ORIGINAL RESEARCH



Acute Treatment Patterns, Migraine Burden, and Healthcare Resource Use in People With Migraine: Results From the OVERCOME (EU) Observational Study

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ABSTRACT

Introduction: The ObserVational survey of the Epidemiology, tReatment and Care Of MigrainE (OVERCOME) European Union (EU) is part of an overarching population-based study program that also includes the United States and Japan. Here, we report data on the migraine/severe headache burden and the use

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Methods: OVERCOME (EU) was an online, non-interventional, cross-sectional survey conducted in adults in Spain and Germany between October 2020 and February 2021. A total migraine cohort was established based on health survey participants who reported headache/migraine in the last 12 months AND identified as having migraine based on modified International Classification of Headache Disorders, third edition criteria OR self-reported physician diagnosis. Data were analyzed for the total migraine cohort and the subcohort with moderate to severe headache attacks, with average pain severity ≥ 5 points, pain duration > 4 h, and at least moderate disability due to migraine [Migraine Disability Assessment (MIDAS) score > 11] over the past 3 months. Results: Pain of moderate or severe intensity was the most frequent symptom in the total migraine cohort (n = 19,103/20,756; 92.0%). Proportions of participants reporting severe disability (MIDAS Grade IV), poorer quality of life (QoL; Migraine-Specific QoL Questionnaire), and higher interictal burden (Migraine Interictal Burden Scale-4), generally increased with number of headache days (HDs)/month. Most participants (92.5%) reported current acute migraine/severe headache medication use, although only 39.0% were using triptans. In the moderate to severe attacks subcohort (n = 5547), 48.4% were using triptans, with

nonsteroidal anti-inflammatory drugs the most common acute medication. The moderate to severe attacks subcohort also reported poorer QoL and greater pain and disability with increasing HDs/month, although severe interictal burden was reported for $\sim 60\%$ of participants regardless of HDs/month. Treatment satisfaction (six-item migraine Treatment Optimization Questionnaire) in those using triptans was generally poor in both total and subcohorts.

Conclusion: High migraine-related burden levels were reported, despite use of acute medication. Although triptans are recommended for moderate to severe migraine attacks in Spanish and German guidelines, less than half of participants were using triptans; treatment satisfaction in those using triptans was generally poor. New tailored treatment options may help address unmet needs in current acute treatment.

Keywords: Migraine; Severe headache; Migraine symptoms; Burden of migraine; Acute treatment patterns; Treatment satisfaction

Key Summary Points

Why carry out this study?

Migraine is associated with disability, pain, lost productivity, healthcare utilization, and reduced quality of life.

As most published real-world migraine treatment and burden data are from observational or registry-based studies, which focus on healthcare settings, they will not capture the experiences of the many people with migraine who do not seek professional medical care.

What did the study ask?/What was the hypothesis of the study?

The population-based ObserVational survey of the Epidemiology, tReatment and Care Of MigrainE (OVERCOME) European Union (EU) survey aimed to further understand current treatment patterns, and the burden and unmet needs of people with migraine/severe headache in Spain and Germany; here, we report data regarding the use of acute medication and healthcare resources.

What was learned from the study?

Our study found high migraine-related burden levels, despite the use of acute medication; the proportions of participants reporting severe disability, poorer quality of life, and higher interictal burden generally increased with the number of headache days/month.

Although triptans are recommended for moderate to severe migraine attacks in Spanish and German guidelines, less than half of participants (39.0% in the overall total migraine cohort, 48.4% of a subcohort with moderate to severe attacks) were using triptans, and treatment satisfaction in those using triptans was generally poor.

New tailored treatment options may help address unmet needs in current acute treatment.

INTRODUCTION

Migraine is a debilitating neurological condition with an overall prevalence of approximately 15% in Europe [1], and 1-year prevalences of 12.6% in Spain [2] and 16.6% in Germany [3]. Reported prevalence estimates vary between studies, possibly due to differences in study design and/or inclusion criteria [4, 5]. Migraine is more prevalent in females (17.6%) than males (8%) in Europe [1], and is most common between the ages of 15 and 49 years, with the highest prevalence (\sim 30%) in females aged 35–39 years [6].

The World Health Organization ranks migraine as the second leading cause of disability overall (based on years lived with disability), and the leading cause of disability in women aged 15–49 years [7]. The pain, physical impairment, and nausea/vomiting associated with migraine attacks, along with the interictal burden and unpredictability of attacks, can result in significant disruption to daily life, missed school/work days, medication overuse, lower health-related quality of life (HRQoL), increased healthcare utilization, and stigma [8–14]. Lost productivity due to migraine in Europe has been estimated to cost \sim 110 billion euros per year [13].

Even though people with migraine may visit a range of healthcare providers (HCPs), many have never seen a physician and use over-thecounter (OTC) or even no medication [14-16]. Triptans are the recommended first-choice treatment for moderate to severe migraine attacks in both Spanish and German guidelines [17, 18], and are available on prescription or OTC in Germany but are prescription-only in Spain. Although the reported proportions of patients using triptans vary between countries, they are generally reported as low [19]. On a general migraine patient population basis, the Eurolight study reported triptan use rates between 3.4% (Lithuania) and 22.4% (Spain, workplace setting), with 11.0% reported for Germany [19]. However, higher rates have been reported for studies among members of lav headache organizations [19].

Most published data on real-world migraine treatment and burden are from observational or registry-based studies, which focus on healthcare settings. However, as many people do not seek professional medical care for migraine, their experiences will not be captured in these studies. New population-based studies are required to further understand current treatment patterns, and the burden and unmet needs of people with migraine in Europe and globally. The ObserVational survey of the Epidemiology, tReatment and Care Of MigrainE (OVERCOME) European Union (EU) is part of an overarching study program that also includes the United States and Japan [20–22]; OVERCOME (EU) reports data from Spain and Germany.

