Disrupted functional connectivity of the periaqueductal gray in chronic low back pain

Citation

Published Version
doi:10.1016/j.nicl.2014.08.019

Permanent link
http://nrs.harvard.edu/urn-3:HUL.InstRepos:13454849

Terms of Use
This article was downloaded from Harvard University’s DASH repository, and is made available under the terms and conditions applicable to Other Posted Material, as set forth at http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA

Share Your Story
The Harvard community has made this article openly available. Please share how this access benefits you. Submit a story.

Accessibility
Disrupted functional connectivity of the periaqueductal gray in chronic low back pain

Rongjun Yu\textsuperscript{a,b,c,*}, Randy L. Gollub\textsuperscript{a,b}, Rosa Spaeth\textsuperscript{a,b}, Vitaly Napadow\textsuperscript{a,b}, Ajay Wasan\textsuperscript{a,b,d}, Jian Kong\textsuperscript{a,b}

\textsuperscript{a}Department of Psychiatry, Massachusetts General Hospital, Harvard Medical School, Charlestown, MA, USA
\textsuperscript{b}Martinos Center for Biomedical Imaging, Massachusetts General Hospital, Charlestown, MA, USA
\textsuperscript{c}Department of Psychology, National University of Singapore, Singapore, Singapore
\textsuperscript{d}Perioperative and Pain Medicine, Department of Anesthesiology, Brigham and Women's Hospital, Harvard Medical School (HMS), Boston, MA, USA

**A B S T R A C T**

Chronic low back pain is a common neurological disorder. The periaqueductal gray (PAG) plays a key role in the descending modulation of pain. In this study, we investigated brain resting state PAG functional connectivity (FC) differences between patients with chronic low back pain (cLBP) in low pain or high pain condition and matched healthy controls (HCs). PAG seed based functional connectivity (FC) analysis of the functional MR imaging data was performed to investigate the difference among the connectivity maps in the cLBP in the low or high pain condition and HC groups as well as within the cLBP at differing endogenous back pain intensities. Results showed that FC between the PAG and the ventral medial prefrontal cortex (vmPFC)/rostral anterior cingulate cortex (rACC) increased in cLBP patients compared to matched controls. In addition, we also found significant negative correlations between pain ratings and PAG–vmPFC/rACC FC in cLBP patients after pain-inducing maneuver. The duration of cLBP was negatively correlated with PAG–insula and PAG–amygdala FC before pain-inducing maneuver in the patient group. These findings are in line with the impairments of the descending pain modulation reported in patients with cLBP. Our results provide evidence showing that cLBP patients have abnormal FC in PAG centered pain modulation network during rest.

© 2014 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/3.0/).

1. Introduction

Chronic low back pain (cLBP) is one of the most common reasons for all physician visits in the USA and is a leading contributor to job-related disability and missed work (Chou and Shekelle, 2010; Hart et al., 1995). The etiology of cLBP is heterogeneous (Ehrlich, 2003). Non-specific cLBP, which represents the majority of cLBP patients, is characterized by a lack of recognizable pathology (Chou et al., 2007; Ehrlich, 2003; Savigny et al., 2009). Although cLBP is a serious health concern, treatment for cLBP has achieved limited success (Bogduk, 2004). Previous studies have shown that PAG stimulation can significantly inhibit behavioral responses to noxious stimuli in both animals (Mayer et al., 1971; Reynolds, 1969) and humans (Baskin et al., 1986; Hosobuchi et al., 1977). Recent studies have shown that the functional connectivity fluctuations and structural connectivity between the PAG and the ventral medial prefrontal cortex (vmPFC) predicted mind wandering away from pain, i.e., spontaneous disengagement of attention from pain (Kucyi et al., 2013). The structural connectivity between these two regions also predicted individual difference in placebo analgesia (Stein et al., 2012). It is now believed that the brainstem (PAG) receives direct projections from regions within the limbic forebrain such as the anterior cingulate cortex (ACC) and limbic-related areas such as the insula and amygdala and modulates pain by descending modulation of the spinal cord neurons (Brooks and Tracey, 2005; Fields, 2004; Heinricher et al., 2009; Ploner et al., 2010; Tracey and Mantyh, 2007). A recent study has shown that the PAG is functionally connected to the vmPFC/ACC, insula and amygdala during resting state (Kong et al., 2010b).

The insula is a key region in pain process (Bernard et al., 1992; Chudler et al., 1993; Craig, 2002; Craig et al., 2000; Kong et al., 2013a; Kong et al., 2006; Schneider and Lidsky, 1981; Wiech et al., 2005). A...
previous study suggested that prestimulus functional connectivity between the insula and pain-modulatory brain regions (e.g., PAG) differed between physically identical trials that were rated as painful and trials perceived as non-painful (Ploner et al., 2010). The amygdala has a central role in regulating emotional responses during acute and persistent pain (Martikainen et al., 2013; Neugebauer et al., 2004). Given the close anatomical connectivity between the PAG–insula–amygdala and their role in pain perception and modulation (Ploner et al., 2010), it is possible that alterations in these pathways may also contribute to the development or maintenance of chronic pain. Recent neuroimaging studies have shown that cLBP is associated with alterations in resting state brain activity (Apkarian et al., 2009; Apkarian et al., 2004; Baliki et al., 2011; Baliki et al., 2006; Kobayashi et al., 2009; Tagliazucchi et al., 2010; Wasan et al., 2011). However, the role of the PAG and the associated networks detected by resting state fMRI in cLBP is still unclear. In the present study, we investigated PAG centered brain resting state functional connectivity (FC) differences between cLBP patients and matched HCs and FC differences when cLBP patients experienced different levels of pain intensity. We hypothesized that cLBP would be associated with abnormal FC between the PAG and other brain regions including the vmPFC, insula, and amygdala, given the close link between these regions and their role in pain modulation.

