The relevance of migraine in the clinical spectrum of mitochondrial disorders

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Recent scientific evidence suggests a link between migraine and brain energy metabolism. In fact, migraine is frequently observed in mitochondrial disorders. We studied 46 patients affected by mitochondrial disorders, through a headache-focused semi-structured interview, to evaluate the prevalence of migraine among patients affected by mitochondrial disorders, the possible correlations between migraine and neuromuscular genotype or phenotype, comorbidities, lactate acid levels and brain magnetic resonance spectroscopy. We explored migraine-related disability, analgesic and prophylactic treatments. Diagnoses were achieved according to International Classification of Headache Disorders, 3rd edition. Lifetime prevalence of migraine was 61% (28/46), with high values in both sexes (68% in females, 52% in males) and higher than the values found in both the general population and previous literature. A maternal inheritance pattern was reported in 57% of cases. MIDAS and HIT6 scores revealed a mild migraine-related disability. The high prevalence of migraine across different neuromuscular phenotypes and genotypes suggests that migraine itself may be a common clinical manifestation of brain energy dysfunction. Our results provide new relevant indications in favour of migraine as the result of brain energy unbalance.

Migraine is the second leading cause of years lived with disability in the world (Global Burden of Disease Study)¹. Its one-year prevalence in women and men is 18% and 6%, respectively, with prevalence peaks between the ages of 25 and 55²-⁴. Migraine is associated with a considerable financial burden⁵-⁶ and its impact on functioning can be evaluated using validated scales as Headache Impact Test (HIT6)⁷, Migraine Disability Assessment (MIDAS)⁸ or the recent Migraine Physical Function Impact Diary⁹. The last decades witnessed a solid progress in understanding the mechanisms underpinning migraine, but much still remains to be explained¹⁰. A pioneering view of primary headaches indicates them as “adaptive behavioural responses”¹¹. Brain ³¹P-magnetic resonance spectroscopy (MRS) findings and mitochondrial energy metabolism markers in blood and muscle supported this theory¹²,¹³. In confirmation of this idea, Lisicki and collaborators suggested an activity-induced rupture of cerebral metabolic homeostasis in migraineurs, using ¹⁸FDG-PET and visual evoked potentials¹⁴. A recent review pointed to a role of energy metabolism in the pathogenesis of migraine, with mitochondria carrying out a central role¹⁵. Some authors speculate that migraine is a “conserved and adaptive response that occurs in genetically predisposed individuals with a mismatch between the brain energy reserve and workload”¹⁶. Under this assumption, we evaluated the prevalence of primary headaches and, in particular, of migraine (migraine without aura (MO), migraine with aura (MA), hemiplegic migraine (HM), chronic migraine (CM), probable migraine without aura (pMO) or probable migraine with aura (pMA)) among patients suffering from mitochondrial disorders, looking for clinical hints supporting the relationship between migraine and energy metabolism.

Results

Prevalence of primary headaches among patients with mitochondrial disorders. Forty-six individuals affected by mitochondrial disorders were included with a mean age of 55±14.6 years (range 17–83 years), 25 females (54.3%) and 21 males (45.7%). Body Mass Index (BMI) ranged from 15.2 to 30.8 kg/m² in females (mean value 22.1 ± 4.2 kg/cm²) and from 16.7 to 34.0 kg/m² in males (mean value 24.1 ± 4.3 kg/cm²).

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Other demographic features and cardiovascular risk factors are reported in Table 1. In this context, “Stroke/TIA” refers to an actual cerebrovascular events, i.e.: stroke in case of clear ischemic brain lesions, and TIA in case of transient acute neurological deficit in patients with multiple cerebrovascular risk factors.

The interview was conducted face-to-face in 57% of cases (26/46), and by telephone in 43% (20/46). A total of 21/46 patients (45.7%) reported at least one first-grade or second-grade relative who had suffered from recurrent headaches. In this group, a maternal inheritance pattern was reported in 12/21 cases (57%); paternal inheritance in two cases, while siblings but not parents were affected in the remaining seven cases (33%).

According to ICHD3 criteria, we considered first the one-year prevalence: 28.3% of our patients (13/46) reported no headache in the last year, 11 patients (23.9%) suffered from MO, one had CM and 4/46 (8.7%) had pMO. MA was reported in 2/46 (4.3%) and one had sporadic HM. Aggregate data show 41.3% of patients (19/46) suffering of headache with migrainous features in the last year. A group of 7/46 patients (15.2%) reported infrequent episodic tension-type headache (ETTH), 5/46 (10.9%) had frequent ETTH, and one had post-ictal headache and one unspecified headache (Fig. 1).

