Exploring Young Adulthood Psychopathology Networks Related to Depression Using Partial Correlation Network and Bayesian Network

Min Seob Kim

Graduate School of Medical Science and Engineering, Korea Advanced Institute of Science and Technology (KAIST)

Bumseok Jeong (✉ bs.jeong@kaist.ac.kr)

Graduate School of Medical Science and Engineering, Korea Advanced Institute of Science and Technology (KAIST)

Research Article

Keywords: broad spectrum of symptoms, acyclic graph, depressive symptoms in young adults, mental health management

Posted Date: January 8th, 2021

DOI: <https://doi.org/10.21203/rs.3.rs-131241/v1>

License: © ⓘ This work is licensed under a Creative Commons Attribution 4.0 International License.
[Read Full License](#)

Abstract

To characterize young adulthood depression is complicated because it is entangled with a broad spectrum of symptoms as well as traumatic experiences during development. However, previous symptom network studies have focused on undirected transdiagnostic association among depression and anxiety symptoms. Our study investigated both undirected and directed connections among variables potentially associated with depression, such as anxiety, addiction, subjective distress caused by traumatic events, perceived emotional adversities, and support systems. Both the regularized partial correlation network analysis and Bayesian network analysis were applied to 579 subjects screened for depression. Anxiety-related symptoms played a role as a hub node in the partial correlation network and Bayesian network. The vulnerability analysis of the partial correlation network showed that verbal abuse, social anxiety, concentration problems, and suicidal ideation had the strongest influence on changes in the network's topology. In the Bayesian network analysis, loss of interest, depressed mood, and parental verbal abuse were located as parent nodes in the directed acyclic graph. In the aspect of disease networks, more attention should be paid to certain variables encompassing various domains as well as depressive symptoms in young adults' mental health management.

Introduction

A characteristic feature of psychopathology in young adulthood depression is a broad spectrum of symptoms. It is also known that the probability of having a family history of mood disorders; being diagnosed with bipolar disorder in the future; and accompanying symptoms, such as substance abuse¹ as well as irritability and anxiety² are more common than in young adults than in other age groups. Therefore, when screening young adulthood depression, it is difficult for clinicians to determine whether particular symptoms are due to depression or other mental disorders, including generalized anxiety disorder, social anxiety disorder, or substance use disorder³

There are important developmental tasks to accomplish in the transitional period between the ages of 20 and 30. However, the experience of depression at the transitional period is likely to impair appropriate functions to accomplish these tasks⁴. Functional impairments during this period appear to persist for a substantial time after the remission of depression^{5,6}. In addition, emotional adversities experienced before young adulthood cause structural⁷⁻⁹, functional^{10,11}, and neurochemical¹² alterations in the brain. Therefore, various variables consisting of trans-diagnostic symptoms, related behavioral patterns, and current and past stresses may be intricately intertwined in young adulthood depression. Those variables can be connected in a subjective causal space (e.g., the experience of being criticized by one's parents as a child seems to make one nervous when a stressful situation arises). However, it was not clear how these variables were connected in past psychopathology studies, and this uncertainty was a factor that confused clinicians. This is partly due to our poor understanding of how variables that can potentially affect depression are associated with certain age groups.

A network analysis does not just consider certain variables to be the phenotypes of unobservable latent variables¹³. Instead, network analysis interprets phenotypes as fundamental components of certain disorders. In terms of network analysis, episodes of mental disorder appear because of interactions between symptoms^{14,15}. In the same vein, comorbidity of mental disorders occurs due to the interactions between psychopathologies' entangled networks. We believed that the network-based analysis method would be suitable for studying the relationships of psychopathologies related to depression in young adulthood that can exhibit a wide range of symptoms. There have been several studies on network analysis of psychopathology in young adulthood. However, most studies were conducted on groups in the general population, such as college students¹⁶, or on the general population focusing on eating disorders¹⁷. In our previous study¹⁸, using the same sample, the effect of verbal abuse on psychiatric symptoms was mainly studied. Other studies have investigated the symptom network of depression, but it seems that there were limitations, such as studying networks limited to only depression or anxiety symptoms^{19,20}. However, in this study, we focused on the high depression symptom (high-DS) group selected from the parent group. Through network analysis, we aimed to identify the associations of not only symptoms of young adulthood depression, but also other variables, such as social anxiety disorder, substance abuse, the number of mentors, the number of concerns, and experience of verbal abuse by parents, which may be potentially related to young adulthood depression.

