Introduction The discovery of mirror neurons, considered to be responsible for empathy, intrigued researchers all over the world. Many studies have been developed associating mirror neurons to the incidence of Autism Spectrum Disorder (ASD).

Objective Identifying a possible influence of mirror neuron in autism.

Aims Reviewing the recent trajectory of neuroscience in relation to the connection of impaired mirror neurons in autism.

Method Bibliographical review of studies in English, published in SciELO and LILACS databases, between 2008 and 2013. The keywords used were: autism, brain, cortex and mirror neuron.

Results Different subjects explored the influence of mirror neurons in autism as shown below (Fig. 1). Among 17 studies, 12 were bibliographical reviews and 5 involved experiments. Seventy-six percent of the studies were favorable to the influence of these neurons, while 24% were not.

Conclusion There was a balance in the distribution of themes explored in the articles and few studies exploring the role of mirror neurons in autism. Even though the current research may not be conclusive, it can be said that currently neuroscientists tend to agree that mirror neurons significantly influence ASD. Recent studies suggest that, if properly stimulated, ASD individuals can develop their social skill and, consequently, be socially inserted. According to most author studied, technological development is needed in order to enable scientific advances involving mirror-neurons and ASD.

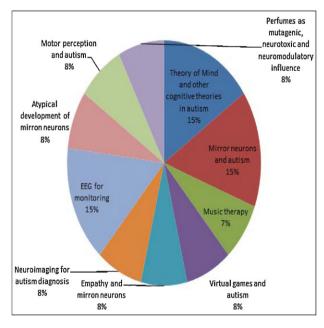


Fig. 1 Percentage of themes explored in the studies.

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Comorbidity/Dual pathologies

EV362

Tardive dyskinesia: When one should suspect of another diagnosis?

R. Almendra ^{1,*}, A.R. Figueiredo ², V. Espirito Santo ¹, A. Almeida ¹, P. Guimarães ¹, A.G. Velon ¹

- ¹ Centro Hospitalar Trás-os-Montes e Alto Douro, Neurology, Vila Real, Portugal
- ² Centro Hospitalar Trás-os-Montes e Alto Douro, Psychiatry, Vila Real, Portugal
- * Corresponding author.

Introduction Tardive dyskinesia is a collection of symptoms related to the side effects of neuroleptic medications that can mimic other types of disorders. Accurate diagnosis can be challenging, as there is no single test for tardive dyskinesia.

Case report Female patient, 64 years old, with personal history of Chronic Myeloid Leukaemia and psychosis since forth decade, currently medicated with quetiapine 350 mg/day, risperidone IM 50 mg 15/15 days and trazodone 150 mg/day (previously medicated with haloperidol, amisulpride and olanzapine). She started with involuntary movements interpreted as tardive dyskinesia after 2 years on neuroleptic treatment. The difficult control of involuntary movements motivated the reference to ambulatory Neurology department. The review of personal history suggested a family history of involuntary movements and psychiatric illness. Physical examination showed generalized choreic movements. The analytical and imagiologic study was unremarkable. The presence of family history and involuntary movements atypical to be classified as tardive dyskinesia supported a genetic test for Huntington's disease who detected a CAG expansion with 43 repetitions in HTT gene. Despite treatment with amantadine and riluzole she maintains disease progression and evident cognitive deterioration.

Conclusion The diagnostic process of involuntary movements may involve more than one physician and requires the review of a detailed medical history, a physical examination and a neuropsychological evaluation in order to determine whether one is indeed suffering from tardive dyskinesia or a different neurological disorder.

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

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EV363

Toxic consumption among patients suffering delusional disorder

C.M. Carrillo de Albornoz Calahorro*, A. Porras, M. Guerrero, J. Cervilla Ballesteros

Hospital Universitario San Cecilio, Unidad de Salud Mental, Granada, Spain

* Corresponding author.

Introduction Several epidemiological studies describe the association between substance abuse and appearance of psychotic symptoms. There is a higher prevalence of psychotic symptoms among cannabis and cocaine consumers compared to the general population.

The cannabinoid receptors regulate the release of dopamine and cocaine has a strong inhibitory action on reuptake of the same. This may explain the greater proportion of subjects moderately or heavily dependent on cocaine or cannabis experience symptoms of psychotic sphere.

Objectives/Aims Describing the profile of drug consumption among a group of patients diagnosed with delusional disorder.

Methods Our data come from a case register study of delusional

disorder in Andalucía (Spanish largest region). By accessing digital

² University of São Paulo, Department of Physical Therapy, Speech-Language Pathology and Audiology, and Occupational Therapy, São Paulo, Brazil

^{*} Corresponding author.