Research Article

Clinical Manifestations, Imaging Features, and Pathogenic/Prognostic Risk Factors for Temporomandibular Disorders (TMD): A Case-Control Study Based on Psychogenic Factors of Patients

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Objective. To survey the clinical manifestations and imaging features of temporomandibular disorders (TMD) and analyze the risk factors for pathogenesis/prognosis through a case-control study based on psychogenic factors of patients. Methods. According to the inclusion criteria, 200 adult patients were randomly enrolled from the maxillofacial department of our hospital from January 2020 to May 2021, including 100 patients with TMD as the study group and 100 healthy patients as the control group. The study group can be assigned into four subgroups according to their clinical manifestations: (1) articular area or/and masticatory muscle pain group, (2) mandibular movement abnormality group, (3) joint murmur group, and (4) two or more symptom groups. Based on the study of psychogenic factors of patients, the clinical manifestations and imaging features of TMD were determined, and the risk factors for pathogenesis/prognosis were analyzed. Results. The distribution of psychological status in the TMD group was higher than that in the control group ($P < 0.05$). The distribution of anxiety, depression, and somatic symptoms in the TMD group was significantly different from that in the control group ($P < 0.05$). Anxiety, depression, and somatic symptoms were the risk factors for TMD. Compared with the control group, the incidence of abnormal MRI images in patients with temporomandibular disorders was significantly different from that in the control group ($P < 0.05$). Anxiety, depression, and somatic symptoms were the risk factors for TMD. Depression was the risk factor for pain ($P < 0.05$). Conclusion. In patients with TMD, MRI can early identify disc abnormalities and other related imaging features, which is helpful for more comprehensive clinical evaluation and treatment of TMD patients. There exhibits no significant difference in psychological status (anxiety, depression, and somatic symptoms) of patients with different clinical symptoms, and abnormal psychological status may be one of the risk factors leading to different clinical symptoms and development of different types of TMD patients.

1. Introduction

Temporomandibular disorders (TMD) describe a group of diseases with the same clinical symptoms, but its etiology remains unknown [1]. Its main clinical symptoms include pain in the temporomandibular joint and/or masticatory muscles, joint ringing, murmur and fragmentation, and abnormal mandibular movement [2]. Epidemiological studies demonstrate that the prevalence of TMD is about 40%, and the annual incidence rate is about 3.9%. Female patients are more common than male patients [3]. TMD occurs in all age groups, especially in young adults, and the highest
prevalence rate is between 20 and 40 years old [4]. Muscle pain and articular disc displacement are common in adolescents, while degenerative osteoarthropathy has a high incidence in middle-aged and elderly people [5]. MRI is regarded as the first choice because of its nonradiation and high soft tissue resolution. It can clearly display the joint structure and especially can accurately judge the shape and position of the articular disc, but it has been reported that articular disc displacement can also occur in normal people, so it is necessary to observe other MRI changes, including joint effusion, thickening of the attachment point of the lateral pterygoid muscle, and abnormal condyle [6]. Traditional X-ray and CT examination can well demonstrate the changes of joint bone structure, but it is not good for showing intra-articular soft tissue and joint cavity effusion. Arthrography is an invasive examination, and the scope of clinical application is limited. MRI can clearly demonstrate the bony structure, articular disc, and soft tissue around the joint and can be used at any level, so it is regarded as the gold standard of TMD imaging. In addition to conventional static imaging, MRI can more directly indicate the motion status of the joint, but dynamic imaging is prone to “jumping” phenomenon or motion artifacts, which needs to be observed in combination with static imaging [7].

