Altered alkaline phosphatase activity in obese Zucker rats liver respect to lean Zucker and Wistar rats discussed in terms of all putative roles ascribed to the enzyme

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Abstract

Biliary complications often lead to acute and chronic liver injury after orthotopic liver transplantation (OLT). Bile composition and secretion depend on the integrated action of all the components of the biliary tree, starting from hepatocytes. Fatty livers are often discarded as grafts for OLT, since they are extremely vulnerable to conventional cold storage (CS). However, the insufficiency of donors has stimulated research to improve the usage of such marginal organs as well as grafts. Our group has recently developed a machine perfusion system at subnormothermic temperature (20°C; MP20) that allows a marked improvement in preservation of fatty and even of normal rat livers as compared with CS. We sought to evaluate the response of the biliary tree of fatty liver to MP20, and a suitable marker was essential to this purpose. Alkaline phosphatase (AlkP, EC 3.1.3.1), frequently used as marker of terms of all putative roles

Introduction

The increased demand of organs for orthotopic liver transplantation (OLT) has stimulated the research of strategies to increase the pool of donors by including the so-called marginal organs, especially fatty livers. Liver steatosis, which is frequently observed in potential donors, has been considered an important risk factor for injury caused by conventional cold storage with a higher incidence of postoperative primary non-function and poor initial function of the graft, and lower patient and graft survival.1-3 As a matter of fact, steatotic livers are more susceptible to ischemia/reperfusion injury after transplantation, with loss of viability of sinusoidal endothelial cells, sinusoidal congestion and rupture of hepatocytes releasing fat droplets, causing microcirculatory failure.4-6

Biliary complications are considered the Achilles’ heel of liver transplantation because of their frequency and potential lethal effect on the survival of both graft and patient.5-11 These complications are a major cause of morbidity and graft failure in patients after OLT and include a wide spectrum of functional and anatomical abnormalities, such as bile leakage and large biliary duct strictures; their incidence is estimated to be 8-20% and may lead to acute and chronic liver injury.2-14 The fast restoration of biliary secretion is an important index of hepatic functional restoration after preservation by cold ischemic storage (CS),15 and bile analysis is a useful tool to assess the integrity of biliary epithelial cells after cold ischemia/reperfusion of rat liver.16 The formation of bile depends on the structural and functional integrity of the biliary tree and its impairment results in the syndrome of cholestasis.

Experimental and clinical studies have indicated that bile formation early after liver transplantation may be disturbed, resulting in more cytotoxic bile with a relatively low phospholipid/bile salt ratio.7 A close relationship between this ratio early after liver transplantation and injury of the small bile ducts in the liver has been found, supporting the hypothesis that early changes in bile composition contribute to the relatively late strictureting of the large bile ducts.14 This data suggest that the different cell types of the biliary tree (hepatocytes, transition cells of Hering canals and cholangiocytes in small and large bile ducts) play different roles in the progressive post-transplantation injury phases to the biliary tract.

The analysis of the biliary tree behaviour towards transplantation has however lagged behind studies concerning hepatocytes and sinusoidal cells, even though bile canaliculi are one of the liver structures that are damaged markedly and early during ischemia/reperfusion occurring in patients undergoing OLT.13 Furthermore, the damage to bile duct cells is important in the long-lasting phase of reperfusion injury,8 the recovery of the biliary tree from preservation injury takes longer compared with hepatocytes or endothelial cells12 and regeneration of its cellular ATP is
that do not present the mutation. They thus cannot respond to the satiety stimulus and became hyperphagic, obese, hyperinsulinenic (insulin-resistant), but have normal blood glucose levels and do not develop diabetes. The lean ZR that are heterozygous for the allele fa maintain a lean phenotype throughout life with normal blood insulin and glucose levels.

Our group has recently developed a machine perfusion (MP) system at subnormothermic temperature, 20°C (MP20) with a low-viscosity perfusion medium based on Krebs-Henseleit buffer and containing N-acetylcysteine, glucose and low concentration of calcium, that appears as a promising strategy to protect the liver of normal rats (Wistar, lean Zucker) and fatty liver (FL) of obese ZR performing biochemical, histochemical and ultrastructural analyses. In particular, we have shown that, compared with traditional cold storage, MP20 leads to FL preservation in terms of enzyme release into the perfusate and bile, energy charge, glycogen stores and reactive oxygen species production. Histochemical analyses revealed that MP20 caused a marked reduction of steatosis through ketogenesis (preliminary results) and of parenchymal and sinusoidal damage to injury have been conducted in genetically susceptible or decreased tolerance of fatty livers and insulin-resistance. As controls, we used obese Zucker rats (ZR), frequently used as models of obesity and diabetes. The lean ZR that are heterozygous for the allele fa maintain a lean phenotype throughout life with normal blood insulin and glucose levels.

