PORTAL HEPATOTROPHIC FACTORS, DIABETES MELLITUS AND ACUTE LIVER ATROPHY, HYPERTROPHY AND REGENERATION


DURING THE LAST DECADE, evidence has been accumulated in our laboratories (24, 25, 37-40) that substances termed hepatotrophic factors in portal venous blood can specifically influence liver function as well as the size, chemical composition and dividing capability of the hepatocytes. In this report, the role of these portal hepatotrophic factors will be examined in the acute regeneration process that follows partial hepectomy.

METHODS
Animal Groups

Seventy mongrel dogs, weighing 8 to 29 kilograms, contributed to the final data. An approximately equal number of experiments failed because of thrombosis of the vascular reconstructions, premature death of the dog from a variety of other complications or because of failure to produce alloxan diabetes.

The various operations, biopsies and sacrifice procedures were performed under anesthesia with pentobarbital sodium supplemented with phencyclidine hydrochloride (Sernylan®) and succinylcholine chloride (Anectine Chloride®).

Group 1. Eleven normal dogs were anesthetized, group 1A. After obtaining several grams of tissue from one of the right and one of the left liver lobes for the various analyses, the dogs were sacrificed (Fig. 1a). These same dogs were the normal controls in another recent publication (40).

Six additional normal dogs of group 1B were submitted to a sham operation that was designed to simulate the procedure in group 2. The portal vein and its bifurcation were dissected free as well as the more inferior junction of the superior mesenteric and splenic veins. The superior mesenteric vein was occluded for 20 to 30 minutes after which the tip of the inferior lobe of the pancreas was resected as shown in Figure 1b. Four days later, the dogs were sacrificed, and tissue was obtained for both the histopathologic and the autoradiographic studies.

Group 2. Five dogs had a procedure called splanchic division, which diverts the nutrient rich intestinal venous blood into the left liver lobes through a reversed external jugular vein graft (37) and which provides perfusion of the right liver lobes by the hormone rich pancreaticogastroduodenal venous blood (Fig. 1b). The tail of the inferior lobe of the pancreas was resected (Fig. 1b) to prevent the experimental artifact that would have been caused by drainage of this small piece of
pancreatic tissue into the intestinal vein and, from there, into the left liver lobes.

**Group 3.** Stable diabetes mellitus was induced with a single injection of 70 to 80 milligrams of alloxan—mesoxalyl urea—per kilogram of body weight. The diabetes mellitus was managed by daily subcutaneous injections of 10 to 15 units of neutral protamine Hagedorn insulin. Three to eight weeks later, two dogs had the same splanchnic division procedure as those in group 2 (Fig. 1c). Insulin therapy was continued in the three and five day periods of postoperative observation.

**Group 4.** Six dogs had total pancreatectomy with preservation of the duodenum and common duct at the same operation as splanchnic venous division (Fig. 1d). Three of the dogs were given insulin subcutaneously for the three to four day period of postoperative study, group 4A. The diabetes mellitus of the remaining three dogs was not treated, group 4B.

**Group 5.** Four normal dogs had complete removal of the most right and the most left liver lobes (Fig. 2a). These two lobes make up an estimated 30 per cent of the total liver weight. The dogs were sacrificed four days later.

**Group 6.** Four dogs had resection of the two most leftward liver lobes, the most right liver lobe and the tip of a small lobe located along the left side of the vena cava. The estimated resection was 60 per cent. The dogs were sacrificed three days later.

**Group 7.** Six dogs had the same splanchnic division procedure as those in group 2. In addition, a 30 per cent hepatectomy was performed (Fig. 2b). The dogs were sacrificed four to five days later.

**Group 8.** Five dogs had the same splanchnic division procedure as those in group 2, plus a 60 per cent hepatectomy. The dogs were sacrificed after two or three days.

**Group 9.** Four dogs had total pancreatectomy on the same day as a 30 per cent hepatectomy and splanchnic division (Fig. 2c). During the four days between operation and sacrifice, two dogs were treated subcutaneously with insulin, group 9A. Two dogs were not treated, group 9B.

**Group 10.** Five dogs had partial portacaval transposition (Fig. 3a). The left portal branch was detached and anastomosed end-to-end to the supra-renal inferior vena cava. Thus, the left lobes were perfused with systemic venous blood, including the renal and adrenal effluent, whereas the right lobes were exposed to the total nonhepatic splanchnic venous return. The dogs were sacrificed five days after operation, except for one dog that was observed for seven days.

