Does Schizophrenia Itself Cause Obesity?

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Background: Schizophrenia (SC) is considered the most serious of all mental disorders. Some antipsychotics are associated with weight gain and metabolic abnormalities. Whether SC itself causes obesity remains uncertain.

Methods: We collected 185 first-episode drug-naive SC and 59 healthy controls (HCs) from the Third People’s Hospital of Foshan, Guangdong, China, and distinguished their course of disease in order to understand the body mass index (BMI) and body fat metabolism of SC.

Results: We found that excluding the drug factors, the longer the course of SC, the more obvious the increase of BMI and the higher the proportion of obesity. BMI was positively correlated with age, course of disease, fasting blood glucose (FBG), low-density lipoprotein (LDL), triglyceride (TG), and total cholesterol (TC), and negatively correlated with high-density lipoprotein (HDL). The results of regression analysis were further proof that age (β = 0.094, p < 0.001), duration (β = 0.081, p = 0.002), FBG (β = 0.987, p = 0.004), and TG (β = 0.918, p = 0.002) were the risk factors for the increase of BMI. HDL (β = –2.875, p < 0.001) was the protective factor.

Conclusion: SC itself can increase BMI and easily lead to obesity. We should pay more attention to the monitoring of blood metabolism indicators, so as to reduce the risk of obesity and improve the quality of life of patients.

Keywords: metabolic indexes, obesity, overweight, body mass index, schizophrenia

INTRODUCTION

Schizophrenia (SC) is considered the most serious of all mental disorders (1, 2). Many patients with SC do not fully recover, and even among those with a good prognosis, the disease changes their lives, including social isolation, stigma, and reduced likelihood of finding a companion (3, 4). Poor eating habits, weight gain, smoking, and substance abuse can reduce life expectancy by 13–15 years (5, 6). Antipsychotics are commonly used to treat many different mental disorders (7). Some drugs are related to weight gain and metabolic abnormalities (8–10). Therefore, some people believe that obesity in SC is mainly caused by antipsychotics (11). However, whether SC itself causes obesity remains uncertain. Hence, we collected the first-episode drug-naive SC from Foshan, Guangdong, China, and distinguished their course of disease, in order to understand the body mass index (BMI) and body fat metabolism of the SC.
MATERIALS AND METHODS

Participants

The outpatients and inpatients with SC, who met the diagnostic criteria of the Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV) from August 2016 to September 2021 in the Department of Psychiatry, The Third People’s Hospital of Foshan, Guangdong, People’s Republic of China, were included. According to the course of disease, they were divided into groups A (duration < 6 months), B (6 months ≤ duration < 24 months), and C (duration ≥ 24 months).

Inclusion criteria were as follows: (1) SC: ① 18–45 years old (in order to avoid physical diseases caused by aging); ② years of education ≥ 6; ③ Han nationality; ④ before test, did not take any antipsychotics, antidepressants, mood stabilizers, sedatives, etc.; (2) healthy control (HC): volunteers recruited through advertising in Foshan from March 2020 to December 2021; ① 18–45 years old; ② no history or family history of psychosis; ③ Han nationality; ④ years of education ≥ 6; ⑤ gender, age, and education were matched with the patient group.

Exclusion criteria were as follows: ① comorbidity other mental disorders, including intellectual disability or other cognitive impairment; ② patients with diabetes, hypertension, severe kidney, liver function damage, cardiac insufficiency, etc.; ③ those who did not cooperate with venous blood drawing due to phobia, etc.; ④ smoking habits (≥ 1 cigarette per day) or drinking habits (≥ 1 unit alcohol per week); 1 unit alcohol = 480–600 ml of beer = 350 ml of low alcohol liquor or red wine, yellow wine = 50 ml of high spirits (40° or more).

Assessment and Blood Sampling

Details such as names, gender, and age were collected through the interviews from subjects who were willing to participate, after receiving the signed informed consent form. Then, the weight, height, and BMI of these subjects were measured with Automatic Measuring Stadiometer BSM370 (https://smitechasia.com). The positive and negative syndrome scale (PANSS) was used to assess the extent of the patient’s mental symptoms.

