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Examining the association between social cognition and functioning in individuals at ultra-high risk for psychosis

Running title: Examining the association between social cognition and functioning in UHR

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Abstract

Objective: Social and role functioning is compromised for the majority of individuals at ultra high risk (UHR) for psychosis and it is important to identify factors that contribute to this functional decline. This study aimed to investigate social cognitive abilities, which have previously been linked to functioning in schizophrenia, as potential factors that impact on social, role and global functioning in UHR patients.

Method: Thirty UHR patients were recruited from an established at risk clinical service in Melbourne, Australia and completed a battery of social cognitive, neurocognitive, clinical and functioning measures. We examined the relationships between all four core domains of social cognition (emotion recognition, theory of mind, social perception and attributional style), neurocognitive, clinical and demographic variables with three measures of functioning (the Global Functioning Social and Role scales and the Social and Occupational Functioning Assessment Scale) using correlational and multiple regression analyses.

Results: Performance on a visual theory of mind task (visual jokes task) was significantly correlated with both concurrent role ($r = .425, p = .019$) and global functioning ($r = .540, p = .002$). In multivariate analyses it also accounted for unique variance in global, but not role functioning after adjusting for negative symptoms and stress. Social functioning was not associated with performance on any of the social cognition tasks.

Conclusions: Among specific social cognitive abilities only a test of theory of mind was associated with functioning in our UHR sample. Further longitudinal research is needed to examine the impact of social cognitive deficits on long-term functional outcome in the UHR group. Identifying social cognitive abilities that impact significantly on functioning is

important to inform the development of targeted intervention programs for UHR individuals.

Keywords: Functioning; Prodrome; Psychosis; Social cognition; Theory of mind; Ultra high risk

Introduction

Social cognition refers to the 'mental operations that underlie social interactions' (Green et al., 2008). It is comprised of four core domains: emotion processing (perceiving and displaying emotions), theory of mind (ToM; the ability to represent the mental states of others), social perception (decoding and interpreting social cues in others) and attributional style (the way in which individuals explain the causes, or make sense, of social events or interactions) (Pinkham et al., 2014). Social cognitive impairments are a common feature of schizophrenia-spectrum disorders (Savla et al., 2013). These impairments are present not only in the established illness, but are evident in the early phase of illness, at the time of the first episode of psychosis (FEP) (Thompson et al., 2012).

Additionally, over the last two decades, it has been possible to identify individuals who are at ultra high risk (UHR) of developing a psychotic disorder such as schizophrenia (i.e. they are putatively prodromal for the illness) (Yung et al., 1996, 1998; Miller et al., 2002), and there is growing evidence that social cognitive deficits are present in this group (Barbato et al., 2015; Lee et al., 2015). Performance in each of the four core social cognitive domains in UHR individuals is generally intermediate between FEP and healthy control groups (Green et al., 2012; Thompson et al., 2012).

Social cognitive impairment is an important determinant of functional outcome in schizophrenia (Fett et al., 2011). However, relatively little is known about the relationship between social cognition and functioning in the UHR population. The few studies that have examined this relationship have all reported inconsistent results. In the largest cross-sectional study conducted to date, facial emotion recognition and ToM task performance were positively correlated with functioning (Barbato et al., 2013). In contrast, a smaller

study reported no associations between functioning and performance on four separate ToM tasks (Stanford et al., 2011). After accounting for negative and depressive symptoms, poorer performance on a vocal but not facial affect recognition task was also reported to be modestly but significantly associated with poorer functioning (Amminger et al., 2013).

To the authors' knowledge, no previous studies have examined the association between functioning and all four core domains of social cognition in a single UHR cohort. Studies have also tended to use functioning measures that combine both social and role functioning into a single global construct. Functioning is a broad concept that constitutes a variety of domains. While global functioning is useful as a marker of overall impairment, it is important to examine social and role functioning as separate constructs on the basis of previous evidence that dysfunction in each of these domains may be driven by different aspects of illness (Fett et al., 2011; Strassnig et al., 2015). Many functioning measures have also been developed for adults with chronic illness and therefore may not be suitable for UHR populations, which consist predominantly of adolescents and young adults who face functional demands largely unique to individuals at this stage of life (e.g. high school or tertiary education, starting work for the first time and dating).

Thus the aim of the current exploratory study is to investigate whether performance on a broad range of social cognitive measures is associated with concurrent social, role or global functioning in individuals at UHR for developing psychosis. On the basis of previous research in those with established psychotic disorders (Fett et al., 2011; Pinkham et al., 2015) and the largest UHR study conducted in this area to date (Barbato et al., 2013), we hypothesized that poorer performance on ToM tasks would be associated with poorer functioning.

Methods

Participants

Data were derived from a larger study, which investigated the degree of social cognitive impairment in UHR individuals in comparison to FEP and healthy control participants. This sample has been previously described elsewhere (Thompson et al., 2012, 2013). Briefly, 30 UHR participants were recruited from Orygen Youth Health, an outpatient service in Melbourne. Inclusion criteria for the UHR participants were: 1) aged between 15-25 years, 2) be help-seeking, 3) present with a drop in functioning (or chronic low functioning for the past year) and 4) meet UHR criteria for either attenuated psychotic symptoms, brief limited intermittent psychotic symptoms (BLIPS) and/or trait vulnerability, according to the Comprehensive Assessment of At-Risk Mental States (CAARMS; Yung et al., 2005). In addition, UHR participants were required to be clinically stable (i.e., not an inpatient or requiring crisis care) and be able to provide informed consent. Patients were excluded if they were being treated for a major neurological disorder, had poor English language skills or had an IQ <70. Those with impaired visual acuity (i.e., blurred vision or less than corrected 20/40 vision) or corrected auditory acuity were also excluded from the study. The study was approved by the local ethics committee. All participants provided written informed consent.