Male Wistar rats (250-300g) (Harlan-Nossan, Corezza, MB, Italy), 11-12 week old obese (fa/fa) (375±15 g) and lean (fa/+) (300±10 g) male ZR (Charles River, Calco, LC, Italy) were used. The animals were allowed free access to water and food in all the experiments. The use and care of animals in this experimental study was approved by the Italian Ministry of Health and by the University Commission for Animal Care. Rats were anesthetized with sodium pentobarbital (40 mg/kg i.p.) and received 250 U of heparin via the inferior vena cava before liver uptake.

Small tissue blocks of about 0.5 cm³ were cut, then inserted in cryovials and snap-frozen immediately in liquid nitrogen. Afterwards, the material was kept at -80°C until further use. Cryostat sections, 8 μm thick, were cut at -24°C on a Leica CM 1850 cryostat.

**Morphology**

Samples of the liver of obese rats were quickly removed, and all small fragments were fixed by immersion in 2.5% glutaraldehyde in 0.13 M Millonig buffer (pH 7.2-7.4) at 4°C for 4 h, rinsed, post-fixed with 1% osmium tetroxide at 4°C for 2 h, washed, dehydrated through graded concentrations of alcohol, and embedded in Epon. Semi-thin sections (1 μm thick) were stained with 1% toluidine blue.

**Microscopy and photomicrography**

The slides were observed with Zeiss Axioskop 2 Plus light microscope (Carl Zeiss Microimaging, Jena, Germany) equipped with differential interference contrast (DIC) system and a Olympus C-4040 Zoom digital camera (Olympus, Tokyo, Japan) with 4 Mpixel of resolution. Digital images were elaborated with Adobe Photoshop 5 (Adobe Systems Inc., San José, CA, USA) and processed with Image Pro Plus 4.7 image analysis software (Media Cybernetics, Inc., Bethesda, MD, USA). A selective discrimination based on colour level was performed in order to highlight with false colour the distribution pattern of AlkP.

**Materials and Methods**

**Chemicals**

Unless otherwise stated, all reagents were of the highest purity grade available and were purchased from Sigma (Milano, Italy).

**Animals**

**Results**

With the indoxyl-tetrazolium salt method, the activity of AlkP is demonstrated by a dark brown final reaction product. A light brown diffuse staining was also observed in the controls lacking the substrate, therefore corresponding to non-specific reaction usually mentioned as nothing dehydrogenase reaction (NDH).

**Wistar rat liver**

In Wistar rats (Figure 1), an irregular but strong staining was present in bile canaliculi, especially in the periportal region but also in the mid-zone (Figure 1a and 1b); a light diffuse staining was seen the cytoplasm of portal hepatocytes. Small bile ducts and Hering canals were intensely coloured, but large intralobular bile ducts, whose morphology is clearly delineated by DIC, were negative (Figure 1a, 1c and 1d). In Hering canals the reaction was cytoplasmic whereas in small bile ducts it was located mainly in the apical and lateral membrane domains (Figure 1c). Reaction was also seen in the adventitia of large arteries, but not of small ones (Figure 1a).

**Lean Zucker rat liver**

Respect to Wistar rat liver, differences in reactivity were noticed only in hepatocytes (Figure 2). Namely, staining was less intense and present not only in bile canaliculi but also in the apical pole of hepatocytes and cholangiocytes and secreted in bile in large amounts that is frequently used as marker of membrane transport in hepatocytes and bile ducts. The determination of AlkP in the bile is used as an index of damage to cholangiocytes. Since no data were available concerning this enzyme in fatty liver, we investigated AlkP activity and its distribution in the liver of obese ZR, frequently used as models of obesity and insulin-resistance. As controls, we used either lean Zucker rats (the usual control for obese Zucker rats), heterozygous for the mutated allele (fa/+) and normal Wistar rats that do not present the mutation.

