

اتحاد الأطباء النفسيين العرب Arab Federation of Psychiatry

The Arab Journal of Psychiatry المجلة العربية للطب النفسي

Vol. 23 No 2. November 2012 المجلد 23، العدد الثاني، نوفمبير 2012



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Papers are submitted in electronic form

- O Title, running head (Max: 40 letters), title of the article in EnglishandArabic, the names of authors should be without their titles and addresses in both languages.
- O Abstract in English (max: 200 words). It should follow a structured format (objectives, method, results and conclusion). It should be followed by key words (max. 5).
- O Declaration of interest after the key words.
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- Acknowledgment of support and persons who have had major contribution to the study can be included after the references.
- Arabic abstract like the English abstract should follow a structured format. And it follows the references section (last page).
- All Pages should be numbered.

Tables

Tables should be typed with double-spaced in separate pages. They should be numbered with Arabic (e.g1, 2, 3) numerals and have a short descriptive headings.

Illustrations

All illustration should be submitted camera-ready; line drawings/diagrams should be approximately twice the size they will appear in print.

Reference List

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Example of references:

- O Zeigler FJ, Imboden, JB, Meyer E. Contemporary conversion reactions: a clinical study. Am. J. Psychiatry 1960: 116:901 10.
- O Mosey AC. Occupational therapy. Configuration of a profession. New York: Raven Press, 1981.

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Editorial Letter

Dear Colleagues

This issue will appear with the 12th pan Arab Conference on psychiatry in Dubai, an occasion that we wait for; to have all the members of the Arab Federation of psychiatrists meet and develop the federation and the Journal.

Continuous efforts to improve the Journal and indexing it are going on.

Walid Sarhan

November 2012

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Letter to the editor

Subject: Re: prevalence and correlates of physical and sexual history in patients with schizophrenia Arab journal of psychiatry 2012, 23 (22-29).

Dear Sir,

This was a very carefully designed study, and in my opinion it corrects a popular misconception regarding a correlation between sexual abuse and the later development of schizophrenia. The results - as stated in the next paragraph – are consistent with my own findings. To quote their findings directly:

"Our findings around the lack of significant association of previous exposure to abuse with psychotic symptoms replicated previous reports. 15,43,44On the contrary, it contradicts previous studies that reported significant associations of abuse with positive symptoms and lowered functioning in patients with schizophrenia. This finding deserves special consideration since it suggests that sexual and physical abuse alone did not show influence over clinical picture or disability in our study. These two kinds of trauma are likely to affect children later in life in relation to other traumas such as emotional and physical neglect and emotional abuse, which is not measured in this study."

Yes. Other studies immediately search for and "find" a correlation between early sex abuse and the later development of schizophrenia, but the original presumption is incorrect. The age of origin of schizophrenia is in the first 18 months of life, schizoaffective disorder is 19-21 months, bipolar hypomania peaks at 22 months, the remainder of the psychotic depressions up to 24 months, and non-psychotic depressions between 24 and 34 months.

There could be co-morbidity, in that other kinds of infant separation traumas at an early age could be associated with abuse at a later age, etc.

What people do not realize is that more overwhelming than war trauma to a soldier is separation from mother to a baby, because, for as long as mammals have populated the earth, separation from mother has meant death. Then years or decades later, instead of a loud noise precipitating the flashback, it is separation or rejection from some other "most important person" (husband, wife, girlfriend, boyfriend) which precipitates the initial step back in time, and instead of combat reality and behavior it is infant reality and behavior that we see.

My earlier studies began in 1966 - and extended over a course of about 20 years. Patients who were on average 20 years old at that time, had mothers who spent 5 days in the hospital following delivery of the next baby -which was overwhelming to the older child. The first two studies were sequential and therefore cumulative. Thirteen with schizophrenia or psychotic depression were traumatized prior to 24 months and 14 with non-psychotic depression were traumatized between 24 and 34 months. That's one over two to the 27th power, or one chance in 134,217,728 by chance alone. An unbelievable number! But PTSD cannot occur without original trauma, and I just happened to identify a trauma for which we knew the exact age when it occurred. Further calculations revealed an estimate of about one chance in 50 that another trauma would have been responsible for the origin of the disorder.

My work further supports the above referenced article in that it reveals twelve precise parallels between posttraumatic stress disorder from adult life and post-traumatic stress disorder from infancy (schizophrenia, depression), and it reveals, for the first time, the derivation of the precursors of schizophrenia, the negative symptoms of schizophrenia and the derivation of the positive symptoms of schizophrenia.

Clancy D. McKenzie, MD

USA

Dear Editor

Re: The prevalence of mental health symptoms among outpatients in the UAE

I was happy to read this article but I have my own reservations about the method and even some of the results.

This study, considering the small number of subjects, can only be considered a pilot one and only a first step before doing a much larger study. However, there is some originality in the way it was conducted.

I believe that using and elaborating on the several standardized assessment instrument used in this study would prove useful particularly for junior mental health care professionals.

Some of the results were rather unexpected and unusual e.g. the very high prevalence of PTSD (42.9%) in a country not known to have stuffed from wars or disasters. Even research from Iraq, a country which has suffered from several wars has not reached that level of prevalence.

