Cognitive Decline Is Associated with Reduced Reelin Expression in the Entorhinal Cortex of Aged Rats

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Abstract: Cognitive deficits that commonly occur during aging are associated with the disruption of specific neural networks. The layer II neurons in the entorhinal cortex, which form a key circuit in the medial temporal lobe memory system, have a particular vulnerability that extends across a spectrum of age-related memory loss, including amnestic mild cognitive impairment (MCI) and Alzheimer’s disease (AD). These neurons are functionally positioned at the intersection of information processing in cortical networks and the hippocampal formation, and they provide the majority of input to the hippocampal formation. Because the cellular basis for entorhinal layer II vulnerability remains undetermined in these conditions, it is of interest to identify specific molecular features of those neurons in cognitive aging, which may give clues about the earliest changes associated with hippocampal network dysfunction.

Keywords: aging, Alzheimer’s disease, lateral entorhinal cortex, learning, mild cognitive impairment

Brain regions and neural circuits differ in their vulnerability to changes that occur during aging and in age-related neurodegenerative diseases. Among the areas that comprise the medial temporal lobe memory system, the layer II neurons of the entorhinal cortex, which form the perforant path input to the hippocampal formation, exhibit early alterations over the course of aging. Reelin, a glycoprotein implicated in synaptic plasticity, is expressed by entorhinal cortical layer II neurons. Here, we report that an age-related reduction in reelin expression in the entorhinal cortex is associated with cognitive decline. Using immunohistochemistry and in situ hybridization, we observed decreases in the number of Reelin-immunoreactive cells and reelin messenger RNA expression in the lateral entorhinal cortex of aged rats that are cognitively impaired relative to young adults and aged rats with preserved cognitive abilities. The lateral entorhinal cortex of aged rats with cognitive impairment also exhibited changes in other molecular markers, including increased accumulation of phosphorylated tau and decreased synaptophysin immunoreactivity. Taken together, these findings suggest that reduced reelin expression, emanating from layer II entorhinal neurons, may contribute to network dysfunction that occurs during memory loss in aging.

Materials and Methods

Animals and Behavioral Characterization

Male Long-Evans rats, obtained from Charles River Laboratories, were 6 (adult) or 24 (aged) months of age at the time of the studies. All rats were individually housed for 2 months to acclimate to the vivarium environment before being sacrificed. For ethical reasons, the vivarium temperature was maintained at 23°C and the lighting was set to a 12:12 light-dark cycle. Rats were allowed ad libitum access to food and water.

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before behavioral testing, which took place as described (Boric et al. 2008). Training occurred over 8 days in sessions of 3 trials per day with a 60-s intertrial interval. During the trials, rats were placed in the water at the perimeter of the pool, with starting locations varied across trials. Each trial lasted for 90 s or until the rat successfully located the platform. Every sixth trial was a probe trial to assess the rat’s spatial bias during its search. Rats were permitted to escape on probe trials when a retracted platform was made available after 30 s for completion of those trials. An index score, derived from the proximity of the rat to the escape platform location during the 30-s free swim on probe trials, was used to characterize performance of the rats in the maze for the purpose of neurobiological analyses. This index is the sum of the weighted proximity scores measured during the probe trials, with lower scores reflecting better spatial memory as indicated by shorter average distances from the platform location (Gallagher et al. 1993). For immunohistochemistry experiments, we analyzed tissue from \( n = 9 \) young, \( n = 9 \) aged-impaired, and \( n = 9 \) aged-unimpaired rats. For in situ hybridization, we used tissue from \( n = 5 \) young, \( n = 6 \) aged-impaired, and \( n = 6 \) aged-unimpaired rats.

**Euthanasia and Tissue Preparation**

For immunohistochemistry, rats were anesthetized with isoflurane and perfused transcardially with sterile saline, followed by 4% paraformaldehyde in phosphate buffer. After 24 h postfixation, brains were moved into progressively increasing concentrations of glycerol in phosphate buffer. Brains were then sectioned on the coronal plane in a 1 in 10 series at 50 \( \mu \)m thickness. Sections were stored in cryoprotectant at -80 °C.

For in situ hybridization, rats were anesthetized, perfused, and brains were postfixed as described above. After postfixation, brains were moved into 4% paraformaldehyde in phosphate buffer containing 20% sucrose for dehydration and cryoprotection. Brains were then frozen on dry ice and stored at -80 °C prior to sectioning. A 1 in 12 series of coronal sections (30 \( \mu \)m) were cut using a freezing microtome and stored in 4% paraformaldehyde at 4 °C.

