Brain Network Alterations in Rectal Cancer Survivors With Depression Tendency: Evaluation With Multimodal Magnetic Resonance Imaging

Wenwen Zhang1†, Ying Zou2†, Feng Zhao3, Yongqing Yang4, Ning Mao5,6, Yuan Li4*, Gang Huang7*, Zhijun Yao7* and Bin Hu7,8,9,10*

Surgery and chemotherapy may increase depression tendency in patients with rectal cancer (RC). Nevertheless, few comprehensive studies are conducted on alterations of brain network induced by depression tendency in patients with RC. Resting-state functional magnetic resonance imaging (rs-fMRI) and diffusion tensor imaging (DTI) data were collected from 42 patients with RC with surgery and chemotherapy and 38 healthy controls (HCs). Functional network (FN) was constructed from extracting average time courses in brain regions, and structural network (SN) was established by deterministic tractography. Graph theoretical analysis was used to calculate network properties. Networks resilient of two networks were assessed. Clinical correlation analysis was explored between altered network parameters and Hamilton depression scale (HAMD) score. This study revealed impaired FN and SN at both local and global levels and changed nodal efficiency and abnormal small-worldness property in patients with RC. On the whole, all FNs are more robust than SN. Moreover, compared with HC, patients with RC show less robustness in both networks. Regions with decreased nodal efficiency were associated with HAMD score. These cognitive dysfunctions are mainly attributable to depression-related brain functional and structural network alterations. Brain network reorganization is to prevent patients with RC from more serious depression after surgery and chemotherapy.

Keywords: rectal cancer, depression tendency, functional network, structural network, multimodal research
INTRODUCTION

Rectal cancer (RC) is a disease characterized by a high mortality rate. Patients with RC usually suffer from tremendous psychological stress, which leads to a series of psychological diseases (1, 2). Some contemporary studies showed that various discomfort symptoms caused by chemotherapy could induce cognitive impairment in patients with cancer, including impaired attention, memory, and executive function (3, 4). The long-term emotional distress of the patient increased the risk of depression (5). Depression-related factors may contribute to the less optimal network topology in the functional network (FN) (6, 7) and structural network (SN) (8) of patients with cancer. Neuroimaging studies reported that a considerable number of patients with cancer with surgery and chemotherapy had morphological variation and functional abnormalities in brain, such as decreased hippocampal volume (9), lower white matter volume (10), memory difficulties (1), and cognitive deficit (11). Therefore, research on FN and SN in this study could bring new insights into the neurophysiological mechanisms in patients with RC, and through the analysis of brain images, the emotional distress of patients with RC could be diagnosed and treated, thereby improving the quality of life of patients (12).

Combining multimodal data can reveal hidden relationships among different data, unifying different findings in brain imaging (13). Therefore, multimodal imaging is a prominent method in cognitive neuroscience research. Graph-based functional and structural brain connectivity analysis is a new method, which provides evidence for the complexity of the brain by modeling the interactions between different brain regions (14, 15). Previous studies have consistently shown that the brain functional network is organized in a small-worldness property, with local specialization and high global information transmission capabilities (16, 17). Bruno et al. (1) reported significantly reduced shortest path length and small-worldness property in the breast cancer group. Several findings point that the functional network of patients with cancer loses its ability to support various cognitive functions following chemotherapy (18, 19). Observational studies found that alterations in brain structural network had an adverse impact on the cognition of cancer survivors (20, 21). Although there are many studies on brain cognitive impairment in patients with cancer, less is known about FN and SN abnormalities in patients with RC. We used multimodal neuroimaging to investigate the alterations in FCs and SCs for gaining insight into the brain cognitive dysfunction in patients with RC.

In view of the poor understanding of psychological disorders and cognitive impairment of RC survivors in existing studies, it is essential to study the depression tendency and related factors in patients with RC. Therefore, this study investigated the abnormalities in FN and SN using graph theory analysis in patients with RC with surgery and chemotherapy characterized by depression tendency compared with healthy controls (HCs). We hypothesized that patients with RC would show altered small-worldness property and topological architecture in the FN and SN due to the effects of depression tendency. We sought to expand our understanding of the resilience of the brain network in patients with RC. The study also explored the potential association between the significant alterations in network properties of patients with RC and severity of depression symptoms.

