

Increased use of illicit drugs in a Dutch cluster headache population

Cephalalgia 2019, Vol. 39(5) 626–634 © International Headache Society 2018 © 0 Article reuse guidelines: sagepub.com/journals-permissions DOI: 10.1177/033102418804160 journals.sagepub.com/home/cep



Ilse F de Coo^{1,2,*}, Willemijn C Naber^{1,*}, Leopoldine A Wilbrink¹, Joost Haan^{1,3}, Michel D Ferrari¹ and Rolf Fronczek^{1,4}

Abstract

Introduction: Many patients with cluster headache report use of illicit drugs. We systematically assessed the use of illicit drugs and their effects in a well-defined Dutch cluster headache population.

Methods: In this cross-sectional explorative study, 756 people with cluster headache received a questionnaire on lifetime use and perceived effects of illicit drugs. Results were compared with age and sex-matched official data from the Dutch general population.

Results: Compared to the data from the general population, there were more illicit drug users in the cluster headache group (31.7% vs. 23.8%; p < 0.01). Reduction in attack frequency was reported by 56% (n = 22) of psilocybin mushroom, 60% (n = 3) of lysergic acid diethylamide and 50% (n = 2) of heroin users, and a decreased attack duration was reported by 46% (n = 18) of PSI, 50% (n = 2) of heroin and 36% (n = 8) of amphetamine users.

Conclusion: In the Netherlands, people with cluster headache use illicit drugs more often than the general population. The question remains whether this is due to an actual alleviatory effect, placebo response, conviction, or common pathophysiological background between cluster headache and addictive behaviours such as drug use.

Keywords

Cluster headache, heroin, GHB, cannabis, psilocybin mushroom

Date received: 15 April 2018; revised: 23 August 2018; accepted: 7 September 2018

Introduction

Cluster headache is a rare primary headache disorder characterized by severe unilateral headache attacks lasting 15 to 180 minutes accompanied by cranial ipsilateral autonomic symptoms (1). Most people suffer from the episodic form, in which attacks occur in clusters of weeks to months alternating with remissions of several months to years (1). About 14% of people with cluster headache suffer from the chronic form and do not have attack-free episodes longer than a month (1,2). Although most people with cluster headache can be successfully treated with regular medication, in 10–20% this is not effective (3). These people are more likely to experiment with alternative treatments (4), including illicit drugs.

Few epidemiologic studies have investigated the use of illicit drugs in cluster headache (4–7). In two French studies, 26-32% of cluster headache patients used cannabis regularly, which is higher than in the general French population (7%) (6,7). The prevalence of cannabis use was considerably lower (10%) in an American cluster headache cohort (7,8). Illicit drug use in cluster headache has been associated with younger age, male gender, and smoking. Use of alternative treatments in cluster headache has also been associated with lower income and higher annual attack frequency (5,9).

Some patients report an acute effect of cannabis, cocaine, psilocybin mushrooms (PSI) and heroin (4,10,11). In a small open study, 22 of 26 patients reported an acute effect on their cluster headache attacks after psilocybin mushrooms use (11). In Italy,

⁴Slaap-Waakcentrum SEIN, Heemstede, the Netherlands

*Ilse F de Coo and Willemijn C Naber contributed equally to this work.

Corresponding author:

Ilse F de Coo, Department of Neurology, K5-Q-104, Albinusdreef 2, 2333 ZA, Leiden University Medical Center, Leiden, the Netherlands. Email: i.f.de_coo@lumc.nl

¹Department of Neurology, Leiden University Medical Center, Leiden, the Netherlands

²Sophia Rehabilitation Center, the Hague, the Netherlands

³Department of Neurology, Alrijne Hospital, Leiderdorp, the Netherlands

7-19% of people with cluster headache reported having tried cannabis as an acute treatment, but self-reported effects were inconclusive (4,7). In 25%, it even seemed to trigger attacks (7).

Illicit drugs are not only described as acute treatment, but in several small studies also as alternative prophylactic treatment. Remission-extending qualities have been attributed to PSI, gamma-hydroxybutyrate (GHB) and lysergic acid diethylamide (LSD) (4,11–13). GHB's sodium salt form (sodium oxybate) reduced attack frequency in five patients with cluster headache (12,13). In a small study, patients reported a perceived prophylactic effect of both LSD and PSI (11).