The etiology of TMD is complex, and the pathogenesis of TMD is still unclear [8]. At present, it is considered to be a disease caused by a variety of pathogenic factors. The main pathogenic factors include social psychology, occlusion, joint anatomy, autoimmunity, genetic factors, environmental factors, and so on [9]. Many scholars believe that psychological factors are strongly related to the occurrence and development of TMD [10]. In addition, whether to regulate psychological factors plays an important role in the treatment of TMD. Mun et al. found that under the same environmental conditions, patients with temporomandibular disorder syndrome were more likely to express severe depression, anxiety, and other abnormal psychological conditions than those without temporomandibular disorder syndrome, indicating that there was a significant correlation between abnormal psychological status and TMD [11]. Owens et al. compared the results of occlusal plate therapy in 80 patients with chronic TMD and found that the therapeutic effect of the occlusal plate on patients with poor psychological status such as anxiety and depression is often not good compared with those with normal psychological status [12]. More scholars agree with the view of TMD from the perspective of bio-psycho-social model [13]. Psychological factors include the influence of individual, crowd, and environmental changes on patients’ adaptability. Because patients have great differences in cultural, economic, and social background, the related psychological factors also demonstrate diversity [14]. The study demonstrates that patients are difficult to adapt to the surrounding environment in terms of personality characteristics and emotional state and are in a state of tension, depression, and anxiety for a long time. Human is a more complex individual, an organ is also affected by other factors of the whole body, and it is difficult to separate simple psychological factors [15]. Many clinical studies start with psychological questionnaires and find that there is a correlation between psychological factors and occurrence. At the same time, the investigation of the mental and psychological status of the patients did find the patients with mental disorders. Some animal experiments have found that the ultrastructure and metabolism of temporomandibular joint change under psychological stress [16]. However, the research on mental and psychological factors is still less and not deep enough, and its influence mechanism is not clear, and it is difficult to treat frequently occurring diseases in clinic. Based on this, this study explores the clinical manifestations and imaging features of TMD and analyzes the risk factors for pathogenesis/prognosis according to the case-control study of psychogenic factors of patients. The results are reported as follows.

2. Patients and Methods

2.1. General Information. According to the inclusion criteria of the experiment, 200 adult patients were randomly enrolled from the maxillofacial department of our hospital from January 2020 to May 2021, including 100 patients with TMD as the study group and 100 healthy patients as the control group. The patients were examined by general data, GAD-7, PHQ-9, and PHQ-15 self-rating psychological scale, and clinical and imaging examination of the temporomandibular joint (CBCT or MRI). There were 46 males and 54 females with an average age of 25.31 ± 5.44 years in the study group and 43 males and 57 females with an average age of 27.32 ± 4.33 years in the control group. With the permission of the Medical Ethics Association of our hospital, all patients signed the informed consent form. The research group can be divided into four groups according to clinical manifestations: (1) articular area or/and masticatory muscle pain group, (2) mandibular movement abnormality group, (3) joint murmur group, and (4) two or more symptom groups.

Diagnostic criteria were as follows: using the study diagnostic criteria of TMD (DC/TMD) issued by the International Dental Federation in 2014, the patients were assigned into the control group and study group according to the diagnostic roadmap in axis I (Figure 1) combined with the clinical symptoms and imaging manifestations of the temporomandibular joint.

Selection criteria were as follows: (1) first visit adults over 18 years old; (2) completing screening scale independently; (3) cooperating with temporomandibular joint function examination and imaging examination; (4) willing to participate in this study with informed consent; (5) no obvious occlusal interference, tumor, dysplasia, and other factors; and (6) no history of orthodontic or orthodontic treatment.

Exclusion criteria were as follows: (1) previous personality and mental retardation, (2) failure to meet the standard due to serious systemic or local disorders, (3) unwillingness or inability to participate in this research, and (4) patients with multiple tooth loss and occlusal factors after orthodontic treatment.

