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**STUDIES ON THE ACETYLCHOLINE-INDUCED  
EXOCRINE RESPONSES IN THE ISOLATED  
AND PERFUSED RAT PANCREAS**

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Continuous stimulation with acetylcholine (ACh,  $3 \times 10^{-7}$  M) induced 3 phases of amylase release: transient maximum release in the first phase; continuous release in the second phase lasting about 40 min; and declining release in the third phase. These 3 phases of amylase release were uniformly nullified when  $5 \times 10^{-6}$  M atropine was added to the perfusing solution 10 min prior to the initiation of continuous ACh stimulation. In a Ca-deficient environment, the amount of amylase release was inhibited, and the release decayed rapidly. The ACh-induced amylase release was little affected by the removal of  $\text{HCO}_3^-$ . The amount of amylase release rose when the concentration of ACh was increased. A dose-response relation was found between the amount of amylase release and the concentration of ACh over the range from  $10^{-8}$  M to  $3 \times 10^{-7}$  M. The addition of a low dose of cholecystinin-pancreozymin (CCK-PZ, 1.0 m-U/ml) shifted the dose-response relation to the left. The addition of a low dose of secretin (Sc, 1.0 m-U/ml) also shifted the dose-response relation to the left. The increase in amylase release during continuous stimulation with ACh was usually accompanied by a concomitant increase in the pancreatic juice flow. These results were analysed using Michaelis-Menten kinetics. These results may be explained by postulating a receptor complex composed of receptor subunits and an ionophore subunit. The receptor subunits may be activated by either one of these secretagogues, and, in turn, may activate the ionophore subunit.