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Enhanced Cognition and Hypoglutamatergic Signaling in a Growth Hormone Receptor Knockout Mouse Model of Successful Aging.

Abbreviated Title: Cognition & Glutamate in Aged GHR-KO Mice

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Running Title: Cognition and glutamate in GHR-KO Mice

Abstract

Growth hormone receptor knockout (GHR-KO) mice are long lived with improved healthspan, making this an excellent model system for understanding biochemical mechanisms important to cognitive reserve. The purpose of the present study was to elucidate differences in cognition and glutamatergic dynamics between aged (20-24 month-old) GHR-KO and littermate controls. Glutamate plays a critical role in hippocampal learning and memory and is implicated in several neurodegenerative disorders, including Alzheimer's disease. Spatial learning and memory were assessed using the Morris water maze (MWM), while independent dentate gyrus (DG), CA3 and CA1 basal glutamate, release, and uptake measurements were conducted in isoflurane anesthetized mice utilizing an enzyme-based microelectrode array (MEA) coupled with constant potential amperometry. These MEAs have high temporal and low spatial resolution while causing minimal damage to the surrounding parenchyma. Littermate controls performed worse on the memory portion of the MWM behavioral task and had elevated DG, CA3 and CA1 basal glutamate and stimulus-evoked release compared with age-matched GHR-KO mice. CA3 basal glutamate negatively correlated with MWM performance. These results support glutamatergic regulation in learning and memory and may have implications for therapeutic targets to delay the onset of, or reduce cognitive decline, in Alzheimer's disease.

Keywords: healthspan, longevity, biosensor, electrode, Alzheimer's disease

1. Introduction

Growth hormone (GH) is secreted by the anterior pituitary gland and binds to its receptor ubiquitously expressed throughout the mammalian reproductive, muscular, endocrine, and nervous systems, where it regulates growth and metabolism often mediated through insulin-like growth factor-I (IGF-1)¹. The GH/IGF-1 axis is regarded as an important regulator in aging such that attenuated signaling increases lifespan², potentially by providing protection against cancer, diabetes, and neurodegeneration³. This is supported by GH receptor knockout (GHR-KO) mice that exhibit increased lifespan (30-36 months) and improved healthspan (reduced and delayed incidence of neoplasia, protection from diet-induced nephropathy, and retained cognitive abilities). Furthermore, inhibition of GH has been shown to increase neuronal differentiation⁴ and GHR-KO mice have an approximate 25% increase in total neuron cell density in the cortex.⁵ However these benefits come at a concession of decreased size with increased adiposity, similar to Laron Syndrome⁶⁻¹⁰.

The increased lifespan of GHR-KO mice make them an excellent model system for studying age-related cognitive reserve and neurotransmitter regulation in comparison to littermate controls. For example, recent data support that GHR-KO mice maintain hippocampal glutamatergic function from 4-22 months of age, including vesicular glutamate transporter (VGLUT) 1 (the predominant hippocampal VGLUT located in classical excitatory terminals¹¹) and 3, glutamate transporter 1 (GLT-1), and the N-methyl-D-aspartate (NMDA) receptor subtype GluN2B¹². Glutamate, the predominant excitatory neurotransmitter in the mammalian CNS, has a strong prevalence in neocortical and hippocampal pyramidal neurons; therefore, playing a critical role in learning and

memory¹³. Because of this, glutamate dysregulation has been implicated in age-related cognitive decline associated with neurodegenerative disorders such as Alzheimer's disease¹³⁻¹⁷.

The purpose of the present study was to elucidate differences in cognition and glutamatergic dynamics between aged (20-24 month-old) GHR-KO and age-matched littermate controls. Spatial learning and memory was assessed using the Morris water maze (MWM) behavioral paradigm¹⁸⁻²⁰. Extracellular glutamate dynamics were studied in the dorsal hippocampus since this region is important for consolidation and retrieval of spatial memory²¹. Glutamate measurements were conducted utilizing an enzyme-based microelectrode array (MEA) coupled with constant potential amperometry to independently measure basal glutamate and stimulus-evoked glutamate release, and uptake in the dentate gyrus (DG), CA3, and CA1 of isoflurane anesthetized littermate controls and GHR-KO mice. These MEAs have high temporal (4 Hz)²² and low spatial resolution (50 x 100 μ m) while causing minimal damage to the surrounding parenchyma (50-100 microns)²³. The results presented here support the importance of glutamatergic regulation for learning and memory in the GHR-KO mouse model of successful aging.

2. Methods

2.1 Animals

Female GHR-KO (20-24 months of age) and age-matched female normal littermate controls, were produced in a breeding colony maintained at Southern Illinois University School of Medicine. This colony was established by crossing 129Ola/BALB/c GHR +/- breeders²⁴ provided by Dr. John J. Kopchick with mice derived from crosses of C57BL/6J and C3H/J strains and maintained as a closed colony with inbreeding minimized by avoiding brother x sister matings²⁵. Protocols for animal use were approved by the *Laboratory Animal Care and Use Committee* at Southern Illinois University School of Medicine. Animals were group housed on a 12:12 hour light: dark cycle, and food and water were available *ad libitum*. Each mouse underwent cognitive assessment, *in vivo* glutamate recordings, and post mortem histological analysis of brain tissue with the exception of a loss of two GHR-KO mice that occurred after MWM due to complications during surgery prior to *in vivo* electrochemistry. Immediately following *in vivo* glutamate recordings, mice were euthanized with an overdose of isoflurane and decapitated. Upon decapitation, 5 mm tail snips were collected and stored at -80°C until shipment to TransnetYX[®], Inc (Cordova, TN) for genotype verification.

2.2 Chemicals

All chemicals were prepared and stored according to manufacturer recommendations unless otherwise noted. L-glutamate oxidase (EC 1.4.3.11) was obtained from Cosmo Bio Co. (Carlsbad, CA) and diluted in distilled, deionized water to make a 1U/ μ l stock

solution that was stored at 4°C. Sodium phosphate monobasic monohydrate, sodium phosphate dibasic anhydrous, 1,3-phenylenediamine dihydrochloride (mPD), sodium chloride, calcium chloride dehydrate, and hydrogen peroxide (H₂O₂, 30% in water) were obtained from Thermo Fisher Scientific (Waltham, MA). L-glutamic acid sodium salt, potassium chloride, bovine serum albumin (BSA), glutaraldehyde, dopamine hydrochloride (DA), L-ascorbic acid (AA), and DPX were obtained from Sigma-Aldrich Co. (St. Louis, MO). Rabbit polyclonal glial fibrillary acidic protein antibody (GFAP) was obtained from Dako. Guinea pig polyclonal VGLUT1 antibody was obtained from Millipore. Biotinylated goat anti-rabbit serum, biotinylated goat anti-guinea pig serum, avidin-biotin complex (ABC) kit, and VIP peroxidase substrate kit were obtained from Vector Laboratories.

