Hippocampal Transcriptomic Profiles: Subfield Vulnerability to Age and Cognitive Impairment

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The current study employed next-generation RNA sequencing to examine gene expression differences related to brain aging, cognitive decline, and hippocampal subfields. Young and aged rats were trained on a spatial episodic memory task. Hippocampal regions CA1, CA3, and the dentate gyrus were isolated. Poly-A mRNA was examined using two different sequencing platforms, Illumina, and Ion Proton. The Illumina platform was used to generate seed lists of genes that were statistically differentially expressed across regions, ages, or in association with cognitive function. The gene lists were then retested using the data from the Ion Proton platform. The results indicate hippocampal subfield differences in gene expression and point to regional differences in vulnerability to aging. Aging was associated with increased expression of immune response-related genes, particularly in the dentate gyrus. For the memory task, impaired performance of aged animals was linked to the regulation of Ca^{2+} and synaptic function in region CA1. Finally, we provide a transcriptomic characterization of the three subfields regardless of age or cognitive status, highlighting and confirming a correspondence between cytoarchitectural boundaries and molecular profiling.

Keywords: aging, hippocampus, cognitive function, transcription, gene expression, Illumina HiSeq, Ion proton

INTRODUCTION

Regional variation in brain aging may explain differential susceptibility to impairment in specific cognitive domains over the course of aging. Deficits in hippocampal-dependent episodic memory emerges in middle-age, and the degree or propensity for impairment increases with advancing age in humans and animal models (Uttl and Graf, 1993; Ronnlund et al., 2005; Cansino, 2009; Foster, 2012). There has been a notable increse in recent efforts to accurately identify boundaries that define distinct hippocampal subfields in humans (Yushkevich et al., 2015; Wisse et al., 2016; Berron et al., 2017) and to link behavioral impairments in older individuals to subfield-specific hippocampal volume or activity changes (Mueller and Weiner, 2009; Yassa et al., 2011; Daugherty et al., 2016). Computational approaches, as well as research from brain damanged humans...
and animal models, suggest that the different subfields of the hippocampus (CA1, CA3, and dentate gyrus-DG) contribute to various aspects of episodic memory (Rolls and Kesner, 2006; Goodrich-Hunsaker et al., 2008; Maass et al., 2014; Wang and Diana, 2016). This includes pattern separation and completion mechanisms, rapid acquisition, and retention across different time scales. In addition, each region may employ different mechanisms to modulate the strength of their synaptic communication to alter connectivity during the formation of episodic memories. Furthermore, the three regions exhibit differential vulnerability to the aging process (Jackson et al., 2009; Wang et al., 2009; Zeier et al., 2011; Chapman et al., 2012). To address the complexity of the molecular mechanisms linked to age-related changes in hippocampal-dependent cognition, it is necessary to consider the variability in region-specific responses to aging and memory mechanisms.

The current study investigated changes in gene expression within each of the three hippocampal regions in relation to both aging and to cognitive decline in aging. Previous work suggests that brain aging is associated with altered genes and proteins linked to biological processes for cell maintenance and repair, reactions to oxidative stress, neuroinflammation, and neuron-specific signaling, including cell excitability and synaptic plasticity (Blalock et al., 2003; Lu et al., 2004; Fraser et al., 2005; Wang et al., 2009; Burger, 2010; Zeier et al., 2011; Cribbs et al., 2012; Haberman et al., 2013; Masser et al., 2014). It is unclear; however, which biomarkers may represent causative factors for cognitive decline, compensation to maintain cognitive function, or epiphenomenon associated with the aging process (Gray and Barnes, 2015). For example, studies that examine gene expression soon after behavioral testing indicate a decrease in expression of neural activity-related genes; however, it is unclear if altered expression represents the cause of the age-related deficit, or whether the genes are not expressed because learning did not occur (Rowe et al., 2007; Penner et al., 2011; Chawla et al., 2013). Similarly, brain aging can be characterized by an increase in molecular markers of oxidative stress and neuroinflammation (Foster, 2006; Droge and Schipper, 2007; Craft et al., 2012) and it is unclear whether such changes represent precursors to cognitive decline, age-related factors independent of cognition, neuroprotective elements in cognitively unimpaired individuals, or compensation due to changes in afferent connections.

Next-generation sequencing technology is a powerful tool for examining complex processes by monitoring the parallel expression of tens of thousands of genes. Along with the advantage of examining large numbers of transcripts, comes the necessity for validation of the differentially expressed genes. Normally this would occur using a different technique, such as qRT-PCR. We have taken advantage of two different next-generation platforms to confirm differential expression. Our study was designed to increase experimental robustness by utilizing two different laboratories to perform the behavioral characterization of the animals and by using two different next-generation sequencing chemistries on the same RNA samples to minimize false positives and negatives in a staged discovery/validation approach. In this case, transcriptome analysis of the same tissue was performed using Illumina fluorescent labeling and Ion Proton semiconductor measures of proton release during sequencing-by-synthesis. Each technology has pros and cons (Jessri and Farah, 2014); however, previous reports indicate a good concordance in the accuracy of expression profiling between the different platforms (Kusko et al., 2014; Li et al., 2015; Reuter et al., 2015). This study was designed to lead to a high probability of independent validation in other laboratories.

**METHODS**

**Animals**

Procedures involving animal subjects have been reviewed and approved by the Institutional Animal Care and Use Committee at the University of Arizona (AZ) and University of Florida (FL) and were in accordance with guidelines established by the U.S. Public Health Service Policy on Humane Care and Use of Laboratory. Male Fischer 344 rats of two ages, young (5–6 months, total n = 10; n = 5 AZ, n = 5 FL) and aged (17–22 months, total n = 24; n = 11 AZ, n = 13 FL) were obtained from National Institute on Aging’s colonies (Taconic, FL; Charles River, AZ). Animals were maintained on a 12:12 h light/dark schedule, and provided *ad libitum* access to food and water. Animal characterization was performed at two sites—University of Arizona and University of Florida—and the resulting data were combined and analyzed as a single cohort.

**Morris Water Maze**

Procedures for the water maze have previously been published (Foster et al., 1991; Bean et al., 2015). Animals were trained in a tank, 1.7 m (FL), 1.8 m (AZ) in diameter, positioned in room that provided cues for the animals. Water (27 ± 2°C) was maintained at a level approximately 8 cm below the surface of the tank. For cue and spatial tasks, training consisted of five blocks with three trials per block and training on each task was massed into a single day. Inter-trial intervals were 20 s and inter-block intervals were ~15 min. Rats remained on the platform between trials and in home cages under the heat lamp after each block. Behavioral data were acquired with either Noldus EthoVision computer tracking software (Noldus Information Technology, (Leesburg, VA) in FL or AnyMaze (Wood Dale, IL) in AZ and included path-length and time in the goal and opposite quadrants.

