

Melatonin prevents neuroinflammation in the hippocampus of aged mice

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Abstract

Aging is accompanied by deterioration in cognitive functions together with a variety of neurobiological changes including neuroinflammation. Low-grade chronic up-regulation of pro-inflammatory cytokines occurs during ageing. Melatonin is a key endogenous indoleamine secreted and released by the pineal gland; it plays a critical role in the regulation of circadian rhythms. Moreover, melatonin acts as a potent free radical scavenger, anti-inflammatory and numerous other functions. In the present study we examined the neuroinflammation status in aged mice. The result showed that the level of integrin α M (CD11b), Glial fibrillary acidic protein (GFAP) and major proinflammatory cytokines significantly increased in hippocampus of 22-month aged mice compared with 2 month-old mice. In order to examine the effect of melatonin on this neuroinflammation status in aged mice, melatonin was given in the drinking water at a dose of 10 mg/kg/day, started from 16-month old for 6 months. Our results revealed that melatonin significantly attenuated the increase of CD11b and GFAP in aged-mouse hippocampus. Moreover we found that melatonin could attenuate reduction of major proinflammatory cytokine levels in aged-mouse hippocampus. The results suggested that melatonin attenuated age-related neuroinflammation through reduction of glial activation.

Keywords: astrocyte, brain aging, inflammatory cytokines, melatonin, microglia,

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