The Study on Mucin Release by Airway Goblet Cells in Primary Culture

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Surface epithelial cells isolated from hamster tracheas and grown on a thick collagen gel become a highly enriched population of mucus-secreting cells. Epithelial cells from tracheas of hamsters were collected using enzymatic procedures and cultured under various conditions. The medium used consisted of a 1:1 mixture of medium 199 and Dulbecco's modified Eagle's (DME) medium which was conditioned before use. Insulin, transferrin, hydrocortisone, epidermal growth factor, and extract from bovine hypothalamus were used as supplement. Due to relatively low basal rates of mucin secretion from in vitro cultures, cultures are generally radiolabeled using 3H-glucosamine as a metabolic precursor. The radiolabeled mucinsreleased are quantitated by precipitation with TCA/PTA. Using this cell culture system, we investigated mucin release of goblet cells by altering the media bathing the apical surface of hamster tracheal surface epithelial(HTSE) cells. Acidic media added sulfuric acid caused sigcificant increases in mucin release (155 ± 20% at pH 4 and 146 ±16% at pH 5). Ammonium hydroxide also increased mucin release at pH  $9.0(156\pm17\%)$  and pH  $10(295\pm9\%)$  respectively. This additional mucin release seems to be associated with cell membrane damage as indicated by release of cellular LDH. SP stimulates secretion of mucin in cultured HTSE cells(154± 16% at  $1 \times 10^{-6}$ M and  $165 \pm 25\%$  at  $1 \times 10^{-5}$ M). PAF at  $5 \times 10^{-6}$ M and  $5 \times 10^{-5}$ M enhanced by HTSE cells in vitro 168 ± 34% and 259 ± 30% of mucin secretion, respectively. The increase in mucin release by PAF and SP was not secondary to cell damage or necrosis. SP and PAF may be in mediating mucous secretion induced by inflammation irritantion and infection.