Rats were first trained on the cue discrimination version of the water escape task. The escape platform was extended approximately 1 cm above the water level and its location was signaled by a visual cue. For each trial, the platform position and start location were randomized. If an animal did not escape the water maze within 60 s, the rat was gently guided to the platform. Three days following cue training, animals were trained on the spatial discrimination task. For spatial discrimination, the escape platform was hidden approximately 1.5 cm beneath the water level and remained in the same location relative to the distal cues in the room for the duration of the initial spatial training. Fifteen minutes following the end of training on block 5, a free-swim probe trial was administered as a measure of learning. For the probe trial, the platform was removed and the animal placed in the tank for 60 s. A spatial discrimination index was computed.
according to the formula \((G − O)/(G + O)\) where \(G\) and \(O\)
represent the percent of time spent in the goal quadrant and
quadrant opposite the goal, respectively.

**Statistical Analysis of Behavior**
The mean distance to find the platform during the first training
block was used as a baseline for the water maze cue and
spatial tasks and the mean percent change from the baseline
was calculated for each subsequent block. Repeated measures
analyses of variance (ANOVA) were used to examine age
and training effects over blocks of trials. One-way ANOVA
were used to examine age effects for the water maze probe
trial discrimination index. Fisher’s protected least significant
difference comparisons, with the \(p\)-value set at 0.05, were used
as post hoc tests to localize differences. Pearson’s correlations
were calculated between the distance measures for the last block
of cue and spatial discrimination trials in order to determine
if differences in sensory-motor function or procedural learning
contributed to spatial learning. Correlations were limited to
within each age group.

**Tissue Collection**
Two weeks following water maze testing, rats were briefly
anesthetized using isoflurane (Piramal Healthcare) and then
quickly decapitated. The brains were rapidly removed, briefly
rinsed in sterile saline, and the hippocampus was dissected out
using a microspatula, razor blade, and surgical scissors. Methods
for dissection of hippocampal subfields was adapted from
previous work, which enabled isolation of CA1, CA3, and the
dentate gyrus (DG) along the rostral and ventral hippocampus.

**RNA Isolation**
The tissue was sent to University of Alabama at Birmingham
for RNA isolation. Total RNA was extracted (Qiagen, miRNeasy
#217004), DNase-treated (Qiagen), and quality was assessed
(Bioanalyzer, Agilent), which indicated a RNA integrity of 8.83
\(±\) 0.04 (mean \(±\) standard error) for the cohort.

**Ion Proton Sequencing and Analysis**
Sequencing and analysis by using Ion Proton was performed at
the University of Florida. Poly-A selection for the Ion Proton
sequencer was performed with 250 ng of total RNA using the
Dynabeads mRNA DIRECT Micro kit (Thermo Fisher, catalog
number 61021) followed by library preparations with the Ion
Total RNA-seq Kit v2 (Thermo Fisher, catalog number 4475936)
with the addition of the Ion Xpress barcodes for multiplex
sequencing (Thermo Fisher, catalog number 4475485). In brief,
poly-A selection was performed by the base pairing of the poly-
A tail of mRNA to the oligo (dT)\(_{25}\) sequence of the magnetic
Dynabeads. Further, the mRNA was enzymatically fragmented by
RNase III, purified, ligated to the Ion adaptor mix, and reverse
transcribed. The cDNA was uniquely barcoded per biological
replicate and amplified with 16 cycles of PCR. The concentration
of the libraries was quantified by the Qubit dsDNA HS Assay
(Thermo Fisher, catalog number Q32851), and size distribution
was evaluated with the High Sensitivity D1000 ScreenTape in
the 2200 Tapestation system (Agilent Technologies).

Template preparation was performed in the Ion Chef system
and sequencing was completed in the Ion Proton (Thermo
Fisher). Low quality reads were removed from the FASTQ files
resulting in reads containing an average length of 134 bp. The
Ion Proton data were aligned to the *rattus norvegicus* (rn5)
genome using the two step alignment method with TopHat2 and
Bowtie2 in the Partek Flow servers (Partek, Inc.). Aligned reads
were summarized as gene-level counts (featureCounts 1.4.4). The
FASTQ files from the Ion Proton have been submitted to NCBI’s
Gene Expression Omnibus (GEO) under the accession number:
GSE97608.
or subfield. For the transcriptional changes due to age and cognitive performance, a significant cut-off was set at \( p < 0.01 \). The resulting data sets for each region represent seed lists of differentially expressed genes. Validation of expression was performed using poly-A mRNA sequenced on the Ion Proton. For Ion Proton mRNA, a significance cut-off was set at \( p < 0.05 \) for one-tailed-tests, with the tail specified by the direction of fold change (FC) determined by the Illumina seed list. Thus, the adjusted \( p \)-value for the combined tests was \( p < 0.0005 \). The following formula was used to calculate the false discovery rate (FDR) = (number of genes identified by Illumina + 0.0005/number genes from this list identified by Ion Proton). Heat maps were generated in R with “ggplots” (3.0.1) and “ComplexHeatmap” (1.14.0) using counts which were standardized to z-scores from genes validated with the Ion Proton, and the box plots were generated with “ggplot2” (2.2.1).

To be labeled subfield-specific in CA1, CA3, or DG, gene counts had to be significantly different (Benjamini-Hochberg adj- \( p < 0.05 \)) from both of the other two subfields with a concordant direction of fold change and validated by Ion Proton sequencing at the same significance threshold. Finally, pathway analysis was conducted with the ToppGene web portal against KEGG, PantherDB, and Reactome databases. An additional analysis was conducted using DAVID (version 6.8), considering Gene Ontology for biological processes and cellular components in the “Direct” and “FAT” categories with the Benjamini FDR set at \( p < 0.05 \) as a cut-off for cluster selection.

RESULTS

Behavior

Cue Discrimination

A repeated measures ANOVA indicated an effect of training \( [F(4, 128) = 5.81, p < 0.0005] \) and age \( [F(1, 32) = 11.57, p < 0.0005] \), and an interaction \( [F(4, 128) = 2.46, p < 0.05] \) (Figure 1A). Despite the age difference, all animals were able to locate the platform by the end of training and an ANOVA in each age group confirmed decreased path length with training (\( p < 0.05 \)). Post hoc analysis indicated that young rats exhibited a rapid rate of learning observed as a decrease in escape path length for blocks 2–5 relative to block 1. For aged animals, learning was slower, such that escape path length decreased on blocks 4–5 relative to blocks 2–3.

