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10.1097/XCE.0000000000000274

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Document Version

Publisher's PDF, also known as Version of record

Citation for published version (Harvard): H Heald, A, Shakespeare, M, Phillipson, A, Cade, J, Netter, P & Higgs, S 2023, 'Does food responsiveness change in people with first-episode psychosis over a period of 3 months after commencing antipsychotics? Preliminary results', *Cardiovascular endocrinology & metabolism*, vol. 12, no. 1, e0274. https://doi.org/10.1097/XCE.000000000000274

Link to publication on Research at Birmingham portal

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Cardiovascular Endocrinology & Metabolism 2023. 12:e10.1097/ XCE.000000000000274:doi: 10.1097/XCE.00000000000274

Does food responsiveness change in people with first-episode psychosis over a period of 3 months after commencing antipsychotics? **Preliminary results**

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Weight gain occurs in many patients with schizophrenia [1], a diagnosis often subsequently made in individuals who first present with psychosis (characterized by culture-dyscongruent beliefs, disordered thinking and abnormal experiences). Frequently weight gain occurs in the first few weeks and months after diagnosis [2] and is related to an increase in appetite and food intake, with often double portions consumed and high-calorie snacks taken between meals. Early weight gain is a predictor of longer-term weight gain [3]. Weight gain in psychosis is associated with an increased likelihood of the development of type 2 diabetes [4] and increased cardiovascular risk with associated shortened life expectancy [5].

Many of the drugs that are effective in treating psychosis are associated with changes in the way that people experience reward when they eat [6].

To date, there has been little research in this area [7], and as such little is known about people's response to food and appetite shortly after commencing antipsychotic medication.

The purpose of this project was to increase our understanding of exactly what are the drivers of weight gain in terms of an individual's experience of food reward and reduced satiety - and therefore how we can help people with schizophrenia to keep their weight down. At this stage, we are looking at the feasibility of applying currently available evaluation tools to people in this situation.

Our aim was to determine if food responsiveness changes in people with first-episode psychosis (FEP) over a period of 3-6 months after commencing antipsychotics. The study was approved by the West Midlands Ethics Committee (Reference 20/WM/0146).

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Received 15 October 2022 Accepted 24 October 2022.

A convenience sample was used with the aim of recruiting service users from early intervention services. This was designed as a feasibility study to provide data to underpin a fully powered, larger investigation.

The study was conducted in early intervention services across the Rotherham, Doncaster and South Humber (RDASH) National Health Service Foundation Trust. Early intervention services provide support and treatment for people with FEP in England. Participation in the study was offered to those who had commenced antipsychotic medication within the previous 6 weeks, in the context of a diagnosis of FEP.

Inclusion criteria: patients diagnosed with FEP, aged 18–35 years and attending the RDASH early intervention service.

Exclusion criteria were: age <18 years or >50 years, unable or unwilling to provide informed consent, pregnancy, non-English speaking, intellectual disability or communication difficulties hindering their ability to complete assessments or mental health or distress levels too high as deemed by their care coordinator.

Rating scales applied were

Dutch Eating Behaviour Questionnaire (DEBQ) [8]: this measures restrained eating, emotional eating and external eating.

Intuitive Eating Scale [9]: this measures an individual's tendency to follow their physical hunger and satiety cues.

Power of food questionnaire [10]: this measures responsiveness to the food environment.

The loss of control over eating scale [11]: this measures a global sense of whether individuals experience loss of control over eating.

The questionnaires were administered through a telephone interview at baseline and 3–6 months later.

In the event, only six participants were recruited because of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) pandemic constraints on recruitment. The six participants (four men and two women) were aged between 17 and 31 years. All were diagnosed with FEP. Time from initiation of treatment with an antipsychotic to the first interview varied from 13 to 39 days.

Three people were started on olanzapine at the dose of 5 or 10 mg daily, one was started on risperidone at 1 mg daily and two were started on aripiprazole at 10 and 15 mg daily, respectively.

Baseline scores are shown in Fig. 1. At the beginning of antipsychotic treatment, there was moderate responsiveness to food and a moderate tendency to follow hunger/satiety cues, with a low emotional drive to eat and low compulsion to eat.

At 3–6-month follow-up (pooled results) (Fig. 1), there was: a trend for a decrease in the DEBQ emotional eating score (P=0.06) reflecting less response to external cues such as the aroma of food, presence of food or watching others eat and a decrease in the intuitive eating score (reflecting a lesser tendency to follow hunger/satiety cues) (P=0.02).

Our preliminary results reveal that at the beginning of antipsychotic treatment there was moderate responsiveness to food and a moderate tendency to follow hunger/satiety cues, with a low emotional drive to eat and low compulsion to eat. At 3- to 6-month follow-up, there was a decreased response to satiety cues (intuitive eating

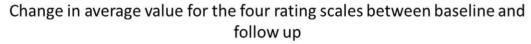
score) and a trend for a decrease in the DEBQ emotional eating score (P=0.06) reflecting less response to external cues. Externality theory describes external eating as eating in response to food-related stimuli, regardless of hunger or satiety [12]. The change to less responsiveness to external food-related stimuli may relate to the effects of the acute illness itself or the effects of antipsychotic treatment on response to food.