Procedure and measures

Demographics

Patient's age, gender and number of years of completed education were recorded at the time of the assessment.

Psychopathology

Positive psychotic symptoms were assessed using the Brief Psychiatric Rating Scale (BPRS) psychotic subscale (Overall and Gorham, 1962), derived from combining scores from the unusual thought content, suspiciousness, hallucinations and conceptual disorganization items. Negative psychotic symptoms were assessed using the Scale for the Assessment of Negative Symptoms (SANS; Andreasen, 1984). Depression, anxiety and stress were measured using the Depression Anxiety Stress Scale (DASS; Lovibond and Lovibond, 1995). This is a 42-item self-report instrument which provides a total score and three subscale scores for depression, anxiety and stress. These were included on the basis that they may be associated with poor functioning and may influence the association between social cognition and functioning (Cotter et al., 2014). All assessments were conducted by two of the authors (AP and CB). Inter-rater reliability was not formally assessed; however, both raters had considerable experience administering psychiatric assessments and undertook reliability checks throughout the study period.

Social cognitive measures

Social cognitive measures were chosen that represent each of the 4 core domains of social cognition (Pinkham et al., 2014), described below.

Emotion recognition. The adult version of the Diagnostic Analysis of Nonverbal Accuracy-2 (DANVA-2) scale for faces and voices (Nowicki and Carton, 1993; Nowicki and Duke, 1994) was used to assess emotion recognition. The faces subtest consisted of 24 colour photographs of an equal number of happy, sad, angry and fearful facial expressions of high and low intensities. The participant was required to choose the correct facial emotion expression from the four options given. The Paralanguage subtest consisted of two alternating professional actors speaking in a way designed to express happy, sad, angry, or fearful feelings by saying a neutral sentence, "I am going out of the room now but I'll be back later". There were 24 voices of an equal number of emotions of high and low intensities. Both the face and voice stimuli were presented in a randomised order using a computer program. Total scores for errors on either task (maximum 24) were computed. The DANVA-2 has good reliability and reasonable construct validity (Nowicki and Duke, 1994).

Theory of Mind. The Hinting Task (Corcoran et al., 1995) was used to assess the ability to infer real intentions based on indirect speech content. Ten short passages were read aloud to the participant. Each involved an interaction between two characters and ended with one of the characters dropping an obvious hint. For each passage the participant was asked what the character 'really' meant based on what they had said/hinted at. A second obvious hint was given if the participant's first response was incorrect. A total score was calculated out of 20. In addition, a version of the visual jokes task (Corcoran et al., 1997) was also administered because it is not reliant on verbal memory or verbal comprehension. Ten cartoon jokes were displayed on a computer screen one at a time. Jokes were designed to elicit either mentalising about a character's thoughts (5 ToM jokes), or mentalising about

physical aspects of the cartoon (5 control jokes). These were presented in a randomised order both within and between categories. Participants were instructed to explain the humour in the joke, and could take as much time as they needed (responses were audio recorded). Scoring was performed by CB, AP and AT in collaboration, and was based on the previously devised validated method (Corcoran et al., 1997). Higher scores were given when the participants' explanation specifically referred to mental states (each joke scored between 0-3). Total mentalising sub-scores were calculated for the control (maximum score of 15) and ToM conditions (maximum score of 15). Only the ToM jokes score was used for the analysis in the current study.

Social perception. The social cognitive measure from the MATRICS battery (Nuechterlein et al., 2008), the managing emotions module (branch 4) of the Mayer-Salovey-Caruso Emotional Intelligence Test (MSCEIT; Meyer et al., 2002), was used to measure social perception. Vignettes are read aloud to participants and they are asked various questions relating to their emotions and the emotions of others. These questions typically involve the perception of social or interpersonal situations and require interpretation of the appropriate response to these situations. The total managing emotions score on the MSCEIT is scored using a web-based scoring program (available from Multi-Health Systems, Inc, Toronto, Ontario, Canada) using unadjusted consensus norms from a large normative sample. Scores are automatically calculated and scaled with a mean of 100 and a standard deviation of 15. Higher scores represent better emotional management. The Schema Component Sequencing Task - Revised (SCST-R; Corrigan and Addis, 1995) was administered to assess social knowledge, another aspect of social perception (Pinkham et al., 2014). Twelve social scenarios (e.g., eating out in a restaurant) are broken down into six

(short sequence) or nine (long sequence) actions. Each action was typed on a laminated card. Prior to each sequence the participant was informed of the scenario with a “header” card. Each of the action cards for that given scenario were then placed in front of the participant in a predetermined random order. Participants were instructed to arrange the cards into the correct order as quickly as possible. Mean “juxtaposition scores” were calculated for each sequence by dividing the number of cards correctly juxtaposed to neighboring cards by the total possible correct juxtapositions. Scores ranged from 0-1 (Corrigan and Addis, 1995).

