Association of Migraine with Its Comorbidities and Food Specific Immunoglobulin G Antibodies and Inflammatory Cytokines: Cross-Sectional Clinical Research

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**Purpose:** The relationship between food allergy caused by food specific IgG antibodies and migraine has received increased attention in recent years. Here, we aimed to evaluate the effects of food specific IgG antibodies on headache, gastrointestinal symptoms, anxiety, depression, sleep disorders, dermatosis, and serum inflammatory cytokines in migraine patients, and to quantitatively assess the effect of IgG levels on the severity of headache and its comorbidities.

**Methods:** Of 89 migraine patients, those who had one or more food specific IgG antibodies ≥50 U/mL were classified into the IgG positive group, which was then further divided into subgroups based on differing numbers of food allergens. All other subjects were classified into the IgG negative group. We compared the frequency and severity of migraine, anxiety, depression, sleep disorders, dermatosis, and inflammatory cytokines between groups. A regression model was performed to further assess the effect of overall positive IgG concentration and the mediation effect of inflammatory cytokines.

**Results:** Participants in the positive IgG group (n = 67) were more likely to have longer time elapsed since diagnosis, more frequent and severe migraine, a higher risk of developing anxiety and gastrointestinal symptoms, along with higher IL-6 and TNF-α. Subgroups with more food allergens generally had worse conditions as well. After adjusting for the inflammatory cytokines, the effect of IgG was reduced.

**Conclusion:** Migraine patients with positive food specific IgG antibodies had worse migraine, anxiety, and gastrointestinal symptoms. Inflammatory cytokines partially mediate the causal pathway between food specific IgG antibodies, migraine, and migraine comorbidities.

**Keywords:** migraine, food, immunoglobulin G antibody, gastrointestinal symptoms, anxiety, interleukin-6, tumor necrosis factor-α

**Introduction**

Migraine is a recurrent headache disease often characterized by a pulsating sensation, and is associated with a variety of comorbidities including gastrointestinal disorders, neuropsychiatric disorders (ie, anxiety, depression, suicide ideation) etc.\(^1\) These symptoms may have a dramatic negative impact on one’s quality of life and thus might be classified as a type of disability, especially when symptoms interfere with the person’s daily activities. Many epidemiological studies have revealed a high prevalence of migraine and its accompanied disorders, as well as...
a burden on individuals and society. Based on a systematic review involving 19 population-based studies from 17 countries, the aggregate weighted estimates prevalence of definite and suspected migraines over a 12-month period was 18.5%. Migraine attacks can be associated with a variety of factors, including dietary consumption. Allergies mediated by food specific IgG antibodies can cause chronic aseptic inflammation in many systems of the body. High circulating serum concentrations of some IgG antibodies have been measured in certain atopic individuals. The relationship between food allergy caused by specific IgG positive food and migraine has received increased attention. Some scholars have proposed that IgG may be a biomarker used to identify foods related to enhanced inflammatory response in vivo, which has been suggested as a crucial role in migraine mechanism and pathology.

Proinflammatory cytokines, interleukin-6 (IL-6) and tumor necrosis factor-α (TNF-α), are considered to be the inducers of visceral pain, and migraine patients have been shown to have higher serum levels of IL-6 and TNF-α. Interleukin-10 (IL-10) is a potent anti-inflammatory cytokine which has been demonstrated to inhibit the production of inflammatory cytokines and attenuate the inflammatory signaling pathway. Though the connection between migraine and inflammatory cytokines has been reported, there are no published studies that quantitatively evaluate the correlation between inflammatory cytokines and food specific IgG antibodies in migraine patients.

The aim of this study was to evaluate the effects of food specific IgG antibodies on headaches, gastrointestinal symptoms, anxiety, depression, sleep disorders, dermatosis, and serum inflammatory cytokines (IL-6, IL-10 and TNF-α), and to explore the quantitative correlation between the concentration and types of serum positive food specific IgG antibodies and symptom scores and the levels of inflammatory cytokines in migraine patients. The findings from this study may improve the integration of evidence regarding the mechanisms of migraine and provide new insights for migraine prevention strategies and treatment plans.