**Immunohistochemistry and Immunofluorescence**

For reelin peroxidase labeling, free-floating sections were incubated in 3.0% H\(_2\)O\(_2\) in phosphate-buffered saline (PBS) to quench endogenous peroxidases. Sections were rinsed, then reacted in 0.1 M citric acid for 10 min. After additional rinses, the tissue was moved into primary antibody solution containing mouse anti-reelin (1:1000, Chemicon) with 0.25% Tween-20. The tissue was reacted in primary antibody solution containing mouse anti-reelin (1:1000, Chemicon) and rabbit anti-phospho-tau antibody was used at 1:500 (antibody kindly provided by Dr Rudolf Leube). An index score, derived from the proximity of the rat to the escape platform location during the 30-s free swim on probe trials, was used to characterize performance of the rats in the maze for the purpose of neurobiological analyses. This index is the sum of the weighted proximity scores measured during the probe trials, with lower scores reflecting better spatial memory as indicated by shorter average distances from the platform location (Gallagher et al. 1993). For immunohistochemistry experiments, we analyzed tissue from \( n = 9 \) young, \( n = 9 \) aged-impaired, and \( n = 9 \) aged-unimpaired rats.

**Antibody Specificity**

Standard immunohistochemical controls included omission of primary antibodies. The specificity of the reelin antibody used in these studies has been determined previously (de Bergeyck et al. 1998). To produce the CP13 antibody against tau phosphorylated at serine 202, mice were immunized with paired helical filament tau purified from AD brain tissue as described by Jicha et al. (1997). The CP13 antibody specifically recognizes the phosphopeptide sequence GYSSPG(phospho-serine 202)PGTPGSR and does not react with any other phosphoserine site on the tau protein (Jicha et al. 1997). The total tau antibody was raised against the C-terminal part (amino acids 243–441) containing the 4 repeated sequences involved in microtubule binding. The anatomical pattern of staining for total tau was identical to that shown using this antibody in previous studies (Planell et al. 2007), and this antibody has previously been shown to label tau protein independently of its phosphorylation state (Biernt et al. 2002). The synaptophysin antibody detects an epitope corresponding to amino acids 221–313 mapping at the C-terminus of synaptophysin of human origin, with crossreactivity for the synaptophysin protein in rats. The spatial pattern of synaptophysin immunoreactivity was identical to that observed in previous studies (Stranahan et al. 2008), and we tested antibody specificity in tissue from synaptophysin knockout mice (kindly provided by Dr Rudolf Leube). No staining was observed in tissue from synaptophysin knockout mice.

**Unbiased Stereology and Anatomical Criteria**

For stereological quantification, the optical fractionator method was implemented using the Stereoinvestigator morphometry system (MicroBrightField), as described (Rapp et al. 2002). Slides were coded prior to analysis and the code was not broken until the analysis was finished. Neuron counts were derived from bilateral sections through the entire rostrocaudal extent of the entorhinal cortex. The anatomical borders of the lateral and medial entorhinal areas were traced under low-power magnification, and subsequent cell counting was performed within these borders as shown schematically in Figure 2A and as described previously for the entorhinal cortex in this study population (Rapp et al. 2002). In detail, the lateral entorhinal cortex was localized anatomically by its position inferior to the rhinal sulcus and by its rostral-lateral position relative to the medial entorhinal cortex, as described (Insausti et al. 1997). The lateral entorhinal cortex is bordered rostrally by the perirhinal cortex. Reelin labeling in layer II was restricted to the lateral entorhinal cortex and was not observed in the superficial layers of the perirhinal cortex (Fig. 2B,C). In this regard, the presence of reelin labeling in layer II is a chemoarchitectonic marker that delineates the lateral entorhinal cortex.

