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Differences in resting corticolimbic functional connectivity in bipolar I euthymia

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Abstract

Objective—We examined resting state functional connectivity in the brain between key emotion regulation regions in bipolar I disorder to delineate differences in coupling from healthy subjects.

Methods—Euthymic subjects with bipolar I disorder ($n = 20$) and matched healthy subjects ($n = 20$) participated in a resting state functional magnetic resonance imaging scan. Low frequency fluctuations in blood oxygen level-dependent (BOLD) signal were correlated in the six connections between four anatomically-defined nodes: left and right amygdala and left and right ventrolateral prefrontal cortex (vlPFC). Seed-to-voxel connectivity results were probed for commonly coupled regions. Following this, an identified region was included in a mediation analysis to determine the potential of mediation.

Results—The bipolar I disorder group exhibited significant hyperconnectivity between right amygdala and right vlPFC relative to healthy subjects. The connectivity between these regions in the bipolar I disorder group was partially mediated by activity in the anterior cingulate cortex (ACC).

Conclusions—Greater coupling between right amygdala and right vlPFC and their partial mediation by the ACC were found in bipolar I disorder subjects in remission and in the absence of a psychological task. These findings have implications for a trait-related and clinically-important imaging biomarker.

Keywords

amygdala; bipolar disorder; euthymia; functional connectivity; resting state; ventrolateral prefrontal cortex

Bipolar disorder is a serious psychiatric illness thought to involve deficits in the neural substrates for emotion regulation because of its characteristic profile of lifetime depressed

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and manic mood episodes (1, 2). Neuroimaging research in healthy subjects has revealed a number of brain areas involved in emotional regulation, including subcortical (e.g., amygdala) and ventrolateral prefrontal cortical (vlPFC) regions (3, 4). Research in both humans and primates has additionally demonstrated strong anatomical projections between these areas, forming the substrate for which functional coupling (connectivity) is possible (5, 6). The amygdala, for example, is functionally responsive to emotional stimuli, while the vlPFC is instrumental in the regulation of that response. In healthy subjects, during down-regulation of emotion, the functional connectivity between these areas has been observed to increase (7, 8).

The above-referenced functional magnetic resonance imaging (fMRI) studies of emotion regulation involve observation of brain activity while subjects perform a task. Recent studies that have evaluated subjects while not performing any cognitive tasks—e.g., resting state studies—have also revealed intrinsic activation patterns in the brain. Resting state fMRI provides complementary information to task-based fMRI in that both provide a platform for examining functional brain networks. Spatially distributed large-scale brain networks can be reliably derived and interrogated by either kind of fMRI experiment (9–11) and one could argue that both contribute different dimensions to a full characterization of brain activity. A variety of methods have been used to analyze such data (12) and have been applied to diverse clinical populations in the effort to delineate disease-related dysfunctions in connectivity (13).

While there are a handful of studies that have investigated functional connectivity in bipolar disorder, few are resting state studies (Table 1). Of these, there is some overlap with the regions used as seeds, such as the medial prefrontal cortex (14, 15) or amygdala (16, 17) but the findings show differences in connectivity in lateralization (left or right hemisphere), sign (positive or negative coupling), and between-group differences (healthy subjects > bipolar disorder or bipolar disorder > healthy subjects). Three of these studies investigated bipolar mania and/or depression (14, 15, 17); however, one combined several mood states into its bipolar disorder sample (16) while another combined bipolar I and II disorder subjects (14). This latter issue is particularly problematic as there is evidence to suggest that patterns of activation and connectivity change with different mood state and subtype of bipolar disorder (17–19). For a clearer picture of the underlying pathophysiology of bipolar disorder, further research that investigates common regions in both hemispheres in single mood states and in single bipolar disorder subtypes is necessary.

The goal of the present resting state study was to elucidate differences in intrinsic functional connectivity that might contribute to mood lability in bipolar I disorder. We therefore took a focused approach to assessing connectivity in two bilateral brain regions strongly implicated in emotion regulation (four regions in total: right and left amygdalae and right and left vlPFC) in a group of euthymic bipolar I disorder subjects. We hypothesized that subjects with bipolar I disorder would demonstrate aberrant functional connectivity relative to healthy subjects between amygdalae and lateral vlPFC—regions that subserve emotion regulation. Furthermore, observed alterations of brain connectivity during the euthymic state may be considered a potential biomarker of trait aspects of the disease.

Methods and materials

Participants

Participants provided written informed consent in accordance with the Institutional Review Boards at the University of California, Los Angeles (UCLA) and the Veterans Affairs (VA) Greater Los Angeles Healthcare System. Subjects with bipolar I disorder were recruited through the outpatient UCLA Mood Disorders Clinic, the outpatient Bipolar Disorders

Clinic of the VA Greater Los Angeles Health Care System, and through local advertising in the community. Healthy subjects were recruited by advertisements in local newspapers and campus fliers. All participants were first interviewed using the Structured Clinical Interview (SCID) (20) for DSM-IV to confirm a bipolar I disorder diagnosis or absence thereof. Participants with bipolar I disorder were excluded if they met criteria for any other current Axis I disorder. Healthy subjects were excluded if they had a current or past psychiatric diagnosis (including history of substance abuse) or were taking any medications for medical reasons. Additional exclusion criteria for all participants included left-handedness, neurological illness, metal implants, and a history of skull fracture or head trauma with loss of consciousness of more than five minutes.

