10.34 (2.45)	15.34 (2.50)

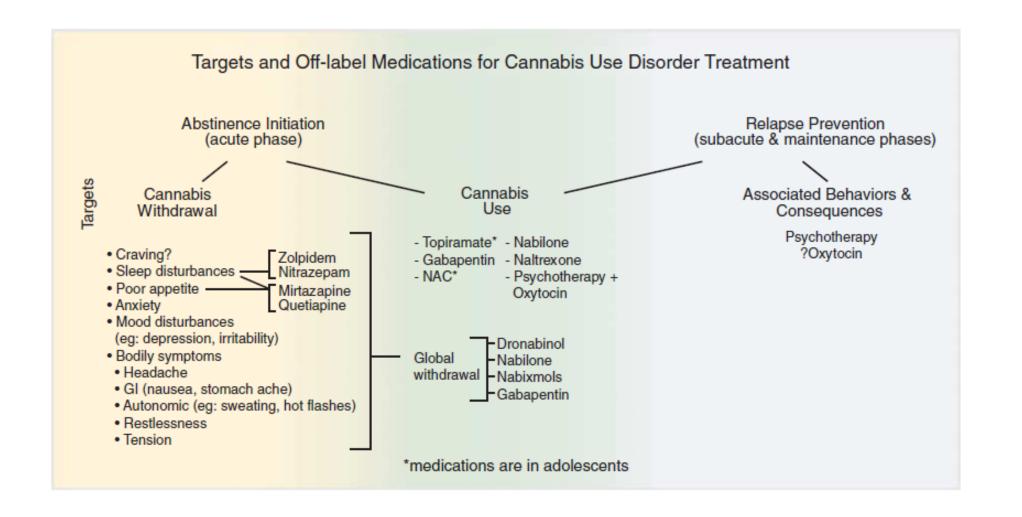


Kerridge BT et al. Predictors of treatment utilization and barriers to treatment utilization among individuals with lifetime cannabis use disorder in the United States. Drug Alcohol Depend. 2017 Dec 1;181:223-228

TABLE 3 Summar	y of Placebo-Controlled	Clinical Trials of	Medications for	CUD Treatment
----------------	-------------------------	--------------------	-----------------	---------------

Medication(s)	Mechanism of action	Reasoning for CUD treatment	Utility in CUD treatment	Publications
Bupropion	NE+DA reuptake inhibition	Withdrawal Cannabis use	Limited, if any	Haney et al, 2001 (L) Carpentar et al, 2009 (T) Penetar et al, 2012 (T)
Nefazodone	NE reuptake inhibition	Withdrawal Cannabis use	Limited, if any; no longer available in the United States	Haney et al, 2003 (L) Carpentar et al, 2009 (T)
Atomoxetine ^a	NE+DA reuptake inhibition	Cognitive symptoms (similarities with ADHD), +ADHD	Limited, if any	McRae-Clark et al (2010) (T
Venlafaxin e ^a	NE+5HT reuptake inhibition	CUD+MDD	Limited, potentially exacerbates cannabis use	Levin et al, 2013 (T)
Mirtazepine	NE+5HT reuptake inhibition	Withdrawal Cannabis use	Specific withdrawal symptoms of insomnia and food intake	Haney et al, 2010 (L)
Buspirone	5HT IA partial agonist	Cannabis use, anxiety	Limited, if any, particularly in women	McRae-Clark et al, 2009 (T) McRae-Clark et al, 2015 (T)
Escitalopram	5HT reuptake inhibition	Cannabis use, withdrawal, anxiety, depression	Limited, if any	Weinstein et al, 2014 (T)
Vilazodone	5HT IA partial agonist+5HT reuptake inhibition	Cannabis use	Limited, if any	McRae-Clark et al, 2016 (T)
Divalproex	Blocks voltage-dependent Na channels, increases GABA	Withdrawal, irritability Cannabis use	Limited, if any	Haney et al, 2004 (L) Levin et al, 2004 (L)
Lithium carbonate	Not fully known, mood stabilizer with impact of depression, stimulates oxytocin release	Withdrawal Treatment completion	Limited, if any	Johnston et al, 2014 (T/I)
Quetiapine	5HT2A, DA2, H1, al, a2 antagonism; 5HT1A partial agonism; NE reuptake inhibition	Withdrawal Cannabis use	Specific withdrawal symptoms, including sleep, food intake, and weight loss; concerns about increases in craving need to be considered	Cooper et al, 2013 (L)
Badofen	GABA-B agonism	Withdrawal Cannabis use	Limited, if any	Haney et al, 2010 (L)
Zolpidem	GABA-A agonism	Withdrawal, insomnia	Withdrawal-related sleep disturbances	Vandrey et al, 2011 (L)