Here, we report the data on the burden of migraine/severe headache and the use of acute medication and healthcare resources in Spain and Germany. The use of preventive medication by participants in this study is reported elsewhere [23]. We also report findings of a subcohort analysis of participants with moderate to severe headache attacks to determine the burden in this population, and whether these people are using triptans in line with Spanish and German guidelines [17, 18].

METHODS

Study Design

OVERCOME (EU) was an online, non-interventional, cross-sectional, observational survey conducted in adults (aged \geq 18 years) in Spain and Germany between October 2020 and February 2021. Full details of the study design and methods, including the survey instrument used, are published elsewhere [23]. Briefly, participants registered in existing opt-in online survey panels [Kantar Profiles (Lightspeed) global panel and its partners] [24] were invited to participate in a health survey without knowledge of the specific health topic. This study was performed in accordance with the Helsinki Declaration of 1964 and its later amendments. Ethical review board approval for the study was granted by the Marqués de Valdecilla University Hospital (Spain) and participants were required to provide informed consent prior to commencing the study by selecting "I agree to participate" on the online form. All survey respondents provided informed consent and all data were anonymized before analysis.

Three phases were used to establish the migraine cohort [23]. In phase I, a random

sample was selected based on pre-specified demographics (e.g., age and sex) to ensure (to the extent possible) that the data collected were representative of the Spanish and German adult population. In phase II, respondents were assigned to two cohorts, a migraine cohort and a non-migraine control cohort. The migraine cohort included people with a reported headache/migraine (any type) in the last 12 months AND identified as having migraine based on the modified International Classification of Headache Disorders, third edition (ICHD-3) screening criteria [25] OR self-reported physician diagnosis of migraine (i.e., where a participant reported having received this diagnosis from a medical professional). In phase III, participants in the migraine cohort were invited to complete further questions to collect data on demographics, clinical features of migraine, patientreported outcomes, and the use of medication.

Data were analyzed for the total migraine cohort and for the subcohort with moderate to severe headache attacks over the past 3 months (moderate to severe attacks subcohort). Moderate to severe headache attacks were defined as being of average pain severity \geq 5 points [based on the Migraine Disability Assessment (MIDAS) question on pain severity], pain duration \geq 4 h, AND moderate or severe disability (MIDAS score \geq 11) due to migraine.

Study Objectives and Analyses

The primary objectives of the OVERCOME (EU) study were to understand diagnosis, barriers to treatment, and treatment patterns of people with migraine in Spain and Germany. A summary of the main categories of survey questions has been published previously [23].

The analyses reported here are focused on symptoms, duration of migraine/severe headache, pain severity, migraine-related disability, and burden/quality of life (QoL), together with HCP visits, acute migraine medication treatment patterns (including the use of acute treatment in combination with preventive medication), and treatment satisfaction.

Migraine-Specific Assessments

Migraine-related data were assessed using patient-reported outcome measures or by asking specific questions to participants. Migraine-related disability was assessed using the MIDAS scale [26-28]. MIDAS is a brief, self-administered, questionnaire containing five questions that quantifies headache disability based on the number of days a person has missed or had reduced productivity at work, home, or social settings over the past 3 months. Two additional questions about the number of headaches and average associated pain level are not used in scoring but can help physicians in better understanding the clinical features of headache for each affected person [27]. Disability grades are then assigned based on these number of days, with higher scores indicating more severe disability: Grade I = little or no disability (MIDAS score 0-5); Grade II = mild (score 6-10); Grade III = moderate (score 11-20): and Grade IV = severe (score > 21). The MIDAS instrument is considered reliable and valid and is correlated with clinical judgment regarding the need for medical care [29, 30]. Spanish and German versions of the MIDAS instrument have also been validated [31, 32]. Pain severity was assessed through the specific question of the MIDAS: "On a scale of 0 to 10 (where 0 = nopain at all and 10 = pain as bad as it can be), on average how painful are your migraine or severe headaches?".

Headache duration was assessed through another specific survey question: "When you have a migraine or severe headache, how long does it usually last?".

Satisfaction with migraine treatment was assessed using the six-item migraine Treatment Optimization Questionnaire (mTOQ-6) [33, 34]. The mTOQ-6 is a validated, self-administered questionnaire that assesses the efficacy of current acute migraine treatment, and asks questions about the medications the patient is currently using to treat headaches (recall period 3 months). The items assess the domains of quick return to function, 2-h pain free, sustained 24-h pain relief, tolerability, comfortable making plans, and perceived control.

	Total	HDs/month				
	(<i>N</i> = 20,756)	$\overline{0-3}$ (<i>N</i> = 13,759)	4-7 (<i>N</i> = 4203)	8-14 (<i>N</i> = 1730)	≥ 15 (N = 1064)	
Mean (SD) age, years	40.4 (13.5)	40.1 (13.5)	40.7 (13.3)	41.2 (13.5)	42.2 (13.9)	
Mean (SD) age at migraine diagnosis, years	24.2 (10.8)	24.0 (10.6)	24.4 (11.0)	25.1 (11.0)	24.3 (11.6)	
Female, n (%)	12,512 (60.3)	7,846 (57.0)	2717 (64.6)	1177 (68.0)	772 (72.6)	
ICHD-3 criteria met, n (%)	18,987 (91.5)	12,541 (91.1)	3872 (92.1)	1611 (93.1)	963 (90.5)	
Previously diagnosed with migraine by healthcare provider, <i>n</i> (%)	11,948 (57.6)	7269 (52.8)	2805 (66.7)	1169 (67.6)	705 (66.3)	
Most frequently reported health conditions (\geq 20% in total m	nigraine cohort),	n (%)			
Allergies/hay fever	8043 (38.8)	5161 (37.5)	1724 (41.0)	716 (41.4)	442 (41.5)	
Anxiety	5430 (26.2)	3257 (23.7)	1204 (28.6)	571 (33.0)	398 (37.4)	
High cholesterol/lipids	5122 (24.7)	3217 (23.4)	1071 (25.5)	502 (29.0)	332 (31.2)	
Depression	4978 (24.0)	2788 (20.3)	1148 (27.3)	592 (34.2)	450 (42.3)	
Hypertension	4875 (23.5)	3034 (22.1)	1004 (23.9)	496 (28.7)	341 (32.0)	
MIDAS grade ^a , total migraine cohort, n (%)						
I—little or no disability	8177 (39.4)	6689 (48.6)	1043 (24.8)	291 (16.8)	154 (14.5)	
II—mild disability	3976 (19.2)	2988 (21.7)	728 (17.3)	180 (10.4)	80 (7.5)	
III—moderate disability	4004 (19.3)	2438 (17.7)	1036 (24.6)	382 (22.1)	148 (13.9)	
IV—severe disability	4599 (22.2)	1644 (11.9)	1396 (33.2)	877 (50.7)	682 (64.1)	

