Partially restored resting-state functional connectivity in women recovered from anorexia nervosa

Ilka Boehm, MSc; Daniel Geisler, MSc; Friederike Tam, MD; Joseph A. King, MSc; Franziska Ritschel, MSc; Maria Seidel, MSc; Fabio Bernardoni, PhD; Julia Murr, MD; Thomas Goschke, PhD; Vince D. Calhoun, PhD; Veit Roessner, MD; Stefan Ehrlich, MD

Introduction

Anorexia nervosa is a severe psychiatric disorder hallmarked by extreme weight loss (or failure to gain weight during growth) due to relentless control of food intake and has been associated with structural1 as well as functional brain alterations.2 Structural neuroimaging studies have reported grey matter atrophy in acutely ill patients to be largely reversible in weight-restored patients. On the level of brain functioning, abnormal neural responses in the lateral–frontal brain circuitry have been associated with excessive cognitive control in acutely ill as well as recovered patients.5–7

Despite these findings, we still lack understanding of the neural mechanisms underlying anorexia nervosa. One reason might be that the mainstay of studies using fMRI has been the identification of associations between specific brain regions with disorder-related dysfunction. However, this approach of mapping dysfunctions to individual brain areas has recently been challenged by a network perspective. It has been argued that aberrant functional connectivity (FC) of widespread brain regions seems more appropriate in explaining heterogeneous phenomena, such as psychiatric disorders.8–10

Investigation of functional connectivity usually involves identifying temporal correlations in blood-oxygen level–dependent (BOLD) signal between spatially distinct brain areas.11 This can be applied to fMRI data acquired during tasks or during rest. The latter method, referred to as resting-state functional connectivity (rsFC), is particularly suitable for clinical application, as it requires little compliance from the participant12 and scanning times are relatively short (5–10 min). During a typical resting-state scan, participants are asked to lie still with closed eyes or to fixate on a crosshair.

Seed-based analysis is a widely used approach that involves selecting a specific brain region and evaluating its correlation...
with time course in other voxels. Beside this, independent component analysis (ICA) is one of the most common techniques used to analyze rsFC data. In contrast to the seed-based approach, it does not require strict a priori hypotheses regarding regions of interest and allows the detection and separation of multiple resting-state networks (RSN) at once. Independent component analysis uses a mathematical algorithm to separate the fMRI data set into maximally spatially independent components. However, the identified components do exhibit temporal correlation, which offers another valuable perspective. The identified RSNs have been found to be highly reliable across individuals and cover functional properties (e.g., motor, salience detection) despite the absence of a task.

Previous studies investigating rsFC in patients with anorexia nervosa have identified abnormalities in several large-scale RSNs. Our own work, in which we used the ICA approach, provided evidence for increased rsFC between the angular gyrus and the frontoparietal network (FPN) in acutely ill patients that was also associated with self-reported persistence, a personality dimension strongly pronounced in this group and associated with cognitive control. This finding is in line with a neurobiological framework proposed by Kaye and colleagues suggesting that enhanced executive functions mediated by prefrontal brain areas play an important role in the etiology of anorexia nervosa. Furthermore we reported increased rsFC strength between the default mode network (DMN) and the anterior insula associated with self-reported problems of interoceptive awareness, a typical symptom of anorexia nervosa. However, our previous study investigating within-network rsFC in acutely ill patients with anorexia nervosa suggested a strong coupling between the DMN and the salience network manifested in an assignment of the anterior insula to the DMN in patients with the disorder. This coupling may indicate an altered interaction among the salience network, DMN and FPN in patients with anorexia nervosa, as proposed by the triple network model.

