Effect of experimental pancreatic growth on the content of xenobiotic-metabolising enzymes in the pancreas

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SUMMARY  The concentration in pancreatic tissue and the total pancreatic content of three xenobiotic metabolising enzymes has been determined in two models of experimental pancreatic growth namely, cholecystokinin-octapeptide injections and soy flour feeding. No significant change in pancreatic concentrations of benzo(a)pyrene hydroxylase or glucuronyl transferase was detected. In both models of pancreatic growth, however, the concentration of glutathione-S-transferase was significantly reduced. It is possible that the reduction in this enzyme may be of some importance in determining the susceptibility of the pancreas to carcinogenesis observed with long term soy flour feeding.

Injections of cholecystokinin (CCK) and its synthetic analogues and the feeding of raw soya flour or soy bean trypsin inhibitor are potent stimuli for pancreatic growth in several species.12 There is some evidence that the trophic response of the rat pancreas to soy flour feeding is, at least in part, mediated by the release of increased amounts of endogenous CCK produced by intraluminal inhibition of trypic activity in the intestine by the soy bean trypsin inhibitor.34 Soy flour feeding in conjunction with the feeding of azaserine, a weak pancreatic carcinogen, enhances the yield of malignant tumours in these animals as compared with the feeding of the carcinogen alone.6 It has also been observed, however, that prolonged feeding of raw soya flour to rats results in development of adenomatous nodules in the pancreas and ultimately pancreatic carcinomas.7

While the prolonged stimulus may, by itself, lead to the development of carcinomas, one possible mechanism may involve modification of the ability of the pancreas to handle the low background levels of carcinogenic xenobiotics present in the environment, most importantly, in the diet. Over a prolonged period of time, a small change in the capability of the pancreas to handle such compounds might be important in the development of malignant change in the gland. To study this, we have examined the ability of the control and hypertrophic pancreas of rats to carry out phase I (arylhydrocarbon hydroxylase) and phase II (glucuronyl transferase and glutathione-S-transferase) reactions.

Methods

ANIMALS

Adult male Sprague-Dawley rats, purchased from Canada Hybrid Farms, NS were maintained, four to a cage, at a minimum 40% relative humidity, approximately 23 °C, 12:12 photoperiod and provided with food and water ad libitum until killed.

TREATMENT

In one experiment two rats (200–300 g when first injected) in each cage received twice daily subcutaneous injections of cholecystokinin-octapeptide (CCK-OP) (Kinevac 300 ng/100 g/day) for 14 days while the other two were similarly injected with sterile 0.9% NaCl. Animals were fed Rat Chow® (Ralston Purina Co.).

In the other experiment half the rats (120–200 g) were maintained for 28 days in a pelleted feed (Ralston Purina Co, Richmond, Ind.) containing 40% raw defatted soya flour while the other half were provided with pellets identical except that the soya flour had been toasted to inactivate most of the trypsin inhibitor (TI) activity (reduced from 29.5 to 5 mg TI/g).
Radioactive determination by measuring enzyme assays.

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0-25 paper headed at 0-4 Teflon to 25% centrifugation was from sequential major blood vessels. The vessels by removed tissues were adhering fat, connective tissue and major blood vessels were removed. Excess fluid was removed by pressing the tissue between two layers of filter paper (Canlab F-2402) under a 200 g weight for 20 seconds. After weighing, the tissue was transferred to 25 ml of cold 50 mM Tris, pH 7.5, containing 0.25 M sucrose. All remaining steps were performed at 0–4 °C. The tissue was homogenised using a motor driven Teflon pestle for six strokes. The supernatant from sequential 650 g and 10000 g 10-min. centrifugation was centrifuged at 105000 g in an angle headed rotor for one hour. The 'microsomal' pellet was resuspended in 9 ml of buffer/sucrose by homogenising six strokes. Samples were frozen in liquid nitrogen and stored at −70 °C.

Cell fractionation was carried out using standard methods. The pancreas was quantitatively freed from the surrounding tissues and transferred to iced 0.9% saline where adhering fat, connective tissue and major blood vessels were removed. Excess fluid was removed by pressing the tissue between two layers of filter paper (Canlab F-2402) under a 200 g weight for 20 seconds. After weighing, the tissue was transferred to 25 ml of cold 50 mM Tris, pH 7.5, containing 0.25 M sucrose. All remaining steps were performed at 0–4 °C. The tissue was homogenised using a motor driven Teflon pestle for six strokes. The supernatant from sequential 650 g and 10000 g 10-min. centrifugation was centrifuged at 105000 g in an angle headed rotor for one hour. The 'microsomal' pellet was resuspended in 9 ml of buffer/sucrose by homogenising six strokes. Samples were frozen in liquid nitrogen and stored at −70 °C.