While these possible alleviatory effects of illicit drugs in cluster headache are intriguing, the studies that have been conducted were all small and uncontrolled. Although restricted use of cannabis is legal in the Netherlands, the use of drugs in cluster headache has not been investigated. Here we systematically assessed (i) the use of illicit drugs in general and (ii) the use of several illicit drugs in a representative Dutch cluster headache population and compared this to the Dutch general population. Additionally, we determined whether illicit drugs influenced cluster headache attack duration and attack frequency in the Dutch cluster headache population.

Material and methods

Study design

This explorative cross-sectional study was conducted as part of the ongoing nationwide Leiden University Cluster Headache neuro-analysis programme (LUCA) and compared to official data of the Dutch general population.

Cluster headache population. The LUCA program is heavily promoted throughout the Netherlands to attract as many potential cluster headache participants of 18 years and older as possible. In addition, participants attending the Leiden University Medical Center and other headache outpatient departments were invited to participate in the LUCA program. All possible participants were invited to fill out a validated, web-based screening questionnaire about cluster headache based on the ICHD-II criteria for cluster headache (14). The screener has been validated and has a diagnostic specificity of 0.89 for cluster headache (15). All people who fulfilled the ICHD-II criteria also fulfilled the latest ICHD-III beta version for cluster headache (16). All people who screened positive received a second, more extensive web-based questionnaire.

For the present study, people who had been screened as positive for cluster headache received an email asking them to fill out a questionnaire concerning illicit drug use. This questionnaire was designed by the authors (IFC and LAW) and included questions about lifetime use of illicit drugs and more specifically the use of cannabis, cocaine, heroin, PSI, 3,4-methylenedioxymethamphetamine (MDMA), LSD, amphetamine and GHB. Also, the questionnaire included two questions about the effect on attack duration and attack frequency of these illicit drugs (there were three answer options: Increased, no effect, decreased). Web-based questionnaires were used, except for people not capable of using the internet, who were allowed to fill out the questionnaires on paper. Those who did not respond to the initial email were reminded twice per email and, when still not responding, they were contacted two more times (once by phone and once by email again). Only people who filled out all items regarding drug use were included.

All LUCA data was securely stored in a web-based database management system of ProMISe (Project Manager Internet Server), which meets ISO 27001 and the requirements for data safety and privacy set by international law.

Dutch general population. Statistics Netherlands (Centraal Bureau voor Statistiek) provided data about the Dutch general population from their annual health survey, which monitors various health and lifestyle aspects including illicit drug use in the Dutch population. Each year, 15,000 randomly selected Dutch citizens of all ages are approached by mail to fill out their online survey. For our study, we only included data from people 18 years old and older. Statistics Netherlands supplies this data as anonymous categorical variables. The information is therefore not traceable to an individual.

For our study, we divided the data into three categories: All, people classified as having headache (migraine or regular severe headache in the previous 12 months), and people classified as having chronic pain (in the previous 12 months).

Ethical approval. The LUCA study was approved by the local medical ethics committee of the Leiden University Medical Center. All participants in the LUCA study provided written informed consent. The survey in the Dutch general population by Statistics Netherlands was exempt from ethical approval according to Dutch laws, therefore no informed consent was needed.

Statistics. We have performed a Chi-square test for categorical variables and independent t-test for continuous variables to describe the population characteristics as shown in Table 1. The Chi-square test with Yates' correction and Fisher's Exact test (n < 5 per group) were used to determine categorical differences in drug use between the cluster headache population and the general population, the headache subgroup and the chronic subgroup of the general population. All data analyses were performed using SPSS 23.0 (SPSS Incorporated, IBM, USA) with statistical significance set at p < 0.05. No power analysis has been performed.