The medial entorhinal cortex abuts the caudal and medial edge of the lateral entorhinal cortex (Fig. 2A). The transition from lateral to medial entorhinal cortex was readily apparent based on greater width of the lamina dessicans (layer IV) in the medial relative to the lateral entorhinal cortex. Layer II of the lateral entorhinal cortex exhibits continuous labeling for reelin, relative to the medial entorhinal cortex, consistent with the cellular architectures in these regions on Nissl-stained preparations (Rapp et al. 2002). Thus, we defined the border between the lateral and medial entorhinal cortices using anatomical criteria with reference to Paxinos and Watson (1998) and cytoarchitectural criteria with reference to the continuity (lateral) or clustering (medial) of layer II neurons. The medial entorhinal cortex is bordered by the parasubiculum on its medial surface, and the transition between the medial entorhinal cortex and the parasubiculum is demarcated by a distinct break in the width of layer II (Rapp et al. 2002). Cells were viewed through a 40× objective and counted as they first came into focus within each optical dissector. Counts derived from these samples were converted to estimates of total labeled cell numbers for each region.

**Confocal Microscopy and Typhoon Imaging**

Analysis of synaptophysin labeling in the entorhinal cortex was carried out as described in Stranahan et al. (2008). Briefly, images of synaptophysin labeling were captured on a Zeiss LSM 510 Meta confocal microscope. The frame size was 71.43 × 71.43 μm with 512 × 512 pixel density. Ten frames per animal were collected from superficial layers of the lateral and medial entorhinal areas, with sampling spaced evenly throughout the rostrocaudal extent of these regions. Quantitative analysis was carried out using LSM 510 software.
To acquire images of tau and phospho-tau labeling that could be sampled across larger regions of the entorhinal cortex, we used a typhoon fluorescent imaging system (GE Healthcare). For this analysis, the 488 excitation channel was captured at the 520 emission setting with bandpass 40 filtering. The 532 excitation channel was captured at 610 emission with bandpass 30 filtering. For both channels, the photomultiplier tube was set to 500 V. This scanning method has a 10 µm minimum resolution and allowed us to measure immunoreactivity for tau and phospho-tau in the same tissue sections using ImageJ. Grayscale optical density values for phosphorylated tau and total tau were averaged across anatomically matched tissue sections to give a single score for each brain region in each animal.

**In Situ Probe Synthesis**

Probe templates were synthesized as described in Haberman et al. (2008). Initial primer sequences were as follows: for reelin, left, agaactcaacagtctggttg, right, ctctgaagcaattcga; for DAB1, left, gaaactcagtgtaccagac, right, agagccaaacacatctgcac. Polymerase chain reaction (PCR) products were verified by restriction endonuclease digestion. Initial PCR products were amplified further with the same PCR primers that had been modified by the addition of T7 or SP6 RNA polymerase-binding sites. PCR products containing T7 and SP6 RNA polymerase-binding sites. PCR products containing T7 and SP6 were purified by SVgel and a PCR cleanup kit (Promega). Digestion of PCR products containing T7 and SP6 RNA polymerase-binding sites. PCR products containing T7 and SP6 were then treated with acetic anhydride solution (11.3% triethanolamine, 0.25% acetic anhydride, 0.04 M acetic acid) for 10 min at room temperature. This was followed by two 15-min washes in 2x sodium citrate). Next, sections were transferred to hybridization buffer containing 20% formamide, 0.4 M sodium citrate/citrate buffer (SSC buffer; 20x concentration, 0.3 M sodium citrate). Next, sections were transferred to hybridization buffer containing 20% formamide, 0.4x Denhardt’s solution, 4% dextran sulfate, and 1.6x SSC) supplemented with 0.25 mg/ml transfer RNA, 0.33 mg/ml sheared salmon sperm DNA, 100 mM dithiothreitol (DTT), and 1 x 10^7 cpm/ml 35S-UTP-labeled probe for overnight reaction at 60 °C. The following day, sections were washed at 60 °C in 4x SSC/0.01 M DTT and 2x SSC/50% formamide. They were then incubated with RNase (20 µg/ml) at 37 °C for 30 min. Sections were washed with progressively decreasing concentrations of SSC before mounting on slides.

**Densitometry of In Situ Hybridization mRNA Labeling**

Slides processed for in situ hybridization were exposed to a phosphoimager screen and quantified by using ImageQuant (GE Healthcare). Digital images were acquired for entorhinal cortical sections from the same levels for all animals. We analyzed (n = 5) sections per animal from lateral entorhinal cortex, with sampling extending from bregma –4.80 to bregma –8.72 (Paxinos and Watson 1998). In the medial entorhinal cortex, we sampled (n = 3) sections per animal from bregma –6.30 to bregma –9.16, as shown in Figure 2A. The subregion of interest was outlined by hand by a researcher who was blind to group conditions. Sections were averaged to obtain a single score for each animal.