Gabapentin	Blocks a2d subunit on voltage gated Na channels, indirect GABA modulator; restores brain CRF-mediated homeostasis	Withdrawal, cognitive performance, cannabis use, problems secondary to cannabis	Encouraging for use in withdrawal, reductions, craving, cognitive functioning, and improvement in problems	Mason et al, 2012 (T)
Topiramate	Blocks Na and Ca channels, potentiates GABA-A; AMPA/kinate glutamate antagonism	Cannabis use	Encouraging for its reduced use in adolescents, not well tolerated, slower titration may help	Miranda et al, 2016 (T)
Dronabinol	CB-I agonist	Withdrawal Cannabis use	Encouraging for reductions in global withdrawal symptoms	Haney et al, 2004 (L) Budney et al, 2007(L) Levin et al, 2011 (T)
Dronabinol +lofexidine	CB-I agonist+a2 agonist	Withdrawal Cannabis use	Lofexidine adds no benefit and is poorly tolerated to dronabinol monotherapy	Haney et al, 2008 (L) Levin et al, 2016 (T)
Nabilone	CB-I agonist	Withdrawal Cannabis use	Encouraging for its reductions in withdrawal and cannabis use	Haney, 2013b (L)
Nabilone +zolpidem	CB-I agonist+GABA-A activity	Withdrawal, withdrawal-related sleep disturbance, cannabis use	Encouraging for reductions in withdrawal and cannabis use	Herrmann et al, 2016 (L)
Nabixmols	CB-I agonist+multi functions (through CBD)	Withdrawal Cannabis use	Encouraging for use in withdrawal	Allsop et al, 2014 (T/I)
Cannabidiol	Multifunctional	Cannabis use	Limited, if any	Haney et al, 2016 (L)
Rimonabant	CB-I antagonism	Cannabis use	Limited if any due to discontinued use/ safety risk	Huestis et al, 2001 (L) Huestis et al, 2007 (L)
Naltrexone	Mu opioid antagonism	Cannabis use	Encouraging for reductions in use when chronically dosed	Cooper and Haney, 2010 (L) Haney et al, 2015 (L)
N-acetylcysteine	Cysteine prodrug (cysteine-glutamate exchanger)	Cannabis use	Encouraging for reductions in use in adolescents; limited use in adults	Gray et <i>al</i> , 2012 (T) Gray, 2017 (T)
Oxytocin	Neural roles in prosocial behavior	Cannabis use (indirectly by enhancing psychosocial treatment)	Encouraging for enhancement of psychosocial treatment	Sherman et al, 2017 (T)





Cochrane Database of Systematic Reviews

Pharmacotherapies for cannabis dependence (Review)

Nielsen S, Gowing L, Sabioni P, Le Foll B

AUTHORS' CONCLUSIONS

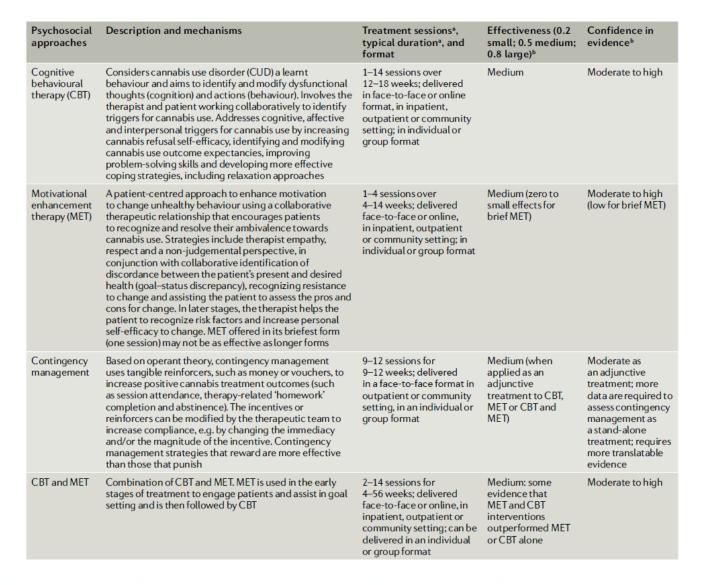
Implications for practice

Studies undertaken to date on pharmacotherapies for cannabis dependence are insufficient to guide clinical practice. There is incomplete evidence for all of the pharmacotherapies investigated in this review.