2. Materials and methods

We briefly describe the experimental procedures below. Please also see a previous published study (Kong et al., 2013b) for more details on the experimental procedure. The data have been used in this previous publication (Kong et al., 2013b), but the analytic methods used here do not overlap. In that study, we compared structure and function difference between the cLBP and controls using structural imaging data with morphometric analysis and resting state MRI data with degree centrality (DC) analysis (Kong et al., 2013b). Degree centrality is a measure of local network connectivity and identifies the most connected nodes by counting the number of direct connections to all other nodes (Buckner et al., 2009). There is no overlap on the results between the previous paper and the current paper.

2.1. Participants

Eighteen cLBP patients and 18 healthy controls, matched for age and gender, completed the study (see Table 1 for demographic details).

<table>
<thead>
<tr>
<th>ID</th>
<th>Gender</th>
<th>Age</th>
<th>Race</th>
<th>BDI</th>
<th>Duration (years)</th>
<th>Pain intensity (Low pain)</th>
<th>Pain intensity (High pain)</th>
<th>BPI (avg)</th>
<th>Gender</th>
<th>Age</th>
<th>Race</th>
<th>BDI</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>F</td>
<td>48</td>
<td>White</td>
<td>13</td>
<td>3</td>
<td>4.5</td>
<td>3.5</td>
<td>7</td>
<td>F</td>
<td>47</td>
<td>White</td>
<td>0</td>
</tr>
<tr>
<td>2</td>
<td>M</td>
<td>41</td>
<td>Asian</td>
<td>8</td>
<td>4</td>
<td>5</td>
<td>6.5</td>
<td>6</td>
<td>M</td>
<td>37</td>
<td>Asian</td>
<td>10</td>
</tr>
<tr>
<td>3</td>
<td>F</td>
<td>49</td>
<td>Black</td>
<td>30</td>
<td>8</td>
<td>6.5</td>
<td>8.75</td>
<td>6</td>
<td>F</td>
<td>50</td>
<td>Black</td>
<td>4</td>
</tr>
<tr>
<td>4</td>
<td>F</td>
<td>47</td>
<td>Hisp.</td>
<td>7</td>
<td>3</td>
<td>10</td>
<td>9.5</td>
<td>3</td>
<td>F</td>
<td>49</td>
<td>Black</td>
<td>0</td>
</tr>
<tr>
<td>5</td>
<td>F</td>
<td>23</td>
<td>White</td>
<td>1</td>
<td>10</td>
<td>3</td>
<td>6</td>
<td>3</td>
<td>F</td>
<td>26</td>
<td>White</td>
<td>11</td>
</tr>
<tr>
<td>6</td>
<td>M</td>
<td>27</td>
<td>White</td>
<td>0</td>
<td>10</td>
<td>4</td>
<td>6.5</td>
<td>3</td>
<td>M</td>
<td>30</td>
<td>White</td>
<td>0</td>
</tr>
<tr>
<td>7</td>
<td>F</td>
<td>23</td>
<td>White</td>
<td>4</td>
<td>3</td>
<td>2</td>
<td>6</td>
<td>3</td>
<td>F</td>
<td>23</td>
<td>White</td>
<td>3</td>
</tr>
<tr>
<td>8</td>
<td>M</td>
<td>38</td>
<td>White</td>
<td>0</td>
<td>2</td>
<td>4.5</td>
<td>6</td>
<td>4</td>
<td>M</td>
<td>39</td>
<td>White</td>
<td>7</td>
</tr>
<tr>
<td>9</td>
<td>M</td>
<td>25</td>
<td>Multi.</td>
<td>0</td>
<td>5</td>
<td>4.5</td>
<td>7</td>
<td>3</td>
<td>M</td>
<td>27</td>
<td>White</td>
<td>0</td>
</tr>
<tr>
<td>10</td>
<td>F</td>
<td>44</td>
<td>White</td>
<td>9</td>
<td>12</td>
<td>2.25</td>
<td>6</td>
<td>4</td>
<td>F</td>
<td>45</td>
<td>White</td>
<td>0</td>
</tr>
<tr>
<td>11</td>
<td>M</td>
<td>30</td>
<td>Multi.</td>
<td>5</td>
<td>10</td>
<td>2</td>
<td>5.5</td>
<td>9</td>
<td>M</td>
<td>34</td>
<td>White</td>
<td>4</td>
</tr>
<tr>
<td>12</td>
<td>F</td>
<td>31</td>
<td>Black</td>
<td>1</td>
<td>2</td>
<td>5.5</td>
<td>8.5</td>
<td>6</td>
<td>F</td>
<td>32</td>
<td>Black</td>
<td>3</td>
</tr>
<tr>
<td>13</td>
<td>F</td>
<td>47</td>
<td>Black</td>
<td>3</td>
<td>5</td>
<td>0</td>
<td>9.5</td>
<td>8</td>
<td>F</td>
<td>47</td>
<td>Black</td>
<td>0</td>
</tr>
<tr>
<td>14</td>
<td>F</td>
<td>46</td>
<td>Black</td>
<td>9</td>
<td>3</td>
<td>3</td>
<td>8.5</td>
<td>6</td>
<td>F</td>
<td>46</td>
<td>Black</td>
<td>3</td>
</tr>
<tr>
<td>15</td>
<td>F</td>
<td>46</td>
<td>White</td>
<td>8</td>
<td>10</td>
<td>3</td>
<td>6.5</td>
<td>5</td>
<td>F</td>
<td>47</td>
<td>White</td>
<td>0</td>
</tr>
<tr>
<td>16</td>
<td>F</td>
<td>34</td>
<td>Black</td>
<td>10</td>
<td>3</td>
<td>7</td>
<td>8.5</td>
<td>8</td>
<td>F</td>
<td>34</td>
<td>White</td>
<td>2</td>
</tr>
<tr>
<td>17</td>
<td>M</td>
<td>26</td>
<td>White</td>
<td>0</td>
<td>1.5</td>
<td>0.5</td>
<td>3</td>
<td>2</td>
<td>M</td>
<td>27</td>
<td>White</td>
<td>0</td>
</tr>
<tr>
<td>18</td>
<td>F</td>
<td>25</td>
<td>Asian</td>
<td>9</td>
<td>0.5</td>
<td>1</td>
<td>4</td>
<td>2</td>
<td>F</td>
<td>28</td>
<td>Asian</td>
<td>9</td>
</tr>
</tbody>
</table>