Another problem is the high prevalence of Alcohol. This is likely to cause selection bias and diagnosis problems.

A third problem is the fact that all participants have had at least 3 sessions of psychological therapy. This is also likely to affect the presentation of symptoms and diagnoses.

However, in spite of all that I believe this study is a useful one and empowers research in UAE.

Dr. Ali Alqam

Amman-jordan

قسيمة الاشتراك

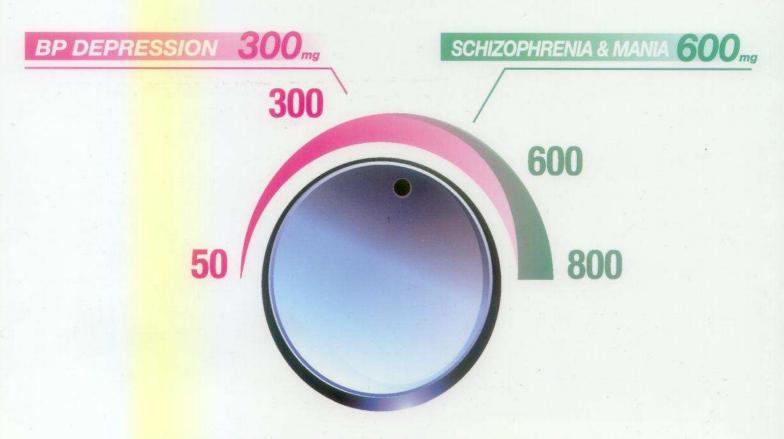
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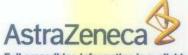
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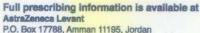
Seroquel XR, a simplified therapy with efficacy in Bipolar Depression, Mania and Schizophrenia



(*) For the treatment of manic episodes associated with bipolar disorder the daily dose of therapy is up to 800 mg after day 2 Reference: Seroquel XR prescribing informations

Indication: Schizophrenia including: preventing relapse in stable schizophrenic patients who have been maintained on Seroquel XR Bipolar disorder including: manic episodes associated with bipolar disorder, major depressive episodes in bipolar disorder, preventing recurrence in bipolar disorder in patients whose manic, mixed or depressive episode has responded to quetiapine treatment. Contraindications: hypersensitivity to the active substance or to any of the excipients of this product. Concentration of cytochrome P450 3A4 inhibitors, such as HIV protease inhibitors, accide antifungal agents, erythromycin, clarithromycin and netazodone, is contraindicated. Special warnings and precaution for use: Suicide / suicidal thoughts or clinical worsening; Somnolence; Cardiovascular disease; Cerebrovascular disease; Conditions predisposing to hypotension; Family history of QT prolongation; Medicines know to increase QTc interval; Concomitant neuroleptics; Congenital long QT syndrome; Congestive heart failure; Heart hypertrophy, Hypokalaemia; Hypomagnesaemia; Seizures; Extrapyramidal symptoms Tardive Dyskinesia; Neuroleptic Malignant Syndrome; Severe neutropenia; Interactions: Hyperghycaemia; Lipids; Withdrawal; Elderly patients with dementia-related psychosis; Hepatic effect; Dysphagia; Aspiration pneumonia; Venous thromboembolism; Lactose intolerance; Children and adolescents (10 to 17 years of age) Interactions: Centrally acting drugs and alcohol; Grapefruit juice; Hepatic enzyme inducers (carbamazepine and phenytoin). Adverse reactions: the most commonly reported Adverse Drug Reactions (ADRs) with quetiapine are somnolence, dizziness, dry mouth, mild asthenia, constipation, tachycardia, orthostatic hypotension and dyspepsia. As with other antipsychotics, weight gain, syncope, neuroleptic malignant syndrome, leucopenia, neutropenia and peripheral oedema, have been associated with quetiapine. Posology and method of administration: Seroquel XR should be administrated once daily, without food (at least one hour before a meal). The tablets should be swallowed whole and not split, chewed or crushed. Adults: for the treatment of schizophrenia the daily dose the start of therapy is 300 mg on day 1 and 600mg on day 2. The recommended daily dose is 600 mg. for the treatment of manic episodes associated with bipolar disorder the daily dose at start of therapy is 300 mg on day 1, 600mg on day 2 and up to 800 mg after day 2 for the treatment of depressive episodes associated with bipolar disorder Seroquel XR should be administrated once daily at bedfirm. The total daily dose for the first four days of therapy is 50 mg (day 1), 100mg (day 2), 200mg (day 3) and 300mg (day 4). The recommended daily dose is 300 mg, for preventing recurrence in bipolar disorder for prevention of recurrence of manic, depressive or mixed episodes in bipolar disorder, patients who have responded to Seroquel XR for acute treatment of bipolar disorder should continue on Seroquel XR at the same dose administrated at bedtime. Switching from seroquel immediate-release tablets: For more convenient dosing, patients who are currently being treated with divided doses of immediate release Seroquel tablets (Seroquel[®]) may be switched to Seroquel XR at the equivalent total daily dose taken once daily. To ensure the maintenance of clinical response, a period of dose titration may be required.





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