In total, 21 bipolar I disorder subjects and 26 healthy subjects were scanned. Mood symptoms were evaluated in all participants on the day of the scan, using the Young Mania Rating Scale (YMRS) (21) and the 21-item Hamilton Depression Rating Scale (HDRS) (22) to assess for any symptoms of mania or depression. Participants were considered appropriate for this study if they had a YMRS score of < 7 (bipolar I disorder: 1.9 ± 1.9 ; healthy subjects: 0.74 ± 1.19 , $p = 0.03$), a 21-item HDRS score of < 7 (bipolar I disorder: 3.55 ± 2.5 ; healthy subjects: 0.79 ± 0.9 , $p < 0.001$), and had been euthymic by self-report and by SCID for at least one month prior to scanning.

Following the removal of seven subjects due to excessive head motion, 20 bipolar I disorder subjects (10 female, age 42.1 ± 11.4 years) and 20 (10 female, 39.8 ± 12.6 years) age- ($p = 0.55$) and gender-matched ($p = 1.0$) healthy subjects were included in the final analysis. Based on their age at first episode, they had been ill an average of 22.7 ± 11 years and currently euthymic an average of 34.5 ± 38.9 weeks. Length of formal education was 14.1 ± 1.9 years for bipolar I disorder subjects and 15.6 ± 2.1 for healthy subjects. Employment rate was 10% for bipolar I disorder subjects and 95% for healthy subjects. Three (15%) of these bipolar I disorder subjects were unmedicated and had been off medications for two months. The remaining 17 subjects were receiving a number of different medications including antipsychotics (15%), antidepressants (75%), and anticonvulsants (valproic acid: 25%, lamotrigine: 20%).

Image acquisition

Subjects were scanned on a 3T Siemens Trio scanner. They were asked to rest with their eyes closed during the scan but to not fall asleep. A structural T1 MPRAGE was acquired with parameters of repetition time (TR) = 1.9 sec, echo time (TE) = 2.26 msec, flip angle = 9° , matrix = 256×256 , field of view (FOV) = 250 mm, voxel size = 1 mm isotropic, and total sequence time 6 min and 50 sec. The resting state fMRI scan was acquired using a T2*-weighted echo planar imaging (EPI) gradient-echo pulse sequence with IPAT, with TR = 2 sec, TE = 25 msec, flip angle = 78° , matrix = 64×64 , FOV = 192 mm, in-plane voxel size = 3×3 mm, slice thickness = 3 mm, 0.75 mm gap, and 30 total interleaved slices. To allow for scanner equilibration, two TRs at the beginning of the scan were discarded. The total sequence time was 7 min and 2 sec, with 208 volumes acquired.

fMRI analysis

Following slice-timing correction of functional volumes, we performed motion realignment, gray matter/white matter/cerebral spinal fluid structural segmentation, normalization of both structural and functional scans to Montreal Neurological Institute (MNI) space, 3 mm isotropic voxel re-slicing, and 6 mm full-width at half-maximum (FWHM) Gaussian smoothing. To perform the statistical analysis, we used the CONN-fMRI functional connectivity toolbox, version 12.p (www.nitrc.org/projects/conn) (23), built on SPM8 (www.fil.ion.ucl.ac.uk/spm/).

Head movement can be a serious confound with resting state studies because it can introduce spurious correlations across the brain (11, 24). We took several precautions to ensure that it would not be a factor in our results. First, at a gross level, subjects were excluded if they had translational movement greater than one voxel (3 mm). Second, we tested whether the maximum head motion values (three translation and three rotation) were different between groups and we found no differences (all p -values > 0.05). Third, for each subject we calculated two different combinations of the six motion parameters that indicated total displacement (TD) along the scan or framewise displacement (FD) (i.e., scan-to-scan). Both metrics took into account each subject's individual head sizes to accurately assess the effects of rotation (25). We tested whether there were differences between groups for these metrics as well. TD did not differ between groups (bipolar I disorder: $\mu = 0.69 \pm 0.3$; healthy subjects: $\mu = 0.89 \pm 0.5$, $p = 0.14$) but FD did (bipolar I disorder: $\mu = 0.14 \pm 0.07$; healthy subjects: $\mu = 0.1 \pm 0.04$, $p = 0.01$). Therefore, subject-specific FD mean values were added as a separate covariate of non-interest during subsequent analysis and were found to exert no significant influence on the final reported results.