According to the diagnostic criteria of overweight and obesity proposed by the China Obesity working group, “BMI<24 kg/m²” was defined as non-obesity, “24 kg/m² ≤ BMI<28 kg/m²” was defined as overweight, and “BMI ≥ 28 kg/m² as obesity” was defined as obesity (12).

Subjects were instructed to be on fasting for more than 8 h before drawing the venous blood and the night before blood drawing, to maintain a normal diet, not to drink any alcohol or coffee after dinner, and to avoid strenuous exercise. FBG, HDL, LDL, TG, and TC were recorded in the subject’s clinical data sheet for subsequent data analysis.

Statistical Methods

We used the Statistical Product and Service Solutions 19 software¹ to analyze the data. Chi-square test was used to compare the differences of general demographic parameters.

The blood metabolic indexes, which correspond to normal distribution, were compared by one-way ANOVA and non-parametric data were compared by Kruskal-Wallis test. The relationships of BMI and variable indexes were analyzed by Pearson correlation. Multiple linear regression was used to analyze the influencing factors of BMI.

RESULTS

Comparison of Demographic Characteristics and Metabolic Indexes

There were 244 participants in this study, including HC (n = 59), A (n = 92), B (n = 45), and C (n = 48), while 10 subjects were excluded due to consumption of breakfast before drawing blood or due to not showing interest in performing blood test.

There were no significant differences in age, gender, height, weight, PANSS, HDL, LDL, TG, and TC (p > 0.05). Also, there were significant differences in BMI and FBG (p < 0.05). The incidence of obesity in groups HG, A, B, and C was 5.26, 7.61, 15.56, and 22.92%, respectively, and the incidence of overweight was 22.81, 27.17, 22.22, and 31.25%, respectively (Table 1 and Figure 1).

Pearson Correlation Between Body Mass Index and Various Indexes in Schizophrenia

The results showed that BMI was positively correlated with age, duration, FBG, LDL, TG, and TC, and negatively correlated with HDL, while PANSS, PANSS (P), PANSS (N), and PANSS (G) were not correlated with BMI (Table 2).

Multiple Linear Regression Analysis of the Influencing Factors of Body Mass Index in Schizophrenia

Taking BMI as the dependent variable (Y) and age, duration, HDL, LDL, TG, and TC as independent variables (X), and gender as a covariate, a stepwise multiple linear regression model (F = 16.394, p < 0.001) was established. Finally, the elements such as age, duration, FBG, HDL, and TG, were considered for the model (Table 3).

DISCUSSION

Our study included first-episode, drug-naive SC with a different course of disease. By comparing their blood metabolic indexes, we found that after excluding the drug factors, the longer is the course of SC, the more obvious is the increase of BMI, and the higher is the proportion of obesity.

A previous study has shown that there was no significant difference in the incidence of obesity in first-episode SC compared with HC (13). However, after using psychotropic drugs for a period of time, the weight of patients would increase to varying degrees, even to the extent of obesity (10, 14). The abovementioned research indicated that SC itself did not

¹https://www.ibm.com/analytics/spss-statistics-software
TABLE 1 | Comparison of demographic characteristics and metabolic indexes.

