

sidering patient's symptoms, such as tachycardia, slightly increased body temperature, subjective chest pain, dyspnea. However, this symptomatology is not always present in a clozapine-related pericarditis. Some authors suggest measuring BNP levels to detect early and asymptomatic cardiac dysfunction. We here report the clinical cases of two women, respectively 22 and 28 years old. They both suffered from an early onset resistant schizophrenia. Clozapine was gradually introduced, at a dose of 200 mg/day, in both patients. After about one month in both cases, while the first patient was nearly asymptomatic, apart from the intermittent fever (only PCR and pro-BNP values were elevated, 16.88 mg/dL and 1004 pg/mL, respectively), the second one showed a classic symptomatology suggestive of pericarditis. Clozapine was discontinued in both patients, resulting in progressive resolution of pericarditis. Interestingly, in the patient in which pro-BNP was elevated, after clozapine cessation, the pro-BNP fell down dramatically. Pro-BNP plasma levels appears to be an interesting test in identifying subjects with asymptomatic cardiac impairment. It would be useful to evaluate if early treatment with beta-blockers and ACE-inhibitors may allow the prosecution of clozapine treatment after developing of mild signs of cardiac toxicity in drug resistant schizophrenic patients responsive to clozapine.

Disclosure of interest The authors have not supplied their declaration of competing interest.

<http://dx.doi.org/10.1016/j.eurpsy.2016.01.2029>

EV1045

Pseudoakathisia in a patient with clotiapine abuse: Report of a case

P. Quandt*, M.D.R. Cejas Méndez

Hospital Universitario de Canarias, Psiquiatría, La Laguna, Spain

* Corresponding author.

Introduction Objective symptoms of akathisia in the absence of subjective symptoms is known as pseudoakathisia, more often diagnosed in older patients with long-term antipsychotic treatment.

Objective To describe a case of pseudoakathisia in a patient with clotiapine abuse.

Aims Pseudoakathisia management.

Methods X is a 47-year-old male with chronic insomnia treated with clotiapine 40 mg/day for four years. He admits abusive neuroleptic consumption in the past eight months (160 mg/day), without any psychiatric control for years. In recent months he has experienced different organic complications, requiring multiple hospitalizations. During psychiatric examinations due to confusional states, repeated lower limbs movements were objectified. X reported he presented these movements for at least six months, without complaints of inner restlessness feeling. Neurological examination showed normal DAT-SCAN result. Clinical progression was evaluated using BARS scale (Barnes Akathisia Rating Scale).

Results Following the results of tests and statements of drug history, X was diagnosed with clotiapine-induced pseudoakathisia. Neuroleptic treatment was suspended, and clonazepam 6 mg/day and propranolol in ascending doses up to 80 mg/day were initiated. In subsequent evaluations, progressive decrease in movement intensity was observed. However, complete remission after four months from clotiapine suspension was not achieved.

Conclusions Pseudoakathisia is a concept not well defined at this moment and different hypotheses about its nature are considered. It has been suggested that it is a form of delayed dyskinesia, or a clinical progression from akathisia, with acquired subjective discomfort tolerance. The most widely used treatment includes benzodiazepines, beta-blockers and anticholinergics, although their effectiveness is limited.

Disclosure of interest The authors have not supplied their declaration of competing interest.

<http://dx.doi.org/10.1016/j.eurpsy.2016.01.2030>

EV1046

Reduction in medication expenditure: Review of strategies at a children's psychiatric facility

B. Reddy*, M.W. Azeem, J. Smiles, L. Carrabetta

Albert J. Solnit Children's Center, Child and Adolescent Psychiatry, Middletown, CT, USA

* Corresponding author.

Prescription drug costs rise about 15% annually. Solnit Center has been exploring ways to reduce overall expenditure on medications while promoting best practice of care. Lack of facility-based pharmacy has posed specific challenges in ordering medications, optimal usage and minimizing wastage of drugs. Each of these areas were examined and reviewed at Pharmacy and Therapeutics Committee of the facility. This information was shared with the ordering physicians and standard prescribing practices were established. This project was aimed at tracking medication costs over a 11-year period while monitoring supplies and destruction of unused medications.

