Influence of Viral Vector–Mediated Delivery of Superoxide Dismutase and Catalase to the Hippocampus on Spatial Learning and Memory During Aging

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Abstract

Aims: Studies employing transgenic mice indicate that overexpression of superoxide dismutase 1 (SOD1) improves memory during aging. It is unclear whether the improvement is due to a lifetime of overexpression, decreasing the accumulation of oxidized molecules, or if increasing antioxidant enzymes in older animals could reduce oxidative damage and improve cognitive function. We used adeno-associated virus to deliver antioxidant enzymes (SOD1, SOD2, catalase [CAT], and SOD1+CAT) to the hippocampus of young (4 months) and aged (19 months) F344/BN F1 male rats and examined memory-related behavioral performance 1 month and 4 months postinjection. **Results:** Overexpression of antioxidant enzymes reduced oxidative damage; however, memory function was not related to the level of oxidative damage. Increased expression of SOD1, initiated in advanced age, impaired learning. Increased expression of SOD1+CAT provided protection from impairments associated with overexpression of SOD1 alone and appears to guard against cognitive impairments in advanced age. **Innovation:** Viral vector gene delivery provides a novel approach to test the hypothesis that increased expression of antioxidant enzymes, specifically in hippocampal neurons, will provide protection from age-related cognitive decline. Further, expression of multiple vectors permits more detailed investigation of mechanistic pathways. **Conclusion:** Oxidative stress is a likely component of aging; however, it is unclear whether increased production of reactive oxygen species or the accumulation of oxidative damage is the primary cause of functional decline. The results provide support for the idea that altered redox-sensitive signaling rather than the accumulation of damage may be of greater significance in the emergence of age-related learning and memory deficits. *Antioxid. Redox Signal.* 16, 339–350.

Introduction

THE BRAIN is highly sensitive to oxidative stress (22), and the accumulation of damaged molecules may contribute to age-related memory impairments (8, 14, 30, 37, 38). Antioxidant molecules and enzymes balance the biological activity of reactive oxygen species (ROS), superoxide ($O_2^{\bullet-}$), and hydrogen peroxide (H_2O_2). Superoxide dismutase (SOD) catalyzes $O_2^{\bullet-}$ into H_2O_2 and catalase (CAT) or glutathione peroxidase (GPx) converts H_2O_2 to water and oxygen. SODs are classified according to their metal cofactors and cellular localization. The Cu/Zn-SOD1 (SOD1) is distributed throughout the cytoplasm, nucleus, and inner membrane space of mitochondria, Cu/Zn-SOD3 (SOD3) is located in the extracellular space, and Mn-SOD (SOD2) is restricted to the mitochondrial matrix (11, 18, 28).

SOD1 overexpression differentially influences brain function over the course of aging (23). Enhanced long-term potentiation (LTP) and spatial memory are observed in aged transgenic

SOD1 (tg-SOD1) mice (25, 27). In contrast, tg-SOD2 mice do not exhibit altered synaptic plasticity or memory (23). It is unclear whether increased SOD1 activity is required over the lifespan to prevent the accumulation of oxidative damage; or whether enhanced SOD1 activity, initiated in aged animals, might be therapeutic. To test this hypothesis, we used adeno-associated virus (AAV) to deliver antioxidant enzymes (SOD1, SOD2, CAT, or SOD1 + CAT) to the hippocampus of young and aged rats. The results demonstrate a dissociation of learning from measures of oxidative stress and suggest that changes in redox-sensitive signaling may mediate cognitive impairment.

Results

Efficiency-specificity of the viral vectors

Following behavioral testing, a subset of animals was killed for examination of vector expression. Figure 1 illustrates the pattern of AAV-mediated transfection. Strong expression was

