

EW0250

Efficacy and safety of MIN-101: A new drug for the treatment of negative symptoms in schizophrenia a 12-week randomized, double blind, placebo-controlled trial

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Objective To compare the efficacy, safety, and tolerability of MIN-101, a compound with high affinities for sigma 2 and 5-HT_{2A} receptors, to placebo in treating negative symptoms, in stabilized patients with schizophrenia.

Methods This multi-national phase 2b trial enrolled 244 patients with schizophrenia who were symptomatically stable for ≥ 3 months prior to entering the trial and had scores ≥ 20 negative subscale of the PANSS. Patients were randomized to monotherapy with MIN-101 32 mg/day, MIN-101 64 mg/day or placebo in a 1:1:1 ratio. The primary endpoint was the PANSS negative symptom score based on the five factors (pentagonal) model.

Results Statistically significant reduction in the primary endpoint score was demonstrated for MIN-101 32 mg and 64 mg compared to placebo ($P \leq 0.022$, ES 0.45 and ≤ 0.003 , ES 0.58, respectively). This was supported by similar effects on most of the secondary measurements including: the PANSS three factors negative symptoms subscale, PANSS total score, CGI, BACS, CDSS, and PSP. There were no statistically significant differences in PANSS positive subscale scores between MIN-101 and placebo. No weight gain or clinically significant changes in vital signs, prolactin levels, routine laboratory values, metabolic indices and extrapyramidal symptom scores (EPS) were observed.

Conclusions Since positive symptoms and EPS did not change, the improvement in negative symptoms was not secondary to improvement in positive symptoms or EPS, suggesting that MIN-101 might be the first specific treatment to have a direct effect on negative symptoms.

Disclosure of interest I have received consultant fees from Minerva Neuroscience the sponsor of this trial and own stock of Minerva Neuroscience

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EW0251

The importance of family in the long-term evolution of psychoses

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Introduction Adherence and tolerance to treatment are important factors, which may predict the long-term evolution of a psychosis. Family members may influence prognosis by modulating emotional expressivity and treatment supervision.

Objectives To assess the role of family members in the long-term evolution of psychoses.

Method The present study is retrospective, conducted on patients with psychosis. Data were obtained from psychiatric records extending for a period of four years. The following parameters were analyzed: socio-demographic data, family relationships (parents, spouses) and clinical/evolutive data (onset age for psychosis, number of recurrences).

Results We analyzed 71 patients, 42 (59.2%) women and 29 (40.8%) men with a mean age of 30.38 years (SD=9.33). The subjects were diagnosed according to ICD 10 criteria with acute and transient psychotic disorder (50 patients, 70.4%), schizophrenia (13 patients, 18.3%), and schizoaffective disorder (8 patients, 11.3%). Patients who reported conflicts between parents had significantly more recurrences ($t = -2.1$, $P = 0.04$), while those who reported

satisfactory relationships in their family of origin had fewer recurrences ($t = 2.58$, $P = 0.01$) and a later onset age ($t = -2.89$, $P = 0.006$). Unmarried/single subjects had the psychosis onset at a significantly earlier age ($t = 4.72$, $P = 0.0001$). In addition, these patients had more conflicts between parents ($Z = -2.02$, $P = 0.04$) in comparison with married ones.

Conclusions Conflicts in the family of origin may predispose to a greater number of recurrences and to an earlier disorder onset. The presence of a spouse may represent a protective factor.

Disclosure of interest The authors have not supplied their declaration of competing interest.

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EW0252

Classification of first-episode schizophrenia spectrum disorders and controls from whole brain white matter fractional anisotropy using machine learning

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Background Schizophrenia is a chronic disorder with an early onset and high disease burden in terms of life disability. Its early recognition may delay the resulting brain structural/functional alterations and improve treatment outcomes. Unlike conventional group-statistics, machine-learning techniques made it possible to classify patients and controls based on the disease patterns on an individual level. Diagnostic classification in first-episode schizophrenia to date was mostly performed on sMRI or fMRI data. DTI modalities have not gained comparable attention.

Methods We performed the classification of 77 FES patients and 77 healthy controls matched by age and sex from fractional anisotropy data from using linear support-vector machine (SVM). We further analyzed the effect of medication and symptoms on the classification performance using standard statistical measures (t -test, linear regression) and machine learning (Kernel-Ridge regression).

Results The SVM distinguished between patients and controls with significant accuracy of 62.34% ($P = 0.005$). There was no association between the classification performance and medication nor symptoms. Group level statistical analysis yielded brain-wide significant differences in FA.

Conclusion The SVM in combination with brain white-matter fractional anisotropy might help differentiate FES from HC. The performance of our classification model was not associated with symptoms or medications and therefore reflects trait markers in the early course of the disease.

Disclosure of interest The authors have not supplied their declaration of competing interest.

