European Psychiatry S479

Here, we discuss an unusual presentation of Huntington's Disease causing diagnostic dilemma.

**Objectives:** Case report discussing the unusual presentation of Huntington's Disease.

**Methods:** Case: Mr X is a 61 year old Caucasian male.He had an uneventful birth and early childhood attaining milestones appropriately. He experienced childhood adversity in the form of sexual abuse between ages 2-14 years. His mental health difficulties started following sexual abuse when he attempted to end his life by hanging and overdosing at age 15. He got married twice, both of which broke down. There is a history of significant alcohol abuse between ages 40-50. Following this, he had a myocardial infarction and a stroke requiring stenting.

He presented to Psychiatric Outpatient Services in 2011 with auditory hallucinations, social anxiety with panic attacks, OCD type rituals, claustrophobia and feeling hot all the time. He was started on an antipsychotic medication for psychosis, but clinically deteriorated. He started having anger outbursts, marching on the spot ,and head banging. He was diagnosed with Huntington's Chorea in 2021 after he had developed chorea. He currently has low mood and is head banging for hours.

**Results:** Psychiatric symptoms in HD can span a variety of domains but most common are symptoms of frontal lobe dysfunction-disinhibition, poor attention, irritability, impulsivity and personality change. Apathy, emotional blandness and social withdrawal are also prominent features.

Mr X had strong family history of Paranoid Schizophrenia (aunt and cousin). There was no family history of HD. His mental health problems started early in life with DSH, Depression and Harmful use of Alcohol. He presented predominantly with psychotic symptoms like auditory hallucinations, social anxiety, paranoia. Motor symptoms started late which he incorporated into voluntary movements like head banging which made it difficult to differentiate from deliberate self harm.

**Conclusions:** Psychiatric symptoms constitute the core of HD. Studies have shown that though depression and personality change are typical of HD, there are number of other psychiatric symptoms that can impair quality of life. Early diagnosis and treatment of these symptoms will help patients and families to cope better with severe symptoms of this progressive disease.

**Disclosure of Interest:** None Declared

#### **EPV0246**

### Concurrent Gender Dysphoria/Incongruence and Autism Spectrum Disorder, a literature review

N. Clementi<sup>1,2</sup>\*

<sup>1</sup>General Adult Psychiatry/ Gender Identity Clinic, Royal Cornhill Hospital, NHS Grampian and <sup>2</sup>University of Aberdeen, School of Medicine, Aberdeen, United Kingdom

\*Corresponding author.

doi: 10.1192/j.eurpsy.2024.995

**Introduction:** Several studies have found that ASD (Autism Spectrum Disorder) and GD (Gender Dysphoria by DSM-V)/GI (Gender Incongruence by ICD-11) tend to co-occur, and in recent years the interest and publications on this comorbidity has increased rapidly.

**Objectives:** To review the prevalence of ASD in individual with a diagnosis of GD/GI.

To better tailor and improve care offered in the National Health Service (NHS) Gender Identity Clinics (GICs) throughout the UK. **Methods:** Systematic literature review was conducted via Pub Med, MEDLINE and PsycINFO by the author, for all English-language articles published between 2018 and 2023, containing keywords as ASD, GD (Gender Dysphoria), GI (Gender Incongruence), transgender, autistic traits, autism, gender diversity, gender variance.

**Results:** Rate of people with ASD appear to be higher in people accessing Gender Identity Clinics (GICs) than in the general population. Results from this literature review show increased prevalence of GD and GI in ASD population.

Conclusions: This comorbidity has highlighted the importance of better tailor transgender healthcare services for people with neurodevelopmental conditions and neurodiversity, to avoid delay in ASD individuals accessing care and gender affirming medical treatments. Services should strive to provide an effective and equitable service. It is also important to better identify potential barriers for ASD people in accessing gender care. Literature also shows the people with ASD have more difficulties in communicating and describing their gender narrative and to express their wishes for gender treatments. Symptoms including problems in communications and social skills, obsession and rigidity can also impact their assessment of GD/GI in gender identity services. Some studies showed that for individuals who have concurrent ASD and GD/GI, assessment in GICs may be extended to better review their wishes for gender identity and for gender affirming treatment. Further research is needed to better investigate and understand factors explaining the relationship between ASD and gender diversity. There is still limited research in the real life experiences of gender diverse and autistic people. There is also a need to improve Gender reassignment protocol nationally to better care for individual with ASD and GD/GI throughout GICs in the UK.

Disclosure of Interest: None Declared

### **EPV0247**

# Osmotic demyelination syndrome (ODS), and psychiatric manifestations

P. Argitis<sup>1\*</sup>, A. Karampas<sup>2</sup>, M. Peyioti<sup>1</sup>, A. Goudeli<sup>1</sup>, S. Karavia<sup>1</sup> and Z. Chaviaras<sup>1</sup>

<sup>1</sup>Psychiatric, General Hospital of Corfu, Corfu and <sup>2</sup>Psychiatric, General Hospital of Ioannina, Ioannina, Greece

\*Corresponding author.

doi: 10.1192/j.eurpsy.2024.996

**Introduction:** Hyponatriemia can be potentially fatal if it is not corrected immediately. The rapid correction of chronic hyponatriemia can cause demyelinating brain lesions.

