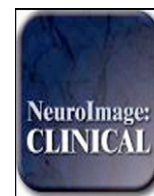Contents lists available at ScienceDirect

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Shared white matter alterations across emotional disorders: A voxel-based meta-analysis of fractional anisotropy

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ABSTRACT

Background: White matter (WM) integrity may represent a shared biomarker for emotional disorders (ED). Aims: To identify transdiagnostic biomarkers of reduced WM by meta-analysis of findings across multiple EDs. *Method:* Web of Science was searched systematically for studies of whole brain analysis of fractional anisotropy (FA) in adults with major depressive disorder, bipolar disorder, social anxiety disorder, obsessive-compulsive disorder or posttraumatic stress disorder compared with a healthy control (HC) group. Peak MNI coordinates were extracted from 37 studies of voxel-based analysis (892 HC and 962 with ED) and meta-analyzed using seed-based *d* Mapping (SDM) Version 4.31. Separate meta-analyses were also conducted for each disorder. *Results:* In the transdiagnostic meta-analysis, reduced FA was identified in ED studies compared to HCs in the left inferior fronto-occipital fasciculus, forceps minor, uncinate fasciculus, anterior thalamic radiation, superior corona radiata, bilateral superior longitudinal fasciculi, and cerebellum. Disorder-specific meta-analyses revealed the OCD group had the most similarities in reduced FA to other EDs, with every cluster of reduced FA overlapping with at least one other diagnosis. The PTSD group was the most distinct, with no clusters of reduced FA overlapping with any other diagnosis. The BD group were the only disorder to show increased FA in any region, and showed a more bilateral pattern of WM changes, compared to the other groups which tended to demonstrate a left lateralized pattern of FA reductions. *Conclusions:* Distinct diagnostic categories of ED show commonalities in WM tracts with reduced FA when compared to HC, which links brain networks involved in cognitive and affective processing. This meta-analysis facilitates an increased understanding of the biological markers that are shared by these ED.

1. Introduction

White matter (WM) alterations have been identified in numerous emotional disorders (ED), including mood and anxiety disorders (AD) (Ayling et al., 2012; Nortje et al., 2013; Murphy and Frodl, 2011). These alterations are important because disruption to WM pathways could reveal a disconnection syndrome within neural circuits (Schmahmann et al., 2008; Thomason and Thompson, 2011), potentially modifying behavior and cognitive control of emotion. Although EDs have distinct mechanisms, they share many phenotypic (Clark and Watson, 1991) and genotypic commonalities (Kendler et al., 2003). Furthermore, they are often comorbid with one another (Brown et al., 2001; Kessler et al., 2005; Yerevanian et al., 2001). For example, 50–60% of individuals with major depressive disorder (MDD) have a comorbid anxiety disorder (Fava et al., 2000; Kessler et al., 1996), such as

social anxiety disorder (SAD), panic disorder (PD), generalized anxiety disorder (GAD), obsessive-compulsive disorder (OCD) or posttraumatic stress disorder (PTSD). There is a high rate of comorbidity of these AD in bipolar disorders (BD) (Freeman et al., 2002; Henry et al., 2003; Pini et al., 1997; McElroy et al., 2001). Furthermore, the ADs are also highly comorbid within themselves (Goisman et al., 1995; Kessler et al., 1995; Pietrzrak et al., 2011). This study therefore sought to conduct a meta-analysis of studies of WM in mood and AD (hereafter emotion disorders, ED), to determine which WM alterations are common or distinct to these classes of psychopathologies. Given the call for transdiagnostic mechanisms put forth by NIMH's Research Domain Criteria (RDoC) initiative (Insel et al., 2010), it is particularly timely to identify shared neural circuitry abnormalities across multiple classes of EDs.

Numerous structural models of psychopathology have reliably reported commonalities among MDD and ADs, with both loading on a broad 'internalizing disorders' factor (a factor which appears to be distinct from 'externalizing' conditions such as substance use disorders (Vollebergh et al., 2001)). It is somewhat unclear where BD falls in this hierarchy, with some studies showing that it loads on a separate

factor from MDD and ADs (Watson, 2005; Kotov et al., 2015). Given the high prevalence of comorbid externalizing disorders such as substance abuse in BD (Cassidy et al., 2001), it is possible that BD includes both internalizing and externalizing features. Furthermore, although most studies of WM abnormalities in BD have reported decreases in integrity similar to studies of MDD, increased measures of integrity in the left uncinate fasciculus and optic radiation have been reported (Versace et al., 2008). By aggregating data from multiple studies of ED, we hypothesize that this meta-analysis will identify WM alterations that are shared across ED. This knowledge will provide important insights into potential mechanisms that are shared across these classes of disorders. To our knowledge, no previous meta-analysis has attempted to examine WM abnormalities compared to healthy controls across MDD, BD and ADs. Therefore the results of this study, which followed the guidelines for Meta-analyses of Observational Studies in Epidemiology, will be important for identifying transdiagnostic biomarkers of WM pathology in ED.

Diffusion tensor imaging (DTI) measures the degree to which randomly diffusing water molecules move in one direction, rather than all directions of tissues (Thomason and Thompson, 2011), providing an indication of the orientation and integrity of WM fibers (Beaulieu, 2002; Kumar et al., 2015). The diffusion tensor is estimated at each voxel, and eigenvalues are calculated which indicate the extent of diffusion. Fractional anisotropy (FA) is a ratio of eigenvalues ranging from 0 (isotropic) to 1 (unidirectional, anisotropic movement) that reflects the degree to which diffusion is confined to a particular direction (Kumar et al., 2015). FA usually decreases in damaged, disorganized or atrophied WM (Thomason and Thompson, 2011), however it is only an indirect marker of WM microstructure (see discussion). Nevertheless, given it is the most widely reported DTI measure, FA was selected as the variable of interest for the present meta-analysis.

2. Materials and methods

2.1. Study selection

The Web of Science database was systematically searched by a post-doctoral researcher (LMJ) on the 16th of September 2014 using search terms ("DTI" OR "diffusion tensor") AND ("anxiety" OR "depress*" OR "bipolar" OR "mania"). Reference lists were also manually searched for suitable papers, resulting in 695 results. A subsequent editorial decision to add OCD and PTSD resulted in a new Web of Science search on the 12th of April 2016, and yielded 228 results. This search included the terms ("OCD" OR "obsessive compulsive" OR "PTSD" OR "post traumatic stress disorder") AND ("DTI" OR "diffusion tensor" OR "fractional anisotropy"). In total, 923 studies were identified by these two systematic searches (Fig. 1). Borderline personality disorder was considered for inclusion in the meta-analysis since this diagnosis also involves emotion dysregulation, however we were unable to locate any studies that met our inclusion criteria. Specifically, we conducted a literature search in Web of Science on April 5th, 2016, using the search terms "Borderline personality" and "DTI or diffusion tensor or fractional anisotropy". This search yielded 25 results. Of these 25 results, none met our inclusion/exclusion criteria for entry into the study. There were 6 reviews, 3 abstracts, 2 studies that did not involve DTI, 4 that did not involve Borderline personality disorder, 5 were not whole-brain studies, 2 that only included adolescents. The remaining 3 studies involved TBSS. Since the main analysis used VBA and the TBSS analysis was used to support the findings of the VBA analysis, we decided not to include borderline personality disorder. As this was a meta-analysis of de-identified published results, institutional ethical approval was not required for this study.