To account for movement and other noise confounds during the preprocessing of the functional images, and to avoid known issues with regressing out the global signal (26), the CONN-fMRI toolbox uses the anatomical component correction (aCompCor) method of flexibly removing physiological noise and movement confounds on a voxel-by-voxel basis (27, 28). The aCompCor approach enhances the sensitivity and specificity of positive correlations and can reveal non-artifactual anticorrelations. We used this method to regress out five principal components from the noise regions of interest (ROI) obtained from the anatomical segmentation as well as the six realignment parameters and their first temporal derivatives for each subject. The functional scans were further band-pass filtered between 0.01 and 0.1 Hz (100- to 10-sec cycles) to investigate low frequency correlations, which are most consistently produced within this range (11).

The CONN-fMRI toolbox implements a default set of ($n = 88$) Tailarach Daemon masks that may be augmented by additional ROIs. Thus, although the present study was mainly concerned with a focused set of connectivity pairings, the averaged time series from each of 88 cortical and two subcortical anatomical ROIs across the brain were extracted and bivariate Pearson correlations were calculated from each of these time courses at the voxel-level. The two additional subcortical ROIs added to the default Tailarach Daemon masks were the 75% probabilistic Harvard-Oxford's left and right amygdalae (Harvard Center for Morphometric Analysis).

We performed an analysis among ROIs (ROI-to-ROI) by calculating statistics for all possible connections for a selected subset of ROIs [right and left amygdalae, right and left Brodmann's Area (BA) 47]. This effectively performs a small volume correction on (Fischer-transformed) correlation values by setting the false discovery rate (FDR) correction to assure that only 5% or less of significant effects could be false positives. Results from this ROI-to-ROI analysis were then checked at the wholebrain level to confirm that they were not driven by a very small number of voxels (the ROI-to-ROI analysis works on the averaged time series of all voxels within the ROI). For this wholebrain (seed-to-voxel) analysis the amygdala was chosen as the seed region from the bi-directional right amygdala-vIPFC finding from the ROI-to-ROI analysis (see results below) to allow comparison with other studies that used the amygdala as a seed in connectivity analyses during rest (15, 16) and in studies of emotion regulation (8,29,30). The toolbox creates t -statistic volumes, which were brought into SPM8 for wholebrain investigations of within- and between-group maps. A voxel statistical height threshold of $p < 0.001$, $k = 10$ with a cluster height of $p < 0.05$ was used to identify significant correspondences between right amygdala wholebrain connectivity in the within- and between-group contrasts. These results were then tabulated

(Table 3), with anatomical and Brodmann labels automatically determined by the Talairach Client (www.talairach.org/client.html) after nonlinear coordinate conversion (www.bioimagesuite.org). These labels were then checked and revised against a probabilistic Anatomy toolbox (31) and an atlas in MNI space (32).

For confidence that findings in the tested model were not simply a reflection of global differences between the bipolar I disorder and healthy subject groups, we ran additional analyses between brain areas in which we did not expect to observe connectivity differences between groups. Specifically, we tested correlations of a number of primary sensory areas with the rest of the brain and compared groups.

Post-hoc associations between correlation values of right amygdala-vlPFC functional connectivity and patient course of illness variables (illness duration, number of depressive and manic episodes) and clinical variables (HDRS and YMRS scores) were assessed through correlations and two-tailed *t*-tests with right amygdala to right vlPFC beta values. These beta values were extracted from each bipolar I disorder subject's individual ROI correlation matrices. Prior to conducting these correlational analyses, an outlier correction was applied in which outlying connectivity parameter estimate values were winsorized to a value of 2.5 standard deviations from the group mean as in recent fMRI work (33). These correlations were performed in PASW version 18.0 (SPSS Inc; Chicago, IL, USA).

To explore regions outside our a priori model that may have driven our within-group bipolar functional connectivity results, a *connectivity conjunction map* of two different statistical maps was performed—the map of whole brain correlations with the right amygdala and the map of wholebrain correlations with right vlPFC—using the toolbox xjView (www.alivelearn.net/xjview8/). Strong correlations common to both connectivity maps outside our ROIs were identified by conjunction (i.e., a logical AND), with each map individually set at $p < 0.0001$, $k = 10$. A third brain region was identified in this manner and was included in a group-level mediation analysis, along with the amygdala and vlPFC nodes, to determine whether that region was responsible for a full or partial mediating effect on the main results. The mean times series from the third region were extracted from each of the bipolar I disorder subjects—preprocessed and filtered as the other ROIs—and were included as the mediator in a mixed-effects group-level mediation analysis. Regressions for the mediation analysis were performed in four steps (34) using the all-in-one model described by Beckmann and colleagues (35) which retains the variance of each subject throughout.