Considering the wide age range (17–83 years) and the high mean age of our patients cohort, data about the lifetime prevalence of migraine are more relevant: 18/46 (39.1%) of patients achieved the diagnosis of MO (in one case together with MA diagnosis), one reported CM, 2/46 (4.3%) had MA, in one case with hemiplegic features.

**Table 1.** Demographic features and cardio-cerebrovascular risk factors distribution in 46 patients, distinguished by sex. CHD coronary heart disease.

|                      | Females | Males |
|----------------------|---------|-------|
| Population           | 25 (54.3%) | 21 (45.7%) |
| Age                  | 53.9 ± 15.8 | 56.4 ± 13.4 |
| Weight (kg)          | 57.1 ± 11.7 | 73.9 ± 15.3 |
| Height (cm)          | 160.7 ± 5.6 | 174.6 ± 7.0 |
| BMI (Kg/cm²)         | 22.1 ± 4.2 | 24.1 ± 4.3 |
| Married              | 17/25 (68.0%) | 13/21 (61.9%) |
| Primary school       | 1/25 (4.0%) | 3/21 (14.3%) |
| Middle school        | 9/25 (36.0%) | 6/21 (28.6%) |
| High school          | 8/25 (32.0%) | 7/21 (33.3%) |
| Graduated            | 5/25 (20.0%) | 4/21 (19.0%) |
| Smokers              | 8/25 (32.0%) | 9/21 (42.9%) |
| Hypertension         | 7/25 (28.0%) | 10/21 (47.6%) |
| Dyslipidaemia         | 7/25 (28.0%) | 6/21 (28.6%) |
| Diabetes mellitus    | 3/25 (12.0%) | 4/21 (19.0%) |
| CHD                  | 2/25 (8.0%) | 2/21 (9.5%) |
| Stroke/TIA           | 3/25 (12.0%) | 2/21 (9.5%) |
| Alcohol              | 14/25 (56.0%) | 13/21 (61.9%) |
| Hormonal contraceptives | 13/25 (52.0%) | / |

**Figure 1.** Prevalence of primary headaches among patients with mitochondrial disorders, according to ICHD3 criteria. One-year prevalence (lower bar); lifetime prevalence of migrainous headache (including migraine without aura, migraine with aura, chronic migraine and probable migraine without aura) by sex (first and second bars) and in the whole study population (third bar).
A group of 7/46 patients (15.2%) reported pMO. Among the pMO patients, diagnostic criterion D ("during headache at least one of the following: 1. Nausea and/or vomiting; 2. Photophobia and phonophobia") was lacking in four cases. Criteria B ("headache attacks lasting 4–72 h, when untreated or unsuccessfully treated") and C ("headache has at least two of the following four characteristics: 1. Unilateral location, 2. Pulsating quality, 3. Moderate or severe pain intensity, 4. Aggravation by or causing avoidance of routine physical activity") were not satisfied in two and one cases, respectively. Overall, 60.9% (28/46) of patients with a mitochondrial disorder suffered from a migrainous headache in their lifetime. Migraine prevalence was 68.0% in females (17/25), and 52.4% (11/21) in male patients (Fig. 1). Only 10/46 patients (21.7% of cases) did not report any sort of recurrent headache in their life, despite a higher mean age (66.0 ± 13.9 years; range 42–83 years).

Clinical features of headache in mitochondrial patients. The median age at onset of headaches was 25.5 years, with 95% confidence interval from 20 to 40 years (Fig. 2A); this model includes "censored" patients who never developed headache. Migraine showed a median clinical onset at about 20 years, where TTH patients had an earlier median onset, at 19 years (Fig. 2B,C); these headache-type-specific models excluded "censored" patients, hence the earlier estimated median values. The difference between the median age at onset among the different headache subtypes was not statistically significant. Nine patients reported the past resolution of a highly-disabling migrainous headache towards a less impacting ETTH (5 cases) at the mean age of 44 years or a complete resolution of their MO (2 cases) or pMO (1 case), at a mean age of 51 years. One patient reverted from CM to MO, while another one stopped suffering from MO, while continuing with his MA attacks.

Aura prevalence reported in only 3 patients (10.7% of the migraineurs): 2 were typical auras, while 1 patient carrying MICU1 mutations experienced hemiplegic migraine.