Criteria for inclusion were: Whole brain (using voxel-based analysis, VBA) or whole WM (using tract-based spatial statistics, TBSS) analysis of FA in human participants with either MDD, BD or AD, reported in standardized 3D space, constituting original data from participants aged 18–65 years, compared with a healthy control (HC) group,
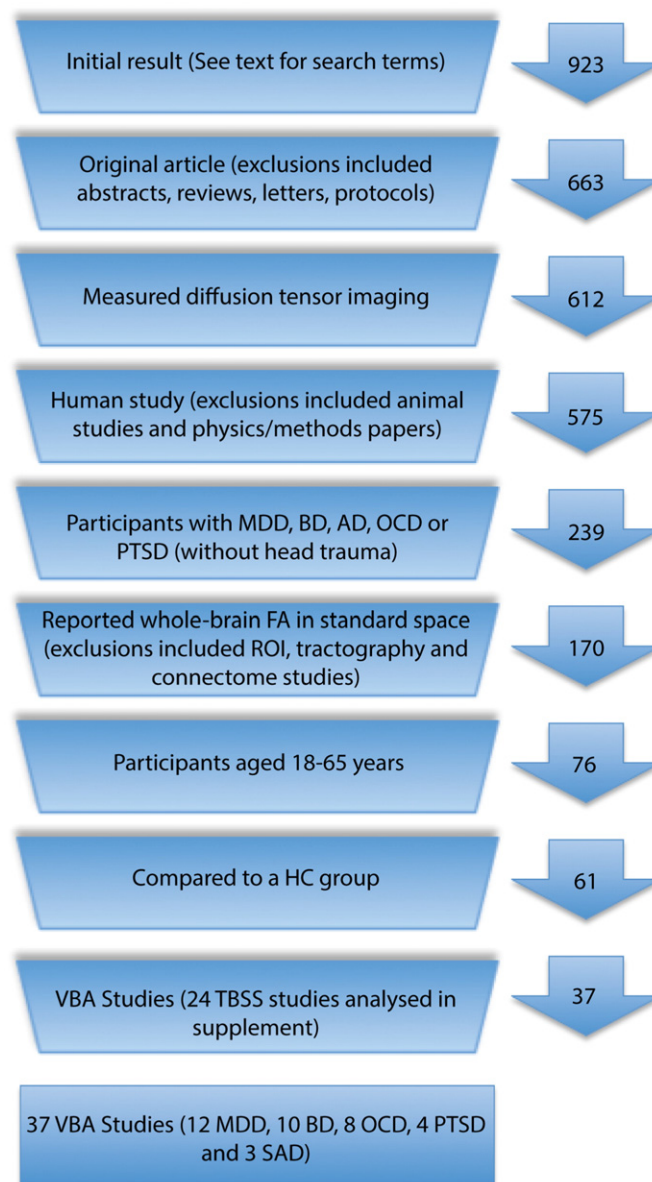

Fig. 1. Flow chart of exclusion of studies that were identified in the literature search.

reported in English in a peer-reviewed journal. Given the common comorbidity of PTSD with Traumatic Brain Injury (TBI), studies of PTSD that did not exclude patients with TBI or provide separate results for a non-TBI sub group were excluded. Fig. 1 shows a flowchart of the selection of studies for inclusion. For the purposes of this meta-analysis, AD were limited to generalized anxiety disorder (GAD), social anxiety disorder (SAD), panic disorder (PD), obsessive compulsive disorder (OCD) or post traumatic stress disorder (PTSD). Our literature search identified only one study of GAD that met our inclusion criteria, so GAD was excluded, since one study is not sufficient to establish within-disorder reliability. All studies of PD used TBSS, which was analyzed separately, thus the meta-analysis combined two classes of mood disorder (BD and MDD), and three classes of AD (SAD, OCD and PTSD) to examine transdiagnostic WM alterations for EDs.

The meta-analysis was conducted using seed-based *d* Mapping (SDM) Version 4.31 (described below). Unlike some other methods, SDM incorporates both positive and negative peaks, and null studies, thus whilst the majority of studies of FA in ED report reduced FA compared to HCs, we were able to include studies in the meta-analyses that reported null results or increased FA. SDM cannot simply or directly

Table 1
Characteristics of VBA studies included in meta-analysis.

Paper	Patients male/female	HCs male/female	Contrast	Illness length (months)	Patient age mean (SD)	HC age mean (SD)	Meds	HRSD score mean (SD) # items	Dx tool	Excluded psychiatric comorbidities	B₀	Vox. size (mm)	Directions	Highest b (s/mm²)	p
MDD															
Arnold et al. (2012)	4 M/13 F	7 M/14 F	rMDD vs HC	N/A	30.41 (11.35)	26.90 (7.82)	No	9.18 (6.82) 17 item	SCID-I mood sect.	Other psychiatric Dx as per the MINI, SUD	1.5	2.5 × 2.5 × 2.5	30	900	Cluster differences of $p < 0.001$, uncorrected
Blood et al. (2010)	10 M/12 F	10 M/12 F	MDD vs HC	N/A	36.3 (12.1)	35.3 (11.6)	Yes	NA, 31 items	SCID	Psychotic, BD, eating, OCD, SUD, GAD, PD, PTSD	3	2 × 2 × 2	6	700	$p < 0.00005$, whole-brain Bonferroni corrected
Choi et al. (2014)	64 M/70 F	28 M/26 F	MDD vs HC	112.08	38.49 (11.09)	34.42 (10.06)	Med. free	19.28 (3.47) – 17 item	SCID	Psychotic, BD, OCD, SUD	3	2 × 2 × 2	60	1000	Permutation based TFCE, and FWE correction ($p < 0.05$, 10,000 permutations) used with estimation of cluster size
Jia et al. (2010)	Suicide attempt = 5 M/11 F; non-attempt = 20 M/16 F	24 M/28 F	1. suicide attempt vs HC; 2. non-attempt vs HC	attempt = 80.00; non-attempt = 21.00	attempt = 34.2 (13.7); non-attempt = 34.7 (12.5)	37.1 (16.0)	Med.-free for at least 2 weeks prior to study	attempt = 24.6 (3.8); non-attempt = 22.3 (4.3)- 17 item	SCID	Any DSM-IV axis I comorbidities, SUD	3	1.9 × 1.9 × 3	15	1000	Voxel-level $p < 0.05$ (FWE corrected), extent threshold $p < 0.05$ (uncorrected), k ≥ 50
Ma et al. (2007)	2 M/12 F	2 M/12 F	1st ep. MDD vs HC	10.30	28.9 (8.0)	27.1 (6.7)	Med. naïve	N/A (BDI = 37.5; 6.9)	SCID	"Any lifetime psychiatric disorder"	1.5	1.9 × 1.9 × 4	13	1000	Voxelwise $p < 0.001$ (uncorrected) & k ≥ 50
Osoba et al. (2013)	8 M/12 F	8 M/12 F	MDD vs HC	N/A	38.3 (11.6)	33.8 (7.2)	All patient antidep	15.1 (6.1) – 21 item	ICD-10	Major medical Dx, seizures, ECT, psychiatric Dx, SUD	3	2 × 2 × 2	12	1000	Voxelwise $p < 0.05$ (corrected) & whole brain αSim correction height $p < 0.001$ (uncorrected) & k ≥ 256
Ouyang et al. (2011)	9 M/9 F	9 M/9 F	1st ep. MDD vs HC	15.10	27.4 (6.4)	27.0 (6.8)	Naïve	24.2 (4.3) 17 item	SCID	Substance abuse Hx	1.5	1.9 × 1.9 × 4	13	1000	$p < 0.001$ (uncorrected), k ≥ 30
Peng et al. (2013)	19 M/11 F	14 M/11 F	TRD vs HC	56.16	26.77 (5.28)	28.24 (4.98)	N/A, resistant to 2+ classes of antidep > 4 weeks at max. dose	N/A BDI: 20.47 (4.45)	DSM-IV	DSM axis I or II Dx	3	1.3 × 1.3 × 3	30	1000	Voxelwise $p < 0.05$ (FDR corrected) & k ≥ 50, cluster level $p < 0.05$ (FDR corrected)
Tha et al. (2013)	12 M/7 F	13 M/6 F	MDD vs HC	18.37	38.6 (13.5)	36.5 (12.5)	Antidep free >6 months	19.0 (4.0) 17 item	DSM-IV-TR and MINI	DSM axis I or II Dx	1.5	1.9 × 1.9 × 5	12	1000	$p < 0.001$ (uncorrected), k ≥ 50
Wang et al. (2013)	5 M/16 F	8 M/14 F	MDD pre-treatment vs HC	14.20	29.6 (12.6)	30.2 (10.2)	Yes	20.4 (6.3) -17 item	DSM-IV	Other psychiatric Dx or Hx of symptoms	3	1 × 1 × 5	14	1000	$p < 0.001$, k ≥ 50, & αSim correction at $p < 0.05$ (height threshold $p < 0.001$, k ≥ 23)
Wu et al. (2011)	10 M/13 F	9 M/12 F	1st ep., treatment-naïve MDD vs HC	2.17	31.4 (8.8)	30.4 (8.2)	Naïve	21.8 (3.8) 17 item	SCID	DSM axis I or II Dx	1.5	1 × 1 × 4	13	1000	$p < 0.001$ (uncorrected), k ≥ 30

