

Decreased mean diffusivity in the brain tissue of aged rats with learning deficits

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Introduction

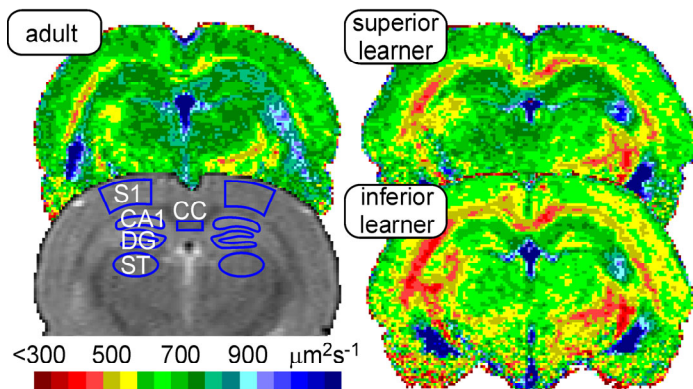
Aging as well as many neurodegenerative diseases is accompanied by serious cognitive deficits, particularly impaired learning and memory loss. Cognitive decline in old age has been linked to changes in brain anatomy, morphology, volume, and functional deficits (1). Nervous tissue, particularly in the hippocampus and cortex, is subject to various degenerative processes, including a decrease in the number and efficacy of synapses, neuronal loss, astrogliosis, and changes in extracellular matrix proteins. These and other changes not only affect the efficacy of signal transmission at synapses, but could also affect extrasynaptic (volume) transmission, mediated by the diffusion of transmitters as well as other substances through the volume of the extracellular space (2). Therefore, we studied diffusion of water in the brain tissue of aged rats with and without a learning deficit.

Subjects and methods

Experiments were performed *in vivo* on 4- (adult) and 22-month-old (aged) Wistar male rats. Diffusion-weighted (DW) MRI was used to determine the mean diffusivity $\langle D \rangle$ and fractional anisotropy FA (3) in different brain regions of behaviorally characterized rats. Prior to MR measurements, the ability to cognitively process spatial information was tested in a Morris water maze (4) and *superior* and *inferior* learners were selected according to their escape latencies (5). DW images were obtained with diffusion weightings (*b*-factor) of 75, 499, 1235 and 1732 s/mm². Diffusion weighting was applied along seven non-collinear directions.

Results

The aged superior learners did not show significantly longer escape latencies (28 ± 2 s) than adult rats (29 ± 1 s), but the latencies were significantly different in inferior learners (45 ± 2 s). We evaluated mean diffusivity and fractional anisotropy in the primary somatosensory cortex, corpus callosum, CA1 region of the hippocampus, dentate gyrus and striatum (Figure 1). In all these regions, the mean diffusivity was significantly lower in inferior learners when compared either to adult rats or superior learners. There were no such differences between adult rats and aged superior learners except in the corpus callosum. The values are summarized in the Table 1. We have not found any change in fractional anisotropy related to learning abilities. However, an age-related decrease in FA in the somatosensory cortex was observed (from 0.179 ± 0.007 , N = 8 in adult to 0.153 ± 0.005 , N = 16 in aged rats).



| | adult $\langle D \rangle \mu\text{m}^2\text{s}^{-1}$ | superior learners $\langle D \rangle \mu\text{m}^2\text{s}^{-1}$ | inferior learners $\langle D \rangle \mu\text{m}^2\text{s}^{-1}$ |
|-----------------|---|---|---|
| cortex | 654 ± 8 | 637 ± 7 | $613 \pm 4^{*†}$ |
| corpus callosum | 602 ± 15 | $546 \pm 5^*$ | $516 \pm 10^{*†}$ |
| CA1 | 705 ± 10 | 719 ± 4 | $685 \pm 6^{*†}$ |
| dentate gyrus | 645 ± 10 | 649 ± 5 | $605 \pm 5^{*†}$ |
| striatum | 670 ± 13 | 658 ± 5 | $611 \pm 8^{*†}$ |

Figure 1: Diffusivity maps acquired in the brain of aged superior and inferior learners and in adult control. Note the decrease of diffusivity in the inferior learner's brain. The mean values of $\langle D \rangle$ were calculated in the delineated areas: primary somatosensory cortex (S1), corpus callosum (CC), CA1 region of the hippocampus, dentate gyrus (DG) and striatum (ST).

Table 1: Mean diffusivity in aged rats, divided into two subgroups according to their learning abilities, and in adult rats (controls). All groups of animals shown in the table consist of 8 rats, data are expressed as mean \pm S.E.M. Significant differences (two-tailed Student's t-test, $p < 0.05$) compared to adult rats or superior learners are marked with asterisks or crosses, respectively.

Discussion and conclusion

Fractional anisotropy is not a sensitive measure of the structural changes found previously in the hippocampus of aged inferior learners (5). Decreased mean diffusivity was found in aged inferior learners but not in aged superior learners, whose mean diffusivity and performance in the Morris water maze were essentially indistinguishable from those of much younger adult animals. Thus, our data suggest that learning deficits are not only an inevitable result of the aging process, but that decreases in mean diffusivity such as those observed in inferior learners can significantly influence extrasynaptic transmission and thereby contribute to age-related learning deficits.

References and acknowledgement

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