Taste Behavior of Ornithyltaurine, a Salty Peptide, and an Attempt to Synthesis of a New NaCl Substitute

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Ornithyltaurine · hydrochloride (OTA · HCl) produce a good saltiness. Last year, a convenient synthesis of OTA was developed and we can obtained pure OTA easily. The product, which was almost tasteless by itself, produced a good saltiness without Na<sup>+</sup> in the presence of HCl. The best quality of saltiness was obtained by adding of 1.2 equiv. of HCl to the OTA solutions. The OTA·1.2HCl was equally salty to NaCl on a molar basis and also had an enhancing effect on saltiness of NaCl. The intake of Na<sup>+</sup> could be cut by 95% in a model system by using OTA·1.2HCl. OTA produced a variety of tastes by a kind of acids for neutralization. A good saltiness similar to OTA·HCl was produced by combinations of inorganic acids and OTA. And combinations of organic acids with big molecular weight produced TNS (Taste of Neutral Salt) and Umami taste as well as saltiness. Especially, OTA-Glu produced a favorable taste mixed OTA·HCl like saltiness with Glu·Na like Umami. OTA · 1.1Glu was the optimum condition of the taste, but the Umami was much stronger than the saltiness. OTA · 0.75HCl · 0.25Glu produced an excellent quality of taste containing *Umami* and saltiness, and the taste was almost the same as that of solutions containing 85.6 mM (0.5%) of NaCl and 14.7 mM (0.25%) of Glu Na. In this model system, the intake of Na<sup>+</sup> from not only NaCl but also Glu·Na could be cut completely using OTA· 0.75HCl 0.25Glu. To develop more effective salty compound, we searched a compound which can enhance tastes. When amino acids, i.e., Ala, Orn, and Glu with sweetness, TNS and *Umami*, respectively, introduced to the 2-NH, on D-Glucosamine, each taste were remained and the sweetness and TNS were enhanced. The sweetness and TNS were much enhanced by introducing Ala and Orn to 6-OH and 2-NH, on D-Glucosamine. sugar-amino acid complex has an enhancing effect of taste of introduced amino acid(s) and we can expect to develop an effective salty compound using sugar-amino acids (or peptides) complex.