KIBRA Polymorphism Is Associated with Individual Differences in Hippocampal Subregions: Evidence from Anatomical Segmentation using High-Resolution MRI

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The KIBRA gene has been associated with episodic memory in several recent reports; carriers of the T-allele show enhanced episodic memory performance relative to noncarriers. Gene expression studies in human and rodent species show high levels of KIBRA expression in the hippocampus and whether specific subfields are differentially affected by KIBRA genotype. High-resolution magnetic resonance imaging (T2-weighted, voxel size = 0.4 × 0.4 mm, in-plane) was used to manually segment hippocampal cornu ammonis (CA) subfields, dentate gyrus (DG), and the subiculum as well as adjacent medial temporal lobe cortices in healthy carriers and noncarriers of the KIBRA T-allele (rs17070145). Overall, we found that T-carriers had a larger hippocampal volume relative to noncarriers. The structural differences observed were specific to the CA fields and DG regions of the hippocampus, suggesting a potential neural mechanism for the effects of KIBRA on episodic memory performance reported previously.

Introduction

Several recent studies have identified genetic variations and molecular pathways that are associated with episodic memory performance (for review, see Papassotiropoulos and de Quervain, 2011), which is the type of memory that enables recollection of specific events (Tulving, 2002). In the first genome-wide association study of human memory, an association between a single nucleotide polymorphism (SNP) of the kidney and brain regulator (KIBRA) gene (C→T) and episodic memory was first identified (Papassotiropoulos et al., 2006). In three cohorts, T-allele carriers (TT/TC) of the rs17070145 SNP showed an advantage in episodic memory performance relative to noncarriers (CC). This finding has been replicated several times, although there have been null findings (for a meta-analysis, see Milnik et al., 2012).

Given the well documented role of the hippocampus in episodic memory (Eichenbaum et al., 2007), the goal of the present study was to determine whether the KIBRA C→T polymorphism is also associated with volumetric differences in the human hippocampus and whether specific subfields are differentially affected by KIBRA genotype. KIBRA expression is high in the hippocampus in both humans and rodents (Papassotiropoulos et al., 2006; Johannsen et al., 2008; Yoshihama et al., 2009). Moreover, KIBRA-dependent differences in blood oxygenation level-dependent (BOLD) response in the hippocampus and medial temporal lobe (MTL) cortices have been observed during episodic remembering (Papassotiropoulos et al., 2006; Kauppi et al., 2011), although these studies actually yielded opposite results (i.e., increased activation in noncarriers vs increased activation in T-carriers, respectively). Yet, null genotype differences in whole hippocampal and MTL cortex volume have been observed in one report that used both manual and automated magnetic resonance imaging (MRI) segmentation (Papassotiropoulos et al., 2006). However, the hippocampus consists of histologically heterogeneous subregions (i.e., cornu ammonis (CA) fields 1–3, dentate gyrus (DG), subiculum) with distinct patterns of connectivity and cellular structure (Amaral and Insausti, 1990; Duvernoy, 2005). Papassotiropoulos et al. (2006) demonstrated that KIBRA is most highly expressed in the CA1 and DG in rodents; likewise Johannsen (2008) observed intense KIBRA expression in DG granule cells and CA pyramidal cells in rodents (also see Yoshihama et
al., 2009), raising the possibility that whole-hippocampal volume may be a less sensitive marker to assess KIBRA-dependent structural variability in humans. To assess regional differences in the effects of KIBRA genotype on hippocampal structure, we conducted MRI-based volumetric analyses of hippocampal and adjacent MTL cortical subregions. We hypothesized that KIBRA-dependent differences in hippocampal volume would be localized within the hippocampal CA fields and DG, parallelizing the expression patterns observed in rodents.

