

A critical review of metacognitive training (MCT) for psychosis: Efficacy, proposed mechanisms of action and significance for functional outcomes

BC Schneider^{1*}, C Andreou¹

Abstract

Introduction

Despite significant advances in antipsychotic medications, schizophrenia continues to be a highly disabling condition. Many patients go untreated, are treatment resistant or receive inadequate treatment. A large body of work has demonstrated that cognitive biases (e.g., jumping to conclusions [JTC], overconfidence in errors, bias against disconfirmatory evidence) are involved in the formation of delusions in schizophrenia and thus represent a possible target for intervention. The following article provides a critical review of metacognitive training (MCT) for psychosis.

Discussion

MCT is a variant of cognitive behavioural therapy (CBT) that seeks to both normalize and increase awareness about cognitive processes involved in the emergence of symptoms utilizing strategies that induce doubt in delusions by addressing cognitive biases, while also instilling hope. Randomized controlled trials using blind raters have mainly yielded moderate effects, which in one study were maintained up to three years after treatment. MCT is positively rated by patients signaling that it is useful, fun and the rationale is understood by patients. Further work is needed to clarify whether the JTC bias represents a unique mechanism of action of MCT that may account, in part, for its effects beyond antipsychotic medication.

*Corresponding author
Email: b.viertel@uke.de

¹ University Hospital Hamburg-Eppendorf, Hamburg, Germany

In contrast, overconfidence in errors may be targeted by antipsychotic medications through dopamine modulation as well as through MCT's cognitive-behavioral strategies. MCT may lead to improvement beyond psychotic symptoms as initial evidence indicates that improvement in the JTC bias contributes to better vocational outcomes and improved social relationships.

Conclusion

We conclude that MCT represents a promising new direction in intervention research for psychosis that can complement standard treatment approaches.

Introduction

Schizophrenia is a chronic disorder that affects approximately 1% of the population worldwide¹ and is among the world's most disabling conditions².

Despite significant improvements in antipsychotics and treatment approaches for schizophrenia, their effect on symptom reduction remains moderate³. Due to factors including limited insight, memory problems, adverse effects, and mistrust of the clinician, discontinuation rates of antipsychotic medication are typically quite high^{4,5}. Additionally, up to 30% of patients are treatment resistant⁶ and less than 15% of all patients recover completely⁷.

These observations have led to increased interest in adjunctive psychological treatments that might reduce symptoms resistant to or not effectively treated by medication. Indeed, treatments such as cognitive-behavioural therapy for positive symptoms (CBTp)⁸ and cognitive remediation (CR)⁹ have yielded small to medium effects on symptoms

beyond antipsychotic medication alone. Metacognitive training (MCT)^{10,11} draws from concepts used in CBT, CR and psychoeducation.

Like these approaches, it aims to target psychotic symptoms, but does so through a "backdoor approach" that seeks to both normalize and increase awareness about cognitive processes involved in the emergence of symptoms. MCT is founded upon a huge body of empirical work that has demonstrated generalized cognitive biases in schizophrenia, including jumping to conclusions^{12,13,14} (Figure 1), bias against discriminatory evidence (BADE)^{15,16}, attributional distortions^{17,18}, and overconfidence in errors¹⁹ (Figure 2). MCT also targets Theory of Mind (ToM) deficits that are common in psychosis²⁰.

In particular, delusions, defined as fixed false beliefs that are held with high conviction, are among the most challenging symptoms to treat and were once thought to be psychologically inaccessible^{21,22}. Rather than viewing delusions as absurd thoughts that emerge without a meaningful context, the concept of cognitive biases implies that delusions result from specific disruptions in the normal cognitive processes for belief generation and evaluation, and thus might be amenable to specific psychological interventions²³.

Though antipsychotic medication creates detachment from delusions, potentially through a "numbing" effect²⁴, it does not directly alter underlying cognitive processes; thus, MCT aims to address delusions in a manner complementary to that of antipsychotic medication. We will present a hypothesis of the proposed "mechanism of action" used by MCT to address cognitive biases, and therefore

delusions, in the second half of this review.

Discussion

The authors have referenced some of their own studies in this review. These referenced studies have been conducted in accordance with the Declaration of Helsinki (1964) and the protocols of these studies have been approved by the relevant ethics committees related to the institution in which they were performed. All human subjects, in these referenced studies, gave informed consent to participate in these studies.