In this study, we sought to determine how psychopathologies or potentially related variables (including psychological or environmental factors) were associated in the young adulthood age group and how the symptoms of the depressive disorder of young adulthood could be relatively various in the aspect of network analysis. For that purpose, we conducted a network analysis of various questionnaires on psychopathologies of 5,615 subjects who were college students and 579 subjects screened for depression from the 5,615 subjects. To the best of our knowledge, this is the first study to explore the psychopathology of depression and potentially related variables associated with depressive symptoms in the young adulthood age group and in the subgroup screened for depression based on the Patient Health Questionnaire-9 (PHQ-9)²¹ which is widely used in the screening and diagnosis of depression²².

Methods

Participants

We used data from self-reported questionnaires, which was part of the annual healthcare screening of 5,685 college students' mental health at Korea Advanced Institute of Science and Technology (KAIST) in Daejeon, South Korea, between April 2014 and February 2015. Respondents knew that their responses would not be used for any purpose other than self-affirmation of their mental health. The responses of 5,615 students between the ages of 18 and 30 were used among the gathered data, and samples with missing data were excluded. We used the PHQ-9 scores to define the subjects screened for depression. The group was defined as respondents with PHQ-9 scores of five or more while reporting that they were experiencing functional impairment due to the depressive symptoms of PHQ-9²¹. The group was

expected to correspond to mild or more severe depression, herein called the high depression symptom severity group (high-DS group). Among the 5,615 subjects, 579 subjects were screened for the high-DS group.

Measures

The survey for this study was conducted through an online website while ensuring anonymity and confidentiality. We conducted surveys of various types of variables that could be associated with depression in the students of KAIST and used the results of the surveys as nodes in our network analysis: Gender; smoking history (Smoking); alcohol abuse (CAGE)²³; the number of concerns (Concerns); the number of mentors within and outside the family (Ment_intrafam, Ment_extrafam); the smartphone addiction scale (SAS)²⁴; the Generalized Anxiety Disorder-7 (GAD)²⁵; Leibowitz social anxiety scale (LSAS)²⁶; the experience of verbal abuse from parents, superiors, and peers (VA_parents, VA_superiors, VA_peers)²⁷; the impact of event scale - revised (IESR)²⁸; and 9 items of the patient health questionnaire-9 (PHQ_01, PHQ_02, PHQ_03, PHQ_04, PHQ_05, PHQ_06, PHQ_07, PHQ_08, PHQ_09)²¹.

To focus on the high-DS group with a sample size of only 10% of the parent group, unlike our previous study¹⁸, which constructed a network using all individual items, the total scores of the each measure were used, excluding PHQ-9. The details of the variables we used are presented in Table 1.

Table 1. List of the variables (nodes) that comprising the networks.

Domains	Variables	Mean (S.D.)
Gender	Gender (0 : Male, 1: Female)	0.3 (0.46)
Depression	PHQ-1 (Lack of interest)	1.29 (0.87)
	PHQ-2 (Depressed mood)	1.27 (0.72)
	PHQ-3 (Sleeping difficulties)	1.38 (0.96)
	PHQ-4 (Fatigue)	1.63 (0.85)
	PHQ-5 (Appetite problems)	1.18 (0.92)
	PHQ-6 (Feelings of Worthlessness)	1.09 (0.88)
	PHQ-7 (Concentration problems)	0.60 (0.81)
	PHQ-8 (Psychomotor agitation/retardation)	0.34 (0.62)
	PHQ-9 (Suicidal ideation)	0.29 (0.59)
Anxiety	GAD (Generalized anxiety)	5.87 (4.10)
	LSAS (Liebowitz social anxiety scale)	35.76 (24.29)
	IESR (Impact of event scale – revised)	18.35 (15.80)
Addiction	CAGE (Alcohol abuse)	0.43 (0.76)
	Smoking (0: Never Smoker 1: ex-smoker 2: current smoker)	1.34 (0.70)
	SAS (Smartphone addiction scale)	78.89 (24.29)
Perceived verbal abuse	VA_parents (perceived verbal abuse from parents)	7.58 (12.94)
	VA_superiors (perceived verbal abuse from superiors)	6.63 (10.22)
	VA_peers (perceived verbal abuse from peers)	8.95 (13.00)
Support systems	Mentor_intrafam_No (The number of mentors inside family)	1.29 (1.33)
	Mentor_extrafam_No (The number of mentors outside family)	3.78 (4.10)
Stresses	Concern (the number of major concerns)	2.49 (1.66)

Analyses

- *Partial correlation networks*

In the first analysis, a graphical Gaussian model was used to construct the networks, in which the edges represent partial correlations between nodes controlling all other nodes' effect. To construct the regularized partial correlation networks, we used a graphical LASSO algorithm²⁹ implemented in the *EBICglasso* function of R package *qgraph*³⁰. The network construction process is briefly summarized as follows. Because the LASSO algorithm makes the network parsimonious while maintaining the partial correlation network's explanatory power, the LASSO penalty was used to zero the edges with insignificantly small partial correlation values. Also, by using the model comparison with the extended Bayesian information criterion, we found the most optimal parameter (λ of the LASSO penalty, and we built a parsimonious network that best describes the model using the parameter. We set the hyperparameter (γ) value to 0.5³¹. The degrees of association between two connected nodes were indicated by the thickness of the edges. The signs of the partial correlations were indicated by the color of the edges.

