Normal Brain Tissue Volumes after Long-Term Recovery in Anorexia and Bulimia Nervosa

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Background: Individuals who are ill with anorexia (AN) and bulimia nervosa (BN) often have increased cerebrospinal fluid (CSF) volumes and decreased total gray and white matter volumes. It is unclear whether such disturbances persist after recovery from an eating disorder.

Methods: Magnetic resonance imaging was performed on 40 women who were long-term recovered (>1 year no binging, purging, or restricting behaviors, normal weight, and menstrual cycles, not on medication) from restricting or binge/purging type AN or BN and 31 healthy control women (CW). Voxel-based morphometry (VBM) was used for data analysis.

Results: Recovered AN and BN subgroups were similar to CW in terms of cerebrospinal fluid (CSF) volume as well as total or regional gray or white matter volume.

Conclusions: These findings suggest that structural brain abnormalities are reversible in individuals with eating disorders after long-term recovery.

Key Words: Voxel based morphometry, magnetic resonance imaging, anorexia nervosa, bulimia nervosa, recovery, structural brain abnormality

There is ample evidence that individuals ill with AN have reduced brain mass. For example, post mortem studies showed reduced cerebral mass with prominent sulci and small gyri (Gagel 1953; Martin 1958). Neuroimaging studies with computerized tomography (CT) confirmed these findings and additionally demonstrated enlarged ventricles (Dolan et al 1988; Krieg et al 1988; Palazidou et al 1990). Magnetic resonance imaging (MRI) studies showed large cerebrospinal fluid (CSF) volumes in association with deficits in both total gray matter (GM) and total white matter (WM) volumes (Katzman et al 1996; Swayze et al 2003) as well as enlarged ventricles (Golden et al 1996; Swayze et al 1996). These abnormalities were less pronounced in ill BN individuals (Krieg et al 1989) who also have decreased cortical mass (Hoffman et al 1989; Husain et al 1992; Laessle et al 1989).

It is less clear whether these abnormalities persist after recovery from AN and BN, or are partially or fully reversible (Katzman et al 1997; Lambe et al 1997; Swayze et al 2003). Thus the goal of the present study was to assess brain tissue abnormalities in a large sample of individuals long-term recovered from eating disorders (EDs). To our knowledge this is the first study looking at differences in AN subtypes as well as BN, in the recovered state. Since little is known about the process of recovery, the findings in this study may shed light on the development of one of the most striking physical consequences of EDs.

Methods and Materials

Sample Collection
Forty recovered ED subjects were recruited, 14 meeting criteria for restricting type AN (RAN), 16 for binge-purging type AN (BAN) and 10 for BN. To be considered “recovered,” for the previous year subjects had to (1) maintain a weight above 85% of average body weight; (2) have regular menstrual cycles; (3) have not binged, purged, restricted food intake or exercised excessively; and (4) not used psychoactive medications. Thirty-one CW were recruited. They had no history of an ED or any psychiatric or medical illness.

The studies were conducted according to institutional review board regulations of the University of Pittsburgh, and all subjects gave written informed consent. The Structured Clinical Interview for DSM-IV Axis I and II Disorders was used to diagnose DSM-IV Axis I and personality disorders (First et al 1996).

MRI Acquisition
T1-weighted volumetric “spooled gradient recalled” (SPGR) MRI scans (echo time [TE] = 5 msec, repetition time [TR] = 25 msec, flip angle 40°, number of excitations [NEX] = 1; section thickness = 1.5 mm) were acquired in the coronal plane on a 1.5-T scanner (GE Medical Systems, Milwaukee, Wisconsin).

Image Analysis
Morphologic changes were investigated using the “optimized” VBM method (Good et al 2001) using SPM2 (Wellcome Department of Cognitive Neurology, London, United Kingdom). The individual MRIs were segmented into GM, WM and CSF images in their native space. The GM images were normalized to Montreal Neurological Institute (MNI) space and the transform applied to the original SPGR image. The normalized SPGR images were segmented once again, and the new GM, WM and CSF images were modulated with the Jacobian determinates derived from the normalization resulting in volumetric GM, WM and CSF images. Total GM and WM volumes were extracted from...
the modulated images. Modulated images were smoothed with a 12 mm Gaussian kernel prior to statistical analysis in SPM2.