Metacognitive Training for Psychosis (MCT)

As MCT has been extensively described elsewhere^{10,25,26}, only a brief description of the modules and aims will be provided here.

The training (2 sets of 8 modules each for most versions) aims to sow the “seed of doubt” in patients in an entertaining, playful and collaborative manner through corrective experiences. Cognitive biases are addressed through the eight modules, including: attributional distortions (module 1), jumping to conclusions (modules 2 and 7; Figure 1), bias against disconfirmatory evidence (module 3), and over-confidence in memory errors (module 5; Figure 2).

Two modules (4 and 6) deal with deficits in theory of mind and the importance of cognitive biases when making social inferences, in light of research suggesting a central role of social cognition deficits for functional outcome in schizophrenia²⁷.

To address the high rates of affective disturbance (approx. 50%)²⁸, depressive cognitive patterns are also discussed (module 8). An example of topics covered in Modules 2 and 7 is presented in table 1.

To avoid directly confronting delusions and dysfunctional coping skills, each module begins with psychoeducational elements and attempts to “normalize” cognitive distortions through examples of the fallibility of human cognition that all people experience. This is an

important feature, as normalization has been demonstrated to foster treatment engagement for psychotherapy²⁹.

Using a dialectic approach, the next step of MCT examines how exaggerated (pathological) fallacies of normal thinking can lead to problems in daily life and possibly delusions in individuals with schizophrenia. These fallacies are illustrated with real-world examples of people with psychosis, and opportunities for group participants to exchange their own experiences are provided.

Therefore, differing from the traditional CBT approach, rather than beginning at the level of symptoms, the “seed of doubt” is sowed in a neutral context with the hope that through multiple examples and discussion of personal experiences, patients will gradually gain insight into the dysfunctional nature of their delusions. The ultimate goal is to provide patients with practical strategies while at the same time instilling hope. An individualized version of the intervention also exists³⁰ (MCT+, free download at www.uke.de/mct_plus) during which patients may explore specific symptoms and target cognitive biases most relevant for them. MCT is currently available at no cost in currently 30 languages.

Current state of the literature

While most studies have examined the standard group training, abbreviated or modified versions of the therapy have also been used, such as those that have blended MCT with other interventions or have used MCT+.

Overall, the effect sizes yielded by these studies are comparable to that of other therapeutic approaches to the treatment of schizophrenia³¹. Providing further evidence of the potential for MCT as a complementary intervention, all studies used MCT as an add-on treatment to antipsychotic medications. This suggests that MCT's effects may target a separate set of processes than antipsychotics or perhaps enhance medication effects. To this end, our group is currently examining the effects of MCT on

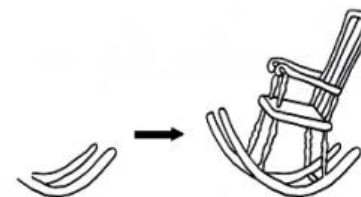


Figure 1: Elephant tusks or a rocking chair? In module 2 (jumping to conclusions I), participants are shown a picture in which an object is depicted with increasing detail. Participants are asked to indicate when they are ready to make a decision about what the object is. Hasty decision-making often leads to incorrect judgments on this task.

unmedicated patients. Following is a brief review of the efficacy of MCT for positive symptoms (for a detailed review of all existing and forthcoming studies on the efficacy of MCT, see Moritz et al.³²).

Of note, as MCT was not designed to target negative symptoms³², results on these symptoms have generally not been reported.

Feasibility/Safety/Subjective Appraisal

Several studies have demonstrated the feasibility and safety of the group MCT^{33,34,35}. Subjective ratings of MCT by patients have been overall positive with patients indicating that they feel as if the training is useful, interesting and an important part of their recovery program^{34,35,36,37,38,39}.

For example, in the study by Moritz, Kerstan et al., most (85% and 92.9%, respectively) participants indicated that they would recommend the training to others and found the training to be useful and sensible³⁹. Given findings that patient ratings of the logicalness of CBT is associated with outcome⁴⁰, this patient feedback regarding their experiences with MCT bodes well for the training's effectiveness.

Positive Symptoms

Several short-term studies have demonstrated that MCT has an effect on positive symptoms in schizophrenia with most studies yielding moderate effect sizes ($d \geq 0.41$)^{34,38,39,41}.

MCT effects include improvements in positive symptoms as measured by the Positive and Negative Syndrome Scale (PANSS)^{34,35,37,39,41}, and fewer delusions, as measured by the Psychotic Symptoms Rating Scale (PSYRATS)^{34,38,39}.