Materials and Methods

Participants. Thirty-two healthy Caucasian individuals, stratified by KIBRA genotype (T-carriers and noncarriers), were recruited from a larger study investigating genetics and memory. Genotype groups were matched on genetic characteristics previously implicated in episodic memory: apolipoprotein E (APOE) ε4 alleles, brain-derived neurotropic factor (BDNF) Val66Met, and catechol-O-methyltransferase (COMT) Val158Met. To complement methodology used by Papassotiropoulos et al. (2006), participants were also one-to-one matched (within 0.4 SDs) between genotype groups on old/new visual recognition memory performance as assessed within 2 years of scanning (for task details, see Rudebeck et al., 2009). Due to equipment failure, imaging data were unavailable for four noncarriers. Thus our sample included 18 T-carriers (22.2 ± 3.7 years old; 15.5 ± 2.8 years of education; 4 male) and 14 noncarriers (20.3 ± 3.0 years old; 13.9 ± 1.6 years of education; 3 male). This did not substantively affect the balance between genotype groups on the aforementioned variables: APOE ε4 alleles (p = 0.37); BDNF Val66Met (p = 0.77); or COMT Val158Met (p = 0.96; χ² tests). Genotype groups also did not significantly differ in visual recognition memory scores (p = 0.78, hp² = 0.003). While it is acknowledged that performance differences attributable to genotype are unlikely to be detected in this sample size, our results may nonetheless help to explain findings from other larger studies where performance effects are observed.

Given previous research that sex modifies the relationship between KIBRA and cognition (Wersching et al., 2011), sex was included as a covariate in all analyses, although this did not change the pattern of results. There was no significant difference in age between groups (p = 0.13, hp² = 0.08). There was a marginal group difference in education (F₁,29 = 3.65, p = 0.07, hp² = 0.11). As education was not found to be associated with the volume of any regions of interest (ROIs) within genotype group, it was not included as a covariate. This study was approved by local ethics. Participants provided written informed consent and were compensated $50 for the MRI.

Genotyping. Participants provided a saliva sample (~2 ml) in an Oragene OG-500 DNA kit (DNA Genotek). Five SNPs across four genes were genotyped using a TaqMan predesign assay; COMT Val158Met (rs4818); BDNF Val66Met (rs243853); KIBRA (rs17070145); and both the ApoE 112 (rs429358) and 158 polymorphism (rs7412; Life Technologies). For each reaction, 20 ng of genomic DNA was amplified and scaled to a total volume of 10 μl in an Applied Biosystems system 2720 thermal cycler. Post amplification products were analyzed on the ABI Prism 7500 Sequence Detection System using the allelic discrimination option and genotype calls were determined manually by comparison to six No Template Controls. For the ApoE markers, the 112 and 158 genotypes were combined to determine participants’ ApoE α diplotyp. Genotyping of 10% of the samples was replicated for quality control with no discrepancies. Following previous studies (Mihnik et al., 2012) we combined all individuals carrying the T-allele (i.e., TT/TC vs CC).

MRI acquisition. Structural images were acquired using a 3 T Siemens Trio scanner. For segmentation, high-resolution T1-weighted images were acquired in an oblique-cortical plane; slices were arranged perpendicularly to the long axis of the hippocampus (TE/TR = 68 ms/3000, 2–28 oblique-cortical slices depending on head size, 512 × 512 acquisition matrix, voxel size = 0.43 × 0.43 × 3 mm, no skip, FOV = 220 mm). The first slice was placed slightly anterior to the collateral sulcus; the last slice was placed just before the full superior–inferior extent of the ventricles. To confirm the placement of slices according to these boundaries, a whole-brain anatomical MRI was first acquired, using a 3D T1-weighted MP-RAGE (TE/TR = 2.63 ms/2000 ms, 176 oblique-axial slices, 192 × 192 acquisition matrix, voxel size = 1 mm³, FOV = 256 mm). The T1-weighted images were also used to obtain a measure of total brain volume (TBV; see below).

Segmentation. ROI segmentation was completed in participants’ native space (coronal plane; Fig. 1). Akin to most previous studies (Mueller et al., 2007; Olsen et al., 2009, 2013; Chen et al., 2011) subregions were not segmented across the entire long axis of the hippocampus. Instead, segmentation was completed only within the middle section, where the dark bands separating hippocampal layers are clear. This corresponded mainly to the body of the hippocampus; however, we also demarcated the most posterior head slices, where the subfields can also be differentiated (Carr et al., 2010; Chen et al., 2011) (for an alternative approach strictly excluding head slices see Mueller et al., 2007). We segmented CA1, subiculum, and DG/CA3, with the latter grouped into a single ROI, as these regions cannot be reliably delineated at this resolution. Combined ROIs (i.e., without subjfield demarcation) were used to define the remaining head and tail hippocampal slices, which afforded a measure of full hippocampal volume. Outside the hippocampus, the MTL cortices were segmented: entorhinal cortex (ERC), perirhinal cortex (PRC), and parahippocampal cortex (PHC).

