

Methods. A 64-year-old woman with an underlying recurrent depressive disorder with psychotic symptoms presented to a psychiatric hospital in June 2023. She exhibited self-neglect, low mood, paranoid delusions, and non-concordance to oral psychiatric medications.

In the first week, she declined all oral medications and was subsequently started on flupentixol decanoate (Depixol) depot injection at 40 mg once every 2 weeks. While showing good improvements in her mental state, she began complaining of akathisia and dystonia since July 2023, consistent with extrapyramidal side effects secondary to flupentixol.

The symptoms improved by lowering flupentixol to 30 mg every 2 weeks and adding procyclidine 5 mg twice daily and propranolol 20 mg three times daily.

In early September 2023, she experienced severe restlessness, stiffness, muscle weakness and felt hot and clammy over 36 hours. Physical observations showed fever, tachycardia, and hypertension. Examination revealed diaphoresis, rigidity in both upper and lower limbs, lower limb weakness, and normal reflexes. Blood tests indicated acute kidney injury (AKI) stage 1, deranged liver function tests, and a creatinine kinase (CK) level of 9405.

She was promptly admitted to the medical hospital for NMS and received extensive intravenous fluid rehydration along with oral Dantrolene. She made a complete recovery, and Depixol was discontinued. Two weeks later, she was started on quetiapine and gradually titrated to 50 mg once daily.

Results. EPSE and NMS are associated with dopamine receptor blockade and commonly occur during the initiation or dosage increment of neuroleptic medications.

NMS is rare but life-threatening, presenting with manifestations of muscle rigidity, pyrexia, altered mental status, sympathetic nervous system lability and elevated CK.

In our case, our patient, who recently started taking neuroleptic medication, experienced EPSE and later deteriorated acutely, raising a high suspicion of NMS. It is essential to consider other possible diagnoses, including serotonin syndrome, malignant hyperthermia, malignant catatonia and electrolyte disturbances.

The commonly used diagnostic criteria include Diagnostic and Statistical Manual of Mental Disorders Fifth Edition (DSM-5) and Levenson's criteria but diagnosis of NMS remains clinical.

The crucial step after identifying NMS is to immediately stop the neuroleptic agent, followed by supportive medical treatment. **Conclusion.** Early recognition and prompt treatment of NMS in our patient led to a full recovery.

Abstracts were reviewed by the RCPsych Academic Faculty rather than by the standard *BJPsych Open* peer review process and should not be quoted as peer-reviewed by *BJPsych Open* in any subsequent publication.

Feeding Two Birds With One Seed: Using Fluoxetine for Pre-Menstrual Dysphoric Disorder and Bulimia Nervosa

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Aims.

Background

This case study describes the use of fluoxetine for reduction of pre-menstrual dysphoric disorder (PMDD) and bulimia nervosa symptoms. The case report also describes an increase in binge

purge symptoms in the pre-menstrual period, along with other mood and cognitive symptoms. This supports a hormonal basis to the exacerbation of eating disorders. Patient consent was obtained prior to the publication of this report.

Methods.

Case report

A 41-year-old lady with significant binge purge behaviours and mood disturbance was referred to our eating disorder service. She met the diagnostic criteria for bulimia nervosa after a thorough assessment, along with a component of mood dysregulation. She was prescribed sertraline for depressive symptoms in primary care. The patient described a worsening of mood symptoms, along with cognitive difficulties before the start of her menstrual cycle. After a medical review, we agreed on tracking these symptoms along with the binge-purge frequency for a period of two cycles. This was done using a PMDD tracker. The tracker reflected a clear diagnosis of PMDD along with an exacerbation of bingeing and purging symptoms before the start of a menstrual cycle. Following this, sertraline was switched to fluoxetine, and titrated up to its maximum dose of 60 mg a day.

Results.

Discussion

Following the commencement of fluoxetine, purging frequency dramatically reduced and subsequently stopped. Although mood symptoms still persisted, the specific mood symptoms along with cognitive symptoms in the pre-menstrual period reduced.

Conclusion. There is some evidence for the use of fluoxetine use for binge purge symptoms in bulimia nervosa. Fluoxetine has also been used either continuously or in the luteal phase for PMDD. This case report reflects the possible correlation between binge-purge symptoms and PMDD symptoms, and the potential use of fluoxetine for dual symptom reduction. PMDD still remains to be a significantly under-diagnosed condition in women. This case report also signifies the importance of exploring PMDD symptoms in eating disorders.

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Antidepressant-Coincident Manic Episode in a Prepubertal Girl Presenting With Obsessive-Compulsive or Related Disorders: A Case Report

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Aims. Obsessive-Compulsive or Related Disorders (OCRDs) comprise a group of disorders characterized by repetitive thoughts and behaviours and are fairly less prevalent among children. The recommended treatment for OCRDs involves high doses of antidepressants, specifically selective serotonin reuptake inhibitors (SSRIs), along with non-pharmacological management. However, evidence suggests that the risk of inducing mania with antidepressants may be especially high in children and adolescents aged 14 years and younger.