2.3 Morris water maze

The MWM tests spatial learning and memory by requiring the mouse to utilize visual clues for locating a static, submerged platform, regardless of the starting quadrant as previously reported²⁰. The MWM paradigm consists of 5 consecutive learning days with three, 90 s trials/day and a minimum of 20 min between trials. After two days without testing, mice are given a single, 60 s probe challenge to test memory. The ANY-maze video tracking system (Stoelting Co., Wood Dale, IL) records and analyzes duration in each quadrant, distance traveled, average speed, and path efficiency for the 5 learning days. Additional parameters analyzed for the single probe trial include the number of platform crosses, time in annulus 40, latency, distance, and path efficiency to first platform cross.

2.4. In Vivo Glutamate Measurements

2.4.1 Enzyme-Based Microelectrode Arrays

Enzyme-based MEAs with platinum (Pt) recording surfaces (Figures 1A & B) were fabricated, assembled, coated, and calibrated for *in vivo* mouse glutamate measurements^{22,26,27}. Briefly, one of the R2 MEA Pt sites was coated with an L-glutamate oxidase, BSA and glutaraldehyde coating solution. BSA and glutaraldehyde increase the adhesion and crosslink L-glutamate oxidase to the MEA surface. L-glutamate oxidase causes the enzymatic breakdown of glutamate to α -ketoglutarate and the electroactive reporter molecule, H_2O_2 . The second Pt recording site (self-referencing or sentinel site) was coated similar to the glutamate recording site, except L-glutamate oxidase was omitted from the coating solution; therefore, the sentinel site was unable to enzymatically generate H_2O_2 from L-glutamate. A potential of +0.7V vs a Ag/AgCl reference electrode was applied to the Pt recording surface, resulting in a two electron oxidation of H_2O_2 and the current was amplified and digitized by the Fast Analytical Sensing Technology (FAST) 16mkIII (Quanteon, LLC; Nicholasville, KY) electrochemistry instrument.

2.4.2 mPD Electropolymerization

A minimum of 72 hrs after enzyme coating, all Pt recording surfaces were electroplated with 5 mM mPD in 0.05 M phosphate buffered saline (PBS)²⁸. FAST electroplating software applied a potential as a triangular wave with an offset of -0.5V, peak-to-peak amplitude of 0.25V, at a frequency of 0.05 Hz, for 20 min to create a size exclusion layer that restricts the passage of AA, DA, uric acid and 3,4-dihydroxyphenylacetic acid.

2.4.3 Calibration

A minimum of 24 hrs after mPD electropolymerization, each MEA was calibrated *in vitro* prior to implantation to generate a standard curve for the conversion of current to glutamate concentration²⁹. The Pt recording sites and a glass Ag/AgCl reference electrode (Bioanalytical Systems, Inc., West Lafayette, IN) were placed in a continuously stirred solution of 0.05 M PBS (40.0 mL) maintained at 37°C with a recirculating water bath (Stryker Corp., Kalamazoo, MI). Final beaker concentrations of 250 µM AA, 20, 40, and 60 µM L-glutamate, 2 µM DA, and 8.8 µM H₂O₂ were used to assess MEA performance (Figure 1C). A total of 18 MEAs (13 unique) were used in the present study. The average ± standard error of the mean (SEM) for glutamate sensitivity was 9.7 ± 1.0 pA/µM ($R^2 = 0.995 \pm 0.003$), selectivity ratio of 805 ± 327 to 1, and limit of detection of 0.45 ± 0.26 µM based on a signal-to-noise ratio of 3.

2.4.4 Microelectrode Array / Micropipette Assembly

A glass micropipette (1.0 mm outer diameter, 0.58 mm internal diameter; World Precision Instruments, Inc., Sarasota, FL) was used to locally apply solutions to the mouse hippocampal subfields. Glass micropipettes were pulled using a vertical micropipette puller (Sutter Instrument Co., Novato, CA) and the tip was 'bumped' to create an internal diameter of 12-15 µm. The tip of the micropipette was positioned between the pair of recording sites and mounted 100 µm above the MEA surface. The micropipettes were filled with sterile filtered (0.20 µm) 70 mM KCl (70 mM KCl, 79 mM NaCl and 2.5 mM CaCl₂, pH 7.4). Fluid was pressure-ejected from the glass micropipette using a Picospritzer III (Parker-Hannafin, Cleveland, OH), with pressure (5-15 psi) adjusted to

consistently deliver volumes between 100-200 nl over 1-2 s intervals. Ejection volumes were monitored with a stereomicroscope (Luxo Corp., Elmsford, NY) fitted with a calibrated reticule³⁰.

2.4.5 Reference Electrode

A Ag/AgCl reference electrode was prepared by stripping 5 mm of Teflon from each end of a silver wire (200 μm bare, 275 μm coated; A-M Systems, Carlsberg, WA). One of the stripped ends was soldered to a gold-plated test connector (Newark element14 Chicago, IL) and the other end was coated with AgCl by placing the tip of the stripped silver wire (cathode) into a 1 M HCl plating bath saturated with NaCl containing a stainless steel wire (anode) and applying +9 V DC using a power supply to the cathode versus the anode for 15 min.

