

The average measurements of vital signs during 2 hours of monitoring for each application can be seen in Tab 2:

Conclusions: The use of SC ketamine showed remission in BDI, BSI and BAI, respectively demonstrated safety in use.

Disclosure of Interest: None Declared

EPV0843

Serotonin reuptake inhibitors and its cognitive burden...or relief.

A brief review

M. J. Amorim*, F. Araújo, P. Passos Perestrelo and M. Maia Marques

Departamento de Psiquiatria e Saúde Mental, Unidade Local de Saude do Alto Minho, Viana do Castelo

*Corresponding author.

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Introduction: The use of selective serotonin reuptake inhibitors (SSRIs) has increased exponentially and worldwide in the last decade. Taking into account their tolerability and safety profile, they constitute the first line, in all age groups and in polymedicated population, for treatment of depressive, anxious, and phobic disorders, among others. Although safe, especially when compared to first generation antidepressants, they are not totally exempt of adverse effects, and may cause, especially in this context, some degree of cognitive impairment, which may or may not be completely reversible. On the other hand, and despite the controversy related to the subject, some studies suggest a possible protective effect of this pharmacologic class regard the development of cognitive disfunction, which, although not very consistent, should not be neglected.

Objectives: To understand the cognitive impact of short- and long-term use of SSRIs, given the increasing use in an aging, polymedicated population.

Methods: Brief sistematic review of selected articles from Pubmed, Medline and Uptodate databases, with the keywords "SSRIs", "cognitive", "impairment", "adverse effects", "dementia".

Results: SSRIs are not entirely exempt from cognitive effects, despite their recognized safety profile. Some of the adverse effects typically related to the class, such as hyponatraemia, especially when insidious and silent course, as well as anticholinergic activity (typically associated with first generation antidepressants, but not exclusive) interfere with global functionality and may clinically present as mild cognitive impairment or even major neurocognitive disorders. Furthermore, given their potential to induce or inhibit the cytochrome P450 system, they are significantly implicated in pharmacokinetic drugs interactions that increase cognitive burden associated with other psychotropic drugs.

In the long term, and in certain patient populations, it has been hypothesized that they may exert some protective effect on physiological and pathological cognitive functions decline, by preventing neuronal death, acetylcholine decrease and amyloidogenesis.

Conclusions: Despite their benign adverse effect profile, when compared with tricyclic antidepressants, SSRIs are not devoid of

adverse effects on cognitive domains. However, and despite contradictory and inconsistent results, when well tolerated, SSRIs may confer benefits in terms of preserving global functionality, far beyond the affective symptoms resolution that motivated their introduction in the first place.

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EPV0844

DOCTOR, I'M PREGNANT. Psychopharmacological treatment of depression in pregnant women. A clinical case of a pregnant woman and major depressive disorder

M. Queipo De Llano De La Viuda*, G. Guerra Valera, C. Vallecillo Adame, C. De Andrés Lobo, T. Jiménez Aparicio, M. Fernández Lozano, I. D. L. M. Santos Carrasco and N. De Uribe Viloria

Hospital clinico Universitario de Valladolid, Valladolid, Spain

*Corresponding author.

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Introduction: Depression during pregnancy can appear with a prevalence of up to 11% of pregnant women. Psychotherapeutic treatment in these cases is considered the first option, but treatment with antidepressants is sometimes required in these cases.

Objectives: To present a clinical case of a pregnant patient diagnosed with depression.

Methods: Literature review of the psychopharmacological treatment of depression during pregnancy and possible complications.

Results: A 25y Year old woman, 22 weeks pregnant, who lives with her partner. She has no background in mental health. Paternal aunt diagnosed with type I Bipolar Disorder. She goes to the Mental Health Center for evaluation, due to anxiety and depressive symptoms of 4 weeks of evolution, she refers sadness and apathy, continuous crying, somatic anxiety and obsessive ruminations in relation to childbirth and inability to care for your child. Suicidal ideation as a resolution of her discomfort. She presents with global insomnia and a significant loss of appetite, with a weight loss of 3 kg. Treatment with sertraline 50 mg/day was started, with good tolerance and clinical response

Conclusions: The psychopharmacological treatment of antenatal depression is a challenge for the psychiatric professional. In all cases, an adequate balance must be made between the risks and complications for the fetus and the psychopathological stability of the pregnant woman. Among the main risks of untreated depression are: preterm delivery and low birth weight, an increased risk of suicide and alterations in the development during the baby's infancy. The most used antidepressants are the SSRIs, with sertraline being a good option. Paroxetine has been associated with cardiac defects in the newborn. There are studies with tricyclics and duals but no specific teratogenic pattern has been seen. They are associated with an increased risk of spontaneous abortion. Exposure during the third trimester may be associated with obstetric complications.

Disclosure of Interest: None Declared