Sexual Dysfunction in Chronically Medicated Male Inpatients With Schizophrenia: Prevalence, Risk Factors, Clinical Manifestations, and Response to Sexual Arousal

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Background: Sexual dysfunction is a common symptom in patients with schizophrenia, especially in chronically medicated patients. However, the relationship between sexual dysfunction and emotional response to sexual arousal in male patients with schizophrenia remains unclear. This study aimed to assess the incidence, risk factors of sexual dysfunction in males, and their clinical correlations to sexual arousal in male patients with schizophrenia in China.

Methods: A total of 162 male patients, aged 18–50 years, with schizophrenia were recruited from a psychiatric hospital in Ganzhou. The clinical symptoms were assessed by the Positive and Negative Syndrome Scale (PANSS). The Arizona Sexual Experience Scale was utilized to evaluate sexual dysfunction. Erotic images were selected from International Affective Picture System (IAPS). Sixty-eight out of the 162 subjects completed the erotic pictures reactivity task.

Results: Overall, 48 (29.6%) patients were measured as having global sexual dysfunction, 72 (44.4%) patients as having strength of sex drive dysfunction, 51 (31.5%) patients as having sexual arousal dysfunction, 55 (34.0%) patients as having penile erection dysfunction, 60 (37.0%) patients as having reached orgasm dysfunction, and 60 (37.0%) patients as having satisfaction with orgasm dysfunction. The sexual dysfunction patients had significantly higher scores on the negative symptoms of the PANSS. The only important predictor of sexual dysfunction was the severity of PANSS negative factor. The sense of pleasure and arousal post viewing erotic images in the sexual dysfunction group were lower compared to the non-sexual dysfunction group. The sense of pleasure and approach motivation were significantly negatively correlated with the severity of sexual dysfunction.
Conclusions: This study shows that nearly one-third of young and middle-aged chronically medicated male inpatients with schizophrenia suffer from sexual dysfunction. The negative factor of the PANSS can be regarded as the risk factor of sexual dysfunction. Schizophrenia patients with sexual dysfunction experienced lower pleasure and higher avoidance motivation than non-sexual dysfunction patients when exposed to erotic stimuli.

Keywords: schizophrenia, sexual dysfunction, pleasure, arousal, motivation

INTRODUCTION

Sexual dysfunction is a serious side effect that commonly manifests in patients with chronic schizophrenia (1–3). Previous studies have shown that the prevalence of global sexual dysfunction in schizophrenia patients ranged from approximately 16% to 96% (4–6). Generally, the illness itself and antipsychotic drugs both can lead to sexual dysfunction in schizophrenia patients (3, 5, 7). Moreover, sexual dysfunction has a negative impact on treatment adherence and intimate relationships, especially in males (3, 6, 8, 9). In addition, due to decreased libido and erectile dysfunction, most males with chronic schizophrenia are accompanied by low self-esteem (10, 11) and depression (3, 12), which reduce their motivation to seek intimacy and damage their marital relationships (13). Therefore, the rate of unmarried patients with schizophrenia is significantly higher when compared to subjects without the disease (14).

A variety of impacting factors, such as sex (14–17), age (12, 14, 18), marital status (14), and smoking status (5, 19, 20), have all been reported to be related to sexual dysfunction in patients with schizophrenia. Moreover, other risk factors for sexual dysfunction include age of onset, duration of illness (15, 17), psychopathology, and prolactin levels (21–24). In addition, antipsychotic medication is a specific and severe risk factor for increased contingency of sexual dysfunction. Some previous studies have supported that long-term use of antipsychotics, such as clozapine, risperidone, and olanzapine, contributed to drug-induced sexual dysfunction (1–3, 5, 7, 14, 17, 25). Although the results regarding sexual dysfunction due to different antipsychotics were inconsistent (7, 17), sexual dysfunction is still regarded as a serious side effect. Some studies have shown that sexual dysfunction is correlated with the severity of clinical symptoms. For example, Huang et al. (6) reported that in a sample of 742 schizophrenia patients, sexual dysfunction was related to the Brief Psychotic Rating Scale (negative domain) score. Similarly, the severity of sexual dysfunction was found to be significantly associated with the negative subscale of the Positive and Negative Syndrome Scale (PANSS) (26). Although most previous studies have examined the association between comorbid sexual dysfunction, demographic variables, and psychopathological symptoms, they did not explore the correlation between sexual dysfunction and emotional response to erotic stimuli (12, 15, 17, 27). Deficits in emotional and motivational function are well-known in patients with schizophrenia, including flat affect, an impaired ability to experience pleasure, and diminished motivation (28). Generally, images depicting erotic contents can evoke a robust motivational state (29). When presenting erotic images, the judgment of pleasure and arousal in individuals with schizophrenia is similar to healthy individuals (30–32). However, the degree of brain activation and desire motivation elicited by erotic stimuli in schizophrenia patients were significantly lower. Heerey and Gold (33) showed that the basic motivational process in patients with schizophrenia may be impaired. However, whether sexual dysfunction is related to the impairment of emotional and motivational systems in schizophrenia patients remains unclear.