Spatial Discrimination

A repeated measures ANOVA on performance in the spatial task indicated an effect of training \( [F(4, 128) = 11.28, p < 0.0001] \) and age \( [F(1, 32) = 16.81, p < 0.001] \), in the absence of an interaction (Figure 1B). An ANOVA in each age group confirmed decreased path length with training (\( p < 0.005 \)), indicating that both groups exhibited improved performance with training. Post hoc analysis indicated that young rats exhibited a rapid rate of learning observed as a decrease in escape path length for blocks 2–5 relative to block 1. For aged animals, learning was observed as a decrease in escape path length for blocks 4–5 relative to blocks 1–2.

Regression analysis performed within each age group indicated no relationship between performance on the cue task (block 5 distance) and the spatial task (block 5 distance) (young: \( r = 0.43, p = 0.22 \); aged: \( r = 0.078, p = 0.72 \)). The results indicate that age differences in sensory-motor function or acquisition of the procedural aspects of the task did not mediate impairments in acquisition of a spatial search strategy.

Probe Trial Discrimination Index (DI) Scores

An ANOVA on the discrimination index scores using age and testing site as factors indicated an age difference \( [F(1, 30) = 8.41, p < 0.01] \) in the absence of a training site difference or interaction (Figure 1C). As expected, from the age range of 17–22 months in this strain of rat, there was considerable variability for aged animals, with some animals performing as well as young.

The observed lowest DI score for young animals was 0.24. This value was used as cut-off to classify aged animals as aged-impaired (AI) (DI < 0.24) or aged-unimpaired (AU; DI > 0.24). A repeated measures ANOVA was computed comparing AI and AU groups for the cue task. No group difference was observed for distance to escape to the platform on the cue task (\( p = 0.82 \)). Thus, the difference for the DI scores is not likely due to sensory-motor function or learning the procedural aspects of the task.

Gene Expression across Regions: Comparison of Illumina and Ion Proton Technologies

Sequence Quality Metrics

We sequenced a total of 17.1 and 18.9 million reads per sample, the total mapping rate was 94.1% and 96.8%, with exonic mapping 63.8% and 58.0%, intronic 15.2% and 13.1% and the percentage of intergenic reads was 21.0% and 28.9% on the Illumina and Ion Proton platforms, respectively.

The correlation of number of counts for the common genes is shown in Figure S1. We conducted pairwise differential expression analysis between the three subfields regardless of age or cognitive status. For each pairwise comparison, we retained the differentially expressed genes (DEGs) that were statistically significant with both platforms (adj- \( p < 0.05 \)) and concordant for FC direction. A summary of the number of genes detected in all the pairwise comparisons in both platforms is reported in Table 1. A total of 24 DEGs were discordant for FC between the two platforms in the 3 comparisons (specifically: 5 for CA1 vs. CA3, 14 for CA1 vs. DG and 5 for CA3 vs. DG). Finally, the significant genes showing concordant FC in both platforms were: 2,379 (CA1 vs. CA3), 4,352 (CA1 vs. DG), and 4,642 (CA3 vs. DG). The complete results of the differential analysis for the genes detected with both platforms are reported in Tables S1–S3, and the overlap is represented in the proportional Venn diagram in Figure 2A.

We computed the correlation coefficient for the Log2 FC between the two platforms in each pairwise comparison for all detected genes. The Pearson coefficients demonstrated a strong positive correlation for all of the comparisons (\( p < 2.2e-16 \)), ranging from \( R = 0.776 \) (CA1 vs. CA3) to \( R = 0.836 \) (CA3 vs. DG). In Figure 2B, we illustrate the corresponding
2,431 (DG) specific genes. The results for all three regions are consistent across both platforms and concordant for fold change across all specific genes that met the criteria of significantly differentially expressed (adj-p < 0.05) and detected in both platforms (Table S5). The enrichment analysis conducted with ToppGene using genes with average FC > |0.50| (n = 379) reveals one significant pathway: cilium movement, outer dynein arm, and potassium ion transmembrane transport (Table S7). The last process was also detected using ToppGene.

### In the CA3 region a total of 1,051 of the differentially expressed genes were protein coding, whereas the remaining 12 were processed transcripts or pseudogenes. A total of 551 genes (51.8%) were overexpressed, whereas 512 were underexpressed (48.2%) with respect to the CA1 and DG (Table S5). The boxplots for all of the specific genes are illustrated in Figure S5, and in Figure S6 we show the Allen Brain Atlas data from mouse for Col6a6 and Nnat. The first 5 genes ranked by absolute average fold change were Lnp1, RGD1562638, Plk5 (underexpressed), and Mmp3, Mxl1 (overexpressed). Pathway and enrichment analysis conducted using genes with average FC > |0.50| (n = 535) revealed 11 significant pathways after Bonferroni correction. Top results were: KEGG calcium signaling (adj-p = 4.27E-04; 17 genes) and REACTOME potassium channel (adj-p = 1.73E-03) (Table S7). The analysis conducted with DAVID allowed us to detect 3 significant GO biological processes after Benjamini correction: cilia movement, outer dynein arm, and potassium ion transmembrane transport (Table S7). The last process was also detected using ToppGene.

In the DG region 2,403 of the differentially expressed genes were protein coding, whereas the remaining 10 were pseudogenes, miRNA, processed transcripts or lincRNA. 975 genes were protein coding, whereas the remaining 12 were processed transcripts or pseudogenes. A total of 551 genes (51.8%) were overexpressed, whereas 512 were underexpressed (48.2%) with respect to the CA1 and DG (Table S5). The boxplots for all of the specific genes are illustrated in Figure S5, and in Figure S6 we show the Allen Brain Atlas data from mouse for Col6a6 and Nnat. The first 5 genes ranked by absolute average fold change were: Vgll3, Mxl1, Gmhr (underexpressed), and Wnt3, Ackr2 (overexpressed). Pathway analysis conducted with ToppGene using genes with average FC > |0.50| (n = 379) reveals one significant pathway after Bonferroni correction (KEGG: neuroactive ligand-receptor interaction; adj-p = 7.40E-04) that included 20 genes (Table S7). The enrichment analysis conducted with DAVID did not identify any significant results after Benjamini correction.
genes with absolute average FC > |0.50| (n = 927) revealed 7 significant pathways after Bonferroni correction. The most significant pathways were: KEGG axon guidance (adj-p = 4.70E-05; 23 genes) and REACTOME adherens junctions interactions (adj-p = 6.20E-04; 10 genes). The remaining significant pathways are reported in Table S7. In this same subfield we also detected the KEGG pathway neuroactive ligand-receptor interaction (adj-p = 4.92E-03; 32 genes), the only one shared with CA1 and CA3. In contrast, REACTOME gastrin-CREB signaling pathway via PKC and MAPK and REACTOME neuronal system were detected in CA3 region. The analysis conducted with DAVID revealed enrichment for 8 biological processes: neural crest cell migration (FDR adj-p = 1.77E-03; 12 genes), semaphorin-plexin signaling (FDR adj-p = 2.11E-03) and myosin complex (FDR adj-p = 7.65E-03). All the remaining processes (FDR adj-p < 0.05) are reported in Table S7.