Methods

Participants

This cross-sectional study involved participants with a neurologist-confirmed migraine diagnosis defined by The International Classification of Headache Disorders, 3rd edition (ICHD-3). All participants were recruited during their visit to the Health Center of Screening and Prevention of Diseases of the First affiliated Hospital of Harbin Medical University, between October 2020 and February 2021. Participants were screened for study eligibility, inclusion criteria were: 1) female or male ages 18 to 62; 2) minimum six months’ time elapsed since migraine diagnosis; and 3) two or more migraine attacks in the past 30 days. Participants whose headaches were caused by another diagnosed disease were excluded (ie, hypertension, stroke, infectious disease, malignant/benign tumors, mental health disorders), participants were also excluded if they had a peptic ulcer, and currently used immunosuppressive drugs.

The ethics and protocol of this research study were assessed and approved by the Ethics committee of the First Affiliated Hospital of Harbin Medical University (No.201829). This study was conducted in accordance with the Declaration of Helsinki. We discussed the research protocol and the potential risks with each participant, all participants provided written informed consent. This study has been registered in the Chinese Clinical Trial Registry (Registration number: ChiCTR2000039278).

Data Collection

A general Health Questionnaire was completed by a trained survey coordinator. Demographic information and past medical history related to migraine were gathered. The major information in this instrument included: 1) age; 2) gender; 3) body height (cm), body weight (kg), and body mass index (BMI), calculated as weight (in kg)/ height² (in m²); 4) migraine with or without aura; 5) episodic migraine or chronic migraine; 6) time elapsed since initial migraine diagnosis. To collect the information regarding the headache frequency and average pain intensity, participants were asked to complete the Migraine Disability Assessment questionnaire (MIDAS), with two supplemental questions: “On how many days in the last 3 months did you have a headache (If a headache lasted more than 1 day, count each day)” and “On a scale of 0–10, on average how painful were these headaches? (Where 0 = no pain at all, and 10 = pain as bad as it can be)” (Visual Analog Scale, VAS).

We also assessed the impact of migraine on participants’ daily activity, ability to work, quality of life, depression, anxiety, sleep quality, and gastrointestinal symptoms through the following five questionnaires: 1) Headache Impact Test-6 (HIT-6), scores range from 36 to 78, with higher scores indicating greater adverse impact; 2) Migraine-Specific Quality of Life Questionnaire version
2.1 (MSQ v2.1), with raw dimension scores ranging from 14 to 84, higher scores indicating worse health status; 3) Self-Rating Anxiety Scale (SAS), with an index score ranging from 25 to 100, a score ≥ 50 is defined as experiencing anxiety; 4) Self-Rating Depression Scale (SDS) which has a similar scoring structure as SAS, and an index score ≥ 50 is defined as experiencing depression; 5) Pittsburgh Sleep Quality Index (PSQI) with questions related to bed partners removed, a PSQI ≥ 8 is defined as poor sleep quality; and 6) Gastrointestinal Symptom Rating Scale (GSRS). Participants’ dermatosis was diagnosed by an experienced dermatologist.

Sample Collection and Measurements

Blood Sample
A fasting 5 mL venous blood sample was drawn from an antecubital vein from each participant, using serum separator tube. The whole undisturbed blood samples were allowed to clot at room temperature for two hours, and were then centrifuged for 20 min at 1000 × g. The supernatant of the serum was stored at −80 degrees Celsius.