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EW0253

Research and practice for ultra-high risk for psychosis: A national survey of early intervention in psychosis services in England

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Objectives Evidence from meta-analyses of randomised clinical trials shows interventions for young people at ultra-high risk (UHR) of developing psychosis are effective both clinically and economically. While research evidence has begun to be integrated into clinical guidelines, there is a lack of research on the implementation of these guidelines. This paper examines service provision for UHR individuals in accordance with current clinical guidelines within the National Health Service (NHS) in England.

Method A self-report online survey was completed by clinical leaders of Early Intervention in Psychosis (EIP) teams ($n = 50$) within the NHS across the UK.

Results Of the 50 EIP teams responding (from 30 NHS Trusts), 53% reported inclusion of the UHR group in their service mandate, with age range predominantly 14–5 years (81%) and service provided for at least 12 months (53%). Provision of services according to NICE clinical guidelines showed 50% of services offered cognitive behavioural therapy (CBT) for psychosis, and 42% offered family intervention. Contrary to guidelines, 50% of services offered antipsychotic medication. Around half of services provided training in assessment by CAARMS, psycho-education, CBT for psychosis, family work and treatment for anxiety and depression.

Conclusions Despite clear evidence for the benefit of early intervention in this population, current provision for UHR within EIP services in England does not match clinical guidelines. While some argue this is due to a lack of allocated funding, it is important to note the similar variable adherence to clinical guidelines in the treatment of people with established schizophrenia.

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e-Poster walk: Schizophrenia and other psychotic disorders—part 2

EW0254

Effects of chronic antipsychotic treatment on neurophysiological correlates of the auditory oddball task in schizophrenia: A preliminary report from a multicentre study

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Introduction The effects of chronic antipsychotic administration on the human brain are debated. In particular, first-generation (FGAs) and second-generation antipsychotics (SGAs) seem to have different impacts on brain function and structure in subjects with

schizophrenia. Few studies have investigated the effect of chronic administration of FGAs and SGAs on indices of brain function, such as event-related potentials (ERP) or neuropsychological performance.

Objectives Within the Italian Network for Research on Psychoses study, subjects stabilized on FGAs or SGAs were compared on P300, an ERP component, thought to reflect attention, working memory and context integration and on neurocognitive indices.

Methods ERPs were recorded in 110 chronic, stabilized patients with Schizophrenia (28 used FGAs) during a standard auditory oddball task. P300 latency and amplitude were assessed at Pz channel. MATRICS Consensus Cognitive Battery (MCCB) was used for cognitive assessment.

Results Compared with the SGAs group, patients on FGAs showed significant increased P300 latency ($P = 0.003$; Cohen's $d = 0.67$) and significant decreased P300 amplitudes ($P = 0.023$; Cohen's $d = 0.38$). The two groups did not differ on psychopathology and MCCB scores. Multiple linear regressions revealed that "FGAs vs. SGAs" ($\beta = 0.298$, $P = 0.002$) and MCCB neurocognitive composite T-score ($\beta = -0.273$, $P = 0.004$) were independent predictors of P300 latency, whereas only age ($\beta = -0.220$, $P = 0.027$) was an independent predictor of P300 amplitude.

Conclusions FGAs seem to affect the functional brain activity more than SGAs, particularly slowing cortical processing. Our results suggest that discrepant findings concerning P300 latency in schizophrenia might be related to the type of antipsychotic treatment used. Longitudinal studies are needed to further address this issue.

Disclosure of interest The authors have not supplied their declaration of competing interest.

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EW0255

Schizophrenia and major depression: Resilience, coping styles, personality traits, self-esteem and quality of life

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Introduction Resilience is commonly defined as positive adaptation to adverse events or as the ability to maintain or regain mental health after exposure to difficulties. According to the bio-psycho-social model, resilience is influenced by self-esteem, coping strategies and personality traits. In schizophrenic patients, resilience seems to affect real-life functioning, while in mood disorders, resilience influences the longitudinal course of the disorder, reducing the frequency of relapses and improving drugs response.

Objectives The aim of this study is to assess levels of resilience and self-esteem, coping strategies, perceived quality of life and temperament characteristics in a sample composed by patients with major depressive disorder and patients affected by schizophrenia.

Methods We collected a sample composed by 40 patients with major depressive disorder and 40 patients affected by schizophrenia patients recruited at the "Maggiore della Carità" Hospital in Novara, Italy. The assessment protocol included: Resilience Scale for Adults (RSA), Coping Orientation to Problems Experienced Inventory—Brief (BRIEF—COPE), Rosenberg Self-esteem Scale (RSES), Paykel List Of Stressful Events, Temperamental and Character Inventory (TCI) and Short form 36 (SF-36). Comparison of qualitative data was performed by means of the χ^2 , a t -test was performed for continuous normal-distribution variables otherwise a non-parametric Mann–Whitney test was performed. Statistical significance was set at $P \leq 0.05$.