METHODS

Participants

A total of 42 patients with RC with surgery and chemotherapy were recruited from the Gansu Provincial Hospital, whereas the 38 age- and gender-matched healthy control participants were recruited through newspaper advertisements. They were recruited from July 2017 to June 2020. All participants were diagnosed according to DSM-IV criteria by two experienced psychiatrists. They have executed the evaluation of 17-item Hamilton Rating Scale for Depression (HAM-D-17). The evaluation results showed that all the patients with RC included in the experiment had depression tendency. All participants were given written informed consent when image scanning. None of the subjects took any psychotropic drugs.

Data Acquisition

Magnetic resonance imaging data were acquired using a 3.0 T Siemens Trio scanner (Siemens Erlangen, Germany). Subjects were asked to relax with eyes closed and not to think about anything. The structural image was acquired with a T1-weighted spin-echo sequence: TR/TE = 2530/2.98 ms, slice thickness = 1 mm, slice gap = 0.8 mm, FOV = 256*256 mm, The resting-state functional images (rs-fMRI) were obtained with the following parameters: TR/TE = 2,000/30 ms, 64*64 matrix, FOV = 224*224 mm, total 240 volumes, 32 sequential ascending axial slices of 3.5 mm thickness. Diffusion tensor imaging (DTI) data were acquired using a single-shot echo-planar imaging-based sequence with the following parameters: TR = 11,600 , TE = 85 , FOV = 256 mm * 256 mm, acquisition matrix = 112*112, axial slices = 32, 64 diffusion directions with b = 1,000 s/mm², and an additional image without b = 0 s/mm².

Data Processing

We used the SPM8 (Statistical Parametric Mapping: http://www.fil.ion.ucl.ac.uk/spm) and DPARSFA (Data Processing Assistant Resting-State: http://www.restfmri.net) on the preprocessing of all the rs-fMRI data (22). The specific preprocessed steps were as follows: (1) the first 10 volumes of the fMRI were removed; (2) we performed the slice timing, head movement correction, and rearrangement; (3) all subjects were excluded if their head movement which was >2.0 mm maximum displacement in any of the x, y, or z directions was >2° (23); (4) all the rs-fMRI data were spatially normalized to the Montreal Neurological Institute (MNI) space using structural image normalization parameters; (5) the smoothing Gaussian kernel of full width at half maximum (FWHM) was 8 mm (24); (6) the 24 head motion parameters, averaged global, white matter signals, and cerebrospinal fluid were processed by nuisance covariates regression (25); (7) removing linear trends; and (8) temporal band-pass filtering (0.01–0.08 Hz) (26).
The DTI data were preprocessed using PANDA (http://www.nitrc.org/projects/panda) (27) in MATLAB2014a. The specific preprocessed steps were as follows: in MATLAB2014a. (1) converting DICOM files into NIfTI images; (2) estimating the brain mask; (3) cropped the image; (4) correction of the eddy current effect and head motions; (5) averaging multiple acquisitions; and (7) FA metrics calculation.

Construction of Brain Networks
The Human Brainnetome Atlas (246 Atlas) were used to demarcate the nodes and edges of the brain network (28). In each subject, 246 atlas were used to construct brain networks for further graph theory analysis.

The GRETNA (a toolbox for analyzing brain network, www.nitrc.org/projects/gretna/) were used to construct the functional brain network. For each subject, Pearson correlation coefficient was used to construct the correlation matrix which the mean time series of each region was represented each node. Fisher Z transform was applied to each matrix to convert the data into Z-scores. A FA-weighted symmetric matrix was constructed for each participant by deterministic tractography as the following analysis basis. Each matrix represented the white matter network of the cerebral cortex, and each row or column in network represented the brain region of 246 atlas.

Threshold Calculation
To construct an undirected binary network and make the generated graph metrics stable, it is necessary to be thresholded for the weight of the brain networks. There is no fixed method to determine the threshold in current research. Therefore, in FN, we used sparsity (26–50%) with a step of 1% (29) to divide the network threshold. Then, we calculated the topological properties of FN in a series of threshold range. In SN, we use FA (0.20-0.42) with a step of 0.2 as the threshold of the network according to the previous study (30). The small-worldness property is related to the threshold of the network (31), so we need to calculate a network threshold to get effective network properties.