Attributional style. We used a locus of control (LOC) task to infer whether participants had a tendency towards an externalising or internalising bias. LOC, which is the extent to which an individual believes he/she can control events that affect them, was assessed with the Adult Nowicki Strickland Internal External (ANSIE) locus of control scale (Nowicki and Duke, 1974). This is a 40-item, self-administered questionnaire requiring yes or no answers (e.g. “Do you believe that wishing can make good things happen?”). Scores range from 0-40. Higher scores represent a more externalized bias (outcome of events are determined by external factors such as the environment) while a lower score represents an internalized bias (outcome of events are related to something the individual did/thought).

Neurocognitive measures

IQ was measured using the vocabulary and matrix reasoning subtests of the Wechsler Abbreviated Scale of Intelligence (WASI; Wechsler, 1999). Information processing speed was assessed using the Trail Making Tests A and B (Reitan, 1955). Verbal working memory was

measured using the letter number span test (Gold et al., 1997). Visual working memory was measured using the Wechsler Memory Scale - Third Edition (WMS-III) spatial span subtest (Wechsler, 1997).

Functioning

Social and role functioning were independently assessed using the Global Functioning: Social (GFS) and Global Functioning: Role (GFR) scales respectively (Cornblatt et al., 2007). Both scales are interviewer-rated, with scores ranging from 1 (extreme dysfunction) to 10 (superior functioning). Anchors are provided for each point on the scale. The GFS scale assesses engagement in social activities and the quality of interpersonal relationships with family, friends and romantic partners. The GFR scale assesses educational or vocational engagement and performance. These scales have previously been validated in the UHR population (Cornblatt et al., 2007). Functioning was also assessed using the Social and Occupational Functioning Assessment Scale (SOFAS; Goldman et al., 1992). This is a brief and well-established interviewer-rated scale that combines assessment of both social and role functioning to provide a single global functioning score ranging from 0 to 100. Specific anchors are included for each ten-point range. Higher scores indicate better functioning.

Data analysis

Data were analysed using SPSS statistical software (IBM SPSS Statistics for Windows, Version 22.0. Armonk, NY: IBM Corp). In addition to the individual social cognitive assessments, a composite social cognitive score was computed by averaging z scores on each of the six

social cognitive assessments from the emotion recognition, ToM and social perception subdomains. The attributional style data was not incorporated into the composite score as this measure was not based on social cognitive performance per se, but instead based on personal perspective of external events. This was further corroborated by the lack of correlation between the attributional data and the other three social cognitive domains, which were each in turn highly correlated with one another (all $p \leq .002$).

The z scores were derived based on data obtained from an aged-matched healthy control group (for further details of this group see Thompson et al., 2012). Mean scores for each of the assessments were calculated from the healthy controls and then subtracted from the scores obtained from the UHR patients for each of the individual tests; these were then divided by the standard deviation obtained from the healthy controls for that respective test. All scores were computed so that higher values indicated better performance. Scores for each of the assessments were then added together and divided by the number of assessments (six) so that equal weightings were given to each of the assessments and therefore the three social cognitive subdomains within the overall composite score. Bivariate correlations were calculated to examine the relationships of the three functioning measures with social cognitive and neurocognitive performance, clinical symptoms and demographic variables. Variables that were significantly correlated with each of the functioning measures ($p < .05$) were input into multiple linear regression analyses using the 'enter' method. Correlations were uncorrected for multiple comparisons due to the exploratory nature of the research on the basis of limited and inconsistent previous findings in this area (e.g., Stanford et al., 2011; Barbato et al., 2013). Separate multivariate analyses were conducted for each of the social cognition measures that were correlated with any of

the functioning assessments. Functioning measures were included as the dependent variables for all multivariate analyses. Model fit was assessed using the adjusted R^2 statistic.

Results

Sample characteristics

An overview of the clinical and demographic characteristics of the sample and performance on the social cognition tasks is provided in Table 1. Of the 30 patients included in the sample, 22 patients met criteria for attenuated positive psychotic symptoms, 2 met trait criteria, 5 met criteria both for the trait and attenuated symptoms groups and 1 met a combination of trait, attenuated symptoms and BLIPS criteria.

[Insert Table 1 about here]

Correlations

The results of the correlational analysis are presented in Table 2. The ToM jokes task was the only social cognition measure that was significantly correlated with any of the functioning measures. This was positively correlated with both the GFR ($r = .425, p = .019$) and SOFAS scales ($r = .540, p = .002$), suggesting that better performance on this task was associated with better role and global functioning. There was also a trend association verging on significance ($r = .360, p = .051$) between performance on the MSCEIT – managing emotions module and SOFAS score, suggesting that better social perception was associated

with better global functioning. This was also included in multivariate analyses. None of the social cognition tasks were significantly correlated with the GFS scale so additional multivariate analyses were not conducted for this measure. Composite social cognitive performance was also unrelated to any of the functioning measures.