Results

Study population

By August 2014, 756 people with self-reported cluster headache had been invited to this study, of whom 85.1% (643/756) filled out all necessary questionnaires. Of these, 613/643 (95.3%) had received a diagnosis of cluster headache by a physician. A total of 14.542/ 24.396 (59.6%) of the controls from the general population from the 2014 and 2015 cohorts of the Statistics Netherlands responded to the questionnaire and were included in this study. In the Dutch general population cohort, 3457 (23.8%) people reported having "chronic pain" and 2269 (15.6%) people reported having "headaches".

People with cluster headache were more often male, higher educated, smokers, and drank alcohol less often compared to the Dutch general population (Table 1).

Cluster headache compared to the general population

Participants with cluster headache more often used illicit drugs (31.7% vs. 23.8%; p < 0.001; Figure 1), cannabis (29.5% vs. 22.7%; p < 0.001), cocaine (8.9% vs. 4.8%; p < 0.001), amphetamine (6.4% vs. 4.2%; p = 0.011), PSI (9.3% vs. 3.9%; p = 0.00) and heroin (1.1% vs. 0.5%; p = 0.037).

Cluster headache compared to people with headache and the chronic pain in the general population

People with cluster headache more often used illicit drugs than people from the general population classified as having headache (31.7% vs. 25.2; p = 0.001) or chronic pain in the general population (31.7% vs. 20.3%; p < 0.001). Compared to the headache subgroup of the general population, all drugs except GHB were used more frequently by cluster headache patients (Table 2) and compared to the chronic pain subgroup cannabis, cocaine, MDMA, amphetamine and PSI were used more often.

Gender differences

Males more often used illicit drugs compared to females, as shown in Table 3 (cluster headache: 34.7% vs. 23.7%; p=0.008; general population: 29.2% vs. 18.5; p < 0.001). When specified for drug kind, only cannabis (33.0 vs. 20.2%; p=0.028) and LSD (3.25 vs. 0.0%; p=0.044) were more often used in males vs. females with cluster headache.

Males with cluster headache more often used illicit drugs than males from the general population (34.7% vs. 29.2%; p = 0.013), mainly cannabis (33.0% vs. 27.9%; p = 0.019), cocaine (10.4% vs. 6.9%; p = 0.006) and PSI (10.6% vs. 5.5%; p < 0.001; Table 3). There was no difference in illicit drug use between females with and

 Table 1. Demographics of cluster headache population and the general population.

	Cluster headache (n = 643)	General population (n = 14542)	p-value
Demographic characteristics			
Male, n (%)	470 (73.1)	7072 (48.6)	<0.001*
Age (years), median \pm SD	$\textbf{49.9} \pm \textbf{12.1}$	$\textbf{48.5} \pm \textbf{18.0}$	0.0048*
Education, n (%):			<0.001*
 Primary education 	10 (1.4)	1299 (8.9)	
 Pre-vocational secondary education 	151 (23.5)	3001 (20.6)	
 Secondary vocational education 	236 (36.7)	6151 (44.2)	
 Higher professional education 	157 (24.4)	2682 (18.4)	
 University education 	80 (24.4)	4 (9.7)	
Smoking, n (%)	342 (53.8)	3778 (26.0)	<0.001*
Alcohol use, n (%)	481 (75.6)	11792 (81.1)	<0.001*
Body mass index, median \pm SD	25.4 ± 3.8	25.5 ± 4.3	0.655
Episodic cluster headache, n (%)	492 (76.50)	N/A	

*p < 0.05 (χ^2 test with Yates' correction for categorical data, which were male, education, smoking and alcohol use. Single sample test for interval data, which were body mass index and age).

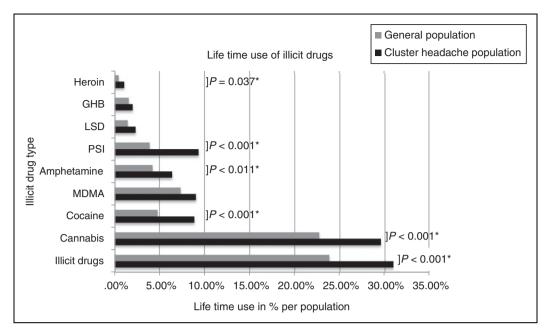


Figure 1. Lifetime use of different types of illicit drugs in cluster headache and general population.
Illicit drugs: All drugs; MDMA: 3,4-methylenedioxymethamphetamine; PSI: psilocybin mushroom; LSD: lysergic acid diethylamide; GHB:
gamma-hydroxybutyrate.