Another neglected aspect in previous studies focusing on rsFC in patients with anorexia nervosa is that abnormal FC can occur on different hierarchical scales. One approach to investigate FC on a superordinate scale is functional network connectivity (FNC). This approach identifies interconnections between large-scale networks and captures weaker, but existing temporal correlations between the components identified by ICA. This method elucidates abnormal functional integration on a macroscopic level, as described by the triple network model of psychopathology. This model suggests that the insula as a central hub of the salience network mediates the switch between the DMN as a network involved in internally oriented mental states to the FPN, which is responsible for externally oriented attention as a response to salient stimuli. Alterations in this process are thought to contribute to dysfunction underlying psychiatric disorders. To date, evidence for changes in FNC in patients with psychiatric disorders is sparse. Jafari and colleagues could show increased FNC among all dominant RSNs in patients with schizophrenia, while von dem Hagen and colleagues demonstrated reduced FNC in patients with autism-spectrum disorders. Yet, to our knowledge, no studies have investigated FNC in patients with anorexia nervosa. However, our previous study investigating within-network rsFC in acutely ill patients with anorexia nervosa suggested a strong coupling between the DMN and the salience network manifested in an assignment of the anterior insula to the DMN in patients with the disorder. This coupling may indicate an altered interaction among the salience network, DMN and FPN in patients with anorexia nervosa, as proposed by the triple network model.

The aims of the present study were 2-fold. First, we compared within-network rsFC between patients recovered from anorexia nervosa and healthy controls to test whether abnormal FPN and DMN rsFC, as found in acutely ill patients, is detectable after recovery and can thus be considered a trait marker of the disorder. To probe effects of recovery, we used a targeted follow-up approach regarding the brain regions that showed group differences in our previous study. We also included the salience network in our analysis, as the anterior insula, a brain region that is often assigned to the salience network, showed group differences in our previous study. Second, we sought to examine FNC among the 3 networks of the triple network model of psychopathology.

**Methods**

**Participants**

We recruited girls and women recovered from anorexia nervosa and pairwise, age- and sex-matched controls for participation in this study. To be considered recovered, they had to have a diagnosis of anorexia nervosa and to have maintained a body mass index (BMI) above 18.5 (if age > 18 yr) or a BMI above the tenth age percentile (if age < 18 yr) for at least 6 months before the study; have regular menses; and have not binged, purged, or engaged in substantial restrictive eating patterns. Control participants had to have a healthy weight, be eumenorrheic and have no history of psychiatric illness. Additional
exclusion criteria for each group are outlined in Appendix 1, available at jpn.ca. Most importantly, we excluded individuals who were taking psychotropic medication within 6 weeks before the study and who had a diagnosis of bulimia nervosa, substance abuse, neurologic, or medical conditions.

This study was approved by our local institutional review board, and all participants or the guardians of underage participants gave written informed consent.

Clinical measures

To ascertain the absence of a current diagnosis of eating disorders the expert form SIAB-EX31 was conducted with all participants. Eating disorder–specific psychopathology, including interoceptive awareness, was assessed using the short version of the Eating Disorders Inventory-2 (EDI-229). We assessed the persistence personality dimension using the German version of the Junior Temperament and Character Inventory (JTCI). Anxiety and depression were measured using the Symptom Check-List-90R (SCL-90-R), and IQ was measured with a short version of the German adaption of the Wechsler Intelligence Scale (WIE) or with a short version of the German adaption of the Wechsler Intelligence Scale for Children (HAWIK; Appendix 1).

Instrumental target reaction task

Participants performed a simple instrumental target reaction task as a measure of goal-directed behaviour to receive monetary rewards depending on their performance. Instrumental responding was defined by reaction time and by number of button presses in response to a target (Appendix 1).

Data acquisition

Data were acquired with a 3 T Siemens Trio scanner. The $T_1$-weighted structural brain scans were acquired using a rapid acquisition gradient echo (MP-RAGE) sequence with the parameters described in Appendix 1. An 8-min resting fMRI scan was acquired using a gradient-echo $T_2^*$-weighted echo planar imaging (EPI) sequence using standard parameters (Appendix 1). During fMRI, participants were instructed to lie still with their eyes closed and without falling asleep.

Image data preprocessing

Functional and structural images were processed using SPM8 (www.fil.ion.ucl.ac.uk/spm/) within the Nipype framework (http://nipy.sourceforge.net/nipype) following standard procedures (Appendix 1).

