

# Association Between Glucose Level And Prevalence of Headache Among Saudi Population: A Cross-Sectional Study

Eman AbdulAziz Balbaid<sup>1</sup>, Jamal Zaid Alshaikh<sup>2</sup>, Amin Abdulrahman Hafiz<sup>3</sup>,Hoda Jehad Abousada<sup>4\*</sup>, Yara Abdulrahman Fatani<sup>5</sup>, Mohammed Khaled Al Sedran<sup>5</sup>, Renad Saeed Nasser<sup>5</sup>, Nasser Abdullah Aljahmi<sup>5</sup>, Muath Abdullah Alqathanin<sup>6</sup>, Abdulrahman Mohammed Al Hadi<sup>6</sup>, Taif Nasser Alharrani<sup>7</sup>, Afrah Barjas Alanazi<sup>8</sup>, Aishah shafi Alenazi<sup>8</sup>, Ebtesam jaber alanzi<sup>8</sup> and Mashael Jaber Alanazi<sup>8</sup>

<sup>1</sup>Department of Family Medicine, Jeddah University Medical Center, Jeddah University, Jeddah, KSA, Saudi Arabia

<sup>2</sup> Department of Family Medicine, Old Airport PHC, MOH, Jeddah, KSA, Saudi Arabia

<sup>3</sup>Department of Clinical Nutrition, Umm Al-Qura University, Kingdom of Saudi Arabia

<sup>4</sup> Department of Obstetrics & Gynecology, Master SA, KSA, Saudi Arabia

<sup>5</sup> Medical Service Doctor, MBBS, KSA

<sup>6</sup> Medical Intern, MBBS, KSA

<sup>7</sup> Medical Student, KSA

<sup>8</sup> Nursing, Riyadh ,KSA

### RESEARCH

Please cite this paper as: Balbaid EA, Alshaikh JZ, Hafiz AA, Abousada HJ, Fatani YA, Al Sedran MK, Nasser RS, Aljahmi NA, Alqathanin MA, Al Hadi AM, Alharrani TN, Alanazi AB, Alenazi AS, Alanzi EJ, Alanazi MJ. Association Between Glucose Level And Prevalence of Headache Among Saudi Population: A Cross-Sectional Study. AMJ 2023;16(12):1089-1111.

https://doi.org/10.21767/AMJ.2023.4007

Corresponding Author: Hoda Jehad Abousada Department of Obstetrics & Gynecology, Master SA, KSA, Saudi Arabia dr.huda1992@outlook.com

## ABSTRACT

#### Objective

To investigate the association between glucose levels and the prevalence of headaches among the adult Saudi population.

### Methods

This research will employ a cross-sectional study design, which involves collecting data from a sample of the Saudi population at a single point in time. This design allows for the investigation of the association between glucose levels and headache prevalence in a cost-effective and timeefficient manner.

#### Results

The study included 574 participants. The most frequent weight among them was 51-65 kg (n= 196, 34.1 Per Cent), followed by 66-75 kg (n= 130, 22.6 Per Cent). The most frequent height among study participants was 151-160 cm (n= 229, 39.9 Per Cent), followed by 161-170 cm (n= 195, 34 Per Cent). The most frequent age among study participants was less than 25 years (n= 203, 35.4 Per Cent), followed by 25-30 years (n= 143, 24.9 Per Cent). The most frequent nationality among study participants was Saudi (n= 547, 95.3 Per Cent), followed by non-Saudi (n= 27, 4.7 Per Cent). The most frequent gender among study participants was female (n= 351, 61.1 Per Cent), followed by male (n= 223, 38.9 Per Cent). The perceived blood sugar level intake varied among study participants, with most having normal blood sugar levels. Participants were asked if they smoked. Most were not smoking (n=482, 84 Per Cent), and nonsmoking were (n=92, 26 Per Cent). Participants were asked about the nature of the headache. The most frequent were they don't have (n= 345, 60.1 Per Cent), followed by sharp (n= 116, 20.2 Per Cent), and the least was continuous (n=113, 19.7 Per Cent). Participants were asked about a kind of headache. The most frequent were they don't have (n=

367, 63.9 Per Cent), followed by stress headaches (n= 116, 20.2 Per Cent), and the lowest cluster (n=15, 2.6 Per Cent).

### Conclusion

The results of the study showed that most of the participants were Saudis. Most of them are women. In addition, the majority of people have normal fasting blood sugar levels. Their physical activity is moderate and the majority are non-smokers. In addition, most of the study participants had good and effective social communication.

#### **Key Words**

Headache

# Introduction

Migraine is a chronic neurological condition characterized by recurrent headache episodes that last 4 to 72 hours and are often accompanied by other symptoms such as nausea, vomiting, sensitivity to light and sound, and phonophobia <sup>1,2</sup>. Commonly occurring between the ages of 25 and 50, migraine is regarded as a complicated neurovascular brain condition<sup>3</sup>. Migraine is the third most prevalent medical condition and the second most debilitating neurological disorder in the world, with a lifetime frequency of 15-20 Per Cent. Migraine has a higher prevalence in females (33 Per Cent lifetime vs. 13 Per Cent annually) than males (6 Per Cent yearly vs. 13 Per Cent lifetime) <sup>4-6</sup>. Migraine is three times more common in women than in males, according to studies done in the United States and Europe<sup>7</sup>. Loss of productivity and increased use of healthcare services contribute significantly to the monetary burden that migraine places on sufferers and their communities<sup>8</sup>. The social and economic burden of migraines is high<sup>9</sup>, with yearly costs of US \$20 billion in the United States and €111 billion in the European Union. An estimated 95 Per Cent of migraine headaches can be placed into one of two major clinical subclasses by the International Headache Society (IHS): migraine without aura (MO), which affects 70-80 Per Cent of migraineurs, and Migraine with Aura (MA), in which patients experience auditory, visual, and sensory hallucinations <sup>10,11</sup>. Migraine with aura, hemiplegic, and retinal migraine are the three primary subtypes of MA. According to the frequency of headache attacks, migraine may be classified as either episodic (EM) or chronic (CM)<sup>12</sup>. Environmental variables have also been observed to have a substantial influence in initiating and maintaining migraine in certain patients, adding another layer of complexity to

the disease. Migraines may be brought on by things like not eating or eating too late, not drinking enough water, hormonal shifts, exposure to strong light or noise, using oral contraceptives, or taking hormone replacement therapy <sup>13</sup>. The evolution of the illness and the effectiveness of treatment and prevention strategies for migraine may be significantly influenced by the presence of many comorbidities, including those of a neurological, cardiovascular, psychiatric, and endocrine nature. Migraine sufferers, particularly those with MA, have a higher risk of cardiovascular complications such stroke <sup>14</sup>, angina <sup>15</sup>, and myocardial infarction <sup>16</sup> than the general population. Multiple studies have sought to identify shared metabolic abnormalities in migraine patients because of their increased risk of multiple cardiovascular illnesses. Glucose metabolic characteristics have been the subject of much research. Hypoglycemia has been linked to migraines for than a century. It has long been recognized that low blood sugar levels may bring on or exacerbate migraine symptoms <sup>17-20</sup>. Preliminary experimental research <sup>21</sup> shows that fasting or the injection of glucose or insulin might cause metabolic

changes that can trigger migraine symptoms. Hypoglycemia may trigger migraine episodes in CM patients after extended fasting because insulin is a key regulator of brain glucose metabolism<sup>22</sup>. Interictal deficits in glucose tolerance and insulin resistance have been seen in several investigations with migraine<sup>23,24</sup>. The hormone insulin, in particular the insulin-sensitive glucose transporter GLUT4, is the primary regulator of glucose homoeostasis by facilitating glucose uptake from the blood into cells, especially fat and muscle.