*p < 0.05 (χ^2 test with Yates' correction for categorical data).

	Cluster headache (n = 643)	General population: Headache subgroup (n = 2269)	p-value ^a	General population: chronic pain subgroup (n = 3457)	p-value ^b
Illicit drugs n (%)	204 (31.7)	571 (25.2)	0.001*	701 (20.3)	<0.001*
Cannabis n (%)	190 (29.5)	549 (24.2)	0.007*	670 (19.4)	<0.001*
Cocaine n (%)	57 (8.9)	101 (4.5)	<0.001*	159 (4.6)	<0.001*
MDMA n (%)	58 (9.0)	137 (6.0)	0.010*	191 (5.5)	0.001*
Amphetamine n (%)	41 (6.4)	76 (3.3)	0.001*	138 (4.0)	0.009*
PSI n (%)	60 (9.3)	75 (3.3)	<0.001*	121 (3.5)	<0.001*
LSD n (%)	15 (2.3)	25 (1.1)	0.030*	54 (1.6)	0.219
GHB n (%)	13 (2.0)	27 (1.2)	0.159	54 (1.6)	0.500
Heroin n (%)	7 (1.1)	9 (0.4)	0.062	22 (0.6)	0.202

Table 2. Lifetime use of different types of illicit drugs in the cluster headache, headache and chronic pain subgroup.

Illicit drugs: all drugs; MDMA: 3,4-methylenedioxymethamphetamine; PSI: psilocybin mushroom; LSD: Lysergic acid diethylamide; GHB: gammahydroxybutyrate; *p*-value^a: *p*-value of comparison cluster headache population to the headache subgroup; *p*-value^b: *p*-value of comparison cluster headache population to the chronic pain subgroup.

*p < 0.05 (χ^2 test with Yates' correction for categorical data).

without cluster headache, except for PSI (5.8% vs. 2.3%; p = 0.008).

Age differences

Age distribution of lifetime prevalence of illicit drug use followed the same pattern in people with cluster headache as in the general population and its subgroups (Figure 2). People with cluster headache of all age cohorts, except between 18-24 (55.6% vs. 41.4%; p=0.502) and 25-30 (58.8 vs. 44.1%; p=0.127), used more illicit drugs than the general population.

Episodic versus chronic cluster headache

There were no differences between episodic and chronic cluster headache general illicit drug use (34.4% vs. 30.9%; p=0.413) or for specific illicit drug use,

	Females			Males			Cluster headache		
	Cluster headache (n = 173)	General population (n = 7470)	p-value	Cluster headache (n = 470)	General population (n = 7072)	p-value	Female (n = 173)	Male (n = 470)	p-value
Illicit drugs n (%)	41 (23.7)	1385 (18.5)	0.105	163 (34.7)	2064 (29.2)	0.013*	41 (23.7)	163 (34.7)	0.008*
Cannabis n (%)	35 (20.2)	1319 (17.7)	0.438	155 (33.0)	1970 (27.9)	0.019*	35 (20.2)	155 (33.0)	0.028*
Cocaine n (%)	8 (4.6)	201 (2.7)	0.192	49 (10.4)	489 (6.9)	0.006*	8 (4.6)	49 (10.4)	0.178
MDMA n (%)	(6.4)	380 (5.1)	0.565	47 (10.0)	681 (9.6)	0.855	11 (6.4)	47 (10.0)	0.799
Amphetamine n (%)	6 (3.5)	187 (2.5)	0.588	35 (7.4)	421 (6.0)	0.224	6 (3.5)	35 (7.4)	0.329
PSI n (%)	10 (5.8)	171 (2.3)	0.008*	50 (10.6)	391 (5.5)	<0.001*	10 (5.8)	50 (10.6)	0.43
LSD n (%)	-	56 (0.7)	0.639	15 (3.2)	154 (2.2)	0.202	-	15 (3.2)	0.044*
GHB n (%)	2 (1.2)	64 (0.9)	0.662	11 (2.3)	165 (2.3)	1.000	2 (1.2)	11 (2.3)	0.661
Heroin n (%)	I (0.6)	19 (0.3)	0.368	6 (1.3)	47 (0.7)	0.143	I (0.6)	6 (1.3)	0.696

Table 3. Lifetime use of different types of illicit drugs in the cluster headache and general population stratified to gender.

