

With regard to biological variables, patients with BD-PS, compared to the counterpart, had a higher Neutrophil to Lymphocyte Ratio (NLR) ($t = 2.776$; $p = 0.006$), lower levels of Gamma-Glutamyl Transferase (γ GT) ($t = 2.249$; $p = 0.026$), higher total bilirubin ($t = 2.348$; $p = 0.019$) and creatine phosphokinase (CPK) ($t = 2.807$; $p = 0.005$), lower total cholesterol ($t = 2.369$; $p = 0.018$) and triglycerides ($t = 2.554$; $p = 0.013$).

Conclusions: Our data appear to be in line with the literature, especially with respect to the occurrence of psychotic symptoms mainly in manic episodes and their association with greater clinical severity, longer hospitalization and worse outcome (Altamura *et al.* Aust N Z J Psychiatry 2019; 53(8) 772-781). From a biological point of view, it seems important to emphasize that patients with lifetime psychotic symptoms presented a higher NLR, revealing more prominent low-grade inflammation in these patients than the counterpart. These data confirm the possibility of using NLR as biomarker of severity in bipolar patients, as proposed previously by other authors (Kulacaoglu *et al.* Nord J Psychiatry 2022). Future multi-center study have to confirm the results of the present study.

Disclosure of Interest: None Declared

EPP0790

Clinical factors associated with unipolar mania: A systematic review and meta-analysis

F. Bartoli¹, C. Nasti^{1*}, D. Palpella¹, S. Piacenti¹, M. E. Di Lella¹, S. Mauro¹, L. Prestifilippo¹, C. Crocamo¹ and G. Carrà^{1,2}

¹Department of Medicine and Surgery, Università degli Studi di Milano - Bicocca, Milan, Italy and ²Division of Psychiatry, University College London, London, United Kingdom

*Corresponding author.

doi: 10.1192/j.eurpsy.2023.1075

Introduction: The existence of a clinical entity on the spectrum of mood disorders characterized by the occurrence of manic episodes without major depressive episodes (Unipolar Mania, UM) is largely debated. Although not classified nosologically, the studies exploring this topic have suggested that UM might differ from bipolar disorder with a manic-depressive course (md-BD), in terms of several clinical characteristics. Individuals with UM might represent a subpopulation with specific clinical profiles and unmet care needs, requiring personalized treatments, as compared with those suffering from md-BD.

Objectives: To identify clinical factors associated with UM, as compared with md-BD.

Methods: We performed a systematic review and meta-analysis of observational studies according to the MOOSE guidelines. We searched for articles indexed up to July 2022 in the main electronic databases. We conducted random-effects meta-analyses of the association between UM and relevant correlates, using odds ratio for categorical variables and standardized mean difference for continuous variables.

Results: Based on data from 21 studies meeting the eligibility criteria, we found that individuals with UM, as compared with md-BD, were more likely to be males ($p = 0.007$) and to have an earlier age at onset ($p = 0.020$). Moreover, UM was significantly associated with a higher number of hospitalizations ($p < 0.001$), the occurrence of psychotic features ($p < 0.001$), as well as hyperthymic temperament ($p = 0.012$). Finally, subjects with UM were less likely to report a family history of depression ($p = 0.006$) and a personal history of suicide attempts ($p < 0.001$).

Conclusions: Our work supports the hypothesis that UM might represent a distinctive diagnostic construct, with peculiar clinical correlates. Additional research is needed to better differentiate UM in the context of affective disorders.

Disclosure of Interest: None Declared

EPP0791

Sleep spindle and slow wave activity in Bipolar Disorder: preliminary observations from a high-density EEG study

C. Sanguineti^{1*}, F. L. Donati¹, M. Sala², C. Carrara¹, C. Casetta³, C. Zangani⁴, A. Mayeli⁵, A. Castelnovo⁶, M. P. Canevini^{1,7} and A. D'Agostino^{1,8}

¹Department of Health Sciences; ²Department of Pathophysiology and Transplantation, Università degli Studi di Milano; ³San Carlo Hospital, ASST SS. Paolo e Carlo, Milano, Italy; ⁴Warneford Hospital, Department of Psychiatry, University of Oxford, Oxford, United Kingdom; ⁵Department of Psychiatry, University of Pittsburgh, Pittsburgh, United States; ⁶Sleep Center, Neurocenter of Southern Switzerland, Civic Hospital of Lugano, Lugano, Switzerland; ⁷San Paolo Hospital, Epilepsy Center - Sleep Medicine Center, Childhood and Adolescence Neuropsychiatry Unit, ASST SS. Paolo e Carlo and ⁸Department of Mental Health and Addiction, ASST Santi Paolo e Carlo, Milano, Italy

*Corresponding author.

doi: 10.1192/j.eurpsy.2023.1076

Introduction: Recent research on Schizophrenia (SCZ) suggests that reduced sleep spindle and slow wave density could be particularly informative of underlying thalamocortical and cortical synchronization mechanisms and dysfunctions. Although sleep disturbances are also highly prevalent across all stages of Bipolar Disorder (BD), the objective evaluation of sleep macrostructure and microstructural oscillatory activity remains understudied in this population.

Objectives: We aimed to investigate sleep EEG activity in BD, with a focus on sleep architecture, sleep spindles and slow waves.

Methods: We recorded high-density EEG (64-channel BrainAmp, Brain Products GmbH, Germany) during sleep in 18 euthymic patients with BD and 18 age/gender-matched healthy control (HC) subjects. After sleep scoring and EEG artifact rejection, several parameters of sleep spindles (12-16 Hz), including density and amplitude, and slow waves (0.1-4 Hz) were identified for the first cycle of sleep using automated algorithms and compared between groups using non-parametric statistics.

Results: BD subjects showed significantly higher Wake After Sleep Onset and lower Sleep Efficiency (Table 1). Total (12 - 16 Hz), slow (12 - 14 Hz) and fast (14 - 16 Hz) sleep spindle parameters of density (Image 1) and amplitude did not differ significantly between groups. On the other hand, slow wave density was reduced in a large frontal cluster of electrodes in the BD group (Image 2).

Image:

Table 1

	BD (n = 18)	HC (n = 18)	Difference (p value)
WASO (min ± sd)	140,61 ± 74,23	84,34 ± 59,84	0,017
Sleep efficiency (% ± sd)	72,47 ± 14,33	82,43 ± 11,58	0,028