**Group 11.** Three dogs had simultaneous partial portacaval transposition and total pancreatectomy (Fig. 3b). Postoperatively, they were not treated with insulin during the four or five days of the experiment.

**Group 12.** Five dogs underwent partial portacaval transposition. From 89 to 95 days later, reoperation was performed. The expected hypertrophy of the right lobes and the atrophy of the left lobes were found. The most far right and the most far left lobes were removed per cent resect were sacrificed.

**Group 13.** Five dogs had total pancreatectomy after partial portacaval transposition of the pancreas and without sham operation. The expected hypertrophy of the right lobes was found. The most far right and the most far left lobes were resected without sham operation.

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**Fig. 1.** A series of control procedures not involving partial hepatectomy. The dogs either were normal or had been submitted to acute splanchnic division. a, Group 1—normal dogs, with or without sham operation. b, Group 2—dogs in which the left liver lobes were perfused with intestinal blood and in which the right liver lobes received pancreatiegastrooduodenosplenic venous blood. Explanation of partial removal of inferior lobe of pancreas is given in text. c, Group 3—same operation as in b, but in dogs with alloxan-induced diabetes. d, Group 4—same operation as in b, but with total pancreatectomy. The duodenum and common duct were left intact.
Formed (Fig. 2b). Five days later, hepatectomy or acute splanchic division was accomplished in groups 5 and 6, plus a 60 per cent hepatectomy in group 9. But, groups 7 and 8—acute splanchic division was carried out plus 30 per cent hepatectomy in group 7 and 60 per cent hepatectomy in group 8, plus total pancreatotomy and 30 per cent hepatectomy on the same day.

Lobes were removed (Fig. 4a), for an estimated 30 per cent resection. After four more days, the dogs were sacrificed.

Group 13. Four dogs had the same 30 per cent hepatectomy as those in group 12, 89 to 95 days after partial portacaval transposition. In addition, the pancreas was totally removed (Fig. 4b). Subcutaneous insulin was given for the next four days until sacrifice.

Deoxyribonucleic Acid Synthesis

About two hours before sacrifice, the 70 dogs were given \((\text{CH}_3^3\text{H})\) thymidine by the intravenous route. The specific activity was 6.4 curies per millimole. Because doses of 1.5 or 4.5 millicuries of \((\text{CH}_3^3\text{H})\) thymidine were used at different phases of the investigation, but without consideration of body weight, the dose spectrum was broad, ranging from 0.05 to 0.41 millicurie per kilogram of body weight.

At the time of sacrifice, liver biopsies were quickly frozen and kept at minus 20 degrees C. until the analyses were completed. Extraction and purification of deoxyribonucleic acid were carried out according to the method of Schneider and Greco (32) with minor modifications as described elsewhere (40). In essence, the method involves first removing the ribonucleic acid and acid soluble fractions with sodium hydroxide and cold perchloric acid, respectively. Deoxyribonucleic acid is then extracted from the residue, including protein and lipid, with hot perchloric acid. The deoxyribonucleic acid content was measured by the diphenylamine method of Burton (5), using calf thymus deoxyribonucleic acid as the standard.

The in Vivo incorporation of \((\text{CH}_3^3\text{H})\) thymidine into deoxyribonucleic acid was estimated by measuring the specific activity of the deoxyribonucleic acid obtained by extraction. The details of the counting were described in another publication.
RESULTS

Normal Dogs

Deoxyribonucleic acid synthesis. Thymidine uptake was equal in the right and left lobes of the livers of the 11 normal dogs of group 1A. The average specific activity was not much different at the doses of 1.5 and 4.5 millicuries (Fig. 5). A 30 per cent hepatic resection resulted in a slight bilateral increase in the specific activity of the residual hepatic tissue of the dogs of group 5 that were administered 1.5 millicuries of tritiated thymidine. After a 60 per cent hepatectomy, the dogs of group 6 that were given 4.5 millicuries of tritiated thymidine had a sixfold increase in specific activity that was equal on both sides (Fig. 5).

Histopathology and autoradiography. There were no significant differences in the sizes of either the lobules or the hepatocytes in the right and left liver lobes of the 11 normal dogs in group 1A. All of these livers lacked fat and contained normal amounts of glycogen and appeared normal ultrastructurally. The other two livers were deficient in glycogen. Autoradiographs of the livers of the dogs in group 1A showed similar low numbers of labeled hepatocytes in the right and left lobes (Table I). The findings were the same in the dogs of group 1B after the sham operation, as shown in Table I.