Our 28 migraineurs had experienced, at least one, 64% (42/66) of the triggers listed in our semi-structured 66-item-questionnaire, including the most frequent triggers of migrainous headache, according to specific scientific literature. Each patient reported up to 15 different triggers of migraine. As expected, among the most reported factors there were stress (39%), work-related stress (36%) or family-related stress (29%). More frequently, 54% of migrainous patients could had attacks triggered by sleeping deprivation, 11% with sleeping more than usual or 18% in case of sleeping in the afternoon. 53% of female migraineurs pointed to female hormonal fluctuations (including premenstrual period, menstrual phase, ovulatory period or oral contraceptives)

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**Figure 2.** (A): Cumulative incidence plot of headache with increasing age. Crosses indicate censored patients. (B): Cumulative incidence plot of migraine, with increasing age. (C): Cumulative incidence plot of tension-type headache, with increasing age. (D): Cumulative incidence plot of headache with increasing age, according to different mitochondrial genotypes. (E): Cumulative incidence plot of headache with increasing age, according to different neuromuscular phenotypes. Dashed lines indicate the 95% confidence interval. Each cross indicates censored patients. Abbreviations: ADOA = autosomal dominant optic atrophy; CPEO = chronic progressive external ophthalmoplegia; MELAS = mitochondrial encephalopathy, lactic acidosis and stroke-like episodes; MM = other mitochondrial myopathy.
of the head (21%–6/28). Moderate physical activity exacerbated headache in 79% (22/28) of cases. Prodromal symptoms were reported by only 3 MO patients (11% of migraineurs), and included tiredness, neck stiffness, fatigue, confusion and blurred vision. Accompanying symptoms consisted of nausea (79%–22/28), vomiting (50%–14/28), photophobia (79%–22/28), phonophobia (54%–15/28) and osmophobia (18%–5/28). Allodynia was reported by 18% (5/28) of migrainous cohort too, with a tendency towards a correlation with a higher number of migraine attacks per month: a mean of 9.2 ± 13.1 attacks per month among allodynic patients, versus a mean of 3.2 ± 3.8 attacks per month among non-allodynic migraineurs (\(p = 0.093\)) (Supplementary Information 1).

Migrainous attacks had a mean duration of 24.1 ± 25.1 h. At the time of the interview, patients reported a mean frequency of 4.1 ± 6.1 migraine attacks per month (range 0.3–30), with a median value of 2.5 attacks per month. 12 patients (43%) reported more than three migraine attacks per month that were often moderately or severely disabling (75% of these sub-sample). Two other patients reported a mean of three highly disabling

| Trigger                                    | Prevalence (%) | Prevalence (n) | N ref |
|--------------------------------------------|----------------|----------------|-------|
| Lack of sleep                              | 53.6           | 15             | 28    |
| Anxiety                                    | 42.9           | 12             | 28    |
| Premenstrual period                        | 41.1           | 7              | 17    |
| Unspecified stress                         | 39.3           | 11             | 28    |
| Anger                                      | 35.7           | 10             | 28    |
| Work-related stress                        | 35.7           | 10             | 28    |
| Warm days                                  | 28.6           | 8              | 28    |
| Family-related stress                      | 28.6           | 8              | 28    |
| Menstrual period                           | 23.5           | 4              | 17    |
| Sun exposure                               | 21.4           | 6              | 28    |
| Strenuous physical exercise                | 21.4           | 6              | 28    |
| Intense lights                             | 21.4           | 6              | 28    |
| Physical fatigue                           | 17.9           | 5              | 28    |
| Daytime sleep                              | 17.9           | 5              | 28    |
| Windy days                                 | 17.9           | 5              | 28    |
| Wet days                                   | 17.9           | 5              | 28    |
| Fasting/skipping meals                     | 14.3           | 4              | 28    |
| Cold days                                  | 14.3           | 4              | 28    |
| Other                                      | 14.3           | 4              | 28    |
| Ovulatory period                           | 11.7           | 2              | 17    |
| Sleeping more than usual                   | 10.7           | 3              | 28    |
| Loud noises                                | 10.7           | 3              | 28    |
| Fever                                      | 10.7           | 3              | 28    |
| Cigarette smoke                            | 10.7           | 3              | 28    |
| Relaxing after stressful period            | 10.7           | 3              | 28    |
| Binge eating                               | 10.7           | 3              | 28    |
| Other climate changes                      | 10.7           | 3              | 28    |
| Scents                                     | 10.7           | 3              | 28    |
| Other emotional reaction                   | 7.1            | 2              | 28    |
| Mood changes                               | 7.1            | 2              | 28    |
| Aeroplane travels                          | 7.1            | 2              | 28    |
| Other food                                 | 7.1            | 2              | 28    |
| Oral contraceptives                        | 5.9            | 1              | 17    |
| Red wine                                   | 3.6            | 1              | 28    |
| Dehydration                                | 3.6            | 1              | 28    |
| Hypoglycaemia                              | 3.6            | 1              | 28    |
| High-altitude                              | 3.6            | 1              | 28    |
| Long travels                               | 3.6            | 1              | 28    |
| White wine                                 | 3.6            | 1              | 28    |
| Other alcoholics                           | 3.6            | 1              | 28    |
| Fruits                                     | 3.6            | 1              | 28    |
| Vegetables                                 | 3.6            | 1              | 28    |