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FIG. 2. Overexpression of antioxidant enzymes in the hippocampus reduces markers of oxidative stress. Western blot analysis of hippocampal lysates from young and aged rats injected with viral vectors to express (A, E) SOD1-myc, (B, F) SOD2, (C, G) CAT, or (D, H) SOD1+CAT. For quantification, band intensities were normalized to the expression of glyceraldehyde 3-phosphate dehydrogenase (GAPDH) and this value was normalized to the mean value of young controls from the same blot. In each case, transduction resulted in an increase in the expression of the antioxidant enzyme. For SOD1-myc (A, D), two bands were observed representing endogenous rat SOD1 (rSOD1) and the human myc-tagged SOD1 (hSOD1). Lipid peroxidation (4-hydroxy-2-nonenal [HNE]) and S-glutathionylated proteins (glutathione [GSH]) were decreased by enzyme overexpression. To determine whether overexpression of antioxidant enzymes would influence expression of downstream antioxidant enzymes, immunostaining was performed for glutathione peroxidase 1 (Gpx1), Gpx4, and CAT. Overexpression of SOD1-myc in aged rats was associated with an increase in Gpx1 (E). Each bar represents the normalized mean $\pm$ SEM ($n=3-6$). Asterisks indicate significant ($p<0.05$) differences determined by Fisher's protected least significant difference (PLSD) *post hoc* test.

FIG. 3. Overexpression of antioxidant enzymes did not affect cue discrimination in water maze. All groups learned to reach the visible platform, as indicated by a significant overall decrease in latency (A, B) and distance (C, D) across blocks. Y=5-month-old and A=20-month-old rats injected with SOD1, SOD2, CAT, or SOD1+CAT. Age-matched controls (Cont) included rats injected with GFP and no-surgery controls. (To see this illustration in color the reader is referred to the Web version of this article at www.liebertonline.com/ars).

was reduced relative to injection of SOD1-myc (*i.e.*, two vs. fourfold).

As shown in Figure 2, SOD1, SOD2, CAT, or SOD1+CAT overexpression was associated with reduced 4-hydroxy-2-nonenal (HNE) staining. Quantification confirmed a decrease in HNE-reactive protein for SOD1 [$F(1,8)=9.26$, $p<0.05$], SOD2 [$F(1,12)=11.45$, $p<0.01$], CAT [$F(1,8)=18.53$, $p>0.01$], and SOD1+CAT [$F(1,8)=16.53$, $p<0.01$]. A decrease in protein S-glutathionylation was observed for rats overexpressing SOD1 [$F(1,12)=15.17$, $p<0.005$], SOD2 [$F(1,12)=7.83$, $p<0.05$], CAT [$F(1,8)=22.54$, $p<0.01$], and SOD1+CAT [$F(1,8)=7.3$, $p<0.05$]. Examination of SOD activity indicated that SOD1-myc rats exhibited a twofold increase in SOD activity compared with green fluorescence protein (GFP)-injected rats ($p<0.05$; $n=4$ /group, data not shown).

SOD overexpression can increase the level of H_2O_2 and the expression of downstream antioxidant enzymes (13, 19, 29, 33). Western blots indicate that GPx4 and CAT levels did not differ among groups; however, there was a tendency for a treatment effect for GPx1 [$F(1,15)=3.46$, $p=0.08$] and *post hoc* comparisons indicated GPx1 expression significantly increased in aged SOD1 rats (Fig. 2E).

Age-dependent influence of enzyme overexpression on spatial learning

Spatial learning in the water maze was tested 1 month following virus injections. Figure 3 shows the mean escape latency and escape path length across the five training blocks during cue discrimination training. All groups exhibited a decrease in escape latency and path length during training, and young animals exhibited superior latency and path length relative to aged rats. An analysis of variance (ANOVA) on latency indicated an effect of training [$F(4,404)=107.21$, $p<0.0001$] and an age effect [$F(1,404)=34.71$, $p<0.0001$] in the absence of a treatment effect (Fig. 3A, B). Similarly, an

ANOVA on path length indicated an effect of training [$F(4,404)=71.23$, $p<0.0001$] and age [$F(1,404)=40.084$, $p<0.0001$] in the absence of a treatment effect (Fig. 3C, D).