**Statistical Analysis**

All data analyses were performed using SPSS 11.0 (SPSS Inc., Chicago, Illinois). For multiple group comparisons analysis of variance was used to test for differences across the groups followed by Tukey post hoc test when the overall difference was statistically significant. All image analysis group comparisons were performed using analysis of variance within SPM2 (p < .05 corrected for multiple comparisons using false discovery rate).

**Results**

Recovered subjects and CW were of similar age and body mass index (BMI) (Table 1). The average length of recovery ranged from 29.8 to 39.5 months. Groups had similar volumes for total GM, WM, and CSF as well as regional values (data not shown). There was a small but not significant increase in recovered subjects with age vs. CSF: r = −.193, p = .232; age vs. WMV: r = .167, p = .304; age vs. GMV: r = .097, p = .552) as well as in CW (age vs. GMV: r = −.160, p = .388; age/WMV: r = .139, p = .457; age/CSF: r = .229, p = .216). The findings remained the same after covarying subjects to CW.

**Discussion**

Recovered AN and BN women were similar to each other and to CW, in total and regional GMV, WMV and CSF volume. Thus, this study suggests that structural brain abnormalities may be reversible after long-term recovery from an ED. To our knowledge, this is the first study to use rigorous criteria to define recovery status based upon the requirements of normal nutrition, stable weight, absence of medication, and normal menstrual cycle for at least one year. Not all studies have compared AN subjects to CW.

Previous research studies found results consistent with the present study, while other studies reported persistent abnormalities. Golden et al. (1996) studied individuals with AN after nutritional rehabilitation and detected normal total ventricular volumes compared to CW. Kingston found enlarged lateral ventricles and dilated sulci in AN compared to CW. However, subjects were rescanned after at least 10% weight gain (Kingston et al. 1996) so that length of recovery was variable.

Swayze rescanned individuals with AN after weight normalization, finding a decrease in total and regional CSF volumes as well as an increase in GMV and WMV compared to pre-scan (Swayze et al. 2003). Degree of recovery may be critical for normalization of brain morphometry. However, there was no comparison to CW, or information about menstrual function or length of recovery. Lambe described an increase in GMV and WMV as well as a decrease in CSF volume in recovered compared to ill AN. Compared to CW, recovered AN had greater CSF volume and smaller GMV, but normal WMV (Lambe et al. 1997). Katzman replicated these findings in a longitudinal study of a subset of 6 weight recovered AN subjects (Katzman et al. 1997). Subjects in both of these studies were weight restored for at least 1 year and had started to menstruate or returned to normal menses at follow-up. More recently Neumarker found a significant volume difference of the lateral ventricles and the Sylvian fissure in AN compared to CW, which abated with weight restoration, whereas deficits of the mesencephalon and pons persisted (Neumarker et al. 2000). Again length of weight restoration is unclear.

All subjects in the present study also participated in positron emission tomography (PET) studies. For PET image analysis partial volume correction (PVC) factors were created by segmenting the SPGR MR image into brain and nonbrain regions, which created a binary image (Meltzer et al. 1996, 1999). After

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**Table 1. Demographics**

<table>
<thead>
<tr>
<th>Group</th>
<th>CW n = 31</th>
<th>RAN n = 14</th>
<th>BAN n = 16</th>
<th>BN n = 10</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean</td>
<td>SD</td>
<td>Mean</td>
<td>SD</td>
<td>Mean</td>
</tr>
<tr>
<td>Study Age (years)</td>
<td>26.8 7.3</td>
<td>23.7 5.3</td>
<td>27.4 7.2</td>
<td>24.0 6.1</td>
</tr>
<tr>
<td>Current BMI (kg/m²)</td>
<td>21.9 2.0</td>
<td>21.2 2.0</td>
<td>21.2 1.5</td>
<td>23.1 2.4</td>
</tr>
<tr>
<td>High Past BMI (kg/m²)</td>
<td>22.8 2.0</td>
<td>21.6 2.3</td>
<td>23.6 2.8</td>
<td>24.7 2.85</td>
</tr>
<tr>
<td>Low Past BMI (kg/m²)</td>
<td>20.1 1.4</td>
<td>14.1 1.4</td>
<td>14.8 2.0</td>
<td>19.2 2.1</td>
</tr>
<tr>
<td>Age of Onset (years)</td>
<td>18.2 4.5</td>
<td></td>
<td>15.7 2.7</td>
<td>15.5 2.3</td>
</tr>
<tr>
<td>Length of Recovery (months)</td>
<td>28.7 20.4</td>
<td>39.5 52.7</td>
<td>29.8 18.1</td>
<td>.753</td>
</tr>
</tbody>
</table>