**Gene Expression Related to Aging**

The data sets for differential expression using the cross platform validation method described above were used to investigate age-related changes in gene expression. The CA1 Illumina poly-A mRNA analysis produced 228 DEGs that passed the statistical filter. From this seed list the number of genes that was detected for Ion Proton poly-A mRNA was 104 (FDR = 1.1E-3). Among these, 90 genes were up regulated and 14 genes were down regulated in aged rats. Enrichment analysis indicated clusters related to immune function including the lysosome (GO:0005764, 14 genes, FDR adj-p = 1.3E-3), regulation of leukocyte mediated immunity (GO:0002443, 14 genes, FDR adj-p = 1.6E-5), immune response (GO:0006955, 28 genes, FDR adj-p = 1.8E-3), and neutrophil activation (GO:0042119, 4 genes, FDR adj-p = 9.8E-3). In addition, clusters related to cell adhesion (GO:0007155, 25 genes FDR adj-p = 2.9E-4) and regulation of ERK1 and ERK2 cascade (GO:0070372, 8 genes, FDR adj-p = 2.2E-2) were also present. Furthermore, a KEGG pathway for the lysosome (rno04142, 8 genes, FDR adj-p = 5.2E-3) was identified. Enrichment analysis using the ToppGene web tool confirmed the enrichment of pathways related to immune function (i.e.: Lysosome, B cell receptor signaling pathway and adaptive immune system). All genes under the indicated immune related functional clusters were up regulated genes (Figure S9).
For region CA3, analysis of Illumina poly-A mRNA indicated 189 DEGs in the seed list and the number validated with the Ion Proton poly-A mRNA was 81 (FDR adj-\(p = 1.2^{-3}\)). All 81 genes were in the NIH DAVID database, with 61 genes up regulated and 20 down regulated genes in aged rats. As with area CA1, clustering analysis was related to immune function including leukocyte activation (GO:0045321, 17 genes, FDR adj-\(p = 4.9^{-5}\)) immune response (GO:0006955, 20 genes, FDR adj-\(p = 8.8^{-5}\)), cytokine production (GO:0001816, 12 genes, FDR adj-\(p = 2.4^{-3}\)), antigen processing and presentation of peptide antigen via MHC class II (GO:0002495, 7 genes, FDR adj-\(p = 4.4^{-7}\)), and interleukin-10 production (GO:0032613, 5 genes, FDR adj-\(p = 3.0^{-5}\)). Additionally, a cluster for biological adhesion (GO:0002610, 21 genes, FDR adj-\(p = 4.4^{-4}\)) and a KEGG pathway for cell adhesion molecules (rn04514, 7 genes, FDR adj-\(p = 2.3^{-5}\)) were found. Enrichment analysis conducted with ToppGene confirmed the presence of pathways correlated with the immune system (i.e., immunoregulatory interactions between a lymphoid and a non-lymphoid cell, adaptive immune system, and phagosome). All genes under the indicated immune related functional clusters were up regulated (Figure S10).

The DG region exhibited similar aging patterns in relation to the CA1 and CA3, with 280 DEGs in the Illumina poly-A mRNA data set. From this list, 162 genes were validated by the Ion Proton (FDR = 8.6^{-4}, and all 161 genes were in the DAVID database. From the validated gene list, 134 genes were up regulated and 27 genes decreased with aging. Enriched clusters included immune response (GO:0006955, 41 genes, FDR adj-\(p = 2.9^{-11}\)), defense response (GO:0006952, 40 genes, FDR adj-\(p = 5.8^{-5}\)), inflammatory response (GO:0006954, 26 genes, FDR adj-\(p = 1.1^{-8}\)), response to wounding (GO:0009611, 15 genes, FDR adj-\(p = 9.3^{-3}\)), response to steroid hormone (GO:0048545, 14 genes, FDR adj-\(p = 1.3^{-5}\)) and cell-cell adhesion (GO:0098609, 30 genes, FDR adj-\(p = 2.7^{-6}\)). KEGG pathways were identified for lysosome (rn004142, 10 genes, FDR adj-\(p = 3.2^{-4}\)), antigen processing and presentation (rn04612, 11 genes, FDR adj-\(p = 5.2^{-7}\)), and cell adhesion molecules (rn04514, 10 genes, FDR adj-\(p = 2.8^{-3}\)). All genes under the indicated immune related functional clusters were up regulated genes (Figure S11). Again, the analysis conducted with ToppGene confirmed the immunologic processes detected with DAVID. In addition, we detected 3 pathways involving the complement system: initial triggering of complement, complement and coagulation cascades, complement cascade, and activation of C3 and C5. Beside the immunological pathways, we detected the cadherin signaling and Wnt signaling.

In examining genes that were differentially expressed with age across regions it is clear that each region contains a number of age-related genes that are unique relative to the other hippocampal regions. For the most part, more genes were up regulated in older rats within each region and several up regulated genes overlapped across regions. Among the genes that overlapped regions, several were related to immune function (Table 2). Only one gene, Col4a5, exhibited a decrease across all three regions. Interestingly, the DG region showed the highest number of gene changes related to the age of the animal (162), including the greatest number of distinct genes (91 genes) (Figure 3). Among the 69 genes that were uniquely up regulated in the DG, 68 matched DAVID for functional annotation clustering analysis with the top clusters involved in immune function. Together, the results indicate that the DG region may be more sensitive to aging effects.