Table 2. Trigger factors for migraine attacks. Complete list of the triggers reported by patients suffering from a migrainous headache; triggers have been selected among the items listed in our semi-structured questionnaire. N. ref. = number of patients representing the total for each item.
migraine attacks per month, whose duration may reach the 48 h. 50% of our active migraineurs did benefit from a prophylactic therapy, but none was regularly assuming it. Considering the whole sample, only 18% (5/28) of our migrainous patients had tried a preventive therapy at least once in their life, including flunarizine (3 cases), followed by topiramate, pizotifen, propranolol, acupuncture and nutraceutical (1 case each).

Mitochondrial phenotype and genotype and migraine. Clinical phenotype (Table 3) was consistent with CPEO in 33 patients (5 CPEO, 25 CPEO plus and 3 KSS), MELAS in 6, ADOA in 1, other MM in 6 patients (a young patient with MICU1 gene mutation and other unspecified MM in 5 cases) (Table 4). 19/33 CPEO patients (58%) suffered from a migrainous headache: 12 MO (36%), 1 CM and 6 pMO. All 10 patients who denied the experience of recurrent headaches in their medical history belonged to CPEO subgroup. 67% of MELAS patients were migraineurs (2/6 MO, 1/6 MA, 1/6 MO plus MA), while 83% of patients with other MM reported a migrainous headache (3/6 MO, 1 pMO, 1 HM). The only ADOA patient had an ETTH (Table 5, Fig. 3a). The absence of any primary headaches results associated with CPEO phenotype with statistical significance (p=0.036). The age of onset of headache was not significantly different across various phenotypic groups (Fig. 2d).

A total of 32/46 patients (70%) had a defined genotype. 11 patients harboured nDNA mutations and represented the largest group: POLG mutations were identified in 7, and OPA1, DGUOK, MGME1 and MICU1 gene mutation in one patient each. Six patients had multiple mtDNA deletions but no nuclear gene mutation identified yet. Thirteen patients showed a single mtDNA deletion, while eight had a mtDNA point mutation: A3243G in 6, a tRNALeu (mt.C3250T) and tRNAMet (mt.A4415G) in in one patient each. 60.5% (23/38) of patients with a defined disease-causing mutation or with multiple mtDNA deletions suffered from a migrainous headache.
Table 5. Distribution of ICHD3 diagnoses with different mitochondrial phenotypes. No no headache, MO migraine without aura, MA migraine with aura, HM hemiplegic migraine, CM chronic migraine, pMO probable migraine without aura, iETTH infrequent tension-type headache, fETTH frequent tension-type headache. ADOA autosomal dominant optic atrophy, CPEO chronic progressive external ophthalmoplegia, KSS Kearns-Sayre syndrome, MELAS mitochondrial encephalopathy, lactic acidosis and stroke-like episodes, MICU1 mitochondrial calcium uptake 1, MM other mitochondrial myopathy.

| Phenotype | No | MO | MA/HM | CM | pMO | iETTH | fETTH | Others | Tot |
|-----------|----|----|-------|----|-----|-------|-------|--------|-----|
| ADOA      | 0  | 0  | 0     | 0  | 0   | 0     | 0     | 0      | 1   |
| CPEO      | 1  | 0  | 0     | 0  | 2   | 2     | 0     | 0      | 5   |
| CPEO plus | 9  | 9  | 0     | 1  | 4   | 1     | 0     | 1      | 25  |
| KSS       | 0  | 3  | 0     | 0  | 0   | 0     | 0     | 0      | 3   |
| MELAS     | 0  | 3  | 1     | 0  | 0   | 0     | 1     | 1      | 6   |
| MICU1     | 0  | 0  | 1     | 0  | 0   | 0     | 0     | 0      | 1   |
| MM        | 0  | 3  | 0     | 0  | 1   | 1     | 0     | 0      | 5   |
| Tot       | 10 | 18 | 2     | 1  | 7   | 4     | 2     | 46     |      |