Table 1 Participant sociodemographic and migraine-related clinical characteristics (total migraine cohort)

HDs headache days, ICHD-3 International Classification of Headache Disorders, third edition, MIDAS Migraine Disability Assessment, SD standard deviation

^aHigher MIDAS scores indicate greater disability: Grade I = score 0–5; Grade II = score 6–10; Grade III = score 11–20; Grade IV = score ≥ 21

Response options utilize a Likert-type scale of "never" or "rarely" (scored as 0), "less than half the time" (scored as 1), or "half the time or more" (scored as 2). Treatment efficacy is categorized as "very poor" = 0, "poor" = 1–5, "moderate" = 6–7, and "maximum" = 8.

Health status (i.e., physical and emotional limitations of specific concern to individuals suffering from migraine headaches) was assessed using the Migraine-Specific QoL Questionnaire (MSQ) Version 2.1. This instrument consists of 14 items that address three domains: (1) Role Function-Restrictive; (2) Role Function-Preventive; and (3) Emotional Function [35].

Raw scores for each dimension are computed as a sum of item responses, the collective sum providing a total raw score that is converted to a 0–100 scale, with higher scores indicating a better HRQoL. The instrument was designed with a 4-week recall period, and is considered reliable, valid, and sensitive to change in migraine [35, 36].

The burden related to headache in the time between attacks was measured using the Migraine Interictal Burden Scale-4 (MIBS-4) instrument. The MIBS-4 consists of four items that address disruption at work and school, diminished family and social life, difficulty

	Total	HDs/month					
	(<i>N</i> = 20,756)	0-3 (<i>N</i> = 13,759)	4-7 (<i>N</i> = 4203)	8–14 (<i>N</i> = 1730)	≥ 15 (N = 1064)		
$\overline{\text{Most frequently reported symptoms}^a (\geq 70\%)}$	in total migrain	e cohort), <i>n</i> (%)					
Pain has moderate or severe intensity	19,103 (92.0)	12,445 (90.4)	3976 (94.6)	1660 (96.0)	1022 (96.1)		
Sound is bothersome	17,366 (83.7)	11,286 (82.0)	3640 (86.6)	1505 (87.0)	935 (87.9)		
Pain is pounding, pulsating or throbbing	17,243 (83.1)	11,146 (81.0)	3679 (87.5)	1505 (87.0)	913 (85.8)		
Light is bothersome	16,896 (81.4)	11,004 (80.0)	3530 (84.0)	1459 (84.3)	903 (84.9)		
Pain worse on just one side	15,926 (76.7)	10,125 (73.6)	3430 (81.6)	1457 (84.2)	914 (85.9)		
Pain is made worse by routine activities such as walking or climbing stairs	15,040 (72.5)	9686 (70.4)	3218 (76.6)	1319 (76.2)	817 (76.8)		

Table 2 Most frequently reported (≥ 70% in total migraine cohort) migraine symptoms (total migraine cohort)

HDs headache days

^aSymptoms occurring with a Migraine Symptom Severity Score of ≥ 3 less than half of the time or more

planning, and emotional difficulty. The questionnaire specifically asks about the effect of the disease over the past 4 weeks on days without a headache attack. Each response has an associated numerical score, with the sum across all four items resulting in a total score ranging from 0 to 12. The level of interictal burden is categorized into the following: 0 for none, 1–2 mild, 3–4 moderate, and \geq 5 severe [12, 37].

Statistical Analyses

Data from the total migraine cohort, moderate to severe attacks subcohort, and specific subgroups [based on different frequencies of headache days per month (HDs/month)] were analyzed using descriptive statistics. Continuous variables are reported as means with standard deviations (SDs), or medians and ranges, as appropriate. Categorical variables are summarized as frequencies and percentages. SAS Version 9.4 software (SAS Institute, Cary, NC, USA) was used to undertake all analyses.

RESULTS

Total Migraine Cohort

Participants

A total of 20,756 participants were included in the total OVERCOME (EU) migraine cohort (Table 1). Respondents had a mean age of 40.4 years, a mean age at diagnosis of 24.2 years, and 60.3% were female. The proportion of participants with comorbidities, especially anxiety and depression, generally increased numerically with the number of HDs/month. Severe disability (MIDAS Grade IV) was more commonly reported in participants with a greater number of HDs/month.