A 30 per cent hepatic resection in the dogs of group 5 resulted, after four days, in a slight, but nonsignificant, enlargement of the hepatocytes in all lobes (Table I) and in an accumulation of vacuoles of various sizes in the liver cell cytoplasm. In one dog, large fat globules were present in the centrilobular hepatocytes. The amount of stainable glycogen was reduced. The mitotic index was normal, but autoradiographs (Table I) showed that the number of labeled hepatocytes was slightly and significantly raised, p < 0.001. Ultrastructurally, the hepatocytes contained more autophagosomes and less glycogen than normal, and the cisternae of the rough endoplasmic reticulum were often dilated.

Three days after 60 per cent hepatectomy in group 6, the hepatocytes were enlarged in both sides of the liver, and their cytoplasm contained many vacuoles and some fat. The amount of stainable glycogen was reduced. Both the mitotic index and the number of labeled hepatocytes (Table I) were markedly raised. Electron microscopic examination of the hepatocytes confirmed the decrease in the amount of cytoplasmic glycogen and showed...
Splanchnic Division

Deoxyribonucleic acid synthesis. The absolute specific activities in the dog livers submitted to splanchnic division were so variable from experiment to experiment that data analysis was limited to comparisons between the right and left lobes, using an analytic technique whereby the side with the greater specific activity was assigned a value of 100 per cent and a proportionally lower percentage calculated for the other side of the liver. The numbers in parentheses correspond to the dog group numbers in parentheses from the normal dogs of group 1 and the dogs of groups 5 and 6 after hepatic resection. Statistical comparisons between the right and left lobes in each group are indicated by p values or by NS, not significant. For comparison, the results are given in the top row from the normal dogs of group 1 and the dogs of groups 5 and 6 after hepatic resection. Note that the advantage in thymidine uptake enjoyed by the right lobes which received pancreatico-gastro-duodenosplenic blood after splanchnic division in groups 2, 7 and 8 was abolished in groups 3 and 4 or even transferred to the left lobes in group 9 by the creation of diabetes, either with alloxan or by pancreatectomy.

Histopathology and autoradiography. Five days after splanchnic division in the dogs of group 2, the hepatocytes in the left liver lobes, which had retained their normal morphology and increased numbers of lipid droplets, showed a slight, but hepatocytes in the right liver lobes, which had lost their normal morphology and decreased numbers of lipid droplets, showed a more pronounced morphological change. The predominant deoxyribonucleic acid synthesis in the dogs of group 2 was in the right lobes which were perfused with pancreatico-gastro-duodenosplenic blood, although the lobular differences were not significant. However, this effect was accentuated in groups 7 and 8 by partial pancreatectomy and reached statistical significance. The dominance of the right lobes in deoxyribonucleic acid synthesis was eliminated by alloxan-induced diabetes, group 3, or after pancreatectomy, group 4. The most clearcut results were in the dogs of group 9 submitted to partial hepatectomy plus total pancreatectomy (Fig. 6). The predominant deoxyribonucleic acid synthesis was transferred to the left lobes to a statistically significant degree, p<0.01.

Histopathology and autoradiography. Five days after splanchnic division in the dogs of group 2, the hepatocytes in the left liver lobes, which had re-
ceived the nutrient rich intestinal venous blood, had undergone a number of changes (Fig. 7). They had decreased in size (Table I), were irregular in shape and were depleted of glycogen. The lobules were shrunken, and the Kupffer cells were increased in number and size and contained hemo-
siderin. There were no fat globules visible by light microscopy. Ultrastructurally, the rough endoplasmic reticulum was reduced in amount and disor-
ganized, and some of the cisternae were dilated.

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**TABLE I.—COMPARISONS OF HEPATOCYTE SIZE AND CELL DIVISION BY AUTORADIOGRAPHY IN SPLANCHNIC DIVISION EXPERIMENTS**

<table>
<thead>
<tr>
<th>Group</th>
<th>Description</th>
<th>No. of labeled hepatocytes per 1,000 hepatocytes</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A</td>
<td>Normal dogs</td>
<td></td>
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<tr>
<td>1B</td>
<td>Normal sham</td>
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</tr>
<tr>
<td>5</td>
<td>30 per cent resection</td>
<td></td>
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<tr>
<td>6</td>
<td>60 per cent resection</td>
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</tr>
<tr>
<td>2</td>
<td>Splanchnic division</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>Splanchnic division + 30 per cent</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>Splanchnic division + 60 per cent</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>Splanchnic division + diabetes</td>
<td></td>
</tr>
<tr>
<td>4A</td>
<td>Splanchnic division + pancreatectomy</td>
<td></td>
</tr>
<tr>
<td>4B</td>
<td>Splanchnic division + pancreatectomy, no insulin</td>
<td></td>
</tr>
<tr>
<td>4A and B</td>
<td>Splanchnic division + pancreatectomy</td>
<td></td>
</tr>
<tr>
<td>9A</td>
<td>Splanchnic division + pancreatectomy, 30 per cent resection</td>
<td></td>
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<tr>
<td>9B</td>
<td>Splanchnic division + pancreatectomy, 30 per cent resection, no insulin</td>
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</tr>
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Mean ± standard deviation.
NS, Not significant.