Study	Patients (M/F)	Controls (M/F)	Comparison	Illness duration	Age patients	Age controls	Medication	Symptom severity	Diagnostic	Comorbidity/exclusion	Tesla	Voxel size	Smoothing	N voxels	Threshold
Zou et al. (2008)	15 M/30 F	15 M/30 F	MDD vs HC	20.00	33.2 (8.9)	31.0 (10.3)	Yes	23.8 (3.3) 17 item	DSM-IV	Major psychiatric Dx including schizophrenia, BD, affective disorder, personality disorder, substance abuse, SUD	3	1.9 × 1.9 × 3	15	1000	Cluster threshold p < 0.001 (uncorrected), p < 0.05 at voxel-level after FDR correction
BD															
Bruno et al. (2008)	13 M/23 F	Total N = 28, # M/F not reported	BD (I&II) vs HC	165.60	39 (no SD)	N/A other than matchBD group (range 21–63)	Yes	N/A	SCID	"Comorbid psychiatric conditions"	1.5	2.5 × 2.5 × 5	7	4 values between 0 and 700	p < 0.05 at cluster level after FWE correction for multiple comparisons
Chaddock et al. (2009)	9 M/10 F	10 M/8 F	Psychotic BD I in remission vs HC	N/A	43.3 (10.2)	41.7 (12.2)	Yes	N/A but BDI mean = 7.9 (7.0)	SADS-LV	SUD	1.5	2.5 × 2.5 × 2.5	64	1300	Voxelwise p < 0.05, the permutation tested with expected number of false positives < 1 per analysis
Chen et al. (2012)	18 M/0 F	27 M/0 F	BD (I & II 1st ep. Mania) vs HC	Median 50.40	32.0 (7.6)	31.3 (6.8)	Yes but not > 5 days prior	3.2 (1.1) – 17 item	SCID	Axis I Dx, Hx of drug or alcohol abuse	3	1.9 × 1.9 × 3	15	1000	p < 0.05 (FDR corrected) and k ≥ 50
Cui et al. (2011)	10 M/8 F	18 M/12 F	BD I (current manic ep.) vs HC	57.60	27.9 (5.7)	23.9 (10.2)	Yes	3.1 (2.2)	SCID	Hx substance abuse	3	1.9 × 1.9 × 3	15	1000	Voxelwise p < 0.001 (uncorrected) and k ≥ 50
Ha et al. (2011)	BD I = 3 M/9 F; BD II = 2 M/10 F	5 M/17 F	1. BD I vs HC; 2. BD II vs HC	BD I = 159.60; BD II = 159.60	BD I = 37.3 (10.59); BD II = 35.6 (7.56)	34.7 (7.12)	Yes	BP I = 5.5 (6.9); BP II = 4.2 (4.43) - 17 item	SCID	Hx substance abuse, current "severe mood episode"	1.5	2 × 2 × 3	15	600	p < 0.001 (uncorrected), k ≥ 40
Liu et al. (2010)	BD I = 7 M/7 F; BD II = 2 M/11 F	8 M/13 F	1. BD I vs HC; 2. BD II vs HC	BD I = 87.60; BD II = 112.8	BD I = 35.6 (10.9); BD II = 35.1 (9.8)	38.3 (11.9)	Yes	BP I = 6.7 (5.8); BP II = 9.5 (6.6) 17 item	SCID	N/A	1.5	2 × 2 × 2.2	13	900	p < 0.001 (uncorrected, k ≥ 50), then applied SVC to significant clusters using 8 mm radius at peak with FDR p < 0.001
Mahon et al. (2009)	15 M/15 F	22 M/16 F	BD (I, II & NOS) vs HC	N/A	33.4 (8.7)	31.9 (8.6)	Yes	N/A	SCID	N/A	1.5	1.7 × 1.7 × 5	25	1000	Voxelwise p < 0.001, k ≥ 50 & FDR corrected p < 0.015
Sussmann et al. (2009)	22 M/20 F	19 M/19 F	BD I vs HC	219.60	39.6 (10.1)	37.2 (11.9)	Yes	2.3 (5.8)	SCID	N/A	1.5	1.7 × 1.7 × 2.8	51	1000	p < 0.001 (uncorrected)
Wessa et al. (2009)	11 M/11 F	12 M/9 F	BD I & BD II (remitted) vs HC	264	45.41 (12.60)	42.95 (13.17)	Yes	1.55 (1.53)	DIGS	N/A	1.5	1.9 × 1.9 × 2	41	700	For VBA: p < 0.05 (FDR corrected), k ≥ 70
Zanetti et al. (2009)	13 M/24 F	12 M/14 F	BD I (active & remitted) vs HC	139.20	34.1 (9.0)	28.8 (9.5)	Yes	9.4 (9.4) 25 item	SCID	Borderline personality disorder, current alcohol or substance abuse or dependence	3	1.6 × 1.6 × 3	6	850	Voxelwise p < 0.001 (uncorrected), k ≥ 50

Table 1 (*continued*)