Declines in delusion conviction and improved delusion awareness³⁴, as well as delusion distress³⁹, are also reported. MCT participants improve significantly more in these areas than participants in active control groups including treatment as usual³⁹ and CR³⁸.

These short-term studies encouraged the design of larger randomized, controlled, rater-blind studies on the long-term effects of MCT. Two studies have been published to date, which used quite different patient samples; while one study included a sample of patients with schizophrenia spectrum disorders of mixed severity ($n = 150$)¹¹, the other used more restrictive criteria that included participants only with schizophrenia or schizoaffective disorders, as well as those with persistent delusions⁴². Both studies reported improvements in indices of delusions and positive symptoms in participants receiving MCT compared to control groups at 6-months ($d=0.51$ to 0.64). Impressively, effect sizes for differences in overall delusions, as well as delusion conviction and delusion awareness between patients in MCT and TAU⁴² and cognitive training¹¹ were maintained or even increased in both studies at 6-months. A recent 3-year follow-up study also found differences in delusions (PSYRATS) and positive symptoms (PANSS); however, for the first time, there was also a significant difference in the PANSS total score⁴³.

It should be made clear that not all studies have found significant group effects^{35,39,41}. Further work is needed to confirm the long-term effects; however, these preliminary findings suggest that the effects of MCT may lead to lasting recovery³². Critically, the robustness of the findings is further supported by the fact that, in



Figure 2: In module 5 (memory), participants are asked to study a complex scene for several seconds and then identify which items on a list were or were not present in the scene. They are also asked to rate how confident they are in their decisions. Patients with psychosis tend to be more confident in their memory errors.

all of these studies, raters were blind to treatment arm allocation.

“Mechanism of Action” of MCT

Despite the demonstrated effectiveness of MCT, its exact mechanism of action has yet to be confirmed. In some^{35,36,38,39,43}, but not all¹¹, of the above studies the beneficial effects of metacognitive interventions on symptoms were accompanied by improvements in delusion-associated cognitive biases, thus providing preliminary support for the hypothesized mechanisms of action of MCT. However, no study has yet directly assessed the effect of the intervention on cognitive biases as a modulator of symptom change. In this section, we first discuss the neurobiological underpinnings of cognitive biases before turning to MCT’s proposed “mechanism of action.”

In contrast to the abundant literature on cognitive biases in schizophrenia, very few studies have examined their neurobiological underpinnings and their connection to the postulated dopaminergic disturbances underlying psychotic symptoms.

These areas may provide insight into the different mechanisms of action for MCT versus antipsychotic medications and clarify MCT’s complementary

effect in medicated patients. Recent evidence suggests that the JTC bias is not affected by antipsychotic medication⁴⁴, although it may act as a moderator of antipsychotic drug response^{44,45}. However, there is evidence that overconfidence in errors correlates negatively with antipsychotic medication dose^{46,47} which might imply that it is modulated by dopaminergic activity.

Indeed, a recent study by our group³⁷ found that a dopaminergic agonist (L-Dopa) and a dopaminergic antagonist (the typical antipsychotic haloperidol) did not affect probabilistic reasoning (which is involved in the JTC bias) in healthy subjects, whereas there was a positive (reducing) effect of haloperidol on error overconfidence compared to L-Dopa.

It is also unknown how cognitive biases relate to the prominent account of delusions that proposes that they result from a (dopaminergically induced) aberrant allocation of salience to random stimuli. The pattern of dopaminergic modulation described above³⁷ is consistent with a dual-disturbance account of delusion formation²³ which postulates that two types of cognitive disturbances are needed to co-occur in order for delusions to emerge: The first disturbance in the so-called “2-Factor

Table 1: MCT Example.

Jumping to Conclusions (Modules 2 and 7)

- **Introduction:** what does “jumping to conclusions” (JTC) mean? What are hasty conclusions? What are the advantages and disadvantages of making hasty conclusions?
- **Examples of JTC in everyday life:** examples of when most people make inferences without having enough proof and discussion of whether this is justified (i.e., does hearing bushes rustling mean there is a burglar outside?); considering multiple pieces of information.
- **Examination of JTC “in action”:** real-world examples of JTCs, such as commonly accepted urban legends (e.g., Paul McCartney is dead); evidence for and against these urban legends are discussed.
- **Why are we doing this?:** explanation of how JTC can lead to misinterpretations, particularly in people with psychosis.
- **Cognitive exercises: examples of JTC:** ‘What do you see in the picture?’ (Figure 1). Gradual presentation of drawings; use of evidence gathering to interpret optical illusions and paintings. Examination of how hasty conclusions can lead to misinterpretations.
- **Learning points:** review of the major themes from the JTC module, transfer to everyday life when making small (e.g., grocery shopping) or crucial (e.g., confronting a neighbor for supposedly spying) decisions.

