Long-Term Treatment of Bile Duct-Ligated Rats with Rapamycin (Sirolimus) Significantly Attenuates Liver Fibrosis: Analysis of the Underlying Mechanisms

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ABSTRACT
Rapamycin is an immunosuppressant with antiproliferative properties. We investigated whether rapamycin treatment of bile duct-ligated (BDL) rats is capable of inhibiting liver fibrosis and thereby affecting hemodynamics. Following BDL, rats were treated for 28 days with rapamycin (BDL SIR) or sham-operated animals served as controls. After 28 days, hemodynamics were measured, and livers were harvested for histology/immunohistochemistry. Liver mRNA levels of transforming growth factor (TGF)-β1, connective tissue growth factor (CTGF), platelet-derived growth factor (PDGF)-β, cyclin-dependent kinase inhibitor p27Kip1 (p27), and cyclin-dependent kinase inhibitor p21WAF1/CIP1 (p21) were quantified by real-time polymerase chain reaction. Liver protein levels of p27, p21, p70 S6 kinase (p70S6K), phosphorylated p70S6K (p-p70S6K), eukaryotic initiation factor 4E-binding protein (4E-BP1), p-4E-BP1 (Thr37/46), and p-4E-BP1 (Ser65/Thr70) were determined by Western blotting. Portal vein pressure was lower in BDL SIR than in BDL CTR animals. Volume fractions of connective tissue, bile duct epithelial, and desmin- and actin-positive cells were lower in BDL SIR than in BDL CTR rats. On the mRNA level, TGF-β1, CTGF, and PDGF were decreased by rapamycin. p27 and p21 mRNA did not differ. On the protein level, rapamycin increased p27 and decreased p21 levels. Levels of nonphosphorylated p70S6K and 4E-BP1 did not vary between groups, but levels of p-p70S6K and 4E-BP1 (Ser65/Thr70) were decreased by rapamycin. Rapamycin had no effect on p-4E-BP1 (Thr37/46) and p-4E-BP1 (Ser65/Thr70) levels. In BDL rats, rapamycin inhibits liver fibrosis and ameliorates portal hypertension. This is paralleled by decreased levels of TGF-β1, CTGF, and PDGF. Rapamycin influences the cell cycle by up-regulation of p27, down-regulation of p21, and inhibition of p70S6K phosphorylation.
genic stimulus for HSC (Friedman, 1999). TGF-β1 levels are increased in liver fibrosis and regulate HSC activity in an autocrine mode. Another important cytokine that is involved in fibrogenesis is connective tissue growth factor (CTGF). CTGF is a downstream mediator of TGF-β1, mediating and potentiating some of the effects of TGF-β1 such as fibroblast proliferation and extracellular matrix production (Frazer et al., 1996) through a receptor that has not yet been defined.

The macrolide fungicide rapamycin, also known as sirolimus, was initially introduced into clinical practice as an immunosuppressive drug. In addition, it possesses potent antimicrobial and antiproliferative properties. The antiproliferative action of rapamycin is due to the ability to interfere with cell cycle progression in response to proliferative stimuli (Wiederrecht et al., 1995). Rapamycin binds intracellularly to FK506 (tacrolimus)-binding proteins (FKBPs). The most important binding protein regarding the rapamycin-sensitive pathway is FKBP12 (Fruman et al., 1995). The complex of FKBPI2 and rapamycin inhibits the mammalian target of rapamycin (mTOR) (Sabatini et al., 1994). It belongs to the family of phosphoinositide 3-kinase related kinases and is involved in the control of crucial growth-related cellular functions. mTOR is not part of a common linear signaling pathway. Signaling through mTOR is activated by the presence of sufficient nutrients (mainly amino acids) and phosphoinositide 3-kinase (PI3K) signaling. However, it is not clear yet whether PI3K directly regulates mTOR or whether the two pathways function independently.

mTOR activates the 40S ribosomal protein S6 kinase (p70S6K) and inhibits the eukaryotic initiation factor (eIF) 4E-binding protein-1 (4E-BP1) (Gingras et al., 1998) by phosphorylation, thus enabling the assembly of the eIF4F complex. The second downstream mediator of mTOR, p70S6K phosphorylates the 40S ribosomal protein S6 and thereby enhances the translation of mRNA of proteins essential for cell cycle progression. In addition, rapamycin decreases the concentration of cyclin D1 and thereby decreases retinoblastoma protein (pRb) phosphorylation (Morice et al., 1993), blocks the elimination of the cyclin-dependent kinase inhibitor p27 (Nourse et al., 1994), and inhibits expression of p21, another cyclin-dependent kinase inhibitor (Gaben et al., 2004).