|                | HC (n = 59) | A (n = 92) | B (n = 45) | C (n = 48) | F/χ² | P   |
|----------------|-------------|------------|------------|------------|------|-----|
| Course of disease (month) | –          | 0–6        | 6–24       | ≥ 24       | –    | –   |
| Duration (month)             | –          | 3.09 ± 1.75| 11.80 ± 1.54| 25.29 ± 6.35| 2.830| 0.065|
| Gender (male/female)          | 25/34      | 40/52      | 18/27      | 19/29      | 0.271| 0.965|
| Age [year]                   | 27.11 ± 7.76| 27.09 ± 12.44| 28.62 ± 12.44| 25.81 ± 7.73| 0.975| 0.405|
| Height (cm)                  | 163.93 ± 7.75| 164.32 ± 9.33| 162.50 ± 10.22| 164.38 ± 7.90| 0.679| 0.692|
| Weight (kg)                  | 60.48 ± 10.19| 60.99 ± 11.97| 61.26 ± 12.31| 66.15 ± 14.39| 2.830| 0.065|
| BMI (kg/m²)ab                | 22.47 ± 3.36| 22.55 ± 3.91| 23.14 ± 3.82| 24.37 ± 4.37| 3.247| 0.040*|
| Obesity (%)                  | 5.26       | 7.61       | 15.56      | 22.92      | –    | –   |
| Overweight (%)               | 22.81      | 27.17      | 22.22      | 31.25      | –    | –   |
| Non-obesity (%)              | 71.93      | 65.22      | 62.22      | 45.83      | –    | –   |
| PANSS –                     | 98.36 ± 13.53| 101.70 ± 15.51| 99.35 ± 13.20| 0.494| 0.612|
| PANSS (P) –                  | 23.50 ± 6.64| 23.51 ± 7.06| 23.91 ± 6.68| 0.238| 0.789|
| PANSS (N) –                  | 24.70 ± 7.72| 25.88 ± 7.44| 24.71 ± 6.68| 0.683| 0.508|
| PANSS (G) –                  | 50.16 ± 6.94| 52.29 ± 8.34| 50.71 ± 8.80| 3.591| 0.006*|
| FBG (mmol/l)bc               | 5.49 ± 0.47| 5.11 ± 0.69| 5.04 ± 0.68| 3.57 ± 1.22| 0.683| 0.508|
| HDL (mmol/l)                 | 1.27 ± 0.29| 1.30 ± 0.39| 1.42 ± 0.50| 1.30 ± 0.36| 0.600| 0.470|
| LDL (mmol/l)                 | 2.64 ± 0.60| 2.47 ± 0.74| 2.62 ± 0.81| 2.58 ± 0.70| 0.314| 0.219|
| TG (mmol/l)                  | 1.12 ± 0.46| 1.43 ± 1.20| 1.38 ± 1.07| 1.57 ± 1.12| 1.539| 0.095|
| TC (mmol/l)                  | 4.72 ± 0.76| 4.33 ± 1.02| 4.59 ± 1.16| 4.62 ± 0.98| 1.629| 0.443|

BMI, body mass index; PANSS, positive and negative syndrome scale; FBG, fasting blood glucose; HDL, high-density lipoprotein; LDL, low-density lipoprotein; TG, triglyceride; TC, total cholesterol.

abHC vs. C, bcA vs. C, p < 0.05. *Indicates the comparison among groups (p < 0.05). Values are expressed as mean ± standard deviation.

cause obesity. According to a large-scale national epidemiological survey, the results show that the prevalence of obesity in urban areas in southern China was 2.8–7.2% from 2010 to 2018 (15). The course of 0–6 months of HC (5.26%) and SC (7.61%) was very close to the above range. However, some researchers believed that SC had metabolic abnormalities first, and then obesity (16). When we focused on SC patients with a longer course of disease (over 6 months), we found that the incidence of obesity was higher than the abovementioned data of the healthy population of the national epidemiological survey. Moreover, when we enrolled the subjects, most patients had a course of disease from 0 to 6 months, and fewer patients had a longer course of disease, which was also the reason that our results were consistent with those of predecessors.

Comparing the FBG of each group, we found that the FBG decreased in patients with 0–6 months course. As we know that SC always starts with negative or positive symptoms (3), psychotic symptoms can lead to eating disorders (17). When patients had positive symptoms such as victim delusion or taste hallucinations, some patients would be afraid to eat (18). They
that excluding the influence of drugs, BMI increased with the mental disorders (10, 20). At the same time, our results suggested the incidence of obesity in SC increased with age, similar to most mental disorders, which meant that the severity of SC had nothing to do with BMI. Serum insulin and had no correlation with the scores of PANSS in each group, showing that BMI was positively correlated with age, course of disease, FBG, LDL, TG, and TC, negatively correlated with HDL, and had no obvious difference (19). Therefore, there was no difference between the members in their daily life, which would reduce food intake, resulting in the decline of FBG the next day. With the migration of the course of disease, the blood sugar detected before breakfast can reflect the function of B cells in the islets, which generally indicates the secretion function of basal insulin, and is the most commonly used indicator for diabetes. FBG is closely related to BMI in the study of diabetes mellitus, which has been proved by extensive research (22, 23). LDL, TG, and TC belong to clinical blood lipid indexes. Their effects on obesity have been unanimously recognized. It is generally believed that their increase is strongly related to the occurrence of obesity (24). Finally, the results of regression analysis were further proof that age, duration, FBG, and TG were the risk factors for the increase of BMI. HDL was the protective factor.