The purposes of our study are to explore: (1) the incidence, risk factors, and clinical correlates of sexual dysfunction in Chinese male inpatients with schizophrenia; (2) response to sexual arousal; and (3) the correlation between sexual dysfunction and emotional response to sexual stimuli.

METHODS

Subjects and Settings

The subjects in the study were inpatients recruited from the Third People’s Hospital in Ganzhou City, China. The survey was conducted from August 1, 2018, to July 31, 2019. All subjects met the criteria, as follows: (1) aged 18 years or above, Han Chinese; (2) self-report of heterosexuality; (3) meeting The Diagnostic and Statistical Manual of Mental Disorders IV, Text Revision (DSM-IV-TR) (2000) criteria for schizophrenia, two psychiatrists independently evaluated the mental status of the subjects based on the Structured Clinical Interview for DSM-IV-TR (SCID); (4) adherence to oral second-generation antipsychotics for at least 1 year; (5) be able to understand the requirements and content of the evaluation; (6) no other somatic diseases, such as diseases of the central nervous system, acute, unstable, or life-threatening diseases (e.g., cancer, infection); (7) normal or corrected-to-normal vision and normal color vision. The sociodemographic and clinical information of the subjects was collected by experienced researchers. Other pieces of information were collected from electronic medical records and ancillary resources. All the subjects underwent physical examination and blood biochemical tests. In total, 162 male patients were recruited in this study. All subjects had no addiction to drugs, alcohol, or other substances.

Ethics Approval and Consent to Participate

The project was approved by the Ethics Committee of the Third People’s Hospital of Ganzhou City (2018002). A trained
researcher explained the purpose of the study, its procedures, and confidentiality of the information to the subjects in a language appropriate to their level of understanding and emotional state. Participants were also informed that the study was only a cross-sectional survey, that there would be no harm to them, and that they would not obtain any intervention. Each subject signed an informed consent form prior to participation in the study.

All methods described in this study were carried out in accordance with the Declaration of Helsinki promulgated by the National Institutes of Health.

**Clinical Measurement**

The PANSS was used by two researchers who received training on how to assess PANSS. The interrater correlation coefficient of repeated measurements for the total score of PANSS was >0.8. The five-factor model of PANSS was used to evaluate the clinical symptoms of the subjects (34). Items P1, P3, P5, and G9 composed positive factor; items N1, N2, N3, N4, N6, and G7 composed negative factor; items G2, G3, and G6 composed depressed factor, which was applied to assess the depressive symptoms; items P2, N5, and G11 composed concrete/disorganized factor; and items P4, P7, G8, and G14 composed excited factor.

**Measurement of Anthropometric Variables**

Sociodemographic information included age, age of onset, education levels, marital status at the time of consultation, and family history of mental illness. Clinical data such as weight, height, smoking status, duration of illness, hospitalization numbers, and the use of antipsychotic and antidepressant medications were collected. Trained research assistants were responsible for collecting the sociodemographic and clinical data.