Methods. Here, we present a case of a nine-year girl, studying in fifth standard, with normal birth and development history, with

no past/family history of psychiatric illness, presented with psychiatric illness of one-year duration and was diagnosed with Trichotillomania, Obsessive-Compulsive Disorder, Skin picking and Onychophagia as per the 11th revision of International Classification of Diseases (ICD-11). After initiating tab. escitalopram 5 mg for 10 days, child developed a manic episode, which leads to a diagnostic dilemma as well as difficulties in her further management. In view of the bipolarity, escitalopram was stopped and the child was started on tab. aripiprazole 2.5 mg which was gradually up-titrated to 7.5 mg/day, following which the manic episode completely resolved and there was also improvement in OCD, hair pulling and skin picking behaviour. Later for the remaining symptoms few sessions of Habit reversal therapy were held. Currently the patient is maintaining well on aripiprazole 7.5 mg for the last six months.

Results. The uniqueness of this case is demonstrated through current limited literature on comorbid OCRDs and antidepressant coincident manic episode, especially in children in whom diagnosing manic episode possess a great challenge owing to various differential diagnosis. While deciding pharmacological therapy in children with OCRDs or Mania the efficacy as well as their safety profile should be considered. Currently there are no medications approved by FDA for treatment of acute manic episode in patients below 10 years of age and use of SSRI which are considered first line for treatment of some OCRDs may exaggerate the manic episode. In literature, second generation antipsychotics such as aripiprazole is found to be useful for the management of both manic episode (as monotherapy) as well as OCD (as an adjuvant). In this case aripiprazole monotherapy led to significant improvement in both groups of symptoms.

Conclusion. Thus, SSRIs should be used cautiously in children with OCRDs and aripiprazole along with other approved non-pharmacological management strategies can be considered as a good treatment option in children with OCRDs and antidepressant coincident manic episode.

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Remission of Symptoms of Functional Neurological Disorder (FND) Utilising Novel Interventions: A Case Report

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Aims. FND can be considered as an umbrella term that includes range of motor and sensory system dysfunctions with genuine symptoms including paralysis, tremors, sensory disturbance, speech disturbance and seizure. Functional seizures usually termed as Non-Epileptic attack disorder (NEAD) can result in profound persisting disability. Brief bouts of unprovoked and uncontrollable laughter, spontaneous in origin, combined with facial contraction in the form of smile, is termed as 'gelastic seizures'. Modafinil is a dopamine modulating molecule for which evidence is accumulating towards its cognitive enhancement

role in multiple domains. Furthermore, it has been shown to promote hippocampal neurogenesis and synaptic plasticity in pre-clinical studies. We report a case of FND in which pharmacological (Modafinil) and non-pharmacological interventions (Brain retraining) resulted in resolution of symptoms of probable gelastic episodes.

Methods. A 50-year-old lady who was referred by consultant neurologist to our Neuropsychiatry pilot service with episodes of uncontrollable laughing, singing, screaming and suffering from staggering and imbalance. Following these episodes, patient described sleeping for hours with fatigue. Her husband first noticed low mood 12 years ago during post-natal period. Treatment with fluoxetine reportedly contributed to 'cyclical highs and mood variations'. One year later, her 'gelastic episodes' started and continued to occur every 2 or 3 months and they were brought on by a range of factors including tiredness, menstrual periods and stress. Patient also reported atypical cognitive deficits such as 'losing vocabulary' and 'stuck every couple of seconds'. Furthermore, detailed history confirmed possible traits of attention deficit and hyperactive thinking since her childhood.

Results. Following a comprehensive assessment, the role of the brain in the manifestation of her symptoms was discussed and agreed upon. Strategies based on Cognitive Behaviour Therapy principles such as active distraction, brain retraining, engaging in therapeutic activities and expressive writing were discussed and agreed upon. Following detailed risk-benefit analysis, modafinil was initiated at 200 mg dose in the morning. Patient made a remarkable recovery nearly back to her baseline with resolution of her gelastic episodes and thus improvement in her mental state. She continues to be stable in the community.

Conclusion. This case highlights the importance of recognising and treating cluster of symptoms which might belong to the impulsive-compulsive spectrum. This further emphasises the role of dopamine-modulating agents such as modafinil along with brain-retraining strategies.

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Nabilone as Part of the Holistic Treatment in Early Onset Alzheimer's Dementia: A Case Study

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Aims. Optimal management of Behavioural and Psychological symptoms of Dementia (BPSD) remains challenging. This report describes using nabilone, a synthetic cannabinoid, in a 61-year-old woman with Alzheimer's dementia (AD) experiencing progressive BPSDs.

Methods. AM was diagnosed with AD in February 2019 and prescribed donepezil and mirtazapine. In August 2021, her behaviour deteriorated, becoming paranoid, repeatedly pacing and developing expressive aphasia. Behaviours further declined leading to an admission to our dementia ward under the Mental Health Act 2007 in January 2022. AM showed limited response to medications including risperidone and mirtazapine which were switched