**Gene Expression Related to Behavior**

For examination of genes linked to cognitive status, only aged animals were included in order to avoid age as a confound. Aged animals were subdivided into AI and AU according to the water maze DI score. Figure 4 illustrates the number of genes that were differentially expressed in CA1, CA3, and the DG according to cognitive status, and were annotated in DAVID. The Illumina analysis of CA1 poly-A mRNA produced 235 sequences that passed the statistical filter. From the seed list, the number of genes that was detected for Ion Proton mRNA was 107 (FDR = 1.1^{-5}) (Table S8). For the 107 genes, 106 were in the NIH DAVID database, with 48 up regulated and 58 down regulated genes in aged impaired animals. The CA3 Illumina mRNA analysis contained 32 sequences in the seed list and the number validated with the Ion Proton poly-A mRNA was 8 (FDR = 2.0^{-3}) (Table S8). All 8 genes were annotated in DAVID (increased: Prl, Pcdh19, Table 2).

**TABLE 2** | Genes, which significantly increased expression across all three regions in aged animals from both the Illumina and Ion Proton data.

| Gene symbol | Gene Name                      | Immune related |
|-------------|-------------------------------|---------------|
| C3          | Complement component 3        | x             |
| Card11      | Caspase recruitment domain family | member 11    |
| Cd4         | Cd4 molecule                   | x             |
| Cd74        | Cd74 molecule, major histocompatibility complex, class II invariant chain | x |
| Cdh1        | Cadherin 1                    | x             |
| Cdh23       | Cadherin 23 (otocadherin)     | x             |
| Csf1r       | Colony stimulating factor 1 receptor | x |
| Cts5        | Cathepsin 5                   | x             |
| Cts2        | Cathepsin Z                   | x             |
| Co2x1       | Chemokine (C-X3-C motif) receptor 1 | x |
| Fgfr2b      | Fg fragment of IgG, low affinity lib, receptor (CD32) | x |
| Glat        | Glaial fibrillary acidic protein | x |
| Gpr183      | G protein-coupled receptor 183 | x |
| Gpr64       | G protein-coupled receptor 84 | x |
| Itgb2       | Integrin beta 2               | x             |
| Ncf1        | Neutrophil cytokotoxic factor 1 | x |
| Nckap1      | NCK associated protein 1 like | x |
| Npc2        | Niemann-Pick disease, type C2 | x |
| Pcdhb4      | protocadherin beta 4          | x             |
| Pkd4        | Phospholipase D family, member 4 | x |
| Slc14a1     | Solute carrier family 14 (urea transporter), member 1 | x |
| Trem2       | Triggering receptor expressed on myeloid cells 2 | x |
| Bpilio4      | BPI Fold Containing Family B, Member 4 | x |
| Wdfy4       | WDFY Family Member 4          |               |

Genes that are related to immune function are indicated.

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region and submitted for cluster analysis. The results indicated that only CA1 exhibited significant enrichment. Clustering was performed using Illumina DEGs with \( p < 0.01 \) validated with Ion Proton mRNA with \( p < 0.05 \).

![Figure 3](image-url) Number of genes altered during aging across hippocampal regions. Summary of the total number of genes across the hippocampal subfields whose expression increased (up arrow) or decreased (down arrow) using Illumina DEGs with \( p < 0.01 \) validated with Ion Proton mRNA with \( p < 0.05 \).

![Figure 4](image-url) Number of genes differentially expressed in regions CA1, CA3, and DG according to cognitive function. In each case, the number of genes down regulated (open portion) and up regulated (filled portion) in aged-cognitively impaired animals are represented.

Pcdh17, Gabrb1, Gda, Ahs2; decreased: Msrb2, Wnt6). The DG Illumina poly-A mRNA analysis contained 97 sequences in the seed list and the number validated with the Ion Proton mRNA was 30 (FDR = 1.6\(^{-3}\)) (Table S8), which included 29 genes that were annotated in DAVID (16 up regulated and 13 down regulated). Only one gene, Prl, was increased across all regions.

Up and down regulated genes were combined within each region and submitted for cluster analysis. The results indicated that only CA1 exhibited significant enrichment. Clustering was observed for cellular components with CA1 genes linked to synapse (GO:0045202, 20 genes, FDR adj-\( p = 2.6^{-3}\)), cell junction (GO:0030054, 18 genes, FDR adj-\( p = 1.2^{-2}\)), neuron part (GO:0097458, 22 genes, FDR adj-\( p = 5.6^{-3}\)), postsynapse (GO:0098794, 10 genes, FDR adj-\( p = 1.3^{-2}\)), ion channel complex (GO:0034702, 9 genes, FDR adj-\( p = 5.9^{-3}\)) and plasma membrane region (GO:0098590, 16 genes, FDR adj-\( p = 6.1^{-3}\)) (Table 3). In addition, enrichment for biological processes included regulation of signaling (GO:0023051, 34 genes, FDR adj-\( p = 4.7^{-3}\)), the KEGG pathways for calcium signaling (rno04020, 7 genes, FDR adj-\( p = 1.7^{-2}\)) and long-term potentiation (rno04720, 4 genes, FDR adj-\( p = 3.5^{-2}\)). ToppGene identified five significant pathways after Bonferroni correction: glutamatergic synapse, neuronal system, heterotrimeric G-protein signaling pathway-Gq alpha and Go alpha mediated pathway, calcium signaling, and phosphatidylinositol signaling system. The results confirmed the presence of neuronal genes, but pointed to more specific processes, including the glutamatergic synapse.

A closer examination of the direction of gene changes suggested that there was decreased expression of genes that regulated Ca\(^{2+}\) entry by modulating the function of NMDA receptors (Neto1, Prickle2), or voltage-gated Ca\(^{2+}\) channels (Cacnb2) (Figure 5). Decreased expression was also observed for genes linked to receptor mediated release of Ca\(^{2+}\) from intracellular Ca\(^{2+}\) stores (Adra1d, Homer3, Itprl). A decline in protein expression for these genes might be expected to decrease intracellular Ca\(^{2+}\). Similarly, increased expression was observed for the membrane Ca\(^{2+}\) ATPase pump (Atp2b4), which would remove intracellular Ca\(^{2+}\). Moreover, decreased expression was observed for genes linked to downstream Ca\(^{2+}\) signaling (Hpc1a, Dclk2, Prkca). Together, the results suggest that in impaired animals, transcription is altered in an attempt to limit intracellular Ca\(^{2+}\) signaling (Figure 5).

Closer examination of the direction of change for genes linked to synaptic activity suggests differential expression in genes for regulating synaptic receptors and K\(^{+}\) channels (Figure 6). For example, there was a decrease in the GABA receptor subunit (Gabra5) and Nptxr, which are involved in clustering of AMPA receptors at the synapse. Increased expression was observed for two K\(^{+}\) channels, Hcn4 and Kcnkl1, and Kcnab2, a cytoplasmic potassium channel subunit that modulates K\(^{+}\) channel activity was down regulated. Finally, there was down regulation of genes for proteins that are downstream of G-protein coupled receptor activation (Mpp3, Ksr1, Akap13) and an increase expression of the glutamate metabotropic receptor (Grm2).