Theory²³ is the generation of an implausible thought, and the second disturbance explains why this thought is uncritically accepted as being true. Thus, it might be assumed that the first type of disturbance corresponds to aberrant salience, whereas the JTC bias comes into play in the second step of this process. It is not clear how overconfidence in errors fits into this account: Conceptually, it would play a role at a later stage of delusion consolidation. However, the association of both overconfidence in errors and salience with dopaminergic activity raises the probability that these disturbances reflect two facets of the same core deficit. Indeed, recent functional neuroimaging studies suggest that subjective confidence may reflect self-generated dopaminergic signals in the reward system⁴⁸. Though not yet tested, this may provide a possible link between overconfidence and salience.

In summary, some initial evidence suggests that the effects of MCT on the JTC bias may be considered unique and independent of those of antipsychotic medication, whereby this has not been consistently supported. In contrast, MCT’s effects on error overconfidence might be similar or complementary to those of medication. However, these hypotheses are preliminary, as a direct comparison of MCT as monotherapy versus antipsychotic drug treatment is lacking.

Functional Deficits

Functional deficits in schizophrenia have been well-established and schizophrenia ranks sixth among the top 10 causes of disability². MCT has been shown to be associated with improvements in social relationships³⁹, depression^{34,43}, quality of life⁴³ and, in a forensic patient population, functional mental capacity (i.e., the ability to exercise legal rights)⁴⁹.

The aforementioned cognitive biases targeted by MCT may represent the link between functional deficits and specific symptoms or behaviours in schizophrenia. For example, in a study by Köther et al.⁵⁰, patients with schizophrenia not only made more false judgments than healthy controls in interpreting the mental state of others, but they also made significantly more high-confident errors and demonstrated decreased insight into their impairments⁵⁰. This may very well have functional implications: A higher conviction in false judgments about the intentions or thoughts of others, especially when accompanied by delusions or paranoia, may lead to behavioural consequences (e.g., withdrawal, confrontation) and emotional distress (e.g., panic, anger)⁵¹.

For example, Moritz & Van Quaquebeke⁵¹ found that conviction in one’s paranoid beliefs (i.e., being pursued by the secret service), directly related to the subsequent actions (i.e., deleting internet traces [low conviction], buying a bullet-proof

vest [high conviction] or even arming oneself [very high conviction]). The assumption that cognitive biases are relevant for functional outcomes is supported by recent work on JTC and functioning by our group demonstrating that improvements in JTC positively affected vocational outcomes⁵². Therefore, though the main aim of MCT is cognitive biases, change in cognitive biases may lead to improved psychosocial functioning.

Conclusion

Taken together, MCT represents a new and promising direction for schizophrenia treatment that can complement existing treatment approaches.

Though a number of studies have demonstrated the feasibility, acceptance and effectiveness of MCT on positive symptoms, findings are preliminary and most sample sizes remain small.

However, moderate effects suggest that MCT leads to increased improvement beyond antipsychotic medication, and preliminary findings from one long-term study indicate that this improvement is long-lasting. Current studies examining the efficacy of MCT in unmedicated patients, as well as studies using updated, shortened or blended versions of MCT and MCT+ will provide clearer evidence of MCT’s effects.

Much remains to be clarified about MCT’s mechanism of action and we hope that future work will be aimed at

Competing interests: None declared. Conflict of interests: None declared. All authors contributed to conception and design, manuscript preparation, read and approved the final manuscript. All authors abide by the Association for Medical Ethics (AME) ethical rules of disclosure.

further identifying the connection between changes in positive symptoms, cognitive biases (particularly, if and how cognitive biases other than jumping to conclusions may be effected by MCT) and functional status.

Funding

C. Andreou was supported by the Brain and Behavior Research Foundation (NARSAD Grant Nr. 18749). The funding body had no role in the design; collection, analysis and/or interpretation of data; in the writing of the manuscript; and in the decision for submission.