BDI = Beck Depression Inventory; BPI = Brief Pain Inventory; Pain Intensity = Average self-reported pain rating before and after resting state fMRI scanning. Multi. = Multiple; Hisp. = Hispanic; SE = standard error.
asked to wait for 10 min in a comfortable position before starting the second half of the scanning session.

During the exercises, cLBP patients were intermittently asked to report their level of pain using the 0–10 pain scale; the exercises were repeated until subjects reported an increase in pain of approximately 3 points on the pain scale. Once this level of pain was achieved, subjects were placed back in the scanner to repeat the same fMRI scans that were acquired before the pain-inducing maneuvers. All maneuvers were applied slowly at a pace that the patients feel acceptable. For healthy controls, structural and resting state scans were only collected once. Before and immediately after each 6 min resting state MRI scan, subjects were asked to rate the intensity of their LBP using a 0–10 pain scale. The average self-reported pain rating before and after these maneuvers was used as an index of endogenous pain intensity.

2.3. Medication

Medication use per self-report was limited to non-steroidal anti-inflammatory drugs (NSAIDs, e.g., ibuprofen, Motrin, Advil, and Naproxen) and acetaminophen (e.g., Tylenol). Additional non-pharmacological methods of self-reported pain management included chiropractic massages, physical therapy, exercises, and acupuncture.

2.4. MRI data acquisition

All MRI data were acquired with a 3 T Siemens whole-body scanner with echo-planar imaging capability using a 32-channel radio-frequency head coil at the Martinos Center for Biomedical Imaging. During the resting state fMRI scan, subjects were asked to keep their eyes open and look at a darkened screen for 6 min. The BOLD fMRI scan acquisition included 47 slices with slice thickness of 3 mm, TR = 3000 ms, TE = 30 ms, and a 3 × 3 mm in-plane spatial resolution. T1-weighted MPRAGE structural images were acquired using the following parameters: voxel size 1.2 × 1.2 × 1.2 mm, TR = 2.2 s, TE = 1.54 ms, flip angle = 7°, slices = 144; field of view = 230.

2.5. PAG seed based functional connectivity analysis

The fMRI data were then preprocessed using SPM8 software (available at: http://www.fil.ion.ucl.ac.uk/spm) implemented in a MATLAB suite (MathWorks, Inc., Natick, Massachusetts). The first 10 volumes were not analyzed to allow for signal equilibration effects. Images were realigned to correct for motion, corrected for errors in slice timing, spatially transformed to standard stereotaxic space (based on the Montreal Neurologic Institute coordinate system). A recent study shows that the motion induced artifacts occur with movements on the order of a few tenths of a millimeter or less and produce systematic but spurious correlations in functional connectivity, such that long distance correlations are decreased by subject motion, whereas many short-distance correlations are increased (Power et al., 2012). ANOVA on each movement parameter with cLBP patients in high pain condition and in low pain condition and healthy control as between-subjects factor revealed no significant differences (P > 0.1). In Table 2, we show that the means for six movement parameters are small and there was no significant group difference on movement parameters. There were no participants with movement greater than 3 mm of translation or 3 degrees of rotation. There were also no significant differences between the total range of movement across any axis of translation or rotation between groups (see Table 2). The data were then smoothed with a 6-mm full-width half-maximum (FWHM) Gaussian kernel. The smooth kernel size was chosen because FWHM resolution usually equals or is greater than twice the voxel size (Miki et al., 2008). Data were then bandpass filtered from 0.01 to 0.08 Hz to remove low frequency noise (including slow scanner drifts) and influences of higher frequencies reflecting cardiac and respiratory signals (Cordes et al., 2001).