**Demonstration of alkaline phosphatase activity**

The indoxyl-tetrazolium salt method utilizes a tissue protectant, the polyvinyl alcohol (PVA), in order to improve the localization of the coloured product in the exact zone in which the enzyme is present. Sections 8 μm thick were cut on a manually-driven Leica CM 1850 cryostat at a cabinet temperature of -24°C. The sections were picked up onto clean glass slides and stored at -80°C until used.

After a 5 min drying at room temperature, the cryosections were incubated with a medium containing 18% PVA dissolved in 100 mM Tris-HCl buffer (pH 9.0), 0.7 mM 5-bromo-4-chloro-3-indolyl phosphate, 0.44 mM 1-methoxy-phenazine methosulphate (mPMS), 10 mM MgCl₂, 5mM sodium azide and 5 mM tetranitroblue tetrazolium (TNBT), previously dissolved in a heated mixture of dimethylformamide and ethanol (1:1, v/v), since it is not possible to dissolve 5mM TNBT in aqueous media. The final concentration of these solvents in the incubation medium was 2%. Incubation lasted 20 min at 37°C. To stop the reaction and to remove the incubation medium the sections were rinsed with hot (45-50°C) tap water and the slide mounted with glycerine-jelly. Control sections were performed in the absence of substrate.
in the basolateral membrane domains (Figure 2a,b,c). As in Wistar rat liver, the AlkP activity was very strong in bile ductules but negative in larger bile ducts (Figure 2a).

**Obese Zucker rat liver**

In small and large bile ducts the AlkP distribution pattern in this animal model of obesity was similar to that seen in lean ZR and Wistar rats (Figure 3a and inset). Marked differences were observed in hepatocytes. In particular, hepatocytes in the perportal area were negative whereas macrosteatotic hepatocytes in the mid-zone displayed intense staining in canaliculi and moderate staining in basolateral membrane domains (Figure 3b). The mid-zone was furthermore characterized by moderately intense NDH reaction. The typical morphology of macro- and microsteatotic hepatocytes is shown in Figure 3c.

A better appraisal of the differences in subcellular distribution of AlkP activity in hepatocytes of Wistar, lean and obese ZR liver is shown in Figure 3 d-f, where the most intense levels of AlkP are evidenced by a blue false colour by means of image analysis.

**Discussion**

The biliary tree, considered the Achilles’ heel of liver transplantation is morphologically and functionally heterogeneous. The primary bile is secreted by hepatocytes into bile canaliculi, which drain into canals of Hering at the ductular-hepatocellular junction, lined in part by hepatocytes and in part by cholangiocytes. In the rat, Hering canals are in direct flow with small bile ducts (lined by cuboidal cells) that in turn empty into large bile ducts (lined by columnar cells). Cholangiocytes play an important role in water and electrolyte secretion, modulating canalicular bile through a sequence of both secretory and absorptive processes aimed at adjusting bile flow and alkalinity to the physiological needs.

Three mechanisms contributing to bile duct injury after liver transplantation have been postulated: injury due to preservation or ischemia/reperfusion, immunological processes and injury induced by cytotoxicity of biliary bile salts. Bile salts are potent detergents that may damage cells by affecting the integrity of cellular membranes, in particular by extracting phospholipids and cholesterol from membranes to form micelles. Normally, these toxic effects are prevented by neutralization of bile salts by phospholipids and the formation of mixed micelles in bile. Phospholipids are secreted into bile in human via the concerted action of canalicular multidrug resistance-3 (MDR3) P-glycoprotein and canalicular bile salts.

Steatotic liver, increasingly frequent among potential liver grafts for OLT, is often discarded. Besides investigating with a histochemical approach various parameters related to the metabolism and injury of parenchymal and sinusoidal cells of livers submitted to MP20, we sought to doc-
ument the response of the biliary tree using the most frequently used functional marker of biliary structures and integrity, alkaline phosphatase. Since no data were available we made a preliminary study, here reported, to describe its distribution and activity in fatty liver, taking normal Wistar rat liver and lean ZR liver as controls.