Acknowledgment

We would like to thank Prof. Dr. Steffen Moritz for his assistance in preparation of the manuscript.

References

1. Saha S, Chant D, Welham J, McGrath J. A systematic review of the prevalence of schizophrenia. *PLoS Med.* 2005 May;2(5):e141.
2. Mathers C, Fat B, Boerma J. The global burden of disease: 2004 update. Geneva, Switzerland: World Health Organization Press; 2008.
3. Leucht S, Arbter D, Engel RR, Kissling W, Davis JM. How effective are second-generation antipsychotic drugs? A meta-analysis of placebo-controlled trials. *Mol. Psychiatry.* 2009 Apr;14(4):429–47.
4. Moritz S, Peters MJV, Karow A, Deljkovic A, Tonn P, Naber D. Cure or curse? Ambivalent attitudes towards neuroleptic medication in schizophrenia and non-schizophrenia patients. *Mental Illness.* 2009;1(1): e2.
5. Lambert M, Conus P, Cotton S, Robinson J, McGorry PD, Schimmelmann BG. Prevalence, predictors, and consequences of long-term refusal of antipsychotic treatment in first-episode psychosis. *J. Clin. Psychopharmacol.* 2010 Oct;30(5):565–72.
6. Kennedy JL, Anothony AC, Taylor DL, Degtiar I, Hornberger JC. The social and economic burden of treatment-resistant schizophrenia: A systematic literature review. *Int. Clin.*

Psychopharmacol. 2014 Mar; 29(2):63–76.

7. Jääskeläinen E, Juola P, Hirvonen N, McGrath JJ, Saha S, Isohanni M, et al. A systematic review and meta-analysis of recovery in schizophrenia. *Schizophr. Bull.* 2013 Nov; 39(6):1296–306.
8. Lincoln TM, Ziegler M, Mehl S, Kesting M-L, Lüllmann E, Westermann S, et al. Moving from efficacy to effectiveness in cognitive behavioral therapy for psychosis: a randomized clinical practice trial. *J. Consult. Clin. Psychol.* 2012 Aug; 80(4):674–86.
9. Wykes T, Huddy V, Cellard C, McGurk SR, Czobor P. A meta-analysis of cognitive remediation for schizophrenia: methodology and effect sizes. *Am. J. Psychiatry.* 2011 May; 168(5):472–85.
10. Moritz S, Woodward TS, Burlon M. *Metacognitives Training für schizophrene Patienten (MKT)*. 1st ed. Hamburg: VanHam Campus; 2003. German. Available as free download: <http://www.uke.de/mct>
11. Moritz S, Veckenstedt R, Bohn F, Hottenrott B, Scheu F, Randjbar S, et al. Complementary group Metacognitive Training (MCT) reduces delusional ideation in schizophrenia. *Schizophr. Res.* 2013 Dec; 151(1-3):61–9.
12. Lincoln TM, Ziegler M, Mehl S, Winfried R. The jumping to conclusions bias in delusions: Specificity and changeability. *J. Abnorm. Psychol.* 2010 Feb; 119(1):40–9.
13. Fine C, Gardner M, Craigie J, Gold I. Hopping, skipping or jumping to conclusions? Clarifying the role of the JTC bias in delusions. *Cogn. Neuropsychiatry.* 2007 Jan; 12(1):46–77.
14. Garety PAM, Hemsley DR, Wessely SR. Reasoning in deluded schizophrenic and paranoid patients: Biases in performance on a probabilistic inference task. *J. Nerv. Ment. Dis.* 1991; 179 (4):194–201.
15. Woodward TS, Moritz S, Menon M, Klinge R. Belief inflexibility in schizophrenia. *Cogn. Neuropsychiatry.* 2008 May; 13(3):267–77.
16. Woodward TS, Buchy L, Moritz S, Liotti M. A bias against

disconfirmatory evidence is associated with delusion proneness in a nonclinical sample. *Schizophr. Bull.* 2007 Jul; 33(4):1023–8.