Whole Brain Network Organization
Graph theoretical analyses of the FN and SN in patients with RC and HC were calculated with routines from the GRETNA toolbox. The network topological properties at the global levels were calculated, including (1) properties that suggest network segregation of brain, such as the normalized clustering coefficient (γ), the local efficiency $E_{loc}(G)$; (2) properties that indicate network integration of the brain, such as the normalized shortest path length(λ), the global efficiency $E_{glob}(G)$; (3) small-worldness (σ) property which evaluates the balance of segregation and integration.

The nodal efficiency ($E_{nod}$) measures the ability of a particular node to propagate information with all other nodes in the network. It is considered as the inverse of the harmonic mean of the minimum path length between an index node and all other nodes in the network.

Network Resilience Analysis
Network resilience refers to the ability to withstand perturbations or failures in the network, which is usually related to the stability of complex networks (32, 33). In FN and SN, we used random or targeted attacks with fixed sparsity or FA values to evaluate the network resilience, so as to ensure that all anatomical regions were involved in the network, thus minimizing the number of false-positive paths (32). In targeted attack analysis, the betweenness value of each node in the network was calculated and sorted in descending order. We deleted the nodes in the network in order of betweenness value and calculated the global efficiency of each network after attack (34). In random attack analysis, we deleted the nodes of network randomly and calculated the global efficiency of each network after attack.

Statistical Analysis
The demographic and clinical characteristics of the patients with RC and HC were analyzed by chi-square test and two-sample $t$-tests using SPSS 21. We performed statistical comparisons of topological measures between the two groups using non-parametric permutation tests with 5,000 iterations for each sparsity and FA value (35). For the $E_{nod}$, the non-parametric permutation tests was repeated at the sparsity $= 26\%$ and the FA value $= 0.42$. FDR correction was conducted for all these results. Besides, we used Pearson correlation analyses to explore the correlations in patients with RC between nodes with significant difference in $E_{nod}$ and the severity of depression (HAMD score).

RESULTS
Demographic Characteristics
Demographic information is summarized in Table 1. There is no significant difference in age ($p = 0.564$) and gender ($\chi^2 = 1.312$, $p = 0.765$) between the two groups. There was a significant difference in HAMD score between the two groups ($p < 0.001$).

The Alterations of FN and SN Properties
In the functional network, patients with RC have a higher shortest path length (λ) than HC (Figure 1B, sparsity $= 26\%$). Since there is no significant difference in the clustering coefficient (γ) between the two groups under the same sparsity (Figure 1A), this leads to the abnormal small-worldness(σ) (Figure 1C) of patients with RC. The local efficiency has increased in patients with RC (Figure 1D, sparsity $= 26\%$), and there is no significant change in global efficiency (Figure 1E).

| Variables (Mean ± SD) | Patients with RC ($n = 42$) | Healthy controls ($n = 38$) | $p$-value |
|-----------------------|-----------------------------|-----------------------------|-----------|
| Gender (M:F)          | 23:19                       | 22:16                       | 0.564#    |
| Age (years)           | 50.89 ± 7.50                | 48.96 ± 7.93                | 0.756*    |
| HAMD                  | 9.94 ± 4.93                 | --                          | --        |

SD, standard deviation; HAMD, Hamilton depression rating scale. # and * indicate p-value for chi-square test and two-sample $t$-test, respectively.

Table 1  | Demographic and clinical characteristics of subjects.
Zhang et al. Rectal Cancer, Depression Tendency, Multimodal Connectivity, Network Properties

FIGURE 1 | Functional connectivity network at different sparsity for patients with RC (the blue line) and controls (the pink line) and their statistical comparison results ($p < 0.05$ 5,000 permutation test, FDR correction). (A) Gamma, (B) lambda, (C) sigma, (D) local efficiency, and (E) global efficiency. The black triangles indicate a significant group difference.

FIGURE 2 | Structural connectivity network at different FA threshold for RC (the blue line) and controls (the pink line) and their statistical comparison results ($p < 0.05$ 5,000 permutation test, FDR correction). (A) Gamma, (B) lambda, (C) sigma, (D) local efficiency, and (E) global efficiency. The black triangles indicate a significant group difference.
TABLE 2 | Brain regions with significant group effect in the nodal efficiency between patients with RC and HC for FN.