Aims 1. Reduce overall medication expenditure while maintaining standard of care. 2. Develop a program to return unused medications for refund.

Methods 1. Monthly review of pharmacy cost by facility, patient and medication. 2. Development and legislative approval of a program to return drugs. 3. Collaborate with contracted pharmacy to explore ways to cut costs. 4. Train nurses and physicians to understand optimal ordering practice. 5. Demonstrate medications wasted with associated financial impact to the facility.

Results The expenditure to the facility over 11 years has gradually decreased despite increase in medication costs. In 2004, the facility spent \$ 712,904 and in 2014, the expenditure was \$ 584,022.

Conclusions Awareness about costs and optimal ordering practices led to significant savings to the facility.

Disclosure of interest The authors have not supplied their declaration of competing interest.

<http://dx.doi.org/10.1016/j.eurpsy.2016.01.2031>

EV1047

Mega-review of meta-analyses investigating the short-term efficacy of pharmacologic augmentation strategies of antipsychotics in patients with schizophrenia

J.M. Rubio^{1,*}, G. Inczedy-Farkas¹, S. Leucht², J.M. Kane^{1,3}, C. Correll^{1,3}

¹ Zucker Hillside Hospital, Department of Psychiatry, Glen Oaks, NY, USA

² Klinikum rechts der Isar, Technischen Universität München, München, Germany

³ Feinstein Institute for Medical Research, Department of Psychiatry, Manhasset, NY, USA

* Corresponding author.

Antipsychotics are the cornerstone of treatment for schizophrenia, but they have limited effectiveness, as most patients require subsequent strategies at some point of their treatment. Despite being widely used, the efficacy of pharmacologic augmentation of antipsychotics is controversial and no combination treatment has been approved for schizophrenia. We conducted a systematic review in PubMed and PsycInfo on June 1st 2015 and a random effects meta-analysis of meta-analyses of short-term, placebo-controlled studies of pharmacological augmentation strategies of antipsychotics in schizophrenia. Methodological quality of meta-analyses was measured using the AMSTAR, plus 6 additional items developed to rate the content quality of the meta-analyzed trials. Out of 3062 publications, we identified 36 eligible augmenting

strategies. For total symptom reduction, 25 strategies augmenting antipsychotics and 5 strategies augmenting clozapine were eligible and examined. Eleven strategies were more efficacious than placebo, none of them augmenting clozapine. Significant effect sizes ranged between SMD -1.03 and -0.23 . Efficacy was not correlated with the quality of the meta-analyses. Only the meta-analysis for NSAIDs augmentation had a score greater than half of the possible points for content quality. Only antipsychotics, azapirones, antidepressants and lithium were less discontinued than placebo. Serotonin-3-receptor antagonists, lamotrigine, mirtazapine/mianserine, minocycline and estrogens had large effect sizes augmenting antipsychotics. However the quality of the content of most meta-analyses was low. The NSAIDs augmentation meta-analysis had the best content quality, yet with a low effect size for efficacy. The evidence for short-term augmentation strategies of antipsychotics in schizophrenia is inconclusive, due to the limited quality of the available trials.

Disclosure of interest The authors have not supplied their declaration of competing interest.

<http://dx.doi.org/10.1016/j.eurpsy.2016.01.2032>

EV1048

California rocket fuel: And what about being a first line treatment?

J. Silva^{1,*}, J. Mota², P. Azevedo¹

¹ Magalhães Lemos Hospital, Inpatient C Unit, Porto, Portugal

² Magalhães Lemos Hospital, Inpatient C Unit, Electroconvulsive therapy Unit, Porto, Portugal

* Corresponding author.

Introduction The association venlafaxine-mirtazapine is currently known as California Rocket Fuel (CRF). Studies show advantage in terms of efficacy and rapid control of depressive symptoms compared to other associations. Venlafaxine is a selective serotonin-noradrenalin reuptake inhibitor and mirtazapine is a noradrenergic-specific serotonergic antidepressant: the result is a potent noradrenergic and serotonergic effect. Studies say that CRF should be performed only for drug-resistant depression; however, there are case reports of its use as a first line treatment, in selected patients.