**Sexual Dysfunction**

The Mandarin Chinese version of the Arizona Sexual Experience Scale (ASEX) (35–37) was used to assess the sexual functioning of each subject with schizophrenia. The Mandarin Chinese version of ASEX has good reliability and validity in China and has been used in patients with schizophrenia (24, 35, 36, 38, 39). ASEX assessed the status of sexual function in the last week. Five aspects of sexual functioning composed of strength of sexual drive, sexual arousal, penile erection, ability to reach orgasm, and satisfaction with orgasm were measured; each category with a score of Likert 1–6. Total score ranges from 5 to 30, with higher scores indicating much more serious sexual dysfunction. A total score ≥19, any single item score ≥5, or any three item scores ≥4 are defined as the presence of clinical sexual dysfunction.

**Erotic Pictures Reactivity Task**

Pictures for the affective reactivity task included 10 erotic images of nude couples in erotic poses and 10 neutral images of dressed people in daily life. The pictures were all selected from the International Affective Picture System (IAPS) and have been used in previous studies (40–42). According to the normative rating program provided by Bradley et al. (29) in the IAPS manual, E-prime was used to edit the evaluation program. After providing instruction consent, the 20 pictures were presented to participants in a random order. The participants watched each image for 6 s, and the subjects were asked to rate pleasure, arousal, and approach motivation at the same time through a computer. Participants rated three dimensions with self-report in a 9-point rating scale. The pleasure dimension is rated as 1 indicating “extreme disgust,” 5 indicating “neutral,” and 9 indicating “extreme pleasure.” Similarly, in the arousal dimension, 1 indicates “not at all arousing,” 5 indicates “moderately arousing,” and 9 indicates “extremely arousing.” Approach motivation evaluated the degree to which subjects want to get involved in the scene or approaching the scene after viewing pictures, with 1 indicating “extremely escaping,” 5 indicating “neutral,” and 9 indicating “extremely approaching.” Before the formal experiment, participants were given practice lists in order to understand the scoring standard. During the formal experiment, the subject was left alone in a private room.

**Data Analysis**

The G*Power 3.1.9.2 program was used to perform a sample size based on effect size derived from a previous study (26). The sociodemographic and clinical variables of sexual dysfunction (SD) and non-sexual dysfunction (NSD) groups were compared. The continuous variables were tested by independent-samples t-test, and the categorical variables were compared by chi-square test. The prevalence of global sexual dysfunction and each factor were analyzed by chi-square test. ANCOVA was performed to control the influence of age, education years, age of onset, hospitalization numbers, and family history of mental illness. Correlations of sexual dysfunction and sociodemographic and clinical variables were conducted by Pearson correlation coefficients. Bonferroni corrections were used to adjust for multiple tests. Then, a stepwise multiple regression analysis was applied to explore significantly predictive variables correlated to sexual dysfunction.

The pleasure, arousal, and approach motivation ratings were analyzed with 2 × 2 mixed-model analyses of variance (ANOVA), and the Greenhouse–Geisser method was run to adjust degrees of freedom. Using the type of pictures (erotic picture and neutral picture) as within-subject factors and sexual dysfunction (NSD and SD) as between-subject factors.

SPSS 22.0 was conducted for all statistical analyses. p-values were double-tailed with significance level at 0.05. The continuous variables were presented in the form of mean ± standard deviation.

**RESULTS**

**Demographic and Clinical Characteristics of Patients**

Based on the effect sizes derived from a previous study (26), the sample size should be 96. Thus, 162 patients in the present study were sufficient to obtain a power of 0.8 at α < 0.05 (two-tailed). Of the 162 male chronic schizophrenia patients, 128 were single (79.0%) and 34 were married (21.0%). The age of the patients ranged from 18 to 50 years, with an average age of 28.41 years (standard deviation = 7.61). The duration of education ranged from 5 to 16 years, with an average of
Prevalence of Sexual Dysfunction in Male Patients With Chronic Schizophrenia

The prevalence of global sexual dysfunction was 29.6% (48/162) in male patients with chronic schizophrenia, strength of sex drive dysfunction was 44.4% (72/162), sexual arousal dysfunction was 31.5% (51/162), penile erection dysfunction was 34.0% (55/162), the ability to reach orgasm dysfunction and satisfaction with orgasm dysfunction was 37.0% (60/162). Seventy-nine patients (48.8%) were active smokers, and 32 patients (19.8%) had a family history of psychiatric disorders.