At this point in time, psychological approaches such as MET and cognitive-behavioural therapy remain the mainstay of treatment for cannabis use disorders.



CUD: TRATTAMENTO APPROCCIO PSICOSOCIALE



- Social support Counselling
- Drug education Counselling
- Relapse prevention
- Mindfulness Meditation
- Mutual help programmes
- Individual, group- based and family-based

Number of studies examining each type of psychotherapy in the two reviews combined.

Type of psychotherapy examined	Number of studies focusing on this intervention	Reference of studies
SS	2	[20,23]
MM	1	[29]
DC	2	[27,31]
DC + CBT	1	[30]
DC+CM	1	[27]
CM alone	3	[26,28,35]
MET	7	[17,19,24,25,36,39,40]
MET + CM	1	[22]
CBT	5	[18-20,23,28]
CBT + CM	2	[26,28]
MET + CBT	9	[17,21,27,32-35,37,38]
MET + CBT + CM	5	[17,21,27,35,37]

CM: Contingency management; CBT: Cognitive-behavioral therapy; DC: Drug counseling; MET: Motivational enhancement therapy; MM: Mindfulness-based meditation; SS: Social support.

THC: UNITA' STANDARD



















First author, year	Reference	Standard cannabis unit	Description	Strengths: accounts for	Limitations: does not account for
Wetherill, 2016	[61]	Gram years	Number of daily grams consumed, multiplied by years of cannabis use	Some different methods of administration	Variation in quantities of THC per gram of cannabis; other methods of administration (e.g. vaporizer, vape pen, edible, liquid)
Casajuana- Kögel, 2017	[62]	Standard joint unit	1 unit = 1 joint, or 0.25 g cannabis, or 7 mg THC, or 1 Euro	The most common method of administration in Europe	Variation in quantities of THC per joint; other methods of administration (e.g. pipe, blunt, bong, dabbing, vaporizer, vape pen, edible, liquid)
Ziesser, 2012	[63]	Standard joint	1 standard joint = 0.5 g cannabis, 10 puffs, or 5 bong hits, or 5 pipe hits	Some different administration methods and/or number of puffs	Variation in quantities of THC; other methods of administration (e.g. blunt, dabbing, vaporizer, vape pen, edible, liquid)
Norberg, 2012	[64]	Cannabis unit	1 unit = 0.25 grams cannabis, or 1 paper joint or 1 blunt, or 2 skinny paper joints/blunt, or 3 cones/water pipes/bongs/ bucket bongs	Some different sizes of joint and methods of administration	Variation in quantities of THC; other methods of administration (e.g. pipe, dabbing, vaporizer, vape pen, edible, liquid)
Hindocha, 2017	[65]	THC/CBD ratios	High THC & low CBD (e.g. 1 unit = 0.25 g) Equal THC & CBD (e.g. 1 unit = 0.50 g) High CBD & low THC (e.g. 1 unit = 0.75 g)	Variation in THC/CBD ratios	Complete variation in quantities of THC per gram of cannabis; other methods of administration (e.g. vaporizer, vape pen, edible, liquid)



Panel: Summary of guidance on the standard THC unit

The standard THC unit:

- is defined as any formulation containing 5 mg of THC;
- applies to both natural and synthetic THC formulations;
- is specific to THC and does not apply to other cannabis constituents for which standard units have not been established;
- should be used to report the dose of THC administered to or consumed by research participants; pharmacokinetic parameters should be reported using conventional units (eg, ng/mL or ng·h/mL).

