Regulation of gene expression in pancreatic adaptation to nutritional substrates or hormones

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Beginning with the work of J Pavlov, published in 1897, it has been known that levels of proteases and glycosidases in pancreatic juice were regulated in direct proportion to levels of protein and starch, respectively, in the diet. Pancreatic adaptation to nutritional substrates has been extensively studied at the level of tissue content and at the level of biosynthesis of specific enzymes. From a study of normal, diabetic, and insulin-treated rats, Desnuelle first suggested that insulin is involved in the adaptation of amylase biosynthesis in response to changes in carbohydrates in the diet. Despite numerous studies however, which have documented the predominant role of secretin and cholecystokinin (CCK) in stimulating fluid, bicarbonate and enzyme secretion in the exocrine pancreas, the role of these peptide hormones in the regulation of protein synthesis and pancreatic adaptation was, until recently, largely unappreciated.

Because of rapid dedifferentiation of pancreatic acinar cells placed in tissue culture, it has not been possible to study under in vitro conditions, adaptive changes on normal isolated cells. By use of continuous intravenous infusion of synthetic caerulein and secretin in optimal concentrations in the conscious rat we have been able to determine the limits of biochemical response to these hormones. After removal of the pancreas and incorporation of radio-labelled amino acids in the pancreatic lobules under in vitro conditions, separation of proteins by two dimensional IEF/SDS-gel electrophoresis has allowed a study of individual rates of synthesis for each of the exocrine products synthesised in the differentiated gland. An initial study varying the content of protein and carbohydrate in the diet between 0 and 82%, respectively, keeping lipid constant, indicated that amylase and the majority of protease zymogens were synthesised in direct proportions to the nutritional substrates (Fig. 1).

Caerulein infusion in the conscious rat resulted in both coordinate and anticoordinate regulation of protein synthesis. Beginning with a latent period of one to two hours synthesis of the majority of protease forms (anionic trypsinogen forms 1 and 2, anionic (C1) and cationic (C2) forms of chymotrypsinogen, anionic proelastase (PE1), and the group of pro-carboxypeptidase A and B (PCP A & B) was increased while the synthesis of amylase forms 1 and 2 was decreased (Fig. 2). Synthesis of lipase and cationic trypsinogen (T3) was not altered. Anticoordinate changes in protein synthesis were also observed with secretin stimulation but the period of latency was longer (two to three hours) and the pattern of response in the synthesis of some enzymes was different. In this case the synthesis of lipase and cationic proelastase (PE2) was increased and the synthesis of amylase forms 1 and 2 was decreased (Fig. 3).

Thus, in the rat it appears that synthesis of functional groups of enzymes (proteases, lipase, amylases) is regulated by specific hormones (CCK, secretin, insulin), respectively. In order to compare
the patterns of regulation of protein synthesis during hormonal stimulation with those observed during dietary adaptation, it was necessary to compare exogenous administration of hormones with conditions under which endogenous release of hormones is obtained. This can be achieved by either feeding a single dose of a new synthetic proteinase inhibitor (Camostate, FOY 305) which releases CCK by interfering with feedback regulation of active trypsin on CCK release in the duodenum. A second experimental approach involves prolonged fasting (72 hours), followed by refeeding, which also stimulates prolonged CCK release. Oral feeding of a single dose of 50 mg/kg FOY 305 resulted in a 10-fold increase in plasma CCK-levels which persisted for three to six hours. Total rate of protein synthesis increased during this time period by 80 to 90% above control levels and returned to normal by nine hours. Changes in individual rates of protein synthesis were observed beginning at nine hours after FOY administration. The changes were similar to the ones previously observed with caerulein infusion (increase in the majority of protease forms, no change in lipase, and a decrease in amylase).

After endogenous release of CCK, however, increase in the synthesis of anionic chymotrypsinogen (C1) was greater than that observed with anionic trypsinogen form 1 and 2. The opposite had been observed with caerulein infusion. It is noteworthy that a similar pattern of response as observed after FOY feeding has been found with the fasting-refeeding protocol.