2.5 *In Vivo* Anesthetized Recordings

At least one week following the MWM probe challenge, mice were anesthetized using 1.5-2.0% isoflurane (Abbott Lab, North Chicago, IL) from a calibrated vaporizer (Parkland Scientific, Inc., Coral Springs, FL) and prepared for *in vivo* electrochemical recordings (Hascup & Hascup, 2014). The mouse was placed in a stereotaxic frame (David Kopf Instruments, Tujunga, CA) fitted with a mouse anesthesia mask and body temperature was maintained at 37°C with a hydrothermal pad connected to a recirculating water bath. A craniotomy was performed to access the DG (AP: -2.0, ML: \pm 1.0, DV: -2.2 mm), CA3 (AP: -2.0, ML: \pm 2.0, DV: -2.2 mm), and CA1 (AP: -2.0, ML: \pm 1.0, DV: -1.7 mm) from Bregma based on the coordinates of Paxinos and Franklin³¹. A Ag/AgCl reference wire

was implanted in the right cortex, remote from the recording area. The MEA / micropipette assembly was lowered into the DG, CA3, or CA1 (hippocampal subfield and hemisphere randomly assigned for each mouse) using a microdrive (Narishige International, East Meadow, NY) attached to the electrode holder of the stereotaxic arm. Constant potential amperometry (4Hz) was performed using the FAST16mkIII and FAST software for multi-channel simultaneous recordings³². MEAs were allowed to reach a stable baseline for 60 min, at which time basal glutamate measures were taken (ten second average) followed by pressure ejection studies. All values for pressure ejections refer to changes versus baseline. After pressure ejection studies, brains were removed and fixed in 4% paraformaldehyde followed by 0.1 M phosphate buffer containing 30% sucrose for at least 24 hours before cryosectioning for immunohistochemistry (45 μ m).

2.6 Immunohistochemical Staining and Semiquantification

Immunohistochemistry in the hippocampus was performed using rabbit polyclonal GFAP (1:2000) or guinea pig polyclonal VGLUT1 antibody (1:1,000). Briefly, primary antibodies were applied to serial sections taken for every sixth section from the hippocampus based on our previous protocols³³. Endogenous peroxidase activity was quenched by treating sections with 10% H₂O₂ in 20% methanol for 10 minutes. Sections were then permeabilized in TBST (Tris-buffered saline with 0.25% TritonX-100) following treatment for 20 minutes with sodium metaperiodate. Non-specific binding was controlled by incubation in 10% normal goat serum for 1 hour. Sections were then incubated overnight in the primary antibody at room temperature. The next day, sections were incubated for 1 hour with the secondary antibody (1:200; biotinylated goat anti-rabbit serum or

biotinylated goat anti-guinea pig serum) and 1 hour with the ABC kit. The reaction was developed using the VIP peroxidase substrate kit to enhance the reaction and produce a color stain. This reaction was stopped using 0.1 M phosphate buffer, and the sections were mounted on glass slides, dehydrated, and cover-slipped with DPX. To control for staining intensity, staining of all sections for each antibody were conducted on the same day, and developed with VIP for the same amount of time (GFAP: 3 minutes, VGLUT-1: 2 minutes).

Staining intensity of GFAP and VGLUT-1 in the hippocampus was determined using National Institutes of Health Image J Software 1.48 to measure a gray scale value within the range of 0–256, where 0 represents white and 256 black. A template for the DG, CA3, and CA1 subregions of the hippocampus was created and used on all brains similarly, and images were captured with a Nikon Eclipse E-600 microscope equipped with an Olympus-750 video camera system, and a Dell Pentium III computer. Measurements were performed blinded and approximately 6 sections were averaged to obtain 1 value per subject. Staining density was obtained when background staining was subtracted from mean staining intensities on every sixth section through the hippocampus.

2.7 Data Analysis

The FAST16MkIII electrochemical instrument and FAST software saves amperometric data, time and pressure ejection events for all Pt recording sites. Calibration data, in conjunction with a MATLAB (MathWorks, Natick, MA) graphic user interface program developed by Jason Burmeister Consulting, LLC (Version 6.1) was used to calculate basal glutamate and 70 mM KCl-evoked glutamate release and uptake. To determine

extracellular glutamate concentration, the sentinel site current (pA) was subtracted from the glutamate recording site current (pA) and divided by the slope (pA/ μ M) obtained during the calibration^{32,34–36}. Basal glutamate was calculated by taking a 10 s baseline average prior to start of pressure ejection in the DG, CA3, and CA1. For stimulus-evoked glutamate studies, five reproducible signals were obtained in each hippocampal subfield. These signals were then averaged into a single data point for each hippocampal subfield per mouse for comparison between genotypes. Glutamate uptake followed first-order-rate kinetics; therefore the uptake rate constant (k_{-1}) was calculated as the logarithmic slope of glutamate concentration versus time (s^{-1}) estimated by use of regression analyses ($R^2 \geq 0.9$). Because of different cell types and afferent inputs, hippocampal subfields were analyzed independently. Prism software (GraphPad Software, Inc., La Jolla, CA) was used for all statistical analyses. A two-way Analysis of Variance (ANOVA) with Tukey's multiple comparison's post-hoc test was used to analyze MWM training data. An unpaired, two-tailed Student's t-test was used to analyze MWM probe, electrochemical, and immunohistochemical data in each hippocampal subfield. Comparisons between MWM and glutamate electrochemical data were established using Pearson correlation. Outliers, determined using Grubbs' test with $\alpha = 0.05$, were removed prior to analysis (no more than one per genotype). Data are represented as mean \pm SEM and significance was defined as $p < 0.05$.

3. Results

3.1 MWM Training and Probe Challenge

Learning and memory was assessed using an 8 day MWM behavioral paradigm as described in section 2.4²⁰. A significant increase by the fifth training day, relative to the first training day, in the path efficiency ($F(4,72) = 13.72$; $p < 0.0001$) to reach the hidden escape platform was observed for the littermate controls and GHR-KO mice with no differences observed between the two genotypes on any day (Figure 2A); indicating that both genotypes were able to consolidate memories and there were no learning-related genotypic differences. During the probe challenge littermate controls took a less efficient path (0.16 ± 0.04 ; $F(9,9) = 2.844$; $p = 0.0280$) to first platform entry compared to GHR-KO mice (0.37 ± 0.07 , respectively); indicating littermate controls had impaired memory (Figure 2B). Representative probe challenge tracks for littermate controls and GHR-KO mice are shown in figures 2C & D, respectively.