Results

Figure 1 (left image) shows the main ROI (ROI-to-ROI) model investigated, with the six labeled functional correlations between the four a priori nodes (left and right amygdalae and left and right vlPFC). Figure 1 (right image) shows the between group ROI-to-ROI positive connectivity result in graphical form. Only the connection 'IV' (right amygdala with right vlPFC) survived FDR-multiple comparison correction [$t(38) = 3.07$, FDR $p = 0.024$]. Adding head movement measurements TD and FD as covariates of non-interest left this result virtually unchanged [TD: $t(37) = 2.93$, FDR $p = 0.035$; FD: $t(37) = 2.74$, FDR $p = 0.028$]. Table 2 lists the within- and between-group results for this ROI-to-ROI analysis. The strength of within-group connectivity between all nodes survived FDR correction in both groups with the exception of weaker 'V' (right amygdala with left vlPFC) and 'VI' (left amygdala with right vlPFC) connections in healthy subjects ($p < 0.05$, FDR corrected). There were no wholebrain connectivity differences between bipolar disorder subjects and healthy subjects in primary somatosensory (BA 1), auditory (BA 41, 42), or visual (BA 17) regions.

As stated, wholebrain connectivity from the right amygdala seed was also assessed to confirm that the ROI-to-ROI results were not driven by a very small number of voxels, as is possible when averaged time series are extracted from an anatomical mask. Figure 2 (top and middle rows) shows within-group axial slices, while Figure 2 (bottom row) shows the between group contrast [circled cluster size =11 voxels, cluster $p < 0.05$; peak at MNI: ($x = 51, y = 41, z = -8$), $p < 0.0001, Z = 3.52$]. Consistent with the ROI-to-ROI result our between-group results (bipolar disorder > healthy subjects) confirmed significant hyperconnectivity between the right amygdala and right vIPFC, but also revealed significant regional hyperactivity between a number of other regions such as the right and left medial frontal gyri (BAs 10 and 11, respectively), the right superior temporal gyrus (BA 41), and the precentral gyrus (BA 6). Table 3 lists these within- and between-group right-amygdala-to-wholebrain connectivity results.

We found no significant correlations between right amygdala with right vIPFC connectivity and any of the five illness or clinical variables collected (illness duration, number of depressions, number of manias, and HDRS and YMRS scores). We also checked for medication effects on connectivity parameter estimates to make sure our few unmedicated bipolar I disorder subjects were not actually outliers driving our results.

Finally, we conducted a post-hoc test for possible *third* regions outside our model that may have driven, through mutual coupling, our right amygdala with right vIPFC result in the bipolar I disorder subjects. A significant cluster in the anterior cingulate cortex (ACC) [BA 32, center of gravity ($x = 1.9, y = 44, z = 0.63$)] was found to be strongly and positively functionally connected to both the right amygdala and the right vIPFC using the *connectivity conjunction map* procedure (Fig. 3A). We constructed our mediation model with the ACC mediating the influence of vIPFC to amygdala, which is what we hypothesized based on previous regression-based connectivity analyses of these regions (8, 29). A diagram and group-level betas for the analysis are shown in Figure 3B. We found the ACC to partially mediate the influence of vIPFC on the amygdala and this mediation effect was significant with the Sobel test ($Z = 7.88$).

Discussion

Bipolar disorder is associated with a trait dysregulation of the corticolimbic network (1, 2), which may in turn result in fluctuations of mood states over time. Here, we investigated the corticolimbic network using a focused, seed-based correlation analysis of resting state data in one mood state of one subtype of bipolar disorder. Our results demonstrate an altered intrinsic connectivity in euthymic bipolar I disorder subjects compared to healthy subjects between brain regions vital for emotion regulation. Specifically, we observed hyperconnectivity of the right hemisphere vIPFC (BA 47) and right amygdala in bipolar I disorder and its partial mediation by the ACC. Here *hyperconnectivity* refers to greater positive correlations in low-frequency fMRI signal in one group than a reference group (36), which in this case are healthy subjects. Our within-group maps clearly support this difference in magnitude; subjects with bipolar I disorder have greater positive connectivity between these two regions in comparison to healthy subjects (Fig. 2 top and middle rows). Additionally, in subjects with bipolar I disorder, we found this resting hyperconnectivity elsewhere in the brain between amygdala and, for example, the ACC and the middle frontal gyrus (BAs 10 and 11) (see also Table 3).

One interpretation of our hyperconnectivity result is that functional coupling between vIPFC and may be especially sensitized in bipolar disorder and that this may contribute to or reflect the characteristic mood lability of the disease. Whether this sensitization stems from differences in right hemisphere white matter structural connections is unclear and warrants

further research (37). The lateralization of differences of brain coupling in bipolar disorder also warrants further exploration and replication in general, however what we report here is consistent with prior neuroimaging research documenting right lateralization for emotion-processing effects (38).

The present findings are also consistent with four other studies in bipolar disorder subjects that have shown increases in seed-based connectivity between or involving at least one of these regions. For example, hyperconnectivity, as measured by mutual information, has been observed in bipolar disorder between ventromedial PFC and right amygdala while labeling sad facial expressions (39). During a psychophysiological interaction, hyperconnectivity was observed in bipolar disorder subjects between right vIPFC and insula (40), and during rest hyperconnectivity was observed during mania between medial and vIPFC (15) as well as between left and right ventral PFC in a variety of mood states (16) (c.f. Table 1.)