Figure 3. Histograms showing the distribution of migraine, ETTH, other types of headache or absence of headache in all phenotypic (a) and genotypic (b) groups of mitochondrial disorders. Legend: light blue = includes migraine without aura, migraine with aura, hemiplegic migraine, chronic migraine, probable migraine without or with aura; dark blue = episodic tension-type headache, black = other types of headache; grey = no headache sufferers. Abbreviations: ADOA = autosomal dominant optic atrophy; CPEO = chronic progressive external ophthalmoplegia; KSS = Kearns-Sayre syndrome; MELAS = mitochondrial encephalopathy, lactic acidosis and stroke-like episodes; MICU1 = mitochondrial calcium uptake 1; MM = other mitochondrial myopathy. DGUOK = deoxyguanosine kinase; MELAS = mitochondrial encephalopathy, lactic acidosis and stroke-like episodes; MGME1 = mitochondrial genome maintenance exonuclease 1; MICU1 = mitochondrial calcium uptake 1; MM = other mitochondrial myopathy; multiple deletions = multiple mtDNA deletions; OPA1 = OPA1 mitochondrial dynamin like GTPase; POLG = DNA polymerase gamma; single deletion = single mtDNA deletion.
test was not significant. A total of 43% of patients with a normal MRS study received a diagnosis of migraine (6/14); Fisher's absence of headaches), in 3 cases (3/17–18%) revealing an increase in lactate levels and a migrainous headache part of a different previous research protocol about mitochondrial disorders, independently from the presence/

Among patients with nDNA mutations, migrainous headache (4 MO, 4 pMO and one HM) was present in 52.9% of patients (9/17). Sixty-two percent (8/13) of patients with single mtDNA deletion were migraineurs (6 MO and 2 pMO). The mtDNA point mutations were associated with migraine in 75% of cases (6/8): 5 MO and one MA (Table 6, Fig. 3b). The distribution of primary headaches with the different genotypes did not differ significantly among subgroups. Age at headache onset and lack of primary headaches was not statistically different according to genotypes (Fig. 2e).

Among 19 patients with active migraine, mean HIT6 score was 55.1 ± 6.8 pts (range 46–65). Eleven patients reported mild (HIT6 score between 50 and 55 points) or no disability (HIT6 score < 50 points) related to their migraine; eight (42%) experienced a moderate (HIT6 score from 56 to 59 points) or severe (HIT6 score > 59 points) migraine-related disability. CPEO plus patients showed a tendency towards a higher degree of disability and KSS patients reported a mild degree of disability, but no statistically significant differences in HIT6 scores could be found across different phenotypes or genotypes. POLG patients showed a tendency towards a higher migraine-related disability (Supplementary Information 2).

The 19 patients with active migraine reported a mean MIDAS score of 11.2 ± 19.6 points (range 0–80 points) with a median of 1 point. A mild (MIDAS score from 6 to 10 points) or negligible migraine-related disability (MIDAS score < 6 points) has been reported by 74% (14/19) of the patients, with only two cases of moderate disability (MIDAS score from 11 to 20 points) and three patients with severe disability related to migraine (MIDAS score > 20 points). The last five patients had a moderate-severe disability score at HIT6, as well.

Table 6. Distribution of ICHD3 diagnoses with different mitochondrial genotypes. No no headache, MO migraine without aura, MA migraine with aura, HM hemiplegic migraine, CM chronic migraine, pMO probable migraine without aura, pMA probable migraine with aura, iETTH infrequent tension-type headache, fETTH frequent tension-type headache, DGUOK deoxyguanosine kinase, MGME1 mitochondrial genome maintenance exonuclease 1, MICU1 mitochondrial calcium uptake 1, OPA1 OPA1 mitochondrial dynamin like GTPase, POLG DNA polymerase gamma, mtM other point mtDNA mutations, MELAS mitochondrial encephalopathy, lactic acidosis and stroke-like episodes.