Furthermore, we estimated the strength centrality, closeness, and the betweenness centrality to explore each node's importance in the obtained network. The first metric was computed by summing the values of the edges connected to each node. The second was computed by summing the length of the shortest paths between each node and all other nodes. The third was computed by counting the number of times a specific node appeared on the shortest path connecting all possible pairs of two different nodes. We assumed that nodes with high values of these metrics would play a more important role in the network. In addition, we used the R package *bootnet*³² to measure the stability of the network. Through 1000 times of bootstrapping, we measured metrics, including the confidence interval of each edge's strength.

- *Bayesian networks (DAG)*

We computed the Bayesian network using the random-restart hill-climbing algorithm implemented in R package *bnlearn*³³. First, it looks for a network structure that optimizes the Bayesian information criterion by adding, removing, or reversing the edge of the network. This process randomly repeated 100 times to prevent the hill-climbing algorithm from falling into the local optima. This procedure only determines whether the edges exist in the network and what the directions of the edges would be.

To obtain stable results, we extracted 1,000 sample networks through bootstrapping with resampling. Then, we checked how often each edge appeared in the networks obtained through bootstrapping. If the edges appeared in more than 85%³⁴ of the total sample networks, we included them in the final averaged DAG. In addition, if at least half of the sampled 1,000 networks had the same direction, then the direction would be illustrated in the final averaged network. Note that while the DAG construction algorithm was applied, edges were excluded toward gender or verbal abuse from any other nodes ("blacklisting")³³.

- *Vulnerability analysis of regularized partial correlation network*

In the field of neuroimaging, there have been studies on how the changes in regional brain networks affect the overall network topology^{35,36}. In our analysis, we investigated how local changes (or a therapeutic approach to the nodes) in nodes affect networks' overall integrity. Artificial intervention was applied by replacing each node with random values that follow a uniform distribution between 0 and 1. The intervened regularized partial correlation network for each node was sampled through jackknife resampling. In other words, sampled intervened regularized partial correlation networks were obtained for every 22 intervened nodes (579×22 networks). The distribution of the intact network was also obtained using jackknife resampling. The distributions of global efficiencies in 579 intervened networks for each node and 579 intact networks were compared through 22 Wilcoxon Rank-Sum tests with multiple comparison correction (Bonferroni correction). This comparison allowed us to evaluate how much intervention to each node (or a therapeutic approach to each node) would affect the entire psychopathology network's global efficiency or clustering coefficient. Through this study, we tried to determine which nodes' preferential treatment is most effective in terms of network topology.

Ethical Standard

The Institutional Review Board at Korea Advanced Institute of Science and Technology (KAIST) approved the current study (IRB approval no. KH-2012-16), and written informed consent was obtained from all subjects after the procedures had been fully explained. All procedures were performed in accordance with the ethical standards of the KAIST IRB on human experimentation and the Helsinki Declaration of 1975, as revised in 2008.

Results

Partial correlation network

The partial correlation network of high-DS subjects is shown in Fig. 1. The number of non-zero edges was 66 (28.6%). Most of the edges on the network showed positive correlation coefficients (83.33%). It was found that nodes belonging to the same symptom domains tended to form a cluster and were connected within the cluster. For example, the PHQ-01, 02, 06, 09, called cognitive/affective symptoms, tended to cluster together, and the PHQ-03, 04, 05, 06, 07, 08, which are somatic symptoms³⁷, tended to cluster together. IESR (subjective distress caused by traumatic events), GAD (generalized anxiety), and LSAS (social anxiety), which are classified as anxiety-related symptoms, were also clustered with strong correlations. Interestingly, nodes related to perceived distress due to verbal abuse by parents, peers, and superiors were also clustered. It seems that PHQ_02, GAD, and IESR act as hub nodes that mediate the psychopathology network's symptoms.

Regarding each edge in the network of high-DS subjects, the five edges with the strongest positive correlations were between PHQ_01 and PHQ_02 (0.45); VA_parents and VA_peers (0.32); GAD and IESR (0.27); VA_superiors and VA_peers (0.26); and PHQ_02 and PHQ_06 (0.25). The five edges with the strongest negative correlations were between gender (female) and smoking (-0.24); LSAS and

Num_mentor_extrafam (-0.12); PHQ_07 and Num_mentor_extrafam (-0.11); LSAS and Num_mentor_intrafam (-0.10); VA_parents and Num_mentor_intrafam (-0.07). Confidence intervals for the strength of edges obtained by 1,000 bootstrapped networks are illustrated in Fig. 2.