We also found that an area of the ACC (BA 32) was significantly coupled with both right amygdala and right vIPFC. The ACC is known to have anatomical and functionally reciprocal connections with the amygdala (41, 42), has been strongly implicated in mood disorders (6), and has been found by one group to mediate the coupling between right vIPFC and amygdala during incidental emotion regulation (3). The functional connectivity between this region and the amygdala has been found by another group to be decreased in bipolar disorder relative to healthy subjects during the processing of fearful and happy faces (43). Therefore, another way we could interpret our results is that the ACC in bipolar I disorder is dysfunctionally mediating the relationship of the vIPFC and amygdala to determine the hyperconnectivity we observed. It is interesting that Chai et al (15) also found hyperconnectivity during rest between the vIPFC and this area in manic bipolar disorder subjects; however, not between this area and the amygdala, as we did. This suggests this connection could be dependent on mood state, and other research that shows the coupling between ACC and amygdala tracking with anxiety (44) supports this. Further investigation is necessary to elucidate the resting effective connections of the ACC.

There are several limitations to the current study. First, the volumes of the anatomical ROIs we used for this seed-based functional connectivity analysis, especially the vIPFC, were based on an atlas and as such will contain certain errors. If, for example, the time-series extracted from an atlas-derived brain region averages over both gray and non-gray matter signals, the resulting series will not properly represent that region's activation, which can result in unreliable measures. However, the aCompCor algorithm, which incorporates a high-resolution, subject-specific anatomic segmentation into its noise reduction procedure, takes such variability into account and attempts to compensate by higher weighting of gray matter voxels, thereby minimizing the effect of this issue [Nieto Castañón, personal communication; (also see 28)]. Nonetheless, there is likely some degree of functional heterogeneity within both the vIPFC and amygdala atlas ROIs used, which would add variability to the extracted time series. The effect of this, however, could likely dilute our results. Thus, our observed significant effects would likely only become more significant if this limitation were corrected.

A second limitation, characteristic of any bivariate correlation, is that significant couplings between two brain regions could be the result of their being driven by a third. We explored this possibility and found that the ACC is a region potentially driving the apparent coupling between right hemisphere amygdala and vIPFC (see results above and Fig. 3A). We investigated this further and found partial mediation of the ACC with the vIPFC to amygdala connection. Despite our standard *arrowed* diagram (Fig. 3B) the causality in such a mediation should be interpreted cautiously, as it represents 'soft' causal links that stem from the 'DV'/IV' regression relation. The mediation analysis does, however, provide a more

mechanistic description of the hyperconnectivity between amygdala and vIPFC. Future studies that systematically expand our four-node model and assess effective connectivity between regions are needed to further elucidate the causal relationship of functional pathology of bipolar disorder (45). Such work is ongoing in our laboratory.

Certain characteristics of our sampled subjects are also limitations. First, many of our bipolar I disorder subjects were medicated. There were too few subjects in each medication group to investigate the effects of specific medications, but this is something future work should address. We did, however, examine the correlation beta values of the three unmedicated subjects in our bipolar I disorder group to make sure their connectivity estimates were not driving the results and found them to be well within one standard deviation of the group mean. Despite the potential, but inconclusive, impact of medications on blood oxygen level-dependent (BOLD) signal in subjects with bipolar disorder (46), we argue that including medicated subjects ensures that our subject population is representative of the bipolar disorder population at large, thereby increasing the generalizability of our results. Second, there was a significant difference in employment between the two groups: the vast majority of bipolar I disorder subjects were unemployed while the majority of healthy subjects were employed. This is not something commonly controlled for in bipolar disorder studies, although future studies would benefit from taking a closer look at this issue. We note that unemployment is often correlated to bipolar disorder severity (47), but the relationship of the variable to resting state functional coupling has, to our knowledge, not been studied.

Finally, and not so much a limitation to our particular study as a reminder to exert interpretive caution with resting state studies, the current state of MRI-based imaging does not yet have a clear explanation of the relationship between task-based and resting state functional brain activity. The extent to which findings from experimental manipulations of cognitive states should and can map to resting states studies is still being actively investigated (10, 11). Many resting state studies have focused on a small set of reliably produced large-scale networks across the brain, and yet, for two decades the majority of cognitive neuroscientists have studied the activity and connectivity in a variety of different networks specific to cognitive processes and which have not been reliably uncovered within resting state analyses. One likely explanation for this discrepancy is that analyses of resting state data are conducted on a much lower and usually non-overlapping end of the fMRI signal frequency spectrum than standard task-based studies (11, 48). Knowledge coming from advances by those researchers actively investigating these important relationships will, no doubt, inform the types of studies, like this one, performed by clinical neuroscientists.