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**Enzyme Assays**

Microsomal mono-oxygenase (BPH) activity was determined by measuring the water soluble metabolites produced during 15 minutes incubation with radioactive benzo(a)pyrene (unmetabolised BP was extracted using ethyl acetate: acetone). Microsomal glucuronontransferase (GT) activity was measured using radioactive L-naphthol as substrate. Glutathione-S-transferase (GSH-T) activity in the 105000 g supernatant ('cystosol') was measured spectrophotometrically using L-chloro-2,4-dinitrobenzene as substrate. Protein concentrations were measured using the Biorad Protein Assay. Microsomal mono-oxygenase (BPH) GT and GSH-T activities were measured in 10 mM phosphate buffer, pH 7.4, containing 1 mM MgCl₂ at 37 °C.

Results are expressed as means ± standard error of the mean. A paired sample t-test was used to identify statistically significant differences. Significance was determined at p < 0.05.

**Results**

The pancreatic weights, enzyme concentrations and total pancreatic content of the enzymes are presented in Figs 1–7. Both CCK-OP and unheated soy flour feeding significantly increased pancreatic wet weight compared with appropriate controls. The pancreata in animals fed the toasted soy flour diet are slightly larger than those of animals fed regular chow. This probably reflects the fact that there is some residual trypsin inhibitor activity in the heat treated flour. With respect to the specific activity, the only significant alteration in the three enzymes studied was a fall in the concentration of glutathione-S-transferase. The decreases observed were 32% in the CCK-OP treated animals and 14% in those fed raw soya flour diets as compared with control values.
Experimental pancreatic growth and xenobiotic-metabolising enzymes in the pancreas

Discussion

The causes of pancreatic cancer in man are presently unknown. Epidemiological evidence points to environmental factors as of considerable importance. Environmental chemical carcinogens in food and tobacco smoke must be considered among these factors. Experimental models of cancer of the pancreas have been most successfully developed in rodents for example, the azaserine induced carcinoma in rats and the nitrosamine induced model in hamsters (see reviews by Longnecker).

Most chemical carcinogens are procarcinogens, being inert until activated in the tissues by metabolic processes, chiefly oxidations. Thus the importance of xenobiotic-metabolising enzymes in target tissues is evident. Of particular interest is the fact that the activity of these enzymes can be induced by a variety of substances and conditions including exposure of the tissues to the substrates of these enzymes - for example, the polycyclic aromatic hydrocarbons.

Two general types of metabolic conversion of xenobiotics occur in tissues, namely, phase I reactions which add reactive groups to the molecule, for example, the hydroxylations of benzo(a)pyrene, and phase II reactions which detoxify reactive metabolites by conjugation with endogenous compounds such as glutathione or glucuronic acid.
example, oxidations, hydroxylations or acetylations, and phase II reactions which involve the conjugation of phase I products to form glucuronides, sulphates and glutathione conjugates (thio-ethers). The latter subsequently undergo further modification as glutamic acid and glycine are cleaved from the glutathione residue and, ultimately, acetylation of the cysteine residue yields mercapturic acid derivatives. In general, phase II reactions are regarded as detoxifying processes by which reactive, potentially toxic products of phase I reactions are converted into water-soluble, readily excretable products.

Of all tissues, the liver contains the greatest amount of enzymes concerned with phase I and II reactions but significant quantities of these enzymes are found in extrahepatic tissues. The pancreas is known to have measurable amounts of aryl hydrocarbon hydroxylase, a phase I enzyme system and glucuronyltransferase, a phase II enzyme. The concentration of these enzymes is much lower than those of the liver. On the other hand, glutathione-S-transferase, another phase II enzyme, is present in relative abundance in the pancreas of rats, being present at approximately 50% of the concentration of the liver enzyme.

It has been shown that trophic influences on the pancreas, notably injections of CCK and soy flour feeding, act as cocarcinogens with chemical carcinogens such as N-nitrosobis(2-oxopropyl)amine and azaserine. Furthermore, prolonged soy flour feeding can ultimately yield not only pancreatic hypertrophy but also adenomas and carcinomas. It is possible that the action of low levels of carcinogens in the diet is enhanced by trophic factors such as soy flour feeding.

Our results show that with the pancreatic hypertrophy associated with either CCK-OP or soy flour, no detectable change in aryl hydrocarbon hydroxylase or glucuronyl transferase specific activity (units/mg protein) occurs, but that a significant fall in glutathione-S-transferase can be demonstrated in both forms of pancreatic hypertrophy. There is evidence that the hypertrophy of soy flour feeding represents the effect of endogenous hypercholesterokininaemia, thus these results may reflect the same phenomenon.

Although the explanation for the relative abundance of glutathione-S-transferase in the pancreas is not known it is possible that this enzyme represents a major detoxifying capacity of the pancreas. In a number of tissues, the concentrations of this enzyme, like other xenobiotic-metabolising enzymes, is increased by various agents present in the human diet, notably the butylated hydroxyanisoles. Agents inducing this enzyme might confer protection on the host against various xenobiotics. Conversely, conditions which depress the level of this enzyme may be of great significance in that they reduce the detoxifying capacity of a tissue. Our results, therefore, may offer one explanation for the carcinogenic activity of prolonged soy flour feeding in rats.

References

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