Symptoms, Duration, Pain Severity, and Burden of Migraine/Severe Headache

Pain of moderate or severe intensity was the most frequent migraine symptom, reported by 92.0% of participants in the total migraine cohort; other frequently reported symptoms are presented in Table 2. The proportion of participants reporting symptoms generally increased with the number of HDs/month. Most migraine/severe headache attacks were of

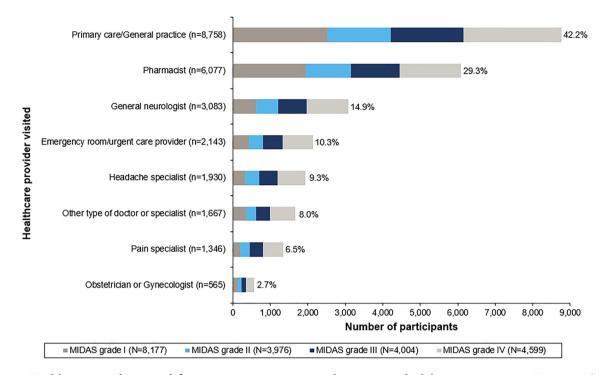


Fig. 1 Healthcare providers visited for migraine or severe headaches at least once in the prior 12 months (total migraine cohort; N = 20,756). Higher MIDAS scores

4–71 h duration, although around 1 in 10 participants in the \geq 15 HDs/month group reported headache durations of \geq 72 h (Table 3). The mean (SD) pain severity score in the total migraine cohort was 6.2 (1.9). Pain intensity increased numerically with a greater number of HDs/month.

Participants with a higher number of HDs/month also reported poorer QoL and higher interictal burden. Mean (SD) MSQ Role Function-Restrictive scores were 47.5 (21.5) in participants with \geq 15 HDs/month, compared with 67.2 (21.3) in those with 0–3 HDs/month. The overall proportion of participants reporting moderate interictal burden (MIBS-4) was 12.6%, with little variation according to HDs/month [range 12.3% (8–14 HDs/month) to 13.8% (\geq 15 HDs/month)], whereas severe interictal burden increased with the number of HDs/month, from 41.2% of participants with 0–3 HDs/month to 57.8% of those with \geq 15 HDs/month.

indicate greater disability: I = score 0–5; II = score 6–10; III = score 11–20; IV = score ≥ 21 . *MIDAS* Migraine Disability Assessment

HCP Visits

Just over half of visits [mean proportion of total HCP visits per participant: 52.5% (SD: 37.9%)] to an HCP in the past 12 months were due to migraine/severe headache. Primary care/general practice was the service accessed by most participants (42.2%) for migraine/severe headache, followed by the pharmacist (29.3%; Fig. 1), although one in ten participants (10.3%) contacted an emergency service/urgent care provider. Participants with higher MIDAS grades were more likely to visit specialists or emergency care, whereas those with lower grades were more likely to visit a pharmacist or primary care/general practice.

Acute Migraine Treatment Patterns and Treatment Satisfaction

Most participants (92.5%; 19,207/20,756) reported that they were currently using acute migraine/severe headache medication, with a mean of 4.9 (SD: 4.0) current acute medications per participant in the total migraine cohort

	Total (N = 20,756)	HDs/month					
		$\overline{0-3} \ (N=13,759)$	4-7 (N = 4203)	8–14 (<i>N</i> = 1730)	$\geq 15 \ (N = 1064)$		
When you ha	When you have a migraine or severe headache, how long does it usually last? <i>n</i> (%)						
<4 h	7853 (37.8)	5838 (42.4)	1271 (30.2)	511 (29.5)	233 (21.9)		
4–71 h	12,458 (60.0)	7773 (56.5)	2826 (67.2)	1155 (66.8)	704 (66.2)		
\geq 72 h	445 (2.1)	148 (1.1)	106 (2.5)	64 (3.7)	127 (11.9)		
Average pain	severity (according to M	(IDAS)					
Mean (SD)	6.2 (1.9)	5.9 (2.0)	6.6 (1.6)	6.9 (1.6)	7.2 (1.6)		

 Table 3 Duration of migraine/severe headache (total migraine cohort)

HDs headache days, MIDAS Migraine Disability Assessment, SD standard deviation

(Table 4). The most common categories of acute medication were OTC analgesics, followed by nonsteroidal anti-inflammatory drugs (NSAIDs) on prescription, and paracetamol. Fewer than half (39.0%) of participants currently using acute migraine medication were using triptans; the use of triptans was generally consistent across subgroups based on HDs/month but was slightly lower in people with 0–3 HDs/month (36.2%) compared with those with 4–7 (44.6%), 8–14 (45.1%), or \geq 15 (41.4%) HDs/month.

Triptan use was more common in participants using preventive migraine medication, with 78.5% of those using current preventive medication using triptans (Table 4; Supplementary Table S1). NSAIDs on prescription, OTC analgesics, and suppositories were also commonly used in combination with preventive medication.

Responses to the mTOQ-6 indicated that total rates of poor and very poor treatment satisfaction were higher in people currently taking triptans compared with those using acute medications excluding triptans (Fig. 2a). Use of triptans and acute medications excluding triptans according to number of HDs/month and MIDAS grade is presented in Table 5. The proportions of patients using triptans and acute medications excluding triptans were both highest in patients with MIDAS Grade IV and > 8HDs/month. However, > 60%of patients with MIDAS Grade IV and either 8-14 (61.0%) or ≥ 15 (68.8%) HDs/month were using triptans, whereas the use of acute medications excluding triptans was 43.4% in those with 8–14 HDs/month and 60.9% in those with \geq 15 HDs/month. Many participants (n = 5538) reported that they had stopped using triptans; the most common reason for stopping triptans being "other medications work better for my migraine/severe headaches" (Fig. 2b).

Analyses of Participants in the Moderate to Severe Attacks Subcohort

Participants and Acute Medication Usage

A total of 5547 of the 20,756 participants in the total migraine cohort (26.7%) experienced moderate to severe headache attacks [average pain severity \geq 5 points (based on the MIDAS question on pain severity), pain duration \geq 4 h, and at least moderate disability due to migraine (MIDAS score \geq 11)] and were included in the moderate to severe attacks subcohort (Table 6).