Objectives To summarize the latest literature about this field and to present a case report.

Aim To explore and critically review the controversies of venlafaxine-mirtazapine association as a first line antidepressant strategy.

Methods A brief review of the latest literature was performed, using PubMed and the keywords “venlafaxine-mirtazapine association”. A case report about a depressed woman is presented.

Results Despite most studies are referent to its utility in drug-resistant depression, there are recent pilot studies that recommend CRF as a first line option.

M., a 64-year-old woman, had her first psychiatric consultation. She had been depressed for 2 years, she lost 10 kg, had total insomnia and suicidal thoughts. CRF was started up to 150/15 mg, daily. An improvement was noticed after two weeks of treatment and the stabilization of depressive symptoms were achieved by the fourth month.

Conclusions CRF seems to be effective and useful. Patients with insomnia and weight loss may benefit from CRF as a first line option. However, more studies are needed.

Disclosure of interest The authors have not supplied their declaration of competing interest.

<http://dx.doi.org/10.1016/j.eurpsy.2016.01.2033>

EV1049

The impact of tobacco smoking in patients taking long-action injection drugs – A retrospective comparative study between haloperidol and risperidone

J. Silva^{1,*}, H. Prata-Ribeiro²

¹ Magalhães Lemos Hospital, Inpatient C Unit, Porto, Portugal

² Júlio de Matos Hospital, General and Transcultural Psychiatry Unit, Lisbon, Portugal

* Corresponding author.

Introduction Smoking rate seems to be higher among patients with schizophrenia, comparing to other psychiatric entities, mainly in those who are on typical antipsychotics. Tobacco is known to have enzyme inducer properties, due to cytochrome P450 complex activity: CYP1A1, CYP1A2, CYP2E1 and CYP2D6. CYP2D6 and CYP1A2 play an important role in antipsychotics metabolism, mainly in the first generation ones, like haloperidol, despite its importance in risperidone metabolism.

Aim To analyze the importance of tobacco smoking in patients taking long-action injections.

Objectives To investigate how sexual dysfunction varies with tobacco smoking, in patients taking long-action injections.

Methods Individuals from both sexes, from 18 to 55 years old, taking antipsychotic long-action injections, answered the Arizona Sexual Experience Scale (ASEX).

Results In the studied population ($n = 44$), there were 20 individuals on haloperidol and 24 individuals on risperidone. In a total of 18 (40.9%) positive results for sexual dysfunction, 6 were on haloperidol (30%), 12 (50%) were on risperidone. Seventeen individuals of the 20 who were on haloperidol were smokers, but only 4 were considered to have sexual dysfunction, 35.3%; 12 of the 24 individuals who were on risperidone were smokers, but only 5 were considered to have sexual dysfunction, 41.7%.

Conclusions Patients treated with haloperidol smoke more, comparing to risperidone. Sexual dysfunction is more frequent in patients taking risperidone than in patients taking haloperidol. This data supports that CYP2D6-CYP1A2 induction by tobacco, mainly interacts with haloperidol, which may be helpful for patients to try less side effects.

Disclosure of interest The authors have not supplied their declaration of competing interest.

<http://dx.doi.org/10.1016/j.eurpsy.2016.01.2034>

EV1050

Interferon-induced psychosis: Myth or reality?

A. Antunes*, L. Sousa

Hospital de Santa Maria, Lisbon Academic Medical Center, Psychiatry and Mental Health, Lisbon, Portugal

* Corresponding author.

Introduction Interferon combined with ribavirin is widely used to decrease the burden of Hepatitis C virus (HCV), but some serious side effects might limit its usefulness. There has been recently a growing awareness about neuropsychiatric complications of many drug treatments. Anecdotal case reports of HCV treatment induced psychosis were published but there seems to be no consensus about the causative relation and no systematic reviews were done to the date.

Objective To describe a paradigmatic case that was managed as an iatrogenic psychotic episode following interferon associated with ribavirin.

Aim Call attention to problems that interfere with the recognition, diagnosis and management of drugs induced psychosis.

Methods Bibliographic research was conducted through the PubMed in the Medline library and clinical information was