| Regions      | Patients with RC | Control | p-value |
|--------------|------------------|---------|---------|
| RC < HC      |                  |         |         |
| vmPu.R       | 0.5580 ± 0.0582  | 0.6190 ± 0.0578 | 0.0002  |
| dCa.L        | 0.5116 ± 0.0694  | 0.5816 ± 0.0592 | 0.0002  |
| GPr          | 0.5575 ± 0.0732  | 0.6195 ± 0.0526 | 0.0002  |
| vmPu.L       | 0.5765 ± 0.0490  | 0.6264 ± 0.0542 | 0.0002  |
| TL.R         | 0.5874 ± 0.0466  | 0.5941 ± 0.0533 | 0.0008  |
| PPHtha.R     | 0.5245 ± 0.0737  | 0.5846 ± 0.0519 | 0.0004  |
| GPL          | 0.5581 ± 0.0661  | 0.6107 ± 0.0620 | 0.0006  |
| lPFtha.R     | 0.5645 ± 0.0698  | 0.6148 ± 0.0589 | 0.0032  |
| vCa.R        | 0.5223 ± 0.0507  | 0.5645 ± 0.0514 | 0.0044  |
| A7m.R        | 0.6101 ± 0.0540  | 0.6404 ± 0.0666 | 0.0060  |
| Oth.R        | 0.5318 ± 0.0491  | 0.5833 ± 0.0610 | 0.0052  |
| mPMththa.R   | 0.5719 ± 0.0416  | 0.6158 ± 0.0542 | 0.0060  |
| msOccG.R     | 0.6029 ± 0.0705  | 0.6429 ± 0.0569 | 0.0098  |
| dCa.R        | 0.5286 ± 0.0650  | 0.5701 ± 0.0647 | 0.0090  |
| mPPtha.R     | 0.5545 ± 0.0615  | 0.6007 ± 0.0778 | 0.0090  |
| RC > HC      |                  |         |         |
| A23v.L       | 0.6719 ± 0.0485  | 0.6601 ± 0.0612 | 0.0018  |
| A40d.L       | 0.6862 ± 0.0527  | 0.6738 ± 0.0686 | 0.0042  |
| A40v.L       | 0.7195 ± 0.0408  | 0.6875 ± 0.0388 | 0.0048  |
| A23v.R       | 0.6742 ± 0.0394  | 0.6260 ± 0.0479 | 0.0064  |
| A4t.R        | 0.7038 ± 0.0857  | 0.6709 ± 0.0631 | 0.0048  |

vmPu, ventromedial putamen; dCa, dorsal caudate; GP, globus pallidus; TL, area TL (lateral PPHC, posterior parahippocampal gyrus); PPHtha, posterior parietal thalamus; lPFtha, lateral prefrontal thalamus; vCa, ventral caudate; A7m, medial area 7(PEp); Othha, occipital thalamus; mPMththa, pre-motor thalamus; msOccG, medial superior occipital gyrus; mPPtha, medial prefrontal thalamus; L, left; R, right.

In the structural network, the clustering coefficient (γ) (Figure 2A, FA = 0.32) and small-worldness(σ) (Figure 2C, FA = 0.32, 0.34, and 0.38) of patients with RC is larger than that of HC, the shortest path length (λ) (Figure 1B) being unchanged. In addition, compared with HC, the global efficiency of patients with RC has increased (Figure 2E, FA = 0.28–0.32).

Regional Efficiency Comparison
In FN, patients with RC showed that significantly decreased nodal efficiency. There were several regions including bilateral basal ganglia, right parahippocampal gyrus, bilateral thalamus, right precuneus, and right lateral occipital cortex. Meanwhile, the increased nodal efficiency was mainly in frontal lobe (orbital gyrus), basal ganglia, left inferior frontal gyrus, left amygdala, bilateral cingulate gyrus, left inferior parietal lobule, and right precentral gyrus in FN and SN for patients with RC (p < 0.05, after 5,000 permutation test, FDR test) (Tables 2, 3; Figure 3).

The Analysis of Network Resilience
With the targeted and random attack, a significantly decreased decline of the global efficiency was found in FN and SN (Figure 4). In both networks, the global efficiency of patients with RC decreased faster over a wide percentage of removal, which reflected that the networks of patients with RC were more fragile.

In all subjects, the resilience of structural network is weaker than that of functional network under the same threshold.

Correlations Between Network Properties and HAMD Scores for RC
In the analysis, there were correlations between the HAMD score and the nodes with significant $E_{\text{nod}}$ in the FN. For FN, mPMththa.R ($r = 0.389, p = 0.023$) (Figure 5B), and for SN, LAmyg.L ($r = 0.440, p = 0.01$) (Figure 5A).