3.2 Basal Glutamate

Prior to stimulus-evoked glutamate release, basal glutamate measures were assessed in each of the hippocampal subfields. Littermate control basal glutamate (Figure 3A) was elevated compared with GHR-KO mice in the DG ($2.6 \pm 0.5 \mu\text{M}$, $0.9 \pm 0.3 \mu\text{M}$; $F(6,6) = 3.582$; $p = 0.0099$), CA3 ($3.1 \pm 0.4 \mu\text{M}$, $1.2 \pm 0.4 \mu\text{M}$; $F(6,6) = 1.232$; $p = 0.0029$) and CA1 ($4.3 \pm 0.8 \mu\text{M}$, $1.2 \pm 0.5 \mu\text{M}$; $F(7,6) = 3.134$; $p = 0.0055$). A negative correlation ($r = -0.5619$; $p = 0.0365$) was observed between CA3 basal glutamate and path efficiency to first platform entry during the MWM probe trial (Figure 3B).

3.3 Stimulus-Evoked Glutamate Release

We locally applied consistent volumes of 70 mM KCl to littermate controls and GHR-KO mice in the DG (158 ± 7 nl, 147 ± 4 nl; $F(9,7) = 3.555$; $p = 0.2008$), CA3 (152 ± 5 nl, 155 ± 10 nl; $F(9,7) = 2.447$; $p = 0.8040$), and CA1 (149 ± 6 nl; 153 ± 5 nl; $F(9,6) = 1.560$; $p = 0.6239$) to evoke glutamate release (Figure 4A). As shown in Figure 4B, local application of 70 mM KCl elicited robust, reproducible glutamate signals in the CA3 of littermate controls and GHR-KO mice. These glutamate signal patterns were observed in all three hippocampal subfields studied. The average maximal stimulus-evoked glutamate release was significantly elevated in the littermate controls versus GHR-KO mice in the DG (4.9 ± 1.0 μ M, 2.3 ± 0.5 μ M; $F(9,7) = 3.984$; $p = 0.0452$), CA3 (9.1 ± 0.9 μ M, 3.7 ± 0.5 μ M; $F(9,7) = 4.772$; $p = 0.0002$) and CA1 (3.7 ± 0.6 μ M, 2.0 ± 0.2 μ M; $F(9,6) = 12.91$; $p = 0.0332$) as shown in Figure 4C. No statistical differences between stimulus-evoked glutamate uptake was observed between the littermate controls and GHR-KO mice in DG (0.4 ± 0.2 μ M/sec, 0.7 ± 0.2 μ M/sec; $F(8,7) = 1.647$; $p = 0.3392$), CA3 (2.0 ± 0.5 μ M/sec, 1.1 ± 0.3 μ M/sec; $F(9,7) = 3.263$; $p = 0.1470$), or CA1 (0.6 ± 0.2 μ M/sec, 0.7 ± 0.3 μ M/sec; $F(8,7) = 2.673$; $p = 0.4228$) (Figure 4D).

3.4 Histological Assessment

Littermate control GFAP (Figure 5A) was decreased compared with GHR-KO mice in the DG (10.6 ± 1.3 , 15.3 ± 1.3 ; $F(8,7) = 1.121$; $p = 0.0236$), CA3 (12.7 ± 0.5 , 15.9 ± 1.0 ; $F(7,8) = 3.089$; $p = 0.0096$), and CA1 (8.7 ± 0.6 , 12.5 ± 0.8 ; $F(7,8) = 1.684$; $p = 0.0016$). Representative images of GFAP staining in whole hippocampus for littermate controls and GHR-KO mice are shown in Figures 5B and C, respectively. Magnified images of the DG, CA3, and CA1 for littermate controls (5D, F, and H, respectively) and GHR-KO mice (Figures 5E, G, and I, respectively) are presented. We observed hippocampal brain

region and genotype dependent changes in VGLUT1 staining. Littermate controls VGLUT1 (Figure 6A) was decreased compared to GHR-KO mice in the DG (4.5 ± 0.6 , 12.3 ± 0.8 ; $F(7,8) = 1.633$; $p < 0.0001$), no change in the CA3 (6.8 ± 0.8 , 8.7 ± 1.0 ; $F(7,8) = 1.163$; $p = 0.1476$), and increased in the CA1 (8.2 ± 0.9 , 5.2 ± 0.5 ; $F(8,7) = 3.997$; $p = 0.0133$). Representative images of VGLUT1 staining in the DG, CA3, and CA1 of littermate controls and GHR-KO mice are shown in Figures 6B-G.

4. Discussion

The glutamatergic system plays an important role in age-related cognitive decline and cognitive disorders, and that tight regulation of glutamate is essential for normal brain/cognitive function. Generally, the two major means of controlling both tonic and phasic glutamate neurotransmission are through glutamate release and uptake. Under normal conditions, glutamate release is primarily accomplished through depolarization of glutamatergic neurons, which can be identified by the presence of VGLUTs located at the terminals. Glial cells, composed of astrocytes and microglia, are predominantly responsible for clearance of glutamate from the extracellular space mediated through surface expression of excitatory amino acid transporters (EAATs). One EAAT, Glt-1 (EAAT2 in humans), is responsible for ~90% of glutamate clearance from the extracellular space³⁷. Additionally, Glt-1 surface expression and function decrease with age, possibly leading to excitotoxicity, which may be exacerbated in age-related cognitive disorders, further supporting the importance of proper glutamate system maintenance for cognitive retention in advanced age¹⁷. We have previously reported on age-related changes in

glutamatergic markers, including elevated mRNA expression of GLT-1 and retained VGLUT1 levels in the hippocampus of long lived GHR-KO mice compared to age-matched littermate controls¹². In the present study, we examine cognition and glutamatergic neurotransmission dynamics to elucidate the role of glutamate in GHR-KO mice that exhibit enhanced cognition in old age compared to age-matched littermate controls.

Our MWM data supports that 20-24 month-old littermate controls and GHR-KO mice learn to locate the hidden escape platform, indicating no differences in memory consolidation. However, after a 48 hour delay, littermate controls present with impaired memory retrieval compared to GHR-KO mice during the probe challenge of the task. While it is well known that memory retention declines with age in mice^{38,39}, previous studies utilizing the inhibitory avoidance, open field, and MWM tasks indicate memory retrieval in GHR-KO mice does not decline with age⁴⁰⁻⁴². This may be the result of more stringent regulation of the glutamate system in the hippocampus, as supported by our previous findings¹². Therefore, a possible explanation for improved memory retrieval in GHR-KO mice is that they experience delayed aging thereby postponing the onset of cognitive decline⁴⁰.