Respondents in the moderate to severe attacks subcohort had a mean age of 39.8 years and 66.7% were female. Just under half of this subcohort (48.4%) were using triptans, and NSAIDs were the most commonly used acute treatment.

Migraine-Related Burden, QoL, Pain, and Disability

Participants in the moderate to severe attacks subcohort generally reported increased burden with increasing headache/migraine severity

	Total	HDs/month						
	(N = 19,207)	0-3 (<i>N</i> = 12,569)	4–7 (<i>N</i> = 3989)	8–14 (<i>N</i> = 1659)	≥ 15 (N = 990)			
Number of current acute medicat	ions per participar	nt						
Mean (SD)	4.9 (4.0)	4.6 (3.7)	5.3 (4.2)	5.6 (4.5)	5.6 (5.0)			
Median (IQR)	4.0 (2.0-6.0)	4.0 (2.0-6.0)	4.0 (2.0–7.0)	4.0 (3.0–7.0)	4.0 (2.0–7.0)			
Number of participants using cur	rent acute medicat	tions per category,	n (%)					
Triptans ^a	7493 (39.0)	4554 (36.2)	1780 (44.6)	749 (45.1)	410 (41.4)			
Ergot alkaloids ^b	412 (2.1)	255 (2.0)	88 (2.2)	50 (3.0)	19 (1.9)			
NSAIDs (on prescription)	10,647 (55.4)	6655 (52.9)	2349 (58.9)	1033 (62.3)	610 (61.6)			
Paracetamol (prescription or OTC)	10,237 (53.3)	6657 (53.0)	2165 (54.3)	893 (53.8)	522 (52.7)			
OTC analgesics ^c	13,532 (70.5)	8884 (70.7)	2873 (72.0)	1154 (69.6)	621 (62.7)			
Other prescription drugs	7094 (36.9)	4199 (33.4)	1614 (40.5)	764 (46.1)	517 (52.2)			
Other OTC drugs	2921 (15.2)	1695 (13.5)	676 (16.9)	340 (20.5)	210 (21.2)			
Suppositories ^d (prescription or OTC)	6042 (31.5)	4104 (32.7)	1203 (30.2)	488 (29.4)	247 (24.9)			
Number of participants using cur	rent preventive me	edication for migra	uine/severe heada	che, n (%)				
Current use	4710 (22.7)	2932 (21.3)	1009 (24.0)	449 (26.0)	320 (30.1)			
Combinations of current acute ar	nd preventive med	ications						
Number of participants using cu	rrent preventive n	nedication combin	ed with: $n (\%)^{e}$					
Any current acute medication	4598	2852	993	444	309			
Triptans ^a	3611 (78.5)	2260 (79.2)	795 (80.1)	346 (77.9)	210 (68.0)			
Ergot alkaloids ^b	280 (6.1)	177 (6.2)	60 (6.0)	29 (6.5)	14 (4.5)			
NSAIDs (on prescription)	3536 (76.9)	2168 (76.0)	777 (78.2)	354 (79.7)	237 (76.7)			
Paracetamol (prescription or OTC)	1954 (42.5)	1101 (38.6)	490 (49.3)	229 (51.6)	134 (43.4)			
OTC analgesics ^c	3149 (68.5)	1993 (69.9)	692 (69.7)	291 (65.5)	173 (56.0)			
Other prescription drugs	2270 (49.4)	1300 (45.6)	526 (53.0)	247 (55.6)	197 (63.8)			
Other OTC drugs	561 (12.2)	276 (9.7)	143 (14.4)	87 (19.6)	55 (17.8)			

Table 4 Current acute migraine/severe headache medication use in participants using acute migraine/severe headache medications (total migraine cohort)

	Total	HDs/month				
	(<i>N</i> = 19,20 7)	0-3 (<i>N</i> = 12,569)	4–7 (<i>N</i> = 3989)	8–14 (<i>N</i> = 1659)	≥ 15 (<i>N</i> = 990)	
Suppositories ^d (prescription or OTC)	3383 (73.6)	2314 (81.1)	666 (67.1)	260 (58.6)	143 (46.3)	

Table 4 continued

HDs headache days, IQR interquartile range, NSAID nonsteroidal anti-inflammatory drug, OTC over-the-counter, SD standard deviation

^aTriptans are available on prescription or OTC in Germany but are prescription-only in Spain

^bErgot alkaloids are only reimbursed in Germany

^ce.g., aspirin and loxoprophen

^dAny suppository used to treat or relieve migraine or severe headache (data were not analyzed according to the type of suppository used). Sumatriptan is available as a suppository in Germany but not in Spain

^ePercentages are based on the number of participants taking each class of preventive medication combined with any current acute medication

(Fig. 3). Patients with a higher number of HDs/month reported poorer QoL (MSQ), higher pain levels (MIDAS), and more severe migraine-related disability (MIDAS) than those with fewer HDs/month, although severe interictal burden (MIBS-4) was reported for $\sim 60\%$ of participants regardless of the number of HDs/month.

Triptan Use and Treatment Satisfaction in Participants Who Sought Care

Although > 80% of participants in the moderate to severe attacks subcohort sought medical care, less than half were diagnosed with migraine and/or used triptans (Fig. 4). Of those who did use triptans, treatment satisfaction was generally poor regardless of HDs/month.

DISCUSSION

This study provides population data for realworld migraine burden and treatment patterns in Spain and Germany and illustrates the negative effects of migraine/severe headache on QoL, pain, disability, and healthcare resource use, despite the availability of current treatment options.

Pain of moderate or severe intensity was the most frequently reported migraine symptom

(92.0% of participants in the total migraine cohort), and pain intensity and the proportion of participants reporting symptoms generally increased with the number of HDs/month. Participants with a higher number of HDs/month also reported poorer QoL (MSQ) higher and interictal burden (MIBS-4). Migraine/severe headache also accounted for just over half of visits to an HCP in the past 12 months; people with higher MIDAS grades were more likely to visit specialists or emergency care, whereas pharmacist or primary care/general practice visits were more likely for those with lower grades.