Paper	Patients male/female	HCs male/female	Contrast	Illness length (months)	Patient age mean (SD)	HC age mean (SD)	Meds	HRSD score mean (SD) # items	Dx tool	Excluded psychiatric comorbidities	B_0	Vox. size (mm)	Directions	Highest b (s/mm^2)	p
SAD															
Baur et al. (2011)	18 M/7 F	18 M/7 F	SAD vs HC	192	32 (10.4)	32 (10.1)	Yes	N/A BDI = 15 (10.8)	MINI	N/A	3	0.9 × 0.9 × 3.2	21	1000	Voxelwise p < 0.00001 (uncorrected), cluster-extent FWE correction p < 0.05, k ≥ 48
Phan et al. (2009)	15 M/15 F	10 M/20 F	SAD vs HC	N/A	27.20 (7.80)	29.90 (8.13)	Free for < 8 weeks (except n = 1)	N/A BDI = 10.7 (6.51)	SCID	OCD, PTSD, BD, psychotic disorder	3	0.9 × 0.9 × 3	12	900	p < 0.01 (uncorrected) volume ≥ 100 mm3
Qiu et al. (2014)	12 M/6 F	12 M/6 F	SAD vs HC	49.22	22.72 (3.85)	21.78 (3.90)	Naïve	N/A	SCID	DSM axis I or II disorders, drug abuse or dependence	3	1.9 × 1.9 × 3	15	1000	p < 0.01, uncorrected
OCD															
Admon et al. (2012)	10 M/3 F	10 M/3 F	OCD vs HC	96	25.5 (1.0)	27.0 (0.5)	All taking	N/A	SCID	Substance abuse	1.5	1.6 × 1.6 × 3	6	1000	Cluster p < 0.005, k ≥ 10, uncorrected
Cannistraro et al. (2007)	3 M/5 F	4 M/6 F	OCD vs HC	N/A	26.4 (5.9)	23.3 (1.5)	Med-free 4+ weeks	N/A	SCID	Current or past axis I Dx	1.5	2 × 2 × 2	6	600	Voxelwise p < 0.05, k ≥ 20, uncorrected, then cluster p < 0.005, uncorrected
Garibotto et al. (2010)	15 M/0 F	16 M/0 F	OCD vs HC	121.56	31.87 (7.92)	29.67 (6.3)	Yes	N/A	DSM-IV	Hoarding behavior, motor tics, Tourette's, axis I or II Dx	1.5	1.9 × 1.9 × 2.3	35	1000	p < 0.005, uncorrected, k ≥ 20
Li et al. (2014)	Both OCD N = 11 (M/F N/A)	N = 11 (M/F N/A)	1. TROCD vs HC; 2. OCD vs HC	TROCD = 23.4; OCD = 18.6	N/A	N/A	All	TROCD = 15.36 (4.30); OCD = 13.55 (2.94)	SCID	Current axis I Dx or psychiatric Dx in past year	3	2 × 2 × 2	30	1000	p < 0.05 FDR corrected, k ≥ 50
Menzies et al. (2008)	9 M/21 F	10 M/20 F	OCD vs HC	N/A	32.2 (11.1)	33.7 (11.2)	Yes	N/A	DSM-IV, MINI	Hoarding or motor tics, Hx of substance abuse, DSM-IV axis I Dx (except PD)	1.5	2.3 × 1.9 × 4	25	1000	Cluster p < 0.017, uncorrected
Nakamae et al. (2008)	9 M/6 F	9 M/6 F	OCD vs HC	121.2	29.7 (6.9)	29.1 (6.0)	All currently taking	N/A	SCID	SUD, schizophrenia, schizoaffective disorder, delusional disorder, brief reactive psychosis, psychotic disorder NOS	1.5	1.8 × 2.2 × 3	15	1000	p < 0.001 (uncorrected), k ≥ 100
Szeszko et al. (2005)	10 M/5 F	10 M/5 F	OCD vs HC	259.2	38.5 (10.9)	38.5 (11.8)	Yes	13.9 (4.9)	SCID	SUD, schizophrenia, schizoaffective disorder, delusional disorder, brief reactive psychosis, psychotic disorder NOS	1.5	1.7 × 1.7 × 5	25	1000	p < 0.005, k ≥ 20 (uncorrected)
Yoo et al. (2007)	8 M/5 F	8 M/5 F	OCD pre-treatment	86.4	27.8 (7.3)	26.9 (7.0)	Naïve	12.5 (7.6)	SCID	Psychosis, BD, SUD, Tourette's	1.5	1.7 × 1.7 × 4	6	600	p > 0.001 (uncorrected),

			vs HC							or other tic-related conditions						k ≥ 20

PTSD

Study																
Kennis et al. (2015)	rPTSD = 15 M/0 F; PTSD = 18 M/0 F	24 M/0 F	Non-blast PTSD vs combat controls prior to PTSD treatment	N/A	rPTSD = 34.38 (9.58); PTSD = 36.61 (8.74)	37.64 (10.97)	Yes	N/A	DSM-IV	Alcohol or substance dependence	3	1.9 × 1.9 × 2	30	1000		TFCE p < 0.05, corrected
Kim et al. (2005)	8 M/12 F	9 M/11 F	PTSD vs HC	N/A	28.4 (8.9)	30.6 (8.9)	N/A	N/A	SCID and CAPS	Premorbid axis I Dx, antisocial or borderline personality disorder, ADHD or lifetime exposure to illicit substances	3	0.9 × 0.9 × 3.5	25	1000		p < 0.05, corrected
Sun et al. (2013)	8 M/13 F	9 M/13 F	Trauma exposed with PTSD vs trauma exposed without PTSD	All scans within 2 days of trauma, Dx made at 1 or 6 months	40.86 (12.26)	40.23 (12.54)	Med-free for 4 weeks prior to MRI	N/A	MINI and CAPS	Axis I Dx, drug or alcohol abuse or dependence within 6 months	3	2 × 2 × 2	20	1000		αSim p < 0.05 corrected, with voxel p < 0.001, cluster k ≥ 76
Zhang et al. (2011)	17 M/0 F	14 M/14 F	First episode PTSD vs HC	N/A	34.06 (4.97)	28.96 (6.22)	Naive	N/A	SCID	Any current or past comorbid psychiatric Dx	1.5	1.9 × 1.9 × 4	13	1000		p < 0.001, uncorrected, k ≥ 100

Note. Abbreviations: αSim = AlphaSim, antidep = antidepressant = BD = bipolar, CAPS = Structured Clinician-Administered PTSD Scale for DSM-IV, DIGS = Diagnostic Interview for Genetic Studies, DSM = Diagnostic and Statistical Manual, Dx = diagnosis, eating = eating disorder, ECT = electroconvulsive therapy, ep. = episode, FDR = False Discovery Rate, FWE = family-wise error, GAD = generalized anxiety disorder, HC = healthy control, HRSD = Hamilton Rating Scale for Depression, Hx = history, k = cluster size threshold, max = maximum, MDD = major depressive disorder, med = medication, MINI = Mini-International Neuropsychiatric Interview, NOS = not otherwise specified, OCD = obsessive compulsive disorder, PD = panic disorder, psychotic = psychotic disorder, PTSD = post-traumatic stress disorder, rMDD = remitted major depressive disorder, rPTSD = remitted post-traumatic stress disorder, SAD = Social Anxiety Disorder, SADS-LV = Schedule for Affective Disorders – Lifetime Version, SCID = Structured clinical interview for DSM disorders, SD = standard deviation, sect. = section, SVC = small volume correction, SUD = substance use disorder, TCFE = threshold-free cluster enhancement, TRD = treatment resistant depression, TROCD = treatment resistant obsessive compulsive disorder, VBA = voxel based analysis, vox size (mm) = voxel size in millimeters.

meta-analyze statistical parametric maps from diffusion-weighted studies that utilized whole brain WM as well as studies that used TBSS. This is due to the fact that the TBSS method restricts its analysis to mean FA tract skeletons, which do not completely overlap across studies. We chose the approach taken by previous researchers who used SDM (Nortje et al., 2013; Radua et al., 2014), and only meta-analyzed the VBA results. However for a supplementary analysis we also extracted data from the TBSS studies and conducted a separate meta-analysis in an attempt to support our results (see Supplement).