Fig. 3 shows three centrality metrics of the network for high-DS subjects. The five nodes with the greatest strength centrality were, in descending order, PHQ_02 (1.75), IESR (1.44), GAD (1.40), PHQ_07 (1.03), and LSAS (0.92). The five nodes with the greatest closeness centrality were IESR (2.23), GAD (2.09), LSAS (1.14), PHQ_06 (0.85), and PHQ_02 (0.67). The five nodes with the greatest betweenness centrality were IESR (2.96), GAD (2.37), LSAS (0.91), PHQ_06 (0.84), and PHQ_02 (0.62).

Bayesian Network

It was found that the core symptoms (PHQ_01 and PHQ_02) of depression were at the top of DAG. It was found that the nodes with greater numbers of edges in the DAG were PHQ_06, IESR, and GAD. It was also found that female gender, known as a risk factor for depressive disorder, had no significant effect on the interaction of depression- or anxiety-related symptom clusters. Instead, the male gender seemed to be related to alcohol and nicotine addiction; however, the cluster was not associated with a depression or anxiety forming 'island'. In our results, these three nodes (Gender, Smoking, and CAGE) seemed not to contribute to maintaining the network of depression and potentially related variables. Among the symptoms of addiction, only smartphone addiction was connected with the depression network.

The edges related to VA_parents were noteworthy as well. Edges from VA_parents to SAS and PHQ_03 were observed in the network of all subjects. However, edges connecting VA_parents to other nodes was not observed in the network of high-DS subjects. See Figures 4 for detailed network information.

Vulnerability analysis of regularized partial correlation network

The vulnerability analysis revealed that all nodes except Smoking, Gender, and CAGE were significant for the decrease in global efficiency of the network. The nodes were significant even after multiple comparison correction (Bonferroni correction). Figure 5A shows each node's median of the differences of global efficiency in the one-sided Wilcoxon Rank-sum test. The ten nodes showing the greatest median of the differences were the following in descending order: LSAS (8.43), VA_parents (8.36), PHQ_07 (7.72), VA_peers (6.88), PHQ_09 (6.76), PHQ_06 (4.27), Ment_intrafam (4.76), PHQ_02 (4.15), SAS (3.90), and PHQ_05 (3.45).

Regarding the impact to the clustering coefficient of each node, the significant nodes were VA_peers, VA_parents, PHQ_09, PHQ_07, PHQ_06, PHQ_04, Ment_intrafam, LSAS, and IESR. Figure 5B shows each node's median of the differences of clustering coefficient in the one-sided Wilcoxon Rank-sum test. The nodes were significant with multiple comparison correction (Bonferroni correction). The significant nine nodes' median of the differences were the following: PHQ_09 (0.034), LSAS (0.028), VA_parents (0.026), PHQ_07 (0.028), VA_peers (0.018), Ment_intrafam (0.007), PHQ_04 (0.007), IESR (0.006), and PHQ_06 (0.006).

Discussion

In this study, we performed a network analysis with depressive symptoms and variables potentially related to depression in subjects screened for depression (N=579). The nodes with high centrality measures identified in regularized partial correlation networks were IESR, GAD, and PHQ_02. It was found that PHQ_01, PHQ_02, PHQ_03, and VA_parents were at the higher part of the Bayesian network. It seems that our results were consistent with the expectation that various variables would be closely linked beyond each domain, and the networks adequately reflect existing medical knowledge. For example, the core symptoms of depression, namely, loss of interest (PHQ_01), and depressed mood (PHQ_02), occupied the higher part in the causal relationship and showed relatively higher centrality scores. In addition, when the connected edges were removed (vulnerability analysis), the five nodes that had the most significant impact on the decrease in the integrity of the entire network were PHQ_09, LSAS, VA_parents, PHQ_07, and VA_peers. Our results suggest that these nodes could become primary treatment target symptoms that can effectively reduce the integrity of the entire psychopathology network of high-DS subjects.