To address head motion concerns in resting-state fMRI analyses, we calculated the voxel-specific mean framewise displacement (FD) for
accounting head motion at group-level analysis (Power et al., 2012, 2013; Power et al., 2014; Van Dijk et al., 2012). FD measure indexes the movement of the head from one volume to the next and is calculated as the sum of the absolute values of the differentiated realignment estimates (by backward differences) at every time point (Power et al., 2012). Then, we repeated the above analyses after removing frames with FD > 0.5 mm (‘scrubbing’). One time point before “bad” time points and two time points after “bad” time points were deleted.

Functional connectivity analysis was carried out by applying a seed-region approach using the right ventrolateral PAG (x = 4, y = −26, z = −14, with 2 mm radius) as the FC seed (see Fig. 1B). The rationale for choosing this location as a seed is: 1) in a previous study, we found that increased levels of heat pain can evoke a significant fMRI signal increase in this region (Kong et al., 2010a); 2) it is located within the ventrolateral PAG, which is believed to be important for opioid antinociception (Bandler and Shipley, 1994); 3) it is consistent with previous PAG seed based FC study showing that the PAG is functionally connected to the ACC/vmPFC and the insula (Kong et al., 2010b). One concern is that signals from other regions may confound the findings due to partial volume effect. We plotted the time course from the PAG seed for all subjects and found no cyclic trends in the data (see Fig. 1C for an example). Given that the PAG seed region sits adjacent to a ventricle with significant pulsatility effect, we also used a seed from the third ventricle (x = 0, y = −3, z = −7, with 2 mm radius) as a control region.

For each seed region, individual participant analyses were carried out using the General Linear Model (GLM) with the time series for the seed region as well as for the nuisance covariates (white matter, cerebrospinal fluid, and six motion parameters) as predictors. These nuisance signals are typically adjusted for in resting-state FC studies because they reflect signal fluctuations of nonneuronal origin (e.g., physiological artifacts associated with variables such as cardiac and respiratory cycles, CSF motion, and scanner drift) (Fox and Raichle, 2007).

Contrast images were generated for each subject by estimating the regression coefficient between all brain voxels and each seed’s time series, respectively. These images were then included in second-level group random effect analyses, adopting a t-test design. We also used regression analyses to examine whether illness duration and/or endogenous pain intensity were related to PAG functional connectivity, when considering the cLBP participants alone. Endogenous pain intensity was defined as the average self-reported pain rating immediately before and after resting state fMRI scanning. The threshold was family-wise alpha level set to P = 0.05, family-wise error (FWE) correction for multiple comparisons for ROIs using small volume correction (svc). Based on the anatomical connection of PAG and previous studies (Brooks and Tracey, 2005; Fields, 2004; Heinricher et al., 2009; Ploner et al., 2010; Tracey and Mantyh, 2007), the ROIs include the vmPFC/ACC, anterior insula, posterior insula, and amygdala, derived from automated anatomical labeling (AAL, see Fig. 1C–E) (Tzourio-Mazoyer et al., 2002). A threshold of P < 0.05 family-wise error (FWE) corrected at the cluster level across the whole brain was used for non-ROI brain regions.

### 3. Results

A total of 18 cLBP patients (6 males) and 18 age- and gender-matched HCs (6 males) completed the study. Demographics, clinical assessments and characteristics for cLBP patients and HCs are presented in Table 1. The age difference between the two groups was not significant (mean ± SE 36.1 ± 2.3 in the patient group versus 37.1 ± 2.2 in the control group), P = 0.71. The duration of illness in the patient group is 5.3 ± 0.9 years. The BPI was used to measure the pain in the preceding week in the patient group, 6.5 ± 0.7.

Of the 18 patients who completed the study, one patient had strong chronic pain at baseline and thus did not perform any exercises. After lying down for 10 min, the patient felt a reduction in low back pain. The patient received the exact same set of scan procedures before and after the 10 minute rest period, comparable to the healthy control condition. This data was included in the data analysis. In this study, the self-reported endogenous LBP intensity recorded before and after resting state fMRI scanning was averaged and then used as an index of pain during scanning. After slow clinical pain-inducing maneuver, the self-reported pain intensity was significantly increased from 3.79 ± 0.58 (mean ± SE) before pain-inducing maneuver to 6.65 ± 0.46, t = 5.38, P < 0.001. In our data analysis, we define the scanning period during which patients’ self-reported pain intensity is lower as the low pain (LP) condition and the scanning period during which patients’ low back pain ratings are higher as the high pain (HP) condition.

During the LP condition (Fig. 2A) and HP condition (Fig. 2B) in the patient group and in the healthy group (Fig. 2C), we found predominantly positive correlations between PAG activity and activity in nearby structures, including the brainstem (mostly the midbrain), thalamus, parahippocampus, amygdala, and cerebellum, and with distant regions, including the anterior cingulate (not in control group) and temporal cortex. These findings were consistent with our previous PAG centered FC study applied in healthy subjects (Kong et al., 2010b) and a previous diffusion tensor imaging study on PAG (Hadjipavlou et al., 2006).