Finally, we examined the specificity of gene changes. In this case, aged animals were separated by performance on the cue task, using a mean split for block 5. Relative to gene expression associated with spatial discrimination performance, fewer genes were associated with performance on the cue task. Analysis of the CA1 region poly-A mRNA using Illumina produced 135 sequences that differed according to cue discrimination performance. From this seed list the number of genes that was detected for Ion Proton poly-A mRNA was 24 (FDR = 2.8\(^{-3}\)). For region CA3, 38 differentially expressed genes were detected by Illumina and 3 genes were confirmed by the Ion Proton (FDR = 6.3\(^{-3}\)). For the dentate gyrus, Illumina detected 71 sequences and 4 were confirmed by the Ion Proton (FDR = 8.8\(^{-3}\)). No gene clusters were observed for any region.
TABLE 3 | Genes of age impaired and age unimpaired animals, which were significantly differentially expressed in region CA1 in both the Illumina and Ion Proton data.

| Gene symbol | Gene Name                                                                 | 
|-------------|--------------------------------------------------------------------------|
| Adra1d      | Adrenergic, alpha-1D-, receptor                                           |
| Akap13      | A kinase (PRKA) anchor protein 13                                        |
| Arpc5       | Actin related protein 2/3 complex, subunit 5                             |
| Dck2        | Doublecortin-like kinase 2                                               |
| Dgkz        | Diacylglycerol kinase zeta                                              |
| Gabra5      | Gamma-aminobutyric acid (GABA) receptor, alpha 5                        |
| Homer3      | Homer homolog 3 (Drosophila)                                             |
| Itpr1       | Inositol 1,4,5-triphosphate receptor, type 1                             |
| Kcnab2      | Potassium voltage-gated channel, shaker-related subfamily, beta member 2|
| Ksr1        | Kinase suppressor of ras 1                                               |
| Mmp3        | Membrane protein, palmitoylated 3 (MAGUK p55 subfamily member 3)         |
| Neto1       | Neurexin (NRX) and tollloid (TLL)-like 1                                 |
| Nptxr       | Chromobox homolog 6; neuronal pentraxin receptor                        |
| Pragmin     | Pragma of Rnd2                                                           |
| Prickle2    | Prickle homolog 2 (Drosophila)                                           |
| Rem2        | RAS (RAD and GEM) like GTP binding 2                                     |
| Slc17a8     | Solute carrier family 17, member 8                                       |
| Vav2        | Avdin 2 guanine nucleotide exchange factor                                |
| Cacnb2      | Calcium channel, voltage-dependent, beta 2 subunit                      |
| Dgka        | Diacylglycerol kinase, alpha                                             |
| Hpca        | hippocalin                                                               |
| Itpr1       | Inositol 1,4,5-triphosphate receptor, type 1                             |
| Prkca       | Protein kinase C, alpha                                                  |
| Scube2      | Signal peptide, CUB domain                                               |
| Sulf2       | Sulfatase 2                                                              |

Genes that are related to Ca^{2+} binding and synaptic function are indicated.

DISCUSSION

Gene Expression across Subfields: Subfield Specific Transcriptome Profiling

We report the results of the transcriptional differences among hippocampus brain subfields (CA1, CA3, and DG) regardless of the animal’s age or cognitive status. Our approach was based on retaining for each subfield, genes significant and concordant in FC in both pairwise comparisons. We were able to detect a high number of specific genes for each subfield, specifically: 908 (CA1), 1,063 (CA3), and 2,431 (DG) genes. These lists include genes highly or lowly expressed in each subfield with respect to the others. In CA1, the top gene with high values of expression compared to the other subfields was Wnt3 (Wnt Family Member 3), which is considered to play a central role in synaptogenesis and adult neurogenesis (Varela-Nallar and Inestrosa, 2013). In area CA1, Wnt signaling has been linked to synaptic plasticity and associated hippocampal-dependent memory processes (Xu et al., 2015; Ivanova et al., 2016). At the pathway level, those genes with higher expression in CA1 exhibited enrichment for Neuroactive ligand-receptor interaction, including Ns (neurotensin), Htr1b (5-hydroxytryptamine receptor 1B) and Gprl61 (G Protein-Coupled Receptor 161).

For subfield CA3, we observed increased expression of genes linked to processes involving potassium channel activity in both ToppGene (Potassium channels, Voltage gated potassium channels) and DAVAD enrichment analysis (GO:001805 Potassium ion transmembrane transport). Potassium channels are central to intrinsic excitability in region CA3 of the hippocampus, which is highly vulnerable to cell death due to seizure activity (Ben-Ari, 1985; Cooper, 2012). Indeed, many of the top overexpressed genes (Mmp3, Pagl1Zac1, Galr1, Gla1, and Nk3r) are regulated by neural activity and linked to epileptogenesis of the hippocampus (Elmslie and Gardiner, 1995; Valente et al., 2004; Mazarati et al., 2006; McColl et al., 2006; Dubey et al., 2017).

Enrichment analysis of genes that are highly expressed in the DG relative to other subfields indicated that the most significant processes were “Axon guidance” (KEGG) and “Neural crest cell migration” (GO:0001755). In general, the genes and the biological processes detected as enriched in the DG may be due to the known neurogenesis activity in this subfield. In the DG, the top genes overexpressed with respect to the region other regions included Bhlhe23 (basic helix-loop-helix family member E23) and Msx3 (msx like homeobox 3), which are involved in neurogenesis and specification of neuronal subtype (Bramblett et al., 2004; Liu et al., 2004; Hesse et al., 2011). EphA8 (EPH receptor A8), a member of the ephrin receptor subfamily and Clq12 (complement C1q like 2) are implicated in neurite growth and establishing synaptic connections (Buchser et al., 2010; Iijima et al., 2010). Plk5 (polo like kinase 5) modulates the formation of neuritic processes upon stimulation of the brain-derived neurotrophic factor (BDNF)/nerve growth factor (NGF)-Ras pathway in neurons (de Carcer et al., 2011). Interestingly, we also found Bdnf specifically overexpressed in this subfield (average FC = 0.948).