The purpose of the present study is to investigate the impact of rapamycin on liver fibrosis and on the hemodynamics in bile duct-ligated (BDL) rats. In addition, the effect of rapamycin on profibrogenic and/or promitogenic cytokines such as TGF-β1, CTGF, and PDGF and on cell cycle regulatory proteins regulated by mTOR is studied.

Materials and Methods

Animals, Induction of Cirrhosis, and Medical Treatment.

Male Wistar rats were kept under a 12-h light/dark cycle with free access to rat chow (Kliiva, Kaiseraugst, Switzerland) and water. The animal experiments had been approved by a state-appointed board on animal ethics and were performed according to international guidelines concerning the conduct of animal experimentation.

To induce biliary cirrhosis, animals underwent double ligation and section of the common bile duct (BDL) under pentobarbital anesthesia (50 mg/kg intraperitoneally). In addition, six animals were sham-operated (Gross et al., 1987).

The rats were divided into three groups: BDL animals in the treatment group (BDL SIR) that were treated with 2.0 mg/kg/day rapamycin (Rapamune; Wyeth-AHP, Zug, Switzerland) in the drinking water; BDL animals in the nontreatment group that received no drug treatment (BDL CTR); and sham-operated animals that received no drug treatment. Treatment for 28 days was started immediately after BDL.

Hemodynamic Measurements and Tissue Conservation.

Hemodynamic measurements were performed under pentobarbital anesthesia (50 mg/kg intraperitoneally). A femoral artery and vein as well as a carotid artery were cannulated with a PE-50 tube (Clay Adams, Parsippany, New Jersey). Then the abdomen was opened, and a PE-50 tube (Clay Adams) was introduced into the portal via an ileocolic vein. Portal vein pressure and mean arterial pressure were measured using Statham transducers (Braun, Melsungen, Germany). Cardiac output and organ blood flow were determined using the microsphere technique (Groszmann et al., 1982). The catheter in the carotid artery was advanced into the left ventricle, 111In-labeled microspheres were injected, and the reference sample was withdrawn using a Harvard infusion pump (Harvard Apparatus Inc., Holliston, MA) set to 1 ml/min. Portosystemic shunting was determined by injection of 125I-Co-labeled microspheres into the portal vein. Then rats were killed by exsanguination while taking blood samples for analyzing the activity of AST, alkaline phosphatase, serum levels of the drug, bile acids, and TGF-β1 immunoassay. The visceral organs were removed and weighed. The liver was cut into pieces. Liver samples were either snap-frozen in liquid nitrogen and stored at −75°C for Western blotting, put into Tissue-Tek (Sakura Finetek Europe, Zoeterwoude, the Netherlands) and stored at −20°C for immunohistochemistry, put into RNAlater (Ambion, Huntingdon, UK) for RNA extraction, or stored in formaldehyde for histology. The remaining part of the liver was used for γ counting. The γ-isotopes in organs were measured on a Packard COBRA-II γ-spectrophotometer (PerkinElmer Life and Analytical Sciences, Hünenberg, Switzerland), with appropriate corrections for isotope spillover. Organ flow and portosystemic shunting was calculated as described by Groszmann et al. (1982).

RNA Isolation and Purification.

At least five randomly chosen samples from different regions of the liver with a total of 100 mg of tissue were taken and put into 1 ml of TRIzol (Invitrogen, Basel, Switzerland). The samples were disrupted and homogenized in TRIzol using an MM 300 vibration mill (Retsch, Haan, Germany), and RNA was extracted following the manufacturer’s guidelines. RNA concentration was measured spectrophotometrically using a Genequant Pro instrument (Amersham Biosciences, Dürendorf, Switzerland).

DNA Digestion and Reverse Transcription.

Total RNA (5 μg) from each liver sample was used. Prior to reverse transcription, samples were DNA-digested by incubation with RQ1 RNase-free DNase (Promega, Wallisellen, Switzerland). Reverse transcription was done using Moloney murine leukemia virus reverse transcriptase (Invitrogen) and random primers (250 ng; Microsynth, Bulgach, Switzerland). Control reactions did not contain reverse transcriptase.