We evaluated the quality of the fMRI data by manual inspection and using artifact detection tools (ART).

Cortical thickness measurement

The $T_1$-weighted images were registered, motion-corrected, realigned, averaged and analyzed using the FreeSurfer software suite (http://surfer.nmr.mgh.harvard.edu) version 5.1.0. Cortical thickness was measured using standard FreeSurfer procedures (Appendix 1). We also extracted averaged thickness measurements for each participant from each of the respective labels of the Destrieux cortical atlas in each hemisphere according to the regions identified to show aberrant rsFC.

Independent component analysis

To identify temporally coherent RSNs, we conducted a spatial group ICA for all participants using the Group ICA fMRI Toolbox (GIFT) implemented in MATLAB (http://mialab.mrn.org/software/gift). The fMRI data were decomposed into maximally independent components according to the following steps. The number of components were estimated using modified minimum description length criteria, then we applied the infomax algorithm to the data. For each participant, we back-reconstructed component spatial maps using group independent component analysis (GICA) and converted them to z values.

Component selection

To identify components of interest for further analysis, we applied a systematic 2-step process by correlating components with white matter and cerebral spinal fluid templates as well as RSN templates created by Ye and colleagues. The evaluation was confirmed by investigating spectral metrics (Appendix 1).

Analyses of independent components

For group analyses, spatial maps of the back-reconstructed components representing the networks of interest for each participant were entered into SPM8. The $z$ values of these maps represent the concordance of the voxel-specific time-course to the averaged components time-course. The analysis of rsFC comprised 2 steps. First, we performed a 2-sample $t$-test for each preselected component, masked with the aforementioned RSN templates. Between-group differences had to exceed $p < 0.05$, family-wise error (FWE)–corrected, to guard against type I errors. Second, to follow up the previous finding of increased rsFC in acutely ill patients in comparison to healthy controls in the FPN and DMN, we examined the contrast “recovered patients > healthy controls” by applying a mask (Appendix 1) that corresponds to the results of our previous study in acutely ill patients (angular gyrus, anterior insula).

Additional analyses

To further assess the association of the magnitude of group differences in rsFC with psychometric parameters, we extracted the b values of the respective clusters at a threshold of $p < 0.001$, uncorrected, using MarsBar and computed the Pearson $r$ correlation using SPSS statistical software version 21.0 (SPSS) for each diagnostic group separately and for both
groups combined. Based on our previous work, we used a hypothesis-driven approach and planned to test the association between interoceptive awareness (EDI-2) and possible clusters showing group differences within the DMN as well as between persistence (JTCI) and the clusters of the FPN. To demonstrate external validity of the hypothesized function of the FPN, we investigated the association of instrumental responding and the magnitude of rsFC of the FPN. To address possible developmental effects, we tested for associations between age and rsFC using the same approach.

Additionally, we investigated the association between the cortical thickness measures and rsFC using partial correlation, with age as a covariate.

Functional network connectivity

The assessment of the FNC, defined by the temporal correlation between the spatially independent components, was conducted with the FNC toolbox (http://mialab.mrn.org/software/fnc). This toolbox computes the maximum lagged correlation, as described by Jafri and colleagues. First, the time courses of the respective components are filtered with cut-off frequencies below 0.017Hz and above 0.32Hz and interpolated to allow the detection of hemodynamic latencies below scanner repetition time. Next, the maximum lagged correlation is computed for all possible pairwise combinations of component time-courses according to the formula reported in Appendix 1. Correlation combinations that exceeded a significance level of $p < 0.05$, false discovery rate (FDR)-corrected, and their corresponding lag values were separately extracted for both groups. Differences in FNC between recovered patients and healthy controls were tested using a 2-sample $t$ test ($p < 0.05$, FDR-corrected).

Results

Participants

The study sample consisted of 62 female volunteers: 31 patients recovered from anorexia nervosa and 31 pairwise age- and sex-matched controls. There were no differences in age, body mass index standard deviation score (BMI SDS) and IQ between recovered patients and healthy controls. Recovered patients still had some residual eating disorder symptoms. The demographic and clinical characteristics of participants are described in Table 1.

