

[Food Hydrocolloids, 4, 323 (1990)]

**The Decrease of Thaumatin's Sweetness Intensity Upon Interaction with Carrageenan.**

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Carrageenan was added at various ratios to thaumatin, a sweet protein, and the interactions between thaumatin and  $\lambda$ -,  $\kappa$ - and  $\iota$ -carrageenan were investigated from the following standpoints: pH, turbidity at 550nm, CD spectral change ( $\Delta\epsilon$ ) and the decrease in sweetness intensity. Decrease in thaumatin sweetness intensity was observed with lower pH (<4) and higher carrageenan concentrations in the following order:  $\lambda \approx \iota > \kappa$ -carrageenan. There was no correlation between sweetness intensity decrease and turbidity. With the increase in carrageenan concentration, CD spectral change ( $\Delta\epsilon$ ) was distinctly observed at pH 3-4. Thaumatin's sweetness intensity decreased simultaneously.

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**Interaction of Thaumatin with Carrageenans. II. Effects of pH, Temperature and Competing Cations studied by Circular Dichroism.**

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Interaction of thaumatin with carrageenans was studied spectroscopically on the basis of circular dichroism (CD) at pH 3-7 and 20-60°C and examined from estimating the sweetness of the complex. The most noticeable reduction in sweetness intensity of thaumatin was pH 3-4 and less sweetness reduction occurred at pH 5-7. Thaumatin was completely dissociated from a thaumatin-carrageenan complex at pH 5-7 when a salt was added to the complex, but no complete dissociation occurred at pH 3-4. The results suggest that the complex formation at pH 5-7 is due to electrostatic bonding (ionic bonds), but that at pH 3-4 another bonding mechanism is also involved.

[Free Radical Biol. Med., 8, 25 (1990)]

**A One-step Enzyme Immunoassay for Human Manganese Superoxide Dismutase with Monoclonal Antibodies.**

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A one-step enzyme immunoassay for the determination of manganese superoxide dismutase in serum has been developed with two kinds of monoclonal antibodies. Proposed method had high sensitivity (assay range, 0.4~200 ng/ml), good recovery (recovery percentage, 102.9~106.2%) and reproducibilities (intraassay, C.V.=1.87~3.66%; interassay, C.V.=3.03~10.4%). From these results, it is possible to apply this method to routine clinical analysis and biochemical research with various purposes.