Segmentation was performed in FSLview (v3.1.2) by a single rater, blind to group status (Fig. 1). Segmentation was guided by standard anatomical guidelines (Amaral and Insausti, 1990; Insauti et al., 1998; Duvernois, 2005; Fig. 1) following the procedures described for MTL subregion segmentation for hi-resolution functional MRI (SMI) by Olsen et al. (2009); which are similar to those used to assess subregion volumetrics (Mueller et al., 2007; Olsen et al., 2013).

Intrarater and inter-rater reliability were established by comparing segmentations of five participants’ brains segmented twice by the same rater (repeated with a 2–6 month interval) and to that of a second rater (Table 1). Reliability was assessed using the Dice overlap metric (Dice, 1945), which was computed for each ROI within each hemisphere (= 2 (intersected region)/(region A + region B); 0 = no overlap; 1 = perfect overlap). Dice values were comparable to those reported previously (Bonnicci et al., 2012). TBV estimates were acquired from the T1-weighted images using an adapted version of the ANIMAL algorithm (Collins et al., 1995). Genotype groups differed marginally in TBV (F₁,29 = 3.00, p = 0.09, hp² = 0.09); noncarriers had a larger TBV relative to T-carriers. TBV was accounted for in each ROI using a regression-based technique; each ROI was regressed on TBV (collapsed across groups); and the residual value (i.e., the structure’s actual size minus its predicted value based on the individual’s TBV) was accounted for in each ROI for each individual (Arndt et al., 1991). All analyses reported below were performed on TBV-adjusted ROI values.

Results

Group differences in MTL subregion volumes (TBV adjusted) were assessed with three mixed-design ANCOVAs (i.e., whole unsegmented hippocampus, segmented hippocampus, segmented MTL cortex) and post hoc tests. ANCOVAs included KIBRA genotype as a between-subjects factor and laterality and ROI as within-subjects factors (ROI modeled only for the latter two ANCOVAs). All models included sex as a covariate, as noted

| Table 1. Dice reliability values for intrarater (intra) and inter-rater reliability (inter) |
|---|---|---|---|---|
| | Intra | | Inter | |
| | L | R | L | R |
| CA₁ | 0.85 | 0.86 | 0.76 | 0.70 |
| DG/CA₃ | 0.89 | 0.90 | 0.84 | 0.81 |
| Sub | 0.80 | 0.77 | 0.70 | 0.67 |
| PRC | 0.84 | 0.88 | 0.73 | 0.77 |
| ERC | 0.83 | 0.85 | 0.69 | 0.72 |
| PHC | 0.92 | 0.91 | 0.78 | 0.83 |

L, left; R, right.
Figure 1. Slices from T2-weighted (0.4 × 0.4 mm) images through the MTLs for one representative participant. Left, Depicts a 3D rendering of the hippocampus. Middle and Right, Show coronal slices through MTL. Subfields were drawn where a clear “C-shape” was discernible, which included all of the body and extended into the most posterior head slices (Zeineh et al., 2000; Olsen et al., 2009; the remaining head slices and the entire tail included all subfields). Demarcation varied across the long axis (Amaral and Insausti, 1990). Anteriorly, the lateral (superior) boundary of CA1 was drawn by bisecting the most lateral undulation of hippocampus (i). Moving posteriorly, the CA1 was drawn 3/4 of the way up the lateral bend of hippocampus (ii), and its medial extension bisected the DG/CA2/3 regions (iii). In the most posterior slices of the body, CA1 was drawn 3/4 of the way up the lateral bend of the hippocampus, and its medial extension was drawn in line with the medial extent of the “tear-drop”-shaped DG/CA2/3 (iv). Regions extending superior and medial to the CA1 were taken as DG/CA2/3 (Zeineh et al., 2000). Anteriorly, the medial portion of the subiculum (sub) extended until the elbow of the isthmus (v) and in more posterior slices the medial subicular border was drawn halfway down the bend of the isthmus (vi). PRC, ERC, and PHC were segmented according to Insausti et al. (1998). R, right; S, superior; A, anterior; P, posterior.
Table 2. Mean volumes (mm$^3$) corrected for TBV are shown for each genotype group for each ROI, hippocampal subregions