Quantitative Real-Time PCR.

Primers and probes were designed using the Primer Express software (Applied Biosystems, Rotkreuz, Switzerland) and custom-synthesized by Microsynth or Applied Biosystems, respectively. Sequences and GenBank accession numbers of the primers and probes (except for p21, which was bought as an “assay on demand” from Applied Biosystems) are given in detail in Table 1.

Real-time PCR was performed using the ABI 7700 Sequence detector (Applied Biosystems) as described previously (Biecker et al., 2004). A dual-labeled fluorogenic probe (labeled with a “reporter” dye at the 5'-end and a second dye, quenching the emission of the “reporter” dye, at the 3'-end) complementary to a sequence within each PCR product was added to the PCR reaction. Cleavage of the probe during elongation by the exonuclease activity of the TaqDNA polymerase separates the reporter from its quencher. Accumulation of PCR products is detected in real time by monitoring the increase
in fluorescence of the reporter dye. The PCR reaction was performed in a volume of 25 μl containing 12.5 μl 2× TaqMan PCR master mix (Applied Biosystems) as well as 1.2 μl (equivalent to 300 ng of total RNA) of cDNA. The concentrations of the primers and probes are given in Table 1. Glyceraldehyde 3-phosphate dehydrogenase (GAPDH) provided as “ready-to-use” primers (100 nM each) and probe (200 nM) by the manufacturer (rodent GAPDH endogenous control, Applied Biosystems) served as the endogenous control.

The results were expressed as the number of cycles (Ct value) at which the fluorescence signal exceeded a defined threshold. The difference in Ct values of the target cDNA and endogenous control (GAPDH) is expressed as ΔCt values. Therefore, lower ΔCt values denote higher mRNA levels. The ΔCt method was used for quantification of the results. For all the used sets of primers and probes, a validation experiment was performed according to the manufacturer’s guidelines. It was demonstrated in those experiments that the efficiencies of the real-time PCR for the target and reference were approximately equal, and the ΔΔCt method could therefore be used for relative quantification.

**TGF-β1 Immunoassay.** Plasma samples of six animals from each group were taken and analyzed using a Quantikine TGF-β1 immunoassay (R&D Systems, Wiesbaden, Germany) following the manufacturer’s guidelines. Results were calculated by construction of a standard curve by plotting the mean absorbance for each standard on the y-axis against the concentration of the x-axis and drawing of a best-fit curve through the points on the graph. The data were then linearized by plotting the log of the TGF-β1 concentrations versus the log of the optical density, and the best-fit line was determined by regression analysis.

**Western Blot Analysis.** At least five randomly chosen samples from different regions of the liver with a total of 500 mg of tissue were taken from each organ and put into 2 ml of 0.25 mol/l sucrose buffer containing a mixture of protease inhibitors (Complete Mini tablets; Roche Diagnostics, Mannheim, Germany) at 4°C and homogenized using an MM 300 vibration mill (Retsch). Samples were then centrifuged for 10 min at 12,000 g to remove cell debris. The supernatant was used in the subsequent experiments. Protein concentration was measured according to Lowry et al. (1951). Proteins from whole-liver homogenate were separated by SDS-PAGE and subsequently transferred to nitrocellulose membranes (Schleicher & Schull, Dassel, Germany). The membranes were blocked with 5% nonfat dry milk in incubation buffer at 4°C overnight and probed for 2 h with the primary antibodies at room temperature. The membranes were washed three times, incubated at room temperature for 1 h with peroxidase-conjugated secondary antibody, washed three times, and the signal was revealed by enhanced chemiluminescence (Western Lightning Chemiluminescence Reagent; PerkinElmer Life and Analytical Sciences). The data of the antibodies and conditions are given in detail in Table 2. Protein extracts of human embryonic kidney (HEK) cells harvested in the exponential growing phase served as positive controls. The intensity of antigenic signals was video-imaged (FujiFilm LAS-1000; FujiFilm, Düsseldorf, Germany) and quantified by using the AIDA software package (Raytest, Überlingen, Switzerland). Densitometric integrated values (area of the band × density of the band) were determined, and the results were expressed as arbitrary units.