**Wistar rat liver**

The distribution of AlkP in Wistar rat liver is in keeping with the histochemical literature on the enzyme, independently of the method used for its visualization, that is the indoxyltetrazolium salt method used in the present research,35,43,44 the metal capture method or the simultaneous coupling method.36,37,45 However, as far as large bile ducts are concerned, our data disagree with data reported on reviews,38,46 according to which in the rat AlkP is expressed by large interlobular bile ducts but not by small bile ducts. In particular, a research showing that exogenous alkaline phosphatase was able to inhibit secretin-stimulated ductal secretion by blocking cystic fibrosis transmembrane regulator (CFTR) channels expressed only in large ducts,47 was said to be

![Figure 2. Alkaline phosphatase activity in lean Zucker rat liver.](image)

![Figure 3. Alkaline phosphatase activity in obese Zucker rat liver and comparison among the highest levels of AlkP activity in Wistar, lean and obese Zucker rat hepatocytes.](image)
in keeping with the expression of AlkP in large but not small ducts.\textsuperscript{40} However, Alvaro et al.,\textsuperscript{52} who used the same enzyme histochemical method we did, reported AlkP activity along bile canaliculi in the perportal region and in cholangiocytes, without mentioning which ducts and showing a photograph at low magnification, where the morphology of the stained bile structures cannot be evaluated. On the other hand, and as demonstrated herein, the observation of sections from all the three animal models under differential interference contrast never showed AlkP activity in cholangiocytes nor in the lumen of large ducts, but only in bile ductules and Hering canals. We suggest that what could be interpreted by Alvaro et al.,\textsuperscript{52} as basal activity in large ducts when sections were observed under bright field and low magnification, was in reality the reaction present in small bile ducts or Hering canals that form a marked plexus around the portal canal.\textsuperscript{48} The best choice for analysing unstained structures is indeed DIC.

Staining for AlkP activity of the adventitia of large arteries is in accordance with observations made on rat myocardium with the metal capture method and with antibodies against rat liver AlkP.\textsuperscript{20} The absence of staining in the sinusoidal endothelial cells has as well been reported before.\textsuperscript{50} It is worth recalling that whereas human liver AlkP activity occurs in the sinusoid and, to a lesser extent, biliary pole of hepatocytes,\textsuperscript{51} in rat liver activity is mainly localized in the canalicular membrane of hepatocytes and in portal triad blood vessels.\textsuperscript{52}

**Lean Zucker rat liver**

Respect to Wistar rat liver, a significant difference in intracellular AlkP activity in lean Zucker rat liver was seen in hepatocytes, whereas similar patterns and reaction intensities were seen in the other segments of the biliary tree. As a matter of fact, hepatocytes in lean ZR displayed AlkP activity not only in bile canaliculi but also (though less intense respect to obese rat liver) in basolateral membrane domains.

**Obese Zucker rat liver**

Respect to Wistar and lean ZR, in the obese rat liver the differences of AlkP activity were once more seen only at the hepatocyte level. In fatty liver, portal hepatocytes were negative whereas the most steatotic cells in the midzone had intense lateral membrane staining and only occasionally canicular activity, distorted by fat droplets.

In order to correlate the differences in AlkP activity in hepatocytes to metabolic features of the three animal models it would have been helpful to know the physiological role of AlkP and the nature of its natural substrate in hepatocytes and cholangiocytes. Unfortunately, these remain controversial, although AlkP has been routinly applied as a marker for liver function for over 70 years.\textsuperscript{27-30} In particular, the function and role of AlkP in the progression of cholestatic diseases are virtually unknown.\textsuperscript{47}

In the following we review and discuss several alternative proposals for AlkP function.

**Phosphorycholine hydrolysis**

Pekarthy et al.,\textsuperscript{35} suggested that at least one of the functions of canicular AlkP in hepatocytes was to hydrolyze phosphorycholine so that choline could be transported across the bile canalicular membrane; this hypothesis was later questioned by studies showing that purified rat liver alkaline phosphatase had no particular affinity for phosphorylcholine and that it could act instead in general as a non-specific ATPase.\textsuperscript{31} However, the speculation that AlkP might be involved in choline transport across cell membranes is still being put forth.\textsuperscript{56} This hypothesis could be relevant for the turnover of biliary phosphatidyl choline that may be degraded in bile by phospholipase C to phosphorylcholine and diacylglycerol.\textsuperscript{55} By catalysing the subsequent removal of phosphate from phosphorylcholine, alkaline phosphatase of hepatocytes would contribute to reabsorption of choline from bile.\textsuperscript{50} To our knowledge, no choline reabsorption in bile ducts has been reported.