Whole brain voxel-by-voxel functional connectivity in HCs and cLBP patients during the LP condition was compared using a two-sample t-test. Results showed that cLBP patients in the LP condition had significantly greater FC between the PAG and the left vmPFC/ACC, x = −6, y = 45, z = −6, Z = 3.20, voxels = 11, P_{FWE} < 0.05 svc and right vmPFC/ACC, x = 6, y = 42, z = −12, Z = 3.18, voxels = 36, P_{FWE} < 0.05 svc, as well as other regions including the superior temporal gyrus and the precentral gyrus (Table 3 and Fig. 2D). The opposite contrast showed no FC differences above threshold. Compared with HCs, patients in the HP condition also showed enhanced FC between the PAG and the left vmPFC, x = −6, y = 42, z = −9, Z = 2.96, voxels = 11, P_{FWE} < 0.05 svc, as well as other regions including the lingual gyrus, superior temporal gyrus, precentral gyrus, dorsal cingulate cortex, and posterior insula (Table 3 and Fig. 2E). The opposite contrast showed no FC differences above threshold. When we compared cLBP patients in the HP and LP conditions, we did not find a significant difference using the prior threshold (Fig. 2).

To test the association between FC and subjective cLBP rating at rest, we applied a regression analysis between FC and low back pain ratings using the average pain rating scores in the LP condition. We believe that this measure captures the immediate pain level during scanning since participants rated their pain before and immediately after each 6 minute resting state MRI scan. The results showed no significant correlations. For the resting state scan in the HP condition, we found significant negative correlations between FC and LPB ratings in the HP condition at the left vmPFC x = −9, y = 57, z = −12, Z = 3.83, voxels = 9, P_{FWE} < 0.05 svc, and no positive correlations were found (Fig. 3A).

To test the association between functional connectivity with the PAG and cLBP illness duration, we applied a regression analysis between FC and low back pain ratings using the average pain rating scores in the LP condition. We believe that this measure captures the immediate pain level during scanning since participants rated their pain before and immediately after each 6 minute resting state MRI scan. The results showed no significant correlations. For the resting state scan in the HP condition, we found significant negative correlations between FC and LPB ratings in the HP condition at the left vmPFC x = −9, y = 57, z = −12, Z = 3.83, voxels = 9, P_{FWE} < 0.05 svc, and no positive correlations were found (Fig. 3A).

To test the association between functional connectivity with the PAG and cLBP illness duration, we applied a regression analysis between FC and low back pain ratings using the average pain rating scores in the LP condition. We believe that this measure captures the immediate pain level during scanning since participants rated their pain before and immediately after each 6 minute resting state MRI scan. The results showed no significant correlations. For the resting state scan in the HP condition, we found significant negative correlations between FC and LPB ratings in the HP condition at the left vmPFC x = −9, y = 57, z = −12, Z = 3.83, voxels = 9, P_{FWE} < 0.05 svc, and no positive correlations were found (Fig. 3A).

To test the association between functional connectivity with the PAG and cLBP illness duration, we applied a regression analysis between FC and low back pain ratings using the average pain rating scores in the LP condition. We believe that this measure captures the immediate pain level during scanning since participants rated their pain before and immediately after each 6 minute resting state MRI scan. The results showed no significant correlations. For the resting state scan in the HP condition, we found significant negative correlations between FC and LPB ratings in the HP condition at the left vmPFC x = −9, y = 57, z = −12, Z = 3.83, voxels = 9, P_{FWE} < 0.05 svc, and no positive correlations were found (Fig. 3A).

To test the association between functional connectivity with the PAG and cLBP illness duration, we applied a regression analysis between FC and low back pain ratings using the average pain rating scores in the LP condition. We believe that this measure captures the immediate pain level during scanning since participants rated their pain before and immediately after each 6 minute resting state MRI scan. The results showed no significant correlations. For the resting state scan in the HP condition, we found significant negative correlations between FC and LPB ratings in the HP condition at the left vmPFC x = −9, y = 57, z = −12, Z = 3.83, voxels = 9, P_{FWE} < 0.05 svc, and no positive correlations were found (Fig. 3A).

To test the association between functional connectivity with the PAG and cLBP illness duration, we applied a regression analysis between FC and low back pain ratings using the average pain rating scores in the LP condition. We believe that this measure captures the immediate pain level during scanning since participants rated their pain before and immediately after each 6 minute resting state MRI scan. The results showed no significant correlations. For the resting state scan in the HP condition, we found significant negative correlations between FC and LPB ratings in the HP condition at the left vmPFC x = −9, y = 57, z = −12, Z = 3.83, voxels = 9, P_{FWE} < 0.05 svc, and no positive correlations were found (Fig. 3A).