2.2. MRI Inspection Method. All patients were informed before MRI and accepted and signed the informed consent form of MRI scan. IDEALIQ software was used in the study
group and the control group, and the scanning sequence and parameters of the two groups were the same. Single channel facet joint coil (loop array) was employed to scan the routine sequence and T2 mapping sequence of TMD using Discovery MR 750 3.0 T (General Electric Company, Boston, US). A special mouth opener should be employed when scanning the sagittal position of mouth opening. In order to ensure the safety of patients and prevent nosocomial infection, a disposable mouth opener should be employed during examination. Patients or volunteers should be informed of the examination instructions before the scan to ensure that there are no contraindications to MRI. Remove all metal magnetic objects before magnetic resonance examination, remove dentures, wear special earplugs for magnetic resonance examination to reduce noise and protect the patient’s hearing, fix the patient’s head with a sponge, and instruct the patient to stay on the scanning bed. And tell the patient to stay still during the scan. When entering the bed, the head is advanced, the loop coil is placed on the affected side of the collar face, close to the temporomandibular joint area and fixed properly, and the scan is also located in the temporomandibular joint area or in the coil. After positioning, it is sent to the examination bed to wait for scanning. T2WI and PDWI sequences in oblique sagittal position and PDWI sequence in oblique coronal position of bilateral temporomandibular joint opening and closing were collected in all cases. The sagittal plane was perpendicular to the long axis of the internal and external diameter of the condyle, and the coronal plane was parallel to the long axis of the internal and external diameter of the condyle.

2.3. Observation Index

2.3.1. Psychological Factor Scale and Diagnostic Criteria. The psychogenic factors of the patients were evaluated by using DC/TMD axis II self-rating scales GAD-7, PHQ-9, and PHQ-15. Generalized Anxiety Disorder 7 (GAD-7) is used to assess the patient’s anxiety state. There are 7 items. According to patient selection, each item is scored 0-3 points, with a total score of 5, 10, and 15 points. The anxiety state of patients was divided into four grades: no anxiety, mild anxiety, moderate anxiety, and severe anxiety. The patient’s depression status was assessed using the Patient Health Questionnaire-9 (PHQ-9), with a total of 9 items. Each item is scored on a scale of 0-3, depending on the patient’s choice. According to the total score of 5, 10, and 20, the degree of depression of the patients was divided into 5 grades: nondepression, mild depression, moderate hesitation, moderate to severe depression, and severe depression. The severity of physical symptoms was assessed by the Patient Health Questionnaire-15 (PHQ-15). There are 15 questions in total, with 0-3 points for each question. According to the total score of 5, 10, and 15 points, the physical
condition of the patients was divided into four grades: asymptomatic, mild, moderate, and severe.

2.3.2. MRI Image Processing. All the scanned images were analyzed and processed on the IDEAL IQ. The T2-star-p2-anatomical anatomical images accurately located the articular disc on the T2-image map. The anterior zone and posterior zone of the Guanye plate were enrolled, respectively, and their T2 values were measured. The points should be taken as far as possible to avoid inflammation, effusion, and other areas, and the points should be enrolled many times to take the average value. Two experienced diagnostic doctors employed the double-blind method to postprocess and analyze the images of each sequence of MRI in all the study groups and the normal control group. The signal strength of the articular disc is the average signal degree measured in the region of interest (ROI). The best plane of the articular disc is enrolled, and the point surface is 1 mm², and the T2 values of the anterior and posterior bands of each articular disc are measured, respectively. In order to reduce the manual measurement error as much as possible, two experienced doctors entered the jack point measurement, and each group of data was measured for 3 times and then the average value was calculated. The scanned images were analyzed and counted by two experienced doctors using the double-blind method. The method. The image observation indexes of the temporomandibular joint in patients with TMD include whether the position of articular disc is displaced or abnormal, whether there is edema in bone marrow, whether there is adhesion of the joint, whether there is effusion in the articular cavity, and so on.

2.4. Statistical Analysis. The data collected from each group were statistically analyzed by SPSS 23.0 software. The scores of GAD-7, PHQ-9, and PHQ-15 in the study group and the control group were statistically analyzed, and the differences of psychological severity and distribution were compared. With or without TMD as dependent variables, logistic regression analysis of anxiety, depression, and somatic symptoms was carried out. $P < 0.05$ shows that the difference is statistically significant.

