S1092 E-Poster Viewing

EPV1028

Clinico-psychopathological characteristics of patients with residual states in long-term schizophrenia

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doi: 10.1192/j.eurpsy.2023.2319

Introduction: Long-term schizophrenia, even in remission, is necessarily accompanied by residual symptoms that may be quite pronounced and may significantly affect many aspects of the patient's life, requiring exploration of specific therapeutic approaches. The alleviation of residual symptoms is an important factor in the patient's better adjustment.

Objectives: Assessment and study clinical characteristics long-term schizophrenia.

Methods: Clinical, statistical, psychometric. A total of 90 patients, mean age $66,6\pm13,3$ years, 26 males, 64 females were examined.

Results: Negative symptoms were predominant in patients with long-term schizophrenia (17,8 \pm 6,7). It was represented by: abstract thinking disorders $(2,8\pm1,0)$, stereotyped thinking $(2,7\pm1,1)$, passive-apathetic social isolation $(2,6\pm1,2)$, avolition $(2,6\pm0,8)$, flattening of affect (2,5 \pm 0,8). It manifested as lack of expressiveness in facial expressions and gestures, deficit of communicative gestures as well as emotional indifference $(2,4\pm1,1)$, limitation of contacts with people, and spontaneous and fluent speech impairments. Positive symptoms were rare, mainly represented by suspiciousness $(2,2\pm1,2)$, sometimes rising to delirium $(1,8\pm1,4)$. Conceptual disorganization was detected in 1,9 \pm 0,7. Agitation and aggression were generally not characteristic of those surveyed. Depression/ anxiety was quite pronounced in patients with long-term schizophrenia. Depression (1,8±0,8) was represented by low mood, hopelessness and loss of social interests. Anxiety (2,9 \pm 1,2) was even more prominent and predominant amongst all symptoms.

Conclusions: The authors expanded our understanding of the clinical characteristics of residual symptoms of long-term schizophrenia to allow timely identification and provision of medical and rehabilitative care.

Disclosure of Interest: None Declared

EPV1027

Addictive behaviors in schizophrenic patients: descriptive and analytical study

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doi: 10.1192/j.eurpsy.2023.2320

Introduction: The association of an addictive disorder (harmful use, abuse or dependence) with a schizophrenic disorder is the rule. Genetic vulnerability and social and economic factors are common to both disorders.

Objectives: determine the impact of addictive behavior on patients suffering from schizophrenia.

Methods: A descriptive and analytical retrospective study of 150 patients with schizophrenia hospitalized in the psychiatry department of Taher Sfar University Hospital in Mahdia from January 2017 to December 2021.

Results: The average age of the patients was 39.8 ± 11.23 years with a predominance of the age group 36-45 years (38.4%). All of the patients were male. Three quarters of the patients (75.5%) were consumers of psychoactive substances (PSA): nearly three quarters (72.8%) were dependent on tobacco, more than a third (39.7%) were dependent on alcohol, more than a quarter (29.1%) dependent on cannabis and almost a quarter (26.5%) dependent on other SPAs. Criminal history, suicide attempts and hospitalization in psychiatry were significantly more frequent in patients who consumed SPA than those who did not consume (39.5% vs 8.1%; p=0.008, 17.5% vs 2.7%; p=0.02, 89.5 % vs 75.7%; p=0.03, respectively). Patients who consumed SPA had significantly more positive signs of schizophrenia (51.8% vs 10.8%; p=0.001) and were significantly less observant to treatment (55.3% vs 16.3%; p=0.001) than those who did not consume. Hetero-aggressiveness, selfaggressiveness and job change were significantly more frequent in patients with addictive behaviors than those without addiction (86.8% vs 48.7%; p=0.001, 23.7% vs 2.7%; p=0.004, 14.9% vs 0%;p=0.015, respectively). Multivariate analysis revealed that criminal history, hetero-aggressiveness, predominant positive symptomatology and work stoppage were the factors independently associated with SPA consumption in patients with schizophrenia in our study (β =14.7 95% CI 3.23-67.01, p=0.001, β =0.099, 95% CI 0.03-0.31, p=0.001, β =7.18, 95% CI 2.09–24.67, p=0.002, β =5.24 95% CI 1.27-21.7; p=0.02, respectively).

Conclusions: According to our study, addictive comorbidities concern three quarters of our patients. They expose them to a higher risk of legal problems, hetero-aggressiveness, predominance of positive signs and instability at work. These results encourage the development of methods for early diagnostic identification of addictive behavior comorbid with schizophrenia as well as integrated care combining psychiatric and addictological care.

Disclosure of Interest: None Declared

EPV1028

DIAGNOSTIC CHALLENGES IN DISTINGUISHING AUTISM SPECTRUM DISORDER FROM PSYCHOSIS: A CASE REPORT

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Introduction: Autism spectrum disorders (ASD) and psychotic disorders have historically considered to be related conditions with a long history of diagnostic confusion. Although DSM-III distinguishes ASD and Schizophrenia Spectrum Disorders as distinct clinical entities, they continue to share overlaps in their clinical symptom presentations leading to diagnostic challenges that may consequentially result in delayed treatment. Prompt diagnosis is crucial in the context of psychosis, where early intervention impacts recovery.

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Objectives: To present the diagnostic challenges encountered in distinguishing ASD from Psychosis.