The resolution obtained in the study of 16 exocrine products and in the pattern of their biosynthesis has allowed us to compare in detail changes observed during dietary adaptation with changes observed during hormonal stimulation. The pattern of response found during administration of a high protein diet (increase in C1 > T1, 2 > C2 > PE1 > PCPA + B, no change in L or T3, decrease in A1 = A2) was largely identical to that observed after endogenous release of CCK and similar to that observed during caerulein stimulation. Although less well studied, changes in
protein synthesis seen in the rat during administration of a high fat diet (increase in lipase and colipase synthesis) correlated reasonably well with changes seen during secretin infusion (personal communication). Thus, in the rat it appears that pancreatic adaptation to changes in nutritional substrates in the diet is regulated through specific hormones. This, however, might not be true for other species and further work is needed to investigate the response of protein synthesis to specific hormones in other species.

Elucidation of the molecular mechanisms which regulate anticoordinate changes in protein synthesis in the exocrine pancreas in response to hormone stimulation is of fundamental importance in the understanding of dietary adaptation. From the work of a number of laboratories, the basic steps in gene expression are now understood. Transcription of a specific gene by RNA polymerase II results in the formation of a primary RNA transcript, which together with a wide variety of nuclear RNA transcripts constitutes heterogenous RNA, so called because of the variation observed in both nucleotide length and sequence specificity. Within the nucleus, the primary RNA transcript is modified or processed largely through (a) the removal of intron sequences by splicing mechanisms and (b) the addition of a poly (A) tail to the 3’ terminus. Processed mRNA is transported from the nucleus into the cytoplasm where, after functional binding of 40S and 60S ribosomal subunits, its coding sequence is translated into protein. Regulation of gene expression may therefore occur at different levels including (1) transcription, (2) RNA processing, (3) mRNA transport (4) efficiency of mRNA translation, or (5) mRNA sequestration or degradation.

Although the information contained in the genetic code flows from DNA to RNA to protein, the events which regulate gene expression involve the interaction of regulatory proteins with specific nucleotide sequences on DNA. Thus synthesis of heterogenous RNA from limited regions of DNA is catalysed by the enzyme RNA polymerase II. This multimeric enzyme initiates the transcription process after binding to a specific region of DNA termed the promoter or TATA sequence, which usually resides 30 nucleotides upstream from the nucleotide at which transcription begins, the mRNA start site. After unwinding the DNA helix at the promoter site, RNA polymerase moves along the DNA and polymerises a single stranded mRNA sequence through the sequential addition of individual ribonucleotides. The efficiency of gene transcription initiated at the promoter site, however, is regulated by enhancer sequences, which usually reside further upstream in the 5’ flanking region of genes. Enhancer elements occur in two types, one of which regulates the expression of genes in specific tissues and the other which modulates gene expression in response to hormones.

An understanding of how hormones regulate gene expression is largely confined to the action of steroid hormones. Studies from several laboratories have indicated that steroid hormone receptor proteins are DNA binding proteins. Subdomains of the steroid receptor include a hormone binding site and a DNA binding site which interacts with specific nucleotide sequences in the enhancer regions of several genes whose expression is regulated by steroid hormones. The available evidence indicates that steroid hormones regulate gene expression at the level of transcription. In this case the changes observed in protein synthesis correlated with significant changes in mRNA levels and these in turn were correlated with changes in transcription rates of specific genes as determined by the use of complementary DNA probes.