3. Results

3.1. Comparison of the Distribution of Patients according to Anxiety, Depression, and Somatic Symptoms. The distribution of patients based on the level of anxiety in the study group is as follows: normal in 15 cases, mild in 64 cases, moderate in 20 cases, and severe in 1 case. According to the degree of depression, the distribution of patients was displayed: normal in 28 cases, mild in 58 cases, moderate in 28 cases, and severe in 7 cases. Using the severity of somatic symptoms, the distribution of cases was shown: normal in 5 cases, mild in 49 cases, moderate in 39 cases, and severe in 7 cases. On the other hand, the distribution of cases in the control group was exhibited: normal in 48 cases, mild in 51 cases, moderate in 1 case, and severe in 0 case according to the level of anxiety. Based on the level of depression, the distribution of cases in the control group was displayed: normal in 54 cases, mild in 45 cases, moderate in 34 cases, and severe in 0 case. By the severity of somatic symptoms, the distribution of cases was displayed: normal in 65 cases, mild in 34 cases, moderate in 1 case, and severe in 0 case. The distribution of the cases via physical symptoms in the control group was as follows: normal $(n=48)$, mild anxiety $(n=51)$, moderate anxiety $(n=1)$, and severe anxiety $(n=0)$. In terms of the control group, the distribution of psychological status in the study group was higher compared to that in control group, and the difference is statistically significant ($P < 0.05$). The results indicated that there exhibited statistical differences in the distribution of anxiety, depression, and somatic symptoms between the study group and the control group. All the data results are indicated in Table 1.

3.2. Multivariate Analysis of Anxiety, Depression, Somatic Symptoms, and TMD. The occurrence of TMD was taken as the dependent variable, anxiety, depression, and somatic symptoms were taken as independent variables into a logistic regression equation, and statistical analysis was carried out. Multiple logistic regression analysis indicated that anxiety, depression, and somatic symptoms were statistically significantly risk factors for TMD, and the difference is statistically significant ($P < 0.05$). All the data results are indicated in Table 2.

3.3. Imaging Features of TMD. Among the patients in the study group, 9 patients indicated bilateral TMD disorder, 73 patients indicated unilateral TMD disorder, and 126 discs in 200 joints were abnormal accounting for about 63%, of which 54 were reducible anterior displacement (Figure 2); 58, irreducible anterior displacement (Figure 3); 6, irreducible anterolateral displacement; and 6, articular disc adhesion. The shape of 48 articular discs was abnormal, accounting for about 48%. Among them, 15 were enlarged posterior band, 8 were elongated, 5 were folded, and 2 were biconvex; of the 100 patients, 47 joints indicated effusion in 84 patients (Figure 2), thickening of the attachment point of the inferior head of the lateral pterygoid muscle in 9 cases (Figure 3), condylar bone degeneration in 22 of the 62 joints, flatness of the top of the condyle in 7 cases, and hyperostogeny in 15 cases (Figure 4).

3.4. Clinical Manifestation of TMD. The correlation between clinical symptoms and MRI findings is indicated in Table 3. Compared with the control group, the incidence of abnormal images in MRI of temporomandibular joint disorders is significantly different ($P < 0.05$). All the data results are indicated in Table 3.

3.5. Differences of Psychological Status among Different Clinical Symptoms of TMD. There exhibited no significant difference in psychological status (anxiety, depression, and somatic symptoms) among the four groups, and the difference is statistically significant ($P < 0.05$). All the data results are indicated in Figure 5.

3.6. Differences in the Distribution of Psychological Status among Different Clinical Symptoms of TMD. There were differences in the distribution of psychological status (anxiety,
depression, and somatic symptoms) among the four groups; the results indicated that $P > 0.05$, indicating that there exhibited no significant difference in the distribution of anxiety, depression, and somatic symptoms among the four groups. All the data results are indicated in Figures 6(a) and 6(b).