There was also a frequent than no vacuoles was incr
There was also proliferation of the smooth endoplasmic reticulum, glycogen granules were less frequent than normal and the number of small fat vacuoles was increased. The right liver lobes, which had been perfused by the hormone rich pancreaticogastroduodenosplenic blood, had larger lobules and hepatocytes than were present in the normal dogs of group 1. The enlarged hepatocytes con-
FIG. 8. Changes in a liver three days after a splanchnic division and total pancreatectomy experiment of group 4B as shown by light microscopy, electron microscopy and autoradiography. The hepatocytes on both sides of the liver are atrophic and contain excess cytoplasmic fat. The amount of glycogen and rough endoplasmic reticulum is reduced. The rate of cell division, as indicated by autoradiography, is bilaterally normal. Upper panels, hematoxylin and eosin, X120; middle panels, electron micrography, X1,700; and lower panels, autoradiography, X300.

Tained normal quantities of glycogen and were free of fat. Binucleate liver cells and proliferating bile ductules were present, and the mitotic index was raised. Ultrastructurally, the enlarged liver cells were essentially normal. Autoradiography showed about three times as many labeled hepatocytes in the enlarged right lobes as in the smaller left lobes (Table I).
The addition of a 60 per cent partial hepatectomy to splanchnic division in group 8 resulted in the kind of histologic appearance shown in Figure 8. There was marked enlargement of the lobules and hepatocytes on both sides of the liver (Table I). The left lobe hepatocytes, instead of being atrophic, were as large as the liver cells on that side after 60 per cent resection alone. Right lobe hepatocytes...
Partial Portalaval Transposition

of the dogs of group four or five days after partial hepatectomy, group 12, was again significant.

Histopathology, days after partial hepatectomy, group 10, which had increased in size (combined with the predominant lobes, the left lobes, left did not achieve

Partial portacaval was not associated with any change in the size of the liver, atrophy was most obvious in the three dogs of group 4B which were not treated with insulin; these dogs showed increased amounts of fat in the cytoplasm of the hepatocytes. The livers of the other dogs in groups 3 and 4 were free of fat visible by light microscopy.

Autoradiography showed more labeled hepatocytes on both sides of the liver than after partial hepatectomy (Table I). The numbers of radioactive nuclei on the left sides of the livers were roughly the same as after the 60 per cent resection alone. The numbers on the right side were higher than after either 60 per cent resection or splanchic venous splanchnic division alone. Ultrastructurally, the hepatocytes on both sides contained fewer glycogen granules than normal and increased numbers of autophagosomes and lipid droplets. The amount of rough endoplasmic reticulum was normal, but the cisternae were a little dilated.

Similar, but much milder, changes were produced by the combination of 30 per cent partial hepatectomy and splanchic division, which was used for group 7 (Table I).

The difference in size of the hepatocytes in the left versus the right lobes, brought about by splanchic division, did not occur in the seven diabetic dogs of groups 3 and 4 (Table I and Fig. 9). After two to five days, the hepatocytes in the right and left lobes were almost equal in size and smaller than those of normal dogs. The hepatocyte area was most obvious in the three dogs of group 4B which were not treated with insulin; these dogs showed abundant fat in the cytoplasm of the hepatocytes. The livers of the other dogs in groups 3 and 4 were free of fat visible by light microscopy.

Ultrastructurally, the hepatocytes on both sides of these livers in diabetic groups 3 and 4 showed varying degrees of disruption and loss of rough endoplasmic reticulum, an increase in the amount of smooth endoplasmic reticulum, a loss of glycogen particles and increased numbers of lipid droplets. Autoradiography showed normal numbers of labeled hepatocytes in both sides of the liver of the three diabetic dogs of group 4B that were not given insulin. In the dogs treated with insulin, there were increased numbers of radioactive liver cells in the left lobes.