2.2. Data extraction

In total, there were 892 healthy controls (HCs) and 962 participants with ED. These included 415 participants with MDD, 273 with BD, 73 with SAD, 110 with OCD and 91 with PTSD. Only whole-brain WM results were reported; furthermore, we only included the whole-brain corrected results, unless the only significant results were uncorrected, then these were included (e.g. Arnold et al., 2012). SDM requires that included studies have the same correction across the whole brain (although it may differ between studies). This is to avoid biasing results towards regions more liberally thresholded (Radua and Matrix-Cols, 2009). When studies reported separate results by illness subtype, these were included as separate studies, i.e. two studies with non-overlapping contrasts for BD I and BD II (Liu et al., 2010; Ha et al., 2011), one study with separate results for suicide attempters and non-attempters with MDD (Jia et al., 2010), and one study with separate results for OCD and treatment-resistant OCD (TROCD) (Li et al., 2014). The 37 VBA studies and 41 contrasts included in the meta-analysis are reported in Table 1.

Data were extracted independently by two researchers (LMJ, AB), including the number of males and females in each group, age, illness duration, diagnostic tool, handedness, medication status, psychiatric co-morbidities, number of diffusion directions, B_0, voxel size, imaging sequence, b value and statistical thresholds. All imaging was performed as spin-echo EPIs. Where necessary authors were contacted for further information. For the meta-analysis, from each contrast we extracted peak MNI coordinates and t values of FA differences between ED and HC groups. Peak coordinates were double-checked by both researchers. Foci reported in Talairach space were converted to MNI space with Brett's transform (http://eeg.sourceforge.net/doc_m2html/bioelectromagnetism/mni2tal_matrix.html). When only z or p values were available, these were converted to t values using the SDM online conversion utilities.

2.3. Meta-analysis

Seed-based d Mapping (SDM), described in more detail elsewhere (Radua and Matrix-Cols, 2009; Radua et al., 2012), is a statistical technique for meta-analysis of neuroimaging data, including diffusion tensor imaging (DTI). SDM uses a random effects model that takes into account sample size, intra-study variability, and between-study heterogeneity, and statistical significance is assessed using a distribution-free permutation test (Radua et al., 2012). A main reason we chose SDM was because it has a specific mask correlation template for FA (Radua et al., 2011). This specific template accounts for the anisotropy in the spatial covariance of the brain, which increases the accuracy of the effect size maps, and has been successful in prior mood disorder studies (Nortje et al., 2013). While other meta-analyses of mood disorders have used different techniques, such as Activation Likelihood Estimation, these techniques are not optimal for use with DTI studies as they do not include a white-matter mask, and they also cannot include clusters of both increased and decreased FA in patients compared to HC.

Similar to Nortje et al. (2013), we employed threshold values of $p < 0.005$, $Z > 1$ and k = 10. This threshold has been shown to optimize sensitivity at the same time as correctly controlling the false positive rate, and was approximately equivalent to a corrected p value of .025 (Radua et al., 2012). Peaks of significant clusters were labeled automatically by the SDM software (Rojkova et al., 2016; Thiebaut de Schotten et al., 2011a, 2011b). A jack-knife analysis was performed to establish reliability of the results (Radua and Matrix-Cols, 2009). Publication bias was assessed with funnel plots (Supplement) and Egger tests. As per a previous SMD meta-analyses (Murphy and Frodl, 2011), a linear meta-regression assessed the influence of illness duration. An additional linear meta-regression assessed the influence of depression severity, as measured with the Hamilton Depression Rating Scale (HDRS).

3. Results

3.1. Transdiagnostic (shared) meta-analysis

The results of the ED meta-analysis are presented in Fig. 2. For all significant clusters, the ED studies reported reduced FA compared to HCs. Cluster peaks are reported in Table 2.

The largest cluster incorporated several white matter tracts, including the left forceps minor of the corpus callosum (CC), anterior thalamic radiations (ATR), inferior fronto-occipital fasciculus (IFOF) and the uncinate fasciculus. The second largest cluster peaked in the left superior longitudinal fasciculus (SLF), and also involved white matter in the parietal lobe. The third largest cluster peaked in the left cerebellum (hemispheric lobule I-IV), and there was a smaller cluster in the right cerebellum (hemispheric lobule VI). Another cluster peaked in the left superior corona radiata and incorporated the corticospinal tract. The smallest cluster peaked in the right SLF. For illustrative purposes, Fig. 3 shows the location of these clusters of reduced FA on the affected fiber tracts, which are otherwise intact. It should be noted that we did not perform tractography, rather Fig. 3 shows the significant clusters found in our meta-analysis superimposed on a standard tractography image. Significant clusters are shown in green, and affected WM tracts in yellow, for illustrative purposes. A three dimensional video is available in the online Supplement.

To support these main findings from the VBA studies, we also extracted data from the TBSS studies and conducted a separate meta-analysis. These results are reported in the Supplement.

3.2. Illness duration analysis

A linear meta-regression was conducted in SDM to assess the influence of illness duration (years). Illness duration information was not available for 12 studies (three MDD, two BD, one SAD, two OCD and four, i.e. all PTSD). A univariate ANOVA found a significant effect of group for illness duration, $F(3, 25) = 4.87$, $p = 0.008$. The BD contrasts ($n = 10$) had the longest mean illness duration $= 11.80$, (SD $= 5.65$); SAD ($n = 2$), mean illness duration $= 10.05$, (SD $= 8.41$); OCD ($n = 7$), mean illness duration $= 8.65$, (SD $= 6.71$); the MDD contrasts ($n = 10$) included patients with the shortest mean illness duration $= 2.91$ (SD $= 3.00$). Post hoc tests found a significant difference in illness duration between the MDD and BD groups ($p = 0.005$).

The linear meta-regression revealed longer illness duration was associated with increased FA in nine diffuse brain regions, including several posterior clusters located in the forceps major, and decreased FA in seven brain regions, of which six were left lateralised. These results are reported in Table 3. The finding of increased FA associated with longer illness duration in several regions is difficult to interpret because of the significant difference in illness duration between the MDD and BD group, which may reflect an interaction.

3.3. Depression severity analysis

A linear meta-regression was also conducted to assess the influence of current depression severity, as measured by the Hamilton Rating Scale for Depression (HDRS). HDRS score was not available for 18 studies (three each of MDD and BD, five of OCD, and three and four, i.e. all, of
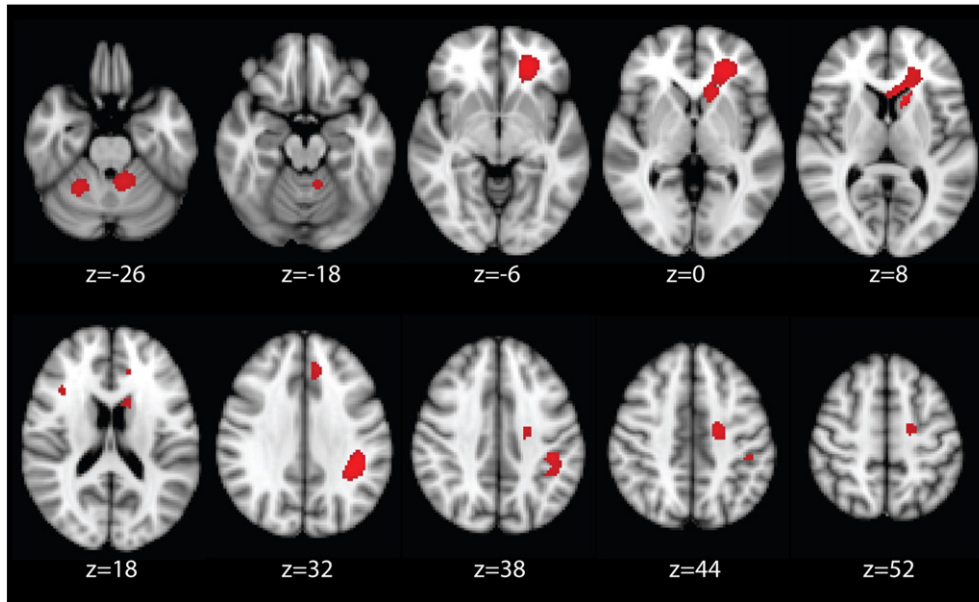
Fig. 2. Regions of decreased fractional anisotropy in emotional disorders compared to healthy controls, identified from the meta-analysis of VBA studies, $p < 0.005$. *Note.* Figure is in radiological convention, i.e. left = right.