Component identification

The dimension estimation revealed 23.76 ± 5.32 dimensions on average. The initial data reduction using principal components analysis (PCA) yielded 36 dimensions that were again reduced to 24 independent components using ICA. Six of these components were identified as RSNs of interest (Fig. 1).

Group comparison of independent components

We found reduced functional connectivity between the right dorsolateral prefrontal cortex (dlPFC) and component 5, associated with FPN ($t_{peak} = 4.87, p = 0.023$, FWE-corrected) in recovered patients compared with healthy controls (Fig. 2). We did not observe any further group differences (below the threshold of $p < 0.05$, FWE-corrected) in the remaining 5 components. Using our targeted follow-up approach, we found increased rsFC of the left angular gyrus with component 5 associated with the FPN ($t_{peak} = 3.16, p = 0.05$, FWE-corrected, in recovered patients > healthy controls; Appendix 1 and Fig. 3). No increased rsFC was found between the left angular gyrus and

Table 1: Demographic and clinical characteristics of study participants

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Group: mean ± SD</th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Recovered* $n = 31$</td>
<td>Control $n = 31$</td>
<td>$t$</td>
<td>$p$ value</td>
<td></td>
</tr>
<tr>
<td>Age, yr</td>
<td>22.27 ± 3.08</td>
<td>21.73 ± 2.99</td>
<td>−0.70</td>
<td>0.49</td>
<td></td>
</tr>
<tr>
<td>BMI†</td>
<td>20.69 ± 1.62</td>
<td>21.37 ± 2.10</td>
<td>1.42</td>
<td>0.16</td>
<td></td>
</tr>
<tr>
<td>BMI SDS</td>
<td>−0.54 ± 0.55</td>
<td>−0.29 ± 0.64</td>
<td>1.60</td>
<td>0.11</td>
<td></td>
</tr>
<tr>
<td>IQ‡</td>
<td>108.52 ± 9.74</td>
<td>110.67 ± 8.80</td>
<td>0.91</td>
<td>0.37</td>
<td></td>
</tr>
<tr>
<td>Age at onset, yr</td>
<td>14.38 ± 1.93</td>
<td>NA</td>
<td>—</td>
<td>—</td>
<td></td>
</tr>
<tr>
<td>Duration of recovery, mo</td>
<td>53.24 ± 33.27</td>
<td>NA</td>
<td>—</td>
<td>—</td>
<td></td>
</tr>
<tr>
<td>Illness duration, mo§</td>
<td>44.63 ± 31.93</td>
<td>NA</td>
<td>—</td>
<td>—</td>
<td></td>
</tr>
<tr>
<td>EDI–2 total score</td>
<td>162.05 ± 44.22</td>
<td>131.92 ± 25.89</td>
<td>−3.26</td>
<td>0.002</td>
<td></td>
</tr>
<tr>
<td>SCL–90–R, depression</td>
<td>6.93 ± 7.41</td>
<td>5.17 ± 8.15</td>
<td>−0.88</td>
<td>0.38</td>
<td></td>
</tr>
<tr>
<td>SCL–90–R, anxiety</td>
<td>3.97 ± 4.50</td>
<td>2.9 ± 6.32</td>
<td>−0.75</td>
<td>0.45</td>
<td></td>
</tr>
</tbody>
</table>

BMI = body mass index; BMI SDS = body mass index standard deviation score; EDI-2 = Eating Disorders Inventory-2; NA = not applicable; SCL-90–R = Symptom Check-List-90R; SD = standard deviation.

*One participant in the sample had been recovered for 9 mo, all other participants were recovered for at least 2 mo.
†BMI is displayed but statistical comparisons are based on BMI-SDS values to ensure comparability across age.
‡IQ was assessed with a short version of the German adaption of the Wechsler Adult Intelligence Scale or a short version of the German adaption of the Wechsler Intelligence Scale for Children for participants aged 15 yr or younger.
§Reported illness duration comprises the total time period from onset of the first episode until recovery following the last episode.
component 8. Regarding the DMN, no increased rsFC between the anterior insula and component 10 or 13 was observed.

