

Last, if compared to healthy controls, children and adolescents after cleft lip and palate (CLP) repair were not at risk reporting sleep difficulties; rather, irrespective of the presence of CLP, sleep was affected by psychological strain.

S04.03

Sleep regulation and cognitive performance in elderly subjects with dementia and depression

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In elderly patients who suffer from depressive symptoms and cognitive impairment the clinical decision between the diagnoses of depression and dementia may be difficult. In addition, patients with dementia and depressed patients frequently show a disturbance of sleep. Sleep EEG registration in depression revealed a characteristic sleep EEG profile concerning distinct alterations of sleep architecture and REM-sleep (reduction of SWS, increase and advance of REM-sleep). In dementia polysomnographic assessment has been done less intensively, mainly in patients with dementia of Alzheimer type (DAT). The most significant polysomnographic finding in DAT is a reduction of REM-sleep, which may reflect impaired cholinergic neurotransmission. Therefore, predominantly REM-sleep variables clearly differ between depressed patients and patients with DAT.

In this presentation polysomnographic data and data of cognitive performance in dementia and depression will be reviewed. In addition, own long term studies in patients with different types of dementia and in depressed patients will be presented. The polysomnographic findings of these studies will be discussed with respect to differential diagnosis, prediction of treatment response and the long term course of both diseases. In addition, the results will be related to the current knowledge of the neurochemical and neuroendocrine regulation of sleep.

S04.04

Neuroendocrine and sleep regulation as predictors of illness course and therapy in depression

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Background and Aims: In depression, changes in EEG sleep measures are well documented findings. However, the predictive value of these alterations for treatment and long-term course of depression still warrants clarification. Therefore, we examined whether the previous course of depression, treatment response during antidepressant therapy, and the long-term outcome in follow-up are associated with sleep regulation. Since the hypothalamic-pituitary-adrenocortical (HPA) system may play a crucial role in depression's neurobiology, we evaluated HPA system function as well.

Methods: 15 patients (4 men, 11 women; age 43–59) with depression were enrolled in the study. HPA system assessment using the combined DEX/CRH test and sleep EEG studies were conducted at baseline, after a 6 week antidepressant treatment period (trimipramine), and at follow-up, i.e., after 2–10 years.

Results: The previous clinical course, i.e., the number of episodes until baseline, correlated significantly with EEG sleep measures i.e. sleep continuity values, slow wave sleep (SWS) and REM latency.

During treatment sleep continuity values improved and the correlation with the previous long-term course disappeared. The correlation with SWS persisted. The only sleep EEG marker at baseline predictive for treatment response was REM latency.

In the prospective long-term outcome SWS and REM density variables were related to the occurrence of recurrences. These sleep EEG markers correlated closely with HPA system regulation.

Conclusions: The long-term outcome of depression is related to the sleep EEG pattern: SWS and REM density measures may reflect predictive markers for the long-term course. These markers are associated with HPA system regulation.

Plenary Lecture: Pathways to integrative care in childhood

PL01.01

Pathways to integrative care in childhood

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Mental health care has traditionally been provided within the framework of a verticle mental health services system. Additionally, mental health care is often provided through a variety of agencies or organizations that have a number of different mandates, responsibilities, authorities and accountabilities. These are not well linked with or to each other.

Furthermore, mental health services for children and youth are often not well integrated into adult mental health services. This profusion of confusion regarding mental health care for young people can be much better defined and operationally developed if a model of care based on population mental health care needs and provider mental health care competencies can be applied. This presentation will present the conceptual framework for such a model that allows for mental health care to be intergrated into all levels of health and other systems that provide interventions for young people suffering from a variety of mental health problems.

State-of-the-art Lecture: Physical illness in persons with severe mental disorders