Of the neurocognitive, clinical and demographic variables, levels of negative symptoms (SANS total) was significantly negatively correlated with all functioning measures and was included in all subsequent multivariate analyses. DASS stress subscale scores and years of education were significantly associated with SOFAS and GFR scores respectively and were included in related multivariate analyses. There was also a trend association between DASS total score and SOFAS ($r = -.352, p = .057$), however this was not included in subsequent multivariate analyses due to high correlation with the DASS stress subscale score ($r = .894, p < .001$).

We performed additional exploratory correlations to examine the strength of the relationships between each of the three functioning measures. Scores on the SOFAS were significantly correlated with both the GFS ($r = .703, p < .001$) and GFR ($r = .435, p = .016$) scales. However, the GFS and GFR scales were not significantly correlated with one another ($r = .158, p = .406$).

[Insert Table 2 about here]

Regression analyses

Global Functioning. Two separate multivariate models were created to examine the relationships between performance on the ToM jokes and MSCEIT tasks with SOFAS score (see Table 3). Performance on the ToM jokes task ($\beta = .381$; $p = .009$) and SANS total ($\beta = -.466$; $p = .003$) remained significantly associated with functioning in the first multivariate model, however, the DASS stress subscale was no longer significantly associated after adjusting for these other variables. The model accounted for 53% of variance in SOFAS scores (adjusted $R^2 = .533$, $F(3,26) = 12.012$, $p < .001$). In the other analysis, investigating performance on the MSCEIT social perception task, total MSCEIT score was no longer significantly associated with SOFAS scores in the multivariate model after adjusting for negative symptoms and stress (adjusted $R^2 = .406$, $F(3,26) = 7.603$, $p = .001$).

[Insert Table 3 about here]

Role Functioning. Performance on the ToM jokes task was not found to be significantly associated with GFR scores in the multivariate model (see Table 4). Though the model was significant, none of the variables entered into the model remained as significant independent predictors (adjusted $R^2 = .281$, $F(3,26) = 4.780$, $p = .009$).

[Insert Table 4 about here]

Discussion

Functioning has become an important outcome of interest across the psychosis-spectrum (Yung et al 2010; Brissos et al., 2011; Lin et al., 2013). Specific investigation of factors associated with and predictive of functional outcome in the UHR population is in its infancy, but rapidly gathering interest as an alternative (or additional) outcome to transition to psychosis (Cotter et al., 2014). This work has been supported by evidence that UHR individuals not only have poor social and occupational functioning at initial presentation (Yung et al., 2004; Velthorst et al., 2010), but that a large proportion continue to function poorly at long-term follow-up regardless of whether they develop a full-threshold psychotic disorder (Addington et al., 2011; Salokangas et al., 2013; Yung et al., 2015). This study sought to examine the association between performance on a broad range of social cognition tasks with social, role and global functioning.

Overview of findings

We found partial support for our hypothesis that poorer performance on ToM tasks would be associated with poorer functioning. A deficit in one measure of ToM (jokes task) was found to be associated with overall functional impairment, even after adjusting for negative symptoms. However it was not associated with specific measures of social or role functioning.

Aside from the significant correlations between the ToM jokes task and functioning assessments, and the trend association between the social perception task and SOFAS score, all other correlations were weak and non-significant, suggesting that social cognitive performance was not generally associated with concurrent social, role or global functioning.

This may be at least partially attributable to the lack of marked social cognitive deficits in this group. When our UHR group was previously compared to healthy controls they performed more poorly on all of the social cognitive measures reported in this paper, however performance was only statistically significantly worse on the ToM hinting task (Thompson et al., 2012). The lack of association between the poor performance on the hinting task and any of the functioning assessments is interesting in light of recent evidence showing it to be one of the social cognitive tests most strongly associated with poor concurrent functioning in a schizophrenia cohort (Pinkham et al., 2015). Mean overall performance on the ToM jokes task did not significantly differ from healthy controls, though the variability in performance was greater in the UHR group (Thompson et al., 2012). This could indicate that the severity of impairments across the other domains were not sufficient to impact on functioning or could be compensated for, unlike what has been reported in patients with full-threshold psychotic disorders who exhibit much more widespread and pronounced social cognitive and neurocognitive deficits (Fioravanti et al., 2012; Thompson et al., 2012). Other factors such as negative symptoms may have a greater influence on functioning when social cognitive impairments are only minor. Alternatively, it may be that sustained negative symptoms worsen social cognitive deficits over time, resulting in further functional decline.

It is also possible that many individuals in our UHR sample were not truly at risk of psychotic disorder. Thus any association between social cognitive impairments and poor functioning in those actually in the prodromal phase of schizophrenia may not be detectable, due to high numbers of people not at risk in whom no association would be expected. It would have been interesting to examine the differences between those who subsequently went on

to develop a full-threshold psychotic disorder and those that did not, however, this was not possible in the current study due to the low number of participants.