The addition of the 30 per cent hepatectomy in the diabetic dogs of group 9 was associated with slightly larger hepatocytes on both sides of the liver. The hepatocyte size was similar to that seen after liver resection alone in group 5. The numbers of radioactive cells were raised on both sides of the liver of the three diabetic dogs of group 4B that were not given insulin. In the dogs treated with insulin, there were increased numbers of radioactive liver cells in the left lobes.

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Partial Portacaval Transposition

Deoxyribonucleic acid synthesis. The right liver lobes of the dogs of group 10 which were perfused for four or five days after operation with the total splanchnic venous return had significantly greater deoxyribonucleic acid synthesis than the left lobes which were receiving the venous return from the kidneys and hindquarters (Fig. 10). This right lobar dominance was not changed by the absence of the pancreas in the dogs of group 11 (Fig. 10).

Three months following partial portacaval transposition and four days after 30 per cent hepatectomy, group 12, deoxyribonucleic acid synthesis was again significantly greater in the right liver lobes. The addition of pancreatectomy at the time of 30 per cent hepatectomy in group 13 switched the predominant deoxyribonucleic acid synthesis to the left lobes, although the lateralization to the left did not achieve significance (Fig. 10).

Histopathology and autoradiography. Four to five days after partial portacaval transposition in the dogs of group 10, the hepatocytes in the left liver lobes, which had received systemic blood, had decreased in size (Table II) and had become depleted of glycogen. The lobules were smaller than normal and the Kupffer cells were increased in size and number and contained hemosiderin. Ultrastructurally, there was loss of rough endoplasmic reticulum, glycogen granules were scarce and there were increased numbers of small lipid droplets. The right liver lobes, which had received splanchnic venous blood, had larger lobules and hepatocytes than were present in the normal dogs of group 1. Ultrastructurally, the enlarged cells were essentially normal. Autoradiographs showed more labeled hepatocytes in the right lobes than in the left lobes (Table II).

The presence of untreated diabetes in the dogs of group 11 changed some of these findings. The enlargement of the hepatocytes in the right lobes failed to occur, and the number of radioactive thymidine-labeled hepatocytes in these lobes was lower than after uncomplicated portacaval transposition, although still greater than on the left side (Table II).

Three months after partial portacaval transposition and four days following 30 per cent hepa­
tectomy, group 12, the hepatocytes of the right lobes were larger than the left, and the incidence of labeled hepatocytes was also greater on that side. The ultrastructure of the liver cells in the right and left lobes did not differ significantly from that seen after partial portacaval transposition alone (40). The addition of diabetes treated with insulin, group 13, was associated with enlargement of the previously atrophic hepatocytes in the left lobes to a size greater than normal and similar to those in the right lobes. The enlarged hepatocytes contained many lipid droplets. Autoradiography showed more labeled hepatocytes in the left lobes than in the right lobes. Ultrastructurally, the rough endoplasmic reticulum was dilated in the hepatocytes of both lobes and reduced in amount in the cells of the left lobes. There were fewer than normal glycogen granules in all lobes.

DISCUSSION

In a classic article published 55 years ago, Rous and Larimore (31) raised the possibility that substances in portal venous blood prevented atrophy and promoted regeneration. In discussing the atrophic effects of portal branch ligation upon the regions of rabbit liver thus deprived of splanchnic blood and the concomitant hypertrophy found in the rest of the liver, they suggested "... that the liver has no essential activity—none on which its maintenance depends—that it (sic) is not intimately connected with substances derived from organs drained by the portal system."

Rous and Larimore (31) did not have evidence in direct support of their suggestion, nor was substantiation provided by subsequent workers including Mann (23) during the next 40 years.
Instead, articles by Child (7) and Fisher (10, 11, 12) and their associates led to the almost universal acceptance of the alternative view that the portal component of total hepatic blood flow was important primarily in proportion to its quantity and that the quality of the blood was not significant.

During the last decade, the pendulum has moved back toward the position that portal venous blood has special properties which profoundly affect hepatocellular structure and function. The revival of the qualitative concept came from our investigations with revascularization of auxiliary liver grafts (24, 25, 39). In these experiments, coexisting whole livers or two portions of the same liver were given different kinds of portal venous inflow. The hepatic tissue perfused with venous return from the splanchnic organs underwent hypertrophy, hyperplasia and glycogenation in comparison with the hepatic tissue deprived of this advantage. The confirmation by other workers of these observations has been summarized elsewhere (37, 38, 40).