Food Specific IgG Antibodies
We used enzyme-linked immunosorbent assays (ELISA) to measure the concentration of 14 food specific IgG antibodies (Bioeurope GmbH, Germany). A standard curve was generated for each targeted IgG antibody by plotting the mean absorbance as the Y axis against the IgG concentration as X axis. The procedure was optimized by using negative control, the concentration of IgG was calculated based on the standard curve. According to the manufacturer’s recommendations, all concentrations ≥ 50 u/mL were considered a positive reaction to a certain food. Based on the results, participants were assigned to the IgG positive group if at least one food specific IgG antibody was positive, otherwise, they were assigned to the IgG negative group. Each participant assigned to the IgG positive group was categorized into subgroups, one positive food allergen, two positive food allergens, and three or more positive food allergens. IgG concentrations were summed up as a total positive IgG, if any IgG concentration ≥50 u/mL.

IL-6, IL-10, and TNF-α
The concentrations of IL-6, IL-10, and TNF-α were measured using ELISA kits purchased from BIM (San Francisco, CA, assay range: 1.5 pg/mL–48 pg/mL), Shanghai Hengyuan Biological Technology Co. Ltd (assay range: 10 ng/L–400 ng/L), and BIM (San Francisco, CA, assay range: 2.5 pg/mL–80 pg/mL), respectively. The procedures were similar as described before. All ELISA assays were performed using Biochrom Anthos 2010 Microplate Reader.

Statistical Analysis
We checked the distribution, outliers, and missingness for the food specific IgG antibodies, three inflammatory cytokines, and data collected from the questionnaires. Data with highly skewed distributions would be log-transformed to be less skewed data. Descriptive statistics were used to compare between (1) the groups with different numbers of food allergens; and (2) positive and negative IgG groups. The continuous and frequency data were described using mean ± SD and proportion (95% CI), respectively. Chi-squared was used to test the difference of the proportions between groups. For the continuous data, the independent sample t-test was conducted if the comparison involved two groups, while ANOVA was used when more than two groups were involved (ie, groups with different number of food allergens).

Consequently, we built a multivariate linear and logistic regression model to further assess the association between each outcome variable of migraine and the related comorbidities with IgG concentration. We then included the three inflammatory cytokines in the regression model to assess the potential mediation effect of the inflammatory cytokines in the association between IgG levels and migraine/comorbidities. The odds ratio (OR) and the 95% CI were reported. The regression analysis was only performed within the group with positive IgG. Data were analyzed using SAS 9.4.

Results
The sample comprised of 89 eligible participants, the majority was female (60 vs 29) with a mean age of 40.3 years (SD = 9.2). Average BMI was 23.5 (SD = 3.7). The time elapsed since initial diagnosis ranged from 12 to 540 months, the median time was 72 months, with 12.4% of the patients reporting chronic headache.

Serum Food Specific IgG Antibodies
IgG antibodies, of varying concentrations, were detected in all participants, 22 of whom were assigned to the IgG negative group, with no food specific IgG concentrations ≥ 50 u/mL (24.7%) and 67 participants had at least one food specific IgG ≥ 50 u/mL and were thus classified to the IgG positive group (75.3%). Participants with one, two, and three or more positive food specific IgG antibodies accounted for 36 (53.7%), 23 (34.3%), and 8 (11.9%) positive IgG participants, respectively.
We found that the top two most prevalent food allergens were eggs and cow’s milk. Among the 67 participants in the positive group, 43.3% had positive egg specific IgG and 26.9% had positive cow’s milk specific IgG. While rice was found to have the lowest frequency of positive cases (Figure 1). For participants in the positive IgG group, total positive IgG concentrations ranged from 54.3 to 577.4 U/mL, with a mean of 212.4 U/mL (STD = 153.9 U/mL), and a median of 178.5 U/mL.

Characteristics Between the IgG Positive and Negative Groups, and Groups with Different Numbers of Food Allergens

Demographic characteristics, migraine characteristics, migraine-related comorbidities, and inflammatory cytokines were compared between the different IgG groups, as seen in Table 1. Demographic characteristics did not significantly differ between the IgG positive and negative groups, nor among the three IgG positive subgroups. Compared to the negative group the time elapsed since diagnosis was significantly longer, additionally, the scores of MIDAS, HIT-6, MSQ, GSRS and SAS were higher for the participants in the IgG positive group. This relationship was also seen with the concentrations of IL-6 and TNF-α, with a higher concentration seen among the IgG positive group. This consistent pattern was observed within the IgG positive group, that is, subgroups with more food allergens generally had worse conditions of migraine (including chronic migraine and the frequency of the migraine attacks), GSRS scores and higher levels of IL-6 and TNF-α.