Our analysis of participants in the moderate to severe attacks subcohort [average pain severity \geq 5 points (based on the MIDAS question on pain severity), pain duration \geq 4 h, and at least moderate disability due to migraine (MIDAS score \geq 11)] also found burden generally increased with increasing headache/migraine severity. Patients with a higher number of HDs/month reported poorer QoL, higher pain levels (MIDAS), and more severe migrainerelated disability (MIDAS) than those with fewer HDs/month, although severe interictal burden was reported for ~ 60% of participants regardless of the number of HDs/month.

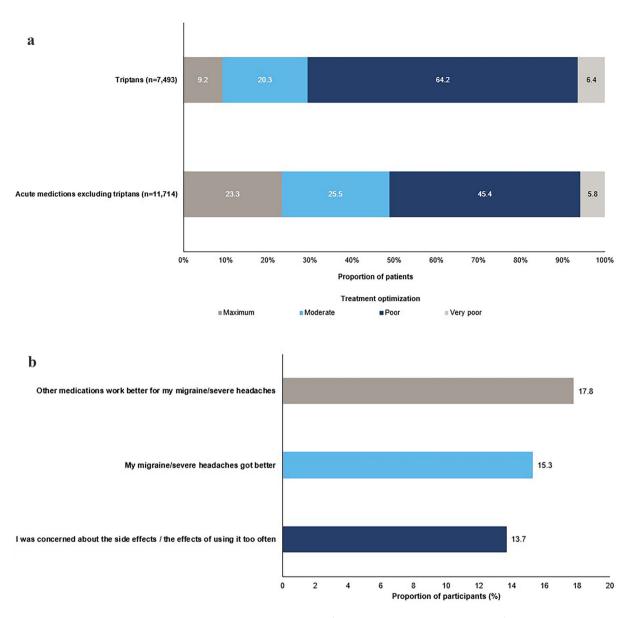


Fig. 2 a Migraine Treatment Optimization Questionnaire (mTOQ-6) responses in participants currently using triptans versus acute medications excluding triptans. b Most common reasons for discontinuing triptan use

(total migraine cohort; n = 5538). Participants could be taking more than one acute medication, and could give more than one reason for no longer taking triptans

Despite the high levels of burden reported, the data also highlight a low use of triptans, even in people with moderate to severe headache attacks, and generally poor treatment satisfaction in those who use triptans. These findings are concerning, given that triptans are the recommended treatment for moderate to severe migraine attacks in both Spanish and German guidelines [17, 18], and illustrate an unmet need for new treatment options for patients with migraine or severe headache.

Most participants in the total migraine cohort reported that they were currently using acute migraine/severe headache medication, although less than half of these were using triptans; the most common categories of acute

	Total	HDs/month						
		0-3	4-7	8-14	≥ 15			
Participants currently using trij	otans							
	<i>n</i> = 7493	n = 4554	n = 1780	n = 749	n = 410			
MIDAS score, median (IQR)	12.0 (6.0–25.0)	10.0 (5.0–17.0)	16.0 (9.0-30.0)	27.0 (14.0-50.0)	35.0 (16.0–78.0)			
Numbers of participants in eac	h MIDAS grade	category, <i>n</i> (% of	population using	triptans)				
I—little or no disability	1,670 (22.3)	1,285 (28.2)	279 (15.7)	72 (9.6)	34 (8.3)			
II—mild disability	1,564 (20.9)	1,190 (26.1)	275 (15.4)	70 (9.3)	29 (7.1)			
III—moderate disability	1,912 (25.5)	1,190 (26.1)	507 (28.5)	150 (20.0)	65 (15.9)			
IV—severe disability	2,347 (31.3)	889 (19.5)	719 (40.4)	457 (61.0)	282 (68.8)			
	Total	HDs/month						
		0-3	4-7	8-14	≥15			
Participants currently using acu	ite medication exe	cluding triptans						
	n = 11,714	n = 8015	n = 2209	n = 910	n = 580			
MIDAS score, median (IQR)	6.0 (2.0–15.0)	4.0 (1.0–10.0)	10.0 (4.0-23.0)	17.0 (8.0-38.0)	33.0 (10.0-80.0)			
Numbers of participants in eac	h MIDAS grade	category, <i>n</i> (% of	population using	acute medication e	xcluding triptans)			
I—little or no disability	5645 (48.2)	4654 (58.1)	692 (31.3)	196 (21.5)	103 (17.8)			
II—mild disability	2175 (18.6)	1606 (20.0)	417 (18.9)	106 (11.6)	46 (7.9)			
III—moderate disability	1875 (16.0)	1100 (13.7)	484 (21.9)	213 (23.4)	78 (13.4)			
IV—severe disability	2019 (17.2)	655 (8.2)	616 (27.9)	395 (43.4)	353 (60.9)			

Table 5 Current acute migraine/severe headache medication use according to MIDAS grade^a (total migraine cohort)

^aHigher MIDAS scores indicate greater disability: Grade I = score 0–5; Grade II = score 6–10; Grade III = score 11–20; Grade IV = score ≥ 21

HDs headache days, IQR interquartile range, MIDAS Migraine Disability Assessment

medication were OTC analgesics, NSAIDs on prescription, and paracetamol, with many people taking multiple acute medications. Very few people were using ergot alkaloids, although these are only reimbursed in Germany and are not reimbursed in Spain. Triptans are available on prescription or OTC in Germany but are prescription-only in Spain. Triptan use was more common in participants who were also using preventive medication, and NSAIDs on prescription, OTC analgesics, and suppositories were also commonly used in combination with preventive medication. It could be speculated that the use of preventive medication may be a sign of a more migraine-specific approach to treatment, or that patients using preventive medication had higher disability and were therefore more likely to use triptans than those using acute medication alone. However, our analysis of triptan use in people with moderate to severe headache attacks (defined as average pain severity \geq 5 points, pain duration \geq 4 h, and MIDAS score \geq 11; n = 5547) also found that less than half (48.4%) of those with