Results of recent studies have indicated the nature of the hepatotrophic substances. The essence of their influence is the interreaction of hormones generated by splanchnic organs and delivered straight to the liver and with a presumably augmented significance because of the concentrated flux of nutritional substrate in the same venous blood (37, 38, 40). Both by inference and by direct testing, insulin was concluded to be the single most important hepatotrophic factor, although by no means the only one (37, 38, 40). This hormonal and biochemical concept has helped to explain earlier observations by Marchioro (25), Pouyet (27), and Ranson (29) and their associates that were made in experiments in which splanchnic venous flow was divided between two livers or two liver fragments.

Although its probable relation to liver regeneration has been obvious, the hepatotrophic concept as it has evolved in our laboratories has been based on chronic experiments not involving partial hepatectomy. The acute studies herein reported were undertaken in diabetic and nondiabetic dogs, periments not involving diabetes or hepatectomy, as it has evolved in our laboratories has been based A special note should be made

The clearing of insulin by the liver has been particularly well studied, as was recently reviewed by Field (9). An unmasking of the role of portal blood constituents is thereby made feasible to an extent not possible if portal blood is bypassed around a single liver but eventually returned to it in diluted form as with Eck's fistula or the portacaval transposition of Child.

Nondiabetic dogs submitted to splanchnic division or partial portacaval transposition showed the remarkable rapidity with which liver tissue was altered after it was denied exposure to either the total splanchnic venous return or its pancreatico-gastroduodenal component. Within four or five days, the hepatocyte atrophy was almost as pronounced as we have reported (24, 37, 40) in the same kinds of experiments after two months. Deglycogenation, depletion or distortion of the rough endoplasmic reticulum, fatty vacuolization and other structural changes also were advanced within four or five days. The ultrastructural abnormalities in these shrunken hepatocytes were similar to those reported by Reaven and his colleagues (30), rat livers within three days after the production of the insulinopenia of alloxan diabetes. At the same time, hypertrophy of the healthy contralateral liver cells had occurred.

These differential effects upon the right and left liver lobes were drastically altered by diabetes mellitus in the splanchnic division experiments. A similar modification of results was caused by the addition of pancreactectomy to partial portacaval transposition. The crucial observation was that the lobes previously protected either by pancreatico-gastroduodenal or by total splanchnic flow now partied, or completely, lost this advantage. The profoundly influential effects of iatrogenic diabetes mellitus were similar to those reported by us previously over the much longer period of two months (38, 40).

A special note should be made about the effect of the blood flow deviations upon over-all hepatocyte replication. In all double liver fragment experiments not involving diabetes or hepatectomy, the deprivation of one portion of the liver of qualitatively optimal blood set into motion a low grade proliferative response which could be consistently detected by autoradiography, which did not occur in the sham operated upon controls and which was shown in a previous report (40) to persist through the first two postoperative months. Although this heightened mitotic activity was predominantly in the hypertrophic liver lobes perfused by the total splanchnic return or its pancreaticoduodenal portion, cell division in the most flow thrc to the two liver since such measure to say that it is liver tissue for the flow with any great, if the div livered to anot This was dem Marchioro and portal vein brat these b: Atrophy of the advented, even t
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sequences of the removal or addition of insulin or other hormones have also been shown by Leffert (18, 19) in hepatocyte monolayer cultures in which flow factors did not exist at all. Earlier, the role of insulin in supporting the growth and replication of other kinds of cells had been well established by Lieberman and Ove (20).

By using cell division as an end point, it was interesting to compare the results in the dogs of group 3 which were submitted to splanchnic division plus alloxan-induced diabetes with those obtained in the dogs of group 4 which had splanchnic division and total pancreatectomy. As in chronic animals reported earlier (38, 40), there was not a potent or even identifiable further effect of the total loss of the pancreas compared with the consequences caused by the simpler hormone defect of alloxan-induced diabetes. This meant that glucagon and other substances from the pancreas singly or together did not have a hepatotrophic role

PORTAL HEPATOTROPHIC FACTORS, DIABETES AND LIVER ATROPHY

Supernormal portal flow in rapidly atrophying auxiliary livers or liver fragments that were revascularized by portacaval transposition has also been reported by Marchioro (24) and Daloze (8) and their associates. Earlier work by us (42) and by Heer and his co-authors (15) had already shown that the total blood flow of whole livers revascularized by portacaval transposition was essentially normal. Finally, Pouyet and his colleagues (27) have calculated in their version of the splanchnic division procedure that the intestinal component of splanchnic venous return was greater than the fraction from the pancreaticoduodenopancreatic area, even though the latter has a much greater hepatotrophic effect.