### Table 1: Comparison of the distribution of anxiety, depression, and somatic symptoms between the two groups (n%).

| Group          | R group (n = 100) | C group (n = 100) | $\chi^2$ | $P$   |
|---------------|------------------|------------------|----------|-------|
| Anxiety       |                  |                  |          |       |
| Normal        | 15 (15.00)       | 48 (48.00)       |          |       |
| Mild          | 64 (64.00)       | 51 (51.00)       |          |       |
| Moderate      | 20 (20.00)       | 1 (1.00)         | 36.945   | <0.01 |
| Severe        | 1 (1.00)         | 0 (0)            |          |       |
| Melancholy    |                  |                  |          |       |
| Normal        | 7 (7.00)         | 54 (54.00)       |          |       |
| Mild          | 58 (58.00)       | 45 (45.00)       |          |       |
| Moderate      | 28 (28.00)       | 1 (1.00)         | 69.992   | <0.01 |
| Moderate to severe | 7 (7.00) | 0 (0)           |          |       |
| Somatic symptoms |                |                  |          |       |
| Normal        | 5 (5.00)         | 65 (65.00)       |          |       |
| Mild          | 49 (49.00)       | 34 (34.00)       |          |       |
| Moderate      | 39 (39.00)       | 1 (1.00)         | 97.240   | <0.01 |
| Severe        | 7 (7.00)         | 0 (0)            |          |       |

### Table 2: Multivariate analysis of anxiety, depression, somatic symptoms, and TMD.

| TMD risk factors | $b$   | SE    | $\chi^2$ | $P$   | OR   | 95% CI for OR |
|------------------|-------|-------|----------|-------|------|---------------|
| Anxiety          | 0.078 | 0.013 | 36.000   | <0.01 | 1.081| 1.054-1.109   |
| Melancholy       | 0.129 | 0.054 | 5.707    | <0.01 | 1.138| 1.023-1.265   |
| Somatic symptoms | 0.134 | 0.044 | 9.275    | <0.01 | 1.143| 1.049-1.246   |

**Figure 2:** The reducible articular disc moves forward, the closed position in (a) demonstrates that the articular disc moves forward, the demarcation angle of the disc is about 30°, and the articular disc is enlarged in the posterior zone. The opening position in (b) demonstrates that the relationship between the articular disc and the condyle returns to normal.

3.7. **Multivariate Analysis of Anxiety, Depression, Somatic Symptoms, and Different Clinical Symptoms of TMD.** Different clinical symptoms of TMD were taken as dependent variables, nonclinical symptoms were taken as reference categories, anxiety, depression, and somatic symptoms were
taken as independent variables into logistic regression equation, and statistical analysis was carried out. It can be noticed that anxiety, depression, and somatic symptoms are the risk factors for abnormal mandibular movement, joint ringing, or murmur; somatic symptoms are the risk factors for all kinds of clinical symptoms of TMD, and depression is the risk factor for pain. All the data results are indicated in Table 4.

4. Discussion

TMD is a common disease, which is more common in young and middle-aged women [15]. The pathogenesis is not clear. It is generally considered to be related to psychological factor, trauma, anatomical abnormalities and joint overload, and other factors. Most patients have multiple pathogenic factors [16]. With the development of living standards and
the continuous improvement of medical standards, more patients seek medical treatment. The main clinical history of this disease is mainly manifested as local muscle dilatation or pain in the temporomandibular joint area and varying degrees of mouth opening or mouth opening limitation. In addition, a small number of patients can also be accompanied by temporal tenderness, dizziness, tinnitus, and other symptoms. The etiology is complex, and the pathogenesis is not fully understood, which may be related to trauma (such as external force impact, biting hard objects, and excessive mouth opening), occlusal disorders, or mental factors [16, 17]. At present, the imaging methods of TMD include X-ray, CT, MRI, arthroscopy, curved tomography, and so on. X-ray can only reflect obvious bone destruction, while most patients have irreversible bone destruction in the temporomandibular joint. The main deficiency of ultrasound is that it cannot demonstrate the articular disc and surrounding soft tissue very clearly. When there is articular cavity effusion or slight forward movement of the articular disc in the temporomandibular joint and the articular disc is displaced to the medial or lateral region, ultrasound is easy to demonstrate blurred images. And ultrasound imaging ability of bone tissue is not good, so it cannot make a clear and effective diagnosis and can even lead to misdiagnosis or missed diagnosis [17]. Although CT can clearly demonstrate the bone structure and articular cartilage, a series of pathological changes such as edema of lateral pterygoid muscle, bone marrow edema of condyle, degeneration of the articular disc, injury, and displacement cannot be clearly demonstrated. Arthrography is the main means of examination of the temporomandibular joint before the appearance of MRI. Arthrography can demonstrate whether there are obvious abnormalities in the joint space and the changes of the joint bone, in order to further check whether there are space-occupying lesions in the joint. In addition, arthrography can also be combined with X-ray to dynamically observe the movement relationship between the articular disc and naked process, but arthrography is an invasive examination. This test should not be performed in patients with bleeding disorders, patients taking anticoagulants, patients with iodine allergy, and patients with localized joint skin infections. However, MRI has high soft tissue resolution and can well display the lesions of the temporomandibular joint and surrounding soft tissue in recent years [18]. According to related studies, in the early stage of TMD, the articular disc has varying degrees of fiber abnormality, articular disc shape deformation, thick and even fiber fracture, articular disc collagen fiber fracture loss of continuity, and increased double-plate fibrosis. The cartilage layer of the condyle became thinner, or chondrocytes decreased. Early detection and early diagnosis and treatment can control or delay the deterioration of the disease, so early diagnosis is particularly important for the treatment and prognosis of patients.