The dorsal hippocampus is important for consolidation and retrieval of spatial memory during the MWM task²¹. Hippocampal inhibition of the EAATs, and therefore decreased glutamate clearance, has been shown to induce long term depression (LTD) mediated through elevated extrasynaptic glutamate binding to the GluN2B-NMDA receptor subtype⁴³ and negatively impacting cognition. In support of this, we observed a negative correlation between CA3 basal glutamate and path efficiency to first platform entry on the MWM task, however, this correlation was not observed with DG or CA1 basal glutamate.

This may be due to the importance of mossy fiber projections to the CA3 subfield for memory consolidation and retrieval in spatial navigation tasks^{44,45}. Therefore, the increased availability of basal glutamate to activate GluN2B in the CA3 may contribute to the cognitive decline observed in littermate controls while the corollary is true in the GHR-KO mice. Furthermore, we observed elevated GFAP staining in the hippocampus of GHR-KO mice indicating an increase in glia, which likely contributed to the overall decrease in glutamatergic tone in these mice.

We observed decreased evoked release of glutamate in GHR-KO mice in all three hippocampal subregions examined compared to age-matched littermate controls. This may be explained by previous reports¹² of an age-related decrease in VGLUT1 mRNA levels from whole hippocampus in littermate controls, but not GHR-KO mice. Further evaluation of 20-24 month old female GHR-KO and littermate control mice presented in the current manuscript using histological techniques revealed hippocampal subregion dependent changes in VGLUT1. However, this is contradictory to our evoked glutamate release data where littermate controls exhibited elevated glutamate release compared to GHR-KO mice. This likely indicates a disconnect between the number of glutamate terminals (VGLUT1) and the inherent excitability of the neurons (stimulated release). It is also possible that the anesthetic used during our glutamate recordings (isoflurane) preferentially suppressed KCl-evoked glutamate release in GHR-KO mice. However, this is highly unlikely because isoflurane dosage was kept consistent (1.5-2.0%) between mice. Furthermore, GHR-KO mice have a higher metabolic rate than their littermate controls and therefore would metabolize isoflurane more quickly, if anything, thereby having less anesthetic-related alterations.

Since basal glutamate levels are regulated by a combination of continuous release and uptake, we are not able to directly measure clearance kinetics as we can with stimulus-evoked glutamate release. However, we did not observe any difference in 70 mM KCl-evoked glutamate uptake between genotypes in any of the hippocampal subfields studied despite a significant difference in GFAP levels, possibly because there was sufficient GLT-1 present in close proximity and functioning adequately enough to quickly clear evoked glutamate released from both genotypes. We observed significantly more GFAP in all three hippocampal subregions examined in GHR-KO mice compared to littermate controls. It is possible that the increased energy demands of GHR-KO mice compared to littermate controls⁴⁶ may lead to an increase in astrocytes (the main location of glycogen in the brain) to ensure that the brain has sufficient energy, with glutamate uptake into astrocytes contributing to the store of available energy^{47,48}. While an increase in GFAP does not always indicate elevated GLT-1, our previous data indicate elevated GLT-1 mRNA in GHR-KO hippocampus¹². Since GLT-1 accounts for 90% of glutamate uptake³⁷, a decrease in this transporter may slow glutamate clearance, leading to the elevated basal glutamate observed in the littermate controls. However, it should be noted that mRNA levels do not necessarily correspond to protein levels or even surface expression and it is possible that experimental design limitations may have contributed to this discrepancy between elevated GFAP and GLT-1, but no differences in the uptake rate of evoked release of glutamate. First, EAATs are electrogenic and membrane depolarization slows glutamate uptake⁴⁹. Second, isoflurane, the anesthetic used in this study, increases glutamate uptake through GLT-1⁵⁰. Although, the amount of stimulus

and anesthetic were consistent in all mice, it is possible that the combination of these two phenomena could alter GLT-1 such that their effects overshadow differences in transporter number when measured with our recording technique.

Additionally, glucocorticoids may contribute to elevated basal and stimulus-evoked glutamate release in the littermate control mice. Glucocorticoids have the potential to increase basal glutamate levels in the hippocampus and impair memory through binding of glutamate to the GluN2B containing NMDA receptor⁵¹, which may also be affected by isoflurane. While Huack and colleagues⁵² demonstrated there was no difference in corticosterone levels in 4-5 month old female GHR-KO and littermate control mice under stressed and non-stress conditions, to date glucocorticoid levels have not been examined in aged female GHR-KO mice. Previous studies support that glucocorticoid levels increase as female mice age contributing to age-related bone loss⁵³. Considering GHR-KO mice are a model of delayed aging, it goes to reason that glucocorticoid levels may be increased in littermate controls compared to GHR-KO mice, potentially contributing to the elevated extracellular glutamate observed in this study.

In summary, we have demonstrated that littermate controls have impaired cognition and hyperglutamatergic signaling in the DG, CA3, and CA1 hippocampal subfields compared to GHR-KO mice. Cognitive performance on the MWM behavioral task was predicted by CA3, but not DG or CA1, basal glutamate and elevated GFAP observed in GHR-KO mice may be neuroprotective by contributing to decreased basal glutamate levels. Taken together, these data and our recent reports of elevated evoked glutamate release in the hippocampus of APP/PS1 mice, a model of Alzheimer's disease²⁰, support an overarching theme whereby elevated hippocampal glutamate is associated with cognitive

impairment and maintenance of the glutamatergic system throughout life is essential for the preservation of cognition in aging and age-related neurodegenerative disorders. Furthermore, these data support the importance of glutamatergic regulation for learning and memory in the GHR-KO mouse model of successful aging that may relate to cognitive resilience and could have implications as therapeutic targets to delay the onset of, or reduce cognitive decline, in several diseases and disorders, including Alzheimer's disease.

Conflicts of Interest

The authors declare no competing financial interests.

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Figure Legends:

Figure 1: MEA and *In Vitro* Calibration. A) Image of the R2 MEA used for anesthetized recordings with a ruler for scale comparison and (B) magnified tip depicting 2 Pt recording sites, each measuring 50 x 100 μm with 100 μm spacing between sites. C) A typical MEA *in vitro* calibration measuring the change in current on a glutamate measuring site (black) and a sentinel recording site (gray) with the addition of multiple analytes, as indicated (\downarrow). The addition of interferents such as AA and DA produced no current change on either site since they are blocked by the mPD exclusion layer. Three glutamate additions showed a stepwise increase of current on the glutamate oxidase / BSA / glutaraldehyde site, but no response on the BSA / glutaraldehyde sentinel site. The addition of H_2O_2 produced a similar increase of current on both recording sites demonstrating equivalent functionality.