In summary, we took a focused approach to exploring resting state brain patterns in subjects with bipolar disorder during the euthymic mood state. Specifically, we investigated intrinsic connectivity between four ROIs that have been implicated in emotion regulation and have been previously shown to be functionally altered in bipolar disorder. Our results of hyperconnectivity between the right vIPFC and the right amygdala in bipolar I disorder subjects are in accordance with the few prior studies of bipolar disorder that have found either hyperactivation or hyperconnectivity in these brain regions. We further observed a mediating effect from the ACC, which was functionally coupled to both amygdala and vIPFC. Together, these findings suggest a maladaptive misallocation of neural resources in bipolar disorder. Intrinsic hyperconnectivity in the bipolar I disorder group between regions, in the 'normal' euthymic state and while at rest, could be evidence of a trait marker of disease pathophysiology.

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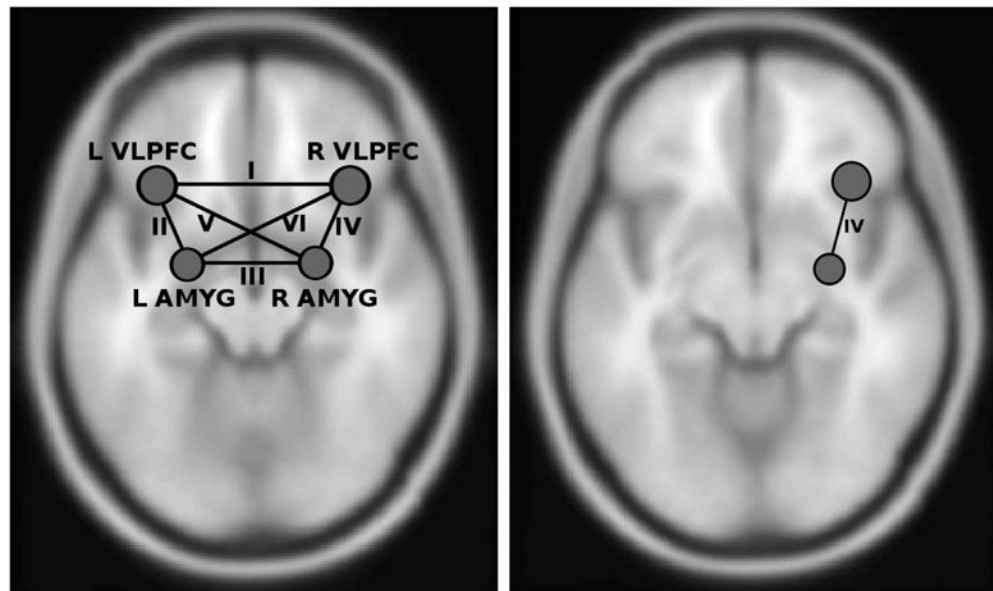


Fig. 1. The four node (six correlation) connectivity model that was tested (**left image**). Node locations at center of mass of anatomical regions of interest. Results of greater positive connectivity between right amygdala (AMYG) and right ventrolateral prefrontal cortex (vLPFC) in bipolar I disorder subjects relative to healthy subjects (**right image**). L = left; R = right.

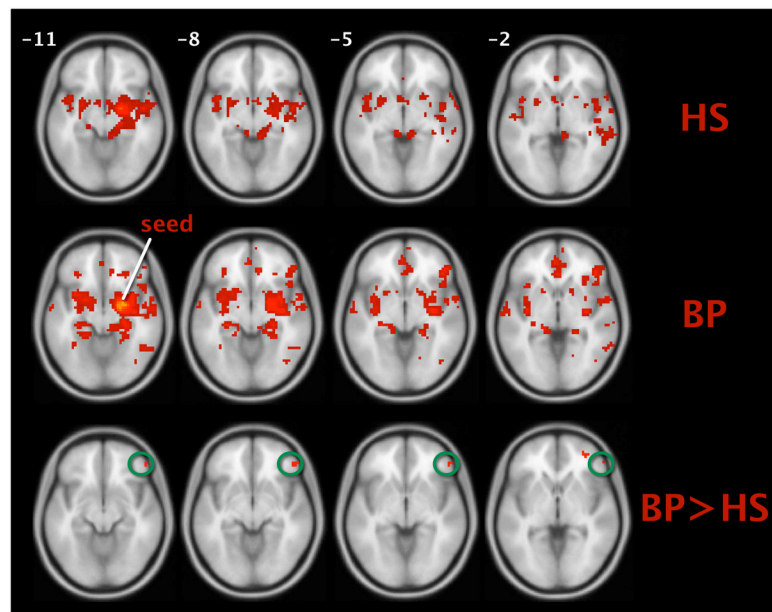


Fig. 2. Positive connectivity with right amygdala. **(Top row)** and **(middle row)**: within-group maps for healthy subjects (HS) and bipolar I disorder subjects (BP). **(Bottom row)**: between group contrast. Right ventrolateral prefrontal cortex (vIPFC) cluster circled. Voxels visualized on an average T1-weighted brain in Montreal Neurological Institute (MNI) space, $p < 0.001$, $k = 10$.