With the continuous development of basic research and other related disciplines, there has been a further understanding of the etiology, outcome, and treatment of this kind of disease; in particular, great progress has been made in etiology [19]. At the sixth symposium on temporomandibular joint diseases and joint studies in 1997, Professor Ma Xichen put forward the necessity of biaxial diagnosis and new diagnostic criteria in China as soon as possible. That is, axis somatic disease assessment and axis-related pain-related loss of function and psychological status emphasized the role of mental and psychological factors [19, 20]. With the changing medical paradigm, the influence of psychiatric and psychosocial factors on TMJ disorders has received increasing attention. Some scholars believe that there is a certain correlation between the occurrence of TMD and physiological

### Table 3: Clinical manifestations of TMD.

|                          | C group (n = 100) | R group (n = 100) |
|--------------------------|-------------------|------------------|
| Reducible disc displacement | 14 (14.00)       | 35 (35.00)       |
| Irreducible disc displacement | 5 (5.00)       | 48 (48.00)       |
| Joint deformation         | 5 (5.00)         | 73 (73.00)       |
| Effusion in articular cavity | 19 (19.00)     | 88 (88.00)       |
| Thickening of the attachment point of the inferior branch of lateral pterygoid muscle | 0 | 22 (22.00) |
| Condylar bone degeneration | 5 (5.00)       | 52 (52.00)       |

**Figure 5:** Psychological differences among different clinical symptoms of temporomandibular joint disorder syndrome.

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**Table 3:** Clinical manifestations of TMD.
factors through clinical research. With the increasingly fierce competition in today’s society, the accelerating pace of life, and the increasing pressure of work, people are more likely to feel anxious, which in turn can easily lead to a series of stress reactions. According to the research report of foreign scholars, psychological changes such as emotional disorders, as a kind of psychological stimulation, can affect the reticular structure of the cerebral cortex, brainstem, and hypothalamus through nerves and stimulate the adrenal gland to release catechins [20]. Then, it causes a large amount of epinephrine and norepinephrine secretion, leading to muscle contraction and muscle spasm, which can easily lead to the disorder of temporomandibular joint structure and joint displacement and further cause occlusal abnormalities, joint deformation, effusion, and other secondary changes [21]. At present, it is believed that rational factors have a certain influence on the incidence of TMD, so it is suggested that stomatologists should pay attention not only to traditional treatment and surgical treatment but also to patients’ psychotherapy in the clinical diagnosis and treatment of TMD.