For spatial discrimination, an ANOVA on escape latency indicated significant main effects of training [$F(5,505)=68.03$, $p<0.0001$] and age [$F(1,505)=37.35$, $p<0.0001$], with a tendency for a treatment effect [$F(4,505)=2.10$, $p=0.08$] (Fig. 4A, B). Analysis of escape path length indicated effects of training [$F(5,505)=81.19$, $p<0.0001$] and age [$F(1,505)=28.83$, $p<0.0001$], with a tendency for a treatment effect [$F(4,505)=2.27$, $p=0.06$] (Fig. 4C, D). To localize treatment effects, the data for the last two training blocks were averaged and an ANOVA was run within each age group. No treatment effects were found in young rats. A tendency [$F(4,50)=1.92$, $p=0.12$] for a treatment effect on latency was observed for aged rats, and *post hoc* analyses indicated that aged SOD1+CAT rats exhibited significantly less time to reach platform compared with aged rats expressing SOD2 or SOD1 alone (Fig. 4E). The results were confirmed for the distance (Fig. 4F). An ANOVA on the mean for the last two blocks revealed tendency for a treatment effect in aged rats [$F(4,50)=2.5$, $p=0.05$] and *post hoc* analysis indicated that aged SOD1+CAT rats had a shorter path length compared with aged SOD2 or SOD1 rats.

A 60 s probe trial was delivered between training blocks 5 and 6. Analyses were performed on two consecutive 30 s segments. An ANOVA on the percent time in the goal quadrant during the first 30 s indicated that young rats spent more time in the goal quadrant than aged rats [$F(1,100)=4.82$, $p<0.05$] and there was a tendency for a treatment effect [$F(4,100)=2.07$, $p=0.08$] (Fig. 5A, B). ANOVAs within each age group revealed a treatment effect for aged animals [$F(4,50)=3.07$, $p<0.05$] and *post hoc* comparisons indicated that aged control rats spent more time in the goal quadrant compared with aged rats with SOD1 or CAT overexpression. Aged SOD1+CAT rats spent significantly more time in the goal quadrant relative to aged SOD1 rats. The percent time in

FIG. 4. Overexpression of SOD1 impaired spatial learning in aged rats. Behavioral measures for young and aged rats during training on the spatial version of the water escape task. Mean latency (A, B) and mean path length (C, D) to escape during spatial discrimination training. No treatment effects were found for young rats. Examination of the mean latency (E) and mean path length (F) averaged across the last two training blocks in aged animals indicated an effect of treatment. The SOD1+CAT group performed better relative to the SOD1 and SOD2 groups. Asterisk indicates a significant ($p < 0.05$) difference. (To see this illustration in color the reader is referred to the Web version of this article at www.liebertonline.com/ars).

the goal quadrant was compared with that expected by chance (*i.e.*, 25%). For the first 30 s of the probe trial, all groups except for aged SOD1 or CAT performed above chance. Examination of time spent searching the goal quadrant for the last 30 s of the probe trial indicated a tendency for an age effect [$F(1,100)=5.06$, $p=0.08$] in the absence of a treatment effect (Fig. 5C, D). An examination of the percent time in the goal quadrant indicated that all groups except aged controls and aged SOD1 rats spent significantly greater time in the goal quadrant relative to chance (Fig. 5C, D). The fact that aged SOD1 rats did not spend a significant portion of their search behavior in the goal quadrant for either segment of the probe trial indicates that these animals did not acquire a spatial search strategy.

The aged SOD1 impairment was confirmed by measuring the latency to the first platform crossing (Fig. 5E, F). An ANOVA indicated an age [$F(1,100)=10.95$, $p < 0.01$] and treatment [$F(4,100)=2.52$, $p < 0.05$] effect. ANOVAs within each age group indicated a tendency for a treatment effect in aged animals [$F(4,49)=2.38$, $p=0.06$] and *post hoc* comparisons indicated that aged SOD1 rats required a longer time to cross the platform location compared with age-matched controls, CAT, and SOD1+CAT expressing groups. There was a tendency ($p=0.08$) for a difference between SOD1 and SOD2 groups. An ANOVA on the total number of platform crossing

indicated an age difference [$F(1,100)=13.15$, $p < 0.001$] (Fig. 5G, H), in the absence of a treatment effect. In sum, the results indicate that SOD1 overexpression for 1 month was associated with impaired spatial learning in aged rats. The fact that the SOD1+CAT exhibited superior performance relative to SOD1 group indicates that the impairment maybe rescued to co-overexpressing CAT.

Overexpression of SOD1+CAT for 4 months improves spatial learning

Our results contrast with studies in tg-SOD1 mice, which indicate that young tg-SOD1 mice exhibit impaired spatial learning while aged tg-SOD1 mice show enhanced spatial memory (20, 27). The difference may result from an interaction of age and the duration of overexpression. To examine this possibility, a subset of animals tested at 1 month was retested at 4 months postinjection. The experimental groups included young rats (8 months) injected with SOD1 ($n=6$), SOD1+CAT ($n=10$), GFP ($n=2$), no-surgery controls ($n=4$), and aged rats (23 months) injected with SOD1 ($n=6$), SOD1+CAT ($n=10$), GFP ($n=2$), or no-surgery controls ($n=4$).