Figure 2: Memory Impairment in Littermate Controls. MWM training sessions indicates path efficiency (A) to locate the hidden platform was significantly increased over time for littermate controls and GHR-KO mice (n=10 per group). By the fifth training session both genotypes took similar durations and distances to reach the platform indicating comparable learning. Two-way ANOVA indicates a significant ($p < 0.001$) effect of training day. $**p < 0.01$, $***p < 0.001$ vs GHR-KO training day 1 and $§§§§p < 0.0001$ vs littermate control training day 1 based on a Tukey's multiple comparison post-hoc. MWM probe challenge indicates littermate controls took a less efficient path (B) to first platform entry compared to GHR-KO mice. Two-tailed Student's t-test (n = 10 mice per group), $*p < 0.05$. Representative MWM probe tracks from littermate controls (C) and GHR-KO mice (D). The small circle represents the previous location of the hidden escape platform while the circle surrounding the platform represents the annulus 40.

Figure 3: Elevated CA3 Basal Glutamate Predicts MWM Impairments. A) Bar graphs depicting elevated basal glutamate in all hippocampal subfields of littermate controls compared to GHR-KO mice. Two-tailed Student's t-test (n = 7-8 mice per group) $**p < 0.01$. B) Correlation of CA3 basal glutamate with MWM probe path efficiency to first platform entry for littermate controls (open circles) and GHR-KO mice (dark squares). Dashed line represents the least squares regression.

Figure 4: Stimulus-Evoked Glutamate Release and Uptake. A) Bar graph depicting a similar range of 70 mM KCl (stimulus) was used to elicit glutamate release in all hippocampal subfields of both mouse genotypes. B) Representative traces of local application (\uparrow) of 70 mM KCl-evoked glutamate release in the CA3 of littermate controls (top, gray) and GHR-KO (bottom, black) mice. C) Bar graphs of average maximal evoked glutamate was elevated in all hippocampal subfields of littermate controls compared to GHR-KO mice. D) Glutamate uptake rate was not significantly different between genotypes in any of the 3 hippocampal subfields. Two-tailed Student's t-test (n = 7-10 mice per group), $*p < 0.05$, $***p < 0.001$.

Figure 5: Hippocampal GFAP Levels. Histological staining of GFAP in the hippocampus of littermate controls and GHRKO mice. A) Bar graph of GFAP average mean density was elevated in all hippocampal subfields of GHR-KO mice compared to littermate controls. Two-tailed Student's t-test (n = 8-9 mice per group), $*p < 0.05$, $**p < 0.01$.

Representative images of GFAP staining in whole hippocampus for littermate controls (B) and GHR-KO mice (C) (scale bar is 100 μ m). Representative magnified images of littermate control and GHR-KO mice DG (D, E, respectively), CA3 (F, G, respectively), and CA1 (H, I, respectively) (scale bar is 50 μ m).

Figure 6: Hippocampal VGLUT1 Levels. Histological staining of VGLUT1 in the hippocampus of littermate controls and GHRKO mice. A) Bar graph of VGLUT1 average mean density in littermate controls was decreased compared to GHR-KO mice in the DG, no change in the CA3, and increased in the CA1. Two-tailed Student's t-test ($n = 8-9$ mice per group), * $p < 0.05$, **** $p < 0.0001$. Representative images of littermate control and GHR-KO mice DG (B, C, respectively), CA3 (D, E, respectively), and CA1 (F, G, respectively). Scale bar is 25 μ m.