SAD and PTSD, respectively). An ANOVA found a significant main effect of group for HDRS, $F(2, 21) = 126.65$, $p < 0.001$. Post hoc tests revealed that the MDD contrasts ($M = 21.93$, $SD = 2.20$) included patients with significantly higher HDRS scores than the BD contrasts ($M = 4.72$, $SD = 2.53$), $p < 0.001$, and the OCD contrasts ($M = 13.83$, $SD = 1.18$), $p < 0.001$. The BD and OCD contrasts also differed significantly, $p < 0.001$. Table 4 reports that higher HDRS score (indicating greater current depression severity) predicted decreased FA in four regions, with the largest clusters peaking in the left ATR and superior longitudinal fasciculus. Comparison of these results to those of Fig. 4 reveals that as expected, given the significantly higher HDRS scores in the MDD group, the regions significantly associated with depression severity were all regions that showed significantly reduced FA in MDD in the disorder-specific meta-analysis, reported below.

3.4. Assessments of reliability and publication bias

The Jack-knife analysis found that the left SLF cluster (-40, -40, 32) remained significant across all 41 sub-analyses, indicating that this was the most robust finding. The largest cluster, located in the left IFOF (-24, 40, -6), remained significant in all sub-analyses except for in five sub-analyses for which the peak of this cluster shifted to a different location within this large cluster (-12, 18, 6). The cluster in the left superior corona radiata (-18, -12, 44) was quite robust, remaining significant in 40 sub-analyses (excluding Sussmann et al., 2009). The clusters in the left (-10, -48, -24) and right cerebellum (26, -52, -26) were both significant in 39 sub analyses, and the right SLF (36, 20, 20) was significant in 38 sub-analyses.

Publication bias was assessed by extracting values from each of the significant peaks in the ED meta-analysis. Funnel plots were created to allow for visual inspection of the inter-study heterogeneity (see Supplement). Egger tests were calculated and revealed no evidence for publication bias for any of the clusters, $p > 0.05$.

3.5. Disorder-specific meta-analyses

We also conducted separate meta-analyses for each ED. These results are presented in Fig. 4. Peak coordinates of these results are reported in the Supplement. The two largest peaks of reduced FA in MDD contrasts were in left hemisphere regions of the ATR and SLF, which were two of the most robust regions identified from the Jackknife analysis. The BD contrasts revealed the only increase in FA (relative to HC), in the right SLF. They also had decreased FA in this same tract more anteriorly. Other regions of decreased FA in the right hemisphere were the forceps minor and IFOF, however the BD group also had decreased FA in several left hemisphere regions, including the SLF, cingulum, forceps minor and cerebellum. The SAD studies showed a primarily left lateralized pattern of decreased FA, which occurred in the SLF and corticospinal tract, and the UF bilaterally (more ventrally on the right). Two of the clusters for SAD overlapped with OCD and/or MDD. The OCD group showed reduced FA in several regions, all of which overlapped with at least one other diagnosis. These included the left SLF, IFOF, cerebellum and right ILF, shared with the BD contrasts, the left superior corona radiata shared with the SAD contrasts, the left IFOF/UF/ATR shared with the MDD and SAD contrasts, and the right cerebellum shared with the MDD group. Finally, like the OCD, SAD and to a lesser extent MDD contrasts, the PTSD contrasts revealed reduced FA in

Table 2
Cluster peaks of meta-analysis of VBA studies of emotional disorders vs HC.

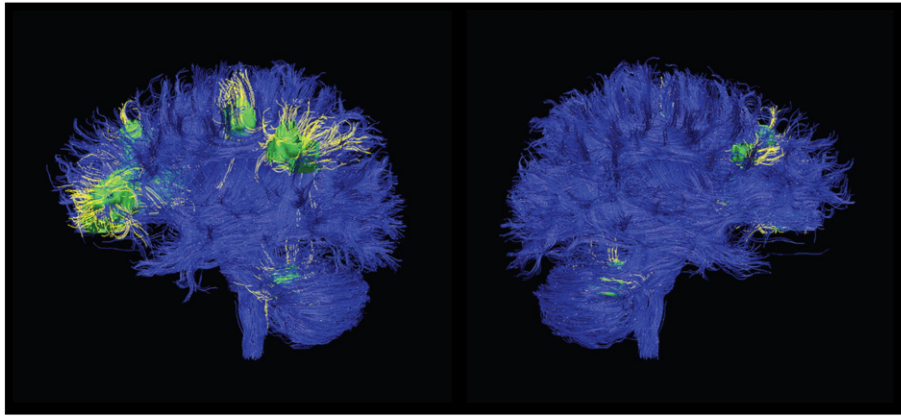| Peak region	x	y	z	Z	p	Voxels
Left forceps minor/ATR/IFOF/UF	-24	40	-6	-1.59	<0.0001	1228
Left SLF	-40	-40	32	-1.82	<0.0001	389
Left cerebellum (hemispheric lobule I–IV)	-10	-48	-24	-1.16	0.0006	237
Left superior corona radiata/corticospinal tract	-18	-12	44	-1.27	0.0002	201
Right cerebellum (hemispheric lobule VI)	26	-52	-26	-1.12	0.0008	109
Right SLF	36	20	20	-1.11	0.0008	69

Note. SLF = Superior longitudinal fasciculus, ATR = anterior thalamic radiation, IFOF = inferior fronto-occipital fasciculus, UF = uncinate fasciculus.

Fig. 3. White matter tracts implicated in VBA meta-analysis of emotional disorders. *Note.* Significant clusters are shown in green, and tracts implicated in yellow. The left hemisphere is shown in the left panel and the right hemisphere in the right panel.

primarily left hemisphere regions. Unlike OCD and to a lesser extent SAD, however, these regions did not overlap with those of other disorders. Reduced FA was found in the left UF, however this was inferior to the region identified in MDD, SAD and OCD. Similarly, the region of the left SLF identified in the PTSD contrasts was inferior to that of the other diagnoses. There was also reduced FA in the left cingulum and the right ILF in PTSD.

4. Discussion

The present meta-analysis revealed that overall, emotional disorders exhibit a shared pattern of reduced FA in fronto-temporal and fronto-parietal white matter tracts compared to HCs. The most robust and replicable finding (determined by Jack-knife analysis), peaked in the left superior longitudinal fasciculus (SLF). The largest cluster identified peaked in the left forceps minor/inferior fronto-occipital fasciculus, and extended to include the uncinate fasciculus (UF) and anterior thalamic radiations (ATR). There was also a significant peak in the region of the left superior corona radiata/corticospinal tract, bilateral cerebellum and right SLF. These tracts link important networks involved in cognition, affect, and the regulation thereof. The discussion therefore focuses on the SLF, IFOF, UF, anterior corpus callosum and tracts within the cerebellum, including evidence for their potential functional roles that may be relevant to emotional disorders. We also present converging evidence for alterations in FA in these tracts in ED from the TBSS

studies listed in the Supplement. Due to space limitation, relatively less discussion of ED differences is provided.