Patients with CM have been found to have greater levels of insulin resistance <sup>25</sup>, despite the fact that there is conflicting evidence about the prevalence of metabolic problems in migraine. In a study of young, normal-weight, migraine-free individuals, Insulin Resistance (IR) was shown to be significantly higher compared to non-migraineurs [26]. On the other hand, IR is a major pathogenetic element in the development of T2D. Different research have shown contrasting findings on the prevalence of type 2 diabetes in sufferers<sup>26-29</sup>. When other T2D migraine related pathogenetic abnormalities are present, IR linked with CM may enhance the risk of T2D <sup>30-32</sup>. Therefore, if -cell insulin secretion is impaired in CM patients with high IR, these individuals may be at an increased risk of developing T2D. IR is linked to many different diseases and conditions<sup>34</sup>, including as dyslipidaemia, obesity, diabetes, high blood



pressure, stroke, and coronary artery disease. Comorbidity between migraine and glucose-related characteristics is not uncommon because of the high prevalence of both illnesses and the fact that they commonly share aetiologies. This narrative review will thus concentrate on observational epidemiology and genetic research to investigate the association between migraine and glucose-related features. The research problem addressed in this study is of significant concern due to its potential impact on public health in Saudi Arabia and beyond. The primary research problem is the prevalence of headaches among the Saudi population, which has been rising in recent years. Headaches can substantially diminish an individual's quality of life and productivity, and understanding their causes and risk factors is crucial for effective prevention and management. This research aims to investigate whether there is a significant association between glucose levels and the occurrence of headaches, seeking to identify a potential relationship that may guide healthcare interventions.

Another aspect of the research problem is the increasing burden of diabetes and metabolic disorders in Saudi Arabia. The nation has witnessed a surge in the incidence of diabetes, which is closely linked to glucose dysregulation. Understanding how glucose levels might contribute to headaches can provide valuable insights into the broader health implications of this growing epidemic. As such, the research problem addresses the urgent need to delve into the interplay between glucose metabolism and headaches, a multifaceted issue that has not been extensively studied in the Saudi context.

Furthermore, this research problem has implications for global health. Diabetes and headache disorders are not confined to Saudi Arabia; they affect populations worldwide. If a substantial association between glucose levels and headaches is discovered in this study, it could have relevance and applications beyond Saudi Arabia's borders. This research seeks to contribute to the growing body of knowledge on the link between metabolic health and headache prevalence, which may have far-reaching implications for healthcare practices and strategies internationally.

# Methods

#### Study design

This research will employ a cross-sectional study design, which involves collecting data from a sample of the Saudi population at a single point in time. This design allows for the investigation of the association between glucose levels and headache prevalence in a cost-effective and timeefficient manner.

#### Study approach

The study will be conducted in various healthcare facilities and communities across Saudi Arabia, ensuring a diverse representation of the population. This will include primary care clinics, hospitals, and urban as well as rural settings to capture a comprehensive view of the Saudi population.

### **Study population**

The target population for this research is the adult Saudi population aged 18 and above, regardless of gender, residing in different regions of Saudi Arabia. Individuals with known chronic conditions that could influence glucose levels or headache prevalence, such as diagnosed diabetes or other significant medical conditions, will be excluded.

### Study sample

A stratified random sampling technique will be employed to ensure that the sample is representative of different age groups and geographical regions. The sample size will be determined based on statistical power calculations, with an estimated level of significance. It is expected to be sufficiently large to detect significant associations.

#### Study tool

For the current study, questionnaire was adopted for data collection, which was also categorized as a study tool.

### Data collection

Data will be collected through online google form questionnaire. Participants will be asked to provide information on demographics, medical history, lifestyle factors, and dietary habits. Glucose levels will be measured through fasting blood tests, and headache prevalence will be assessed through structured questionnaires and medical records review.

#### Data analysis

Statistical analysis will involve the use of appropriate tests such as chi-square, t-tests, and regression analyses to investigate the association between glucose levels and headache prevalence. Data will be analyzed using statistical software (SPSS), and p-values less than 0.05 will be considered statistically significant.

#### **Ethical considerations**

The study will adhere to ethical principles, including obtaining informed consent from all participants. Ethical approval will be sought from the relevant institutional review board or ethics committee. Confidentiality of participant data will be strictly maintained, and participants



will be informed of their right to withdraw from the study at any point without consequences. The study will also ensure that all procedures are conducted in accordance with the Declaration of Helsinki and other applicable ethical guidelines.

### Results

The study included 574 participants. The most frequent weight among them was 51-65 kg (n= 196, 34.1 Per Cent), followed by 66-75 kg (n= 130, 22.6 Per Cent). Figure 1 shows the weight distribution among study participants. The most frequent height among study participants was 151-160 cm (n= 229, 39.9 Per Cent), followed by 161-170 cm (n= 195, 34 Per Cent). Figure 2 shows the height distribution among study participants. The most frequent age among study participants was less than 25 years (n= 203, 35.4 Per Cent), followed by 25-30 years (n= 143, 24.9 Per Cent). Figure 3 shows the distribution of age among study participants.

The most frequent nationality among study participants was Saudi (n= 547, 95.3 Per Cent), followed by non-Saudi (n= 27, 4.7 Per Cent). Figure 4 shows the distribution of nationality among study participants.

The most frequent gender among study participants was female (n= 351, 61.1 Per Cent), followed by male (n= 223, 38.9 Per Cent). Figure 5 shows the distribution of gender among study participants.

The perceived blood sugar level intake varied among study participants with most of them had normal blood sugar levels. Perceived blood sugar level intake is presented in Figure 6.

Participants were asked if they smoked. Most were not smoking (n=482, 84 Per Cent), and non-smoking were (n=92, 26 Per Cent). Participants were asked to assess their diseases. Their results are presented in Table 1.

Participants were asked about the nature of the headache. The most frequent were they don't have (n= 345, 60.1 Per Cent), followed by sharp (n= 116, 20.2 Per Cent), and the least was continuous (n=113, 19.7 Per Cent). Figure 7 shows the participants' nature of the headache.

Participants' activity levels during the day were frequently inactive (n= 51, 8.9 Per Cent), light active per day (n= 160, 27.9 Per Cent), middle active (n=290, 50.5 Per Cent), and high active 73 participants reported that high active per day (12.7 Per Cent).

Participants were asked about a kind of headache. The most frequent were they don't have (n= 367, 63.9 Per Cent),

followed by stress headaches (n= 116, 20.2 Per Cent), and the lowest cluster (n=15, 2.6 Per Cent).