The results of this study both support and conflict with previous research in the UHR group. Barbato et al. (2013) identified a significant association between another measure of visual ToM, the Reading the Mind in the Eyes task, which is corroborated by the findings in the current study. They also found no links between vocal emotion recognition and functioning. However, they did report a significant correlation between facial emotion recognition and functioning which was not found in the current study. This may be due to their much larger sample size ($n=137$) and a lack of power in the current study to replicate this association. The measure of facial emotion recognition used in the current study also included a narrower range of expressions and may have been less sensitive than the measure used in the previous study (Barbato et al., 2013). The results of the current study also conflict with the findings of Stanford et al. (2011) who reported no associations between four different measures of ToM and functioning, despite using the same visual measure of ToM used by Barbato et al. (2013). Amminger and colleagues also reported no association between facial emotional recognition and functioning in a UHR sample, though they did identify a modest association between poor vocal affect recognition and poor global functioning (Amminger et al., 2013). However, this study used the Global Assessment of Functioning (GAF) as the outcome variable, which conflates symptoms with functioning. It is therefore unclear whether clinical symptoms or functional impairment drove this association as results from another UHR study that used the same measure of vocal affect recognition found poor performance on this task to be significantly correlated with more severe negative symptoms (Addington et al., 2012).

Methodological considerations

This is the first study to examine the relationship between functioning and all four core domains of social cognition in a single UHR cohort. This was also the first UHR study to examine the association between social cognition and discrete measures of social and role functioning. Global functioning was correlated with both social and role functioning as would be anticipated. However, the discrete social and role functioning measures were weakly and non-significantly correlated, strengthening the argument that these should be examined as separate constructs (Cornblatt et al., 2007; Strassnig et al., 2015).

Limitations of this study include the small sample size and use of only cross-sectional data which did not permit causal links to be made. The small sample size may have led to a lack of power to detect associations as evidenced by the non-significant correlations between neurocognitive measures such as processing speed and verbal memory which have previously been associated with functioning in larger and longitudinal UHR studies (Carrión et al., 2011; Lin et al., 2011). Additionally, because this was an exploratory as opposed to confirmatory study, we did not correct for multiple comparisons which may have increased the potential for Type I error. We also did not account for other variables that have previously been linked to poor functioning in this population, such as childhood trauma (for a review see Cotter et al., 2014; Yung et al., 2015). Longitudinal studies are needed to determine whether ToM is predictive of long-term functional disability in the same way that

negative and disorganized symptoms, cognitive deficits and childhood trauma have been shown to be in the UHR group (Lin et al., 2011; Meyer et al., 2014; Yung et al., 2015).

Clinical implications

Efforts to date to improve functioning among UHR patients have proven largely ineffective (Cotter et al., 2014). Results from this study suggest that social cognitive remediation specifically targeting ToM may be beneficial for improving functioning in this group. Despite ToM involving a series of complex mental operations, efforts to improve ToM in patients with schizophrenia have demonstrated encouraging results (Kurtz and Richardson, 2012; Bechi et al., 2015). Trials are already underway to try and replicate these findings and assess their impact on functioning in the UHR population (Glenthøj et al., 2015). Oxytocin has also been shown to improve performance on higher order ToM tasks in patients with schizophrenia (Woolley et al., 2014; Guastella et al., 2015), though this effect is not always consistent (Cacciotti-Saija et al., 2015). No work has yet been published examining this effect in the UHR group, though it is a promising avenue for future research (Bartholomeusz et al., 2015).

Conclusions

In conclusion, among specific social cognitive abilities only ToM was related to concurrent functioning in our UHR sample. Though these findings should be interpreted with caution due to the small sample size and exploratory nature of the research, the results support previous psychosis literature that has reported ToM as being the strongest social cognitive

predictor of poor functioning. Further longitudinal research into the different ways in which social cognitive skills, as well as other factors, impact on functional outcome in UHR is needed. This will allow for the development of targeted intervention programs, which may be best delivered during the putatively prodromal period to reduce or prevent further functional decline.

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References

Addington J, Cornblatt BA, Cadenhead KS, et al. (2011) At clinical high risk for psychosis: outcome for nonconverters. *American Journal of Psychiatry* 168: 800-805.

Addington J, Piskulic D, Perkins D, et al. (2012) Affect recognition in people at clinical high risk of psychosis. *Schizophrenia Research* 140: 87-92.

Amminger GP, Allott K, Schlögelhofer M, et al. (2013) Affect recognition and functioning in putatively prodromal individuals. *Schizophrenia Research* 147: 404-405.

Andreasen NC (1984) *Scale for the Assessment of Negative Symptoms*. Iowa: University of Iowa Press.

Barbato M, Liu L, Cadenhead KS, et al. (2015) Theory of mind, emotion recognition and social perception in individuals at clinical high risk for psychosis: findings from the NAPLS-2 cohort. *Schizophrenia Research: Cognition* 2: 133-139.

Barbato M, Liu L, Penn DL, et al. (2013) Social cognition as a mediator between neurocognition and functional outcome in individuals at clinical high risk for psychosis. *Schizophrenia Research* 150: 542-546.

Bartholomeusz CF, Ganella EP, Labuschagne I, et al. (2015) Effects of oxytocin and genetic variants on brain and behaviour: implications for treatment in schizophrenia. *Schizophrenia Research*. Epub ahead of print 26 June 2015. DOI: 10.1016/j.schres.2015.06.007.

Bechi M, Bosia M, Spangaro M, et al. (2015) Combined social cognitive and neurocognitive rehabilitation strategies in schizophrenia: neuropsychological and psychopathological influences on theory of mind improvement. *Psychological Medicine* 45: 3147-3157.

Brissos S, Molodynski A, Dias VV, et al. (2011) The importance of measuring psychosocial functioning in schizophrenia. *Annals of General Psychiatry* 10: 18.

