Poster Session 2: OBSESSIVE-COMPULSIVE DISORDERS

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Gambling and suicidal behaviours: A Swiss study in a treatment center in Switzerland

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Background and aim: Excessive gambling touch between 1 and 3% of the adult population (Shaffer et al. 1999).

Studies of treatment-seeking gamblers establish a relationship between gambling and suicide. We investigated clinical characteristics in excessive gamblers of a Swiss University Hospital(CHUV).

The aim of this study is to compare gamblers with prior suicide attempts (GPSA) with gamblers without prior suicide attempts (Non-GPSA) and with the international literature.

Hypothesis:

- GPSA are confronted with a higher problem load than Non-GPSA.
- GPSA lack social networks and family support as compared to Non-GPSA.

Method: Patients treated for gambling disorders typically show a high level of co-morbidity. Bourget, Data are based on medical files of our treatment center. Among our consecutively admitted patients (2002-2006), we identified pathological gamblers who reported prior suicide attempts directly or not directly linked with gambling.

Results and discussion:

- GPSA were more likely to be women, separated or divorced, referred by the forensic network.
- GPSA were more likely to be disabled and had a history of alcohol abuse. GPSA showed no difference with respect to age at intake or employment status as compared to NON-GPSA.

Discussion: Further research is needed to find out whether the higher proportion of women with prior suicide attempts is due to the fact that men are more likely to complete suicide.

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Young adults: What about gambling disorders?

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Background: Gambling was included in DSM III since 1980 as a psychiatric disorder. Compared to other forms of (compulsive) behavior for example substance abuse), the gambling problems have hardly solicited public and scientific concern. In particular a new generation of young adults raised in an environment of video and internet games has been neglected by researchers. Our knowledge of this population segment with respect to gambling disorders is wanting.

Aims: The present study aims to obtain a clearer description of gambling behavior in this segment of the Swiss population. The

present project should allow us to gain a better understanding of problem burden and will help to identify the different forms of games that are used by young adults age 18 to 25 in Switzerland.

Study design: In a first step we have been planning an explorational and descriptive pilot study. 120 men age 18 to 25 are recruited from Universities, Technical Colleges, Professional Training Schools, Occupational Centers, and newspaper ads as well as via the Internet.

Participants will be evaluated in terms of the following instruments: socio-demographic questionnaire, structured interviews and tools to identify types and characteristics of gambling behavior and concomitant problems, namely SOGS (South Oaks Gambling Survey), Internet Dependence (YOUNG), the Fagerström scale (tobacco), AUDIT (alcohol), Impulsive Behavior Scale (UPPS), BDI II (Beck), the Hamilton Scale for anxiety assessment.

Discussion: The present study serves as the basis for a large scale population based study.

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Long-term treatment of obsessive-compulsive disorder: A prospective follow-up study

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This study aimed to evaluate the long-term course of obsessive-compulsive disorder (OCD) in patients treated with serotonin reuptake inhibitors (SRIs) and to identify predictors of clinical outcome. Seventy-nine patients fulfilling DSM-IV criteria for OCD were followed prospectively for 3 years. Baseline information was collected on demographic and clinical characteristics, using standardized instruments. During the follow-up period, the clinical status of each patient was evaluated monthly in the first year and bimonthly thereafter by means of the Yale-Brown Obsessive Compulsive Scale (Y-BOCS) and the Hamilton Rating Scale for Depression (HDRS). The cumulative probability of achieving an at least partial remission from obsessive-compulsive (OC) symptoms during the 3-year period was 65%. The probability of full remission was 38%. In subjects who achieved an at least partial remission, the probability of subsequent relapse was 60%. Significant predictors of poor outcome included a longer duration of illness, a greater severity of OC symptoms at intake, and the presence of comorbid schizotypal personality disorder. The findings confirm that the course of illness in OCD is usually continuous with fluctuations in the intensity of OC symptoms. Despite adequate SRI therapy, relatively few patients achieve a completely asymptomatic state and, of those who achieve at least a partial remission, a substantial proportion subsequently relapse. One third of OCD patients is treatment-resistant. Further studies with large samples are required to adequately identify predictors of long-term outcome of OCD in order to optimize the choice among the existing treatment modalities.

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Seeking out candidate endophenotypes for OCD. Neurocognitive findings in unaffected relatives.

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Background: Obsessive compulsive disorder (OCD) is a highly heritable neuropsychiatric disorder. Attempts to elucidate contributing genes have met with limited success. Intermediate markers of brain

dysfunction (cognitive endophenotypes) may help focus the search for genetic contributions. Such markers should be present in people at risk of developing OCD in the absence of clinical symptoms. In prior work, OCD patients showed impairment on tests of response inhibition and cognitive flexibility (Chamberlain et al., 2005, 2006).

Methods: First-degree relatives of OCD patients, patient probands, and matched healthy volunteers without a family history of OCD undertook neuropsychological assessment (n=20 per group).

