

Introduction: Metabolic syndrome (MetS) is of primary clinical interest because of its harmful impact on the general health and quality of life of patients with psychotic disorders. Paradoxically, MetS is associated with impaired cognitive functions in patients receiving antipsychotics primarily shown to improve cognition (e.g., clozapine and olanzapine).

Objectives: In this study, we aimed to investigate the relationship between MetS, cognitive functions, and peripheral inflammation.

Methods: The participants were 154 patients with schizophrenia. Fifty-seven patients met the criteria of MetS. We evaluated cognitive functions with the Repeated Battery for the Assessment of Neuropsychological Status (RBANS). The Positive and Negative Syndrome Scale (PANSS) quantified the clinical symptoms. We also measured the plasma levels of IL-6 and C-reactive protein (CRP). In addition to conventional statistics, we also calculated Cohen's effect size (d) and Bayes Factors (BF10).

Results: Results revealed that patients with MetS exhibited worse cognitive function relative to patients without MetS in attention ($d = 0.19$, $BF10 = 2.3$) and delayed memory ($d = 0.25$, $BF10 = 5.7$). No significant between-group differences existed in immediate memory, visuospatial functions, and language. The MetS and non-MetS groups did not differ in positive, negative, or general symptoms. Higher IL-6 levels were associated with worse delayed memory ($r = -0.56$, $BF10 = 34.6$).

Conclusions: Our results suggest that MetS-associated cognitive dysfunctions are less severe than reported in the literature: it was confined to two cognitive domains, the effect size was small, and the Bayesian evidence level was weak. Peripheral inflammation may mediate the association between MetS and long-term memory dysfunctions.

Disclosure of Interest: None Declared

EPP0341

Sociodemographic profile and prescribing pattern of antipsychotic medication in patients with Schizophrenia

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Introduction: Schizophrenia is a complex psychiatric disorder that changes the patient's life by influencing how they think, behave, express emotions, percept reality and their interpersonal relationships.

Objectives: The aim of this study was to evaluate sociodemographic and therapeutic factors that act as risk and protective factors in the clinical outcomes of patients diagnosed with schizophrenia.

Methods: This was an observational retrospective study including patients diagnosed with schizophrenia, treated at the "Xhavit Gjata" Psychiatric Hospital, Tirane, Albania, who were discharged between May 1- October 30, 2022. The follow-up period was six months. Data on further hospitalizations during the follow-up were obtained from the Department of Statistics, QSUT, and confirmed

by family members for hospitalizations in other psychiatric hospitals in the country. Univariate and multivariate analyses were conducted to identify potential factors associated with emergency room stays, length of stay, and time until the next admission.

Results: A total of 158 patients were included in the study, 63 women and 95 men ($p = 0.03$). The average age of the patients was 42.9 years, with women averaging 45.3 years and men 40.6 years ($p = 0.01$). 43.7% of them had elementary education. The average age of disorder onset was 24.7 years. Haloperidol was the ambulatory therapy used in 54.3% of patients, while atypical antipsychotics were used in 75.1% of patients. The most commonly used atypical antipsychotic was Risperidone in 34.1% of patients, followed by Olanzapine in 17% of cases. Depo antipsychotics were used in 35.1% of patients. Clozapine was administered to 29.3% of patients, where 12.8% for the first time. 54.2% of patients starting Clozapine for the first time had three or more admissions. Clozapine was more frequently used in men, showing a significant difference from women ($p = 0.05$). In 44.7% of cases, monotherapy was prescribed. The average hospital stay was 21.9 days, ranging from 2-68 days. Living with a family member, male gender, and being "married" helped reduce the length of hospital stay. In the 6-month follow-up period, 31.4% were re-hospitalized. Significant factors affecting the reduction of time spent outside the hospital until the next hospitalization were social problems, the number of previous hospitalizations, civil status "not married," living arrangements, negative symptoms, and alcohol use (nearly significant). Protective factors included Clozapine, which reduced the prevalence of hospitalization by 57% compared to patients not taking it. Additionally, the use of Clozapine and Haloperidol increased the time spent outside the hospital.

Conclusions: Social and family support, positive compliance, and antipsychotic therapy such as Clozapine serve as protective factors for patients diagnosed with schizophrenia.

Disclosure of Interest: None Declared

EPP0342

The impact of affective and negative symptoms on the development of psychosis in a six-year follow-up of a community-based population

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Introduction: The Clinical High Risk (CHR) group for transition to psychotic disorders (PD) is usually defined by the severity of positive symptoms, help-seeking and impairment in level of functioning. However, the CHR concept has a limited transition risk to PD. Recent studies have shown that some of the risks might be attributable to other symptoms.

Objectives: This study investigates the association between affective and negative symptoms and the risk of transition to PD in a community-based population of 2185 participants in Turkey.