4.1. Superior longitudinal fasciculus

The left SLF was the most replicable tract identified in the transdiagnostic meta-analysis. The SLF is a major association fiber tract connecting parieto-temporal association areas and the frontal lobe, and links aspects of default mode (medially) and cognitive control (laterally) networks. The SLF II terminates in the dorsolateral PFC (Makris et al., 2005), which is closely associated with executive functioning (Tekin and Cummings, 2002). Thus the SLF II may offer a means for the PFC to control the focusing of spatial attention by the parietal lobe (Schmahmann et al., 2008; Makris et al., 2005). The SLF III is located in the parietal and frontal opercula and reaches from the supramarginal gyrus to ventral prefrontal and premotor areas (Makris et al., 2005). It provides higher order somatosensory input to the ventral premotor region and pars opercularis. It may also be important for monitoring hand and facial action, as well as articulatory and phonemic aspects of language (Schmahmann et al., 2008; Makris et al., 2005). These functions are consistent with the impairments in perception of and attention to emotional information in ED, including prosody and facial expressions of emotion. Our findings of reduced FA in the SLF are also supported by the TBSS studies reported in the Supplement, including findings from studies of reduced FA in the left SLF in OCD (Spalletta et al., 2014) and PTSD (Fani et al., 2012), as well of studies of BD

Table 3
Results of linear meta-regression of VBA studies of illness duration vs HCs.

Peak region	Change of FA	x	y	z	Z	p	Voxels
Superior longitudinal fasciculus	+	−30	−48	50	1.43	0.0001	138
Corticospinal tract	+	18	−26	64	1.33	0.0003	134
Forceps minor	+	−20	54	18	1.42	0.0002	128
Forceps major	+	−4	−80	22	1.45	0.0001	99
Anterior thalamic radiation	+	−14	18	8	1.15	0.0012	109
Forceps major	+	24	−80	30	1.26	0.0006	92
Corticospinal tract	+	6	−46	64	1.45	<0.0001	75
Forceps major	+	−34	−80	30	1.25	0.0007	82
Forceps major	+	4	−80	24	1.32	0.0004	59
Superior corona radiata	−	−20	−10	50	−2.06	0.0001	285
Uncinate fasciculus	−	−24	34	−8	−1.26	0.0021	113
Inferior frontooccipital fasciculus	−	−32	−30	14	−1.30	0.0018	72
Cingulum	−	−8	12	36	−1.48	0.0010	66
Superior longitudinal fasciculus	−	−48	−40	40	−1.30	0.0018	53
Cingulum	−	8	18	36	−1.20	0.0025	35
Forceps major	−	−18	−80	4	−1.10	0.0035	17

Note. FA is increased or decreased with longer illness duration.

Table 4
Regions of decreased FA associated with greater Hamilton Depression Rating Scale score.

peak region	x	y	z	Z	p	voxels
Anterior thalamic radiation	−14	20	4	−3.00	<0.0001	730
Superior longitudinal fasciculus	−38	−42	32	−2.88	<0.0001	354
Cerebellum (right VI)	28	−54	−24	−2.26	<0.0001	247
Corticospinal tract	16	−18	54	−1.48	0.0025	53

Note. FA is decreased with greater Hamilton Depression Rating Scale score.

(Ambrosi et al., 2013), particularly in the left SLF (Versace et al., 2010), as well as in the right hemisphere in MDD (Murphy et al., 2012) and OCD (Benedetti et al., 2013), and bilaterally in treatment-resistant depression (de Diego-Adelino et al., 2014).

We also found evidence for increased FA in BD in the right SLF. Increased FA in BD has been reported previously (Versace et al., 2008). Whilst this may seem counter-intuitive, it is important to note that higher FA reflects more numerous, myelinated, dense, and coherently orientated fibers in many circumstances, but it does not always signify healthier WM or beneficial outcomes. On the contrary, higher FA could suggest, for example, compensatory mechanisms, as WM structure is shaped by experiences (Thomason and Thompson, 2011). Therefore, when evaluating results of FA studies, the interpretation that decreased FA is maladaptive and increased FA is adaptive may be too simplistic an explanation.

4.2. Inferior fronto-occipital fasciculus

The largest cluster peaked in the left IFOF, connecting the dorsolateral and inferolateral frontal cortex with the occipital lobe and posterior temporal cortex (Catani et al., 2002). As such, the IFOF is an intra- and cross-network connection of cognitive control and salience networks often implicated in EDs. The IFOF traverses the sagittal striatum, and progresses posteriorly through the medial temporal lobe, passing through the anterior floor of the external capsule where it is parallel to the inferiorly located UF fibers (Catani et al., 2002). The fronto-occipital fasciculus comprises part of the dorsal visual stream and is implicated in peripheral vision and processing visuospatial information (Schmahmann et al., 2008) and the IFOF is located near a region important for processing of facial emotions (Nakamura et al., 1999). The finding of reduced FA in the left IFOF in the VBA studies is supported by the TBSS studies listed in the Supplement, which have reported reduced FA in this region in patients with OCD (Benedetti et al., 2013), MDD

(Kieseppa et al., 2010), melancholic MDD (Korgaonkar et al., 2011), and in patients with MDD who were homozygous for the A allele of NTRK2 (Murphy et al., 2012), a neurotrophic factor hypothesized to be a risk factor for MDD. Reduced FA bilaterally has also been reported in the IFOF in patients with TRD (de Diego-Adelino et al., 2014) in BD (Ambrosi et al., 2013; Chan et al., 2010), PD (Lai and Wu, 2013) and OCD (Fan et al., 2016).

4.3. Uncinate fasciculus

The UF joins the orbital and polar frontal cortex to the anterior temporal lobe, as it reaches out from the amgydala, hippocampal gyrus, uncus and temporal pole (Catani et al., 2002). Considered part of the limbic system (emotion and salience networks), the UF may be a pathway via which the PFC exerts control over limbic regions involved in emotion processing, an ability that is reduced in MDD (Anand et al., 2005). The UF is an important part of the system for regulating emotional responses to auditory stimuli (Schmahmann et al., 2008), and could also play a role in attaching emotional valence to visual stimuli. Reduced FA in the UF might support a mechanism of decreased salience and emotional network connectivity and activity in limbic regions, including the amygdala and subgenual anterior cingulate cortex, which lie at both ends of the UF.

The strongest evidence for reduced FA in the UF in ED is from studies of MDD. For example, tractography studies (excluded from our analysis as they do not report stereotactic coordinates) have reported reduced FA in the UF in MDD (Zhang et al., 2012; de Kwaasteniet et al., 2013). Murphy et al. (2012) found reduced FA in patients with MDD compared to HC in the left UF. Another FA study with an upper age limit beyond 65 years also found reduced FA in the left UF in MDD (Steele et al., 2005). However Lai and Wu (2013) failed to demonstrate reduced FA in panic disorder, and results from individual studies of BD are more mixed, with Ambrosi et al. (2013) finding reduced FA in the UF of patients with BD, but Versace et al. (2008) reporting increased FA. Further research is necessary to determine the impact of hypo/manic symptoms on FA in BD.