DISCUSSION
In this study, different topological organizations of FN and SN in patients with RC and HC were explored. The findings pointed patients with RC displayed altered small-worldness property and global topological organization compared with HC. Moreover, there were regions with significant abnormal $E_{\text{nod}}$ being mainly distributed in frontal region, subcortical regions, and central region in patients with RC. In addition, patients with RC showed vulnerable network resilience in both networks, and FN would be more stable than SN across participants.

Network Properties
Many studies have used fMRI (36, 37) and DTI (21) images to explore the global and regional brain network properties of patients with breast cancer and lung cancer, but there are still few studies on rectal cancer. Compared with HC, the functional networks of patients with RC displayed a higher shortest path length (λ) and decreased small-worldness (σ), reflecting reduced global integration and disrupted organization balance (1, 21). Our results also revealed increased local efficiency in patients with RC. It is a measure of local information transmission among adjacent nodes and therefore an indication of network segregation (38, 39). Previous studies demonstrated reduced local efficiency, a common measure of the brain network’s response to computational attack, associated with patients with breast cancer (21, 40). Due to brain structural damage, decreased local efficiency would affect the fault tolerant ability of brain network. More detail, the result of weakening network fault tolerance is that if a node in the brain is damaged, the connection between previously linked nodes would be greatly affected (41). Therefore, reduced local efficiency is a risk factor for patients with RC.
Recently, researchers use graph theory to analyze complex brain functional networks after chemotherapy. It has been proved that chemotherapy-related cognitive deficits were associated with abnormal topological alterations of brain functional and structural network (42, 43). In this study, increased shortest path length and decreased local efficiency in patients with RC with surgery and chemotherapy could be seen as a brain compensation mechanism, which included changing the global pathway and adjusting regional activity to preserve a seesaw-like balance of the brain network.

Patients with RC showed increased clustering coefficients ($\gamma$), small-worldness($\sigma$), and global efficiency in SN (Figure 2). Abnormal small-worldness property of SN indicated that the local specialization and global integration of brain in patients with RC were disrupted, where the SN tended to be more randomized (44). Global efficiency is the inverse of the average shortest path between nodes. When nodes could interact directly, the efficiency is high (21). Therefore, global efficiency is an indicator of network function integration and parallel information processing capability (41). The present results of abnormal network properties reflected the undesired topological organization in SN, which exhibited that the deficits of emotional and cognitive processing in patients with RC might result from network damages. Besides, the increased network properties of SN in patients with RC might suggest that local nerve fibers reconstructed in response to the abnormalities in brain functional network. The compensatory response of the SN is activated for maintaining brain functional integrity to compensate the cognitive impairment caused by chemotherapy to patients with RC (24). Aforementioned evidence illuminated that cognitive deficit related to patients with RC may act via disrupted coordination between global and regional networks.

**Nodal Efficiency of Networks**

To explore the functional and structural characteristics of the human brain more accurately and quantitatively, our study employed a new standard brain atlas, containing 246 brain regions. This atlas would allow brain network analysis to use predefined nodes in an informed manner (45). Therefore, more detailed division of brain regions provides better help in multimodal data analysis. We observed decreased $E_{\text{nod}}$ only in FN of patients with RC. The significantly changed regions were located in bilateral basal ganglia, bilateral thalamus, right parahippocampal gyrus, right precuneus, and right lateral occipital cortex. The basal ganglia is not only related to motor control, but also related to the cognitive and limbic functions (46). Moreover, basal ganglia is the collection of subcortical nuclei surrounding the thalamus (46). Abnormal activation of basal ganglia/thalamus was found in the depressive studies (47), suggesting that abnormalities in these brain regions may lead to abnormal emotional processing mechanisms. Prior studies reported that parahippocampal gyrus and precuneus were associated with memory function, so alterations in these regions might affect memory decline (48, 49). Task fMRI study of memory factors found that the occipital cortex of patients with cancer was more significantly correlated with vigor and fatigue scores (50). Frequent fatigue is a common symptom of patients with cancer (51). Aforementioned evidence indicated that the decreased $E_{\text{nod}}$ of FN in this study represented alterations in regional characteristics of the brain network, which further affected the cognitive impairment of patients with RC.