Subfield Vulnerability to Aging

It is likely that regional differences in transcription during aging are associated with differences in connectivity and molecular make-up of each subfield. For example, relative to CA3 pyramidal neurons, CA1 pyramidal neurons are more sensitive to metabolic perturbations (Jackson et al., 2009). Blalock and associates have suggested that metabolic changes precede neuroinflammation and have linked increased vulnerability of gene changes in CA1, relative to CA3, to biomarkers of metabolic syndrome in nonhuman primates (Blalock et al., 2010). In considering...
the increase in age-related shift in transcription in the DG, it is important to note that neurogenesis in the DG exhibits a robust decline between adult and middle-age (Kuhn et al., 1996; Lemaire et al., 2000; Driscoll et al., 2006). Similarly, the rat DG exhibits a loss of perforant path input (Geinisman, 1979; Barnes and McNaughton, 1980), and loss of neuronal connections underlie the altered expression of genes related to cell-cell adhesion or axon guidance and increased markers of
inflammation (Ianov et al., 2016). Interestingly, older memory-impaired humans also show reduced white matter volume in the vicinity of the major axon input pathway to the hippocampus, the perforant pathway (Rogalski et al., 2012). Thus, while we observed that the DG exhibits robust changes in transcription during aging, it remains to be discovered how these changes specifically relate to DG function and to anatomical or physiological changes associated with aging. The subfield-specific changes observed in activity of the human dentate gyrus/CA3 region, however, does suggest that these alterations occur across species, and may contribute to memory deficits during aging in humans and other animals (Yassa et al., 2011).

Consistent with a number of previous studies, we observed that brain aging was associated with an increase in markers of neuroinflammation including expression of immune response genes (Rowe et al., 2007; Kadish et al., 2009; Blalock et al., 2010; Zeier et al., 2011; Cribbs et al., 2012; Ianov et al., 2016). Typical genes include Gfap, complement genes (C3, C4b), and genes associated with antigen processing through the lysosome (Cd74, Cisd, Ctss, Laptm5, Dnase2) or phagosome (Fcg2b, Nfj1). In the current study, the DG region showed the highest number of gene changes related to the age of the animal. This is in contrast to previous microarray studies, which suggest that for BNxF344 rats examined between middle-age (12–18) and advanced age (26–28 months), region CA1 is more vulnerable than DG (Zeier et al., 2011; Masser et al., 2014). In this case the differences likely relate to the animal model, age of onset for gene changes, and differences in the age range examined (Ianov et al., 2016). In F344 rats, for ages similar to those employed in the current study (5–6 and 17–22 months), the CA1 region of older animals exhibited relatively smaller transcriptional changes compared to the prefrontal cortex and white matter (Ianov et al., 2016). Similarly, a decline in interneuron markers occurs in all hippocampal regions from adult to middle-age in F344 rats and the decline is observed in CA1 but not the DG when analysis is limited to middle-age and aged BNxF344 rats (Shi et al., 2004; Stanley and Shetty, 2004). Thus, one possibility is that changes in the DG emerge earlier than in CA1, minimizing age-related changes between middle-age and older animals. Alternatively, we have observed that the majority of gene changes in region CA1 of F344 rats occurs between adult and middle-age (Blalock et al., 2010), suggesting possible species differences.

The increase in immune response genes suggests activation of astrocytes and microglia. However, it is important to note that the age-related increase in expression of inflammation genes does not necessarily predict memory impairment or the stage of neurodegeneration (Blalock et al., 2003; Cribbs et al., 2012). In the case of neurodegeneration, neuroinflammation may initially have beneficial and neuroprotective effects (Chakrabarty et al., 2010; Streit et al., 2014; Sochocka et al., 2017). Glial activation during aging is associated with a decline in synaptic density (Rozovsky et al., 2005) and the ability to maintain or form new synapses may underlie resiliency in the face of aging or neurodegenerative disease (Arnold et al., 2013; Ianov et al., 2016, 2017).

### Genes Associated with Impaired Episodic Memory

The identification and specificity of genes related to cognition is a function of the brain region sampled and the type of cognitive process examined (Ianov et al., 2016). In the current study, within each region, more genes were correlated with episodic memory scores relative to sensory-motor performance on the cue discrimination task, suggesting specificity of gene changes. While the hippocampus is required for spatial memory, different hippocampal-dependent processes depend on specific regional and molecular mechanisms. For example, the DG is involved in spatial pattern separation and region CA1 with memory for topological relationships in space (Kesner et al., 2004; Goodrich-Hunsaker et al., 2008; Kesner and Rolls, 2015). Thus, if the task had focused on spatial pattern separation we would expect a greater correspondence between behavior and transcription in the DG. Regardless, aging of the DG likely contributes to impaired spatial episodic memory. While no functional gene clusters were observed, several of the DG genes that were down regulated in impaired animals (Bcl6, Crim1, Dapk1, Mts1, Oas1a) have been linked to cell growth/apoptosis and neurogenesis or neuronal development.

The results on CA1 transcriptional changes associated with cognitive decline are in confirmation of previous studies employing microarray technology, which indicate that impaired cognition is associated with altered expression of synaptic genes, genes involved in Ca$^{2+}$ regulation, and glutamatergic synapses (Toescu et al., 2004; Burger, 2010; Uddin and Singh, 2013; Volk et al., 2015). However, rarely are the same genes observed to change in the same direction across studies. The difference may be due to different cognitive processes examined including reference memory and episodic memory, differences in ages, and the time after behavioral training in which transcription is examined. Previous work has established that Ca$^{2+}$ dysregulation and impaired Ca$^{2+}$-dependent synaptic plasticity in region CA1 is associated with poor performance on episodic spatial memory (Landfield, 1988; Foster and Norris, 1997; Foster, 1999, 2007; Burke and Barnes, 2010; Oh and Disterhoft, 2010) and these processes are influenced by behavioral training. We examined transcription 2 weeks after testing. Thus, the differences observed here are not likely due to differential response to training and may represent an underlying and chronic shift in transcription for this region. In our experiments, the changes in expression of genes involved in the glutamatergic synapses in aging are concordant with the evidence that P301Tau expression increased hippocampal glutamate release and decreased glutamate uptake, and these alterations in glutamate signaling correlated with cognitive deficits in the hippocampal-dependent Barnes maze task (Hunsberger et al., 2015).