To test the association between functional connectivity with the PAG and cLBP illness duration, we applied a regression analysis between FC and low back pain ratings using the average pain rating scores in the LP condition. We believe that this measure captures the immediate pain level during scanning since participants rated their pain before and immediately after each 6 minute resting state MRI scan. The results showed no significant correlations. For the resting state scan in the HP condition, we found significant negative correlations between FC and LPB ratings in the HP condition at the left vmPFC x = −9, y = 57, z = −12, Z = 3.83, voxels = 9, P_{FWE} < 0.05 svc, and no positive correlations were found (Fig. 3A).
no significant difference between the HP and LP conditions \((P > 0.2)\) and between HP and HC or LP and HC \((P\) values > 0.2). The repeated analyses without removing frames with framewise displacement \((FD) > 0.5 \text{ mm} \) (**scrubbing**) yielded similar results (results not reported here).

4. **Discussion**

In this study, we investigated differences in the PAG centered resting state functional connectivity in cLBP patients relative to age and gender...
matched controls. We found that PAG–vmPFC/rACC functional connectivity was enhanced in cLBP patients, compared with HCs. Interestingly, we found that the functional connectivity between the PAG and the vmPFC/rACC decreased as endogenous back pain intensity increased after pain-inducing maneuver, suggesting the dynamic character of functional connectivity at the PAG. Moreover, cLBP duration was negatively correlated with PAG–posterior insula and PAG–amygdala FC before any pain-inducing maneuver. These functional changes of PAG point out that the PAG in particular may play an important role in the pathophysiology of cLBP.

4.1. PAG–vmPFC/rACC functional connectivity

Abnormal vmPFC/rACC activity has been found in cLBP in previous studies. Compared to HCs, cLBP patients showed increased high-frequency BOLD oscillations (0.12–0.20 Hz) circumscribed mainly to the vmPFC and brain regions within the default network (Baliki et al., 2011). More recently, using arterial spin labeling, it has been found that provoked increases in endogenous LBP ratings were positively associated with statistically significant increases in regional cerebral blood flow in a widespread network of cortical areas, including the bilateral vmPFC in cLBP patients (Wasan et al., 2011). In addition, compared with healthy controls, patients demonstrated stronger default mode network connectivity to the pregenual anterior cingulate cortex, the left inferior parietal lobule, and the right insula (Loggia et al., 2013). It has been shown that the prefrontal cortex exerts active control on pain perception by modulating corticosubcortical and corticocortical pathways (Lorenz et al., 2003). The current study further highlights the role of vmPFC/rACC functional connectivity abnormality in cLBP and links the vmPFC/rACC to the PAG.

Table 3
Main group difference results from group analysis cLBP patients in high pain and cLBP patients in low pain condition and healthy controls.

<table>
<thead>
<tr>
<th>Contrast</th>
<th>Voxels</th>
<th>Brain area</th>
<th>Peak coordinate (x, y, z)</th>
<th>Z value</th>
</tr>
</thead>
<tbody>
<tr>
<td>LP &gt; HC</td>
<td>36</td>
<td>R vmPFC/ACC</td>
<td>6, 42, −12</td>
<td>3.18 svc</td>
</tr>
<tr>
<td></td>
<td>11</td>
<td>L vmPFC/ACC</td>
<td>−6, 45, −6</td>
<td>3.20 svc</td>
</tr>
<tr>
<td>HC &gt; LP</td>
<td>No brain region above the threshold</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HP &gt; HC</td>
<td>2401</td>
<td>R precentral gyrus</td>
<td>39, −15, 33</td>
<td>4.68</td>
</tr>
<tr>
<td></td>
<td></td>
<td>L superior temporal gyrus/operculum</td>
<td>−36, −6, 54</td>
<td>4.17</td>
</tr>
<tr>
<td></td>
<td></td>
<td>R superior temporal gyrus/operculum</td>
<td>57, 12, −9</td>
<td>4.09</td>
</tr>
<tr>
<td></td>
<td></td>
<td>L dorsal cingulate cortex</td>
<td>−6, −9, 36</td>
<td>4.04</td>
</tr>
<tr>
<td></td>
<td></td>
<td>R posterior insula</td>
<td>48, −18, 15</td>
<td>3.86 svc</td>
</tr>
<tr>
<td></td>
<td>300</td>
<td>L lingual gyrus</td>
<td>−3, −63, 15</td>
<td>3.95</td>
</tr>
<tr>
<td></td>
<td>11</td>
<td>L vmPFC/ACC</td>
<td>−6, 43, −9</td>
<td>2.96 svc</td>
</tr>
<tr>
<td>HP &gt; HC</td>
<td>No brain region above the threshold</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LP &gt; HP</td>
<td>No brain region above the threshold</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HP &gt; LP</td>
<td>No brain region above the threshold</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HP with pain intensity</td>
<td>9</td>
<td>L vmPFC</td>
<td>−9, 57, −12</td>
<td>3.83 svc</td>
</tr>
<tr>
<td>LP with illness duration</td>
<td>3</td>
<td>L amygdala</td>
<td>−27, −9, −18</td>
<td>3.18 svc</td>
</tr>
<tr>
<td></td>
<td>19</td>
<td>R posterior insula</td>
<td>39, −12, 9</td>
<td>3.04 svc</td>
</tr>
</tbody>
</table>

LP = low pain before pain-inducing maneuver; HP = high pain after pain-inducing maneuver. HC = healthy controls. L = left, R = right. For regions of interest (ROIs), results were significant at P_{FWE} < 0.05 after small volume correction (svc). Other results were significant at P < 0.05 family-wise error (FWE) corrected at the cluster level.