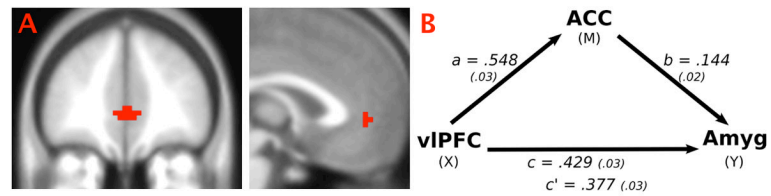


Fig. 3. (A) Connectivity conjunction map shows a cluster in the anterior cingulate cortex (ACC) (BA 32) that is strongly positively functionally connected to both the right amygdala (Amyg) and right ventrolateral prefrontal cortex (vIPFC) regions in bipolar I disorder subjects. (B) Group-level analysis of ACC mediating vIPFC to Amyg. Path coefficients labeled with traditional mediation nomenclature and standard errors in parentheses. All paths were significant ($p < 0.0001$), indicating partial mediation of vIPFC to Amyg by ACC.

Table 1
A selection of functional magnetic resonance imaging (fMRI) connectivity studies in bipolar disorder

Study	Task/rest	BP moodstate	Method	Seed/ROI	BP > HS	HS > BP
Foland-Rosset al. 2008 (29)	Labeling emotional faces	Manic	PPI	L AMYG	Negative FC: ACC (BA32 and 24) and BA10	Negative FC: LH amygdala—bilateral vPFC
Pompeiet al. 2011 (40)	Color-word Stroop	Euthymic	PPI	R/vPFC	Positive FC: RH vPFC—insula	Negative FC: RH vPFC—LH vACC
Townsend et al. 2012 (30)	Explicit emotion regulation	Euthymic	PPI	L AMYG		Negative FC: L AMYG—L vPFC
Versace et al. 2010 (39)	Labeling sad and happy faces	Depressed and euthymic	Seed-based FC	omPFC (BA 11) and AMYG	Sad: RH AMYG—vmOFC	Happy: LH AMYG—vmOFC
Anand et al. 2009 (14)	Rest	Manic and depressed	Seed-based FC	pgACC		LH and RH pgACC—dmTHAL, AMYG, and striatum
Chai et al. 2011 (15)	Rest	Manic	Seed-based FC	mPFC, AMYG, dlPFC, insula, vPFC	mPFC—LH insula mPFC—RH BA47	
Chen et al. 2010 (16)	Rest	Rapid cyclers, euthymic, depressed, and mixed	Seed-based FC	vPFC, AMYG	LH vPFC—RH vPFC	Negative FC: LH AMYG—vPFC
Cerullo et al. 2012 (17)	Steady state	Manic and depressed (longitudinal)	Seed-based FC	L and RAMYG	N/A	N/A

BP = bipolar disorder; ROI = region of interest; HS = healthy subjects; PPI = psychophysiological interaction; FC = functional connectivity; L = left; R = right; AMYG = amygdala; PFC = prefrontal cortex; vPFC = ventrolateral PFC; omPFC = orbitomedial PFC; BA = Brodmann's Area; pgACC = pregenual ACC; mPFC = medial PFC; dlPFC = dorsolateral PFC; ACC = anterior cingulate cortex; RH = right hemisphere; LH = left hemisphere; — = correlation (coupling); vACC = ventral ACC; vmOFC = ventromedial orbitofrontal cortex; vPFC = ventral PFC; dmTHAL = dorsomedial thalamus; N/A = these direct contrasts were not performed.

Table 2

Within- and between-group ROI-to-ROI analysis results with connections labeled as in Figure 1 (left image)

Connection	Healthy subjects						Bipolareuthymic						BP>HS	
	I	II	III	IV	V	VI	I	II	III	IV	V	VI	IV	VI
Betas ^a	0.52	0.11	0.31	0.07	0.04	0.04	0.51	0.12	0.26	0.2	0.14	0.1	0.12	0.12
t-scores	10.74	3.19	6.76	2.71	1.67	1.08	11.81	4.28	5.71	6.89	3.89	3.66	3.07	3.07
Z-scores	7.24	2.98	5.44	2.58	1.63	1.07	7.61	3.84	4.82	5.52	3.55	3.37	2.88	2.88
FDR p-values	0.0000	0.0096	0.0000	0.0207	0.1335	0.2947	0.0000	0.0006	0.0000	0.0000	0.0012	0.0017	0.0237	0.0237

ROI = region of interest; BP = bipolar disorder; HS = healthy subjects; FDR = false discovery rate.

^aFisher-transformed correlation coefficients.

Align values by the decimal point.