CW, healthy control women; RAN, recovered anorexic women, restricting type; BAN, recovered anorexic women, binge/purging type; BN, recovered bulimic women; BMI, body mass index.

**Table 2. Distribution of Segmented Tissues**

<table>
<thead>
<tr>
<th>Group</th>
<th>CW n = 31</th>
<th>RAN n = 14</th>
<th>BAN n = 16</th>
<th>BN n = 10</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean</td>
<td>SD</td>
<td>Mean</td>
<td>SD</td>
<td>Mean</td>
</tr>
<tr>
<td>GMV (ml)</td>
<td>729.3 45.4</td>
<td>749.9 53.0</td>
<td>740.2 55.6</td>
<td>770.2 54.8</td>
</tr>
<tr>
<td>WMV (ml)</td>
<td>393.6 35.3</td>
<td>383.6 26.0</td>
<td>403.1 42.3</td>
<td>402.7 25.7</td>
</tr>
<tr>
<td>CSF (ml)</td>
<td>313.4 55.6</td>
<td>309.3 61.9</td>
<td>342.2 42.9</td>
<td>334.1 47.5</td>
</tr>
<tr>
<td>TIV (ml)</td>
<td>1436.3 109.2</td>
<td>1442.8 115.1</td>
<td>1485.4 107.5</td>
<td>1507.0 106.6</td>
</tr>
</tbody>
</table>

CW, healthy control women; RAN, recovered anorexic women, restricting type; BAN, recovered anorexic women, binge/purging type; BN, recovered bulimic women; GMV, gray matter volume; WMV, white matter volume; CSF, cerebrospinal fluid; TIV, total intracranial volume.

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smoothing, regions of interest (ROI) were placed on the binary image to determine the fraction of nonbrain within each ROI. PVC values were similar for CW and recovered subjects (Bailer et al 2004). Thus 2 different methods showed similar conclusions (Good et al 2002).

While emaciation and malnutrition are known to be associated with altered brain morphometry abnormalities in AN, the physiology underlying changes in brain structure remains elusive. Because brain morphometry abnormalities appear to be reversible, they may be related to fluid shifts, perhaps caused by altered hormonal function. For example, Katzman reported that urinary free cortisol levels in ill AN were positively correlated with total CSF volume and inversely correlated with GMV (Katzman et al 1996). Free serum cortisol levels of CW and recovered subjects in this study were similar (CW 16.1 ± 7.0, all REC. 17.1 ± 6.2, p = .994). Other studies suggest that low triiodothyronine is associated with ventricular size in AN and BN (Krieg et al 1989). In addition, decreased serum proteins could result in decreased colloidal osmotic pressure and a shift of fluid from the intravascular space into the subarachnoid spaces (Heinz et al 1977). In summary, it is possible that alterations in brain volume in the ill state are a consequence of starvation-induced changes in fluid compartments which are reversed by improved nutrition.

The study is limited by its cross-section design. However, subjects met DSM-IV criteria and had elevated EDI-2 scores (not shown) when ill. Moreover, AN subjects had a history of low BMI similar in magnitude to that of ill subjects with altered brain volumes in past studies. There was a selection bias as only subjects whose MRs showed a high quality of brain tissue segmentation were included. In addition, subjects may not be representative of all who recover. While we depend on self-reports of the participants regarding their state of recovery, their normal plasma β-hydroxybutyrate levels (Bailer et al, in Press) support the likelihood of normal nutritional status.

In summary, these data indicate that brain tissue abnormalities might be reversible in ED patients after long-term recovery.

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