Of course, it is best that we conduct a longitudinal follow-up study on the first-episode, drug-naive SC, and do not use any drugs to intervene, which obviously violates ethics and humanitarianism. So, we had to test this hypothesis by selecting untreated SC patients with a different course of disease. In addition, patients with a long course of disease tend to shrink back, including diet and living habits, which would lead to the deviation of our study. Overall, SC itself can increase BMI and easily lead to obesity. We should pay more attention to monitoring of blood metabolism indicators, so as to reduce the risk of obesity and improve the quality of life of patients.

**TABLE 2** Pearson correlation between BMI and various indexes in SC.

| BMI              | $r$   | Education | $r$   | Duration* | $r$ |
|------------------|-------|-----------|-------|-----------|-----|
| Age*             | 0.314 | 0.028     | 0.244 |
| PANSS            | 0.045 | -0.064    | 0.001 |
| PANSS (P)        | 0.673 | 0.549     | 0.538 |
| PANSS (N)        | 0.073 | 0.043     | 0.313 |
| LDL*             | -0.273 | 0.448     | 0.256 |
| Serum insulin    | 0.089 | 0.001     | 0.001 |

BMI, body mass index; PANSS, positive and negative syndrome scale; FBG, fasting blood glucose; HDL, high-density lipoprotein; LDL, low-density lipoprotein; TG, triglyceride; TC, total cholesterol. *Indicates $p < 0.05$.

**TABLE 3** Multiple linear regression analysis of influencing factors of BMI in SC.

| Model            | $B$      | Standard error | Standard coefficient | $t$    | $p$   |
|------------------|----------|----------------|----------------------|--------|-------|
| (Constant)       | 16.394   | 1.772          |                      | 9.254  | <0.001* |
| Age              | 0.094    | 0.026          | 0.239                | 3.609  | <0.001* |
| Duration         | 0.081    | 0.026          | 0.195                | 3.081  | 0.002* |
| FBG              | 0.987    | 0.341          | 0.260                | 2.899  | 0.004* |
| HDL              | -2.875   | 0.804          | -0.297               | -3.576 | <0.001* |
| LDL              | 0.572    | 0.933          | -0.107               | 0.613  | 0.541 |
| TG               | 0.918    | 0.285          | 0.263                | 3.218  | 0.002* |
| TC               | 0.573    | 0.773          | 0.150                | 0.741  | 0.460 |

BMI, body mass index; FBG, fasting blood glucose; HDL, high-density lipoprotein; LDL, low-density lipoprotein; TG, triglyceride; TC, total cholesterol. *Indicates $p < 0.05$.

often consult doctors during the acute exacerbation period, resulting in lower blood glucose measured in the blood test the next day. With the migration of the course of disease, the patient adapted to the psychotic symptoms, and then the FBG gradually recovered. If patients started with negative symptoms, they would shrink back and need to be supervised by family members in their daily life, which would reduce food intake, resulting in the decline of FBG the next day. In addition, patients with SC with a course of more than 2 years were accompanied by undetectable mental symptoms, and the abnormal diet was not obvious (19). Therefore, there was no difference between the results of their FBG.