**Statistics**

Behavioral data were compared across young and aged rats using repeated-measures analysis of variance (ANOVA) with trial block as a repeated measure and age as a fixed factor, followed by Bonferroni post hoc analysis. Learning indices and physiological data were compared across young, aged-unimpaired, and aged-impaired rats using ANOVA with a planned post hoc comparison. The planned comparison was designed to characterize differences between behaviorally impaired aged rats (AI) and young and aged behaviorally intact rats (Y+AU). Correlations between physiological measures and learning indices among the rats in the aged cohort were made using Pearson’s correlation analysis. Our rationale for including both aged-impaired and aged-unimpaired in the correlation analyses is described in detail by Baxter and Gallagher (1996). For all analyses, statistical significance was set at P < 0.05.

**Results**

**Aged Rats Exhibit a Range of Cognitive Abilities on a Task that Depends on Medial Temporal Lobe Function**

Twenty-four-month-old male Long-Evans rats navigate less effectively to the hidden platform in the Morris water maze, relative to young rats, during training trials (Fig. 1A; F1,12 = 12.64, P = 0.0001). The difference in performance on the first block of training trials represents greater improvement across those initial trials in young rats, as the aged rats performed on par with the young rats during the first trial of maze training (t_{12} = 0.25, P = 0.81). As reported elsewhere, aged rats in this study population exhibit individual differences as assessed using an index score derived from the mean distance from the platform location during probe trials (Fig. 1B; Gallagher et al. 1993). While a considerable number of the aged rats were unimpaired with low index scores similar to young adults, others in the aged cohort fell outside the range of young performance. Note that the scores for the young rats, index <240, in this study were in the normative range previously

**Figure 1.** Behavioral characterization on a task that recruits temporal lobe structures identifies a subpopulation of aged rats that are impaired relative to young adults. (A) Aged male Long-Evans rats follow a more circuitous path to the hidden platform in the water maze over the course of training. Differences in performance on the first block of trials are not detectable during the first trial (T1). Data points represent the mean of blocks of 5 trials, and error bars depict the standard error of the mean. (B), The learning index score, derived from the average distance from the platform location during interpolated probe trials (described in Materials and Methods and in Gallagher et al. 1993), distinguishes between aged-unimpaired (AU) and aged-impaired (AI) rats.
observed with this protocol (Gallagher et al. 1993). Using this index, we characterized the aged rats as either impaired or unimpaired relative to young for immunohistochemical analysis.

**Aged-Impaired Rats Have Fewer Reelin Immunoreactive Cells in the Lateral Entorhinal Cortex**

The lateral and medial subdivisions of the entorhinal cortex are readily apparent on coronal sections (shown schematically in Fig. 2A). Immunoreactivity for reelin delineates the layer II neurons of the entorhinal cortex (Figs 2B–D and 3A,B). Stereological analysis revealed that aged cognitively impaired rats have fewer reelin-positive cells in the lateral entorhinal cortex, relative to young rats and aged rats with preserved cognition (Fig. 3B,C; \( F_{2,26} = 4.51, P = 0.022 \)). There was no significant difference in reelin-positive cell numbers in the medial entorhinal cortex (\( F_{2,26} = 0.001, P = 0.99 \)).

Changes in the number of reelin-immunoreactive cells are unlikely to be attributable to loss of neurons, as previous work in this model (Rapp et al. 2002) and in Fischer rats (Merrill et al. 2001) has shown no change in total neuron number in the lateral entorhinal cortex with age-related cognitive impairment. Taken together with the current report, these observations would indicate a loss of reelin expression in neurons in the lateral entorhinal cortex of aged rats that are cognitively impaired. We also looked at the relationship between reelin-immunoreactive cell counts and behavioral performance among rats in the aged cohort. While spatial memory impairment, indicated by higher index scores, was generally related to reduced reelin immunoreactive neuron numbers in the lateral entorhinal cortex, this trend was not statistically significant (Pearson's \( r = -0.43, P = 0.075 \)).