Figure 1

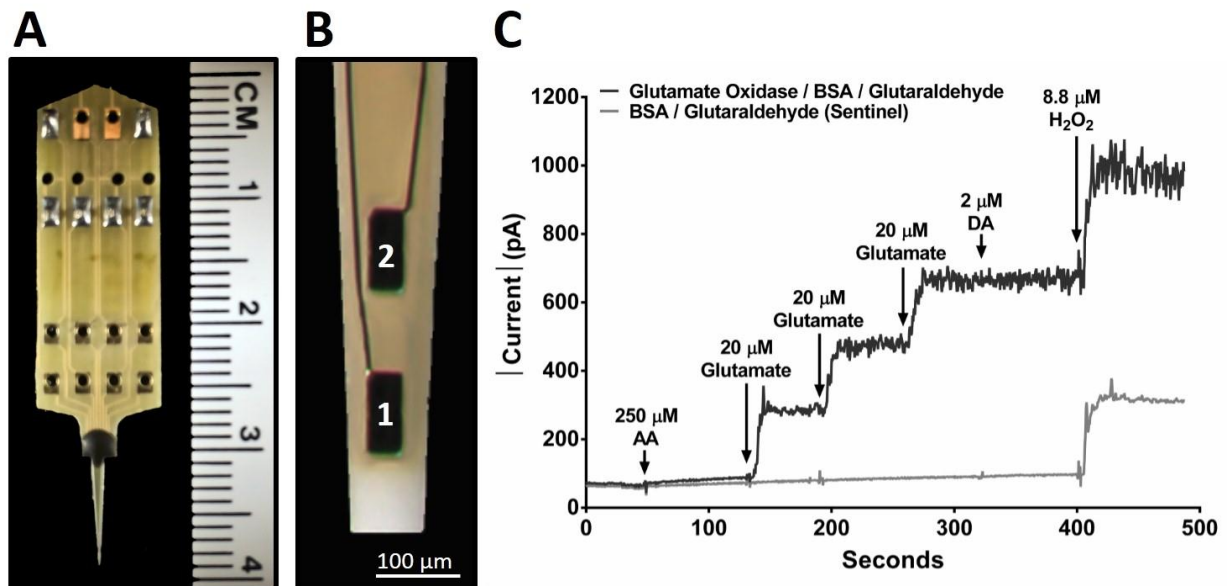


Figure 2

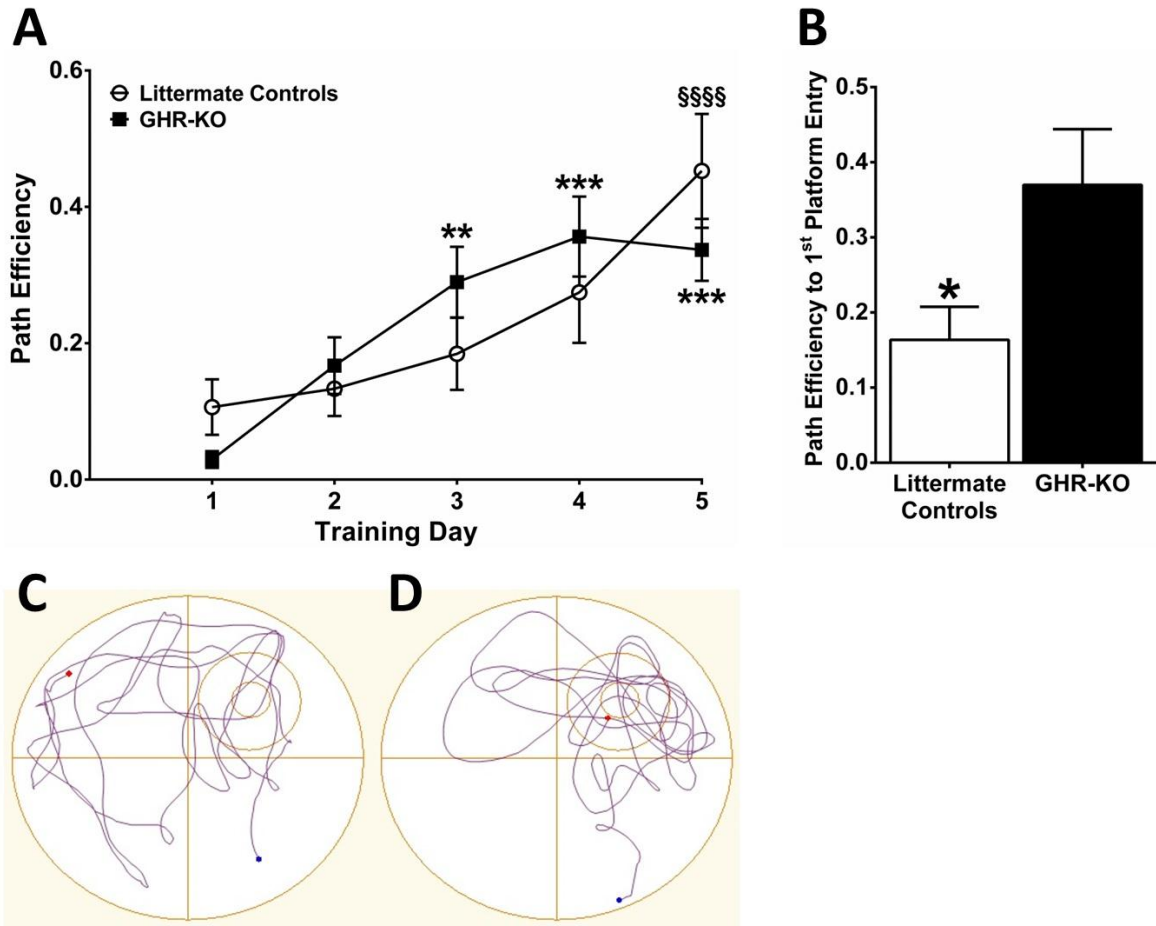


Figure 3

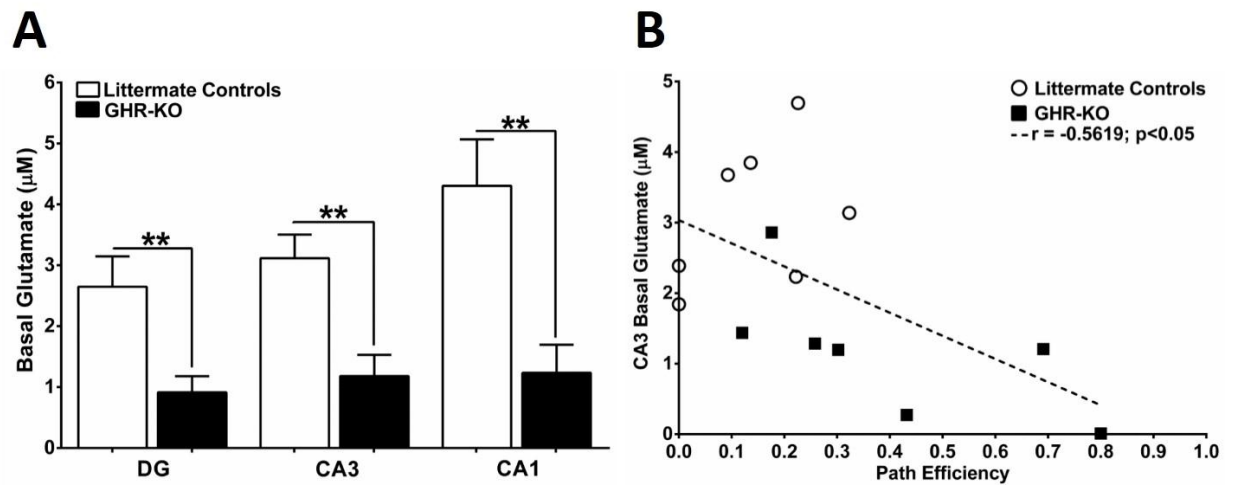