It is well known that depression at a younger age is associated with more anxiety symptoms³⁸. It is also common for these two symptoms to occur together³⁹. Some researchers have even argued that anxiety and depression can be explained by a single factor model^{3,40}. As a matter of fact, the specifier “with anxious distress” was added to the *Diagnostic and Statistical Manual for Mental Disorders, 5th Edition* (DSM-5)⁴¹ to describe the symptoms of anxiety that frequently accompany depression. The fact that patients become anxious when depressed and depressed when anxious is a phenomenon often experienced in clinical settings. Corresponding results were also observed in our study; the symptoms of anxiety and depression were intertwined, according to the results of our network analysis. Impressively, anxiety-related symptoms (IESR, GAD, and LSAS) appeared to be a hub node in the psychopathology networks. The finding was commonly observed in both the regularized partial correlation network and the Bayesian network. Generalized anxiety seemed to act as a bridge node between cognitive/affective (PHQ_01, 02, 06, 09) and somatic factors (PHQ_03, 04, 05, 07, 08) of depression³⁷ in the Bayesian network. GAD was also the node linking subjective distress caused by traumatic events (IESR) and depressive symptoms (PHQ_02, PHQ_06). These findings are consistent with those of previous studies showing that depressed mood and anxiety are associated with somatic symptoms⁴². In addition, there have been studies on the effects of generalized anxiety disorder, panic disorder, and major depressive disorder on somatic complaints using structural equation modeling⁴³. No direct associations between emotional awareness and somatic complaints were found; however, there were direct associations among depression, anxiety, and somatic complaints. Also, recent studies on heartbeat evoked potential (HEP) have also found that generalized anxiety⁴⁴ or social anxiety⁴⁵ were associated with an inadequate increased HEP. These results suggest that anxiety symptoms are associated with abnormally increased somatosensory sensitivity of body sensation. Our study seems to be in line with these findings as well because our network analysis results obtained from young adult subjects can be interpreted as showing psychopathology in which cognitive/affective symptoms spread to somatic symptoms of depression and

other associated symptoms of depression through generalized anxiety both in regularized partial correlation network and Bayesian network.

One notable point was that the depressive symptoms were more strongly associated with SAS (smartphone addiction) than CAGE, Smoking in the network of high-DS subjects. In the regularized partial correlation network, SAS was connected to LSAS, Concerns, VA_parents, VA_peers, and PHQ_07. In the Bayesian network, it was connected to Concerns and LSAS. However, CAGE and Smoking were not connected to depression or anxiety symptoms. This may reflect the bias of the college students' sample. However, some studies have reported that smartphone addiction was related to shyness, loneliness⁴⁶, low self-esteem, and aggressive behaviors⁴⁷. These studies commonly mentioned that smartphone addiction might promote the development of depressive disorders. Our study suggests that smartphone addiction might be linked to stresses (the number of concerns, verbal abuses), social anxiety symptoms (LSAS), and concentration problems (PHQ_07). Hence, it could be a facilitating factor for depression in young adults. Contrary to what we have previously known, it may be important to consider that smartphone addiction could be more associated with depressive symptoms than substance addiction in young adulthood depression.

In addition, in our study, we examined changes in the whole network's topology by comparing it with the intact network after damage to each node. We were able to determine which node was more effective in reducing the integrity of the whole psychopathology network. Inducing a local change in a network and observing a change in the overall topology is different from determining a node's importance through a centrality measure in an intact network³⁶. The former makes it possible to observe the topology change of the whole variable network due to the change in each node, and the latter only represents the importance of the nodes that make up the whole psychopathology network. The analysis allowed us to identify symptoms that require intervention to reduce the connectivity of the entire disease network. It was expected that if intervention for certain variables was prioritized, such as social anxiety (LSAS), verbal abuse (VA_parents and VA_peers), concentration problems (PHQ_07), and suicidal ideation (PHQ_09), it would be possible to stabilize the overall psychopathological network more efficiently in terms of both global efficiency and clustering coefficient.

The advantage of our study was that we aimed to observe the psychopathology network through various statistical aspects of the network, such as partial correlation network, Bayesian network, and how the topology of the whole network changes when each node of the network is intervened. In addition, rather than using only the scales limited to depression or anxiety, our study had the strength of using various scales, including generalized anxiety, social anxiety, subjective distress due to traumatic events, addiction (alcohol, nicotine, smartphone), the number of concerns, the number of mentors, and perceived verbal abuse. This is expected to be more advantageous than previous analysis using only depression or anxiety symptoms in that both social and environmental factors were included to explain the psychopathology networks. This enabled more appropriate network analysis in that it utilized as many variables as possible that could affect the network of psychopathology. However, our study was still limited in that we used data collected cross-sectionally at a particular time point. To compensate for this

limitation, we used not only a graphical Gaussian model but also Bayesian network analysis, which may represent the information of causal relationships because Bayesian network (DAG) analysis is relatively useful in inferring the causal relationship of symptoms in situations where time-series data are not available. However, it is worth noting that the directions of arrows in the DAG does not necessarily indicate causal relationships. The graph from A to B to C and the graph from C to B to A are identical in terms of conditional independence. Certainly, the DAG represents at least the associations between nodes; however, it is challenging to be sure that DAG represents the causal relationships between nodes. In the future, we anticipate that network analysis of psychopathology should be conducted using information gathered at various time points.