Additionally:

- Examples of reporting are as follows: 10 mg THC = 2 standard THC units, 16 mg THC = 3.2 standard THC units; a suggested abbreviation is STU.
- Research studies can administer more (or less) than 5 mg THC as necessary, but results should be expressed in multiples (or fractions) of the standard THC unit.
- Investigators are expected to report the standard THC unit in funding applications, progress reports, and dissemination of results through publications, posters, and other materials. This does not preclude additional reporting of results using other measures (eq, mg of THC).
- There is no requirement to report any other new information other than the standard THC unit.
- Other cannabis constituents can be reported as they typically would be, if relevant.
- If it is not possible to report the standard THC unit, researchers should provide justification for this (eg, a specific dataset or survey instrument might not provide the necessary information needed to convert the amount of THC into standard THC units).

Freeman TP, Lorenzetti V. 'Standard THC units': a proposal to standardize dose across all cannabis products and methods of administration.

Addiction. 2020 Jul;115(7):1207-1216; Freeman TP, Lorenzetti V. A standard THC unit for reporting of health research on cannabis and cannabinoids. Lancet Psychiatry. 2021 Sep 7:S2215-0366(21)00355-2

RIDUZIONE DEL DANNO: Lower-Risk Cannabis Use Guidelines



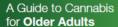




General Precaution A: People who use cannabis (PWUC) need to know that there is no universally safe level of cannabis use; thus, the only reliable way to avoid any risk for harm from using cannabis is to abstain from its use	General Precaution B: Frequent cannabis use, and especially intensive use over longer periods, can lead to a 'cannabis use disorder' (CUD) or cannabis dependence, that may require treatment	General Precaution C: PWUC should exercise social consideration and responsibility in avoiding cannabis use that may result in harm-to-other
Recommendation #1: The initiation of cannabis use should be delayed until after late adolescence, or the completion of puberty, to reduce development-related vulnerabilities for harm	Recommendation #2: PWUC should use 'low-potency' cannabis products, i.e., cannabis products with ideally lower total THC content, or a high CBD/THC content ratio	Recommendation #3: All main available modes-of-use options come with some risk for harm; PWUC should refrain from cannabis 'smoking' and employ alternative routes-of-use for pulmonary health protection
Recommendation #4 : If use occurs by inhalation, PWUC should avoid "deep inhalation", prolonged breathholding, or similar inhalation practices	Recommendation #5: PWUC should refrain from frequent (e.g., daily or near-daily) or intensive (e.g., binging) cannabis use, and instead limit themselves to less frequent or occasional use	Recommendation #6: Where circumstances allow, PWUC should use legal and quality-controlled cannabis products and use devices
Recommendation #7: PWUC who experience impaired cognitive performance should consider temporarily suspending or substantially reducing the intensity (e.g., frequency/potency) of their cannabis use	Recommendation #8: PWUC should avoid driving a motorvehicle or operating machinery while under the influence of cannabis because of acute impairment and elevated risk of crash involvement, including injury or death; however, the severity and duration of impairment vary depending on multiple factor	Recommendation #9: It is prudent for people who intend to procreate and for women who are pregnant or breastfeeding to abstain from cannabis use towards reducing possible risks for reproduction and of health harm to offspring, respectively
Recommendation #10: PWUC should exercise general caution in combining other psychoactive substances with cannabis use	Recommendation #11: Some specific groups of people are at elevated risk for cannabis use-related health problems because of biological pre-dispositions or comorbidities. They should accordingly (and possibly on medical advice as required) avoid or adjust their cannabis use	Recommendation #12: The combination of risk-factors for adverse health outcomes from cannabis use further amplifies the likelihood of experiencing severe harms and should be avoided

Fischer B et al. Lower-Risk Cannabis Use Guidelines (LRCUG) for reducing health harms from non-medical cannabis use: A comprehensive evidence and recommendations update. Int J Drug Policy. 2021 Aug 28:103381

PREVENZIONE E RIDUZIONE DEL DANNO



What Is Cannabis?

Cannabis, commonly called marijuana, pot or weed, is a product made from the cannabis plant. It is often used for non-medical purposes because of its intoxicating effects or the "high" it produces. Cannabis can also be used for medical purposes. However, for most health conditions, more research is still needed to determine whether it is effective. Cannabis contains over 100 chemical compounds called cannabinoids. The two best-known cannabinoids are:

- · Tetrahydrocannabinol (THC) is the primary "psychoactive" component of cannabis. It is the ingredient in cannabis that produces the high. Research suggests that THC might have some medical benefits for certain health conditions.
- · Cannabidiol (CBD) does not produce a high, but can cause drowsiness. Scientists are also studying CBD for its medical applications.