# Discussion

Migraines and other types of headaches may be brought on or made worse by hypoglycemia (low blood sugar/glucose) <sup>35,36</sup>. When meal intake is inadequate, blood glucose levels may fall too low. If you don't eat, don't change your diet, and don't fast, you'll end up in the same place. One of the early suggestions is that low blood sugar, or hypoglycemia, is a contributing factor in fasting headaches. Migraine attacks may be significantly and uniquely triggered in certain people by elevated blood glucose levels. Migraine episodes are reported to be precipitated or made worse by hypoglycemia<sup>37</sup>. Long-term fasting is associated with an increased risk of migraines, and fasting is one of the most often reported triggers of migraines (39 Per Cent-66 Per Cent across 13 studies, 38 studies, and 40 studies). Fasting headache has been linked to changes in pain receptors in the brain, according to some specialists<sup>38-41</sup>. This is especially true for those who are genetically prone to experiencing such changes. Fasting for religious causes, such as during Ramadan or Yom Kippur, has also been studied in relation to migraine <sup>42-45</sup> with similar results. Some individuals have migraines when they fast overnight [48]. Migraines may be worse by fasting and skipping meals. In one research, a 38-year-old obese lady with migraines saw her episodes lessen when she started drinking orange juice to treat hypoglycemia. The similar correlation between low blood sugar and headaches was also shown by the researchers in a further four subjects. These findings suggest that low glucose levels may have a direct or indirect role in causing headaches associated with fasting. Similarly, six out of twelve patients with migraines reported experiencing pain while fasting, as reported by Blau and Cumings. So, it's possible that hypoglycemia plays a role in the attacks, and eating something is an easy method to avoid headaches if that's the case. Furthermore, Blau and Pyke  $^{\rm 46\text{-}49}$  reiterated that the correction of hypoglycemia in T2D and migraine patients led to a significant decrease in migraine. Of the 36 individuals with migraine and T2D studied, 5 had their migraines alleviated or lessened following blood glucose correction, and 4 had their migraines altered by nocturnal hypoglycemia. Fasting or skipping meals was linked to migraine attacks in 6 of the remaining 27 participants.



The insulin receptor has been demonstrated to be activated by fasting in clinical research. An unexpected trigger for migraine aura is intravenous insulin infusion. Research has shown that a decline in blood glucose levels brought on by insulin treatment might cause headache-like symptoms. However, more evidence was supplied ten years later with the discovery that reactive hypoglycemia generated by sugar might cause migraines . Rarely, a high-sugar meal may cause low blood sugar, a condition known as reactive hypoglycaemia, owing to a sudden rise in blood sugar levels (Hyperglycemia) followed by an overproduction of insulin and a subsequent rapid decrease in blood sugar levels. Hypoglycemia may cause a migraine episode in those who suffer from migraines after extended fasting, and insulin is a crucial regulator for brain glucose metabolism. Migrainerelated changes in brain glycogen metabolism were reviewed by Dalkara and Kilic. During the intense synaptic activity of a headache episode, astrocytes store plasma glucose as glycogen and rapidly breakdown it for glutamate and potassium absorption. A prolonged state of low blood glucose and continuous sympathetic activity during longterm fasting has been hypothesized by the authors to deplete presynaptic astrocytes' ability to produce glucose from glycogen, hence eliciting aura and headache [41]. Studies using magnetic resonance spectroscopy (MRS) have shown, for instance, that people who suffer from migraines have abnormalities in mitochondrial Oxidative Phosphorylation (OXPHOS) <sup>50</sup>, resulting in hypometabolism or low ATP levels. Migraine attacks may be triggered by fasting or the injection of glucose or insulin, according to early research that corroborated these findings. Headache is not a typical complaint in people with symptomatic hypoglycemia, however this is countered by arguments in ICHD-3beta. For instance, most migraine sufferers did not experience headaches when their blood sugar was lowered with insulin. While observing 20 people with migraines for 2 hours, Pearce et al. found that 2 of them had headaches owing to insulin-induced hypoglycemia. Migraine episodes, he reasoned, might result from a combination of low blood sugar and other metabolic processes. Migraine episodes and other metabolic disturbances are thus thought to be linked to an abrupt drop in blood glucose levels.

Hemiplegic migraine and migraine with aura have been linked to a genetic condition called GLUT1 (glucose transport protein type 1) deficient syndrome <sup>51</sup>. The GLUT1 gene (encoded by the SLC2A1 gene) has been reported to have a number of heterozygous mutations. Patients with

this condition have a variety of neurological disorders due to impaired glucose transport across the blood-brain barrier. Migraine headaches have been linked to glucose shortage, which is supported by the GLUT1 deficiency syndrome. Also, in contrast to hypoglycemia's influence on CSD durations<sup>52</sup>, hyperglycemia makes the cortex more resistant to the onset of CSD and speeds up CSD recovery. Migraine aura and headaches may be generated by insulininduced hypoglycemia since it greatly increases the duration of CSD in experimental animals. Experimental research released in 2017 indicated that the metabolic alterations brought on by the injection of insulin, glucagon, or leptin significantly modify neuronal activity the in trigeminovascular system, a key mechanism in the etiology of migraine headaches <sup>53-65</sup>. This demonstrates the potential neurobiological connection between migraine and changes in glucose homoeostasis. One of these SNPs (rs1024905, minor allele G) has been linked to increased migraine risk and lower expression of the C12orf5 gene in the brain's blood, cerebellum, and temporal cortex, according to recent bioinformatics analysis <sup>65-75</sup>. Inhibitory regulation of glucose breakdown (glycolysis) in human cells has recently been attributed to the C12orf5 gene, which has been identified to encode the TP53-inducible glycolysis and apoptosis regulator (TIGAR). As a consequence, lower TIGAR expression leads to enhanced glucose breakdown as a result of lessened inhibition of glycolysis. People with the rs1024905-G risk allele for migraines may also have a higher rate of glucose breakdown <sup>80-107</sup>.

# Conclusion

The results of the study showed that most of the participants were Saudis. Most of them are women. In addition, the majority of people have normal fasting blood sugar levels. Their physical activity is moderate and the majority are non-smokers. In addition, most of the study participants had good and effective social communication.

### References

- Gerring ZF, Powell JE, Montgomery GW, et al. Genome-wide analysis of blood gene expression in migraine implicates immune-inflammatory pathways. Cephalalgia. 2018;38(2):292-303. Doi: https://doi.org/10.1177/0333102416686769
- 2. Vos T, Allen C, Arora M, et al. Global, regional, and national incidence, prevalence, and years lived with

disability for 310 diseases and injuries, 1990–2015: a systematic analysis for the Global Burden of Disease Study 2015. The lancet. 2016;388(10053):1545-602.Doi:

https://doi.org/10.1016/S0140-6736(16)31678-6

- 3. Feigin VL, Abajobir AA, Abate KH, et al. Global, regional, and national burden of neurological disorders during 1990-2015: a systematic analysis for the Global Burden of Disease Study 2015. The Lancet Neurology. 2017;16(11):877-97.Doi: https://doi.org/10.1016/S1474-4422(17)30299-5
- 4. Vos T, Barber RM, Bell B, et al. Global, regional, and national incidence, prevalence, and years lived with disability for 301 acute and chronic diseases and injuries in 188 countries, 1990–2013: A systematic analysis for the Global Burden of Disease Study 2013. The lancet. 2015;386(9995):743-800.Doi: https://doi.org/10.1016/S0140-6736(15)60692-4
- 5. Vos T, Flaxman AD, Naghavi M, et al. Years lived with disability (YLDs) for 1160 sequelae of 289 diseases and injuries 1990-2010: a systematic analysis for the Global Burden of Disease Study 2010. The lancet. 2012;380(9859):2163-96.Doi: https://doi.org/10.1016/S0140-6736(12)61729-2
- 6. Dodick DW. Migraine. Lancet. 2018;391:1315-1330.
- 7. Kowalska M, Prendecki M, Kozubski W, et al. Molecular factors in migraine. Oncotarget. 2016;7(31):50708. Doi: https://doi.org/10.18632 Per Cent2Foncotarget.9367
- Bloudek LM, Stokes M, Buse DC, et al. Cost of 8. healthcare for patients with migraine in five European countries: results from the International Burden of Migraine Study (IBMS). J Headache Pain. 2012;13:361-78.
- 9. Linde M, Gustavsson A, Stovner LJ, et al. The cost of headache disorders in Europe: the Eurolight project. Eur J Neurol. 2012;19(5):703-11.Doi: https://doi.org/10.1111/j.1468-1331.2011.03612.x
- 10. Nyholt DR, Borsook D, Griffiths LR. Migrainomicsidentifying brain and genetic markers of migraine. Nat Rev Neurol. 2017;13(12):725-41.
- 11. Launer LJ, Terwindt GM, Ferrari MD. The prevalence and characteristics of migraine in a population-based cohort: the GEM study. Neurology. 1999;53(3):537-.
- 12. The International Classification of Headache Disorders, 3rd edition (beta version) Cephalalgia. 2013;33:629--808.
- 13. Kelman L. The triggers or precipitants of the acute migraine attack. Cephalalgia. 2007;27(5):394-402.
- 14. Li L, Schulz UG, Kuker W, et al. Age-specific association of migraine with cryptogenic TIA and stroke: population-based study. Neurology.

2015;85(17):1444-51.

https://doi.org/10.1212/WNL.000000000002059

- 15. Gray PA, Burtness HI. Hypoglycemic headache. Endocrinology. 1935;19(5):549-60.Doi: https://doi.org/10.1210/endo-19-5-549
- 16. Gross EC, Lisicki M, Fischer D, et al. The metabolic face of migraine—from pathophysiology to Neurology. treatment. Nature Reviews 2019;15(11):627-43.
- 17. Blau JN, Cumings JN. Method of precipitating and preventing some migraine attacks. British medical 1966;2(5524):1242.Doi: journal. https://doi.org/10.1136 Per Cent2Fbmj.2.5524.1242
- 18. Pearce J. Insulin induced hypoglycaemia in migraine. Journal of neurology, neurosurgery, and psychiatry. 1971;34(2):154.Doi: https://doi.org/10.1136 Per Cent2Fjnnp.34.2.154
- 19. Jacome DE. Hypoglycemia rebound migraine. Headache: The Journal of Head and Face Pain. 2001;41(9):895-8.Doi:

https://doi.org/10.1111/j.1526-4610.2001.01163.x

- 20. Goadsby PJ, Holland PR, Martins-Oliveira M, et al. Pathophysiology of migraine: a disorder of sensory processing. Physiological reviews. 2017.
- 21. Fava A, Pirritano D, Consoli D, eta I. Chronic migraine in women is associated with insulin resistance: a cross-sectional study. Eur J Neurol. 2014;21(2):267-72.Doi: https://doi.org/10.1111/ene.12289
- 22. Dexter JD, Roberts J, Byer JA. The five hour glucose tolerance test and effect of low sucrose diet in migraine. Headache. 1978;18(2):91-4.Doi: https://doi.org/10.1111/j.1526-4610.1978.hed1802091.x
- 23. Cavestro C, Rosatello A, Micca G, et al. Insulin metabolism is altered in migraineurs: a new pathogenic mechanism for migraine?. Headache: Journal of Head and The Face Pain. 2007;47(10):1436-42.Doi:

https://doi.org/10.1111/j.1526-4610.2007.00719.x

- 24. Casucci G, Villani V, Cologno D, et al. Migraine and metabolism. Neurological Sciences. 2012;33:81-5.
- 25. Rainero I, Limone P, Ferrero M, et al. Insulin sensitivity is impaired in patients with migraine. 2005;25(8):593-7.Doi: Cephalalgia. https://doi.org/10.1111/j.1468-2982.2005.00928.x
- 26. Bigal ME, Kurth T, Hu H, et al. Migraine and cardiovascular disease: possible mechanisms of interaction. Neurology. 2009;72(21):1864-71.
- 27. Buse DC, Manack A, Serrano D, et al. Sociodemographic and comorbidity profiles of chronic migraine and episodic migraine sufferers. J Neurol Neurosurg Psychiatry. 2010.





- 28. Burch RC, Rist PM, Winter AC, et al. Migraine and risk of incident diabetes in women: a prospective study. Cephalalgia. 2012;32(13):991-7.
- Hamed SA, Hamed EA, Eldin AM, et al. Vascular risk factors, endothelial function, and carotid thickness in patients with migraine: relationship to atherosclerosis. J Stroke Cerebrovasc Dis. 2010;19(2):92-103.Doi: https://doi.org/10.1016/j.jstrokecerebrovasdis.200 9.04.007
- 30. Scher AI, Terwindt GM, Picavet HS, et al. Cardiovascular risk factors and migraine: the GEM population-based study. Neurology. 2005;64(4):614-20.
- Etminan M, Takkouche B, Isorna FC, et al. Risk of ischaemic stroke in people with migraine: systematic review and meta-analysis of observational studies. Bmj. 2005 J;330(7482):63.Doi:
  - https://doi.org/10.1136/bmj.38302.504063.8F
- 32. Merikangas KR, Fenton BT, Cheng SH, et al. Association between migraine and stroke in a large-scale epidemiological study of the United States. Arch. Neurol. 1997;54(4):362-8.
- Sacco S, Pistoia F, Degan D, et al. Conventional vascular risk factors: Their role in the association between migraine and cardiovascular diseases. Cephalalgia. 2015;35(2):146-64.
- 34. Torelli P, Evangelista A, Bini A, et al. Fasting Headache: A Review of the Literature and New Hypotheses. Headache J. Head Face Pain. 2009;49:744–752.
- Hufnagl KN, Peroutka SJ. Glucose regulation in headache: implications for dietary management. Expert review of neurotherapeutics. 2002;2(3):311.
- 36. Macdonald C. Migraine. Lancet. 1933;1:123–126.
- Finocchi C, Sivori G. Food as trigger and aggravating factor of migraine. Neurological Sciences. 2012;33:77-80.
- 38. Spierings EL, Ranke AH, Honkoop PC. Precipitating and aggravating factors of migraine versus tension-type headache. Headache: The journal of head and face pain. 2001;41(6):554-8.
- 39. Rose FC. Trigger factors and natural history of migraine. Functional neurology. 1986;1(4):379-84.
- Dalkara T, Kılıç K. How does fasting trigger migraine? A hypothesis. Curr. Pain Headache Rep. 2013;17:1-7.
- 41. Robbins L. Precipitating factors in migraine: a retrospective review of 494 patients. Headache: The Journal of Head and Face Pain. 1994;34(4):214-6.
- 42. Turner LC, Molgaard CA, Gardner CH, et al. Migraine trigger factors in a non-clinical

Mexican-American population in San Diego county: implications for etiology. Cephalalgia. 1995;15(6):523-30.