Choline is an essential nutrient needed for the structural integrity and signalling functions of cell membranes; for normal cholinergic neurotransmission; for normal muscle function; for lipid transport from liver; it is the major source of methyl groups in the diet, a choline deprived diet may induce steatosis of the liver and eventually cause hepatocarcinogenesis.\textsuperscript{59} As far as membranes are concerned, choline is required for the biosynthesis of phosphatidylcholine, sphingomyelin and choline plasmalogens. The concentration of sphingomyelin is higher in the canicular domain of hepatocytes plasma membrane respect to the sinusoidal domain.\textsuperscript{60} Within the canicular domain, sphingolipids are particularly concentrated in lipid rafts.\textsuperscript{61} Plasmalogens have a relatively low concentration in the liver, but are known to be present mainly associated to sphingolipids in lipid rafts.\textsuperscript{62}

**Non-specific ectonucleotidases**

Earlier studies suggested that AlkP might act as an ecto-phosphatase regulating extra-cellular concentrations of phosphate compounds such as pyridoxal-5'-phosphate (the cofactor form of vitamin B6), phosphoethanolamine\textsuperscript{23} and some phosphoproteins.\textsuperscript{64} Other researches suggested that AlkP might play an important role in the modulation of purinergic signalling, that is, that of an ectonucleotidase capable of degrading extracellular ATP (or its derivatives ADP and AMP) to adenosine.\textsuperscript{51,65-67} Alkaline phosphatases remove phosphate groups in the 5’ and 3’ positions from several types of molecules including nucleotides.\textsuperscript{66} Extracellular nucleotides and, in particular, ATP, act as important autocrine/paracrine signalling molecules regulating hepatobiliary functions such as hepatocyte glycochen metabolism, cell volume, bile formation, and other cell functions. These effects are mediated by the activation of purinergic receptors. Hepatocytes release ATP in venous blood and in the bile, and cholangiocytes secrete ATP in the bile.\textsuperscript{68} Purinergic receptors have been identified in the plasma membrane of hepatocytes and cholangiocytes, their activation contributes to the regulation of metabolism, ion channel activation, coordination within the liver lobule of cell-to-cell Ca\textsuperscript{2+} signalling, cell volume regulation, secretion, and coupling of the separate hepatocyte and cholangiocyte contributions to bile formation.\textsuperscript{69} Nucleotides released in the extracellular space are rapidly inactivated by ectonucleotidases.\textsuperscript{70} Each liver cell type expresses its own repertoire of purinoreceptor subtypes and ecto-ATPases.\textsuperscript{61,71} Immunohistochemical studies of ecto ATP diphosphohydrolase (nucleoside triphosphate diphosphohydrolase) performed on pig liver showed strong reactivity in the bile canaliculi of hepatocytes (especially in the perportal region) in bile ducts, in the endothelium of the portal vein and in smooth muscle cells in the portal artery wall.\textsuperscript{72} This topological distribution colocalizes with the distribution of AlkP activity observed in the Wistar rat liver.

**Ecto- protein phosphatase**

Ecto-phosphatase is still another putative function for alkaline phosphatase.\textsuperscript{33,34} AlkP, being anchored to the plasma membrane, could dephosphorylate soluble cellular substrates, or cell surface proteins. Ectokinases were shown to phosphorylate both soluble substrates and membrane-bound proteins.\textsuperscript{73} These processes were though to regulate ligand binding, signal transduction and cell-to-cell interactions. Both protein kinases and phosphatases are thought to be required for reversible control of extracellular phosphorylation processes acting in a manner similar to that in which they control cytoplasmatic phosphorylation-dephosphorylation systems.\textsuperscript{29,30,35} The physiologic function of AlkP as an ecto-phosphatase remains controversial, mainly because early studies with purified AlkP describe the enzyme as exhibiting a non-physiological alkaline pH optimum.\textsuperscript{77} However, a pH optimum of 7-8 was determined for purified and plasma membrane-bound human liver AlkP.\textsuperscript{35,37} It must be
recalled that, although the main control of bile alkalinitation is performed by cholangiocytes, b) bicarbonate secretion takes place already at canalicular level and thus that the bile pH is compatible with AlkP activity.