Cannabis comes in many forms, including dried cannabis, cannabis extracts and cannabis topicals. Edible cannabis products, commonly known as edibles, are cannabis-infused food (e.g., chocolate) and drinks (e.g., tea).

Did You Know?

Concentrations of THC in dried cannabis have increased over the past few decades. In 1995, dried cannabis contained about 4% THC. Nowadays, it averages about 15-20% THC.

Some cannabis extracts, called concentrates. can contain more than 80% THC.

There are also cannabis products that are made up mostly of CBD and contain only small

Always read the label to check the amount and concentrations of THC and CBD.







What Can I Expect When Using Cannabis? Cannabis affects everyone differently.

Smoking, vaping or eating cannabis products can lead to feelings of euphoria (happiness) and relaxation. It can also cause impairment, which can include:

- Concentration and memory
- Impaired motor skills and slower reaction time that can increase the risk of falls and injury

Cannabis can impair your ability to drive safely. Cannabis-impaired driving is illegal.

Consuming too much cannabis or cannabis with higher levels of THC can lead to over-intoxication. Symptoms can include

- · Anxiety, panic and elevated heart rate
- Nausea and vomiting
- Paranoia and psychosis (i.e., hallucinations and delusions)

If you or someone you know has consumed too much cannabis and is not feeling well, seek medical attention.



COVID-19 and CANNABIS How to reduce your risk





If you plan to use or are using cannabis, there are things you can do to help reduce the spread of COVID-19 and the risks to your health.



Don't share joints, vaping devices, pipes or bongs.

 COVID-19 spreads through respiratory droplets released when people speak, cough or sneeze. You can pick it up if you put items contaminated by droplets — which can include joints, vaportzers. pipes and bongs - in your mouth or touch them and then touch your eyes, nose or mouth.



Wash your hands before putting any cannabis product to your mouth.

. Wash your hands often with soap and water for at least 20 seconds. If soap and water are not avallable, use alcohol-based hand sanitizer



If you plan to smoke or vape cannabis, know that it can negatively affect

- . Try to limit cannable smoking or vaping to once a week, if possible.
- · Avoid smoking or vaping cannabls if you feel sick. · Avoid taking deep inhalations and try not to hold your breath.
- . Use products that contain no more than 100 mg/g (10%) THC content.
- . Follow Canada's Lower-Risk Cannabis Use Guidelines to reduce your risk of experiencing harms when using cannabls.



Purchase your cannabis from licensed and regulated retailers.

. Purchasing cannabls from Illegal sources has negative health and safety risks, as the quality of the product cannot be guaranteed and it can be contaminated with other substances (metals, pesticides, fundi, other drugs, etc.).



Cannabis can interact with your medication and negatively

- . If you take medications, whether prescribed or over the counter, talk to your healthcare provider before using cannabls.
- . If you are using cannabls for medical reasons, talk to your healthcare provider about your use during the COVID-19 pandemic



For more information, visit our COVID-19 resources page at www.ccsa.ca

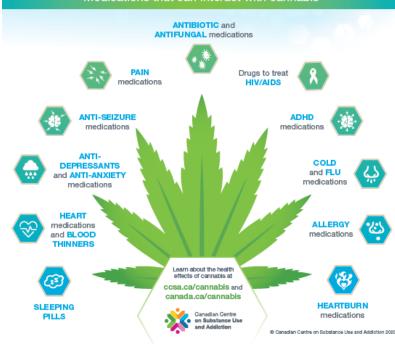
© Canadian Centre on Substance Use and Addiction 2020

Cannabis and **Your Medication**

Cannabis can interact with your medications and affect your health.

For more information, talk to your healthcare provider.