- Scharff L, Turk DC, Marcus DA. Triggers of headache episodes and coping responses of headache diagnostic groups. Headache. 1995;35(7):397-403.
- 44. Peroutka SJ. What Turns on a Migraine? A Systematic Review of Migraine Precipitating Factors Curr Pain Headache Rep. 2014;18:454.
- 45. Abu-Salameh I, Plakht Y, Ifergane G. Migraine exacerbation during Ramadan fasting. J Headache Pain. 2010;11(6):513-7.
- 46. Drescher MJ, Elstein Y. Prophylactic COX 2 inhibitor: An end to the Yom Kippur headache: CME. Headache: The Journal of Head and Face Pain. 2006;46(10):1487-91.Doi: https://doi.org/10.1111/j.1526-4610.2006.00609.x
- 47. Peroutka SJ. Serum glucose regulation and headache. Headache. 2002;42:303–308. Doi: 10.1046/j.1526-4610.2002.02083.x
- 48. Blau JN, Pyke DA. Effect of diabetes on migraine. The Lancet. 1970;296(7666):241-3.Doi: https://doi.org/10.1016/S0140-6736(70)92588-2
- Gross EC, Putananickal N, Orsini AL, et al. Mitochondrial function and oxidative stress markers in higher-frequency episodic migraine. Sci Rep. 2021;11(1):4543.
- Mohammad SS, Coman D, Calvert S. Glucose transporter 1 deficiency syndrome and hemiplegic migraines as a dominant presenting clinical feature. J Paediatr Child Health. 2014;50(12):1025-6.
- Hoffmann U, Sukhotinsky I, Eikermann-Haerter K, et al. Glucose modulation of spreading depression susceptibility. J Cereb Blood Flow Metab. 2013;33(2):191-5.Doi:

https://doi.org/10.1038/jcbfm.2012.132

- 52. Martins-Oliveira M, Akerman S, Holland PR, et al. Neuroendocrine signaling modulates specific neural networks relevant to migraine. Neurobiology of disease. 2017;101:16-26.Doi: https://doi.org/10.1016/j.nbd.2017.01.005
- Westra HJ, Peters MJ, Esko T, et al. Systematic identification of trans eQTLs as putative drivers of known disease associations. Nature genetics. 2013;45(10):1238-43.
- 54. Zou F, Chai HS, Younkin CS, et al. Brain expression genome-wide association study (eGWAS) identifies human disease-associated variants. PLOS Genetics.
  2012 Jun 7;8(6):e1002707.Doi: https://doi.org/10.1371/journal.pgen.1002707
- 55. Bensaad K, Tsuruta A, Selak MA, et al. TIGAR, a p53-inducible regulator of glycolysis and apoptosis. Cell. 2006;126(1):107-20.



- 56. Guldiken B, Guldiken S, Taskiran B, et al. Migraine in metabolic syndrome. The neurologist. 2009;15(2):55-8.
- 57. Bic Z, Blix GG, Hopp HP, et al. In search of the ideal treatment for migraine headache. Medical hypotheses. 1998;50(1):1-7.Doi: https://doi.org/10.1016/S0306-9877(98)90170-0
- 58. Altamura C, Corbelli I, De Tommaso M, et al. Pathophysiological bases of comorbidity in migraine. Front Hum Neurosci. 2021:131.
- 59. Del Moro L, Rota E, Pirovano E, et al. Migraine, brain glucose metabolism and the "neuroenergetic" hypothesis: a scoping review. The Journal of Pain. 2022;23(8):1294-317.Doi: https://doi.org/10.1016/j.jpain.2022.02.006
- 60. Arnold SE, Arvanitakis Z, Macauley-Rambach SL, et a I. Brain insulin resistance in type 2 diabetes and Alzheimer disease: concepts and conundrums. Nature Reviews Neurology. 2018;14(3):168-81.
- 61. Hamer JA, Testani D, Mansur RB, et al. Brain insulin resistance: A treatment target for cognitive impairment and anhedonia in depression. Experimental neurology. 2019;315:1-8.Doi: https://doi.org/10.1016/j.expneurol.2019.01.016
- 62. Gruber HJ, Bernecker C, Pailer S, et al. Hyperinsulinaemia in migraineurs is associated with nitric oxide stress. Cephalalgia. 2010;30(5):593-8.
- 63. Wang X, Li X, Diao Y, et al. Are glucose and insulin metabolism and diabetes associated with migraine? a community-based, case-control study. J Oral Facial Pain Headache. 2017;31(3).
- 64. Bernecker C, Pailer S, Kieslinger P, et al. GLP-2 and leptin are associated with hyperinsulinemia in nonobese female migraineurs. Cephalalgia. 2010;30(11):1366-74.Doi: https://doi.org/10.1177/0333102410364674
- 65. Sachdev A, Marmura MJ. Metabolic syndrome and migraine. Front Neurol. 2012;3:161.
- 66. Bhoi SK, Kalita J, Misra UK. Metabolic syndrome and insulin resistance in migraine. J Headache Pain. 2012;13(4):321-6.
- 67. Sacco S, Altobelli E, Ornello R, et al. Insulin resistance in migraineurs: Results from a casecontrol study. Cephalalgia. 2014;34(5):349-56.Doi: https://doi.org/10.1177/0333102413511155
- Porte Jr D, Baskin DG, Schwartz MW. Insulin signaling in the central nervous system: a critical role in metabolic homeostasis and disease from C. elegans to humans. Diabetes. 2005;54(5):1264-76.
- Schwartz MW, Figlewicz DP, Baskin DG, et al. Insulin in the brain: a hormonal regulator of energy balance. Endocrine reviews. 1992;13(3):387-414.Doi: https://doi.org/10.1210/edrv-13-3-387

- Cetinkalp S, Y Simsir I, Ertek S. Insulin resistance in brain and possible therapeutic approaches. Curr Vasc Pharmacol. 2014;12(4):553-64.
- McCarthy LC, Hosford DA, Riley JH, et al. Singlenucleotide polymorphism alleles in the insulin receptor gene are associated with typical migraine. Genomics. 2001;78(3):135-49.Doi: https://doi.org/10.1006/geno.2001.6647
- Netzer C, Freudenberg J, Heinze A, et al. Replication study of the insulin receptor gene in migraine with aura. Genomics. 2008;91(6):503-7.Doi:

https://doi.org/10.1016/j.ygeno.2008.03.006

- Lee J, Pilch PF. The insulin receptor: structure, function, and signaling. Am J Physiol. 1994;266(2):C319-34.
- Adashi EY, Hsueh AJ, et al. Disparate effect of clomiphene and tamoxifen on pituitary gonadotropin release in vitro. American Journal of Physiology-Endocrinology and Metabolism. 1981;240(2):E125-30.
- 75. Cortelli P, Pierangeli G. Hypothalamus and headaches. Neurological Sciences. 2007;28:S198-202.
- Holland P, Goadsby PJ. The hypothalamic orexinergic system: pain and primary headaches: CME. Headache: The Journal of Head and Face Pain. 2007 J;47(6):951-62.
- 77. Alstadhaug KB. Migraine and the hypothalamus. Cephalalgia. 2009;29(8):809-17.Doi: https://doi.org/10.1111/j.1468-2982.2008.01814.x
- Schulte LH, Allers A, May A. Hypothalamus as a mediator of chronic migraine. Neurology. 2017;88:2011.
- Denuelle M, Fabre N, Payoux P, et al. Hypothalamic activation in spontaneous migraine attacks. Headache: The Journal of Head and Face Pain. 2007;47(10):1418-26.
- Kim C, Siscovick DS, Sidney S, et al. Oral contraceptive use and association with glucose, insulin, and diabetes in young adult women: the CARDIA study. Diabetes Care. 2002;25(6):1027-32.
- Favoni V, Giani L, Al-Hassany L, et al. CGRP and migraine from a cardiovascular point of view: what do we expect from blocking CGRP?. The journal of headache and pain. 2019;20:1-7.
- Edvinsson L. The Trigeminovascular Pathway: R ole of CGRP and CGRP Receptors in Migraine. Headache: The Journal of Head and Face Pain. 2017;57:47-55.Doi:

https://doi.org/10.1111/head.13081

 Lassen LH, Haderslev PA, Jacobsen VB, et al. CGRP may play a causative role in migraine. Cephalalgia. 2002;22(1):54-61.