Furthermore, the increased nodal efficiency was mainly in frontal cortex, left amygdala, bilateral cingulate gyrus, left inferior parietal lobule, and right precentral gyrus in FN and SN for patients with RC. In experiments with high-demand condition, the right inferior frontal gyrus and other components of the two hemisphere working memory circuitry in patients with cancer were found greater activation than the control group in a prior study (52). These abnormalities might be a compensation mechanism to preserve normal thinking and responsiveness in patients with cancer. In addition, chemotherapy affects estrogen levels in patients with cancer. Estrogen levels are thought
to have neuroprotective effects in the brain, thus helping to maintain cognitive function (53). Therefore, female patients with cancer are more likely to develop cognitive impairment in brain regions related to learning and memory after chemotherapy, such as hippocampus and amygdala (54). The anterior cingulate cortex is involved in attention control, response selection, and error monitoring (55). Abnormal brain activity patterns in the attention-controlled regions, including the anterior cingulate gyrus, are related to anxiety (56). The emotional fluctuation caused by excessive psychological stress in patients with RC could induce abnormal activation of cingulate gyrus. Saykin et al. (57) revealed that the activation of frontal and parietal lobes increased during the speech working memory task 1 month after chemotherapy. Compared with controls, the cancer group showed significantly greater activation in right precentral gyrus and right cingulate gyrus (19). Moreover, in the SN, the nodal efficiency was only increased. We speculated that after surgery and chemotherapy, the node efficiency of SN showed more obvious activation to maintain the robustness of overall network at the expense of other network property, such as integration. These results improved the understanding of chemotherapy-induced cognitive impairment in patients with RC from the perspective of brain node efficiency.

As shown in Figure 5, the patients with RC showed a positive relationship between HAMD and decreased nodal efficiency in mPMth a.R of FN, as well as a positive relationship between HAMD and increased nodal efficiency in LAmyg.L of SN. The correlation between the changed node efficiency and HAMD score may indicate impaired cognitive control combined with abnormal affective processing in patients with RC (29). A prior study suggested that regions sensitive to negative emotions were hyperactive in processing negative information (36), and it was not surprising to find a significant positive correlation between increased nodal efficiency and HAMD in the amygdala. Moreover, the positive relationship between HAMD and decreased nodal efficiency revealed that abnormal activation...
of FN in patients with RC might cause cognitive impairments and depressed mood (58). Therefore, we speculated that alterations in brain network properties assist us to study the depression tendency in patients with RC after chemotherapy and surgery.

**Comparison of Network Resilience**

In both networks, a key finding of significantly decreased resilience to targeted and random attack was found (Figure 4). Being more effective than other network properties to measure network integration performance, global efficiency of the FN and SN was utilized to explore network resilience quantitatively (34). In this study, both networks of patients with RC were more vulnerable and SN is less resilient than FN, which were consistent with our previous research results (59). This finding enhanced the conclusion that lower brain resilience was associated with progressive deterioration of cognitive impairment in breast cancer survivors (21). Similar results were investigated in other neurological diseases such as major depressive disorders (33) and temporal lobe epilepsy (32). A previous study showed that the degree distribution of brain network followed the exponentially truncated power law (60). This exponentially truncated power law distribution may be helpful in resisting the targeted attack of the hubs, which means that the brain networks of two groups were almost constant when deletion rate was low (61). The deletion ratios reaching 50%, and the decline rate of global efficiency in networks began to exhibit obvious differences. Exploring the resilient of networks actually simulated the process of cognitive decline in all participants. In detail, as the important nodes were deleted, the functional and structural integrity of brain networks were impaired. Additionally, the FN was more resilient than the SN this study, which were similar to these findings in the previous studies (30, 62). A prior study discovered that there was commonly a functional connectivity between regions that have no direct structural connectivity, implying that functional network was a more stable system in brain network (63). Therefore, functional networks were more robust to node removal. Our results may provide a new direction for studying cognitive impairment in patients with RC after surgery and chemotherapy.

**CONCLUSION**

This study explores the effects of depression tendency on brain functional and structural network in patients with RC with surgery and chemotherapy through multimodal brain connectivity analysis. Patients with RC show the abnormal small-worldness property and network topological organization in FN and SN. The alterations in nodal parameter are mainly observed in the limbic and parietal lobes as well as the subcortical nuclei in patients with RC. The patients with RC demonstrate significant cognitive impairment compared with HC, and this impairment may be associated with lower network attack tolerance. The discovery of functional and structural networks is critical for understanding the neurobiological mechanism associated with depression tendency in patients with RC with surgery and chemotherapy.