![Figure 1](https://doi.org/10.2147/JPR.S316619)

**Figure 1** The frequency of the allergenic foods among 67 participants with positive IgG.
Table 1 Comparison of the Participants’ Characteristics Between Different Food Specific IgG Groups

| Characteristics          | IgG Negative (n = 22) | Overall (n = 67) | IgG Positive With 1 Allergen (n = 36) | IgG Positive With 2 Allergens (n = 23) | IgG Positive With ≥3 Allergens (n = 8) | p value
|--------------------------|-----------------------|------------------|--------------------------------------|---------------------------------------|--------------------------------------|----------
| Age                      | 42.7 (10.4)           | 39.6 (8.8)       | 41.3 (9.6)                           | 38.3 (7.6)                            | 35.8 (6.8)                           | 0.19     |
| Gender (%Male)           | 45.5%                 | 28.4%            | 30.6%                                | 34.8%                                 | 0                                    | 0.18     |
| BMI (kg/m²)              | 22.7 (2.8)            | 23.8 (3.9)       | 24.1 (3.7)                           | 23.1 (3.9)                            | 24.6 (4.6)                           | 0.53     |
| With aura                | 4.6%                  | 16.4%            | 8.3%                                 | 34.8%                                 | 0                                    | 0.017    |
| Chronic migraine         | 4.6%                  | 14.9%            | 8.3%                                 | 13.0%                                 | 50%                                  | 0.023    |
| Time elapsed since diagnosis (months) | 61.0 (44.1) | 107.8 (94.4) | 109.7 (113.5) | 107.8 (70.3) | 99.5 (63.6) | 0.96     |
| MIDAS                    | 33.3 (15.6)           | 50.5 (35.7)      | 40.6 (23.1)                          | 57.1 (37.2)                           | 76.3 (59.9)                          | 0.019    |
| Days with migraine in past 12 weeks | 15.5 (17.2) | 22.0 (19.4) | 17.1 (13.2)                           | 22.1 (16.7)                           | 44.0 (33.5)                          | 0.0011   |
| VAS                      | 6.5 (1.5)             | 7.1 (1.3)        | 6.9 (1.5)                            | 7.2 (1.2)                             | 7.5 (1.1)                            | 0.37     |
| HIT-6                    | 57.9 (5.5)            | 61.7 (6.1)       | 59.9 (6.5)                           | 63.1 (4.5)                            | 65.8 (6.0)                           | 0.019    |
| Days with migraine in past 4 weeks | 4.9 (5.3) | 7.5 (6.5) | 6.3 (4.9)                             | 6.9 (5.1)                             | 14.6 (11.5)                          | 0.0030   |
| MSQ                      | 25.2 (11.3)           | 32.5 (12.8)      | 28.6 (11.6)                          | 35.4 (12.8)                           | 41.4 (12.5)                          | 0.013    |
| GRSR                     | 26.9 (6.4)            | 31.6 (11.0)      | 28.3 (8.1)                           | 31.7 (8.9)                            | 45.9 (16.7)                          | <0.0001  |
| SDS                      | 40.1 (10.7)           | 43.5 (11.0)      | 42.6 (9.5)                           | 43.1 (10.7)                           | 48.8 (17.2)                          | 0.36     |
| Depression (%)           | 18.2%                 | 28.4%            | 22.2%                                | 30.4%                                 | 50%                                  | 0.25     |
| SAS                      | 38.8 (7.7)            | 43.2 (10.5)      | 42.0 (8.1)                           | 43.7 (9.3)                            | 47.5 (20.2)                          | 0.39     |
| Anxiety (%)              | 13.6%                 | 23.9%            | 16.7%                                | 34.8%                                 | 25%                                  | 0.33     |
| PSQI                     | 7.4 (3.5)             | 8.0 (3.5)        | 8.1 (3.0)                            | 7.7 (3.5)                             | 8.5 (5.3)                            | 0.84     |
| Poor sleeping (%)        | 45.5%                 | 46.3%            | 50%                                  | 43.5%                                 | 37.5%                                | 0.83     |
| Dermatosis (%)           | 22.7%                 | 37.9%            | 27.8%                                | 63.6%                                 | 12.5%                                | 0.0081   |
| IL-6 (pg/mL)             | 8.6 (3.8)             | 12.6 (7.1)       | 10.0 (5.1)                           | 13.8 (5.2)                            | 20.5 (11.9)                          | 0.0002   |
| IL-10 (ng/L)             | 173.2 (78.3)          | 136.4 (96.0)     | 159.7 (95.2)                         | 121.0 (99.8)                          | 75.9 (15.0)                          | 0.050    |
| TNF-α (pg/mL)            | 12.0 (5.4)            | 20.3 (12.1)      | 15.3 (6.4)                           | 23.5 (11.7)                           | 33.5 (19.5)                          | <0.0001  |

