(p<0.001), and α 1-PI activity was much higher (p<0.001) in C2. Patients of C1 and C2 did not differ in age, diagnosis, severity of the disease, but differed in clinical features of the course of the disease: significantly more patients with very late onset schizophrenia (76%) were met in C1 (χ 2=13.41, p<0.001). Also, different clinical-biological correlations were found in these clusters. Particularly, negative correlations of baseline NE activity with PANSS general psychopathology subscale scores (R=-0.39, p<0.05) and with total PANSS scores (R=-0.39, p<0.05) were found in C1. Positive correlation of GST activity with PANSS positive subscale score was found in C2 (R=0.43, p<0.05).

Conclusions: The revealed clusters differ in the extent of the glutathione antioxidant system impairment and in levels of the immune response markers. The revealing of the patient subgroups on the basis of biological markers reflecting impairments in metabolic and immune systems can represent interest in the search for individual treatment approaches.

Disclosure of Interest: None Declared

Others

EPP0600

Prevalence of ADHD in Adults: An Umbrella Review of International Studies

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Introduction: Attention deficit hyperactivity disorder (ADHD) is a neurodevelopmental disorder commonly diagnosed in school-age children. However, it can affect individuals of all age groups. This study aimed to provide a comprehensive analysis of the prevalence of ADHD in adults by conducting an umbrella review of systematic reviews and meta-analyses.

Objectives: To provide a comprehensive synthesis of published evidence on the prevalence of Attention Deficit Hyperactivity Disorder (ADHD) in adults through an umbrella review of systematic reviews and meta-analyses, with the aim of highlighting the significance of addressing and managing ADHD in the adult population. **Methods:** To conduct this study, we adhered to the guidelines outlined in the Preferred Reporting Items for Systematic Review and Meta-Analysis (PRISMA). We systematically searched databases such as PsychINFO, Web of Science, PubMed, and Scopus to identify relevant studies. Our review protocol was registered with PROS-PERO (registration number: CRD42023389704). The quality of the studies included in our analysis was assessed using the A Measurement Tool to Assess Systematic Reviews (AMSTAR). For the purpose of conducting a meta-analysis, we employed a random-effects model.

Results: Our umbrella review examined findings from five systematic reviews that encompassed data from 57 unique international primary studies undertaken between 2009 and 2021. These studies involved a total of 21,142,129 adult participants. The meta-analysis, employing an inverse variance-weighted random effect model, yielded a pooled prevalence estimate for ADHD in adults of 3.10% (95% confidence interval: 2.60%–3.60%). Regarding ADHD subtypes, our analysis revealed that ADHD-I (inattentive type) remained the most prevalent among adults, followed by ADHD-HI (hyperactive type) and ADHD-C (combined type).

Conclusions: Our results underscore the relatively high prevalence of ADHD among adults, with ADHD-I emerging as the most common subtype. These findings emphasize the need for proactive measures to prevent, mitigate, identify, and effectively manage ADHD in the adult population.

Disclosure of Interest: None Declared

EPP0601

Investigation of the Effect of Curcumin on Metabolic Dysfunction Caused by Clozapine in Rats

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Introduction: Antipsychotics disrupt intracellular cholesterol traffic and prevent the exit of low-density lipoprotein (LDL)-derived cholesterol from the endosome/lysosome compartment. It was showed that curcumin accelerated the release of cholesterolcontaining exosomes from cells with impaired intracellular cholesterol traffic due to antipsychotic treatment and suggested that curcumin may help minimize the negative metabolic effects associated with chronic antipsychotic treatment.

Objectives: This study aimed to investigate the effectiveness of orally administered curcumin to rats in preventing and treating metabolic syndrome-related side effects such as weight gain and dyslipidemia caused by clozapine.

Methods: In our research, a total of 32 male rats (Wistar Albino), 12 weeks old, produced at Selçuk University Experimental Research and Application Center, were used. All animals divided into 4 groups. Venous blood collection and weight measurements were taken from all groups at the beginning. 32 rats were randomly divided into 4 separate groups: control, only oral clozapine, oral clozapine + 50 mg/kg curcumin, and the oral clozapine + 100 mg/kg curcumin group. Groups II-III-IV were given 15 mg/kg clozapine orally daily for 3 weeks. AST, ALT, glucose, total cholesterol, Triglyceride, HDL, LDL and insulin were studied from the blood samples taken at the beginning and at the end of the experiment. Results: There was no statistically significant difference in comparisons of weight and insulin measurements between the groups at the end of the experiment (p>0.05). In glucose measurements at the end of the experiment, the control group was found to have significantly higher glucose values compared to the other groups (p <0.001). As a result of posthoc analyses, LDL measurements in the control group were found to be lower than those in the CLZ and CLZ +50 c groups (p<0.05). AST value of the control group was significantly higher than the CLZ+100c group (p=0.011). Measurements of the control group for ALT were found to be higher than those of the CLZ+50c and CLZ+100c groups (p<0.05). There was