Illicit drugs: all drugs; MDMA: 3,4-methylenedioxymethamphetamine; PSI: psilocybin mushroom; LSD: lysergic acid diethylamide; GHB: gammahydroxybutyrate.

*p < 0.05 (χ^2 test with Yates' correction for categorical data and Fisher's Exact test for n < 5).

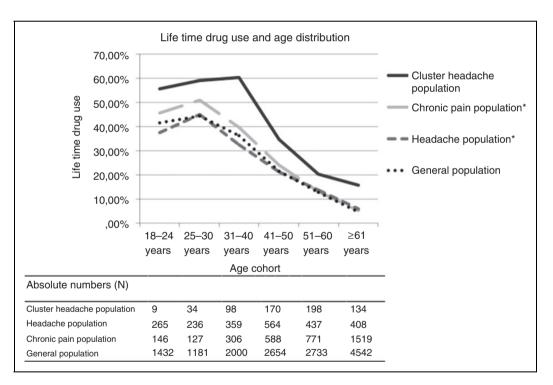


Figure 2. Prevalence of lifetime use of illicit drug stratified to age cohorts. *The headache and chronic pain population are subgroups of the general population.

except for a higher use of MDMA (13.9% vs. 7.5%; p = 0.027) and GHB (4.6% vs. 1.2%; p = 0.015) in chronic cluster headache.

Perceived effects of illicit drug use

Those who used illicit drugs during a cluster headache episode were asked about perceived effect on attack frequency and duration of an individual cluster headache attack. A decrease in attack frequency was reported for LSD (60.0%; n=3), followed by PSI (56.4%; n=22; Table 4). An increased attack frequency was reported for GHB (18.2%; n=2). PSI (46.2%; n=18) and heroin (50.0%; n=2) were most often reported to decrease the duration of individual cluster headache attacks. In contrast, cocaine (10.3%, n=3),

	Cannabis n = 86 (%)	Cocaine n = 29 (%)	MDMA n = 30 (%)	Amphetamine n = 22 (%)	PSI n = 39 (%)	LSD n = 5 (%)	GHB n = I I (%)	Heroin n = 4 (%)
Frequency, n (%)								
Decrease	15 (17.4)	8 (27.6)	11 (36.7)	8 (36.4)	22 (56.4)	3 (60.0)	2 (18.2)	2 (50.0)
No effect	58 (67.4)	18 (62.1)	15 (50.0)	12 (54.6)	17 (43.6)	2 (40.0)	7 (63.6)	2 (50.0)
Increase	13 (15.1)	3 (10.3)	4 (13.3)	2 (9.1)	0 (0.0)	0 (0.0)	2 (18.2)	0 (0.0)
Unknown	_	_	_	_	_	-	_	-
Duration, n (%)								
Decrease	13 (15.1)	7 (24.1)	7 (23.3)	8 (36.4)	18 (46.2)	l (20.0)	l (9.1)	2 (50.0)
No effect	57 (66.3)	17 (58.6)	20 (66.7)	11 (50.0)	18 (46.2)	2 (40.0)	7 (63.6)	2 (50.0)
Increase	5 (5.8)	3 (10.3)	l (3.3)	_	_	-	l (9.1)	-
Unknown	11 (12.8)	2 (6.9)	2 (6.7)	3 (13.6)	3 (7.7)	2 (40.0)	2 (18.2)	-

Table 4. Perceived effects of use of different illicit drugs on duration of individual attacks and attack frequency for those who used illicit drugs during a cluster headache episode.

Illicit drugs: all drugs; MDMA: 3,4-methylenedioxymethamphetamine; PSI: psilocybin mushroom; LSD: lysergic acid diethylamide; GHB: gammahydroxybutyrate.