![Figure 2](image2.png)

**Figure 2.** Reelin is a marker expressed by neurons in the superficial layers of the lateral entorhinal cortex. (A) Sagittal view of the rat brain, showing the anterior and inferior anatomical position of the lateral entorhinal cortex (LEC) and the posterior situation of the medial entorhinal cortex (MEC). The LEC is visible on coronal sections containing the dorsal hippocampus. As sections become more caudal, the LEC occupies more of the temporal cortex. On caudal sections that do not contain hippocampus, the MEC is visible at the medial edge of the cortex. Eventually, the MEC ascends to encompass the temporal cortical field. Image in (A) adapted from Paxinos and Watson (1998). (B) Low-magnification image showing reelin immunoreactivity in the superficial layers of the LEC. The diagram superimposed on the image is from Rapp et al. (2002). (C) Reelin staining in layer II is confined to the LEC, located inferior to the rhinal sulcus on coronal sections. No reelin staining is detected in layer II of perirhinal cortex (PER) surrounding the superficial portion of the rhinal sulcus. (D) Immunoreactivity for reelin among neurons in layer II of the lateral entorhinal cortex.

![Figure 3](image3.png)

**Figure 3.** Aged rats that are cognitively impaired have fewer reelin-immunoreactive cells in the lateral entorhinal cortex. (A) Representative photomicrograph showing reelin labeling in the entorhinal cortex of a young rat. (B) Higher magnification image showing reelin immunoreactivity in layer II of the entorhinal cortex in a representative young rat (Y), a representative aged-unimpaired rat (AU), and a representative aged-impaired rat (AI). (C) Aged-impaired rats have significantly fewer reelin-positive cells in the lateral entorhinal cortex (LEC), relative to young rats. No changes were apparent in the medial entorhinal cortex (MEC). Asterisk indicates significance at \( P < 0.05 \) following one-way ANOVA with planned post hoc comparison between behaviorally impaired rats (AI) and behaviorally intact rats (AU+Y). Error bars represent the standard error of the mean.

**Reelin mRNA Expression Is Reduced in the Lateral Entorhinal Cortex of Aged-Impaired Rats**

In agreement with the observed decrease in reelin-positive cells in lateral entorhinal cortex, a corresponding reduction in reelin mRNA expression was observed in a separate set of aged rats. Behavioral performance for this cohort of aged rats distinguished between aged rats that are cognitively impaired or unimpaired, relative to young rats (Fig. 4A). Aged rats that are cognitively impaired exhibit reduced reelin mRNA expression in the lateral entorhinal cortex (Fig. 4B,C; \( F_{2,16} = 19.94, P = 0.001 \)) with no change in reelin mRNA expression in the medial entorhinal cortex (\( F_{2,12} = 1.76, P = 0.21 \)). Reelin mRNA expression again followed an anatomical gradient, such that greater signal was detected in the lateral entorhinal cortex, relative to the medial entorhinal cortex (Fig. 4C). There was no effect of aging or cognitive status on mRNA for reelin’s intracellular signaling target, “disabled”-1 (Supplementary Fig. 1). No significant
correlations were found between reelin or DAB1 mRNA expression and the learning index among rats in this cohort of aged subjects.

**Alterations in Tau Phosphorylation with Age-Related Cognitive Impairment**

Tangles composed of hyperphosphorylated tau form earlier in the lateral entorhinal cortex, relative to the medial entorhinal cortex, during aging and neurodegenerative disease in humans (Arriagada et al. 1992; Braak H and Braak E 1995). While reelin negatively regulates tau phosphorylation (Hiesberger et al. 1999), there are multiple tau kinases and several modulatory factors that could underlie the regional susceptibility of the lateral entorhinal cortex during aging and AD. To evaluate another cellular event known to be a characteristic of the aging lateral entorhinal cortex, we measured phosphorylation of tau at serine 202 using immunohistochemistry.