Previous studies have proven that TMD is a kind of disease with complex etiology, psychological factors are

![Differences in psychological distribution among different clinical symptoms of temporomandibular joint disorder syndrome](image-url)

**Figure 6:** Differences in psychological distribution among different clinical symptoms of temporomandibular joint disorder syndrome. Note: group 1: abnormal mandibular movement; group 2: articular bounce or tenderness; group 3: pain; and group 4: two or more symptoms.
The results indicated that subjects with depressive symptoms, compared to that in normal people [26]. Byun et al. conducted a prospective cohort study of more than 3000 ordinary people without temporomandibular disorder syndrome. The results indicated that subjects with depressive symptoms had a significantly increased risk of temporomandibular joint pain, and anxiety symptoms were a significant risk factor for joint and muscle pain [27]. Meanwhile, some studies have indicated that the abnormal psychological status of patients with TMD may be related to a series of clinical symptoms such as pain and abnormal mandibular movement caused by TMD [28]. These uncomfortable clinical symptoms may lead to long-term anxiety and depression. Therefore, it is necessary to take painkillers to relieve pain in patients with TMD. The abnormal psychological status of some patients can be significantly improved [29]. Badel et al. suggested that nonsteroidal anti-inflammatory drugs (NSAIDs) for patients with joint pain and diazepam for muscle pain patients can not only effectively relieve pain but also play a positive role in improving the psychological status of TMD patients and the outcome of TMD. In addition, further studies have indicated that some genes that lead to abnormal psychological status are more common in patients with temporomandibular disorder syndrome, proving that these people are susceptible [30]. Studies have indicated that the catechol-O-methyltransferase (COMT) gene may be associated with abnormal psychogenic factors such as anxiety and depression in patients with chronic temporomandibular joint disease. Long-term follow-up studies have found that patients with the COMT genotype have a 2-3 times higher risk of developing TMD than normal people [31]. The above results suggest that clinicians should pay close attention to their psychological status when receiving and treating patients with TMD and refer them to a professional psychologist if necessary.

In conclusion, MRI can identify disc abnormalities and other related imaging features of TMD patients early, which is helpful for more comprehensive clinical evaluation and treatment of TMD patients. There was no significant difference in the psychological state (anxiety, depression, and somatic symptoms) of patients with different clinical symptoms, and an abnormal psychological state may be one of the risk factors for the different clinical symptoms and development of different types of TMD in patients.

### Table 4: Multivariate analysis of anxiety, depression, somatic symptoms, and different clinical symptoms of TMD.

| Different clinical symptoms of TMD | TMD risk factors | b    | SE   | \( \chi^2 \) | P    | OR   | 95% CI for OR |
|-----------------------------------|-----------------|------|------|-------------|------|------|---------------|
| Abnormal mandibular movement      | Anxiety         | 0.156| 0.053| 8.664       | <0.01| 1.169| 1.054-1.297   |
|                                   | Melancholy      | 0.093| 0.033| 7.942       | <0.01| 1.097| 1.029-1.171   |
|                                   | Somatic symptoms| 0.089| 0.042| 4.490       | <0.01| 1.093| 1.007-1.187   |
| A ringing or murmur in the articular area | Anxiety   | 0.083| 0.021| 15.621      | <0.01| 1.087| 1.043-1.132   |
|                                   | Melancholy      | 0.259| 0.042| 38.028      | <0.01| 1.296| 1.193-1.407   |
|                                   | Somatic symptoms| 0.154| 0.035| 19.360      | <0.01| 1.166| 1.089-1.249   |
| Pain                              | Anxiety         | -0.053| 0.046| 1.328       | >0.05| 0.948| 0.867-1.038   |
|                                   | Melancholy      | 0.423| 0.057| 55.072      | <0.01| 1.527| 1.365-1.707   |
|                                   | Somatic symptoms| -0.036| 0.012| 9.000       | <0.01| 0.965| 0.942-0.988   |
| Have two or more symptoms         | Anxiety         | 0.058| 0.032| 3.285       | >0.05| 1.060| 0.995-1.128   |
|                                   | Melancholy      | 0.044| 0.031| 2.015       | >0.05| 1.045| 0.983-1.110   |
|                                   | Somatic symptoms| 0.294| 0.042| 49.000      | <0.01| 1.342| 1.236-1.457   |
Data Availability

No data were used to support this study.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

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