Cacciotti-Saija C, Langdon R, Ward PB, et al. (2015) A double-blind randomized controlled trial of oxytocin nasal spray and social cognition training for young people with early psychosis. *Schizophrenia Bulletin* 41: 483-493.

Carrión RE, Goldberg TE, McLaughlin D, et al. (2011) Impact of neurocognition on social and role functioning in individuals at clinical high risk for psychosis. *American Journal of Psychiatry* 168: 806-813.

Corcoran R, Cahill C and Frith CD (1997) The appreciation of visual jokes in people with schizophrenia: a study of 'mentalizing' ability. *Schizophrenia Research* 24: 319-327.

Corcoran R, Mercer G and Frith CD (1995) Schizophrenia, symptomatology and social inference: investigating "theory of mind" in people with schizophrenia. *Schizophrenia Research* 17: 5-13.

Cornblatt BA, Auther AM, Niendam T, et al. (2007) Preliminary findings for two new measures of social and role functioning in the prodromal phase of schizophrenia. *Schizophrenia Bulletin* 33: 688-702.

Corrigan PW and Addis IB (1995) The effects of cognitive complexity on a social sequencing task in schizophrenia. *Schizophrenia Research* 16: 137-144.

Cotter J, Drake RJ, Bucci S, et al. (2014) What drives poor functioning in the at-risk mental state? A systematic review. *Schizophrenia Research* 159: 267-277.

Fett AK, Viechtbauer W, Dominguez MD, et al. (2011) The relationship between neurocognition and social cognition with functional outcomes in schizophrenia: a meta-analysis. *Neuroscience and Biobehavioral Reviews* 35: 573-588.

Fioravanti M, Bianchi V and Cinti ME (2012) Cognitive deficits in schizophrenia: an updated meta-analysis of the scientific evidence. *BMC Psychiatry* 12: 64.

Glenthøj LB, Fagerlund B, Randers L, et al. (2015) The FOCUS trial: cognitive remediation plus standard treatment versus standard treatment for patients at ultra-high risk for psychosis: study protocol for a randomised controlled trial. *Trials* 16: 25.

Gold JM, Carpenter C, Randolph C, et al. (1997) Auditory working memory and Wisconsin Card Sorting Test performance in schizophrenia. *Archives of General Psychiatry* 54: 159-165.

Goldman HH, Skodol AE and Lave TR (1992) Revising axis V for DSM-IV: a review of measures of social functioning. *American Journal of Psychiatry* 149: 1148-1156.

Green MF, Bearden CE, Cannon TD, et al. (2012) Social cognition in schizophrenia, part 1: performance across phase of illness. *Schizophrenia Bulletin* 38: 854-864.

Green MF, Penn DL, Bentall R, et al. (2008) Social cognition in schizophrenia: an NIMH workshop on definitions, assessment, and research opportunities. *Schizophrenia Bulletin* 34: 1211-1220.

Guastella AJ, Ward PB, Hickie IB, et al. (2015) A single dose of oxytocin nasal spray improves higher-order social cognition in schizophrenia. *Schizophrenia Research*. Epub ahead of print 3 July 2015. DOI: 10.1016/j.schres.2015.06.005.

Kurtz MM and Richardson CL (2012) Social cognitive training for schizophrenia: a meta-analytic investigation of controlled research. *Schizophrenia Bulletin* 38: 1092-1104.

Lee TY, Hong SB, Shin NY, et al. (2015) Social cognitive functioning in prodromal psychosis: a meta-analysis. *Schizophrenia Research* 164: 28-34.

Lin A, Wood SJ, Nelson B, et al. (2011) Neurocognitive predictors of functional outcome two to 13 years after identification as ultra-high risk for psychosis. *Schizophrenia Research* 132: 1-7.

Lin A, Wood SJ and Yung AR (2013) Measuring psychosocial outcome is good. *Current Opinion in Psychiatry* 26: 138-143.

Lovibond SH and Lovibond PF (1995) *Manual for the Depression Anxiety Stress Scales*. Sydney: Psychology Foundation.

Meyer EC, Carrión RE, Cornblatt BA, et al. (2014) The relationship of neurocognition and negative symptoms to social and role functioning over time in individuals at clinical high risk in the first phase of the North American Prodrome Longitudinal Study. *Schizophrenia Bulletin* 40: 1452-1461.

Meyer JD, Salovey P and Caruso DR (2002) *Mayer-Salovey-Caruso Emotional Intelligence Test (MSCEIT): User's Manual*. Toronto: Multi-Health Systems Publishers.

Miller TJ, McGlashan TH, Rosen JL, et al. (2002) Prospective diagnosis of the initial prodrome for schizophrenia based on the Structured Interview for Prodromal Syndromes: preliminary

evidence of interrater reliability and predictive validity. *American Journal of Psychiatry* 159: 863-865.

Nuechterlein KH, Green MF, Kern RS, et al. (2008) The MATRICS Consensus Cognitive Battery, part 1: test selection, reliability, and validity. *American Journal of Psychiatry* 165: 203-213.

Nowicki S and Carton J (1993) The measurement of emotional intensity from facial expressions. *Journal of Social Psychology* 133: 749-750.

Nowicki S and Duke MP (1974) A locus of control scale for noncollege as well as college adults. *Journal of Personality Assessment* 38: 136-137.