**LIMITATION**

The lack of follow-up data limited the ability of studying the causal relationship between alterations in brain network and depression tendency of patients with RC. The statistical power is restricted by small sample size to some extent. Finally, this study lacks the joint analysis for multimodal data. It is very meaningful to use different modal data for fusion research.
DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

ETHICS STATEMENT

The studies involving human participants were reviewed and approved by Medical Ethics Committee of Gansu Provincial People's Hospital. The patients/participants provided their written informed consent to participate in this study.

AUTHOR CONTRIBUTIONS

Conceived and designed the experiments: WZ, YZ, FZ, NM, GH, ZY, and BH. Analyzed the data and wrote the paper: WZ, YY, and YZ. Contributed reagents/materials/analysis tools: WZ, YZ, YL, GH, ZY, and BH. All authors contributed to and have approved the final manuscript.

FUNDING

This work was supported in part by the National Key Research and Development Program of China (Grant No. 2019YFA0706200), in part by the National Key Research and Development Program of China (Grant No. 2016YFC1307203), in part by the National Natural Science Foundation of China (Grant Nos. 61632014, 61627808, 62176140, and 82001775), in part by the Natural Science Foundation of Gansu Province of China (Grant No. 20JR5RA292), and in part by the Department of Education of Gansu Province: Innovation Star Project for Excellent Postgraduates (2021CXZX-121), in part by Gansu Provincial Hospital Youth Research Fund Project, 18GSSY5-5, in part by the Natural Science Foundation of Shandong Provincial of China (Grant Nos. ZR2021MH120 and ZR2020MG013), in part by the National Social Science Foundation of China (Grant No. 20BSH151), in part by Special Fund of Shandong Medical Association (YXH2021ZX055), in part by the Doctoral Scientific Research Foundation of Shandong Technology and Business University (Grant No. BS 202016), in part of Horizontal project of Yantai Vocational College (No. HX2021041).

ACKNOWLEDGMENTS

We thank the staff of the Department of Radiology of the Gansu Provincial Hospital for their assistance in collecting the data. We thank all participants for their contributions to this article.