Such tedious discussions and counterarguments about potential artifacts introduced by blood flow disparities in double liver fragment models no longer serve a useful purpose, since they have been rendered moot by our acute experiments herein described as well as by our reported chronic experiments (40). This work showed that differential hepatotrophic effects found in the two coexisting liver fragments were canceled or remarkably diminished by diabetes mellitus. In both the long and short term experiments, the results with stable alloxan-induced diabetes were particularly enlightening, since alloxan is not known or suspected to change splanchnic flow. Thus, the only conceivable explanation for the results was that a fundamental change in the relative chemical environment of the two liver sides had been wrought by removal of endogenous insulin. Dramatic consequences of the removal or addition of insulin or other hormones have also been shown by Leffert (18, 19) in hepatocyte monolayer cultures in which flow factors did not exist at all. Earlier, the role of insulin in supporting the growth and replication of other kinds of cells had been well established by Lieberman and Ove (20).

By using cell division as an end point, it was interesting to compare the results in the dogs of group 3 which were submitted to splanchnic division plus alloxan-induced diabetes with those obtained in the dogs of group 4 which had splanchnic division and total pancreatectomy. As in chronic animals reported earlier (38, 40), there was not a potent or even identifiable further effect of the total loss of the pancreas compared with the consequences caused by the simpler hormone defect of alloxan-induced diabetes. This meant that glucagon and other substances from the pancreas singly or together did not have a hepatotrophic role

Several times normal, as Zuidema and his associates have reported (47).
approaching in importance that of insulin. These observations have greatly weakened our earlier speculation (37) which was subsequently supported by Bucher and Swaffield (4) that the insulin to glucagon ratio might be more important than insulin alone. Our more recent findings are also a frontal challenge to the contention by Price (28) and Whittemore (45) and their colleagues that glucagon is the principal hepatotrophic factor.

Our emphasis on insulin as the single most important hepatotrophic influence and, for that matter, the only easily identifiable one in our test system, does not imply that it is the sole factor. In a number of recent publications (37, 38, 40, 41), information was reviewed favoring multiple portal and extraportal factors. These presumably include other hormones than insulin and probably also nutrients.

The acute experiments herein reported have further strengthened the multifactorial hypothesis as it applies to hepatocyte mitosis. In the splanchic division experiments of groups 3 and 4, the creation of diabetes eliminated the dominance of cell division in the previously favored right lobes and transferred this to the left, providing there was treatment with subcutaneous exogenous insulin which was eventually distributed to both fragments. However, if the replacement insulin was omitted as in group 4B, the rate of cell division became bilaterally equal. Confirmatory observations were made in the dogs of groups 9 and 13, which were submitted to total pancreatectomy plus 30 per cent hepatic resections. In all these dogs with diabetes of one kind or another, the combination of exogenous insulin plus intestinal blood apparently supported cell renewal in liver tissue to a greater extent than exogenous insulin plus gastroduodenosplenic or inferior vena caval blood. A subtle, but significant, hepatotrophic effect of intestinal venous blood was seemingly unmasked by the special conditions of these experiments. Analogous conclusions have been reached from observations after a two month interval (37, 40).

Until now, the discussion has focused almost entirely upon double liver fragment preparations exclusive of partial hepatectomy. The addition of a 30 or 60 per cent hepatic resection in normal, and otherwise unaltered, dogs evoked the well known changes of healthy hepatocyte hyperplasia and hypertrophy through the whole liver. In nondiabetic dogs with splanchic division or partial portacaval transposition, these findings were most prominent in the optimally perfused liver lobes. However, the lobes deprived of total splanchic flow or its pancreaticogastroduodenosplenic component also participated in the regeneration efforts, although at a lower rate. It was interesting that the disadvantaged lobes appeared generally more healthy than in the companion nonhepatectomy experiments, as if these lobes had benefited or been provided protection by something about the over-all regeneration response. In the same kinds of experiments, the hepatic regeneration response was markedly inhibited bilaterally if diabetes was superimposed and, to an even greater extent, if insulin therapy was omitted postoperatively. A clearer demonstration of the participation of portal blood factors in regeneration and the permissive role of insulin could hardly be imagined. Yet, the same collection of experiments has shown the inexorability of regeneration and its lack of total dependence upon any single known factor. The work of Younger (46) and Short (34) and their associates has made the same point.