Finally, it is important to point out the advantages or disadvantages of the experimental design. The current study was conducted across multiple testing sites and employed two different methods for transcriptional analysis. The advantage of multiple testing sites includes the ability to pool data across sites. However, there is the challenge of equating data due to differences in hardware or protocols. Episodic memory is sensitive to age-related cognitive decline across species; however,
there is considerable variability in cognition with advancing age. Furthermore, aged animals can exhibit sensory-motor impairment and acquire the procedural aspects of the task much more slowly than young animals, which could influence the validity of cognitive measures. Thus, in the current study possible sensory-motor and procedural learning contributions to behavior on the spatial task were controlled by first testing the animals on the cue discrimination task. Characterization of spatial episodic memory was based on a DI score, which would tend to standardize scores for animals tested on mazes of different sizes. The testing procedure and use of a DI score likely contributed to the good correspondence of behavioral results across the different testing sites, such that all young animals performed above chance (DI score = 0) and approximately half the aged animals at each site exhibited impaired performance. Finally, behavior on the spatial task was not related to behavior on the cue discrimination task.

Due to the large numbers of transcripts examined, the possibility of false positives or false negatives should be controlled. In order to reduce the instance of false discovery we were able to increase the power of the analysis by pooling a relatively large number of animals that were tested across the two sites (young = 10; aged unimpaired = 12; aged impaired = 12). In addition, it is advised that transcriptional profiling studies should use another method to confirm suspected changes. Each method has its benefits and limitations and there are major differences in the nature of the approaches. In the current study, we have taken advantage of two different next-generation platforms, Illumina fluorescent labeling and Ion Proton semiconductor measures of proton release, to confirm differential expression of thousands of genes (Kusko et al., 2014; Li et al., 2015; Reuter et al., 2015). Consistent with previous reports, we observed a good concordance in the gene counts between the different platforms (R = 0.909), increasing the confidence in the observed transcriptional changes. We now expand this idea by showing considerable concordance between the two platforms in detecting differences in expression. Thus, coefficients for the Log2 FC between the two platforms across regions was highly significant (Figure 2B), with the correlation coefficients ranging from 0.776 (CA1 vs. CA3) to 0.836 (CA3 vs DG). Indeed, for genes that changed in relation to age or cognitive function, the probability of false discovery across the platforms was generally <2.0^{-3}.

CONCLUSION

The two next-generation sequencing platforms were able to validate variability in gene expression associated with several variables. Due to the high number of specific genes detected in each subfield, our findings confirm previous studies conducted on the same hippocampal subfields in other animal models with the microarray technology, demonstrating a correspondence between cytoarchitectural boundaries, transcriptome distribution, and underlying differences in physiology and vulnerability to insults (Datson et al., 2004; Lein et al., 2004; Newrzella et al., 2007). The results confirm that brain aging is associated with increased expression of genes linked to the immune response; however, we noted regional differences with more robust changes observed in the DG. Finally, the literature suggests several physiological and transcriptional processes that may underlie cognitive decline; however, there is considerable diversity in the literature concerning which specific genes are altered. Our results confirm that cognitive decline is associated with differential expression of CA1 genes linked to Ca^{2+} homeostasis and synaptic plasticity. Furthermore, due to the time between testing and examination of transcription, we would suggest that these changes represent an underlying and chronic shift in transcription.

AUTHOR CONTRIBUTIONS

LI performed experiments, analyzed data, wrote the manuscript, and constructed figures; AR, MC, AS, and AJK, performed experiments; AK and IP performed experiments, contributed in writing the manuscript and figures. MD analyzed data and constructed figures. MH and TF designed the experiments, analyzed the data, constructed figures, wrote the manuscript. JD, CB, and JS designed the experiments, contributed to the manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fnagi.2017.00383/full#supplementary-material

Supplementary Figure 1 | Correlation (R^2 = 0.938) of gene counts across sequencing platforms, Illumina and Ion Proton.

Supplementary Figure 2 | Heatmap of hippocampal subfield changes in gene expression. Plotted for each hippocampal subfield, are genes found with both sequencing platforms to be differentially expressed compared to the other two subfields at a significance of adjP < 0.05. Each row represents the z-scores from the DEGs, and the columns are the biological replicates for each region.

Supplementary Figure 3 | Boxplots for the CA1 specific genes reported in Table S4. The normalized counts are from the Illumina experiment, and the genes are ranked according the average Fold Change computed from the pairwise comparisons CA1 vs. CA3 and CA1 vs. DG. (specific to CA1 in our study) as reported in the Allen Brain Atlas (Mouse).

Supplementary Figure 4 | Expression of genes Wsfl1, Nbx, Ndst4, and Oprl161, (specific to CA1 in our study) as reported in the Allen Brain Atlas (Mouse).

Supplementary Figure 5 | Boxplots for the CA3 specific genes reported in Table S5. The normalized counts are from the Illumina experiment, and the genes are ranked according the average Fold Change computed from the pairwise comparisons CA3 vs. CA1 and CA3 vs. DG. (specific to CA3 in our study) as reported in the Allen Brain Atlas (Mouse).
**Supplementary Figure 7** | Boxplots for the DG specific genes reported in Table S6. The normalized counts are from the Illumina experiment, and the genes are ranked according the average Fold Change computed from the pairwise comparisons DG vs. CA1 and DG vs. CA3.

**Supplementary Figure 8** | Expression of genes Pdln, Pkhd, C1qtl2, and Dsp (specific to DG in our study) as reported in the Allen Brain Atlas (Mouse).

**Supplementary Figure 9** | Heatmap of age-related changes in gene expression in the CA1 region. Each row represents a DEG (Illumina p < 0.01; Ion Proton p < 0.05) associated with aging. Gene-level counts were standardized to z-scores and the color represents the standard deviation increasing (yellow) or decreasing (blue) relative to the mean (black). The age-related gene enrichment clusters (FDR adj-p < 0.05) in the CA1 region included immune related GOs, neutrophil activation, cell adhesion and regulation of ERK1 and ERK2 cascade.

**Supplementary Figure 10** | Heatmap of age-related changes in gene expression in the CA3 region. Each row represents a DEG (Illumina p < 0.01; Ion Proton p < 0.05) associated with aging. Gene-level counts were standardized to z-scores and the color represents the standard deviation increasing (yellow) or decreasing (blue) relative to the mean (black). The age-related gene enrichment clusters (FDR adj-p < 0.05) in the CA3 region included immune related GOs such as leukocyte activation, cytokine production, and antigen processing and presentation.

**Supplementary Figure 11** | Heatmap of age-related changes in gene expression in the DG region. Each row represents a DEG (Illumina p < 0.01; Ion Proton p < 0.05) associated with aging. Gene-level counts were standardized to z-scores and the color represents the standard deviation increasing (yellow) or decreasing (blue) relative to the mean (black). The age-related gene enrichment clusters (FDR adj-p < 0.05) in the DG region included immune related GOs such as defense response, and inflammatory response. Cell-cell adhesion GO was also detected.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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