Methods: We present a case report demonstrating the challenges of distinguishing ASD from Psychosis.

Results: This is a case of a gentleman who initially presented to psychiatric services at 18 years old for conflicts with his mother related to his inflexibility to change. Further psychological evaluation revealed that he had a history of restricted social interaction with his peers, difficulties in non-verbal communications and identifying emotional states, stereotyped interests and obsessions that isolated him from his peers. He was diagnosed with ASD.

In subsequent presentations, there were symptoms of excessive preoccupation of his facial appearance, excessive concern over contracting HIV, obsessions with arranging objects in a particular order and avoiding words starting with the letter "S" out of fears of blasphemy. While these symptoms had qualities of cognitive inflexibility, they could not fully be explained by ASD. Additional diagnoses of Body Dysmorphic Disorder, Borderline Personality Disorder, Obsessive Compulsive Personality Disorder and At-Risk Mental State were considered.

A psychiatric admission was necessitated at 21 years old, when he presented with a 2-year history of repetitive banging of furniture in the middle of the night to communicate his frustrations towards his parents for their perceived acts of blasphemy. He also began to isolate himself, fearing that his parents would be able to look into his soul and reveal his sins. This paranoia towards his parents worsened to the point of urinating and defecating in his room to avoid his parents. His school performance declined as well.

A unifying diagnosis of psychosis was made. His previous diagnosis of ASD was challenged as a misdiagnosis, with the impression that he likely had attenuated psychotic symptoms in his adolescent years, disguised as autistic traits. The diagnosis of psychosis was confirmed when the patient's symptoms were observed to respond to antipsychotic treatment.

Conclusions: This case report illustrates the challenges in distinguishing ASD from psychosis. A prior diagnosis of ASD may result in diagnostic overshadowing and subsequent delays in diagnosing psychosis. Further research in diagnostic tools would be helpful for diagnostic precision, thereby enabling prompt treatment for better recovery outcomes.

Disclosure of Interest: None Declared

EPV1029

Amisulpride Augmentation in Schizophrenia Patients with Poor Response to Olanzapine: A 4-week, Randomized, Rater-Blind, Controlled, Pilot Study

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doi: 10.1192/j.eurpsy.2023.2322

Introduction: Olanzapine (OLA) is a common first-prescribed antipsychotic and has shown favorable efficacy in acutely exacerbated patients with schizophrenia. The mixed receptor activity of OLA and its greater affinity for serotonin 5-HT2A rather than

dopamine D2 receptors are similar to those of clozapine. Pharmacokinetically, OLA is metabolized mainly by hepatic cytochrome enzyme P450 1A2 (CYP1A2). Because risks of antipsychotic polypharmacy include increased drug-drug interactions, pharmacokinetic considerations are important for selection of antipsychotics to be combined. Due to its pharmacological characteristics, amisulpride (AMI), another atypical antipsychotic with proven efficacy, is a promising adjuvant agent of special interest. AMI is unlikely to interact with other drugs due to the low plasma protein binding and metabolism and does not affect the activity of the CYP system. Furthermore, AMI is highly selective for dopamine D2/D3 receptors; has minimal or no affinity for D1, D4, or D5 receptors. Despite the potential benefits of the combination of OLA and AMI, only a few open-label studies have been conducted, and no randomized clinical trial has been performed to date to examine the efficacy and tolerability of the combination. Hence, the goals of this study were to test the hypothesis that AMI augmentation would improve psychotic symptoms and be well tolerated in schizophrenic patients who showed poor response to OLA monotherapy.

Objectives: The purpose of this study was to compare the efficacy and tolerability of continued olanzapine (OLA) versus amisulpride (AMI) augmentation in schizophrenic patients with poor response to OLA monotherapy.

Methods: The present 4-week, randomized, rater-blinded study included 25 patients with schizophrenia who were partially or completely unresponsive to treatment with OLA monotherapy. Eligible subjects were randomly assigned at a 1:1 ratio to continuation of OLA monotherapy (OLA group) or OLA with AMI augmentation (AMI group). Efficacy was primarily evaluated using the Positive and Negative Syndrome Scale (PANSS) at baseline and at 1, 2, and 4 weeks.

Results: The changes in PANSS total score and PANSS-positive subscale score were significantly different (p < 0.05) between the OLA and AMI groups. The differences between the two groups in PANSS-negative subscale, PANSS-general subscale, Brief Psychiatric Rating Scale, and Clinical Global Impression-Severity (CGI-S) scale scores were not statistically significant.

Conclusions: AMI augmentation could be an effective strategy for patients with schizophrenia who show inadequate early response to OLA monotherapy.

Disclosure of Interest: W.-M. Bahk Grant / Research support from: Handok Pharmaceuticals, Seoul, Korea, Y. S. Woo: None Declared, S.-Y. Park: None Declared, B.-H. Yoon: None Declared, S.-M. Wang: None Declared, M.-D. Kim: None Declared

EPV1030

Jobs Stress and Prodromal Psychosis among Employees with Different Job Occupations Abstract

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Introduction: As social stress includes social isolation, urban living, trauma and many other stressful events but social stress in context of workplace or job stress includes different factors. As in the case of social stress present at job place can be identified as a stress that is caused by planned social isolation or lack of social