Comparison of Demographic and Clinical Variables Between Sexual Dysfunction and Non-sexual Dysfunction Patients

Sociodemographic and clinical variables of all patients are shown in Table 2. Compared with non-sexual dysfunction patients, sexual dysfunction patients had significantly higher negative symptoms of PANSS (p < 0.01). After controlling for age and age of onset, a significant difference was observed in negative symptoms (F = 5.30, p < 0.05) between non-sexual dysfunction and sexual dysfunction groups.

The Effect of Antipsychotics on Sexual Dysfunction

Among all participants, the most frequently used antipsychotic drug was risperidone [97 (59.87%)], followed by clozapine [46 (28.40%)] and other second-generation antipsychotics [19 (11.73%), e.g., olanzapine 14, quetiapine 5]. ANOVA showed no significant effect of antipsychotics on sexual dysfunction (all p > 0.05) (Table 3). Eight participants were prescribed citalopram in their medications. There was no significant effect of antidepressant on sexual dysfunction (all p > 0.05) (Table 2).

Correlation of Sexual Dysfunction and Demographic and Clinical Measures

Pearson correlation analysis showed that sexual dysfunction was correlated with negative symptoms (r = 0.16, p = 0.04; Bonferroni corrected p > 0.05). Furthermore, a multiple stepwise regression was performed to predict sexual dysfunction with all sociodemographic and clinical variables. Only negative symptoms predicted sexual dysfunction in a statistically significant manner [F(1,160) = 6.39, p < 0.01, adjusted R² = 0.03]. The coefficients of the variable are shown in Table 4.

Response to Erotic and Neutral Pictures Between Sexual Dysfunction and Non-sexual Dysfunction Patients

Sixty-eight subjects were administered the affective reactivity tasks. The pleasure, arousal, and approach motivation of the erotic pictures and neutral pictures in each group are shown in Table 5. Two-way ANOVAs showed significant interaction effects between sexual dysfunction group (SD vs. NSD) and picture types (Erotic Picture vs. Neutral Picture) on pleasure and arousal evaluation (all p < 0.05). However, there was no significant effect of sexual dysfunction group, picture types, and sexual dysfunction group × picture types on approach motivation evaluation (Table 5).

Correlation of Sexual Dysfunction and Response to Erotic Picture

Pearson correlation analysis showed that there was a significantly negative correlation between the severity of sexual dysfunction and the scores of pleasure (r = −0.28, p = 0.02) and approach motivation (r = −0.28, p = 0.02) on sexual picture, and all passed the Bonferroni correction.

DISCUSSION

To the best of our knowledge, this is the first study to investigate the prevalence of sexual dysfunction and its related risk factors among young and middle-aged male Chinese medicated chronic inpatients with schizophrenia, as well as the relationship between sexual dysfunction and the response to erotic pictures. The main findings of this study included: (1) the percentage of global sexual dysfunction was 29.6% in young and middle-aged male medicated chronic inpatients with schizophrenia; (2) negative symptoms were a risk factor for sexual dysfunction; (3) sexual dysfunction patients experienced lower pleasure and approach motivation when viewing erotic pictures.

Our cross-sectional clinical study indicated that nearly one-third of young and middle-aged chronically medicated male inpatients with schizophrenia suffer from sexual dysfunction, which was lower than those of the previous studies that used ASEX in schizophrenia patients in different sociocultural environments (30.7–84.5%) (2, 15, 26, 38, 43) but significantly higher compared to that of the general population, which was estimated between 8 and 15% (8, 15). In a previous study in China, the incidence of sexual dysfunction was 60.7% in male schizophrenia patients treated in a primary care setting (38), while it was 46% in outpatients with schizophrenia in Turkey males (15). Because sex is a private issue, sexual culture and sexual attitude are more conservative in Asia than in western countries. Talking about sex can make subjects feel shame, and reporting sexual dysfunction can damage their self-esteem. Therefore, people in Asia, especially in rural areas, are more likely to underestimates their sexual dysfunction (6, 44). In addition, age is another important risk factor for sexual dysfunction in the general population (45) and schizophrenia patients (15, 38). In our study, the mean age was 28.4 years, which was younger than those in the study by Hou et al. (38).
TABLE 1 | Frequency of sexual dysfunction in each domain of ASEX.