In terms of psychopathology, our research revealed that not only variables in a limited domain should be considered significant; rather, variables in multiple domains should be considered comprehensively. In addition, the analysis of the network showed that certain variables were more important in terms of centrality, causal relationship, and the potential to lower the integrity of the network. And the specific variables were not limited to depressive symptoms but encompassed various domains. We suggest that the understanding centered on the hub or the bridge node of the network and the treatment centered on the node that can significantly lower the integrity of the network would be helpful in the diagnosis and treatment of young adults' depression.

Declarations

Acknowledgments and Disclosures

We thank GeumSook Shim for recruiting subjects and collecting self-report measures. This research was supported by the Brain Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Science & ICT (NRF-2016M3C7A1914448 and NRF-2017M3C7A1031331). 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.

Contributions

M.S.K. and B.S.J. designed the study and prepared the manuscript. B.S.J. recruited subjects and collected self-report measures. All authors reviewed the manuscript.

Data Availability

All data generated and analyzed during this study are available from the corresponding author on a reasonable request.

References

1. Parker, G., Roy, K., Hadzi-Pavlovic, D., Mitchell, P. & Wilhelm, K. Distinguishing early and late onset non-melancholic unipolar depression. *Journal of affective disorders* **74**, 131-138, doi:[https://doi.org/10.1016/S0165-0327\(02\)00002-2](https://doi.org/10.1016/S0165-0327(02)00002-2) (2003).
2. Roza, S. J., Hofstra, M. B., van der Ende, J. & Verhulst, F. C. Stable prediction of mood and anxiety disorders based on behavioral and emotional problems in childhood: a 14-year follow-up during childhood, adolescence, and young adulthood. *The American journal of psychiatry* **160**, 2116-2121, doi:10.1176/appi.ajp.160.12.2116 (2003).
3. Waszczuk, M. A., Zavos, H. M. S., Gregory, A. M. & Eley, T. C. The Phenotypic and Genetic Structure of Depression and Anxiety Disorder Symptoms in Childhood, Adolescence, and Young Adulthood. *JAMA Psychiatry* **71**, 905-916, doi:10.1001/jamapsychiatry.2014.655 %J JAMA Psychiatry (2014).
4. Paradis, A. D., Reinherz, H. Z., Giaconia, R. M. & Fitzmaurice, G. Major Depression in the Transition to Adulthood: The Impact of Active and Past Depression on Young Adult Functioning. **194**, 318-323, doi:10.1097/01.nmd.0000217807.56978.5b (2006).
5. Lewinsohn, P. M., Rohde, P., Klein, D. N. & Seeley, J. R. Natural course of adolescent major depressive disorder: I. Continuity into young adulthood. *Journal of the American Academy of Child and Adolescent Psychiatry* **38**, 56-63, doi:10.1097/00004583-199901000-00020 (1999).
6. Christie, K. A. *et al.* Epidemiologic evidence for early onset of mental disorders and higher risk of drug abuse in young adults. *The American journal of psychiatry* **145**, 971-975, doi:10.1176/ajp.145.8.971 (1988).
7. Choi, J., Jeong, B., Rohan, M. L., Polcari, A. M. & Teicher, M. H. Preliminary evidence for white matter tract abnormalities in young adults exposed to parental verbal abuse. *Biol Psychiatry* **65**, 227-234, doi:10.1016/j.biopsych.2008.06.022 (2009).
8. Choi, J., Jeong, B., Polcari, A., Rohan, M. L. & Teicher, M. H. Reduced fractional anisotropy in the visual limbic pathway of young adults witnessing domestic violence in childhood. *Neuroimage* **59**, 1071-1079, doi:10.1016/j.neuroimage.2011.09.033 (2012).
9. Lee, S. W. *et al.* Hippocampal Subfields Volume Reduction in High Schoolers with Previous Verbal Abuse Experiences. *Clin Psychopharmacol Neurosci* **16**, 46-56, doi:10.9758/cpn.2018.16.1.46 (2018).
10. Lee, S. W. *et al.* Aberrant function of frontoamygdala circuits in adolescents with previous verbal abuse experiences. *Neuropsychologia* **79**, 76-85, doi:<https://doi.org/10.1016/j.neuropsychologia.2015.10.029> (2015).
11. Lee, S. W. *et al.* Altered Function of Ventrolateral Prefrontal Cortex in Adolescents with Peer Verbal Abuse History. *Psychiatry Investig* **14**, 441-451, doi:10.4306/pi.2017.14.4.441 (2017).
12. Kim, D. *et al.* Anatomical and Neurochemical Correlates of Parental Verbal Abuse: A Combined MRS-Diffusion MRI Study. *Front Hum Neurosci* **13**, 12-12, doi:10.3389/fnhum.2019.00012 (2019).
13. Borsboom, D. & Cramer, A. O. Network analysis: an integrative approach to the structure of psychopathology. *Annual review of clinical psychology* **9**, 91-121, doi:10.1146/annurev-clinpsy-050212-185608 (2013).