Additional analyses

Since acutely ill patients are believed to exert excessive cognitive control that may be reflected in the persistence personality dimension, we tested this association in the FPN clusters, as described in our previous study.18 We found no association between persistence and reduced rsFC in the right dIPFC or increased rsFC in the angular gyrus in either of the groups or in both groups combined. To validate the function of the FPN, we tested the association between rsFC of the FPN and performance on an instrumental responding task. There was an association between the instrumental responding parameters, number of button presses and reaction time and increased rsFC between the angular gyrus and the rest of the FPN (component 5) in both groups combined (reaction time was also associated with rsFC in the angular gyrus within recovered patients; Table 2).

Age was not associated with any finding of aberrant rsFC reported above. The magnitude of rsFC in brain regions showing group differences for FPN connectivity (dIPFC, angular gyrus) was neither associated with cortical thickness of the right dIPFC (middle frontal gyrus, Destrieux atlas) nor the left angular gyrus (superior parietal gyrus, Destrieux atlas). For means with standard deviations and statistics of group differences of persistence, instrumental responding parameters and cortical thickness measures, see Appendix 1.

Functional network connectivity

As expected, components identified to cover the same network were found to correlate significantly in both groups (all p < 0.048, FDR-corrected). In healthy controls we found significant FNC among the components of the DMN, the FPN and salience network. In recovered patients, only components of the FPN and the salience network significantly correlated; we found no correlation with the components of the DMN (Fig. 4). However, the group comparison of FNC revealed no difference between recovered patients and healthy controls (all p > 0.08, FDR-corrected).

Discussion

The aim of this study was to investigate within- and between-network rsFC (or FNC) in patients recovered from anorexia nervosa compared with healthy controls. We found differences in within-network rsFC in the FPN. Specifically, our exploratory approach revealed reduced rsFC between the dIPFC and the FPN in recovered patients, while the targeted follow-up approach is in line with our previous finding (in acutely ill patients) of increased rsFC between the angular gyrus and the remaining parts of the FPN. Aberrant coupling between the anterior insula and the DMN, as found in acutely ill patients, was not detected. Regarding FNC, we found no group differences in interactions among the DMN, FPN and salience network.

Studying within- and between-network rsFC in patients recovered from anorexia nervosa enabled us to probe whether rsFC characteristics are trait or state markers of the disease. In line with the results of McFadden and colleagues,24 we found that aberrant rsFC in the DMN seems to normalize with recovery and therefore constitutes a state marker of the disease. In contrast, Cowdrey and colleagues22 reported increased rsFC in the DMN for recovered patients, but no difference in the FPN. Reasons for these discrepant findings may include the fact that ICA was performed independently in each group, which

---

Fig. 1: Spatial maps of 6 independent components (IC) of interest grouped by network: frontoparietal network (FPN), default mode network (DMN) and salience network. Spatial maps are plotted as t statistics thresholded at p < 0.05, family-wise error–corrected.
results in group-specific components, as well as heterogeneity in the population of patients with anorexia nervosa.

Likewise, task-based studies in patients with anorexia nervosa also indicate that some neural signatures persist into recovery, although discrepancies exist. Some studies investigating food cue reactivity in acutely ill and recovered patients found that increased neural activity in regions involved in the food motivation brain circuit (e.g., amygdala, hypothalamus, orbitofrontal cortex [OFC], insula) reflect trait vulnerability of the illness, while other studies found reduced neural activity in the insula in recovered patients. We in turn found no altered rsFC in the anterior insula (part of the DMN) or salience network (FPN; peak coordinates in Montreal Neurological Institute space: x, y, z = 42, 46, 14).

**Fig. 2:** Group difference at a threshold of \( p < 0.05, \) family-wise error-corrected, between patients recovered from anorexia nervosa (recAN) and healthy controls (HC) in component 5, representing the frontoparietal network (FPN; peak coordinates in Montreal Neurological Institute space: x, y, z = 42, 46, 14).