SOA01.01

Physical illness and access to medical services in people with schizophrenia

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The physical well-being of people with schizophrenia is remarkably neglected. Physical illnesses in these people are underdiagnosed and undertreated. A recent study in Australia showed that, although people with schizophrenia suffer more frequently from cardiovascular problems than the general population, they receive catheter much more rarely. People with schizophrenia have been also reported to be less likely than the general population to receive HbA1c and cholesterol monitoring, to receive a retinal examination for diabetes screening, and to be treated for osteoporosis. They have been also found to be more likely to be treated for physical illnesses only when the latter become life threatening. Among the factors contributing to this underdiagnosis and undertreatment of physical illnesses in people with schizophrenia are a low motivation of patients and their relatives to access medical services, the isolation of psychiatric services from other medical facilities, and a tendency of psychiatrists to overlook physical health problems in their patients. However, the most important factor is likely to be the stigma surrounding schizophrenia. The neglect of physical health in people with schizophrenia should be regarded as an expression of discrimination and disregard for their dignity and their rights as human beings and citizens. Due to the lack of prevention and intervention strategies, people with schizophrenia and their families bear the costs of the mental disorder and those of the concomitant physical illnesses, which can exacerbate psychopathological manifestations and impair the subjects' ability to adhere to treatment. Access to physical health care of the same quality as that available to the rest of the population should be considered a basic right of people with schizophrenia and a crucial dimension on which their quality of life has to be evaluated.

Satellite Symposium: Realizing the potential of new antipsychotics: Practical advice for optimizing schizophrenia care. Sponsored by Bristol Myers Squibb

SS01.01

The art and science of switching in patients with schizophrenia: Strategies for achieving a smooth transition

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Many patients with schizophrenia experience persistent symptoms or side effects on their current antipsychotic regimen. Such patients, particularly those treated with conventional antipsychotic agents may benefit from switching to atypical agents, which offer broader efficacy and improved tolerability compared with earlier counterparts. In addition, patients already receiving treatment with an atypical agent may benefit from switching to an alternative atypical, given that there is great variation in (1) individuals' response to different atypical antipsychotics, and (2) the side-effect profile of the atypicals. With switching from one antipsychotic to another becoming increasingly common, there is an urgent need to define optimal switching

strategies. The main goal when switching antipsychotics is to improve or (in stable patients) maintain the symptomatic and functional level, while improving (or not worsening) tolerability. It is important to identify patients who would be likely to benefit from switching and to discuss with them and their carers the advantages and potential problems of the switching process. To date, four strategies have been effective in controlled studies of switching to atypical antipsychotics: therapeutic dose initiation of the new antipsychotic and abrupt discontinuation of the first ('abrupt switch'); gradual dose escalation of the new antipsychotic and abrupt discontinuation of the first ('ascending switch'); therapeutic dose initiation of the new antipsychotic and gradual discontinuation of the first ('descending switch'), and; gradual dose escalation of the new antipsychotic and gradual discontinuation of the first ('cross-titration'). An individualized approach is key to the success of switching, as are patient cooperation and carer support.

SS01.02

Shifting schizophrenia treatment paradigms: The scope for adjunctive therapies

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Combination therapy is well established in bipolar disorder; however, the evidence for this approach in schizophrenia is less robust. Despite the absence of clear guidance, antipsychotic combinations are commonly used in real-life practice, with estimates suggesting that 20–60% of patients receive multiple antipsychotics concurrently. Such polypharmacy may be clinically useful, combining diverse pharmacological actions. However, a clear pharmacological rationale for specific antipsychotic combinations has not yet been elucidated. In this presentation, we consider the varying pharmacological profiles of agents currently used for schizophrenia and explore how best this pharmacology may be exploited to maximize the newer atypicals in clinical practice. For decades schizophrenia has been treated with some success using typical antipsychotics, which are antagonists at dopamine D2 receptors. Atypical antipsychotics were then developed, having D2 antagonism with additional affinity for other receptors, such as serotonin 5-HT_{2A} and 5-HT_{1A} receptors. Most recently, partial D2 agonists have been developed with efficacy to treat schizophrenia and bipolar disorder. These agents have lower intrinsic activity than full agonists, so can act either as functional agonists or antagonists. Additionally, actions to increase noradrenergic function in the prefrontal cortex may be implicated in the efficacy of some antipsychotics. Given the rich pharmacology of antipsychotics, can the combined use of agents with synergistic mechanisms of action provide a true clinical advance in schizophrenia treatment? Polypharmacy is a complex challenge that requires further study in well-designed, randomized, controlled studies. We will review the pharmacological rationale for antipsychotic combination therapy and recent clinical evidence for their benefits.

SS01.03

Meeting the need for efficacy without over-sedation in patients with schizophrenia

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Although the induction of sleep was originally considered to be a desirable therapeutic endpoint for the rapid control of agitation, it is