![Fig. 3](image-url)Brain–behavioral correlation results. (A) The association between PAG centered functional connectivity with the mPFC in the HP condition and endogenous pain intensity in the HP condition. (B–C) The association between PAG centered functional connectivity with the insula and amygdala in the LP condition and illness duration.
can evoke a significant fMRI signal increase in the PAG (Kong et al., 2010a). Moreover, using midbrain/brainstem specific imaging, an increased PAG activity in response to pain has been demonstrated (Eippert et al., 2009). The involvement of the PAG in pain processing and modulation has been known for a long time. Brain imaging studies (Bushnell et al., 1999; Eippert et al., 2009; Erpelding et al., 2012; Kong et al., 2010a; Kong et al., 2006; Teutsch et al., 2008; Wasan et al., 2011) have found that the PAG is activated during the presentation of noxious stimuli as well as in association with pathological pain states such as chronic low back pain. Studies have shown that distraction tasks reduce the subjective pain sensation (Tracey et al., 2002; Valet et al., 2004). Activation in the periaqueductal gray was significantly increased during the distraction condition (Tracey et al., 2002; Valet et al., 2004), and the total increase in activation was predictive of changes in perceived pain intensity (Tracey et al., 2002).

The PAG also plays a role in pain facilitation. In accordance with a pain facilitation role, the magnitude of PAG activation has been shown to correlate with the degree of patients’ neuropathic pain symptoms (Freyhagen et al., 2006). In irritable bowel syndrome (IBS) patients, activity in the vmPFC has been shown to disrupt a functional connection between the lateral PFC and the PAG, suggesting that the vmPFC and the PAG are involved in enhancing clinical pain (Mayer et al., 2005). Previous studies have shown a general hypersensitivity to painful stimuli in cLPB patients (Farzsay and Meeseusen, 2005; Giescke et al., 2004; Puts et al., 2012). Thus, the enhanced PAG–vmPFC/ACC FC in cLPB patients may indicate an enhanced pain inhibition or facilitation. Further studies are needed to clarify the role of this FC change.

4.2. Relationship between PAG functional connectivity and endogenous pain

In the present study, in the vmPFC, the strength of intrinsic connectivity with the PAG was negatively correlated with pain intensity in high pain condition. The vmPFC has been implicated in regulating affective responses by manipulating the contextual evaluation of sensory events (Rolls and Grabenhorst, 2008). vmPFC activation was associated with decreases in pain unpleasantness ratings induced by mindfulness meditation in healthy subjects (Zeidan et al., 2011). In this context, we speculate that a decrease in the intrinsic correlations between the PAG and the vmPFC in relation to the pain intensity may indicate a failed pain modulation in the brain in response to worsening of pain. An alternative explanation is that both the vmPFC and the PAG reflect arousal, which is high in HP condition. Interestingly, such correlation was significant only in the HP condition in which the pain level was enhanced by clinical pain-inducing maneuvers, suggesting that the vmPFC–PAG network is more related with higher pain states. Whether the observed association between spontaneous pain intensity and changes in PAG connectivity reflects either an adaptive mechanism or an abnormal state in cortical excitability that predisposes individuals to chronic pain needs to be further clarified. Our study suggests that regionally specific FC changes within the PAG–vmPFC networks may be a new locus of dysfunction in cLPB.

4.3. Relationship between PAG functional connectivity and illness duration

We found that illness duration was negatively correlated with PAG–posterior insula and PAG–amygdala FC, only in the LP condition. A previous study has found that the prestimulus FC between the insula and the PAG predicted perceived painfulness (Ploner et al., 2010), suggesting that these two regions interact to determine pain perception. It has been reported that activation in the contralateral posterior insula was positively correlated with temperature level, whereas subjective intensity related more to activation of the right anterior insula (Craig et al., 2000). It has been suggested that the posterior insula may provide a primary ‘interoceptive cortex’, specialized for perception of internal bodily states incorporating pain, temperature, and autonomic arousal (Craig, 2003; Critchley et al., 2002). We found that the longer an individual is in the cLPB state, the weaker the functional connectivity between the PAG and the posterior insula. We speculate that this may suggest that after long-term cLPB suffering, the body is adapted to the situation, and thus the modulation mechanism is somehow weakened. A previous study also found that verum acupuncture induced a higher level of correlations among the amygdala-associated network including the insula and the PAG (Qin et al., 2008), suggesting that this network may be involved in pain modulation. However, it is currently unknown whether altered FC with the PAG is the consequence of or the cause of cLPB. A longitudinal study would be required to determine the order of events. Interestingly, we did not observe significant correlation between PAG–insula FC after pain-inducing maneuver and pain duration in patients. It is possible that unlike in the natural low pain condition, the PAG–insula FC was altered in high pain condition.