	Total	HDs/month					
	(<i>N</i> = 5547)	$\overline{0-3}$ (<i>N</i> = 2329)	4-7 (<i>N</i> = 1697)	8–14 (N = 884)	≥ 15 $(N = 637)$		
Mean (SD) age, years	39.8 (12.8)	38.5 (12.5)	40.2 (12.6)	40.5 (12.9)	42.4 (13.4)		
Female, n (%)	3701 (66.7)	1404 (60.3)	1183 (69.7)	631 (71.4)	483 (75.8)		
Current acute medication usage, n (%)							
Triptans ^b	2687 (48.4)	1149 (49.3)	846 (49.9)	429 (48.5)	263 (41.3)		
Ergot alkaloids ^{c,d}	618 (11.1)	306 (13.1)	166 (9.8)	93 (10.5)	53 (8.3)		
NSAIDs	4943 (89.1)	2079 (89.3)	1532 (90.3)	792 (89.6)	540 (84.8)		
Paracetamol	2940 (53.0)	1229 (52.8)	933 (55.0)	470 (53.2)	308 (48.4)		
Combination analgesics	1385 (25.0)	548 (23.5)	421 (24.8)	232 (26.2)	184 (28.9)		
Combination of triptans/ergot alkaloids/ NSAIDs/paracetamol	4059 (73.2)	1733 (74.4)	1263 (74.4)	640 (72.4)	423 (66.4)		

 Table 6 Participant sociodemographic and characteristics and current acute medication usage (moderate to severe attacks subcohort^a)

^aModerate to severe headache attacks were defined as being of average pain severity \geq 5 points (based on the MIDAS question on pain severity), pain duration \geq 4 h, and at least moderate disability due to migraine (MIDAS score \geq 11) *HDs* headache days, *MIDAS* Migraine Disability Assessment, *NSAID* nonsteroidal anti-inflammatory drug, *OTC* over-the-counter, *SD* standard deviation

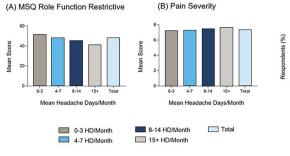
^bTriptans are available on prescription or OTC in Germany but are prescription-only in Spain

^cErgot alkaloids are only reimbursed in Germany

^dData for ergot alkaloid use were missing for 1187 (21.4%) patients in the total moderate to severe attacks subcohort

moderate to severe attacks were using triptans, despite both Spanish and German guidelines recommending their use in this population [17, 18]. This was observed even though > 80%of participants in this subcohort sought medical care, with NSAIDs being the most commonly used acute medication, reflecting an unmet need for people with migraine. Despite seeking care, symptoms may go unrecognized or be underestimated by HCPs, resulting in patients going undiagnosed and not receiving the recommended treatment. Although the proportions of participants using triptans in our study is higher than some other reports from European countries [19, 38], these other reports are based on the general migraine population and migraine severity is not reported in these studies. The Eurolight study also showed considerable variation in medication use according to the population sample, with higher usage reported for general practitioner-based samples and those from lay organizations compared with population-based samples [19]. Triptan use was estimated at < 6% for people with migraine in an Austrian study, but this was based on a general population sample and estimated migraine prevalence, and migraine severity was not reported [38].

A high proportion of participants using triptans reported poor treatment satisfaction, and many people reported that they had stopped using triptans. Treatment satisfaction (according to mTOQ-6) was generally poor regardless of migraine/headache severity, with poor satisfaction being reported by most triptan-using participants in both the total migraine cohort and the moderate to severe attacks subcohort. However, we cannot exclude the possibility that other factors related to migraine severity or the potential concomitant



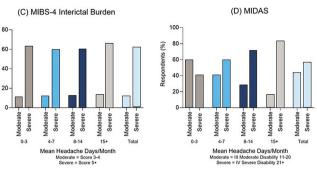


Fig. 3 Migraine characteristics and burden in the moderate to severe attacks subcohort^a. ^aModerate to severe headache attacks were defined as being of average pain severity \geq 5 points (based on the MIDAS question on pain severity), pain duration \geq 4 h, and at least moderate disability due to migraine (MIDAS score \geq 11). Higher scores for MSQ indicate a better QoL; higher scores for MIBS-4 indicate greater burden; higher scores for MIDAS indicate greater disability. *N* numbers for MSQ and MIDAS pain severity survey question (**A**, **B**) = 0–3

use of triptans and other migraine medications may have impacted the mTOQ-6 score, which is not medication specific. A global real-world study of people using triptans as their only acute migraine medication reported considerably higher rates of satisfaction than ours, even in those considered to be insufficient responders to triptans [59% of insufficient responders (i.e., freedom from headache pain within 2 h of treatment in $\leq 3/5$ migraine attacks) reported being satisfied or extremely satisfied, compared with 92% of triptan responders (those with pain freedom within 2 h in $\geq 4/5$ attacks)] [39]. However, that study did not use the mTOQ-6, and data for response based on headache freedom after using triptans were not collected in the OVERCOME (EU) study, so the results are not directly comparable.