*Not all patients used the drugs during an attack.

GHB (9.1%; n=1), cannabis (5.8%; n=5) and MDMA (3.3%; n=1) were reported to extend the duration of individual cluster headache attacks.

Discussion

In this study, Dutch people with cluster headache more often used illicit drugs during their lifetime than people from the general Dutch population. In particular, cannabis, amphetamine, heroin and cocaine were more used by cluster headache patients. Although it would be thought that this is due to alleviatory effects, most users with cluster headache reported no effect of these illicit drugs on their cluster headache attack frequency or duration when used only during a cluster headache episode. A very limited number reported a positive effect of PSI and heroin on the duration of their individual cluster headache attacks and a positive influence of PSI and LSD on attack frequency. These possible influences of PSI, heroin and LSD should be interpreted with caution, as these drugs were used in only a small fraction of the cluster headache population.

Both in our cluster headache population and in the general Dutch population a higher prevalence of illicit drug use was seen in males, which is in line with earlier studies (5). Males more often exhibit risk-taking behaviour (17). Being a female cluster headache patient has been associated with a decreased response to acute treatment and with more painful nocturnal headache attacks (18). In contrast, it has been suggested that bout frequency and duration are lower in females compared to males (5). These gender differences might influence the use of illicit drugs in women. However, we did not observe an increased prevalence of illicit drug use in female patients.

An increased prevalence of PSI use was found in both females and males with cluster headache. This might be due to the fact that PSI has received Dutch media attention as an alternative cluster headache treatment (19). Its efficacy, however, is limited: In a small retrospective study an acute effect of PSI, and even a termination of a cluster headache period, was only found in half of the few patients studied (11). Further research is therefore needed to shed more light on the acute and prophylactic effects of PSI in cluster headache.

Each cluster headache age cohort in our study used more illicit drugs than their age matched cohort in the Dutch general population, except for the 18–24 and 25–30 age cohorts, which were, however, too small to reach significance. Overall lifetime use of illicit drugs was increased in the younger age groups independent of having cluster headache. This increased prevalence of illicit drug use in younger Dutch generations has been described before and has not been seen in other European countries except for Switzerland (20,21). This seems to confirm the role of cultural differences between countries in drug use.

The increased prevalence of illicit drug use in people with cluster headache compared to people with chronic pain, or further unspecified headache, suggests that increased use is specific for cluster headache and not linked to headache or chronic pain per se. There are several possible explanations for this finding.

First, the question remains whether some illicit drugs actually have alleviatory effects on cluster headache. It

is possible that certain illicit drugs may interact with the unknown process that causes cluster headache. Cannabis acts on cannabinoid receptors that are widespread throughout the brain (22). The hypothalamus has cannabinoid receptors and has been implied in the pathophysiology of cluster headache (23,24). In contrast to the sedative qualities of cannabis, cocaine is a strong stimulant, also known for its capacity as a local anaesthetic and vasoconstrictor (25,26). Intranasal cocaine administration is reported to block pain caused by a nitroglycerin-induced cluster headache attack in about 30 minutes (27). However, since the majority of respondents described illicit drugs to have no effect on their cluster headache attacks, it remains questionable whether the possible alleviatory effect of illicit drugs on cluster headache is the actual reason for the increased prevalence of use.

Second, there could be an association between cluster headache and a tendency for addictive behaviour, as suggested before (28,29). This would also be in line with our finding that people with cluster headache are more inclined to smoke.

Third, the reputation of illicit drugs among people with cluster headache combined with the attention that these substances receive on cluster headache blogs, social media, and in some recent publications, may stimulate more patients to try these illicit drugs to treat their cluster headache (30). Last, the placebo effect could have overestimated the effects attributed to the various illicit drugs.

Limitations of this study include the fact that the data analysis needed to be stratified for age groups and gender, because the population data received from the Statistics Netherlands consisted of categorical age data. Age and gender were thus not full co-variates in the analysis and it was not possible to correct for other variables. We therefore tried to give the reader insight into age (Figure 2) and gender distribution (Table 3). Because of the small number of cluster head-ache patients who used certain drugs, we expect that for

those drugs the comparison between females and males could have been underpowered. It is possible that differences were larger than we could demonstrate. The higher education level of cluster headache patients may be a bias of the internet-based recruitment. As higher education is protective towards drug use this might have negatively influenced the results. This could mean that the difference between drug use in the cluster headache versus the general population would be even larger than the difference we found.