REFERENCES

1. Bruno J, Hosseini SH, Kesler S. Altered resting state functional brain network topology in chemotherapy-treated breast cancer survivors. *Neurobiol Dis.* (2012) 48:329–38. doi: 10.1016/j.nbd.2012.07.009
2. Li M, Gu J. Changing patterns of colorectal cancer in China over a period of 20 years. *World J Gastroenterol.* (2005) 11:4685. doi: 10.3748/wjg.v11.i30.4685
3. Wefel JS, Kesler SR, Noll KR, Schagen SB. Clinical characteristics, pathophysiology, and management of noncentral nervous system cancer-related cognitive impairment in adults. *CA Cancer J Clin.* (2015) 65:123–38. doi: 10.3322/caac.21258
4. Morean DF, O’Dwyer L, Cherney LR. Therapies for cognitive deficits associated with chemotherapy for breast cancer: a systematic review of objective outcomes. *Arch Phys Med Rehabil.* (2015) 96:1880–97. doi: 10.1016/j.apmr.2015.05.012
5. Linden W, Vodermaier A, MacKenzie R, Greig D. Anxiety and depression after cancer diagnosis: prevalence rates by cancer type, gender, and age. *J Affect Disord.* (2012) 141:343–51. doi: 10.1016/j.jad.2012.03.025
6. Kesler SR, Rao A, Blayney DW, Oakley-Girvan IA, Karuturi M, Palesh O. Predicting long-term cognitive outcome following breast cancer with pre-treatment resting state fMRI and random forest machine learning. *Front Hum Neurosci.* (2017) 11:555. doi: 10.3389/fnhum.2017.00555
7. Zhao F, Zhang X, Thung K-H, Mao N, Lee S-W, Shen D. Constructing Multi-view High-order Functional Connectivity Networks for Diagnosis of Autism Spectrum Disorder. *IEEE Trans Biomed Eng.* (2021) 69:1237–50. doi: 10.1109/TBME.2021.3122813
8. Billiet T, Emsel L, Vandenbulcke M, Peeters R, Christiaens D, Leemans A, et al. Recovery from chemotherapy-induced white matter changes in young breast cancer survivors? *Brain Imaging Behav.* (2018) 12:64–77. doi: 10.1007/s11862-016-9665-8
9. Eberling JL, Wu C, Tong-Turnbeaugh R, Jagust WJ. Estrogen-and tamoxifen-associated effects on brain structure and function. *Neuroimage.* (2004) 21:364–71. doi: 10.1016/j.neuroimage.2003.08.037
10. McDonald BC, Saykin AJ. Alterations in brain structure related to breast cancer and its treatment: chemotherapy and other considerations. *Brain Imaging Behav.* (2013) 7:374–87. doi: 10.1007/s11862-013-9256-x
11. Janelins MC, Kohli S, Mohile SG, Usuki K, Ahles TA, Morrow GR. An Update on Cancer-and Chemotherapy-Related Cognitive Dysfunction: Current Status, Seminars in Oncology. Elsevier (2011). p. 431–8. doi: 10.1053/j.seminoncol.2011.03.014
12. Vahdaninia M, Omidvari S, Montazeri A. What do predict anxiety and depression in breast cancer patients? a follow-up study. *Soc Psychiatry Psychiatr Epidemiol.* (2010) 45:355–61. doi: 10.1007/s00127-009-0068-7
13. Plis SM, Weisend MP, Damaraju E, Eichele T, Mayer A, Clark VP, et al. Effective connectivity analysis of sMRI and MEG data collected under identical paradigms. *Comput Biol Med.* (2011) 41:1156–65. doi: 10.1016/j.compbiomed.2011.04.011
14. Spisz T, Opposits G, Kis S, Pohubi L, Jakab A, Puskás S, et al. BrainCON: graph theory based multimodal brain connectivity analysis and visualization software. *Eur Cong Radiol.* (2013) 2013:2013. doi: 10.1594/ecr2013/C-2588
15. Li Y, Yao Z, Yang Y, Zhao F, Fu Y, Zou Y, et al. A study on PHF-Tau network topology in chemotherapy-treated breast cancer survivors. *Front Aging Neurobiol.* (2017) 109:djx085. doi: 10.1093/jnci/djx085
16. Bullmore ET, Bassett DS. Brain graphs: graphical models of the human brain connectome. *Ann Rev Clin Psychol.* (2011) 7:113–40. doi: 10.1146/annurev-clinpsy-040510-143994
17. Sporns O. The human connectome: a complex network. *Ann N Y Acad Sci.* (2011) 1224:109–25. doi: 10.1111/j.1749-6632.2010.05888.x
18. Cinmrich B, Reuter-Lorenz P, Nelson J, Clark PM, Therrien B, Normolle D, et al. Prechemotherapy alterations in brain function in women with breast cancer. *J Clin Exp Neuropsychol.* (2010) 32:324–31. doi: 10.1080/13803390903032537
19. Kesler SR, Bennett FC, Mahaffey ML, Spiegel D. Regional brain activation during verbal declarative memory in metastatic breast cancer. *Clin Cancer Res.* (2009) 15:6665–73. doi: 10.1158/1078-0432.CCR-09-1227
20. Amidi A, Hosseini SH, Leemans A, Kesler SR, Agerbæk M, Wu LM, et al. Changes in brain structural networks and cognitive functions in testicular cancer patients receiving cisplatin-based chemotherapy. *J Natl Cancer Inst.* (2017) 109:djx085. doi: 10.1093/jnci/djx085
21. Kesler SR, Watson CL, Blayney DW. Brain network alterations and vulnerability to simulated neurodegeneration in breast cancer. *Neurobiol Aging.* (2015) 36:2429–42. doi: 10.1016/j.neurobiolaging.2015.04.015
61. Friedman EJ, Landsberg AS. Hierarchical networks, power laws, and neuronal avalanches. *Chaos.* (2013) 23:013135. doi: 10.1063/1.4793782
62. Damoiseaux JS, Rombouts S, Barkhof F, Scheltens P, Stam CJ, Smith SM, et al. Consistent resting-state networks across healthy subjects. *Proc Nat Acad Sci.* (2006) 103:13848–53. doi: 10.1073/pnas.0601417103
63. Honey C, Sporns O, Cammoun L, Gigandet X, Thiran J-P, Meuli R, et al. Predicting human resting-state functional connectivity from structural connectivity. *Proc Nat Acad Sci.* (2009) 106:2035–40. doi: 10.1073/pnas.0811168106

**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

**Publisher's Note:** All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

Copyright © 2022 Zhang, Zou, Zhao, Yang, Mao, Li, Huang, Yao and Hu. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.