4.4. Heterogeneity in pain modulation system across pain conditions

In a recent study of migraine pain patients (Mainiero et al., 2011), investigators have found that patients had greater FC between the PAG and the ventral prefrontal cortex compared to the control group. Migraineurs who develop pain in response to normally innocuous stimulation (i.e., migraineurs with a history of allodynia) exhibited decreased FC with the PAG in the insula and the mPFC, compared with migraineurs without allodynia. This result is consistent with findings observed in LPB patients. In another study on a fibromyalgia patient, investigators have found that patients with fibromyalgia (a chronic pain disorder characterized by chronic widespread pain and allodynia (a heightened and painful response to pressure)) display less functional connectivity between the ACC and the PAG (Jensen et al., 2012). We speculate that this may reflect the heterogeneity of different chronic pain conditions. Taken together, these findings may suggest that different roles of pain descending control system underlie the localized chronic pain (e.g., LPB, migraine) and widespread pain (e.g., fibromyalgia pain).

4.5. Study limitations

There are some potential limitations in this study worth mentioning. The first potential limitation is the order effects between the high endogenous LBP and low endogenous LBP conditions. One challenge of cLPB studies is that once LBP is provoked, it is hard to control without any pharmacological intervention. In this study, we used exercise to provoke the patients’ LBP; thus, the high pain condition tended to follow the low pain condition. Secondly, our fMRI data was not acquired with cardiac-gating which minimizes physiological motion artifact due to pressure wave pulsatility in arteries within and around the brain (Napadow et al., 2009). Future studies may compare the fMRI data with and without cardiac gating. Third, a control for the movement exercises corresponding to the pain-inducing maneuvers is missing since healthy controls underwent only one resting state session. The goal of clinical maneuvers is to induce pain in patients with cLPB. The pain-inducing maneuvers include a set of slow movements such as sit-ups, lumbar flexion/extension, and lumbar rotation. These daily life activities are not intense for healthy controls, although they can elicit pain in patients with cLPB. Thus, it is unlikely that movement exercises in healthy controls would evoke low back pain or induce functional connectivity changes in pain modulation network since there would be no pain expected in healthy controls. Previous studies also indicate that the intrinsic resting state functional connectivity is reliable across different sessions (Birn et al., 2013; Liao et al., 2013; Sheliadz et al., 2009). Nevertheless, the contrast between HP and LP conditions should be interpreted with caution. Fourth, the seed region is relatively small. One concern is that the signal to noise ratio (SNR) might not be good enough to detect PAG activity. However, previous resting state MRI studies have successfully used small seed regions, such as amygdala subregions (Roy et al., 2009), nucleus accumbens (Cauda et al., 2011), and red nucleus (Nioche et al., 2013).
2009), and found valid FC results. Finally, we did not include medication in the model, and although it is unlikely that medication exposure accounts for our results, we cannot completely rule out the effect of medication. It is important to note, however, that we excluded all patients using opioids in the study, as a previous study found that daily oral administration of morphine for 1 month can cause anatomical changes in the brain (Younger et al., 2011).

In summary, the present study showed that cLB patients have increased PAG–vmPFC FC and that the FC between the PAG and the vmPFC decreases as endogenous pain intensity increases in high pain condition. These findings may not only deepen our understanding of pain modulation and the development of chronic pain but also ultimately help inform mechanism-based therapies for treating different types of acute and chronic pain.

Conflict of interest
There is no conflict of interest to claim for all authors.

Acknowledgments
This work was supported by R01AT006364 (NCIAM) to Jian Kong, R01AT005280 (NCAM) to Randy Gollub, P01-AT02048 (NCAM) to Bruce Rosen, M01-RR-01066 and UL1 RR02575-01 for Clinical Research Center Biomedical Imaging Core from National Center for Research Resources (NCRR), and P41RR14075 for Center for Functional Neuroimaging Technologies from NCRR.

Appendix A. Supplementary data
Supplementary data to this article can be found online at http://dx.doi.org/10.1016/j.neuroimage.2014.08.019.

References

Acknowledgments
This work was supported by R01AT006364 (NCIAM) to Jian Kong, R01AT005280 (NCAM) to Randy Gollub, P01-AT02048 (NCAM) to Bruce Rosen, M01-RR-01066 and UL1 RR02575-01 for Clinical Research Center Biomedical Imaging Core from National Center for Research Resources (NCRR), and P41RR14075 for Center for Functional Neuroimaging Technologies from NCRR.

Appendix A. Supplementary data
Supplementary data to this article can be found online at http://dx.doi.org/10.1016/j.neuroimage.2014.08.019.

References

Acknowledgments
This work was supported by R01AT006364 (NCIAM) to Jian Kong, R01AT005280 (NCAM) to Randy Gollub, P01-AT02048 (NCAM) to Bruce Rosen, M01-RR-01066 and UL1 RR02575-01 for Clinical Research Center Biomedical Imaging Core from National Center for Research Resources (NCRR), and P41RR14075 for Center for Functional Neuroimaging Technologies from NCRR.

Appendix A. Supplementary data
Supplementary data to this article can be found online at http://dx.doi.org/10.1016/j.neuroimage.2014.08.019.

References

Acknowledgments
This work was supported by R01AT006364 (NCIAM) to Jian Kong, R01AT005280 (NCAM) to Randy Gollub, P01-AT02048 (NCAM) to Bruce Rosen, M01-RR-01066 and UL1 RR02575-01 for Clinical Research Center Biomedical Imaging Core from National Center for Research Resources (NCRR), and P41RR14075 for Center for Functional Neuroimaging Technologies from NCRR.