| Genotype | No | MO | MA/HM | CM | pMO | pMA | iETTH | fETTH | Others | Tot |
|----------|----|----|-------|----|-----|-----|-------|-------|--------|-----|
| nDNA     |    |    |       |    |     |     |       |       |        |     |
| DGUOK    | 1  | 0  | 0     | 0  | 0   | 0   | 0     | 0     | 0      | 1   |
| MGME1    | 1  | 0  | 0     | 0  | 0   | 0   | 0     | 0     | 0      | 1   |
| MICU1    | 0  | 0  | 1     | 0  | 0   | 0   | 0     | 0     | 0      | 1   |
| OPA1     | 0  | 0  | 0     | 0  | 0   | 0   | 0     | 1     | 0      | 1   |
| POLG     | 2  | 2  | 0     | 2  | 0   | 0   | 0     | 0     | 1      | 7   |
| mtDNA point |    |    |       |    |     |     |       |       |        |     |
| mtM      | 0  | 2  | 0     | 0  | 0   | 0   | 0     | 0     | 0      | 2   |
| MELAS    | 0  | 3  | 1     | 0  | 0   | 0   | 0     | 1     | 1      | 6   |
| mtDNA deletion |    |    |       |    |     |     |       |       |        |     |
| Single deletion | 3  | 6  | 0     | 0  | 2   | 0   | 2     | 0     | 0      | 13  |
| Total    | 8  | 15 | 2     | 0  | 6   | 3   | 2     | 2     | 38     |     |

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Treatment strategies: the great majority of patients (79%–22/28) took non-steroidal anti-inflammatory drugs (NSAIDs), with paracetamol and ibuprofen as the leading drugs (29% of cases, each), combination-analgesic or metamizole in three cases, while three patients (11%) did not assume any acute treatment for their migraine attacks. None of our 28 mitochondrial patients treated their migraine with triptans or ergot alkaloids, even if two patients (7% of migraineurs) reported previous Medication Overuse Headache (one paracetamol-overuse headache and a combination-analgesic overuse headache).

A logistic regression analysis of risk of the onset of migraine was performed including variables such as sex, age, BMI, cardiovascular risk factors, vision loss, parkinsonism (defined as the presence of clinical signs suggesting a dysfunction of the extrapyramidal system), gastrointestinal dysmotility, respiratory involvement, family history, coffee use, bruxism, sleep disturbances (including snoring, obstructive sleep apnoea or insomnia) and muscular weakness. This multivariate analysis showed that female sex (logistic regression coefficient 0.44, 95% CI 0.076–0.80, p = 0.005) and sleep disturbances (coefficient 0.431032, 95% CI 0.068–0.80, p = 0.029) were significantly associated with a lifetime diagnosis of migraine (Fig. 4). BMI showed a trend to association with migraine onset (coefficient 0.050, 95% CI 0.0007–0.099, p = 0.058).

Among possible biochemical markers, blood lactate was available only in 67% (31/46) of patients. The mean value was 2.5 ± 3.9 mmol/L (range 0.5–20.0 mmol/L; normal level < 2.2 mmol/L), with higher levels among the 17/31 migrainous patients (mean 3.5 ± 5.1 mmol/L; range 0.8–20.0 mmol/L) compared to the 14/31 non-migrainous patients (mean 1.2 ± 0.7 mmol/L; range 0.5–2.5 mmol/L). Differences in blood lactate levels were statistically significant (p = 0.012 Supplementary Information 3). Seventeen patients (37%) had a brain 1H-MRS (as part of a different previous research protocol about mitochondrial disorders, independently from the presence/absence of headaches), in 3 cases (3/17–18%) revealing an increase in lactate levels and a migrainous headache (100%–3/3). A total of 43% of patients with a normal MRS study received a diagnosis of migraine (6/14); Fisher’s test was not significant.
Discussion