Notes: *Numeric data were described using mean (STD) and categorical data were described by proportion.* **p** values were obtained from ANOVA global test and Fisher’s exact test for the numeric and categorical data, respectively, for comparing the difference among three subgroups with different numbers of food specific allergens. **p** values were from two independent t-tests and Fisher’s exact test for the continuous and categorical data, respectively, for comparing the difference between the IgG positive group vs IgG negative group.

**Abbreviations:** BMI, body mass index; MIDAS, Migraine Disability Assessment questionnaire; VAS, Visual Analog Scale; HIT-6, Headache Impact Test-6; MSQ, Migraine-Specific Quality of Life Questionnaire; GSRS, Gastrointestinal Symptom Rating Scale; SDS, Self-Rating Depression Scale; SAS, Self-Rating Anxiety Scale; PSQI, Pittsburgh Sleep Quality Index; IL-6, interleukin-6; IL-10, interleukin-10; TNF-α, tumor necrosis factor-α.

Characteristics of the IgG Positive Group, and the Association Between Migraine and Comorbidities and the Overall Positive IgG Concentration Before and After Adjusting for the Inflammatory Cytokines

Significant positive associations were observed between the IgG concentration and MIDAS score (p < 0.0001), number of days with a migraine in the previous 12 and 4 weeks (p<0.0001 and p = 0.0001), and GSRS scores (p = 0.0002). With the exception of the GSRS score, after adjusting for the ratio of IL-6 to IL-10, and TNF-α, the strength of the association between the IgG level and each of the outcomes previously mentioned tended to decrease. In other words, the coefficients of the IgG concentration were reduced. Higher IgG was also found to be related to slightly higher odds of having chronic migraine instead of episodic migraine (OR = 1.08, p = 0.0015). This association became non-significant.
after including inflammatory factors in the model, though the OR remained relatively unchanged (Table 2).

**Discussion**

This is the first clinical evidence about food allergy mediated by food specific IgG antibodies of migraine patients in northern China, all 89 participants in this study were from Harbin, which is the capital city with the highest latitude in China. Milk and eggs were found to have the highest IgG antibodies positive rate in the study, these foods are included in the daily diet of local residents. These animal products have strong sensitization, along with being exposed in early life and frequent consumption, leads to the ability to accumulate high levels of serum IgG antibodies in the body and produce chronic antigen antibody reactions.

This study was the first to evaluate the effect of food allergy mediated by food specific IgG antibodies on migraine and its comorbidities by analyzing the numbers of IgG positive foods and overall positive IgG levels. The consumption of sensitized allergenic food triggers specific IgG antibodies that are produced by the immune system; those combined with the allergens form an immune complex, this process could activate multisystem inflammatory reactions, including migraine, depression, anxiety, sleep disorders, irritable bowel syndrome, eosinophilic GI diseases (EGIDs), allergic proctocolitis (AP), urticaria, and eczema.