<table>
<thead>
<tr>
<th>CA$^+$</th>
<th>Hippocampal subregions</th>
<th>Sub</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CC</td>
<td>TT/TC</td>
</tr>
<tr>
<td>L</td>
<td>594 (18)</td>
<td>659 (15)</td>
</tr>
<tr>
<td>R</td>
<td>595 (29)</td>
<td>692 (26)</td>
</tr>
</tbody>
</table>

Volumes are shown for the left (L) and right (R) hemispheres. Standard error of the mean is shown in parentheses.

Table 3. Mean volumes (mm$^3$) corrected for TBV are shown for each genotype group for each ROI, MTL cortices

<table>
<thead>
<tr>
<th>PRC</th>
<th>MTL cortices</th>
<th>ERC</th>
<th>PHC$^+$</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CC</td>
<td>TT/TC</td>
<td>CC</td>
</tr>
<tr>
<td>L</td>
<td>2904 (159)</td>
<td>3254 (254)</td>
<td>1100 (50)</td>
</tr>
<tr>
<td>R</td>
<td>2611 (163)</td>
<td>2647 (170)</td>
<td>1035 (51)</td>
</tr>
</tbody>
</table>

Within MTL cortex, ERC, PRC, and PHC were segmented. *p < 0.05.

Discussion

We investigated the effects of KIBRA genotype on hippocampal and MTL volume using structural imaging. We provide preliminary novel evidence that the KIBRA T-allele is specifically associated with larger CA1 and DG/CA2/3 volume in young adults, suggesting a putative neural mechanism for the effects of KIBRA genotype on episodic memory reported in the literature (Milnik et al., 2012). In terms of subregion specificity, rodent work suggests functional dissociation within the hippocampus, with CA1 implicated in late retrieval and consolidation processes, while DG/CA2/3 is implicated in encoding and early retrieval (for review, see O’Reilly and Rudy, 2001; Rolls and Kesner, 2006). Similar differentiation has been demonstrated in healthy older adults (Yassa et al., 2011) and clinical populations (Mueller et al., 2011, 2012; Kerchner et al., 2012). However, because these studies examined older adults and patients, it is not known whether they extend to young, healthy individuals.

Interestingly, Papassotiropoulos et al. (2006) observed effects of KIBRA on delayed memory in all three cohorts (ranging from 5 min to 24 h), yet no effects on immediate recall, aligning the effect with the functioning of CA1. A similar pattern was observed by Bates et al. (2009) in the largest sampled KIBRA study (>2000 subjects). The interpretation was that KIBRA is not important for processes related to early memory formation, but instead relates to consolidation or delayed retention. These findings corroborate well with the present observation that KIBRA-related differences were evident in this subfield and with prior reports demonstrating high levels of expression in CA1 (Papassotiropoulos et al., 2006; Johannsen et al., 2008). Yet, other studies have observed KIBRA-related performance differences at immediate recall only (Schaper et al., 2008) or both retention intervals (Vassos et al., 2010), suggesting the effects of KIBRA on memory are nuanced.

We found no significant effects of KIBRA genotype on MTL cortex volume, suggesting that the effects of KIBRA genotype on episodic memory previously reported in the literature may be driven by differences in the hippocampus proper. However, fMRI studies have also implicated a major role of the MTL cortex in episodic memory. For example, PRC and PHC are held to encode item and context information, respectively, while the hippocampus integrates this information (Diana et al., 2007). Since contextual information is critical for recollection, PHC is thought to be especially important for this process. Moreover, both fMRI studies of KIBRA (Papassotiropoulos et al., 2006; Kauppi et al., 2011) observed group differences in activation that extended into the MTL cortices, although the specific localization is unclear.