In contrast little is known regarding the molecular mechanisms involved in gene expression by peptide hormones. In this area we have focused our research in the past years on studies relating to CCK and secretin stimulation in the rat exocrine pancreas. A consecutive study analysing functional mRNA concentrations in an in vitro translation system demonstrated that pronounced changes in individual enzyme biosynthesis were not accompanied by changes in levels of mRNA at least during the initial six hours of caerulein stimulation. Consequently, we have suggested that during the early periods of caerulein stimulation, the anticoordinate changes observed in protein synthesis occur at the level of efficiency of mRNA translation. With continuation of caerulein stimulation for 12 or 24 hours, minor changes were observed in mRNA levels which paralleled those observed in protein synthesis. Further research is needed to determine whether changes in mRNA levels during these late periods or after chronic hormone administration for several days are mediated by changes in transcription rates, mRNA turnover rates or both. Thus, with caerulein stimulation, there appears to be at least two distinct and overlapping phases in the regulation of gene expression. It is impressive that the anticoordinate changes observed in protein synthesis in both phases are similar and that regulations which occur in the second period serve to augment those occurring in the initial period. Sequential and overlapping regulation in this manner represents, at the biochemical level, increased commitment to adaptive changes.

The observation of a regulation of protein synthesis in the absence of changes in mRNA levels by peptide hormones poses the question about the regulatory mechanisms. Because of the relatively long half lives of mRNAs for exocrine pancreatic products, four to
six hours, it is reasonable to expect that the early changes observed in protein synthesis, minutes to hours, are mediated at the translational level. Little is known at present, however, regarding the molecular mechanism by which changes in efficiency of mRNA translation may occur in the cell. Presumably such changes are mediated at the level of efficiency in the initiation of mRNA translation, which represents the rate limiting step in protein synthesis. Changes at this level may involve nucleotide signals in the 5'-nontranslated regions of mRNAs and the interaction of these signals or sequences with either regulatory proteins or regulatory RNA molecules.

The majority of mRNAs coding for pancreatic enzymes in the rat and dog have now been sequenced\(^6\) and it is therefore possible to analyse mechanisms which may be responsible for modulation of initiation of mRNA translation. A comparison of the 5'-non-translated sequences of anionic and cationic trypsinogen mRNAs has suggested a mechanism by which the synthesis of the anionic but not the cationic isoenzyme is increased during caerulein stimulation.\(^22\) The 5'-non-translated regions of dog and rat anionic trypsinogen mRNAs are short (14 and 12 nucleotides, respectively) and show a region of conserved sequence involving nine contiguous and identical residues (Fig. 4). In contrast, this conserved sequence was not observed among 29 nucleotides in the 5'-non-translated region of cationic trypsinogen mRNA. The conserved element in the two anionic trypsinogen mRNAs also showed potential base pairing to a region at the 3' end of 18S ribosomal RNA which exists in a stem-loop structure (Fig. 4). This region of the 18S rRNA is known to occur near the site of initiation of mRNA translation. The initiation process in eukaryotic cells involves at
least two interactions between mRNA and the small (40S) ribosomal subunit: (a) binding of a protein complex associated with the 5' cap of mRNA to the ribosomal subunit and (b) binding of the AUG initiation codon to the UAC anticodon of the initiation methionine tRNA, which is associated with activated 40S subunits. We have proposed that base pair interactions between anionic trypsinogen mRNA and 18S rRNA represent, under certain circumstances, a third interaction between mRNA and the 40S ribosomal subunit. Because the stem loop structure in the 18S rRNA exists in the closed configuration under basal conditions as judged by analysis of isolated 40S subunits, we have hypothesized that during hormonal stimulation, the stem loop is disrupted. In the open configuration, potential base pairing may occur between anionic trypsinogen mRNA and 18S rRNA, which may serve to promote mRNA alignment and facilitate the initiation of mRNA translation (Fig. 5). In contrast, alternative configurations in this region of 18S rRNA will have little effect on the binding of cationic trypsinogen mRNA. Under conditions in which the stem loop is open, the efficiency of translation of anionic but not cationic trypsinogen mRNA may increase, resulting in increased synthesis of anionic but not cationic trypsinogen.

Fig. 5  Hypothetical role of stem loop structure at the 3' terminus of 18S rRNA in the regulation of protein synthesis. For details see text.

References


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