**Immunohistochemistry.** Immunohistochemistry for desmin and α-smooth muscle actin (SMA) was performed on one randomly selected piece of rat liver per animal as described previously (Tieche et al., 2001). In brief, desmin was demonstrated by a mouse-anti-desmin monoclonal antibody (clone D3; Dako, Glostrup, Denmark) at a concentration of 5.3 μg/ml and an incubation time of 60 min. Following the washing steps, a rabbit-anti-mouse IgG antibody, absorbed with rat serum (Dako) diluted 1:50, served as the link antibody to the alkaline phosphatase-anti-alkaline phosphatase mouse monoclonal antibody (1:50 in Tris-buffered saline, 45 min; Dako) that was applied as the third layer. SMA was detected using a mouse-anti-SMA (clone 1A4; Sigma-Aldrich, St. Louis, MO) diluted 1:200 in Tris-buffered saline for 60 min. After the primary antibody, a biotinylated rabbit-anti-mouse IgG antibody, absorbed with rat serum (Dako), was applied (1:200, 45 min), followed by a streptavidin-biotin complex/alkaline phosphatase mouse (1:200, 45 min; Dako). Finally,
slides were developed in new fuchsin-naphthol AS-BI (Sigma-Aldrich), counterstained with hematoxylin, and mounted.

**Chromotrope-Anilin Blue Staining.** Three randomly selected pieces of rat liver per animal were fixed in neutral buffered 4% formaldehyde and embedded in paraffin. Sections (2–3 μm) were deparaffinized and fixed in Bouin’s solution (saturated picric acid, 40% formaldehyde, and 96% acetic acid; 15:5:1). Cellular nuclei were visualized using Weigert’s ferrum-hematoxylin. After incubation with 1% phosphomolybic acid, the sections were stained by chromotrope-anilin blue (CAB).

**Morphometry.** To estimate the volumetric density of different tissue components, a point-counting procedure was carried out on liver sections using a microscope with a sampling stage connected to a semiautomatic advance (Gross et al., 1987). In CAB-stained sections, 1500 points per animal were counted, and each of them was classified as overlying either hepatocyte, bile duct epithelial cell, or connective tissue.

Immunohistochemistry was used to distinguish hepatic stellate cells (desmin-positive) from their activated counterparts expressing α-smooth muscle actin (Tieche et al., 2001). Five hundred points per animal were counted, and hepatic stellate cells were further classified as either portal/portal or parenchymal (surrounded by hepatocytes). Results are presented as volume fraction (percentage of specific cells over total number of counted cells).

**Statistical Analysis.** The three groups (sham-operated rats, BDL rats with rapamycin treatment, and BDL rats without treatment) were compared by the analysis of variance/Bonferroni/Dunn method. The p-value was set to 5%, and p < 0.05 was considered statistically significant.

**Results**

**Animal Characterization.** Table 3 shows the animal characteristics. The body weight of BDL CTR rats was significantly higher than the body weight of the sham-operated animals. No significant differences were obtained comparing BDL SIR rats with sham-operated and BDL CTR animals. Liver weight was highest in the BDL CTR group, lower in the BDL SIR rats, and lowest in the sham-operated animals (p < 0.001 between all groups). Spleen weight of the sham animals and BDL SIR rats did not vary significantly but was significantly higher in the BDL CTR animals (p < 0.001). These results demonstrate that rapamycin treatment attenuates the increase in liver weight that typically succeeds BDL. Furthermore, the normal spleen weight in the BDL SIR rats is an indicator for the absence of portal hypertension. The laboratory values for AST, bilirubin, and bile acids reflect the positive effect of the rapamycin treatment. AST, bilirubin, and bile acids were markedly elevated in the BDL CTR group (p < 0.01 compared with BDL SIR and sham-operated rats), whereas values for BDL SIR and sham-operated animals did not differ significantly.

**Hemodynamic Studies.** As already indicated by the normal spleen weight, the hemodynamic parameters (Table 4) were favorably influenced by the rapamycin treatment. Mean arterial pressure was not significantly different between groups but showed a tendency for higher pressure in the BDL SIR compared with BDL CTR animals. Portal venous pressure was highest in the BDL CTR rats, lower in the BDL SIR animals, and lowest in the sham-operated rats (p < 0.01 between all groups). Cardiac output was higher in the BDL CTR rats than in BDL SIR rats and sham-operated animals; however, only the difference between sham-operated animals and BDL SIR rats was statistically significant (p < 0.05). Hepatic artery blood flow was significantly lower in BDL SIR than in sham-operated or BDL CTR (p < 0.001). Portal vein blood flow was highest in sham-operated animals and only slightly lower in BDL SIR animals. The lowest values were obtained in BDL CTR rats (p = 0.01 comparing BDL CTR rats with BDL SIR rats and p < 0.001 comparing BDL CTR rats with sham-operated animals).