14. Eaton, N. R. Latent variable and network models of comorbidity: toward an empirically derived nosology. *Social Psychiatry and Psychiatric Epidemiology* **50**, 845-849, doi:10.1007/s00127-015-1012-7 (2015).
15. Cramer, A. O., Waldorp, L. J., van der Maas, H. L. & Borsboom, D. Comorbidity: a network perspective. *The Behavioral and brain sciences* **33**, 137-150; discussion 150-193, doi:10.1017/s0140525x09991567 (2010).
16. Yun, J.-Y. *et al.* Hubness of strategic planning and sociality influences depressive mood and anxiety in College Population. *Scientific Reports* **7**, 17856, doi:10.1038/s41598-017-18189-x (2017).
17. Sahlan, R. N. *et al.* Disordered eating, self-esteem, and depression symptoms in Iranian adolescents and young adults: A network analysis. *n/a*, doi:<https://doi.org/10.1002/eat.23365>.
18. Yun, J.-Y., Shim, G. & Jeong, B. Verbal Abuse Related to Self-Esteem Damage and Unjust Blame Harms Mental Health and Social Interaction in College Population. *Scientific reports* **9**, 5655-5655, doi:10.1038/s41598-019-42199-6 (2019).
19. Park, S.-C. & Kim, D. The Centrality of Depression and Anxiety Symptoms in Major Depressive Disorder Determined Using a Network Analysis. *Journal of affective disorders* **271**, 19-26, doi:<https://doi.org/10.1016/j.jad.2020.03.078> (2020).
20. An, M. H. *et al.* Depressive Symptom Network Associated With Comorbid Anxiety in Late-Life Depression. **10**, doi:10.3389/fpsyt.2019.00856 (2019).
21. Kroenke, K., Spitzer, R. L. & Williams, J. B. W. The PHQ-9. *Journal of General Internal Medicine* **16**, 606-613, doi:10.1046/j.1525-1497.2001.016009606.x (2001).
22. Levis, B., Benedetti, A. & Thombs, B. D. Accuracy of Patient Health Questionnaire-9 (PHQ-9) for screening to detect major depression: individual participant data meta-analysis. *BMJ* **365**, l1476, doi:10.1136/bmj.l1476 (2019).
23. Ewing, J. A. Detecting Alcoholism: The CAGE Questionnaire. *JAMA* **252**, 1905-1907, doi:10.1001/jama.1984.03350140051025 %J JAMA (1984).
24. Kwon, M. *et al.* Development and validation of a smartphone addiction scale (SAS). *PloS one* **8**, e56936, doi:10.1371/journal.pone.0056936 (2013).
25. Spitzer, R. L., Kroenke, K., Williams, J. B. W. & Löwe, B. A Brief Measure for Assessing Generalized Anxiety Disorder: The GAD-7. *Archives of Internal Medicine* **166**, 1092-1097, doi:10.1001/archinte.166.10.1092 %J Archives of Internal Medicine (2006).
26. Osório, F. d. L., Crippa, J. A. d. S. & Loureiro, S. R. Instruments for the assessment of social anxiety disorder: Validation studies. *World J Psychiatry* **2**, 83-85, doi:10.5498/wjp.v2.i5.83 (2012).
27. Jeong, B. *et al.* The psychometric properties of the korean version of the verbal abuse questionnaire in university students. *Psychiatry Investig* **12**, 190-196, doi:10.4306/pi.2015.12.2.190 (2015).
28. Weiss, D. S. in *Assessing psychological trauma and PTSD, 2nd ed.* 168-189 (The Guilford Press, 2004).