Aged-impaired rats exhibit increased tau phosphorylation in the lateral entorhinal cortex, relative to young rats (Fig. 5A,B; $F_{2,26} = 11.03$, $P = 0.004$). Accumulation of phosphorylated tau was also correlated with behavioral performance among aged rats, such that a greater increase in tau phosphorylation was associated with worse behavioral performance (Fig. 5C; Pearson’s $r = 0.64$, $P = 0.004$). By contrast, there was no effect of aging or cognitive status on total tau immunoreactivity in the lateral entorhinal cortex ($F_{2,26} = 0.74$, $P = 0.49$; Supplementary Fig. 2). Insofar, as total tau expression reflects neuronal integrity, the absence of change in total tau immunoreactivity is consistent with other evidence that total neuron number is unchanged in the lateral entorhinal cortex of aged-impaired rats (Merrill et al. 2001; Rapp et al. 2002).
We observed no significant change in total tau expression ($F_{2,26} = 1.73, P = 0.20$; Supplementary Fig. 2) in the medial entorhinal cortex. There was a strong nonsignificant trend for the main effect of aging, independent of behavioral performance and cognitive status, on tau phosphorylation in medial entorhinal cortex (Fig. 5B; $F_{2,26} = 3.29, P = 0.055$). As expected based on that trend, tau phosphorylation in the medial entorhinal cortex did not correlate with behavioral alterations in the water maze among the aged rats (Pearson’s $r = 0.26, P = 0.31$).

**Loss of Reelin Immunoreactivity in the Lateral Entorhinal Cortex Is Accompanied by Local Reductions in Synaptic Marker Expression**

While neuron numbers are largely preserved throughout the hippocampal formation and medial temporal cortex in aged rats with cognitive impairment (Rapp and Gallagher 1996; Rapp et al. 2002), a circuit-specific loss of synapses has been found in aged rats with cognitive impairment involving the innervation to the hippocampus originating in layer II entorhinal cortex (Smith et al. 2000). Local intrinsic connections involving layer II collaterals also exist within the entorhinal cortex itself, through a network of ascending fibers traversing superficial layers. (Supplementary Fig. 3, and van Strien et al. 2009). We used synaptophysin immunoreactivity, which labels presynaptic terminals, to provide a preliminary estimate for loss of synaptic integrity that might be associated with intrinsic connections of layer II neurons.

The expression of reelin in neurons in the superficial layers of the entorhinal cortex using immunofluorescence is shown in Figure 6A. Fluorescence labeling for synaptophysin is also evident in layer II of the entorhinal cortex, surrounding the dendritic fields of reelin-labeled cells (Fig. 6A,B). Analysis of this labeling for synaptophysin showed a significant reduction in the lateral entorhinal cortex of aged-impaired rats (Fig. 6C; $F_{2,26} = 4.59, P = 0.02$). In contrast, there was no statistically significant effect of aging or cognitive status on synaptophysin immunoreactivity in the medial subregion ($F_{2,26} = 1.30, P = 0.29$). These data suggest that inputs to the superficial layers of the lateral entorhinal cortex are diminished, although the source of those inputs, either extrinsic or intrinsic, remains to be determined. However, we did observe that phospho-tau accumulation was inversely correlated with synaptophysin immunoreactivity (Pearson’s $r = -0.60, P = 0.008$; Fig. 6D). This is notable because tau phosphorylation has been linked to synaptic loss (Spires-Jones et al. 2009).

**Discussion**

We have identified a number of changes associated with individual differences in neurocognitive aging in the lateral entorhinal cortex. These changes include a decrease in synaptophysin immunoreactivity, which is associated with a decrease in reelin expression. This suggests a circuit-specific loss of synapses in the lateral entorhinal cortex that might be associated with intrinsic connections of layer II neurons.
entorhinal cortex. Reelin expression is reduced at both the protein and mRNA level in aged rats that are cognitively impaired. Changes in reelin expression occurred in the context of increased tau phosphorylation and reduced synaptic marker immunoreactivity. These changes followed a consistent anatomical pattern, such that the lateral entorhinal cortex emerged as a focal region in aged rats with impaired cognitive status.

Alterations in reelin expression, synaptophysin immunoreactivity, and tau phosphorylation occurred in the lateral entorhinal cortex of aged rats with spatial memory impairment. By contrast, aged rats with preserved behavioral performance exhibited few alterations compared with young adults. However, individual differences in this model of cognitive aging are not confined to spatial tasks. Aged rats that are behaviorally impaired in the water maze also exhibit reduced odor recognition memory (Robitsek et al. 2008), a function that involves the lateral entorhinal cortex (Young et al. 1997). In this regard, neurological alterations observed in the lateral entorhinal cortex in the current experiments could directly contribute to olfactory recognition deficits reported in aged rodents (LaSarge et al. 2007; Robitsek et al. 2008) and in humans with MCI and AD (Djordjevic et al. 2008). These changes might additionally exert downstream effects on hippocampal networks by disrupting entorhinal connectivity, which would alter the fidelity of information transmitted to the hippocampus.