| ASEX Items               | N (%)/SD patients | N (%)/NSD patients | t     | p    |
|-------------------------|-------------------|-------------------|-------|------|
| Strength of sex drive   | 72/162(44.4)      | 2.60 ± 0.83       | 19.84 | <0.001 |
| Sexual arousal          | 51/162(31.5)      | 2.40 ± 0.74       | 18.39 | <0.001 |
| Penile erection         | 55/162(34.0)      | 2.49 ± 0.64       | 19.76 | <0.001 |
| Ability to reach orgasm | 60/162(37.0)      | 2.43 ± 0.68       | 21.09 | <0.001 |
| Satisfaction with orgasm| 60/162(37.0)      | 2.31 ± 0.68       | 21.82 | <0.001 |
| Total score ≥19         | 48/162(29.6)      | 13.67 ± 2.82      | 19.30 | <0.001 |

Scores of 4 or greater on the 1–6 Likert scale were considered “dysfunctional.” Individual with one or more of the following: total ASEX score ≥19; score on any of 5 items ≥5; score on 3 or more items ≥4.

SD, sexual dysfunction group; NSD, non-sexual dysfunction group.

TABLE 2 | Characteristics of schizophrenia patients with or without sexual dysfunction ASEX.

|                     | All patients (N = 162) | SD patients (N = 48) | NSD patients (N = 114) | t or χ² | p    |
|---------------------|------------------------|----------------------|------------------------|---------|------|
| Age (years)         | 28.41 ± 7.61           | 27.98 ± 8.73         | 28.56 ± 7.18           | 0.44    | 0.66 |
| Education (years)   | 9.29 ± 2.74            | 8.98 ± 3.60          | 9.07 ± 2.65            | 0.17    | 0.87 |
| Marital status      |                        |                      |                        |         |      |
| Single/married      | 128/34, 79.01%/20.99%  | 129/44, 77.09%/22.91%| 99/56, 67.21%/32.79%   | 1.88    | 0.19 |
| Family history of mental illness (yes/no) | 32/130, 19.75%/80.25% | 31/47, 19.23%/80.77% | 99/56, 67.21%/32.79% | 0.51    | 0.52 |
| Smoking behavior (yes/no) | 79/83, 48.77%/51.23% | 78/52, 48.77%/51.23% | 101/60, 64.17%/35.83% | 0.06    | 0.09 |
| BMI (kg/m²)         | 22.70 ± 3.80           | 22.19 ± 4.24         | 22.78 ± 3.63           | 0.45    | 0.66 |
| Duration of illness (years) | 6.51 ± 5.92        | 6.08 ± 6.09          | 6.54 ± 6.01            | 0.44    | 0.66 |
| Age of onset (years) | 22.12 ± 7.20           | 22.23 ± 8.22         | 22.20 ± 6.99           | −0.02   | 0.99 |
| Hospitalization numbers | 3.88 ± 2.70        | 2.20 ± 3.38          | 3.70 ± 4.45            | 1.95    | <0.05 |
| Psychopathology PANSS |                       |                      |                        |         |      |
| Positive symptom    | 18.31 ± 6.64           | 18.27 ± 5.67         | 18.33 ± 7.03           | 0.55    | 0.96 |
| Negative symptom    | 18.72 ± 7.80           | 21.02 ± 8.81         | 17.75 ± 7.15           | −2.48   | <0.01 |
| General psychopathology | 39.48 ± 11.71       | 40.90 ± 10.65        | 38.89 ± 12.13          | −1.00   | 0.32 |
| Total score         | 76.51 ± 22.36          | 80.19 ± 21.47        | 74.96 ± 22.64          | −1.38   | 0.18 |
| Positive factor      | 11.69 ± 7.28           | 12.31 ± 5.40         | 11.43 ± 5.23           | −0.97   | 0.33 |
| Negative factor      | 15.28 ± 7.36           | 18.00 ± 8.24         | 14.14 ± 6.66           | −3.13   | <0.01 |
| Cognitive factor     | 8.38 ± 3.38            | 8.17 ± 3.61          | 8.46 ± 3.30            | 0.51    | 0.61 |
| Excited factor       | 9.13 ± 3.90            | 8.60 ± 3.60          | 9.35 ± 4.02            | 1.11    | 0.27 |
| Depressive factor    | 7.31 ± 3.20            | 7.54 ± 3.27          | 7.21 ± 3.18            | −0.60   | 0.55 |
| On antipsychotics    |                       |                      |                        |         |      |
| Clozapine, (n, %)    | 46/28.40               | 13/27.08             | 33/28.95               | 0.54    | 0.80 |
| Risperidone, (n, %)  | 97/59.87               | 28/58.34             | 69/60.52               | 0.09    | 0.77 |
| SGAs except Clozapine and Risperidone, (n, %) | 19/11.73              | 7/14.58              | 12/10.53               | 0.09    | 0.77 |
| On antidepressants, (n, %) | 8/4.94               | 2/4.17               | 6/5.26                 | 0.09    | 0.77 |