For examination of spatial learning 4 months after viral vector injections, the platform location in the maze was shifted to a new quadrant. An ANOVA on escape latency

FIG. 5. Overexpression of SOD1 impaired acquisition of a spatial search strategy in aged rats. A single probe trial was given between block 5 and block 6 of spatial training. Percent time spent searching the goal quadrant during the first 30 s for (A) young and (B) aged rats. The search behavior of aged rats with overexpression of SOD1 or CAT was not different from chance levels (25% dashed line). Aged SOD1 rats spent significantly less time compared with aged control and aged rats with overexpression of SOD1+CAT. Aged rats with overexpression of CAT spent significantly less time than aged control rats. Examination of the percent of time spent in goal quadrant during the second 30 s of probe trial for (C) young and (D) aged rats indicated that aged rats with overexpression of SOD1 continued to perform at chance levels. Examination of the latency for first platform crossing for (E) young and (F) aged rats indicated a treatment effect for aged animals due to an extended latency for the SOD1 group. Age differences were observed for total number of platform crossings between (G) young and (H) aged rats. Asterisk indicates group difference ($p < 0.05$). Pound sign (#) indicates a significant ($p < 0.05$) difference from chance. (To see this illustration in color the reader is referred to the Web version of this article at www.liebertonline.com/ars).

across training blocks indicated main effects of training [$F(5,185) = 40.53, p < 0.0001$] and age [$F(1,185) = 5.19, p < 0.05$], with a tendency for a treatment effect [$F(2,185) = 5.19, p = 0.07$] (Fig. 6A, B). Analysis of escape path length indicated effects of training [$F(5, 185) = 36.12, p < 0.0001$] and age [$F(1,185) = 5.20, p < 0.05$] (Fig. 6C, D). To localize treatment effects, the data for the last two training blocks were averaged and an ANOVA for treatment effects was run within each age group. An ANOVA for young animals indicated a tendency for a treatment effect [$F(2,18) = 7.12, p = 0.14$]; however, *post hoc* comparisons did not reach significance. Although there was a trend for young SOD1 rats to exhibit an increase in the escape latency relative to young control

rats ($p = 0.05$). An ANOVA for aged animals indicated a tendency for a treatment effect [$F(2,19) = 3.24, p = 0.06$] and *post hoc* comparisons indicated that aged SOD1+CAT rats required less time to reach platform compared with aged SOD1 rats (Fig. 6B). Analysis of the mean distance for the last two training blocks also indicated a tendency for a treatment effect in aged rats [$F(2,19) = 2.67, p = 0.09$] (Fig. 6C, D) and *post hoc* comparisons confirmed that aged SOD1+CAT rats had a shorter path length to escape compared with aged SOD1 rats (Fig. 6D).

Examination of the first 30 s of the probe trial delivered between blocks 5 and 6 indicated a tendency for young rats to spend more time in the goal quadrant relative to aged rats

FIG. 6. Long-term over-expression of SOD1+CAT improved learning in spatial trials in aged rats. Behavioral measures for young and aged rats during training on the spatial version of the water escape task 4 months after viral injections. Mean latency (A, B) and mean path length (C, D) to escape during spatial discrimination training. The bar graphs represent the mean latency averaged across the last two training blocks. For aged animals, better performance was observed for the SOD1+CAT group relative to the SOD1 group. Asterisk indicates a significant ($p < 0.05$) difference. (To see this illustration in color the reader is referred to the Web version of this article at www.liebertonline.com/ars).