**Fig. 3:** Targeted follow-up approach: group difference at a threshold of \( p = 0.05, \) family-wise error-corrected, between patients recovered from anorexia nervosa (recAN) and healthy controls (HC) in component 5, representing the frontoparietal network (FPN; peak coordinates in Montreal Neurological Institute space: x, y, z = 36, –64, 44). Applied mask covering the previous finding at angular gyrus (AG) is displayed in white.
Table 2: Correlations between the magnitude of the cluster-specific rsFC and psychometric parameters, cortical thickness, age, duration of illness, anxiety and depression for both groups and each group separately

<table>
<thead>
<tr>
<th>Analysis</th>
<th>rsFC at dIPFC</th>
<th>rsFC at AG</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Recovered</td>
<td>Control</td>
</tr>
<tr>
<td>Confirmatory analyses</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Persistence</td>
<td>r = 0.35</td>
<td>r = –0.03</td>
</tr>
<tr>
<td></td>
<td>p = 0.06</td>
<td>p = 0.86</td>
</tr>
<tr>
<td>Instrumental responding: #bp</td>
<td>r = 0.41</td>
<td>r = 0.04</td>
</tr>
<tr>
<td></td>
<td>p = 0.03</td>
<td>p = 0.81</td>
</tr>
<tr>
<td>Instrumental responding: RT</td>
<td>r = –0.27</td>
<td>r = –0.17</td>
</tr>
<tr>
<td></td>
<td>p = 0.14</td>
<td>p = 0.37</td>
</tr>
<tr>
<td>Exploratory analyses</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age</td>
<td>r = 0.05</td>
<td>r = –0.12</td>
</tr>
<tr>
<td></td>
<td>p = 0.78</td>
<td>p = 0.54</td>
</tr>
<tr>
<td>Depression</td>
<td>r = 0.09</td>
<td>r = –0.10</td>
</tr>
<tr>
<td></td>
<td>p = 0.63</td>
<td>p = 0.80</td>
</tr>
<tr>
<td>Anxiety</td>
<td>r = 0.03</td>
<td>r = –0.05</td>
</tr>
<tr>
<td></td>
<td>p = 0.89</td>
<td>p = 0.80</td>
</tr>
<tr>
<td>Illness duration</td>
<td>r = –0.13</td>
<td>—</td>
</tr>
<tr>
<td></td>
<td>p = 0.55</td>
<td>—</td>
</tr>
<tr>
<td>Cortical thickness*</td>
<td>r = 0.04</td>
<td>r = 0.11</td>
</tr>
<tr>
<td></td>
<td>p = 0.82</td>
<td>p = 0.56</td>
</tr>
</tbody>
</table>

AG = angular gyrus; dIPFC = dorsolateral prefrontal cortex; rsFC = resting-state functional connectivity; RT = reaction time.
*A partial correlation with age as covariate was calculated for associations between rsFC and cortical thickness.
†Significant when adjusting for multiple testing using the Bonferroni method.

Fig. 4: Functional network connectivity in (A) patients recovered from anorexia nervosa (recAN) and (B) healthy controls (HC). Arrows show the significant correlation between resting-state networks. The direction of the arrow corresponds with the direction of the delay between 2 components. DMN = default mode network; FPN = frontoparietal network; IC = independent component; SN = salience network.
nervosa as a disorder characterized by altered cognitive control processes underpinned by dysconnectivity within the FPN, which seems to persist into recovery, supporting the recently highlighted role of the FPN in psychiatric disorders. Nonetheless, conclusions from rsFC data with respect to task-related interpretations remain somewhat speculative. Alterations observed in the FPN could be interpreted as heightened cognitive control in patients with anorexia nervosa, as inefficient cognitive control mechanisms, or even as independent from any control processes. However, previous work has been able to demonstrate a direct link between rsFC and task-related neural activity.