and other studies. Another important factor was the type of antipsychotic drugs used by all patients, and there were no significant differences in the effects of antipsychotic and antidepressant drugs on sexual dysfunction, which is consistent with a previous study showing that no differences were found in the prevalence of sexual dysfunction between the drug groups (olanzapine, risperidone, aripiprazole, haloperidol) (6, 43). However, in other studies, there were significant differences in the incidence of sexual dysfunction between different antipsychotic drugs (7, 46, 47). Taken together, the diverse rates of sexual dysfunction are likely to be related to the patients’ treatment setting (inpatient, community or outpatient), sample size, different sociocultural contexts, clinical symptoms (especially negative symptoms), and possibly antipsychotic treatment.

Most previous studies have shown that sexual dysfunction is correlated with the severity of psychopathology, and higher scores on the negative symptoms were correlated with the severity of sexual dysfunction (6, 48). In our present study, multiple regression analysis showed that sexual dysfunction was positively correlated with negative symptoms of PANSS, which is consistent with the results of a previous study (26). Malik et al. (49) reported that negative symptoms predicted decreased libido. In addition, our present study did not observe the relationship
between sexual dysfunction and duration of illness and age of onset, which was consistent with a previous study (6).

Notably, in our present study, compared with the non-sexual dysfunction group, the sexual dysfunction group showed lower pleasure and arousal induced by sexual stimuli. Furthermore, our study also found that sexual dysfunction was negatively correlated with pleasure and approach motivation elicited by erotic pictures, which is in line with previous investigations that indicated that hedonic response and intrinsic motivation were impaired in schizophrenia (33, 50–52). As pointed out by Najas-Garcia et al. (53), anhedonia, apathy, avolition, and motivational deficits are negative symptoms in schizophrenia, and sexual dysfunction was significantly correlated with negative symptoms. Compared with non-sexual dysfunction patients, when presenting erotic stimuli, schizophrenia patients with sexual dysfunction experienced lower pleasure and sexual arousal and enhanced motivation to avoid sexual stimulation (53). In addition, sexual dysfunction has a negative effect on the self-esteem of schizophrenia patients, reduces the motivation of actively seeking and maintaining intimate relationships, and worsens their quality of life (44, 54).

Our study had several limitations. First, as a cross-sectional study, no causal relationship of sexual dysfunction can be concluded. Second, the sexual dysfunction and emotion were all self-assessment, which may lead to biased results. Third, this study may not be representative because most of the participants in the current study were young and middle-aged men, and the average age of subjects was younger (28.41 years). In addition, our results were only applicable to hospitalized patients, and our findings may not be generalized to other subjects recruited from other locations in China. Fourth, although we did not find significant differences in the effects of antipsychotic drugs on sexual dysfunction, we did not collect the antipsychotic dose data. Also, we did not measure medication adherence. Therefore, we do not know whether there was a correlation between antipsychotic dose or medication adherence and sex dysfunction in this study, which should be remedied in future investigation. Fifth, the subjectivity of the responses should be considered as a limitation. In this study, the mean age of our patients was below 30, and for them, reporting sexual dysfunction might be an embarrassment. Moreover, some patients might have thought that the study would lead to a treatment for sexual dysfunction, since we did not write this down on the consent form. Sixth, homosexual preferences would affect response to the erotic stimuli and also the reporting of the response. In this study, whether a patient was heterosexual or non-heterosexual only depended on his self-report, and we did not have any objective measure. In particular, Chinese men are reluctant to admit non-heterosexuality, even if they are homosexual. Therefore, the real impact of homosexual preference to the erotic stimuli and reported responses warrant further investigation. Seventh, adherence to oral second-generation antipsychotics for at least 1 year, it is unclear that patients on treatment for 1 year is considered as chronic as one on treatment for 20 years, where inevitably long-term effects are added to sexual dysfunction. Finally, the sample size was small, especially the numbers of affective reactivity task analysis, which may limit the reliability of the findings.