The headache subgroup of the general population could have included cluster headache patients. We included data from a representative sample of the Dutch general population (n = 14542), and cluster headache has a known prevalence of 1 in 1000 patients (31). As such, the sample could include 14–15 cluster headache patients. We expect that all these possible cluster headache patients are listed in the headache subgroup, which would amount to 14–15 out of 2268 people. Even if this were so, we expect that this small number of potential cluster headache patients did not influence the outcome of the headache subgroup.

Our questionnaire did not ask about motives for drug use and the time between filling out the questionnaire and drug use itself. Furthermore, it should be noted that our findings on effects of illicit drugs on cluster headache were all self-reported and should thus be met with caution, since the placebo effect could have overestimated the effects.

In conclusion, in the Dutch cluster headache population there is a higher prevalence of illicit drug use compared to the general Dutch population. This might be due to an actual acute or prophylactic effect, but also to a common pathophysiology between cluster headache and sensitivity for drug use. Another explanation could be a false conviction in people desperately seeking relief of their cluster headache and/or to the almost mythological reputation of illicit drugs in the cluster headache community.

Article highlights

- People with cluster headache have a higher prevalence of illicit drug use compared to the general Dutch population.
- Most illicit drug users report no effect of illicit drugs on attack duration and attack frequency.
- A limited number report a positive effect of PSI and heroin on attack duration. A positive influence on attack frequency is reported for PSI and LSD use. However, these drugs were used in only a small fraction of the cluster headache population studied.

Declaration of conflicting interests

The authors declared the following potential conflicts of interest with respect to the research, authorship, and/or publication of this article: MD Ferrari reports grants and consultancy or industry support from Medtronic, Amgen, Lilly and Electrocore, and independent support from NWO, ZonMW, NIH, European Community, Dutch Heart Foundation, and Fonds Nuts Ohra. This study was supported by grants from the Netherlands Organisation for Scientific Research (NWO) and the European Union's Seventh Framework programme (2007-2013) under grant agreement no. 602633 to MDF. IF de Coo reports support for conference visits from Electrocore. WC Naber, J Haan and R Fronczek report no conflict of interest.

Funding

The authors received no financial support for the research, authorship, and/or publication of this article.

ORCID iD

Ilse F de Coo (D) http://orcid.org/0000-0001-8169-4451

References

- 1. Headache Classification Committee of the International Headache Society (IHS). The International Classification of Headache Disorders, 3rd edition (beta version). *Cephalalgia* 2013; 33: 629–808.
- Fischera M, Marziniak M, Gralow I, et al. The incidence and prevalence of cluster headache: A metaanalysis of population-based studies. *Cephalalgia* 2008; 28: 614–618.
- May A. Cluster headache: Pathogenesis, diagnosis, and management. *Lancet* 2005; 366: 843–855.
- Di Lorenzo C, Coppola G, Di Lorenzo G, et al. The use of illicit drugs as self-medication in the treatment of cluster headache: Results from an Italian online survey. *Cephalalgia* 2016; 36: 194–198.
- Rossi P, Allena M, Tassorelli C, et al. Illicit drug use in cluster headache patients and in the general population: A comparative cross-sectional survey. *Cephalalgia* 2012; 32: 1031–1040.
- Donnet A, Lanteri-Minet M, Guegan-Massardier E, et al. Chronic cluster headache: A French clinical descriptive study. *J Neurol Neurosurg Psychiatry* 2007; 78: 1354–1358.
- Leroux E, Taifas I, Valade D, et al. Use of cannabis among 139 cluster headache sufferers. *Cephalalgia* 2013; 33: 208–213.
- Robbins MS, Bronheim R, Lipton RB, et al. Depression and anxiety in episodic and chronic cluster headache: A pilot study. *Headache* 2012; 52: 600–611.
- Rossi P, Torelli P, Di Lorenzo C, et al. Use of complementary and alternative medicine by patients with cluster headache: Results of a multi-centre headache clinic survey. *Complement Ther Med* 2008; 16: 220–227.