In the 11 years since the first papers from our laboratory reopened the possibility that portal blood contents affect liver structure and function, a number of workers have tested the role of hepatotrophic substances in the acute regeneration after liver resection and often with conflicting conclusions. After a first negative report by Lee and Edgington (16) consistent support for the concept of hepatotropic portal factors has come from surgical research laboratories of the University of California, San Diego, by Lee (17), Chandler (6), Sgro (33) and Broelsch (1) and their associates with studies using double liver fragment preparations or techniques of selective visceral extirpation. Fisher (10, 11, 12), an earlier opponent of the hepatotropic hypothesis, became one of its adherents (13), although incorrectly localizing the liver supporting splanchic factors to the intestine (14), as has been shown by our studies and those of Sgro (35) and Broelsch (6) and their associates and of Poirier and Cahow (26). The contention of Price (28) and Whittemore (45) and their associates that glucagon is the principal hepatotropic substance now seems indefensible for reasons already stated. Weinbren, who long ago showed that liver regeneration can occur after Eck fistula (43), remains the major spokesman against the role of portal blood factors in regeneration (44).

A recent article by Bucher and Swaffield (4) is apt to have a conciliatory effect on these divergent points of view. These authors had previously minimized the role of portal blood factors in the governance of regeneration (3) but in their later work was described an important role by insulin and glucagon as regulators of the rate and extent of the regeneration process. Although our data reported herein and elsewhere in support of our broader vi view of the broader view, the authors, in an elegantly manner, time by Bucher and Swaffield (4), agree to the villainous agent or trigger in the several effectors or circumstances.

SUMMARY

The acute intestinal factors upon regeneration which provide the portal circulation with enriched ileal venous blood and other factors has been shown in the dog to be the primary hepatotrophic influence and by the previous experiments with the acute islets of the 11-year period, the role of portal blood has been greatly weakened. The contribution of the intestine to liver regeneration is now clearly established, and the role of insulin and glucagon in the process is being reassessed.

The acute regeneration process is a complex interaction of multiple factors, including portal blood factors, insulin, glucagon, and other nutrients. The role of insulin in promoting cell division is well documented, and recent studies have suggested that glucagon may also play a significant role. The contribution of the intestine to liver regeneration through the portal circulation is a critical factor, and further research is needed to fully understand the interplay between these factors.
its, although at a disadvantageously in comparison to the authors in this study, as if these data had been collected under similar conditions, the results are still consistent with previous observations made by Buschke and Malt (20) who stated, "... nor is there any basis for choosing between a unitary agent or trigger mechanism versus an interplay of several effector processes with shifts in balance according to circumstances."

**SUMMARY**

The acute influence of portal blood hepatotropic factors upon the canine liver and upon hepatic regeneration was studied after surgical operations which provided qualitatively different portal venous perfusion to the right and left liver lobes. With one such procedure called splanchic division, the nutrient rich venous return from the intestines was directed to the left lobes, whereas the hormone rich blood from the pancreas and other splanchic organs of the upper part of the abdomen passed to the right lobes. Within three to five days, the rate of cell division on both liver sides was increased as judged by autoradiography, but the hormone influenced right lobes exhibited hypertrophy and hyperplasia relative to the nutrient enriched left lobes. In the latter, the hepatocytes underwent pronounced atrophy, deglycogenation, depletion or distortion of the rough endoplasmic reticulum, fatty vacuolization and other structural changes. When 30 or 60 per cent hepatic resection was carried out at the same time as splanchic division, the regeneration of the hormone dominated hepatic tissue after three to five days was greater than that of the hepatic tissue receiving the intestinal venous effluent, as judged by multiple criteria, although both liver sides participated in the regeneration process.

The advantage enjoyed by the right liver lobes in relation to the left liver lobes both in the resting or in the regenerating state after splanchic division was reduced or eliminated by pre-existing alloxan-induced diabetes or after concomitant total pancreatectomy. Similar, but less complete, observations about the effect of pancreatectomy were made in dogs submitted to the procedure of partial portacaval transplantation, in which all the splanchic venous blood passed to the right lobes, whereas the left lobes were revascularized with systemic venous blood from the vena cava.

These observations have added to the recent torrent of evidence that insulin is the most easily demonstrable and, therefore, probably the most important specific hepatotropic factor in portal venous blood. At the same time, further subtle support has been added to our previously proposed hypothesis that multiple other hormonal and possibly nonhormonal factors from the splanchic viscera and other sources also contribute to the essence of the hepatotropic effects. These effects were evident and quite advanced within a few days. A prominent hepatotropic role of glucagon was not identifiable.

**REFERENCES**