29. Friedman, J., Hastie, T. & Tibshirani, R. Sparse inverse covariance estimation with the graphical lasso. *Biostatistics* **9**, 432-441, doi:10.1093/biostatistics/kxm045 %J Biostatistics (2007).
30. Epskamp, S., Cramer, A., Waldorp, L., Schmittmann, V. & Borsboom, D. qgraph: Network Visualizations of Relationships in Psychometric Data. *Journal of statistical software* **48**, doi:10.18637/jss.v048.i04 (2012).
31. McNally, R. J., Mair, P., Mugno, B. L. & Riemann, B. C. Co-morbid obsessive-compulsive disorder and depression: a Bayesian network approach. *Psychological medicine* **47**, 1204-1214, doi:10.1017/s0033291716003287 (2017).
32. Epskamp, S., Borsboom, D. & Fried, E. I. Estimating psychological networks and their accuracy: A tutorial paper. *Behavior Research Methods* **50**, 195-212, doi:10.3758/s13428-017-0862-1 (2018).
33. Scutari, M. Learning Bayesian Networks with the bnlearn R Package. *2010* **35**, 22 %J Journal of Statistical Software, doi:10.18637/jss.v035.i03 (2010).
34. Sachs, K., Perez, O., Pe'er, D., Lauffenburger, D. A. & Nolan, G. P. Causal protein-signaling networks derived from multiparameter single-cell data. *Science (New York, N.Y.)* **308**, 523-529, doi:10.1126/science.1105809 (2005).
35. Alstott, J., Breakspear, M., Hagmann, P., Cammoun, L. & Sporns, O. Modeling the impact of lesions in the human brain. *PLoS computational biology* **5**, e1000408, doi:10.1371/journal.pcbi.1000408 (2009).
36. Henry, T. R. *et al.* Bridging global and local topology in whole-brain networks using the network statistic jackknife. *Netw Neurosci* **4**, 70-88, doi:10.1162/netn_a_00109 (2020).
37. Beard, C., Hsu, K. J., Rifkin, L. S., Busch, A. B. & Björgvinsson, T. Validation of the PHQ-9 in a psychiatric sample. *Journal of affective disorders* **193**, 267-273, doi:<https://doi.org/10.1016/j.jad.2015.12.075> (2016).
38. Alpert, J. E. *et al.* Patterns of axis I comorbidity in early-onset versus late-onset major depressive disorder. *Biological Psychiatry* **46**, 202-211, doi:[https://doi.org/10.1016/S0006-3223\(99\)00017-7](https://doi.org/10.1016/S0006-3223(99)00017-7) (1999).
39. McLaughlin, T. P., Khandker, R. K., Kruzikas, D. T. & Tummala, R. Overlap of anxiety and depression in a managed care population: Prevalence and association with resource utilization. *The Journal of clinical psychiatry* **67**, 1187-1193, doi:10.4088/jcp.v67n0803 (2006).
40. Kroenke, K. *et al.* Patient Health Questionnaire Anxiety and Depression Scale: Initial Validation in Three Clinical Trials. *Psychosom Med* **78**, 716-727, doi:10.1097/PSY.0000000000000322 (2016).
41. American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders (5th ed.)*. (American Psychiatric Publishing, 2013).
42. Löwe, B. *et al.* Depression, anxiety and somatization in primary care: syndrome overlap and functional impairment. *General Hospital Psychiatry* **30**, 191-199, doi:<https://doi.org/10.1016/j.genhosppsych.2008.01.001> (2008).
43. van der Veek, S. M. C., Nobel, R. A. & Derkx, H. H. F. The relationship between emotion awareness and somatic complaints in children and adolescents: Investigating the mediating role of anxiety and

depression. *Psychology & Health* **27**, 1359-1374, doi:10.1080/08870446.2012.685738 (2012).

44. Pang, J. *et al.* Altered Interoceptive Processing in Generalized Anxiety Disorder-A Heartbeat-Evoked Potential Research. *Front Psychiatry* **10**, 616-616, doi:10.3389/fpsyt.2019.00616 (2019).
45. Judah, M. R. *et al.* The relationship between social anxiety and heartbeat evoked potential amplitude. *Biological Psychology* **139**, 1-7, doi:<https://doi.org/10.1016/j.biopsycho.2018.09.013> (2018).
46. Bian, M. & Leung, L. Linking Loneliness, Shyness, Smartphone Addiction Symptoms, and Patterns of Smartphone Use to Social Capital. *Social Science Computer Review* **33**, 61-79, doi:10.1177/0894439314528779 (2014).
47. Lee, J. *et al.* Psychological Factors Associated With Smartphone Addiction in South Korean Adolescents. *The Journal of Early Adolescence* **38**, 288-302, doi:10.1177/0272431616670751 (2016).

Figures

Figure 1

Regularized partial correlations network of high-DS subjects, depicting the regularized partial correlation between nodes representing depressive symptoms and nodes potentially related to depression. Blue

indicates edges that have positive correlation coefficients. Red indicates edges that have negative correlation coefficients. The thickness of edges indicates the magnitude of the correlation between two nodes.

Figure 2

Confidence intervals for the strength of edges obtained by 1,000 bootstrapped networks in high-DS subjects.

Figure 3

Z-scored centrality of the nodes (strength, closeness, and betweenness) of regularized partial correlation network in high-DS subjects

Figure 5

Results of vulnerability analysis of the regularized partial correlation network of high-DS subjects. Panel A. Result of the change in global efficiency when an intervention is applied to each node. Panel B. Result of the change in clustering coefficient when an intervention is applied to each node. The median values of the differences of nodes in the one-sided Wilcoxon Rank-sum tests were plotted. Statistical significances of variables were defined as $p \leq 0.05/22$ (* $p \leq 0.05/22$, ** $p \leq 0.01/22$, *** $p \leq 0.001/22$). Note

that the nodes with negative median values of differences in the one-sided Wilcoxon Rank-sum tests of clustering coefficients were not plotted in Panel B.