[$F(1,37)=3.33$, $p=0.07$] in the absence of a treatment effect (Fig. 7A, B). The percent time searching the goal quadrant was significantly above chance for all groups, indicating that all groups had acquired a spatial search strategy. A tendency for a treatment effect [$F(2, 37)=2.47$, $p=0.09$] in the absence of an age difference was observed for the second half-minute of the probe trial (Fig. 7C, D). Again, all groups spent a significantly longer time in the goal quadrant than expected by chance (Fig. 7C, D). An ANOVA on the latency to the first platform crossing indicated a significant age effect [$F(1,37)=12.73$, $p<0.01$], and a tendency for a treatment effect [$F(2,37)=2.24$, $p=0.1$]. An ANOVA within each age group indicated a tendency for a treatment effect only in the aged group [$F(2,19)=3.29$, $p=0.05$]. *Post hoc* comparison indicated that aged SOD1+CAT rats required less time for the first platform crossing compared with aged SOD1 rats and there was a tendency ($p=0.08$) for SOD1+CAT rats to cross quicker than aged controls (Fig. 7E, F). Finally, an ANOVA on total platform crossing indicated an age effect [$F(1,37)=12.67$, $p<0.005$] and a tendency for a treatment effect [$F(2,37)=2.54$, $p=0.09$]. An ANOVA within each age group indicated a tendency for a treatment effect in aged rats [$F(2,19)=3.45$, $p=0.05$] and *post hoc* comparisons indicated that aged SOD1+CAT rats exhibited significantly more platform crossings than aged controls (Fig. 7G, H). In fact, aged SOD1+CAT rats exhibited performance similar to young animals.

Figure 8 shows Western blot data on enzyme expression and lipid peroxidation in the hippocampus for animals that overexpressed viral vectors for ~5 months. As shown in Figure 8A, the two SOD1 bands (19 and 23 kDa) were observed for SOD1-myc-injected rats. Densitometry confirmed that rats injected with SOD1-myc exhibited approximately threefold increase in total SOD1 [$F(1,12)=133.08$, $p<0.0001$] (Fig. 8A, C). Again, the 19 kDa band was not influenced by SOD1-myc expression (data not shown). Injection of SOD1+CAT increased the expression of SOD1 [$F(1,12)=31.31$, $p<0.001$] and CAT [$F(1,8)=10.40$, $p<0.05$] (Fig. 8B, D). Similar to 2-month overexpression, the level of expression for SOD1

under this condition was reduced relative to injection of SOD1-myc alone (*i.e.*, 2 vs. 2.5-fold).

A decrease in HNE-reactive protein was observed for rats that overexpress SOD1 [$F(1,12)=24.23$, $p<0.001$] or SOD1+CAT [$F(1,12)=7.63$, $p<0.05$] (Fig. 8). For SOD1 rats, there was an age effect on CAT expression [$F(1,12)=6.58$, $p<0.05$] in the absence of a treatment effect. *Post hoc* tests within each treatment group indicated an age-related decrease in expression in animals overexpressing SOD1 (Fig. 8C). Examination of GPx1 indicated no effect of age or treatment for SOD1 animals and SOD1+CAT animals exhibited an effect of age [$F(1,16)=15.00$, $p<0.005$] and treatment [$F(1,16)=4.27$, $p<0.05$]. *Post hoc* analyses indicated that GPx1 expression was reduced in aged rats that overexpressed SOD1+CAT [$F(1,8)=7.9$, $p<0.05$] compared with aged controls (Fig. 8B, D). Levels of protein carbonyls were measured; however, the results did not reveal any information concerning the effect of SOD1 overexpression, possibly due to the lack of specificity of carbonyl measures (1) or the fact that H_2O_2 is a poor mediator of stable oxidative damage for protein carbonyls (43). Further, previous research indicates that lipid peroxidation may be more sensitive since SOD overexpression in mice markedly reduces lipid peroxidation in the absence of a significant effect on protein carbonyls (24). Consistent with the lipid peroxidation results, overexpression of SOD1 resulted in a decrease in Oxo8dG [$F(1,21)=16.58$, $p<0.0005$] in the absence of an age difference (Fig. 9), confirming that SOD1 overexpression decreased oxidative damage.

Discussion

While it is clear that oxidative stress is a likely component of aging, it is unclear whether increased production of ROS or the accumulation of oxidative damage is the primary cause of functional decline. Consistent with previous work in tg-mice, overexpression of SOD1, SOD2, and CAT reduced markers of oxidative stress (41). However, decreased oxidative damage was not associated with improved cognitive function.

Moreover, SOD1 overexpression impaired spatial learning in aged rats. The impairment was specific to SOD1 and spatial learning in aged animals. The results indicate that accumulation of oxidative damage is not the only mechanism for cognitive decline. Other processes influenced by oxidative

stress may be more relevant to age-related cognitive decline, including mitochondrial function, redox signaling, and inflammation.