Some scholars have proposed the potential use of IgG as a biomarker to identify foods linked to an increased inflammatory response, which in turn may be linked to migraine headaches. The hyperinflammatory state is enhanced, manifested by the increased concentration of proinflammatory cytokines, which promotes the activation of the trigeminal nerve. A growing body of evidence has shown that intracranial meningeal neuroinflammation is

| Migraine and Comorbidities | Coefficient (95% CI)* | p value | Adjusting for IL-6, IL-10, and TNF-α | p value |
|----------------------------|----------------------|---------|-----------------------------------|---------|
| Time elapsed since diagnosis (months) | 0.58 (−0.94, 2.09) | 0.45 | 0.36 (<−2.26, 2.99) | 0.78 |
| MIDAS | 1.31 (0.84, 1.78) | <0.0001 | 0.63 (−0.17, 1.42) | 0.12 |
| Days with migraine in past 12 weeks | 0.70 (0.44, 0.96) | <0.0001 | 0.53 (0.14, 0.91) | 0.0083 |
| VAS | 0.002 (<−0.019, 0.023) | 0.85 | 0.011 (<−0.026, 0.048) | 0.55 |
| HIT-6 | 0.084 (<−0.011, 0.18) | 0.083 | 0.0049 (<−0.16, 0.17) | 0.95 |
| Days with migraine in past 4 weeks | 0.19 (0.099, 0.29) | 0.0001 | 0.15 (0.0074, 0.29) | 0.04 |
| MSQ | 0.18 (<−0.020, 0.38) | 0.077 | −0.022 (<−0.36, 0.31) | 0.89 |
| GRSR | 0.31 (0.16, 0.47) | 0.0002 | 0.36 (0.11, 0.62) | 0.0053 |
| SDS | 0.17 (<−0.00061, 0.34) | 0.051 | 0.16 (<−0.13, 0.44) | 0.27 |
| SAS | 0.14 (<−0.024, 0.31) | 0.093 | 0.13 (<−0.12, 0.38) | 0.32 |
| PSQI | −0.0084 (<−0.064, 0.047) | 0.76 | 0.01 (<−0.084, 0.10) | 0.83 |

| OR (95% CI)* | Not Adjusting for IL-6, IL-10, and TNF-α | p value | Adjusting for IL-6, IL-10, and TNF-α | p value |
|--------------|------------------------------------------|---------|-----------------------------------|---------|
| With aura | 1.003 (0.96, 1.05) | 0.91 | 1.00 (0.93, 1.07) | 0.97 |
| Chronic migraine | 1.08 (1.03, 1.13) | 0.0015 | 1.07 (0.99, 1.16) | 0.099 |
| Depression | 1.03 (0.99, 1.06) | 0.11 | 0.99 (0.93, 1.06) | 0.75 |
| Anxiety | 1.01 (0.98, 1.05) | 0.48 | 0.98 (0.91, 1.06) | 0.59 |
| Poor sleeping | 0.98 (0.95, 1.01) | 0.22 | 1.01 (0.95, 1.07) | 0.84 |
| Dermatosis | 1.01 (0.98, 1.05) | 0.42 | 0.99 (0.94, 1.05) | 0.75 |

Note: *Associations were all reported as every 10 u/mL increase in the overall positive IgG level.

Abbreviations: MIDAS, Migraine Disability Assessment questionnaire; VAS, Visual Analog Scale; HIT-6, Headache Impact Test-6; MSQ, Migraine-Specific Quality of Life Questionnaire; GRSR, Gastrointestinal Symptom Rating Scale; SDS, Self-Rating Depression Scale; SAS, Self-Rating Anxiety Scale; PSQI, Pittsburgh Sleep Quality Index; IL-6, interleukin-6; IL-10, interleukin-10; TNF-α, tumor necrosis factor-α.

