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Mar 29, 2018 - download 0 0 1 2 5 8 9 8 9 8 9 9 9 9 9 8 9 9 8 8 9 8 9 8 9 8 9 9 8 8.The anti-inflammatory effect of 2-acetoxy-4-tetrahydro-4-methyl-7-(4-hydroxy-3-methoxy-5-methyl-2-pyrrolidinyl)-dihydro-isoquinoline-3-carboxylic acid, a cyclooxygenase-2 selective inhibitor. 2-Acetoxy-4-tetrahydro-4-methyl-7-(4-hydroxy-3-methoxy-5-methyl-2-pyrrolidinyl)-dihydro-isoquinoline-3-carboxylic acid (BW755C, R(+)-1, R(+)-2, and R(+)-3), a cyclooxygenase (COX)-2 selective inhibitor, suppressed xylene-induced ear oedema in rats in a dose-dependent manner. Treatment with R(+)-1, R(+)-2, and R(+)-3 also attenuated carrageenan-induced paw oedema in rats, and the anti-inflammatory effects were as potent as those of indomethacin, a classic COX inhibitor. The effective dose of R(+)-1 for mouse ear oedema and rat paw oedema (48 and 30 h after carrageenan treatment, respectively) was less than that of indomethacin. The in vivo antinociceptive effect of R(+)-1 was also investigated in various mouse models. R(+)-1 potently inhibited acetic acid-induced writhing in mice, but not in rats. In a hot plate test, R(+)-1, R(+)-2, and R(+)-3 had an antinociceptive effect on male ICR mice, although the effect was less than that of the positive control, morphine. On the other hand, in a tail-flick test, R(+)-1, R(+)-2, and R(+)-3 did not show any antinociceptive effect on male ICR mice. These results suggest that COX-2 and cyclo-oxygenase-1 (COX-1), which is responsible for the inhibitory effect f678ea9f9e

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