Nowicki S and Duke MP (1994) Individual differences in the nonverbal communication of affect: the diagnostic analysis of nonverbal accuracy scale. *Journal of Nonverbal Behavior* 18: 9-35.

Overall JE and Gorham DR (1962) The brief psychiatric rating scale. *Psychological Reports* 10: 799-812.

Pinkham AE, Penn DL, Green MF, et al. (2014) The social cognition psychometric evaluation study: results of the expert survey and RAND panel. *Schizophrenia Bulletin* 40: 813-823.

Pinkham AE, Penn DL, Green MF, et al. (2015) Social cognition psychometric evaluation: results of the initial psychometric study. *Schizophrenia Bulletin*. Epub ahead of print 4 May 2015. DOI: 10.1093/schbul/sbv056.

Reitan RM (1955) The relation of the trail making test to organic brain damage. *Journal of Consulting Psychology* 19: 393-394.

Salokangas RK, Nieman DH, Heinimaa M, et al. (2013) Psychosocial outcome in patients at clinical high risk of psychosis: a prospective follow-up. *Social Psychiatry and Psychiatric Epidemiology* 48: 303-311.

Savla GN, Vella L, Armstrong CC, et al. (2013) Deficits in domains of social cognition in schizophrenia: a meta-analysis of the empirical evidence. *Schizophrenia Bulletin* 39: 979-992.

Stanford AD, Messinger J, Malaspina D, et al. (2011) Theory of Mind in patients at clinical high risk for psychosis. *Schizophrenia Research* 131: 11-17.

Strassnig MT, Raykov T, O'Gorman C, et al. (2015) Determinants of different aspects of everyday outcome in schizophrenia: the roles of negative symptoms, cognition, and functional capacity. *Schizophrenia Research* 165: 76-82.

Thompson A, Papas A, Bartholomeusz C, et al. (2012) Social cognition in clinical "at risk" for psychosis and first episode psychosis populations. *Schizophrenia Research* 141: 204-209.

Thompson A, Papas A, Bartholomeusz C, et al. (2013) Externalized attributional bias in the Ultra High Risk (UHR) for psychosis population. *Psychiatry Research* 206: 200-205.

Velthorst E, Nieman DH, Linszen D, et al. (2010) Disability in people clinically at high risk of psychosis. *British Journal of Psychiatry* 197: 278-284.

Wechsler D (1997) *Wechsler Memory Scale - Administration and Scoring Manual*. San Antonio, Texas: Harcourt Assessment.

Wechsler D (1999) *Wechsler Abbreviated Scale of Intelligence (WASI)*. New York: The Psychological Corporation.

Woolley JD, Chuang B, Lam O, et al. (2014) Oxytocin administration enhances controlled social cognition in patients with schizophrenia. *Psychoneuroendocrinology* 47: 116-125.

Yung AR, Cotter J, Wood SJ, et al. (2015) Childhood maltreatment and transition to psychotic disorder independently predict long term functioning in young people at ultra high risk for psychosis. *Psychological Medicine* 45: 3453-3465.

Yung AR, McGorry PD, McFarlane CA, et al. (1996) Monitoring and care of young people at incipient risk of psychosis. *Schizophrenia Bulletin* 22: 283-303.

Yung AR, Nelson B, Thompson A, et al. (2010) The psychosis threshold in Ultra High Risk (prodromal) research: is it valid? *Schizophrenia Research* 120: 1-6.

Yung AR, Phillips LJ, McGorry PD, et al. (1998) Prediction of psychosis. A step towards indicated prevention of schizophrenia. *British Journal of Psychiatry Supplement* 172: 14-20.

Yung AR, Phillips LJ, Yuen HP, et al. (2004) Risk factors for psychosis in an ultra high-risk group: psychopathology and clinical features. *Schizophrenia Research* 67: 131-142.

Yung AR, Yuen HP, McGorry PD, et al. (2005) Mapping the onset of psychosis: the Comprehensive Assessment of At-Risk Mental States. *Australian and New Zealand Journal of Psychiatry* 39: 964-971.

Table 1. Sample characteristics

	<i>M (SD)</i>
Age (years)	19.1 (2.8)
Years of education	12.0 (1.9)
BPRS psychotic subscale	8.8 (3.2)
SANS total	23.6 (15.0)
DASS total	57.2 (28.0)
DASS – depression subscale	19.5 (12.8)
DASS – anxiety subscale	15.0 (8.6)
DASS – stress subscale	22.7 (10.4)
<i>Functioning assessments</i>	
GFR	6.3 (2.07)
GFS	6.3 (1.27)
SOFAS	60.7 (11.1)
<i>Social cognitive measures</i>	
DANVA-2 – faces (total errors)	5.2 (2.47)
DANVA-2 – voices (total errors)	6.6 (2.42)
Hinting task	16.5 (3.11)
Visual jokes – ToM subset	9.9 (3.25)
MSCEIT – managing emotions module	89.0 (10.33)
SCST-R (juxtaposition score)	0.85 (0.09)
ANSIE – locus of control score	18.3 (7.2)
Composite social cognitive score	-0.50 (1.02) ^a
<i>Neurocognitive measures</i>	
Estimated IQ (WASI)	103.3 (16.0)
Trail Making A (total time)	26.6 (8.6)
Trail Making B (total time)	63.6 (22.3)
Spatial span test (total score)	16.7 (2.1)
Letter number sequencing test	15.2 (3.8)
	<i>N</i>
Sex (male/female)	14/16
<i>Medication use</i>	
Antidepressants	6
Antipsychotics	3
Mood stabilizers	1