Results: Compared to matched controls without a family history of OCD, unaffected first-degree relatives of OCD patients showed impaired response inhibition (p<0.05) and cognitive flexibility (p<0.05). These deficits were comparable to those in the patients themselves.

Conclusions: Brain-based cognitive markers of inhibitory functions may be of utility in the search for OCD endophenotypes. Examination of relationships between these abnormalities, genetics, and structural/functional brain changes, will help to elucidate aetiological contributions to OCD and putative spectrum disorders.

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Relapse prevention in patients with obsessive-compulsive disorder (OCD)

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Purpose: To compare the efficacy of escitalopram 10 or 20 mg/day with placebo in preventing relapse during 24 weeks in outpatients with obsessive-compulsive disorder (OCD) who had responded to an initial 16-week open-label treatment with escitalopram.

Methods: A multinational, randomised, double blind, placebocontrolled, flexible to fixed dose relapse prevention study with escitalopram in outpatients with OCD. The study consisted of a 16-week open-label period with 10 to 20 mg escitalopram followed by a 24 week double blind, placebo-controlled period, and a 1 week taper period. Patients who had responded to treatment (≥25% decrease in the Yale-Brown Obsessive Compulsive Scale (Y-BOCS) total score) by the end of the 16-week open-label period were eligible for randomisation to either escitalopram or placebo for a further 24 weeks.

Results: 468 patients with OCD were treated with open-label escitalopram (10 mg or 20 mg) for 16 weeks. There were 320 responders (68%) who were randomised to change to placebo (n=157) or to continue with escitalopram (at the assigned dose) for further 24 weeks (n=163). The primary analysis (time to relapse) showed a clear beneficial effect of escitalopram relative to placebo (log-rank test, p<0.001). The proportion of patients who relapsed was statistically significantly higher in the placebo group (52%) than in the escitalopram group (23%) (p<0.001, chi-square test). The risk of relapse was 2.74 times higher for placebo- than for escitalopram-treated patients (chi-square test, p<0.001). Escitalopram was well tolerated.

Conclusion: Escitalopram was effective in preventing relapse of OCD and was well tolerated as continuation treatment.

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The treatment of obsessive-compulsive disorder with escitalopram

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Purpose: The efficacy and tolerability of escitalopram in obsessive-compulsive disorder (OCD) were investigated in a 24-week, randomised, placebo-controlled, active-referenced, double blind study.

Methods: 466 adults with OCD were randomised to escitalopram 10mg/day (N=116), escitalopram 20mg/day (N=116), paroxetine 40mg/day (N=119), or placebo (N=115) for 24 weeks. The prespecified primary efficacy endpoint was the mean change in the Yale-Brown Obsessive-Compulsive Scale (Y-BOCS) total score from baseline to Week 12 based on the intent-to-treat population and last observation carried forward (LOCF) using analysis of variance (ANCOVA).

Results: Escitalopram 20mg/day was superior to placebo on the primary endpoint. After 12 weeks, on the primary efficacy endpoint, there was a statistically significant difference from placebo for 20mg escitalopram and paroxetine. In the escitalopram 20mg/day group, the Y-BOCS total score was significantly lower than in the placebo group as early as Week 6. At Week 24, the proportion of remitters (Y-BOCS \leq 10, LOCF, pre-defined) was significantly greater (p \leq 0.05) for 20mg escitalopram (41.2%) than placebo (27.4%), but not for 10mg escitalopram (36.6%) or paroxetine (37.9%). The response rate (>25 decrease from baseline Y-BOCS, LOCF, pre-defined) was significantly greater than placebo (50.4%) for 20mg escitalopram (70.2%) and paroxetine (67.2%). Statistically significantly more patients withdrew from the placebo group (18%) due to lack of efficacy, than paroxetine (8%) or escitalopram 20mg/day groups (6%). More paroxetine-treated patients withdrew due to adverse events than escitalopram- or placebo-treated patients.

Conclusion: Escitalopram was efficacious and well tolerated in the treatment of OCD, with 20mg escitalopram showing statistically significant superiority at the primary efficacy endpoint.

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Amis subito: Assessment, measurement, intervention and studies for the prevention of suicidal behaviour in individuals, inclined to gamble excessively

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Background and aim: Actually, the suicidal risk in people with gambling problems is insufficiently evaluated; this risk is all the more hard to specify within a population which underreports gambling behaviour and associated co-morbidities. Estimations of suicidal behaviour vary between studies, suicide attempts were observed in 4% to 40% of gamblers studied. Suicidal thoughts were reported for 25% to 92% of people with gambling problems. 64% of gamblers that committed suicide did neither inform family or friends nor health professionals about their suicidal intents. In the context of a pilot study, we wish to study suicidal behaviour in people with gambling problems.

Method: The goal of the study consists in the early identification of gambling problems associated with suicidal behaviour. A short intervention, specifically targeted towards the prevention of suicide will be compared with the current treatment for gambling problems. Gambling and suicidal behaviour will be monitored over 6 meetings during 12 months.