17. Bentall RP, Corcoran R, Howard R, Blackwood N, Kinderman P. Persecutory delusions: a review and theoretical integration. *Clin. Psychol. Rev.* 2001 Nov; 21(8):1143–92.
18. Randjbar S, Veckenstedt R, Vitzthum F, Hottenrott B, Moritz S. Attributional biases in paranoid schizophrenia: Further evidence for a decreased sense of self-causation in paranoia. *Psychosis.* 2011; 3(1):74–85.
19. Moritz S, Woodward TS. Metacognitive control over false memories: A key determinant of delusional thinking. *Curr. Psychiatry Rep.* 2006; 8(3):184–90.
20. Savla GN, Vella L, Armstrong CC, Penn DL, Twamley EW. Deficits in domains of social cognition in schizophrenia: a meta-analysis of the empirical evidence. *Schizophr. Bull.* 2013 Sep; 39(5):979–92.
21. Jaspers K. *Allgemeine Psychopathologie*. Berlin, Germany: Springer; 1946. German.
22. Walker C. Delusion: What did Jaspers really say? *Br. J. Psychiatry.* 1991 Nov, 159 (14):94–103.
23. Langdon R, Ward PB, Coltheart M. Reasoning anomalies associated with delusions in schizophrenia. *Schizophr. Bull.* 2010 Mar;36(2):321–30.
24. Mizrahi R, Kiang M, Mamo DC, Arenovich T, Bagby RM, Zipursky RB, et al. The selective effect of antipsychotics on the different dimensions of the experience of psychosis in schizophrenia spectrum disorders. *Schizophr. Res.* 2006 Dec;88(1-3):111–8.
25. Moritz S, Vitzthum F, Randjbar S, Veckenstedt R, Woodward TS. Detecting and defusing cognitive traps: metacognitive intervention in schizophrenia. *Curr. Opin. Psychiatry.* 2010 Nov; 23(6):561–9.
26. Moritz S, Veckenstedt R, Bohn F, Köther U, Woodward TS. Metacognitive training in schizophrenia. Theoretical rationale and administration. In: Roberts DL, Penn DL, editors. *Soc. Cogn. Schizophr* 1st ed. New York, NY: Oxford University Press; 2013. p358–83.

Competing interests: None declared. Conflict of interests: None declared. All authors contributed to conception and design, manuscript preparation, read and approved the final manuscript. All authors abide by the Association for Medical Ethics (AME) ethical rules of disclosure.