The functional consequences of reduced reelin expression with age-related cognitive decline have yet to be elucidated, but the findings in the current study point to the lateral entorhinal cortex as a relevant network for such studies. Reelin is transported along perforant path axons and released within the terminal zones of entorhinal layer II afferents (Martínez-Cerdeño et al. 2003). During aging, synaptic plasticity at lateral perforant path synapses in the dentate gyrus is reduced (Froc et al. 2003), but the question of whether changes in reelin expression contribute to synaptic plasticity deficits has not been addressed. Moreover, although there is significant support for a reelinergic contribution to long-term potentiation at Schaffer collateral synapses in CA1 (for review, see Herz and Chen 2006), the association between reelin expression and synaptic plasticity at lateral perforant path synapses has not been characterized.

Reelin regulates neuronal migration during development and synaptic number and function during adulthood (Herz and Chen 2006). Transgenic overexpression of reelin in neurons during adulthood increases the number of synapses in regions of the hippocampus that receive input from layer II neurons (Pujadas et al. 2010). In contrast, heterozygous reeler mice, which exhibit a nearly 50% reduction in reelin levels, have fewer hippocampal synapses (Niu et al. 2008). Functionally, reelin enhances long-term potentiation and protects against deficits induced by application of β-amyloid (Durakoglugil et al. 2009). This opens the possibility that reelin produced by the entorhinal cortex could confer neuroprotection on target cells in the hippocampus, which would be compromised with loss of reelin expression during age-related cognitive decline.

The observation that AD mice have fewer reelin immunoreactive principal neurons in the entorhinal cortex (Chin et al. 2007) without overt loss of entorhinal neurons (Irizarry et al. 1997) suggests that reduced reelin expression may contribute to early synaptic alterations in AD. While previous studies examining differences in reelin expression in AD mouse models did not differentiate between the medial and lateral entorhinal cortices, the topography of changes in the terminal zones innervated by entorhinal subdivisions suggests particular lateral entorhinal sensitivity. The outer molecular layer, which receives input from the lateral entorhinal cortex, shows the earliest synaptic atrophy (Dong et al. 2007) and accumulation of amyloid deposits (Reilly et al. 2003). This synaptic zone is similarly subject to loss of connectivity in the same population of aged rats used in the current study (Smith et al. 2000). Here, we show that reduced reelin expression is also seen in the context of changes in synaptic marker expression in the lateral entorhinal cortex during normal age-associated cognitive decline.

Reduced reelin expression occurred in concert with increased tau phosphorylation and reduced immunoreactivity for the synaptic marker synaptophysin. The lateral entorhinal cortex exhibited specific vulnerability to these molecular alterations during cognitive impairment in aging. A lateral to medial entorhinal topography has long been recognized in human aging, as tangles initially appear in the transentorhinal cortex with progressive involvement of entorhinal layer II neurons along a lateral to medial gradient (Braak stages 1–2) (Braak H and Braak E 1995). Although neurofibrillary tangles do not occur in aged rats, increased phosphorylation of tau emerges in the lateral entorhinal region in older animals exhibiting cognitive decline. The current study is, to the best of our knowledge, the first report of increased tau phosphorylation in the aged rat. A prior report described a redistribution of phospho-tau from neuronal processes to the soma in aged rats (Niewiadomska et al. 2005) but did not quantify phosphorylated tau immunoreactivity in relation to cognitive status. Mice, in contrast with rats, do not display phosphorylated tau accumulation with aging unless expression of wild-type human tau is present under a transgene (Kimura et al. 2007). This distinction may be attributable to differential expression of tau isoforms between the rat and mouse (Takuma et al. 2003), with the rat more closely resembling human tau isoforms.

The current findings have revealed a particular susceptibility of the lateral entorhinal cortex in age-related cognitive impairment. While these observations are best viewed at present as correlates of behavior rather than mechanisms, we have begun to establish a framework for understanding lateral entorhinal vulnerability in neurocognitive aging. In our model, cognitive impairment with aging is associated with reduced reelin expression, increased tau phosphorylation, and synaptic loss in the lateral entorhinal cortex. Future studies will address the mechanistic interrelationships among these changes and their impact on cognitive function.

Supplementary Material
Supplementary material can be found at: http://www.cercor.oxfordjournals.org/.

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