Our 46 patients cross-sectional study represents the fourth largest study in scientific literature, regarding primary headaches in adult patients with mitochondrial disorders. We found a one-year prevalence of migrainous headache of 41%, significantly greater than the prevalence reported in general population in the European countries\(^20,21\), and higher than the 25.9% prevalence reported in a recent large Italian population study\(^22\). A total of 61% of mitochondrial disease patients suffered or had suffered of a migrainous headache. Since our sample has a high mean age and migraine prevalence is known to decrease with increasing age\(^20\), the prevalence in our patients’ cohort is more than threefold higher than expected in the general population\(^20\) in both sexes, reaching 68% in female. Our estimates are higher than those reported in a recent large Dutch study (55%, mainly focused on mt.3243A > G patients\(^23\)) and in a previous study with more heterogenous subtypes of mitochondrial patients (35.5\%)\(^24\). Given the well-known high prevalence of migraine among mt.3243A > G patients\(^25,26\), the limited number of MELAS patients in our cohort minimizes selection bias. CPEO and CPEO plus are the main phenotypes in our study population, with a subgroup prevalence of migraine of 53%, which is notably higher than the 21–40% prevalence previously reported among CPEO patients\(^27,28\). We found higher migraine prevalence also in POLG patients (57 vs. 23%)\(^30\), whereas the prevalence was similar to that of literature in MELAS patients (67\%)\(^26\). Age at onset, evolution over time, prodromal symptoms, trigger, features and duration of headache are in line with those of the general population with migraine, confirming that the clinical entity is likely the same. However, a few differences from the general population emerged. First, phonophobia shows 54% prevalence and osmophobia involves only 18% of migraineurs, which are frequencies lower than those literature 17,31. This could be partially due to the frequent hearing loss (reported up to 43% of our cohort) or to the decrease in olfactory perception induced by mitochondrial pathology (nearly 10% of our sample). In fact, in our 7 pMO patients the diagnostic criterion D ("During headache at least one of the following: (1) nausea and/or vomiting; (2) photophobia and phonophobia")\(^16\) was absent more often than other definitory features. Furthermore, the 11% prevalence of prodromal symptoms is significantly lower than the 72% reported in literature\(^9\), with a probable bias due to memory recall. These data suggest that we did not document a mere migraine-like headache: migraine, as coded in ICHD3, appears indeed a core clinical manifestation of the clinical spectrum of mitochondrial disorders.

Considering the familiar occurrence of migraine in general population, a maternal inheritance pattern is by far the most frequent\(^32,33\), providing another hint in favour of a possible role of mitochondria in migraine pathogenesis. Interestingly, isolated stress (50\%), lack of sleep (54\%), oestrogen fluctuations (53\%), fatigue (24\%), vigorous physical exercise (18\%), fasting or skipping meals (15\%) are some of the consistently reported migraine triggers which can be connected to failure of energy metabolism, mitochondrial dysfunction and oxidative stress. Accordingly, and in contrast with previous literature data\(^24\), we observed a statistically significant association between migraine and increased blood lactate levels, likewise our 3 MO patients all had increased lactate levels detected by \(^1H-MRS\).

We failed to find any significant difference in age at onset and migraine prevalence, nor was migraine prevalence significantly different across various mitochondrial genotypes, with a prevalence ranging from 53% (nDNA mutation) to 75% (mtDNA point mutation). This lack of correlation between migraine and neuromuscular phenotype or genotype is in line with previous literature\(^24\) and suggests that migraine is likely associated with dysfunction of brain energy metabolism, rather than with a specific mutation or clinical pattern.

Figure 4. Forest plot showing the odds ratios, confidence intervals 95% and direction of association between migraine and demographic features, cardiovascular risk factors and other clinical conditions. Multivariate analysis showed that female sex, age, and sleep disturbances were significantly associated with the diagnosis of migraine.
Migraine showed a statistically significant association with female sex and sleep disturbances (and a tendency towards a correlation with increasing BMI), in accordance with previous studies on migraine\textsuperscript{24–27}, strengthening the theorized central role of energy unbalance as the fulcrum of migraine pathogenesis.

Concerning the results of HIT6 and MIDAS scales, the mean values reported by the only previous work on migraine in mitochondrial disorders\textsuperscript{28} point to a mild level of disability. In our study, 42% of our active-migraine individuals reported an at least moderate migraine-related impairment on HIT6 scale, compared to the 62% prevalence found by Tiehuis and collaborators in a younger Dutch population\textsuperscript{29}. 26% of our patients had a disabling migraine according to MIDAS, versus 34% in the scientific literature\textsuperscript{23}. No association between HIT6/MIDAS scores and mitochondrial phenotypes or genotypes has been found. However, “CPEO plus” and POLG patients showed a tendency towards a higher degree of disability.

Data on migraine treatment were surprising. None of our patients was being administered a preventive therapy and only 18% of patients with a lifetime diagnosis of migraineous headache had tried prophylaxis before our examination, nor acute treatment was previously considered: no one had tried triptans in their life, whereas common analgesics were commonly used in a few cases. Treatment of migraine was far less than optimal in patients with mitochondrial disorders, often without exploiting the most appropriated available pharmaceutical tools. This under-treatment of migraine is in line with the few available data in scientific literature\textsuperscript{23}: it could be partially due to the (wrong) underevaluation, both by patients and physicians, of the role played by headache in the syndromic output of the patients, when compared to the systemic and sometimes life-threatening dysfunctions associated with myopathy or mitochondrial pathology.