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a crucial factor leading to the increased sensitivity of trigeminal nociceptors in migraine patients. In this study, migraineurs with an increased number of positive food allergens and/or higher overall positive IgG concentration had poorer disability measurements and greater related functional impairments, quality of life, disability questionnaire results, and slightly greater odds of having chronic migraine, likely related to inflammation triggered by food specific IgG antibodies and a variety of inflammatory factors produced in this process.

We also found that the GSRS score of the IgG positive group was higher than that for the IgG negative group. Moreover, a significant positive association was observed between not only MIDAS scores but also GSRS scores and the positive IgG concentration. After adjusting for the ratio of IL-6 to IL-10, and TNF-α, the coefficients of the positive IgG concentration were reduced. Migraine can often be accompanied by aura of the gastrointestinal (GI) and autonomic nervous symptoms, such as nausea and vomiting. Previous studies on the relationship between migraine and GI disease found that GI disorders have high comorbidity with migraine. Mechanisms explaining how the gut and the brain may interact in patients with migraine may be explained by the gut-brain axis. The interaction between the gut and the brain seems to be influenced by multiple factors such as inflammatory mediators (IL-1β, IL-6, IL-8, and TNF-α), gut microbiota profile, and nutritional substances. A clinical trial in patients with migraine and irritable bowel syndrome also confirms this view. Food specific allergens were identified by measuring the serum IgG antibodies, then an elimination diet was implemented to avoid the identified allergenic foods, removal of these foods resulted in the alleviation of migraine and IBS disease symptoms.

In addition to migraine conditions, anxiety index scores (SAS) in the positive IgG group were significantly higher than that in negative IgG group, along with a higher prevalence of anxiety disorders. Rammohan et al proposed that the association between migraine, psychiatric comorbidities, and sleep disorders might be explained by the migraine neuro-limbic theory, which suggests that the central sensitization process might be responsible for the threshold changes in pain sensibility. Long-term, repeated, severe migraine attacks, anxiety, depression, sleep disorders, and other mental and psychological symptoms are mutually causal and reinforcing, promoting a vicious cycle. Nevertheless, our study found an unanticipated nonsignificant difference in depression and sleep disorder between the IgG positive and negative groups. Mercante et al reported nearly half of their subjects presented with a triple diagnosis of anxiety, primary headaches, and depression. The results found supported the progressive disorder model in which anxiety disorders, often at a young age, proceeds to migraine and is followed by depressive episodes in adulthood. Therefore, we speculate that more promising results may be obtained with longer observation time and more migraine cases.

Additionally, we observed that participants in the IgG positive group were more likely to experience dermatosis than the IgG negative group, though these differences were not statistically significant. The “aseptic inflammation” state of food allergy mediated by IgG may lead to the release of neuropeptides and inflammatory factors, leading to vasodilation, plasma extravasation secondary to capillary leakage, edema, and mast cell degranulation, resulting in allergy-related skin diseases such as urticaria and eczema.

In this study, levels of IL-6 and TNF-α in the IgG positive group were higher than those in the IgG negative group, the association between IgG and MIDAS lost the significance after including the ratio of IL-6 to IL-10 and TNF-α in the model, which indicated that these three inflammatory cytokines might partially mediate the causal pathway between food specific IgG and migraine frequency/intensity. Cytokines play an important role in the inflammation process and mediation of pain sensation of migraine patients. These have been reported to be involved in the modulation of pain threshold, by promoting or suppressing the sensitization of trigeminal nerve fibers. A number of studies have found a pro-inflammatory state in chronic migraine and episodic migraine patients, compared with the healthy controls, which has been demonstrated by an increased concentration of pro-inflammatory cytokines including TNF-α and IL-6. Togha et al also noted that headache is more likely to be chronic with increased levels of inflammatory factors. We hypothesized that increased levels of pro-inflammatory cytokines such as TNF-α and IL-6 are probable indicators of a low grade chronic inflammatory state caused by IgG mediated food allergy.