To address this issue, we performed exploratory post hoc analyses, which revealed no group differences for PRC ($p = 0.44$, $h^2 = 0.02$) or ERC ($p = 0.84$, $h^2 = 0.001$), while PHC volume was slightly larger in T-carriers relative to noncarriers, with a marginally significant group difference ($F_{1,29} = 3.77$, $p = 0.06$, $h^2 = 0.12$). This subtle difference in volume may either reflect direct effects of KIBRA on PHC or a downstream neuroplastic effect resulting from KIBRA’s effects on the hippocampus and associated mnemonic advantage in T-carriers observed previously. While Papassotiropoulos et al. (2006) reported that the highest levels of KIBRA expression were in the hippocampus and the temporal lobes (encompassing the entire temporal lobe and hippocampus) in humans, the specific localization of expression within the temporal lobes was not examined.

An important caveat is that our protocol does not differentiate between CA1 and CA2/3. Hence, it is possible that we diluted specific effects of one of these subregions. Akin to previous investigations (Zeineh et al., 2000; Mueller et al., 2007; Olsen et al., 2009; Das et al., 2012) we used relatively thicker slices resulting in anisotropic voxels, motivated, in part, by the hippocampal atlastes used for landmark demarcation that employ ~2–4 mm thick slices (Amaral and Insausti, 1990; Duvernoy, 2005). Moreover, the layered structure of the hippocampus is best appreciated coronally, where the resolution was high. Nonetheless, given the resolution, our segmentation scheme was limited to the hippocampal body and most posterior head slices. Accordingly, future studies using higher resolution to segment across the entire hippocampal long axis are needed to assess the specificity of the KIBRA-subregion association reported here and confirm these preliminary findings. Complementary data from human histological studies of KIBRA are also required.
Given differences in MTL segmentatation reported in the literature, methodological factors cannot be ruled out as contributing to the results. Indeed, we observed KIBRA effects on whole hippocampus, while Papassotiropoulos et al. (2006) failed to observe effects at a similar resolution. Likewise, the subregion-specific findings reported here may be due to the segmentation scheme adopted and, therefore, may not be observed with a different labeling approach used by other researchers. Additionally, the null effect of KIBRA genotype on subiculum volume in the present study may have been related to the difficulty in demarcating this structure, where reliability was low relative to the other regions, a finding reported previously (Mueller et al., 2007; Bonnici et al., 2012). Finally, while subfield protocols rely heavily on the use of atlases to determine the placement of landmarks, for reliability, arbitrary landmarks are also sometimes used as a “best guess” in situations where certain borders are less clear, representing a trade-off between validity and reliability. This represents a challenge in this field that may be aided by future studies mapping subregion segmentation from postmortem (ex vivo) MRI onto that of in vivo imaging.

Another limitation is that our sample was predominantly female. Previous research has shown that sex modifies the relationship between KIBRA and cognition with larger genotype effects in females (Wersching et al., 2011). Given the small number of males, we were not able to address this issue, although covarying sex in the analysis did not alter the pattern of results. KIBRA-related differences were observed in the absence of behavioral differences, as groups were deliberately performance matched. Likewise, two previous fMRI studies have reported KIBRA-related differences in hippocampal and MTL cortex BOLD response, in the absence of performance differences (Papasotriopoulos et al., 2006; Kauppi et al., 2011; due to matching). The purpose of performance matching is to ensure that neural differences are not driven by behavioral effects per se (Rasch et al., 2010). As noted by Kauppi et al. (2011), this approach does not necessarily eliminate all behavioral differences; accordingly, the genotype-dependent differences in brain structure or BOLD response may reflect qualitative differences in the integrity of the memory trace (e.g., deeper encoding, greater vividness), which are not necessarily captured by the particular task used for performance matching.

Identifying a putative neural mechanism for KIBRA’s effects on human memory may have clinical utility. Recent research suggests that the hippocampal subfields are differentially associated with various neurological conditions. For example, CA1 is an early target site of pathology in Alzheimer’s disease (AD; Kerchner et al., 2012). Moreover, a recent meta-analysis also suggests an association between KIBRA and risk of AD (Burgess et al., 2011) although the findings have been somewhat mixed with respect to the direction of the allelic-specific effect (Hayashi et al., 2010). Future studies are needed to explore further the complex relationship between KIBRA, cognition, and disease-related neuropathology within the hippocampus. It would also be useful to examine hippocampal volume longitudinally, to determine whether the effects of KIBRA accumulate with age.

References