Our study sheds light on the role of migraine in mitochondrial diseases, with some limitations. First, the cross-sectional design of our study may introduce a recall bias. Second, the still limited size of our cohort of patients makes it difficult to reach statistically sound associations between migraine and neuromuscular phenotypes or genotypes. Third, we lack a matched control group for a direct comparison. Fourth, we did not perform any measure of cognitive and/or memory performance in order to quantify the consistency and completeness of reported data, and finally, psychiatric comorbidities have not been considered in our study.

In this observational study, our preliminary clinical data converge to define migraine and its subtypes as a typical clinical manifestation of mitochondrial disorders. The lifetime prevalence of migraine was 61%, which is markedly higher than the values found in both the general population and the previous literature concerning mitochondrial disorders. In agreement with a previous article\textsuperscript{34}, the prevalence of migraine was high across different neuromuscular phenotypes and genotypes suggesting that migraine itself may be a common clinical manifestation of brain energy dysfunction. Our results provide new relevant indications in favour of migraine as the result of brain energy unbalance.

Materials and methods

Ethical approval. This study was approved by the “Ethical Committee for Clinical Studies of the Padova district” and have therefore been performed in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki and its later amendments.

Patients. We performed an observational study on a series of patients affected by mitochondrial disorders followed at the Neuromuscular Center of the Neurological Clinic of the University of Padova. We included patients with a genetically confirmed diagnosis of mitochondrial disorder and patients in whom the combination of clinical (muscle w/o multisystem involvement), biochemical (OXPHOS dysfunction), neuroimaging (MR abnormalities) and histopathological (ragged red fibers, COX negative/SDH positive fibers, subsarcolemmal rims at SDH) data could strongly suggest the diagnosis of mitochondrial encephalomyopathy, even in absence of a known pathogenic mutation. Patients were clinically classified in “CPEO” (progressive bilateral ptosis and ophthalmoplegias), “KSS” (Kearns-Sayre syndrome, early onset CPEO plus cardiac conduction block and pigmentary retinopathy w/o multisystem involvement), “CPEO plus” (CPEO associated to multiple features of neuromuscular and multisystem involvement), autosomal dominant optic atrophy (“ADOA”), mitochondrial encephalopathy lactic acidosis and stroke-like episodes (“MELAS”) and other mitochondrial myopathy (“MM”). Patients were genotypically classified in individuals carrying nuclear DNA (nDNA) mutations, mitochondrial DNA (mtDNA) mutations and accumulation of multiple mtDNA deletions. All patients underwent neurological examination. Brain MRs and blood lactate levels were collected in a subset of patients.

Headache evaluation. A semi-structured headache-focused interview was performed by a neurologist with specific training in the headache field; whenever possible it was carried out face-to-face, otherwise a telephone interview was performed. We asked for relatives suffering from headache, the time of onset of headache in the personal medical history and its evolution over time, the prodromal and associated symptoms, possible triggers (through a detailed 66-items semi-structured retrospective questionnaire Supplementary Information 4) and therapy. MIDAS and HIT6 scales were administered to each patient to measure migraine-related disability. Diagnoses of primary headache were codified according to International Criteria of Headache Disorders 3rd edition (ICHD3) criteria\textsuperscript{36}. Possible correlations between migraine and neuromuscular phenotype or genotype were investigated. All patients had given written informed consent to the scientific use of their clinical data, which have been treated in a completely anonymous way. We excluded patients aged <16 years or with severe cognitive decline and no reliable caregiver that made impossible the administration of the semi-structured interview, patients with intracranial space-occupying lesion or with a history of recent (less < 1 year) stroke.

Statistical analyses. Results were displayed as mean and standard deviation (SD) or median and range when appropriate. Age at headache onset was reported using the Kaplan–Meier method, and compared between
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Author contributions
Conceptualization, E.P., L.B.; methodology, A.T., F.M.; data acquisition A.T., M.L.V., L.C., G.S., V.C.; formal analysis, L.B., A.T.; writing—original draft preparation, A.T.; writing—review and editing, E.P., M.Z., V.C.; supervision, E.P. All authors read and approved the final manuscript.

Competing interests
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Additional information
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