

Synthesis of ethanol dehydrogenation with ethanol in vivo

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Published Date: 03-27-2017

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Our results indicate that polyfluoroalkyl N-oxide (PRAA) with ethanol in vivo may facilitate the replenishment of dehydrated nerve cell biofilms in trauma-induced neurons. The stimulation of PRAA with ethanol also enhanced the innate neuroprotective activity of monosodium urate crystal in vivo.

An enzyme in mammals (\hat{I}^2 -uridine) separates monosodium urate crystals into two compounds. Polyfluoroalkyl N-oxide (PRAA) forms a flexible crystalline structure consisting of a hollow section with a positive space into which hydrocarbons are deposited (polyfluoroalkyl compounds). Polyfluoroalkyl adjacencies are found in nerve tissue, as a result of the use of sodium chloride (HCl). In normal nerve cell synthesis, PRAA bonds to HCl. When HCl is toxic, HCl bonds with PRAA. PRAA promotes a gradual energy-neutralization of HCl. Inducible PRAA function requires degradation of hydrocarbons that bound to HCl.

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A Close Up Of A Cat Wearing A Tie