^a The mean value for the composite score is presented as a z score relative to a healthy control group

Abbreviations: ANSIE – Adult Nowicki Strickland Internal External scale; BPRS – Brief Psychiatric Rating Scale; DANVA-2 – Diagnostic Analysis of Nonverbal Accuracy-2; DASS - Depression Anxiety Stress Scale; GFR – Global Functioning Role scale; GFS – Global Functioning Social scale; MSCEIT – Mayer-Salovey-Caruso Emotional Intelligence Test; SANS – Scale for the Assessment of Negative Symptoms; SCST-R – Schema Component Sequencing Task Revised; SOFAS – Social and Occupational Functioning Assessment Scale; ToM – Theory of Mind; WASI – Wechsler Abbreviated Scale of Intelligence

Table 2. Correlations between functioning measures, clinical and demographic variables and social cognitive and neurocognitive performance

	GFR		GFS		SOFAS	
	<i>r</i>	<i>p</i>	<i>r</i>	<i>p</i>	<i>r</i>	<i>p</i>
Age	.190	.315	-.131	.491	.174	.357
Years of education	.440	.015*	.123	.518	.325	.080
BPRS psychotic subscale	.125	.512	-.267	.154	-.191	.311
SANS total	-.454	.012*	-.437	.016*	-.651	<.001**
DASS total	-.020	.915	-.355	.054	-.352	.057
DASS – depression subscale	.028	.884	-.316	.089	-.303	.103
DASS – anxiety subscale	.110	.563	-.261	.163	-.242	.198
DASS – stress subscale	-.181	.339	-.349	.058	-.372	.043*
DANVA-2 – faces (total errors)	.020	.918	-.015	.939	.185	.327
DANVA-2 – voices (total errors)	-.166	.380	-.221	.240	-.176	.351
Hinting task	.062	.746	.193	.308	.206	.275
Visual jokes – ToM subset	.425	.019*	.276	.140	.540	.002**
MSCEIT – managing emotions module	-.117	.539	.277	.139	.360	.051
SCST-R (juxtaposition score)	.027	.889	.013	.946	-.170	.369
ANSIE – locus of control score	.034	.859	-.218	.247	-.315	.090
Composite social cognitive score	.145	.446	.240	.202	.235	.211
Estimated IQ (WASI)	.173	.361	.117	.539	.233	.215
Trail Making A (total time)	-.321	.084	-.081	.671	-.143	.450
Trail Making B (total time)	-.342	.064	-.016	.934	-.181	.339
Spatial span test (total score)	.306	.100	-.155	.413	-.029	.881
Letter number sequencing test	.264	.158	.223	.236	.335	.070

**p* < .05

***p* < .01

Abbreviations: ANSIE – Adult Nowicki Strickland Internal External scale; BPRS – Brief Psychiatric Rating Scale; DANVA-2 – Diagnostic Analysis of Nonverbal Accuracy-2; DASS - Depression Anxiety Stress Scale; GFR – Global Functioning Role scale; GFS – Global Functioning Social scale; MSCEIT – Mayer-Salovey-Caruso Emotional Intelligence Test; SANS – Scale for the Assessment of Negative Symptoms; SCST-R – Schema Component Sequencing Task Revised; SOFAS – Social and Occupational Functioning Assessment Scale; ToM – Theory of Mind; Wechsler Abbreviated Scale of Intelligence

Table 3. Multivariate analyses of variables associated with the SOFAS

Model 1	<i>B (SE)</i>	β	<i>t</i>	<i>p</i>
SANS total	-.346 (.105)	-.466	-3.280	.003**
DASS – stress subscale	-.206 (.145)	-.192	-1.425	.166
Visual jokes – ToM subset	1.304 (.458)	.381	2.845	.009**
Model 2				
SANS total	-.414 (.117)	-.557	-3.548	.002**
DASS – stress subscale	-.156 (.166)	-.145	-.937	.357
MSCEIT – managing emotions module	.151 (.166)	.140	.909	.372

p* < .05*p* < .01**Note:** β : Standardized regression coefficients; B: Unstandardized regression coefficients**Abbreviations:** DASS - Depression Anxiety Stress Scale; MSCEIT – Mayer-Salovey-Caruso Emotional Intelligence Test; SANS – Scale for the Assessment of Negative Symptoms; SOFAS – Social and Occupational Functioning Assessment Scale; ToM – Theory of Mind**Table 4.** Multivariate analyses of variables associated with the GFR scale

	<i>B (SE)</i>	β	<i>t</i>	<i>p</i>
Years of education	.296 (.188)	.274	1.576	.127
SANS total	-.044 (.023)	-.320	-1.908	.067
Visual jokes – ToM subset	.135 (.113)	.211	1.188	.245

Note: β : Standardized regression coefficients; B: Unstandardized regression coefficients**Abbreviations:** GFR – Global Functioning Role scale; SANS – Scale for the Assessment of Negative Symptoms; ToM – Theory of Mind