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ACTIVATION OF SEROTONIN METABOLISM AND BDNF CHANGES IN DEPRESSED PATIENTS WITH MULTIPLE SCLEROSIS DURING INTERFERON-BETA THERAPY

S. Bonaccorso¹, A. Ricciardi¹, F. Matrisciano², F. Nicoletti², S. Scaccianoce², A. Ruberto², P. Girardi², R. Tatarelli², F. Pompei², R. Manciocchi², C. Pozzilli², R. Verkerk³, S. Scharpe³, L. Wang⁴, R. Shelton⁴

¹*Institute of Psychiatry at King's College, London, UK*, ²*University of Rome 'La Sapienza', Rome, Italy*, ³*University of Antwerp, Antwerpen, Belgium*, ⁴*Vandebilt University, Nashville, USA*

Background: Indoleamine-2, 3-dioxygenase is responsible for tryptophan catabolism; IFN- β is a treatment for multiple sclerosis (MS) and it has been associated with depression. IDO activation might play a role in IFN- β induction of depressive symptoms. Depressive symptoms associated with MS illness and IFN- β treatment have been treated with pharmacological and non-pharmacological intervention.

Aims: Evaluation of the kynurenine pathway, IDO activation and neurotoxic agents, serum BDNF in MS patients during IFN- β intervention.

Methods: 14 study subjects, aged 18-64 years with major depressive disorder and MS treated with IFN- β , before and after pharmacologic and non-pharmacologic intervention, and 34 age matched healthy controls were enrolled at the University of Rome "La Sapienza" and at the University of Antwerp. Depressed participants were randomized to sertraline-treatment or interpersonal psychodynamic psychotherapy.

Results: There were significant improvements in both depression and anxiety symptoms with medication and psychotherapy groups, although the effect with sertraline was more robust. Sertraline treatment was associated with a decline in serotonin. In the psychotherapy group, a significant increase in 3-hydroxyanthranilic acid (HAA) was observed after 4 months treatment ($p = .04$) with a significant decrease in tryptophan levels at 6 weeks ($p = .02$) and a trend towards a significant decrease in BDNF after 6 weeks ($p = 0.06$), neither of which were seen in the medication condition.

Discussion: The increase in HAA among the psychotherapy-treated patients raises the possibility of a neurodegenerative challenge for patients with multiple sclerosis during treatment with IFN- β which appeared to be prevented with pharmacological treatment.