

Cologne, Cologne, Germany <sup>2</sup> Department of Psychiatry and Psychotherapy, University of Bochum, Bochum, Germany <sup>3</sup> Department of Psychiatry, University of Turku, Turku, Finland <sup>4</sup> Department of Psychiatry, University of Amsterdam, Amsterdam, The Netherlands <sup>5</sup> Early Intervention Service, University of Birmingham, Birmingham, UK <sup>6</sup> Department of Psychology, University of Manchester, Manchester, UK <sup>7</sup> all departments

**Background:** One aim of the European prediction of psychosis study (EPOS) has been to evaluate the clinical course of putatively prodromal patients in terms of psychopathology.

**Methods:** 245 patients at risk for psychosis defined by attenuated positive symptoms, brief limited psychotic symptoms, a state/ trait combination or cognitive-perceptive basic symptoms was recruited in six centres in four countries. The Structured Interview for Prodromal Syndromes (SIPS) and the Bonn Scale for the Assessment of Basic Symptoms – Prediction List (BSABS-P) were employed. Follow-up was scheduled after 9 months (t1) and 18 months.

**Results:** In total, 40 patients developed a psychosis (P). Compared to those without a transition (NP), P showed significantly higher SIPS scores at baseline. The same applied to the BSABS-P sub-scores 'cognitive perception disturbances' and 'cognitive motor disturbances'. The P sub-group developing psychosis after t1 showed no significant change of the SIPS positive (SIPS-P) sub-score or of any BSABS-P score from baseline to t1, whereas all scores improved in the NP group. At t1, SIPS-P and BSABS-P sub-score 'cognitive thought disturbances' were significantly lower in those later becoming psychotic.

**Conclusion:** Patients at risk showing a transition to psychosis during exhibited a pronounced psychopathology at baseline. Also, the positive symptom scores did not significantly improve during 1st follow-up, whereas those patients with no transition during the complete follow-up showed an improvement of all scores. As EPOS is a naturalistic study, different treatments have been performed in a considerable portion of the patients and association with course awaits further analysis.

## S55.02

Subjective quality of life and its changes in patients at risk of psychosis

R.K. Salokangas <sup>1</sup>, M. Heinimaa <sup>1</sup>, T. Svirskis <sup>1</sup>, J. Klosterkoetter <sup>2</sup>, S. Ruhrmann <sup>2</sup>, H. Graf von Reventlow <sup>2</sup>, D.H. Linszen <sup>3</sup>, P.M. Dingemans <sup>3</sup>, M. Birchwood <sup>4</sup>, P. Patterson <sup>4</sup>, G. Juckel <sup>5</sup>, A. Morrison <sup>6</sup>, the EPOS Group <sup>1,6</sup>. <sup>1</sup> Psychiatric Clinic, University of Turku, Turku University Hospital, Turku, Finland <sup>2</sup> Department of Psychiatry and Psychotherapy, University of Cologne, University Hospital of Cologne, Cologne, Germany <sup>3</sup> Akademisch Psychiatrisch Centrum, Academic Medical Centre, Adolescentenkliniek, Amsterdam, The Netherlands <sup>4</sup> Early Intervention Service, University of Birmingham, Birmingham, UK <sup>5</sup> Department of Psychiatry, Ruhr-University of Bochum, LWL-Klinik, Bochum, Germany <sup>6</sup> Department of Psychology, Victoria University of Manchester, Manchester, UK

**Objectives:** The European Prediction of Psychosis Study (EPOS) aimed to study a large sample of young patients who are at risk of psychosis and to estimate their conversion rate to psychosis during 18 months follow-up. This presentation describes quality of life and its changes in patients at risk of psychosis.

**Methods:** In six European centres, 16 to 35 year old psychiatric patients were examined. Risk of psychosis was defined by occurrence of basic symptoms, attenuated psychotic symptoms, brief, limited or

intermittent psychotic symptoms or familial risk plus reduced functioning. Quality of life (QoL), measured by the Modular System for Quality of Life, was assessed at baseline and at 9 and 18 months' follow-ups. Psychiatric patients without prodromal symptoms and healthy subjects were comparison groups.

**Results:** In all, 245 risk patients were included. At baseline, they reported lower QoL than non-risk patients and healthy controls. Basic symptoms associated negatively with QoL, and there were differences between the study centres. During the follow-up, QoL raised less in risk patients than in non-risk patients. Baseline QoL did not predict transition to psychosis. However, its development was poorer in patients with than in those without transition to psychosis.

**Conclusions:** Those of the psychiatric patients who are at risk of psychosis have lower QoL than other psychiatric patients or healthy controls. QoL does not predict transition to psychosis, but its changes correlates with changes in clinical state. The results indicate that there is a need for comprehensive intervention with the patients at risk of psychosis.

## S55.03

Transition to psychosis: Neuropsychological test results of the epos study

D.H. Linszen <sup>1</sup>, D.H. Nieman <sup>1</sup>, H.E. Becker <sup>1</sup>, J.R. van de Vliert <sup>1</sup>, P.M. Dingemans <sup>1</sup>, J. Klosterkoetter <sup>2</sup>, M. Birchwood <sup>3</sup>, R.K. Salokangas <sup>4</sup>, S. Ruhrmann <sup>2</sup>, G. Juckel <sup>5</sup>, A. Morrison <sup>6</sup>, P. Patterson <sup>3</sup>, M. Heinimaa <sup>4</sup>, H. Graf von Reventlow <sup>2</sup>, the EPOS Group <sup>1,6</sup>. <sup>1</sup> Akademisch Psychiatrisch Centrum, Academic Medical Centre, Adolescentenkliniek, Amsterdam, The Netherlands <sup>2</sup> Department of Psychiatry and Psychotherapy, University of Cologne, University Hospital of Cologne, Cologne, Germany <sup>3</sup> Early Intervention Service, University of Birmingham, Birmingham, UK <sup>4</sup> Psychiatric Clinic, University of Turku, Turku University Central Hospital, Turku, Finland <sup>5</sup> Department of Psychiatry, Ruhr-University of Bochum, LWL-Klinik, Bochum, Germany <sup>6</sup> Department of Psychiatry, Victoria University of Manchester, Manchester, UK

**Introduction:** Both schizophrenia and ultra high risk (UHR) patients show reduced neurocognitive performance compared to matched healthy control subjects. In the current study we compared neurocognitive performance at baseline and follow up between UHR patients who made the transition to psychosis and patients who did not.

**Method:** Patients were eligible for the study when they met criteria for one or more of the following groups: Attenuated symptoms or brief limited intermitted psychotic symptoms or a first degree family member with a psychotic disorder and reduced functioning or basic symptoms. We assessed 216 UHR patients (166 males, mean age: 22,6 SD 5,2) with a neuropsychological test battery composed of the National adult reading test (premorbid IQ), California verbal memory test (verbal memory), spatial working memory test, verbal fluency first letter and categories (executive functioning), finger tapping test (motor speed) and continuous performance test (sustained attention). Data were collected in 7 participating centres of EPOS. Follow up was at 9 months.

**Results:** 37 UHR patients made the transition to psychosis (25 males, mean age 21,5 SD 4,8). The only test that showed a significant difference between the transition and non transition group at baseline was verbal fluency categories ( $t=2.79$ ,  $p=0.006$ ).

**Conclusion:** Patients who later make the transition to psychosis perform significantly worse on verbal fluency categories than patients