4.4. Anterior corpus callosum

The CC is critical for interhemispheric communication, particularly the integration of high-level cognitive, emotional, perceptual and linguistic functions across numerous networks (Gazzaniga, 2000). The forceps minor of the CC was implicated in the largest cluster in the ED
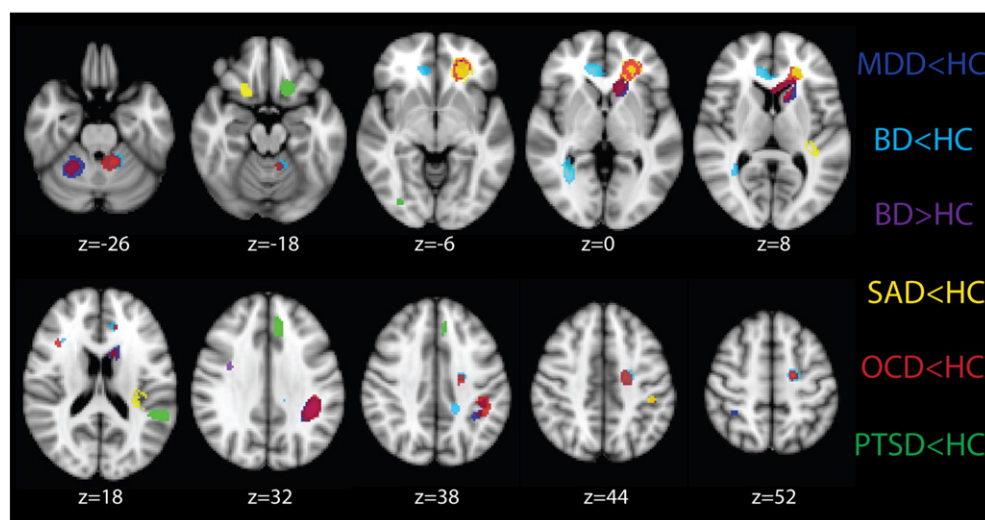
Fig. 4. Regions of differences in fractional anisotropy between MDD, BD, SAD, OCD, PTSD and HC identified from separate meta-analyses of VBA studies for each disorder. *Note.* Figure is in radiological convention, i.e. left = right. MDD = major depressive disorder, BD = bipolar disorder, SAD = social anxiety disorder, OCD = obsessive-compulsive disorder, PTSD = post-traumatic stress disorder, HC = healthy controls.

analysis. Some, but not all studies (Yurgelun-Todd et al., 2007) that were not included because they exceeded the upper age limit to 65 have also reported reduced FA in the forceps minor and genu of the CC in individuals with MDD (Cole et al., 2012; Seok et al., 2013) and BD (Benedetti et al., 2011). Numerous TBSS studies included in the Supplement reported reduced FA in the forceps minor/genu of the CC, including for MDD (Murphy et al., 2012), young adults with first-episode, treatment-responsive depression (Guo et al., 2012) and patients with treatment resistant depression (de Diego-Adelino et al., 2014). One TBSS study of OCD also found reduced FA in the forceps minor bilaterally (Benedetti et al., 2013), consistent with Fig. 4 that showed this region to be affected in the left hemisphere in OCD in the VBA meta-analysis.

4.5. Cerebellum

Tracts within the cerebellum are known to connect with the PFC, subcortical limbic nuclei, and brainstem structures that produce mono-amines (Schmahmann, 2000). The anterior lobe has a primary sensori-motor region, and the posterior lobe contains association areas important for higher order behavior, via connections with basis pontis and thalamic nuclei (Schmahmann and Caplan, 2006). Disrupted cerebellar and corticocerebellar connectivity has been proposed as a main neurobiological mechanism of emotion dysregulation (Schmahmann, 2000), as lesions confined to the cerebellum can result in a cerebellar cognitive affective syndrome that includes dysregulation of affect (Schmahmann and Caplan, 2006).

4.6. Distinctiveness of each finding across EDs

Notably, the results of the transdiagnostic meta-analyses are primarily left lateralized. Severity of post-stroke depression has also been correlated with proximity of stroke lesion to the left frontal pole, which may align with the shared negative affect across these EDs. However this association has been found both in cortical lesions and lesions restricted to subcortical regions (Starkstein and Robinson, 1987), thus depression following brain lesions cannot be attributed to disruptions to WM alone. Furthermore, stroke lesions impair signal transport by destroying nerve fiber bundles locally, in contrast to decreased signal transport due to a more extensive impairment of neurons. Whilst our focus was largely on shared WM alterations across all three disorders, there is some evidence of differences. Most notably, the PTSD findings were the most distinct, demonstrating no overlap with any other EDs for any of the significant clusters in the separate meta-analyses for each disorder. The OCD group showed overlap with both categories of mood disorders, including for clusters in the right hemisphere. OCD also showed overlap with SAD for two of the clusters, one of which it also shared with MDD (the largest cluster, i.e. the IFOF/ATR/UF/forceps minor cluster). The BD findings were more bilateral relative to the tendency for left laterality observed across the other groups. This laterality finding harkens back to early lesion research suggesting right hemisphere stroke is associated with a hypomanic state characterized by inappropriate cheerfulness, loss of interest, anxiety, slowness and agitation (Robinson et al., 1983). Left prefrontal hypometabolism has also been reported in patients with BD in the depressed phase (Baxter et al., 1989). However, subsequent studies have found little consistent support for lateralization of mood or emotion, and only limited support for the valence hypothesis of emotion in the frontal cortex (Wager et al., 2003). BD also differed from the other disorders in that it was the only one to find increased FA in any region. This finding supports previous findings that BD loads on a separate factor from MDD and AD (Watson, 2005; Kotov et al., 2015); however it may also suggest BD is the addition of a separate manic disorder to MDD and/or AD. Future studies might examine the relationship between increases in FA for contrasts between internalizing and externalizing symptoms.

4.7. Limitations

The present study had several limitations. First, there were insufficient GAD studies using VBA that met our inclusion criteria, and all studies of panic disorder used TBSS thus were only able to be included in the Supplemental analysis. Second, we chose to exclude studies with *any* subjects outside of the age range 18–65 to limit the potential influence of degenerative conditions and development, thus numerous studies primarily involving adults were excluded (Steele et al., 2005; Seok et al., 2013; Benedetti et al., 2011). The included studies differed in diffusion imaging methods, which could limit the interpretability of the findings. For example, more recent studies tend to use correcting procedures for EPI distortions that more dated studies did not, which could impact findings in proximity of brain regions that are sensitive to B_0 inhomogeneities. There are also limitations in the interpretation of FA, as it is only an indirect marker of WM microstructure. Although commonly interpreted to reflect decreased myelination or progressive myelin degradation due to a loss of oligodendrocyte cells, there are other interpretations of reduced FA. For instance, low FA could also be the result of axonal damage, reduced fibers (axonal density), or disorganized fibers (Thomason and Thompson, 2011; Kumar et al., 2015). It could all be the result of inadequate stimulation of anatomic structures or pathways. Furthermore, microstructural alterations in the brain may be an antecedent and/or a consequence of disease. It is unclear at this point how length and severity of illness might affect DTI measurements. The diffusion tensor method has received criticism for underestimating FA in regions where fibers cross. Thus findings in such areas of crossing fasciculi should be interpreted with caution. Newer models such as mode of anisotropy avoid such limitations and allow a more meaningful interpretation of FA results (Kumar et al., 2015). Finally, our discussion of the functionality of the decreased FA in each area in relation to the EDs are speculative and require additional research to verify whether these functions are indeed reduced in these disorders, and whether they precede or are a consequence of them.

5. Conclusions

Consistent with the NIMH RDoC initiative, we attempted to identify WM alterations common across different types of ED compared to HCs. Reduced FA occurred in tracts that connect emotion, salience and cognitive control networks. Future studies would benefit from looking at transdiagnostic symptom, cognitive, and affective domains, given the high co-occurrence of many depressive and/or anxious disorders. Through these multi-method studies the field will develop a more complete understanding of common and specific symptoms and associated phenotypes and develop improved treatment targets.

Supplementary data to this article can be found online at http://dx.doi.org/10.1016/j.nicl.2016.09.001.

Conflict of interest

The authors disclose no conflicts of interests, including financial interests, activities, relationships and affiliations.

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