- Iyengar S, Johnson KW, Ossipov MH, et al. CGRP and the trigeminal system in migraine. Headache: The Journal of Head and Face Pain. 2019;59(5):659-81.Doi: https://doi.org/10.1111/head.13529
- 85. Gram DX, Ahrén B, Nagy I, et al. Capsaicin-sensitive sensory fibers in the islets of Langerhans contribute to defective insulin secretion in Zucker diabetic rat, an animal model for some aspects of human type 2 diabetes. Eur J Neurosci. 2007;25(1):213-23.Doi: https://doi.org/10.1111/j.1460-9568.2006.05261.x
- 86. Walker CS, Li X, Whiting L, et al. Mice lacking the neuropeptide α-calcitonin gene-related peptide are protected against diet-induced obesity. Endocrinology. 2010;151(9):4257-69.Doi: https://doi.org/10.1210/en.2010-0284
- Hosseinpour M, Maleki F, Khoramdad M, et al. A systematic literature review of observational studies of the bilateral association between diabetes and migraine. Diabetes Metab Syndr Clin. Res Rev. 2021;15(3):673-8. Doi: https://doi.org/10.1016/j.dsx.2021.03.018
- Dodick DW, Goadsby PJ, Spierings EL, et al. Safety and efficacy of LY2951742, a monoclonal antibody to calcitonin gene-related peptide, for the prevention of migraine: a phase 2, randomised, double-blind, placebo-controlled study. The Lancet Neurology. 2014;13(9):885-92.Doi: https://doi.org/10.1016/S1474-4422(14)70128-0
- Melnyk A, Himms-Hagen J. Resistance to aging-associated obesity in capsaicin-desensitized rats one year after treatment. Obesity Research. 1995;3(4):337-44.Doi: https://doi.org/10.1002/j.1550-8528.1995.tb00159.x
- 90. Pettersson M, Ahren BO, Böttcher G, et al. Calcitonin gene-related peptide: Occurrence in pancreatic islets in the mouse and the rat and inhibition of insulin secretion in the mouse. Endocrinology. 1986;119(2):865-9.Doi: https://doi.org/10.1210/endo-119-2-865
- 91. Gram DX, Hansen AJ, Wilken M, et al. Plasma calcitonin gene-related peptide is increased prior to obesity, and sensory nerve desensitization by capsaicin improves oral glucose tolerance in obese Zucker rats. Eur J Endocrinol. 2005153(6):963-9.
- 92. Saeedi P, Petersohn I, Salpea P, et al. Global and regional diabetes prevalence estimates for 2019 and projections for 2030 and 2045: Results from the International Diabetes Federation Diabetes Atlas. Diabetes Res Clin Pract. 2019;157:107843.Doi:

https://doi.org/10.1016/j.diabres.2019.107843

93. Haghighi FS, Rahmanian M, Namiranian N, et al. Migraine and type 2 diabetes; is there any association?. J Diabetes Metab Disord. 2015;15:1-7.

- Split W, Szydlowska M. Headaches in non insulindependent diabetes mellitus. Functional neurology. 1997;12(6):327-32.
- 95. Aamodt AH, Stovner LJ, Midthjell K, et al. Headache prevalence related to diabetes mellitus. The Head-HUNT study. Eur J Neurol. 2007;14(7):738-44.Doi: https://doi.org/10.1111/j.1468-1331.2007.01765.x
- 96. Berge LI, Riise T, Fasmer OB, et al. Does diabetes have a protective effect on migraine?. Epidemiology. 2013:129-34.
- 97. Burn WK, Machin D, Waters WE. Prevalence of migraine in patients with diabetes. Br Med J. 1984;289(6458):1579.Doi: https://doi.org/10.1136/bmj.289.6458.1579-a
- Hagen K, Åsvold BO, Midthjell K, et al. Inverse relationship between type 1 diabetes mellitus and migraine. Data from the Nord-Trøndelag Health Surveys 1995–1997 and 2006–2008. Cephalalgia. 2018;38(3):417-26.Doi:

https://doi.org/10.1177/0333102417690488

 Antonazzo IC, Riise T, Cortese M, et al. Diabetes is associated with decreased migraine risk: A nationwide cohort study. Cephalalgia. 2018;38(11):1759-64.Doi:

https://doi.org/10.1177/0333102417748573

- 100.Fagherazzi G, El Fatouhi D, Fournier A, et al. Associations between migraine and type 2 diabetes in women: Findings from the E3N cohort study. JAMA neurology. 2019 Mar 1;76(3):257-63.
- 101.Farsani SF, Souverein PC, van der Vorst MM, et al. Chronic comorbidities in children with type 1 diabetes: a population-based cohort study. Arch Dis Child. 2015:archdischild-2014.
- 102.López-de-Andrés A, Luis del Barrio J, et al. Migraine in adults with diabetes; is there an association? Results of a population-based study. Diabetes Metab Syndr Obes Targets. 2018:367-74.Doi: https://doi.org/10.2147/DMSO.S170253
- 103.Tantucci C, Bottini P, Fiorani C, et al. Cerebrovascular reactivity and hypercapnic respiratory drive in diabetic autonomic neuropathy. J Appl Physiol. 2001;90(3):889-96.Doi: https://doi.org/10.1152/jappl.2001.90.3.889
- 104.Wijnhoud AD, Koudstaal PJ, Dippel DW. Relationships of transcranial blood flow Doppler parameters with major vascular risk factors: TCD study in patients with a recent TIA or nondisabling ischemic stroke. J Clin Ultrasound. 2006;34(2):70-6.Doi: https://doi.org/10.1002/jcu.20193

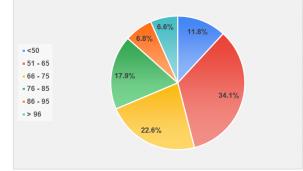


- 105.Ziegler D. Treatment of diabetic polyneuropathy: update 2006. Ann N Y Acad. Sci. 2006;1084(1):250-66.Doi: https://doi.org/10.1196/annals.1372.008
- 106. Rivera-Mancilla E, Al-Hassany L, Villalón CM, et al. Metabolic aspects of migraine: association with obesity and diabetes mellitus. Frontiers in neurology. 2021;12:686398.
- 107.Amin FM, Aristeidou S, Baraldi C, et al. The association between migraine and physical exercise. J Headache. 2018;19:1-9.