27. Fett A-KJ, Viechtbauer W, Dominguez M-G, Penn DL, van Os J, Krabbendam L. The relationship between neurocognition and social cognition with functional outcomes in schizophrenia: A meta-analysis. *Neurosci. Behav. Rev.* 2011 Jan; 35(3):573–88.
28. Buckley PF, Miller BJ, Lehrer DS, Castle DJ. Psychiatric comorbidities and schizophrenia. *Schizophr. Bull.* 2009 Mar;35(2):383–402.
29. Lüllmann E, Lincoln TM. The Effect of an educating versus normalizing approach on treatment motivation in patients presenting with delusions: An experimental investigation with analogue patients. *Schizophr Res Treatment.* 2013.
30. Vitzthum FB, Veckenstedt R, Moritz S. Individualized metacognitive therapy program for patients with psychosis (MCT+): introduction of a novel approach for psychotic symptoms. *Behav. Cogn. Psychother.* 2014 Jan;42(1):105–10.
31. Turner DT, van der Gaag M, Karyotaki E, Cuijpers P. Psychological interventions for psychosis: A meta-analysis of comparative outcome studies. *Am. J. Psychiatry.* 2014 May; 171(5):523–538.
32. Moritz S, Andreou C, Schneider BC, Wittekind CE, Menon M, Balzan RP, et al. Sowing the seeds of doubt: A narrative review on metacognitive training in schizophrenia. *Clin. Psychol. Rev.* 2014 June; 34(4):358–366.
33. Moritz S, Woodward TS. Metacognitive Training for Schizophrenia Patients (MCT): A pilot study on feasibility, treatment adherence, and subjective efficacy. *German. J. Psychiatry.* 2007 ;10 (3): 69–78.
34. Favrod J, Maire A, Bardy S, Pernier S, Bonsack C. Improving insight into delusions: a pilot study of metacognitive training for patients with schizophrenia. *J. Adv. Nurs.* 2011 Feb;67(2):401–7.
35. Aghotor J, Pfueller U, Moritz S, Weisbrod M, Roesch-Ely D. Metacognitive training for patients with schizophrenia (MCT): feasibility and preliminary evidence for its efficacy. *J. Behav. Ther. Exp. Psychiatry.* 2010 Sep;41(3):207–11.
36. Balzan RP, Delfabbro PH, Galletly CA, Woodward TS. Metacognitive training for patients with schizophrenia: Preliminary evidence for a targeted, single-module programme. *Aust. N. Z. J. Psychiatry.* Forthcoming 2014.
37. Andreou C, Moritz S, Veith K, Veckenstedt R, Naber D. Dopaminergic Modulation of Probabilistic Reasoning and Overconfidence in Errors: A Double-Blind Study. *Schizophr. Bull.* Forthcoming 2014.
38. Moritz S, Veckenstedt R, Randjbar S, Vitzthum F, Woodward TS. Antipsychotic treatment beyond antipsychotics: metacognitive intervention for schizophrenia patients improves delusional symptoms. *Psychol. Med.* 2011 Sep; 41(9):1823–32.
39. Moritz S, Kerstan A, Veckenstedt R, Randjbar S, Vitzthum F, Schmidt C, et al. Further evidence for the efficacy of a metacognitive group training in schizophrenia. *Behav. Res. Ther.* 2011 Mar; 49(3):151–7.
40. Carter JD, Luty SE, McKenzie JM, Mulder RT, Frampton CM, Joyce PR. Patient predictors of response to cognitive behaviour therapy and interpersonal psychotherapy in a randomised clinical trial for depression. *J. Affect. Disord.* 2011 Feb; 128(3):252–61.
41. Kumar D, Zia Ul Haq M, Dubey I, Dotivala KN, Veqar Siddiqui S, Prakash R, et al. Effect of meta-cognitive training in the reduction of positive symptoms in schizophrenia. *Eur. J. Psychother. Couns.* 2010 Jun; 12(2):149–58.
42. Favrod J, Rexhaj S, Bardy S, Ferrari P, Hayoz C, Moritz S, et al. Sustained antipsychotic effect of metacognitive training in psychosis: A randomized-controlled study. *Eur. Psychiatry.* 2014 June;29(5):275–281.
43. Moritz S, Veckenstedt R, Bohn F, Hottenrott B, Leighton L, Scheu F, et al. Immediate, delayed and “sleeper” effects of metacognitive group training (MCT) in psychosis. A three-year follow-up investigation. *Arch. Gen. Psychiatry.* Forthcoming.
44. So SH, Garety P a, Peters ER, Kapur S. Do antipsychotics improve reasoning biases? A review. *Psychosom. Med.* 2010 Sep;72(7) :681–93.
45. Menon M, Mizrahi R, Kapur S. “Jumping to conclusions” and delusions in psychosis: relationship and response to treatment. *Schizophr. Res.* 2008 Jan; 98(1-3):225–31.
46. Moritz S, Woodward TS, Jelinek L, Klinge R. Memory and metamemory in schizophrenia: a liberal acceptance account of psychosis. *Psychol. Med.* 2008 Jun;38(6):825–32.
47. Moritz S, Woodward TS, Ruff CC. Source monitoring and memory confidence in schizophrenia. *Psychol. Med.* 2003 Jan;33(1):131–9.
48. Schwarze U, Bingel U, Badre D, Sommer T. Ventral striatal activity correlates with memory confidence for old- and new-responses in a difficult recognition test. *PLoS One.* 2013; 3(3):e54324.
49. Naughton M, Nulty A, Abidin Z, Davoren M, O’Dwyer S, Kennedy HG. Effects of group metacognitive training (MCT) on mental capacity and functioning in patients with psychosis in a secure forensic psychiatric hospital: a prospective-cohort waiting list controlled study. *BMC Res.;* 2012 Jan;5(1):302.
50. Köther U, Veckenstedt R, Vitzthum F, Roesch-Ely D, Pfueller U. Scheu, F et al. „Don’t give me that look“ – overconfidence in false mental state perception in schizophrenia. *Psych. Res.* 2012 March; 196(1): 1-8.
51. Moritz S, Van Quaquebeke N. Are you sure? Delusion conviction moderates the behavioural and emotional consequences of paranoid ideas. *Cogn. Neuropsychiatry.* 2014 Jan;19(2):164–80.
52. Andreou C, Treszl A, Roesch-Ely D, Köther U, Veckenstedt R, Moritz S. Investigation of the role of jumping-to-conclusions bias for short-term functional outcome in schizophrenia. *Psychiatry Res.* 2014 Aug;218(30): 341–347.

Competing interests: None declared. Conflict of interests: None declared.
All authors contributed to conception and design, manuscript preparation, read and approved the final manuscript.
All authors abide by the Association for Medical Ethics (AME) ethical rules of disclosure.