**Objectives:** A fifty-six year old female was brought to the emergency department of the psychiatric clinic by her daughter, with incomprehensible speech and psycomotor agitation. She was diagnosed several years ago with bipolar disorder, with valproic acid and quetiapine being her current medication. She has been living alone, in a small suburban city. Approximately twenty four hours

S480 e-Poster Viewing

before her admission to the hospital she visited her daughter, which aligns with the onset of symptoms.

**Methods:** After both the brain CT scan and the lab results came back normal, the patient was admitted to the psychiatric clinic of the General Hospital of Corfu. On the fourth day of the patient's hospitalization - when both her speech and the psycomotor agitation showed signs of improvement- we were informed that three days before her admission to the clinic she visited the emergency department of another hospital where she was treated for hyponatriemia. The patient's hyponatriemia was corrected over the span of twelve hours by 35 mEq.

**Results:** After receiving this information, we ordered a brain MRI scan which revealed a central pontine myelinolysis. The result can explain the clinical symptoms that our patient showcased before her admission and could have been caused by the rapid correction of hyponatriemia.

**Conclusions:** The patient's speech was fully restored after four weeks and there were no symptoms consistent with any psycho emotional disorder.

Disclosure of Interest: None Declared

#### **EPV0248**

## QTc prolongation in patients hospitalized in enclosed psychiatric facilities in Corfu

P. Argitis<sup>1</sup>\*, A. Karampas<sup>2</sup>, M. Peyioti<sup>1</sup>, S. Karavia<sup>1</sup> and Z. Chaviaras<sup>1</sup>

<sup>1</sup>Psychiatric, General Hospital of Corfu, Corfu and <sup>2</sup>Psychiatric, General Hospital of Ioannina, Ioannina, Greece

\*Corresponding author.

doi: 10.1192/j.eurpsy.2024.997

**Introduction:** An undeniably significant amount of psychotropic medication can evidently affect the corrected QT (QTc) interval, which puts patients' lives at risk. More specifically, certain antipsychotic medication can increase the risk of QTc prolongation and by extension the risk of a potentially fatal arrhythmia or sudden cardiac death.

**Objectives:** Electrocardiograms (ECG) were contacted in one hundred and four (104) chronic patients, with psychosis, through out their hospitalization in several enclosed psychiatric facilities in Corfu. Almost the entirety of the patients along side their antipsychotic medication were also taking various other medication for their individual pathological issues. We observed any changes that might have occurred on the ECG in comparison with each patient's medication and it's potential effect on the QTc.

**Methods:** The measurements of the QT interval were made manually in lead V5 and the mathematical conversion was contacted using the Hodges correction formula.

**Results:** At least one ECG (n = 104) was performed. Among them 29,8% (n=31) had ECG abnormalities, including 13,5% (n=13) with a prolonged Qtc (481.2  $\pm$  26,8 ms). Covariates significantly associated with the QTc were gender (+17.2 ms if female, p < 0.0001) and age (+0.4 ms/year, p = 0.0001).

**Conclusions:** The QTc prolongation that was evident in a notable number of patients, emphasizes the importance of QTc monitoring in patients who are taking anti-psychotic medication. QTc prolongation risk factors should be assessed before the administration or prescription of any anti-psychotic medication.

Disclosure of Interest: None Declared

#### **EPV0249**

Persistent Adult-Onset Attention-Deficit/ Hyperactivity Disorder (ADHD) Manifesting as Occupational Impairment: Highlighting the Therapeutic Potency of Methylphenidate

P. S. Gopan

Psychiatry, Surabhi medical college, Siddipet, India doi: 10.1192/j.eurpsy.2024.998

**Introduction:** This case study emphasizes the significance of considering unrecognized adult-onset ADHD, particularly in patients with chronic forgetfulness and occupational inefficiencies refractory to standard treatment options. The case outlined involves a 33-year-old male with enduring cognitive impairments, leading to Extreme Anxiety Disorder with detrimental consequences on his professional progression and personal wellbeing.

**Objectives:** This necessitates the need for advanced research initiatives and broader awareness programs to facilitate improved diagnostic accuracy and optimization of therapeutic outcomes. Emphasizing ADHD as a potential cause of such symptomatology in adults and integrating effective treatment options can potentially pave the way to personalized therapeutic protocols.

**Methods:** The patient was approached via meticulous reconsideration of previous unsuccessful treatment paradigms that primarily included antidepressants and anxiolytics, which yielded cyclical patterns of negligible amelioration, compounded by intermittent emergence of suicidal ideation. Given the limited response, a differential diagnosis of Adult-Onset ADHD was entertained.

**Results:** The therapeutic intervention involving Methylphenidate administration led to a remarkable enhancement in the patient's mental health and occupational efficiency. Progress was also evidenced in the patient's improved confidence and self-esteem, with critical implications for his professional and personal life dynamics.

**Conclusions:** This case study underscores the transformative potential of precise ADHD management in adults with chronic cognitive impairments. Further research studies involving larger cohorts are warranted to enhance the understanding of adult ADHD, its prevalence, and therapeutic strategies, which could serve as key elements in improving the overall quality of life for these patients.

Disclosure of Interest: None Declared

### **EPV0250**

## The difficulties of Adult ADHD management within a Community Mental Health Team

S. Haugh\* and H. Belay

Department of Psychiatry, Connolly hospital, Dublin, Ireland \*Corresponding author. doi: 10.1192/j.eurpsy.2024.999