To further explore the factors affecting BMI, we correlated the metabolic indexes and PANSS scores with BMI. The results showed that BMI was positively correlated with age, course of disease, FBG, LDL, TG, and TC, negatively correlated with HDL, and had no correlation with the scores of PANSS in each group, which meant that the severity of SC had nothing to do with BMI. The incidence of obesity in SC increased with age, similar to most mental disorders (10, 20). At the same time, our results suggested that excluding the influence of drugs, BMI increased with the increase in the course of disease. The research team of Solmi supported our results that there were multiple risk-related genes between SC and eating behavior disorder and BMI, which were closely related to each other (21). FBG means that after fasting overnight (at least 8–10 h without any food, except drinking water), the blood sugar detected before breakfast can reflect the function of B cells in the islets, which generally indicates the secretion function of basal insulin, and is the most commonly used indicator for diabetes. FBG is closely related to BMI in the study of diabetes mellitus, which has been proved by extensive research (22, 23). LDL, TG, and TC belong to clinical blood lipid indexes. Their effects on obesity have been unanimously recognized. It is generally believed that their increase is strongly related to the occurrence of obesity (24). Finally, the results of regression analysis were further proof that age, duration, FBG, and TG were the risk factors for the increase of BMI. HDL was the protective factor.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in this 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 Foshan, China. The patients/participants provided their written informed consent to participate in this study.

**AUTHOR CONTRIBUTIONS**

JL, YC, and YY made great contributions to the conception, design, and writing of the article. XX, XL, ZL, CX, and GX provided assistance in the acquisition, analysis, and interpretation of data. All authors approved the publication of the manuscript.

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REFERENCES

1. Whiting D, Gulati G, Geddes J, Fazel S. Association of schizophrenia spectrum disorders and violence perpetration in adults and adolescents from 15 countries: a systematic review and meta-analysis. JAMA Psychiatry. (2022) 79:120–32. doi: 10.1001/jamapsychiatry.2021.3721

2. Buckley P. Evidence suggests complex links between violence and schizophrenia. Interview by Bridget M. Kuehn. JAMA. (2012) 308:658–9. doi: 10.1001/jama.2012.9364

3. Jauhar S, Johnstone M, McKenna P. Schizophrenia. Lancet. (2022) 399:473–86.

4. Green M, Horan W, Lee J. Nonsocial and social cognition in schizophrenia: current evidence and future directions. World Psychiatry. (2019) 18:146–61. doi: 10.1002/wps.20624

5. Hjorthøj C, Stürup A, McGrath J, Nordentoft M. Years of potential life lost and life expectancy in schizophrenia: a systematic review and meta-analysis. Lancet Psychiatry. (2017) 4:295–301. doi: 10.1016/S2215-0366(17)30078-0

6. Nielsen R, Banner J, Jensen S. Cardiovascular disease in patients with severe mental illness. Nat Rev Cardiol. (2021) 18:136–45.

7. Olsson M, Blanco C, Wang S, Laje G, Correll C. National trends in the mental health care of children, adolescents, and adults by office-based physicians. JAMA Psychiatry. (2014) 71:81–90. doi: 10.1001/jamapsychiatry.2013.3074

8. Vancampfort D, Correll C, Galling B, Probst M, De Hert M, Ward C, et al. Diabetes mellitus in people with schizophrenia, bipolar disorder and major depressive disorder: a systematic review and large scale meta-analysis. World Psychiatry. (2016) 15:166–74. doi: 10.1002/wps.20309

9. Vancampfort D, Stubbs B, Mitchell A, De Hert M, Wampers M, Ward C, et al. Risk of metabolic syndrome and its components in people with schizophrenia and related psychotic disorders. Bipolar disorder and major depressive disorder: a systematic review and meta-analysis. World Psychiatry. (2015) 14:339–47. doi: 10.1002/wps.20252

10. Li X, Shi X, Tan Y, Yu Y, Tang C, Xu G, et al. Metabolic indexes of obesity in patients with common mental disorders in stable stage. BMC Psychiatry. (2022) 22:91. doi: 10.1186/s12888-022-03752-2