SOD1 can be toxic to motor neurons (7). However, in the current study, the deficit was limited to aged SOD1 animals, indicating that it was not simply due to SOD1 overexpression. Further, one would expect that toxicity should increase over time. On the contrary, we observed that the spatial learning deficit decreased following 4 months of SOD1 overexpression in our oldest rats. Although there was a tendency for the emergence of impairment as young SOD1 rats approached middle age (Fig. 6). SOD1 catalyzes $O_2^{\bullet -}$ into H_2O_2 and the levels of $O_2^{\bullet -}$ and H_2O_2 in brains of Tg-SOD1 mice are decreased and increased, respectively (35), and tg-SOD1 overexpressing mice exhibit age-dependent effects on cognition and hippocampal synaptic plasticity that are thought to result from elevation of ambient H_2O_2 levels (26). Thus, the specificity may be due to ambient H_2O_2 levels. One indication that H_2O_2 was increased in the current study is that aged SOD1 animals exhibited increased GPx1 expression (Fig. 2A). In other tissues, H_2O_2 can provoke an increase in GPx activity (39, 42, 45). Finally, aged SOD1+CAT rats did not exhibit cognitive impairments, suggesting improved cognition resulted from better homeostatic regulation of redox signaling due to CAT processing of excess H_2O_2 .

How can SOD1 overexpression reduce oxidative damage if H_2O_2 levels increase? H_2O_2 is a relatively mild oxidizing agent. Irreversible oxidative damage results from $O_2^{\bullet -}$; SOD1 removes $O_2^{\bullet -}$ and overexpression of SOD1 reduces the level of $O_2^{\bullet -}$ in the brain (35). If free iron is available, H_2O_2 can produce hydroxyl radicals ($OH^{\bullet}$) through Fenton-type reactions. However, a large body of evidence indicates that H_2O_2 *per se* may not induce irreversible oxidative damage (10, 31, 32, 43). In contrast, H_2O_2 can readily and reversibly influence redox-sensitive signaling cascades. Age-related changes in redox state disrupt calcium homeostasis, increasing the release of calcium from internal stores, decreasing CaMKII activity, and impairing LTP (5, 6, 15). H_2O_2 application to hippocampal slices mimics age-related changes, promoting calcium release from internal stores (21), decreasing synaptic CaMKII activity (44), and impairing LTP (3, 26, 40). The LTP impairment in tg-SOD1 mice is attenuated by CAT, suggesting the involvement of elevated H_2O_2 (20). Together the

FIG. 7. Overexpression of SOD1+CAT for 4 months improves spatial learning in probe test. A single probe trial was given between block 5 and block 6 of spatial training. Mean percent of time spent searching the goal quadrant during the first 30s for (A) young and (B) aged rats. Mean percent of time spent in goal quadrant during the second 30s of probe trial for (C) young and (D) aged rats. Examination of the latency for first platform crossing for (E) young and (F) aged rats indicated a treatment effect for aged animals due to a superior performance by the SOD1+CAT group compared with the SOD1 group. Examination of the total number of platform crossings for (G) young and (H) aged rats indicated that aged SOD1+CAT rats exhibited significantly more platform crossings compared to aged control. Asterisk indicates group difference ($p < 0.05$). Pound sign (#) indicates a significant ($p < 0.05$) difference from chance. (To see this illustration in color the reader is referred to the Web version of this article at www.liebertonline.com/ars).

FIG. 8. Antioxidant enzymes and oxidative stress markers in hippocampus with 5-month SOD1 and SOD1+CAT overexpression. Western blot analysis of hippocampal lysates from young and aged rats injected with viral vectors to express (A) SOD1 and (B) SOD1+CAT. For SOD1 (A, C), two bands were observed representing endogenous rat SOD1 (rSOD1) and the human myc-tagged SOD1 (hSOD1). CAT level did not change in SOD1 rats but increased in SOD1+CAT rats. Lipid peroxidation measured with anti-HNE antibody was decreased by SOD1 and SOD1+CAT overexpression. The expression of SOD1 did not change the level of GPx1 but the expression of SOD1+CAT in aged rats was associated with a decrease in GPx1 (D). GAPDH was used as a loading control. Asterisk indicates a significant ($p < 0.05$) difference in treatment. ω indicates a significant ($p < 0.05$) difference in age.