However, there was no significant difference in the level of IL-10 between the two groups, despite the IgG positive group averaging a lower IL-10. In fact, the level of IL-10 in migraine patients is inconsistent with previously published studies. Perini et al found that the plasma concentration of IL-10 in migraine patients was similar to that in healthy participants, but other research reported lower IL-10, compared to healthy participants. Oliveira et al suggested that...
fluctuating IL-10 levels may be associated with head pain and higher psychiatric comorbidities/symptoms. IL-10 is an effective inhibitor of pro-inflammatory cytokines, which can promote antinociception and prevent depressive-like/anxiety behaviors. In addition, IL-10 is a cytokine produced by helper T2 cells (Th2), and is negatively regulated by TNF-α and IL-12.

Our study had strengths but also limitations. First, participants were enrolled during late fall and winter time. Corresponding low temperatures during this time of year result in differing fresh food sources, food processing, and cooking methods than during the spring and summer months. Moreover, our study found a higher prevalence of depression among participants than other studies, presumably resulting from the potential adverse impact of the fall and winter seasons on depression severity. Second, the sample size for this exploratory study may not be sufficient, potentially compromising the power to detect a statistically significant correlation. The insufficient power was reflected by the lack of significant difference in the prevalence of sleep disorders and dermatosis between the IgG positive and negative groups, while the effect sizes for these two measurements were scientifically meaningful. Third, this study was a cross-sectional study and thus could not establish the direct casualty between diet and migraine. We expect to conduct an interventional study in the future to explore the effect of eliminating IgG positive foods on migraine and the related comorbidities.

Conclusion
This study shows that positive food specific IgG antibodies seem to be related to migraine and its comorbidities, and inflammatory cytokines might partially impact this process. Further studies are warranted to reveal the immunological pathogenesis of migraine and the effect of dietary modification on migraine.

Abbreviations
IgG, immunoglobulin G; BMI, body mass index; MIDAS, Migraine Disability Assessment questionnaire; VAS, Visual Analog Scale; HIT-6, Headache Impact Test-6; MSQ, Migraine-Specific Quality of Life Questionnaire; GSRS, Gastrointestinal Symptom Rating Scale; SDS, Self-Rating Depression Scale; SAS, Self-Rating Anxiety Scale; PSQI, Pittsburgh Sleep Quality Index; IL-6, interleukin-6; IL-10, interleukin-10; TNF-α, tumor necrosis factor-α; ELISA, enzyme linked immunosorbent assay.

Data Sharing Statement
Data can be requested from the corresponding author by E-mail after publication.

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Author Contributions
All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure
The authors report no conflicts of interest in this work.

References
1. Headache Classification Committee of the International Headache Society (IHS). The international classification of headache disorders, 3rd edition. Cephalalgia. 2018;38(1):1–211. doi:10.1177/0333102417738302
2. Rammohan K, Mundayadan SM, Das S, Shaji CV. Migraine and mood disorders: prevalence, clinical correlations and disability. J Neurosci Rural Pract. 2019;10(1):28–33. doi:10.4103/jnrp.jnrp_146_18
3. Talafi Noghani M, Namdar H. Migraine associated with gastrointestinal disorders: a pathophysiological explanation. Med Hypotheses. 2019;125:90–93. doi:10.1016/j.mehy.2019.02.041
4. Han SM, Kim KM, Cho S-J, et al. Prevalence and characteristics of cutaneous allodynia in probable migraine. Sci Rep. 2021;11(1):2467. doi:10.1038/s41598-021-82080-z
5. Merikangas KR. Contributions of epidemiology to our understanding of migraine. Headache. 2013;53(2):230–246. doi:10.1111/head.12038
6. Zaeem Z, Zhou L, Dilli E. Headaches: a review of the role of dietary factors. Curr Neurol Neurosci Rep. 2016;16(11):101. doi:10.1007/s11910-016-0702-1
7. Gazerani P. Migraine and diet. Nutrients. 2020;12(6):1658. doi:10.3390/nu12061658