The multidrug resistance-1 (MDR1) P-glycoprotein (ABC1) ATP-binding Cassette transporter) that transports cationic and neutral compounds and steroids out of cells was specifically investigated as a possible target of the above mentioned ectophosphorylation-dephosphorylation processes. MDR1/P-glycoprotein/ABC1 has been reported in bile canaliculi throughout the lobule and in the apical pole of cholangiocytes. Membrane-bound AlkP concentration in different tissues was shown to be positively correlated with the extent of exchange surface per unit volume of the tissue, suggesting an association between AlkP and transport systems. AlkP is located in close association with transporters that play a major role in the process of bile formation both in hepatocytes and cholangiocytes. P-Glycoprotein modulators were shown to significantly affect the activity of hepatic-AlkP and thus the two processes seem to be metabolically coupled.

Modulation of endotoxin toxicity

A further and physiologically completely different role proposed for AlkP is defence mechanism against endotoxin toxicity. Indeed human placental alkaline phosphatase (HPLAP) attenuates the lipopolysaccharide (HPLAP) attenuates the lipopolysaccharide (LPS) effect on endogenous lactate and nicotinamide adenine dinucleotide (NAD+) and to endogenous thiols groups of glutathione, cysteine and other tissue proteins. The higher levels of NDH observed in the liver of obese ZR without AlkP might not be incompatible, but instead refer to different anatomic locations within the biliary tree.

Discussion of histochemical patterns in terms of the putative functions of alkaline phosphatase

The non-specific light staining (NDH) is ascribed to lactate dehydrogenase activity acting on endogenous lactate and nicotinamide adenine dinucleotide (NADH) and/or to endogenous thiols groups of glutathione, cysteine and other tissue proteins. The higher levels of NDH observed in the liver of obese ZR respect to Wistar and lean ZR are in keeping with higher lactate dehydrogenase activity observed in the former (unpublished study) and with data indicating higher concentration of lactate in the liver of obese ZR respect to their lean controls.

The different behaviour of hepatocytes of lean ZR respect to Wistar rats, and in particular, the basolateral reactivity, similar to that observed in macrosteatotic hepatocytes of obese rats, might be a consequence of the fact that lean ZR are phenotypically normal but genotypically heterozygous for a recessive mutated allele.

The distribution of AlkP in hepatocytes of the steatotic liver of obese rats was markedly different from that of normal liver. The absence of canalicular enzyme activity in most hepatocytes suggests a substantial decrease in choline reabsorption and/or decreased modulation of purinergic signalling, and/or modulation of the activity of extracellular kinases. As a matter of fact, a marked reduction of bile salt-dependent and bile salt-independent bile secretion with significant functional and molecular alterations consistent with mild cholestasis were reported in obese ZR respect to lean animals. In particular was decreased the expression of some hepatocyte transporters such as the basolateral organic anion transporting polypeptide-2 (OATP2) and the canalicular multidrug resistance-associated protein 2 (MRP2/ABCC2). The conclusion of these papers, based exclusively on biochemical determinations, was that a defective hepatobiliary transport capacity could contribute to the higher susceptibility of obese ZR to liver injury. To our knowledge, no study has correlated AlkP activity to the activity of these transporters. Further important characteristics of the liver of obese ZR are compromised microcirculation that reduces the organ oxygenation and energy balance (low [ATP/ADP]) with decreased mitochondrial activity respect to lean animals. A lower energy charge and glycogen content of obese ZR liver respect to lean ZR was also reported by our group. Therefore, ATP-dependent transport across the canalicular membrane might be impaired due to low ATP availability.

The intense AlkP activity patterns in basolateral membrane domains in macrosteatotic hepatocytes are typical of cholestasis and were presumed to indicate the formation of extra-canaliculare sites for bile salt transport out of the hepatocytes in conditions where bile secretion is hindered. In cholestasis, a strong up-regulation of apical MDR1/ABC1 was reported, as a presumed defence reaction against toxic metabolites.

In conclusion, the absence of AlkP activity in the canaliculare of hepatocytes of obese Zucker rat liver indicates an impairment of processes related to primary bile formation, extracellular signalling modulation and/or choline reabsorption. By contrast, the processes mediated by AlkP in bile ductules and ducts are apparently not modified in these animals respect to normal ones. As lean ZR are concerned, the AlkP patterns, different from those of normal Wistar rats and are more similar to the patterns of obese rats, suggest that the functioning of their liver is not normal.

Alkaline phosphatase activity had been initially selected as a possible marker of the whole biliary tree response to the stresses endured by the fatty liver during the various phases of transplantation. This study demonstrates that it might be used only for documenting damage bile ducts.

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