

## **P-362 - BRAIN DERIVED NEUROTROPHIC FACTOR, COGNITION AND FUNCTIONING OF PATIENTS WITH FIRST PSYCHOTIC EPISODE**

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**Introduction:** Brain-derived neurotrophic factor (BDNF) promotes growth and maintenance of connections and participates in plasticity mechanisms. The cognition and the functioning of patients with a first episode of psychotic (FEP) is altered.

**Objectives:** We analyze the relation between the BDNF, the cognitive performance and prognosis in patients with FEP.

**Design and Methods:** 45 patients with a FEP from the Basque Country, diagnosed using the SCID-I and DSM-IV. Plasma BDNF levels were measured using the BDNF Sandwich ELISA Kit. All patients were assessed clinically three times over a year using PANSS, GAF and Strauss Carpenter scales. Battery of cognitive tests (Wechsler Memory Scale and WAIS-III) was applied six months after the acute episode.

**Results:** Positive correlation between BDNF levels after six months of treatment and five cognitive domains: abstract verbal reasoning ( $r=0.468$ ), motor and processing speed ( $r=0.397$ ), learning capacity ( $r=0.559$ ), immediate memory ( $r=0.409$ ) and delayed memory ( $r=0.382$ ). Also, the patients with lower BDNF plasma levels at baseline, at 6 months follow-up had worse social activity (0.61 vs. 0.89;  $p=0.041$ ) and functioning (0.69 vs. 0.93;  $p=0.044$ ). The BDNF levels increased along the follow up, after the pharmacological treatment (basal-1 month:  $Z= -2.88$ ;  $p\leq 0.004$  and 1-6 months:  $Z= -2.23$ ;  $p\leq 0.05$ ).

**Conclusions:** Our results suggest that BDNF is associated with cognitive impairment seen after a FEP and their prognosis. After the pharmacological treatment, the BDNF levels increase significantly and at 6 months of treatment there were normal levels. Further investigations of the role of this neurotrophin in the symptoms associated with onset of psychosis are warranted.