Regarding the interaction among the RSNs, we found no alterations among the DMN, FPN and salience network in patients recovered from anorexia nervosa. Based on these findings, we suggest that altered integration of information across these networks, as suggested by the triple network theory, may not be considered a trait marker of the disorder. Given that the most important evidence of abnormal FNC originates from research in patients with schizophrenia, one might speculate whether only a few very marked psychiatric symptoms, such as hallucinations, are associated with changes in rsFC between networks.

Limitations

Our study has to be evaluated in the light of the following limitations. First, owing to age differences we compared results obtained in acutely ill and recovered patients only across studies, but we did not include them into a single statistical model. Second, studying recovered patients allowed us to exclude the effects of acute undernutrition, but the cross-sectional nature of the study does not allow us to determine whether the observed differences in rsFC constitute a true trait marker or a possible scar effect of the disease. It is also possible that patients who eventually recover belong to a different subgroup than chronic or relapsing-remitting patients. Future research applying a longitudinal study design would help to clarify whether those neural substrates are a consequence or a potential antecedent of pathologic eating behaviour.

Strengths of our study include the large, homogeneous sample consisting of young unmedicated patients recovered from anorexia nervosa, who are consistently of the restrictive subtype, as well as the fact that all participants were scanned in the morning after an overnight fast.

Conclusion

Our study demonstrates that some abnormal rsFC patterns found in patients with anorexia nervosa persist even after long-term weight restoration. While abnormal rsFC within the DMN seems to normalize, the increased rsFC between the angular gyrus and the FPN, which might indicate alterations in cognitive control functions, are still present after recovery, and abnormalities regarding rsFC between the dIPFC and the FPN re-emerge. Moreover, we found no altered FNC between networks in patients recovered from anorexia nervosa. Studies evaluating the predictive potential of differences in rsFC for treatment outcome may help to pave the way for potential clinical applications of resting-state fMRI.

Acknowledgments: This work was supported by Deutsche Forschungsgemeinschaft (367/5-1 & SFB 940) and Swiss Anorexia Nervosa Foundation grants to S. Ehrlich. The authors thank all members of the research team for their assistance and thank all participants for their cooperation.

Affiliations: From the Eating Disorder Services and Research Center, Department of Child and Adolescent Psychiatry, Faculty of Medicine, Technische Universität Dresden, Dresden, Germany (Boehm, Geisler, Tam, King, Ritschel, Seidel, Bernardoni, Roessner, Ehrlich); the MGH/MIT/HMS Martinos Center for Biomedical Imaging, Massachusetts General Hospital, Charlestown, MA (Ehrlich); the Harvard Medical School, Department of Psychiatry, Massachusetts General Hospital, Boston, MA (Ehrlich); the Department of Psychotherapy and Psychosomatic Medicine, Faculty of Medicine, Technische Universität Dresden, Dresden, Germany (Murr); the Department of Psychology, Technische Universität Dresden, Dresden, Germany (Goschke); the The Mind Research Network, Albuquerque, New Mexico (Calhoun); and the Department of ECE, the University of New Mexico, Albuquerque, New Mexico (Calhoun).

Competing interests: V. Roessner has received payment for consulting and writing activities from Lilly, Novartis and Shire Pharmaceuticals; lecture honoraria from Lilly, Novartis, Shire Pharmaceuticals and Medice Pharma; and support for research from Shire and Novartis. He has carried out (and is currently carrying out) clinical trials in cooperation with Novartis, Shire and Otsuka. No other competing interests declared.

Contributors: S. Ehrlich designed the study. I. Boehm, D. Geisler, F. Tam, J. King, F. Ritschel. M. Seidel, F. Bernardoni and J. Murr acquired the data, which I. Boehm, D. Geisler, T. Goschke, V. Calhoun, V. Roessner and S. Ehrlich analyzed. I. Boehm, J. King and S. Ehrlich wrote the article, which all authors reviewed and approved for publication.

References

Partially restored rsFC in women recovered from anorexia nervosa


34. Li YO, Adali T, Calhoun VD. Estimating the number of independent components for functional magnetic resonance imaging data. *Hum Brain Mapp* 2007;28:1251-66.