In conclusion, this study revealed that the prevalence of global sexual dysfunction was higher in young and middle-aged male schizophrenia patients than that in the general population. The negative symptom PANSS was the risk factor of sexual dysfunction. Furthermore, upon exposure to sexual stimulation, male schizophrenia patients with sexual dysfunction experienced lower pleasure and approach motivation. Thus, our findings in this study may provide new insights into understanding the psychopathological mechanism of sexual dysfunction so as to find potential solutions for sexual dysfunction in patients with schizophrenia. Clinicians should assess the sexual functioning of patients with schizophrenia and recommend appropriate antipsychotic medications in combination with physical therapy to improve negative symptoms, such as repetitive

### TABLE 3 | The effect of antipsychotics on sexual dysfunction ASEX.

| ASEX Items                  | Clozapine       | Risperidone    | Olanzapine     |
|-----------------------------|-----------------|----------------|----------------|
| Strength of sex drive       | 3.74 ± 1.54     | 3.60 ± 1.38    | 3.63 ± 1.01    |
| Sexual arousal              | 3.07 ± 1.50     | 3.24 ± 1.40    | 3.11 ± 1.15    |
| Penile erection             | 3.48 ± 1.47     | 3.25 ± 1.32    | 3.11±1.20      |
| Ability to reach orgasm     | 3.28 ± 1.54     | 3.43 ± 1.46    | 3.63 ± 1.61    |
| Satisfaction with orgasm    | 3.17 ± 1.66     | 3.34 ± 1.44    | 3.58 ± 1.61    |
| Total score                 | 16.74 ± 6.77    | 16.86 ± 5.69   | 17.05 ± 4.63   |

### TABLE 4 | Predictors generated by Multivariate Logistic Regression with ASEX total score as dependent variables.

| Coefficients Standardized coefficients | t | p | 95.0% confidence interval for B |
|----------------------------------------|---|---|------------------------------|
| (Constant)                             | 14.452 | 1.050 | 13.766 | <0.001 | 12.379 | 16.526 |
| Negative factor                        | 0.157 | 0.062 | 0.198 | 2.529 | 0.012 | 0.034 | 0.279 |
transcranial magnetic stimulation (rTMS) and transcranial direct current stimulation (tDCS). In addition, clinicians should implement concurrent psychotherapy to improve medication adherence, self-esteem, and self-confidence; promote interpersonal communications; adjust attitudes; and improve sexual experiences.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/supplementary material, further inquiries can be directed to the corresponding author/s.

**ETHICS STATEMENT**

The studies involving human participants were reviewed and approved by the Ethics Committee of the Third People’s Hospital of Ganzhou City. The patients/participants provided their written informed consent to participate in this study.

**AUTHOR CONTRIBUTIONS**

XZ: conceptualization, methodology, writing—reviewing, editing, and supervision. DL: formal analysis, writing—original draft, and funding acquisition. HD, HG, WL, DZ, ZM, DaH, DoH, QM, and LQ: investigation, resources, and data curation. SL, MX, NZ, and HW: conceptualization, writing—review, and editing. All authors have contributed to and have approved the submitted version.

**FUNDING**

This research was supported by Key R&D Program of Jiangxi Province (2020BBGL73106), Key R&D Program of Science and Technology in Ganzhou (201926), the Planning Program of Health Committee of Ganzhou (201911), the Planning Program of Health Committee of Jiangxi Province (20202089), and grants 81771139 and 81371477 from the CAS Pioneer Hundred Talents Program and the National Natural Science Foundation of China.

**ACKNOWLEDGMENTS**

We want to express special gratitude to the participants for their cooperation and patience across this study. We also thank the medical staff of the Third People’s Hospital of Ganzhou City for their help in data collection.

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