Table 3

Within- and between-group maps of voxel-level functional connectivity from right amygdala in healthy and bipolar I disorder subjects^a

	Brodmann's Area	Cluster size	Cluster p-value	Peak Z	MNI		
					x	y	z
Healthy subjects right amygdala connectivity							
<i>Positive</i>							
R Amygdala		2391	0.000	Inf	21	-4	-17
R Temporal pole	BA 38			0.000	45	23	-26
L Superior temporal gyrus				0.000	-54	-16	7
L Medial temporal pole				0.000	-33	8	-32
L Insula				0.000	-42	8	-8
R Parahippocampal gyrus	BA 36			0.000	39	-19	-32
L Hippocampus				0.000	-27	-22	-17
L Superior temporal gyrus	BA 13			0.000	-45	5	-14
L Superior temporal gyrus				0.000	-39	2	-20
R Postcentral gyrus	BA 3			0.000	63	-1	22
R Superior temporal gyrus	BA 22			0.000	48	-19	1
L Middle temporal gyrus	BA22	12	0.027	4.56	-66	-55	10
L Middle temporal gyrus	BA 39	10	0.041	3.51	-60	-64	10
Anterior cingulate gyrus		10	0.041	3.57	0	35	1
R Insula	BA 13	53	0.000	4.51	36	-13	19
R Thalamus		16	0.013	3.66	3	-10	7
<i>Negative</i>							
L Middle frontal gyrus	BA 10	70	0.000	4.72	-24	47	25
L Inferior parietal lobule	BA 40	33	0.001	4.6	-48	-52	40
L Precuneus	BA 7	14	0.019	4.07	-9	-64	49
L Superior parietal lobule	BA 7	29	0.002	3.85	-39	-49	61
L Middle orbital gyrus		16	0.013	3.5	-42	56	-11
R Precuneus	BA 7	18	0.009	4.28	6	-70	58
R Middle frontal gyrus		18	0.009	3.84	33	50	13

	Brodmann's Area	Cluster size	Cluster p-value	Peak Z	MINI		
					x	y	z
Bipolar euthymic right amygdala connectivity							
<i>Positive</i>							
R Amygdala		2020	0.000	Inf	27	-4	-20
R Inferior temporal gyrus				5.2	51	-55	-17
R Hippocampus				5.18	33	-13	-35
R Parahippocampal gyrus				5.14	24	-25	-14
R Inferior frontal gyrus	BA 45			5.04	54	32	10
R Ventrolateral prefrontal cortex	BA 45			4.93	45	35	-2
R Fusiform gyrus	BA 36			4.8	24	-40	-17
R Middle temporal gyrus				4.77	60	-7	-14
R Inferior frontal gyrus	BA 47			4.66	48	38	-5
R Temporal pole	BA 38			4.3	42	20	-38
R Thalamus (somatosensory)				4.18	21	-19	4
R Superior temporal gyrus	BA 41	67	0.000	4.58	66	-22	4
R Posterior insula	BA 13	58	0.000	4.33	42	-28	22
R Angular gyrus	BA 39	32	0.001	4.2	54	-67	34
R Posterior cingulate	BA 30	32	0.001	4.12	15	-58	7
R Insula	BA 13	17	0.011	4.08	39	5	10
R Superior temporal gyrus		95	0.000	4.03	54	-46	19
R Middle frontal gyrus	BA 11	42	0.000	4.01	18	29	-14
R Precuneus	BA 19	13	0.022	3.99	36	-85	37
R Precentral gyrus	BA 6	15	0.015	3.99	51	2	37
R Precentral gyrus	BA 6	14	0.018	3.84	54	-7	31
R Middle occipital gyrus		16	0.013	3.81	36	-73	25
R Middle frontal gyrus		12	0.027	3.66	45	20	28
R Inferior temporal gyrus		10	0.041	3.65	45	-70	-5
L fusiform gyrus	BA 37	40	0.000	4.22	-42	-49	-26
L Middle occipital gyrus	BA 19	29	0.002	4.2	-42	-85	25

	Brodmann's Area	Cluster size	Cluster p-value	Peak Z	MNI		
					x	y	z
L Rolandic operculum		13	0.022	4.03	-51	-4	7
L Inferior frontal gyrus		36	0.001	3.82	-30	35	-8
L Posterior cingulate		11	0.033	3.64	-18	-64	13
L Medial frontal gyrus	BA 10	10	0.041	3.59	-6	62	16
Cingulate gyrus	BA 24	50	0.000	5.29	0	2	34
BP > HS right amygdala connectivity							
<i>Positive</i>							
L Middle frontal gyrus	BA 10	16	0.021	3.5	-30	62	13
R Middle frontal gyrus	BA 9	14	0.029	4.3	45	20	40
R Inferior temporal gyrus	BA 20	11	0.049	3.58	33	-4	-47
R Ventral lateral prefrontal cortex	BA 47	11	0.049	3.52	51	41	-8

MNI = Montreal Neurological Institute; R = right; L = left; BP = bipolar disorder; HS = healthy subjects.

⁴There were no significant negative connectivity results with bipolar I disorder subjects. Rows in gray represent subclusters within a large cluster. Within-and between-group maps thresholded at $p < 0.001$, $k = 10$.