FIG. 9. DNA oxidative damage is reduced by overexpression of SOD1. The level of 8-oxo-7,8-dihydro-2-deoxyguanosine (8-oxodGuo) was measured and normalized by young controls (Y-Cont). Bars represent the mean and SEM. Asterisk indicates significant ($p < 0.05$) differences in treatment.

results suggest that SOD1 overexpression could impair learning through increased H_2O_2 acting on redox signaling pathways involved in memory formation.

Adaptation to altered redox state

The effect of SOD overexpression on memory changes with age (23). SOD1 overexpression impaired or had no effect on learning in young mice and enhanced spatial memory in older mice (25, 27). The authors suggest that aged mice adapt to elevated H_2O_2 . Transcription of genes regulating redox homeostasis increases during middle age (2, 4, 47). Thus, the increase in GPx1 expression in aged SOD1 mice may represent an adaptive response. Other adaptive changes may underlie the improvement in behavior following 4 months of SOD1 overexpression. Finally, aged SOD1+CAT rats exhibited superior water maze performance, indicating that treatment to reduce $O_2^{\cdot -}$ and H_2O_2 may be more beneficial than simply enhancing SOD1. Indeed, administration of a SOD/CAT mimetic improved memory in aging mice (34). Future studies should employ redox proteomics to identify specific changes related to cognitive decline.

In summary, memory function was not related to the level of oxidative damage. Increased SOD1 expression impaired learning in older rats. In contrast, increased expression of SOD1+CAT provided protection from age-related cognitive impairments. The results extend a growing body of evidence indicating the importance of H_2O_2 and the redox signaling for regulating cognitive function during aging.

Materials and Methods

Animals

All treatments were approved by the Institutional Animal Care and Use Committee and were in accordance with guidelines established by the United States Public Health Service Policy on Human Care and Use of Laboratory Animals. Young tg-SOD1 mice exhibit impaired spatial learning (20); therefore, young rats were included to examine age effects and specificity of gene overexpression. AAV was injected into the hippocampus of young (4 months) and aged (19 months) male Fischer 344/Brown Norway F1 rats

(Harlan Laboratories) to express SOD1 (young: $n=14$; aged: $n=13$), SOD2 (young: $n=9$; aged: $n=9$), CAT (young: $n=10$; aged: $n=9$), mix of SOD1 + CAT (young: $n=11$; aged: $n=12$), or GFP (young: $n=8$; aged: $n=8$). We also include young ($n=4$) and aged ($n=4$) no-surgery controls. No behavioral difference was observed between GFP and no-surgery controls; therefore, these animals were combined for behavioral analysis.

AAV viral vectors

Virus particles were produced and quantified by dot blot analysis, through the Powell Gene Therapy Center at the University of Florida (Gainesville, FL). Genes were cloned into pseudotyped AAV5 capsid, self-complementary AAV vectors, containing a cytomegalovirus (CMV)/chicken β -actin promoter with a CMV-immediate early (CMV-IE) enhancer. Genes included GFP, human SOD1 with a c-terminal myc tag (myc). The SOD2 and CAT vectors were gifts from Dr. Nicolas Muzyczka.

Rats were anesthetized with ketamine/xylazine (90/10 mg/kg) and virus was stereotactically injected at two sites bilaterally in the hippocampus using glass pipettes. Each injection consisted of 2 μ l of GFP (dot blot titer: 1.02×10^{13} vg/ml), SOD1 (dot blot titer: 1.14×10^{13} vg/ml), SOD2 (dot blot titer: 6.99×10^{12} vg/ml), CAT (dot blot titer: 2.53×10^{13} vg/ml), or 3 μ l 2:1 mix of SOD1 and CAT.

Behavior testing

Methods for water maze testing have been published previously (16). Rats were first trained to find a visible platform using 15 trials separated to five blocks. Rats that did not reach the platform within 60 s on all trials during the fifth block were removed from the experiment. Standard water maze testing across several days is insensitive to cognitive decline across this age span in Fischer 344/Brown Norway F1 rats (46). Therefore, the task difficulty was increased by employing a single-day massed training protocol, which is sensitive to age (9). Spatial discrimination testing occurred 3 days later and consisted of 18 trials separated into six blocks. A free swim probe trial was inserted between blocks 5 and 6. For the probe trial, the number of platform crossings was counted and the time spent in both goal and opposite quadrants was recorded.