In the current study, treatment satisfaction in the total migraine cohort was lower in participants currently taking triptans compared with those using acute medications excluding triptans. The most common reasons for discontinuing triptans [in the 5538 patients who had discontinued triptans (8686 patients reported ever having used triptans)] were that other medications were considered more effective, migraine/severe headaches improved, and

HDs/month (n = 2224), 4–7 HDs/month (n = 1653), 8–14 HDs/month (n = 859), 15 + HDs/month (n = 601), total (N = 5337). N numbers for MIBS-4 and MIDAS (**C**, **D**) = 0–3 HDs/month (n = 2329), 4–7 HDs/month (n = 1697), 8–14 HDs/month (n = 884), 15 + HDs/month (n = 637), total (N = 5547). HDs headache days, MIBS-4 Migraine Interictal Burden Scale-4, MIDAS Migraine Disability Assessment, MSQ Migraine-Specific Quality of Life Questionnaire, QoL quality of life

concerns about side effects (although the incidence of actual adverse events was not reported) and frequent use. Other studies have reported that around 30–40% of people with migraine do not respond to triptans, although some people who do not respond to one triptan find they do respond to an alternative triptan [40, 41]. However, as data on the number of different triptans participants had used, or the sequence of treatments, were not collected in the OVER-COME (EU) study, we were unable to further evaluate any influence of this.

The lack of triptan use may partly be due to the high use of primary care and pharmacist services compared with specialist headache or pain services, even by people with higher MIDAS grades, which again may result from a lack of referrals due to under-recognition of symptoms. Migraine burden in the moderate to severe attacks subcohort generally increased with increasing headache/migraine severity, and interictal burden was high regardless of the number of HDs/month, indicating high levels of burden in patients using existing medications. The unmet treatment need for people with severe headache/migraine was previously highlighted in the American Migraine Prevalence and Prevention study, which found no

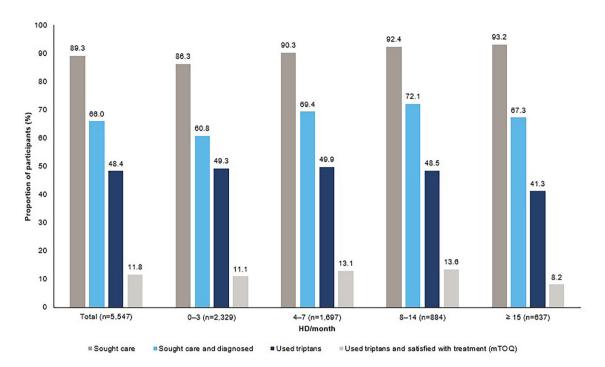


Fig. 4 Migraine characteristics and burden in the moderate to severe attacks subcohort^a. ^aModerate to severe headache attacks were defined as being of average pain severity \geq 5 points (based on the MIDAS question on pain severity), pain duration \geq 4 h, and at least moderate disability due to migraine (MIDAS score \geq 11). ^bTriptan treatment as per Spanish and German guidelines [17, 18].

significant improvement in headache-related disability in people who switched from one triptan to another, an opioid, or a barbiturate, compared with those with consistent triptan use [42]. That study also found that switching from a triptan to an NSAID was associated with an increase in disability, especially among those with \geq 10 HDs/month [42].

Strengths of the OVERCOME (EU) study include its provision of previously lacking realworld evidence from a population-based sample, allowing further insight into the unmet needs of people with migraine adding to previous data from other settings. The use of an internet-based survey also allowed for a large number of study participants, regardless of whether medical care was sought. These data are important as many people with migraine do not seek professional medical care so would not Triptans are available on prescription or OTC in Germany but are prescription-only in Spain. *HDs* headache days, *MIDAS* Migraine Disability Assessment, *mTOQ-6* six-item Migraine Treatment Optimization Questionnaire, *OTC* over-the-counter

be represented in registry or observational studies.

However, surveys such as this are subject to limitations. As the survey data are self-reported, they are susceptible to recall, misinterpretation, and prioritization biases. It was also not possible to confirm if participant responses regarding migraine symptoms, severity, or duration were based on treated or untreated migraine attacks, and data regarding the sequence of treatments used were not collected. Although internet access is widely available in Germany and Spain, as with other online surveys the cohort is more likely to include people with greater technical abilities who were willing to be involved in computer surveys. In order to limit bias, the participants were invited and agreed to take part in the study without knowledge of which health topic was being researched. The health topic (migraine) was then disclosed at the screening stage. However, we acknowledge the potential for differences between survey respondents and the broader migraine population in attitudes and tolerance to pain and other symptoms to influence findings regarding treatment satisfaction.

Further analyses of data from the OVER-COME (EU) study will provide important information on essential topics required to improve care for people with migraine, for example the need for targeted care, and in the case of severely affected patients, specialized care.

CONCLUSIONS

These data provide important insights into the real-world experience of migraine treatment and burden. Less than half of participants in this study were using triptans, despite their recommendation in Spanish and German guidelines [17, 18], and treatment satisfaction in those who did use triptans was generally poor.

High levels of participant burden were experienced despite the use of a range of acute medications, including the use of multiple concurrent medications. It is important to understand the current status of migraine healthcare delivery and reasons why people with migraine do not seek professional help, even though medical advice and treatment is accessible in their country. This understanding, together with new tailored acute treatment options expected to enter the market, may help to address the current unmet needs of people with migraine.

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Data Availability. The data sets generated during and/or analyzed during the current study are not publicly available due to the need for patient data protection.

Declarations

Conflict of Interest. Stefan Evers has received honoraria for lectures and consulting fees from Eli Lilly, Lundbeck, Novartis, Perfood, Rehaler, and Teva (in the last 3 years). Grazia Dell'Agnello, Diego Novick, Saygin Gonderten, and Tommaso Panni are employees and minor shareholders of Eli Lilly and Company. Julio Pascual has participated in Symposia and Advisory Boards of Allergan, Amgen-Novartis, Biohaven, Teva, and Lilly.

Ethical Approval. This study was performed in accordance with the Helsinki Declaration of 1964 and its later amendments. Ethical review board approval for the study was granted by the Marqués de Valdecilla University Hospital (Spain) and participants were required to provide informed consent prior to commencing the study by selecting "I agree to participate" on the online form. All survey respondents provided informed consent and all data were anonymized before analysis.

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