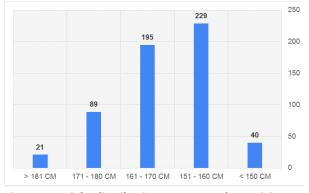
# **Tables & Figures**

# Table 1: Participant disease in the study.

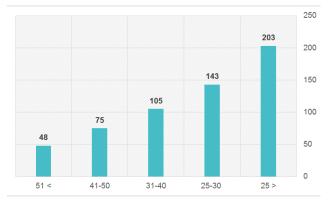
Survey item	Yes	No
	148	426
Do you have frequent urination?	25.80 Per Cent	74.20 Per Cent
	158	416
Do you have blurred eyes?	27.50 Per Cent	72.50 Per Cent
	88	486
Do you have frequent urinary tract infections?	15.30 Per Cent	84.70 Per Cent

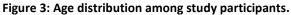












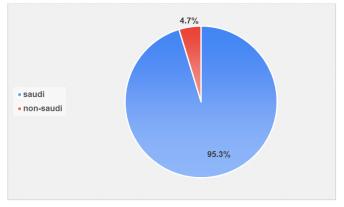


Figure 4: Nationality distribution among study participants.



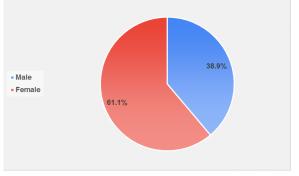


Figure 5: Gender distribution among study participants.

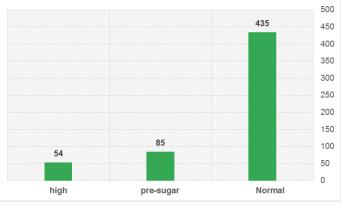


Figure 6: blood sugar level distribution among study participants.

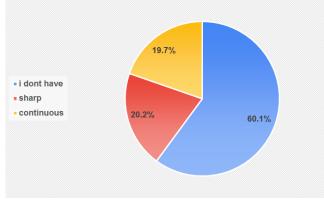


Figure 7 shows the participants' nature of the headache.

# **ANNEX 1: Data Collection Tool**

- 1. How old are you?
- 25 years and less
- 25-30 years
- 31-40 years
- 41-50 years
- 50 years and more
- 2. What is your gender?
- Male
- Female
- 3. What is your nationality?

- Saudi
- Non-Saudi
- 4. What is your height?
- <150 cm
- 151-160 cm
- 161-170 cm
- 171-180 cm
- >181 cm
- 5. What is your weight?
  - <50 Kg

•

•

•

•

6.

.

.

•

7.

•

•

•

.

8.

- 51-65 Kg
- 66-75 Kg
- 76-85 Kg
- 86-95 Kg
- >96 Kg
  - What is your Measuring blood sugar?
- Don't suffer from diabetes
  - Yes, type 1
  - Yes, type 2
  - Do you smoke?
  - Yes
  - No
- What is your marital status?
- Single
- Married
  - Absolute/divorced
- idower/widow
- 9. Do you smoke?
- Yes
- No
- 10. What is your activity level during the day?
- Inactive
- Light
- Middle
- High
- 11. Do you suffer from headaches?
- I do not suffer from headaches
- Migraine
- Cluster
- Stress
- **12.** How long have you been suffering from headaches?
- I don't suffer from headaches
- Less than two years
- 2-5 years
- 5-10 years



•	Furthermore	•	Once a month	
13.	How many headaches do you have during a	•	2-3 times a month	
month	?	•	4-5 times a month	
•	I don't suffer from headaches	•	More than 5 times a month	
•	One a month	19.	Do you eat your daily meals regularly?	
•	2-3 times a month	•	Yes	
•	4-5 times a month	•	No	
•	More than 5 times a month	20.	How many meals do you eat a day?	
14.	The nature of the headache	•	One meal	
•	I don't suffer from headaches	•	2-3 meals	
•	Sharp	•	4-5 meals	
•	Continuous	•	More than one meal	
15.	Does she suffer from any of these symptoms	21.	Do you suffer from headache attacks during the	
before	the headache?	same time of the day?		
•	Nausea	•	I don't suffer from headaches	
•	Vomiting	•	Yes, during the same time	
•	Facial numbness	•	No, at different time	
•	Tinnitus	22.	How many meals do you eat a day?	
•	None of the above	•	One meal	
16.	Does she suffer from any of these symptoms	•	2-3 meals	
before	the headache?	•	4-5 meals	
•	Nausea	•	More than one meal	
•	Vomiting	23.	Do you have a frequent urination?	
•	Facial numbness	•	Yes	
•	Tinnitus	•	No	
•	None of the above	24.	Do you have a blurred eye?	
17.	Do you suffer from polycystic ovary syndrome?	•	Yes	
•	Does not apply	•	No	
•	Yes	25.	Do you have a frequent urinary track infection?	
•	No	•	Yes	
18.	How often do you eat sugary meals?	•	No	
•	Don't eat			

# Appendix 2: Participants responses to scale items

variable		Frequency	Percent
	25 >	203	35.4 Per Cent
	25-30 years	143	24.9 Per Cent
Age	31-40	105	18.3 Per Cent
	41-50	75	13.1 Per Cent
	51 <	48	8.4 Per Cent
Candar	Male	223	38.9 Per Cent
Gender	Female	351	61.1 Per Cent
	Saudi		95.3 Per Cent
nationality	non-Saudi	27	4.7 Per Cent



	Normal	435	75.8 Per Cent
blood sugar level	pre-sugar	85	14.8 Per Cent
	high	54	9.4 Per Cent
	<50	68	11.8 Per Cent
	51 - 65	196	34.1 Per Cent
woight	66 - 75	130	22.6 Per Cent
weight	76 - 85	103	17.9 Per Cent
	86 - 95	39	6.8 Per Cent
	> 96	38	6.6 Per Cent
	< 150 CM	40	7.0 Per Cent
	151 - 160 CM	229	39.9 Per Cent
high	161 - 170 CM	195	34.0 Per Cent
	171 - 180 CM	89	15.5 Per Cent
	> 181 CM	21	3.7 Per Cent

what are the things that relieve headaches		
	Frequency	Percent
Pain killers	185	40.0 Per Cent
Sleep	149	32.2 Per Cent
lce pack	13	2.8 Per Cent
Coffee	84	18.1 Per Cent
Salty food	18	3.9 Per Cent
Sweet food	10	2.2 Per Cent
Fasting	2	0.4 Per Cent
Hunger	2	0.4 Per Cent
Drought	0	0.0 Per Cent

what are the make headaches worse			
	Frequency	Percent	
Pain killers	9	2.1 Per Cent	
Sleep	24	5.5 Per Cent	
Ice pack	15	3.4 Per Cent	
Coffee	33	7.6 Per Cent	
Sweet food	26	6.0 Per Cent	
Fasting	81	18.6 Per Cent	
Hunger	145	33.3 Per Cent	
Drought	103	23.6 Per Cent	



Г

	Frequency	Percent
Nausea	99	32.7 Per Cent
Vomiting	31	10.2 Per Cent
Facial numbness	26	8.6 Per Cent
Tinnitue	60	19.8 Per Cent
Non of above	87	28.7 Per Cent
Have you cha	anged your lifestyle to reduce headaches? Frequency	Percent
		Percent 13.6 Per Cent
Have you cha Playing sports Reduce weight	Frequency	
Playing sports	Frequency 65	13.6 Per Cent
Playing sports Reduce weight	Frequency           65           19	13.6 Per Cent 4.0 Per Cent
Playing sports Reduce weight Sleep regulation	Frequency           65           19           113	13.6 Per Cent 4.0 Per Cent 23.6 Per Cent
Playing sports Reduce weight Sleep regulation Eat sugar	Frequency           65           19           113           9	13.6 Per Cent 4.0 Per Cent 23.6 Per Cent 1.9 Per Cent