Western immunoblotting

The hippocampi were frozen in liquid nitrogen, and stored at -80°C . Tissue was prepared in RIPA buffer supplemented with protease inhibitor, phosphatase inhibitor, and EDTA (Thermo Scientific). The lysates were assayed for protein content using BCA kit (Pierce) and then separated on polyacrylamide gels (Bio-Rad Laboratories) and transferred to PVDF membranes (GE Healthcare) for Western blotting. Primary antibodies (SOD1, 1:5000; SOD2, 1:6000; HNE, 1:100; CAT, 1:1000; GPx1, 1:600; GPx4, 1:1000 [Abcam, Inc.]; GAPDH, 1:6000 [EnCor Biotechnology, Inc.]; glutathione, 1:1000 [Arbor Assays]) were diluted in blocking buffer (5% milk in tris-buffered saline Tween-20) and applied to the membrane overnight at 4°C . Membranes were then incubated with horseradish peroxidase-conjugated secondary antibodies directed against the primary antibody (Cell Signaling Technology, Inc.). Membranes were reacted with and enhanced chemiluminescent

substrate (Pierce Biotechnology). A medical film processor (SRX-101A; Konica Minolta Medical Imaging U.S.A., Inc.) was used to image the film.

Immunohistochemistry

Brains were postfixed in 4% paraformaldehyde followed by 30% sucrose in PBS (4°C). Sections (10–20 μm) were incubated with primary antibodies (SOD1, 1:1000; MAP2, 1:10,000; CAT, 5 $\mu\text{g}/\text{ml}$; SOD2, 1:500 [Abcam, Inc.]; myc-tag, 1:2000 [Cell Signaling Technology]; NeuN, 1:1000 [Chemicon/Millipore]; Iba1, 1–2 $\mu\text{g}/\text{ml}$ [Wako]; GFAP, 1:500 [DakoCytomation]; anti-COX, 1:300 [Invitrogen]) overnight at 4°C . The brain sections were then washed and incubated in corresponding Alexa 488 or 594 secondary antibodies (Molecular Probes) for 1 h at room temperature. Sections were washed and counterstained with 4'-6-diamidino-2-phenylindole solution (0.1 $\mu\text{g}/\text{ml}$ in PBS) before mounting. Expression was confirmed using fluorescent microscopy (Zeiss Axioplan 2 upright fluorescent microscope, equipped with a QImaging Retiga 4000R Camera with RGB-HM-5 Color Filter and QImaging QCapture Pro 6.0 software; QImaging Surrey).

SOD activity was measured using the HT SOD assay kit (Trevigen, Inc.) and protein carbonyls were measured using a commercial ELISA (Zentech PC Test, Protein Carbonyl Enzyme Immuno-Assay Kit; Biocell Corp.) according to the manufacturer's instructions. 8-oxo-7,8-dihydro-2-deoxyguanosine (8-oxodGuo) levels were determined according to our previously published methods (12).

Statistical analysis

ANOVAs were used to establish main effects and interactions. Follow-up ANOVAs and/or Fisher's protected least significant difference *post hoc* comparisons with $p < 0.05$ were employed to determine specific differences. Student's *t*-tests were used to determine whether quadrant search behavior was different than that expected by chance.

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Author Disclosure Statement

No competing financial interests exist.

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Abbreviations Used

8-oxodGuo = 8-oxo-7,8-dihydro-2-deoxyguanosine
 AAV = adeno-associated virus
 ANOVAs = analyses of variance
 CAT = catalase
 COX = OxPhos complex IV subunit I
 GAPDH = glyceraldehyde 3-phosphate dehydrogenase
 GFAP = glial fibrillary acidic protein
 GFP = green fluorescence protein
 GPx = glutathione peroxidase
 GSH = glutathione
 HNE = 4-hydroxy-2-nonenal
 H₂O₂ = hydrogen peroxide
 LTP = long-term potentiation
 NeuN = neuronal nuclei
 PLSD = protected least significant difference
 ROS = reactive oxygen species
 SOD = superoxide dismutase
 tg-SOD = SOD transgenic mice