- Robbins MS, Tarshish S, Solomon S, et al. Cluster attacks responsive to recreational cannabis and dronabinol. *Headache* 2009; 49: 914–916.
- Sewell RA, Halpern JH and Pope HG Jr. Response of cluster headache to psilocybin and LSD. *Neurology* 2006; 66: 1920–1922.
- Hidalgo H, Uhl V, Gantenbein AR, Sandor PS, et al. Efficiency of sodium oxybate in episodic cluster headache. *Headache* 2013; 53: 1490–1491.
- 13. Khatami R, Tartarotti S, Siccoli MM, et al. Long-term efficacy of sodium oxybate in 4 patients with chronic cluster headache. *Neurology* 2011; 77: 67–70.
- Headache Classification Committee of the International Headache Society. The International Classification of Headache Disorders: 2nd edition. *Cephalalgia* 2004; 24: S9–S160.
- Wilbrink LA, Weller CM, Cheung C, et al. Stepwise web-based questionnaires for diagnosing cluster headache: LUCA and QATCH. *Cephalalgia* 2013; 33: 924–931.
- de Coo IF, Wilbrink LA, Haan J, et al. Evaluation of the new ICHD-III beta cluster headache criteria. *Cephalalgia* 2016; 36: 547–551.
- Malmberg M, Overbeek G, Monshouwer K, et al. Substance use risk profiles and associations with early substance use in adolescence. *J Behav Med* 2010; 33: 474–485.
- Rozen TD and Fishman RS. Female cluster headache in the United States of America: What are the gender differences? Results from the United States Cluster Headache Survey. J Neurol Sci 2012; 317: 17–28.
- Beemsterboer T. Hiervan gaat je hoofdpijn weg. Het is alleen verboden. NRC, https://www.nrc.nl/nieuws/2012/ 11/26/hiervan-gaat-je-hoofdpijn-weg-het-is-alleen-verboden-12582678-a598677 (2012, accessed 26 September 2018).
- Monshouwer K, Smit F, de Graaf R, et al. First cannabis use: Does onset shift to younger ages? Findings from 1988 to 2003 from the Dutch National School Survey on Substance Use. *Addiction* 2005; 100: 963–970.
- Muller S and Gmel G. Changes in the age of onset of cannabis use: Results of the 2nd Swiss Health Survey 1997. Soz Praventivmed 2002; 47: 14–23.
- 22. Matsuda LA, Lolait SJ, Brownstein MJ, et al. Structure of a cannabinoid receptor and functional expression of the cloned cDNA. *Nature* 1990; 346: 561–564.
- 23. Iversen L. Cannabis and the brain. *Brain* 2003; 126: 1252–1270.
- May A, Bahra A, Büchel C, et al. Hypothalamic activation in cluster headache attacks. *Lancet* 1998; 352: 275–278.
- Goldstein RA, DesLauriers C and Burda AM. Cocaine: History, social implications, and toxicity – a review. *Dis Mon* 2009; 55: 6–38.
- Harper SJ and Jones NS. Cocaine: What role does it have in current ENT practice? A review of the current literature. J Laryngol Otol 2006; 120: 808–811.

- 27. Costa A, Pucci E, Antonaci F, et al. The effect of intranasal cocaine and lidocaine on nitroglycerin-induced attacks in cluster headache. *Cephalalgia* 2000; 20: 85–91.
- 28. Govare A and Leroux E. Licit and illicit drug use in cluster headache. *Curr Pain Headache Rep* 2014; 18: 413.
- 29. May A. Illicit drugs and cluster headache: An inevitable discussion. *Cephalalgia* 2012; 32: 1021–1022.
- Andersson M, Persson M and Kjellgren A. Psychoactive substances as a last resort – a qualitative study of selftreatment of migraine and cluster headaches. *Harm Reduct J* 2017; 14: 60.
- 31. Nesbitt AD and Goadsby PJ. Cluster headache. *BMJ* 2012; 344: e2407.