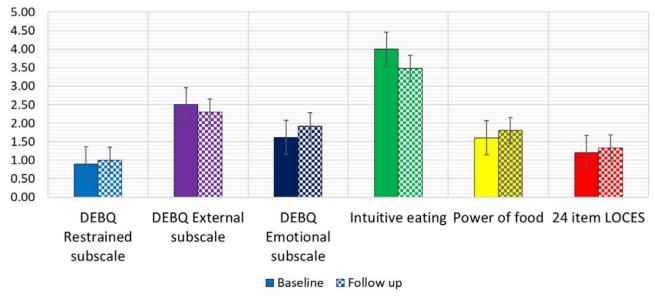
It is important to point to a recent report that even in the absence of antipsychotic medication relative to matched controls, patients with first-episode psychosis who were not taking antipsychotic medication consumed more saturated fat and showed an altered association between BMI and neural response to food cues [13]. Therefore what we have observed here is likely the consequence of multiple interacting factors including prescribed medication.

Although we had few participants and were not able to differentiate responses by antipsychotic agents, we feel that this is a relevant contribution to the literature given the limited work so far in this area [7]. These findings, although preliminary highlight how maladaptive eating patterns and alterations in the association between BMI and neural responses to food cues may be established early in the course of a psychotic illness which may be subsequently diagnosed as schizophrenia.

The difference between the baseline and the follow-up eating behavior scores may provide important clues as to the change in eating behavior with antipsychotic

Fig. 1





Change in DEBQ scores, intuitive eating, power of food and LOCES scales from baseline to 3- or 6-month follow-up interview. DEBQ, Dutch Eating Behaviour Questionnaire; LOCES, loss of control over eating scale.

treatment. Greater knowledge of this will enable more effective targetting of interventions to reduce the likelihood of weight gain with its attendant consequences of elevation of cardiometabolic risk [4,5].

We accept that the numbers here are too small to draw any definite conclusions. The study was very much constrained by the recruiting period starting and completing in a period when face-to-face consultations in mental health services and interactions between healthcare professionals were occurring on a very limited basis. Furthermore, not all participants were contactable for the interview at 6 months so we pooled the 3-month results with the 6-month results where there was no 6-month follow-up. Nevertheless, we feel that our findings provide the basis for a future larger and more comprehensive study, which takes account of the practicalites of recruiting people who face the challenges of FEP.

Acknowledgements

This study was funded by the University of Manchester N8 Pump Priming Fund. Grant Code: AA16858.

The data that support the findings of this study are available from the corresponding author upon reasonable request.

A.H. and A.P. led on the conceptualization, design and writing up of this project. M.S. conducted the data analysis and contributed to all aspects of the article. J.C. and S.H. advised on all aspects of the methods including the specific rating scales employed and also contributed to the Discussion. P.N. acted as a senior leader of the article.

Conflicts of interest

There are no conflicts of interest.

References

- Heald AH, Martin JL, Payton T, Khalid L, Anderson SG, Narayanan RP, et al. Changes in metabolic parameters in patients with severe mental illness over a 10-year period: a retrospective cohort study. Aust N Z J Psychiatry 2017; **51**:75-82.
- Allison DB, Mentore JL, Heo M, Chandler LP, Capelleri JC, Infante MC, Weiden PJ. Antipsychotic induced weight gain: a comprehensive research synthesis. Am J Psychiatry 1999; 156:1686-1696.
- Parsons B, Allison DB, Loebel A, Williams K, Giller E, Romano S, et al. Weight effects associated with antipsychotics: a comprehensive database analysis, Schizophr Res 2009: 110:103-110.
- Cooper SJ, Reynolds GP, Barnes T, England E, Haddad PM, Heald A, et al.; With expert co-authors (in alphabetical order). BAP guidelines on the management of weight gain, metabolic disturbances and cardiovascular risk associated with psychosis and antipsychotic drug treatment. J Psychopharmacol 2016; 30:717-748.
- Howell S, Yarovova E, Khwanda A, Rosen SD. Cardiovascular effects of psychotic illnesses and antipsychotic therapy. Heart 2019; **105**:1852-1859.
- 6 Benarroch L, Kowalchuk C, Wilson V, Teo C, Guenette M, Chintoh A, et al. Atypical antipsychotics and effects on feeding: from mice to men. Psychopharmacology (Berl) 2016; 233:2629-2653.
- Stogios N, Smith E, Asgariroozbehani R, Hamel L, Gdanski A, Selby P, et al. Exploring patterns of disturbed eating in psychosis: a scoping review. Nutrients 2020: 12:3883
- van Strien T, Frijters JER, van Staveren WA, Defares PB, Deurenberg P. The predictive validity of the Dutch restrained eating scale. Int J Eat Disord 1986: **5**:747-755.
- Tylka TL, Kroon Van Diest AM. The intuitive eating scale-2: item refinement and psychometric evaluation with college women and men. J Couns Psychol 2013: 60:137-153.
- 10 Lowe MR, Butryn ML, Didie ER, Annunziato RA, Thomas JG, Crerand CE, et al. The power of food scale. a new measure of the psychological influence of the food environment. Appetite 2009; 53:114-118.
- 11 Latner JD, Mond JM, Kelly MC, Haynes SN, Hay PJ. The loss of control over eating scale: development and psychometric evaluation. Int J Eat Disord 2014: 47:647-659
- Schachter S, Goldman R, Gordon A. (1968). Effects of fear, food deprivation and obesity on eating. J Pers Soc Psychol 1968; 10:91-97.
- Borgan F, O'Daly O, Hoang K, Veronese M, Withers D, Batterham R, et al. Neural responsivity to food cues in patients with unmedicated first-episode psychosis. JAMA Netw Open 2019; 2:e186893.

DOI: 10.1097/XCE.0000000000000274