٦

What is your activity level during the day?			
	Frequency	Percent	
inactive	51	8.9 Per Cent	
light	160	27.9 Per Cent	
middle	290	50.5 Per Cent	
high	73	12.7 Per Cent	
Total	574	100.0 Per Cent	

Do you have headaches?			
	Frequency	Percent	
don't have	367	63.9 Per Cent	
migraine	76	13.2 Per Cent	
cluster	15	2.6 Per Cent	
stress	116	20.2 Per Cent	
Total	574	100.0 Per Cent	



How long you have been suffering from? Headache			
	Frequency	Percent	
Don't have	367	63.9 Per Cent	
Less than 2 years	94	16.4 Per Cent	
2-5 years	59	10.3 Per Cent	
5-10 years	30	5.2 Per Cent	
More than that	24	4.2 Per Cent	
Total	574	100.0 Per Cent	

What is the nature of the headache?			
	Frequency	Percent	
dont have	345	60.1 Per Cent	
sharp	116	20.2 Per Cent	
continuous	113	19.7 Per Cent	
Total	574	100.0 Per Cent	

How many headache do you have during month			
	Frequency	Percent	
l dont have	260	45.3 Per Cent	
One time a month	95	16.6 Per Cent	
2-3 times amonth	124	21.6 Per Cent	
4-5 times a month	59	10.3 Per Cent	
More than 5 times a month	36	6.3 Per Cent	
Total	574	100.0 Per Cent	

Do the menstrual cycle effect to headaches						
Frequency Percent						
Does not apply	276	48.1 Per Cent				
Yes, it increases	152	26.5 Per Cent				
Yes. It reduces	10	1.7 Per Cent				
Don't affect	136	23.7 Per Cent				
Total	100.0 Per Cent					

How many su	ugary meals do you eat?	
	Frequency	Percent



Don't eat	45	7.8 Per Cent	
Once a month	39	6.8 Per Cent	
2-3 times a month	139	24.2 Per Cent	
4-5 times a month	112	19.5 Per Cent	
More than 5 times a month	239	41.6 Per Cent	
Total	574	100 Per Cent	

	Do you have polycystic ovary syndrome?					
	Frequency Percent					
does not apply	251	43.7 Per Cent				
yes	56	9.8 Per Cent				
no	267	46.5 Per Cent				
Total	574	100.0 Per Cent				

	How many meals do you eat a daily					
	Frequency	Percent				
One meal	61	10.6 Per Cent				
2-3 meals	417	72.6 Per Cent				
4-5 meals	83	14.5 Per Cent				
More than 5 meals	13	2.3 Per Cent				
Total	574	100 Per Cent				

Do you have headaches attacks during the same time of the day?								
	Frequency Percent							
Don't have headaches	301	52.4 Per Cent						
Yes, during the same time	57	9.9 Per Cent						
No, at different time	216	37.6 Per Cent						
Total	574	100 Per Cent						

SPSS

Chi-square

		Crossta	ab			
Count						
			Do you have	headaches		
		I dont have	Migraine	Cluster	Stress	Total
Do you have diabetes	I don't have	337	71	11	98	517
	Yes, type 1	15	1	2	10	28
	Yes, type 2	15	4	2	8	29



Г

Total         367         76         15         116         574						
	Total	367	76	15	116	574

	Chi-Square Tests		
			Asymptotic Significance (2-
	Value	df	sided)
Pearson Chi-Square	12.566ª	6	.050
Likelihood Ratio	11.439	6	.076
Linear-by-Linear Association	5.109	1	.024
N of Valid Cases	574		

			Crosstab				
Count							
			Hov	v long heada	che		
less than 2							
		i dont have	years	2-5 years	5-10 years	more than that	Total
Do you have diabetes	I dont have	338	85	49	23	22	517
	yes, type 1	15	4	4	4	1	28
	yes, type 2	14	5	6	3	1	29
Total		367	94	59	30	24	574

	Chi-Square Tests		
	Value	df	Asymptotic Significance (2- sided)
Pearson Chi-Square	12.176 <sup>a</sup>	8	.144
Likelihood Ratio	9.853	8	.276
Linear-by-Linear Association	4.968	1	.026
N of Valid Cases	574		

			Crossta	ıb			
Count							
			F	low many heada	iche		
	one time a 2-3 times a 4-5 times a more than 5						
		i dont have month month month times a month			Total		
Do you have diabetes	I don't have	238	86	110	51	32	517
	yes, type 1	12	5	5	4	2	28
	yes, type 2	10	4	9	4	2	29
Total		260	95	124	59	36	574

Chi-Square Tests						
	Asymptotic Significance (2-					
	Value	df	sided)			
Pearson Chi-Square	3.343 <sup>ª</sup>	8	.911			
Likelihood Ratio	3.204	8	.921			
Linear-by-Linear Association	1.790	1	.181			
N of Valid Cases	574					



Crosstab							
Count							
	Nature headache						
		dont have	sharp	continuous	Total		
Do you have diabetes	I dont have	318	106	93	517		
	yes, type 1	13	6	9	28		
	yes, type 2	14	4	11	29		
Total		345	116	113	574		

Chi-Square Tests						
	Value	df	Asymptotic Significance (2- sided)			
Pearson Chi-Square	10.217 <sup>a</sup>	4	.037			
Likelihood Ratio	9.022	4	.061			
Linear-by-Linear Association	7.356	1	.007			
N of Valid Cases	574					

Crosstab							
Count							
Menstrual cycle headaches							
		does not apply	yes, it increases	yes. it reduces	don't affect	Total	
Do you have diabetes	I dont have	243	136	9	129	517	
	yes, type 1	16	9	0	3	28	
	yes, type 2	17	7	1	4	29	
Total		276 152 10 136					

Chi-Square Tests						
			Asymptotic Significance (2-			
	Value	df	sided)			
Pearson Chi-Square	6.187 <sup>a</sup>	6	.403			
Likelihood Ratio	7.199	6	.303			
Linear-by-Linear Association	3.810	1	.051			
N of Valid Cases	574					

Regression

Model Summary					
Model	R	R Square	Adjusted R Square	Std. Error of the Estimate	
1	0.155 <sup>a</sup>	0.024	0.014	0.475	

	ANOVA <sup>a</sup>								
Model		Sum of Squares	df	Mean Square	F	Sig.			
1	Regression	3.140	6	.523	2.319	.032 <sup>b</sup>			
	Residual	127.974	567	.226					
	Total	131.115	573						

	Coefficients <sup>a</sup>						
				Standardized			
		Unstandardized Coefficients		Coefficients			
Model		В	Std. Error	Beta	t	Sig.	
1	(Constant)	1.115	0.057		19.607	0.000	



Do you have headaches	0.017	0.024	0.042	0.715	0.475
How long headache	0.021	0.025	0.048	0.815	0.415
How many headache	-0.021-	0.025	-0.057-	-0.863-	0.388
Nature headache	0.073	0.042	0.121	1.728	0.085
Menstrual cycle headaches	-0.031-	0.017	-0.079-	-1.881-	0.060
Headaches attacks during same time day	-0.020-	0.029	-0.039-	-0.695-	0.487

Received: 27-Nov-2023, Manuscript No. AMJ-23-4007; Editor assigned: 30-Nov-2023, PreQC No. AMJ-23-4007(PQ); Reviewed: 14-Dec-2023, QC No. AMJ-23-4007; Revised: 19-Dec-2023, Manuscript No. AMJ-23-4007(R); Published: 26-Dec-2023