Medications that can interact with cannabis





PREVENZIONE E RIDUZIONE DEL DANNO















How To Safely Store Your Cannabis



Labels on all cannabis products

- · All legal cannabis products will have a label
- · Make sure the label is always clearly visible
- · If you make your own edible cannabis products, be sure to properly label them

Label examples







Store in child-safe containers

- · All legal cannabis products must be in child-resistant packages
- · Make sure they are appropriately re-sealed after use
- · Consider investing in a lockbox



Keep out of reach and sight of children and pets

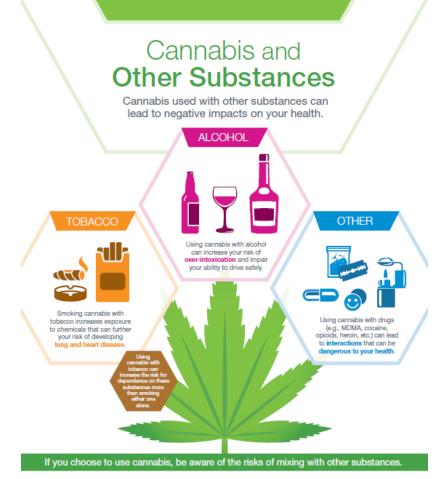
 Make sure you properly store and dispose of cannabis products in a place that is not easily visible and accessible by children or pets



Stay informed

ccsa.ca/cannabis canada.ca/cannabis







Learn about the health effects of cannable at ccsa.ca/cannabis and canada.ca/cannabis

Canadian Centre on Substance Use and Addiction 2020

Know the Health Risks of Cannabis



Mental Health

Daily or near-daily use of cannabis can contribute to dependence and mental health problems over time.



Driving

Cannabis can impair your motor coordination, judgment and other skills required for safe driving.



Respiratory Effects

Toxic and carcinogenic chemicals found in tobacco smoke are also found in cannabis smoke, and can affect the lungs and airways.



Pregnancy

Substances in cannabis are transferred from mother to child and can affect your baby. Not using cannabis if pregnant or breastfeeding is the safest option.



Edible Cannabis

Consuming too much THC can lead to over-intoxication, which includes intense anxiety, vomiting and symptoms of psychosis (paranoia).



Cannabis Extracts

Cannabis extracts with high THC content increase the risk of overintoxication and addiction.



Stav Informed ccsa.ca/cannabis canada.ca/cannabis































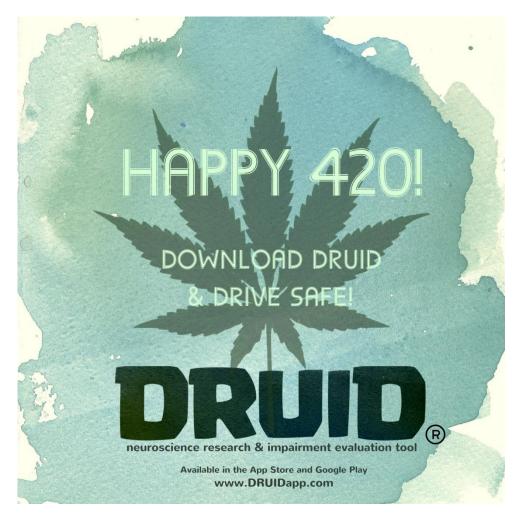








PREVENZIONE E RIDUZIONE DEL DANNO







Trends in attitudes toward cannabis legalisation and perceptions about cannabis	
Support for cannabis legalisation	^
Perceived harmfulness of cannabis use	Ψ
Trends in cannabis prices and potencies	
Prices of legal cannabis	Ψ
Potencies of cannabis products	^
The impact of cannabis legalisation on prevalence of cannabis use and CUD	
Adolescent cannabis use	MIXED
Adolescent cannabis use disorder	Ψ
Adult cannabis use	^
Adult cannabis use disorder	MIXED
Adult subgroups (gender, race, socioeconomic status, marital status) and Specific populations (pregnant, older, disabled)	^
The impact of cannabis legalisation on prevalence of related adverse health effects	
Cannabis-related motor vehicle accidents	MIXED
Cannabis-related health service presentations	^

CONCLUSIONI



Fenomeno in aumento e in mutamento Nuove modalità di consumo

Nuovi approcci di ricerca

Individuare le traiettorie psico-patologiche Individuare il target del trattamento Nuovi strumenti di trattamento

Nuove modalità di presa in carico trattamento

Attività di riduzione del danno



