Figure 4

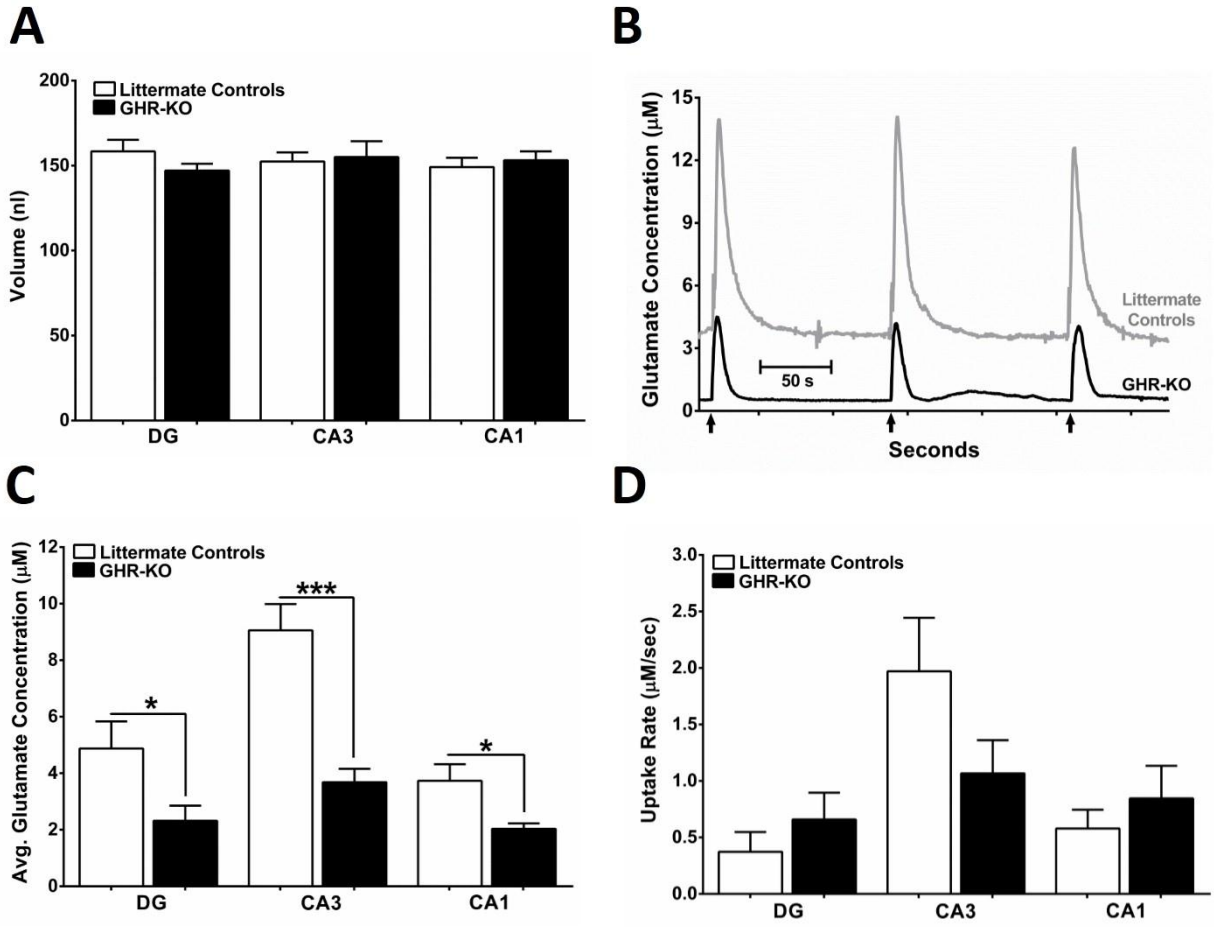


Figure 5

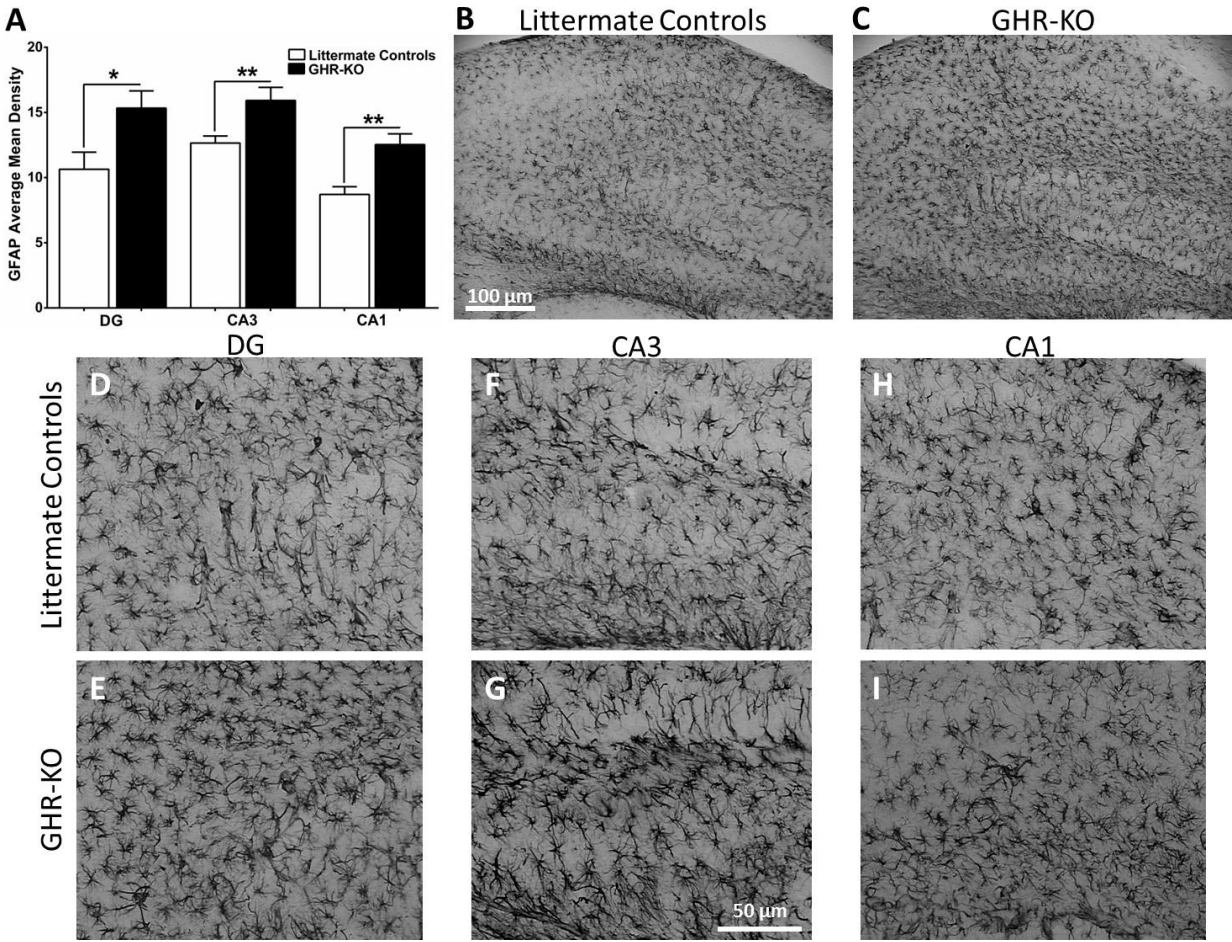


Figure 6