11. Correll C, Sikich L, Reeves G, Johnson J, Keeton C, Spanos M, et al. Metformin add-on vs. antipsychotic switch vs. continued antipsychotic treatment plus healthy lifestyle education in overweight or obese youth with severe mental illness: results from the IMPACT trial. World Psychiatry. (2020) 19:69–80. doi: 10.1002/wps.20714

12. Zeng Q, Li N, Pan X, Chen L, Pan A. Clinical management and treatment of obesity in China. Lancet Diabetes Endocrinol. (2021) 9:393–405. doi: 10.1016/S2213-8587(21)00047-4

13. Tian Y, Wang D, Wei G, Wang J, Zhou H, Xu H, et al. Prevalence of obesity and clinical and metabolic correlates in first-episode schizophrenia relative to healthy controls. Psychiatry Research. (2021) 287:745–53. doi: 10.1016/j.psychres.2021.02024

14. Correll C, Detraux J, De Lepeleire J, De Hert M. Effects of antipsychotics, antidepressants and mood stabilizers on risk for physical diseases in people with schizophrenia, depression and bipolar disorder. World Psychiatry. (2015) 14:119–36. doi: 10.1002/wps.20204

15. Wang L, Zhou B, Zhao Z, Yang L, Zhang M, Jiang Y, et al. Body mass index and obesity in urban and rural China: findings from consecutive nationally representative surveys during 2004-18. Lancet. (2021) 398:53–63. doi: 10.1016/S0140-6736(21)00798-4

16. Sánchez-Ortí J, Balanzá-Martínez V, Correa-Ghisays P, Selva-Vera G, Vila-Francés J, Magdalena-Benedito R, et al. Specific metabolic syndrome components predict cognition and social functioning in people with type 2 diabetes mellitus and severe mental disorders. Acta Psychiatr Scand. (2020) 100:1–12.

17. Kouidrat Y, Amad A, Stubbs B, Louhou R, Renard N, Diouf M, et al. Disordered eating behaviors as a potential obesogenic factor in schizophrenia. Psychiatry Res. (2018) 269:450–4. doi: 10.1016/j.psychres.2018.08.083

18. Malaspina D, Walsh-Messinger J, Brunner A, Rahman N, Corcoran C, Kimhy D, et al. Features of schizophrenia following premorbid eating disorders. Psychiatry Res. (2019) 278:275–80. doi: 10.1016/j.psychres.2019.06.035

19. Fawzi M, Fawzi M. Disordered eating attitudes in Egyptian antipsychotic naïve patients with schizophrenia. Compr Psychiatry. (2012) 53:259–68. doi: 10.1016/j.comppsych.2011.04.064

20. De Hert M, Correll C, Bobes J, Cetkovich-Bakmas M, Cohen D, Asai I, et al. Physical illness in patients with severe mental disorders. I. Prevalence, impact of medications and disparities in health care. World Psychiatry. (2011) 10:52–77. doi: 10.1002/wps.2051-5545.2011.tb00014.x

21. Solmi F, Mascarello M, Zammit S, Kirkbride J, Lewis G. Polygenic risk for schizophrenia, disordered eating behaviours and body mass index in adolescents. Br J Psychiatry. (2019) 215:428–33. doi: 10.1192/bjp.2019.39

22. Chiavaroli L, Lee D, Ahmed A, Cheung A, Khan T, Blanco S, et al. Effect of low glycaemic index or load dietary patterns on glycaemic control and cardiometabolic risk factors in diabetes: systematic review and meta-analysis of randomised controlled trials. Bmj. (2021) 374:n1651.

23. Zaharia O, Strassburger K, Strom A, Bönhof G, Karusheva Y, Antoniou S, et al. Risk of diabetes-associated diseases in subgroups of patients with recent-onset diabetes: a 5-year follow-up study. Lancet Diabetes Endocrinol. (2019) 7:684–94. doi: 10.1016/S2213-8587(19)30187-1

24. Zhang Y, Yang J, Ye J, Guo Q, Wang W, Sun Y, et al. Separate and combined associations of physical activity and obesity with lipid-related indices in non-diabetic and diabetic patients. Lipids Health Dis. (2019) 18:49. doi: 10.1186/s12944-019-0987-6

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