acute therapy studies b) Maintenance (3-month) controlled studies, with proper stratification of stabilized patients and c) Long-term (12-month) manic/depressive episode-preventive studies. Lithium, carbamazepine, sodium Valproate are examples of drugs that could serve as suitable active comparators and standard validated rating scales for the assessment of clinical states are also available. Examples of decision trees for manic or depressed index cases are possible and these are out-lined.

P04.08

Obstetric complications as risk factors for bipolar disorder

B. Kirk Øgendahl*, R.W. Licht, P.B. Mortensen. *National Centre for Register-Based Research, University of Aarhus, Denmark*

Objective: To study whether obstetric complications increase the risk of developing bipolar disorder.

Method: The Danish Psychiatric Central Register contains data on all psychiatric hospitalizations in Denmark since 1969. The Danish Medical Birth Register covers all births in Denmark since 1973. The registers were linked using the CPR-number. In the study we have identified 161 persons born between 1973–1983 being diagnosed with bipolar disorder (after the ICD-10 classification F30 or F31 or the ICD-8 classification 296.19 or 296.39) before 1999. To each of the cases we have matched 50 controls. The controls are born the same year as the case, and are alive on the day the case has been diagnosed. All the cases and controls have an identifiable mother. The controls have no psychiatric diagnosis.

Results: The data will be present on the possible role of low birth weight and gestational age and other obstetric factors as risk factors for bipolar disorder. It will be adjusted for family history for mental disorders as well as social variables.

Conclusion: In case of a correlation between obstetric complications as risk factors for bipolar disorder it can be used as prevention.

P05. Brain imaging – structural

P05.01

A diffusion tensor and metabolite spectroscopic imaging study of white matter connectivity in schizophrenia

J.K. Burns[1]*, M. Bastin[1], S. Lawrie[2], I. Marshall[1], T. McGillivrary[2], H. Whalley[3], D. Job[1], E. Johnstone[1]. *[1]University of Edinburgh, Department of Psychiatry, Scotland; [2]University of Edinburgh, Department of Medical Physics, Scotland; [3]Welcome Trust Clinical Research Fellow, University of Edinburgh, Scotland, UK*

Background: Structural and functional imaging of the brain has demonstrated abnormalities of both grey and white matter. Functional studies (fMRI and PET) suggest aberrant fronto-temporal connectivity, but further evidence is required. Diffusion Tensor Imaging (DTI) measures diffusion anisotropy, an indicator of the structural integrity of neuronal tracts. MR Spectroscopy studies (MRS) have found reduced N-acetyl aspartate (NAA) concentrations in frontal and temporal regions, indicating reduced neuronal density or viability. We used both techniques in an attempt to identify the structural correlates of impaired functional connectivity in schizophrenia.

Methods: Thirty patients with DSM-IV schizophrenia were compared with thirty healthy controls. DTI, MRS and sMRI were performed on all subjects. A symptom scale (PANSS) and IQ score (NART) was recorded. Data analysis included both whole

brain voxel by voxel analysis using Statistical Parametric Mapping (SPM), and a region of interest approach.

Results: Data from previous smaller studies show reduced diffusion anisotropy and reduced NAA concentrations in schizophrenia. We expect to replicate these findings in our larger study and specifically localise abnormalities to fronto-temporal white matter tracts.

Conclusions: DTI and MRS have the capacity to identify the structural correlates of impaired functional connectivity in schizophrenia.

P05.02

Magnetic resonance tomography (MRT) of brain of serial sexual sadists

O.A. Bukhanovskaya[1]*, A.O. Bukhanovsky[2]. *[1]Scientific Center for Cure and Rehabilitation, Rostov-on-Don; [2]Department of Psychiatry, Rostov State Medical University, Russia*

Objectives: a study of the state of the terminal brain and its deep parts, as well as the skull in serial sexual sadists (SSS) with a revelation of a possible dysgenetic origin.

Methods: clinical and pathopsychological, magnetic resonance tomography.

Summary of the results obtained: The principal group included 22 SSS, the control group consisted of 28 males between 18 and 66. All were heterosexual, did not show any signs of paraphilia or sexual aggression and were in general law-abiding. In each case (100%) the following pathological MRT symptoms were revealed: dilation of subarachnoid fissures of frontal, frontal-temporal and frontal-temporal-parietal parts; dilation and flatting of contours of grooves and disordered differentiation between the grey and white substances of the frontal and frontal-temporal parts; dilation of the lateral ventricles and a significant asymmetry due to large right against left; dilation or narrowed fissure-like third ventrile; transparent partition pathology (left displacement up to 7 mm and cysts diameter up to 9 mm; dysgenetic of the callous body; congenital skull dysraphy.

Conclusions: the revealed and described cerebral deviations, changing a number of neurodynamic and psychological characteristics, create a cerebral predisposition to SSS, significantly raising the risk of serial sexual crime appearance.

P05.03

Progression of hippocampal atrophy is associated with clinical deterioration in Alzheimer's disease

P. Schönknecht[1]*, J. Pantel[1], M. Essig[2], M. Amann[2], L. Schad[2], J. Schröder[1]. *[1]Department of Psychiatry, University of Heidelberg; [2]German Cancer Research Center, Heidelberg, Germany*

Introduction: Using quantitative magnetic resonance imaging (MRI), recent studies found atrophic changes of the medial temporal lobe structures already in early stages of Alzheimer's disease (AD). These changes were cross-sectionally correlated with the severity of dementia which led to the hypothesis that progressive medial temporal lobe atrophy might be used as a marker of disease progression.

Method: We investigated the progression rate of hippocampal atrophy with respect to rate of clinical deterioration in 13 AD patients and 8 healthy controls using volumetric MRI.

Results: Already at baseline, the AD patients showed significantly smaller hippocampal volumes than controls. While AD – in comparison to healthy ageing – was characterized by a rapid decline of hippocampal volumes (-8.1%/year) only a moderate decrease of