**Morphometry.** To prove that the positive effect of the rapamycin treatment on the hemodynamic parameters was paralleled by an inhibition of the histological changes that follow BDL, we performed microscopic examination and morphometry (Fig. 1 and Table 5) of liver sections. The BDL SIR group had a significantly higher volume fraction of hepatocytes compared with the BDL CTR group. In contrast, the volume fractions of connective tissue, bile duct epithelial cells, and desmin- and actin-positive cells were significantly lower in the BDL SIR group than in the BDL CTR group. The comparison of actin- to desmin-positive cells gave mean fractions of 0.34 in the BDL SIR group and 0.71 in the BDL CTR group.

**Growth Factors.** Because we were able to show that long-term treatment of BDL rats with rapamycin blocks the development of fibrosis as well as bile duct proliferation, we studied the underlying mechanisms. As a first step, we investigated the liver mRNA steady-state levels of the most important profibrogenic and promitogenic factors. Figure 2a shows the results of the quantitative PCR for TGF-β1. Compared with sham-operated animals, BDL CTR rats exhibited a 5.0-fold increase (p < 0.001), whereas BDL SIR rats exhibited only a 2.5-fold (p < 0.05) increase in TGF-β1 mRNA steady-state levels. This inhibitory effect on the mRNA expression was paralleled by an inhibitory effect of rapamycin treatment on TGF-β1 plasma levels (Fig. 3). Compared with BDL CTR rats, rapamycin treatment decreased TGF-β1 plasma levels by 20%

Liver CTGF mRNA steady-state levels (Fig. 2b) were mas-

### Table 3
Animal characteristics of sham-operated, BDL CTR, and BDL SIR rats

<table>
<thead>
<tr>
<th>Treatment Group</th>
<th>Body Weight*</th>
<th>Liver Weight*</th>
<th>Spleen Weight*</th>
<th>AST†</th>
<th>Bilirubin**</th>
<th>Bile Acids***</th>
</tr>
</thead>
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<tr>
<td>Sham (n = 6)</td>
<td>303.8 ± 45</td>
<td>12.9 ± 2.9</td>
<td>0.8 ± 0.1</td>
<td>78.3 ± 8.7</td>
<td>84.1 ± 23.2</td>
<td>14.0 ± 8.5</td>
</tr>
<tr>
<td>BDL SIR (n = 8)</td>
<td>335.9 ± 28</td>
<td>20.3 ± 3.4</td>
<td>0.9 ± 0.1</td>
<td>92.6 ± 27.3</td>
<td>423.2 ± 167.0</td>
<td>58.2 ± 37.7</td>
</tr>
<tr>
<td>BDL CTR (n = 8)</td>
<td>362.1 ± 43.8</td>
<td>27.5 ± 0.7</td>
<td>2.3 ± 0.7</td>
<td>271.4 ± 79.0</td>
<td>1298.6 ± 686.1</td>
<td>279.5 ± 108.5</td>
</tr>
</tbody>
</table>

* p < 0.05 comparing BDL CTR rats with sham-operated rats; † p < 0.01 between all groups; ‡ p < 0.001 comparing BDL SIR rats with BDL CTR rats and between BDL CTR rats and sham-operated animals; § p < 0.001 comparing BDL CTR with BDL SIR and sham-operated rats; *** p < 0.01 comparing BDL CTR with BDL SIR and sham-operated rats; ** p < 0.001 comparing BDL CTR with BDL SIR and sham-operated rats.
with sham-operated animals (p < 0.05 comparing sham rats with BDL CTR rats; b, p < 0.001 comparing PDGF-β-chain ΔC_T values of sham-operated and BDL CTR rats; c, p < 0.001 comparing TGF-β1 ΔC_T values of BDL SIR and BDL CTR rats and p < 0.001 comparing sham-operated rats with BDL CTR or BDL SIR rats. Circles represent ΔC_T values of single animals; horizontal lines represent the mean mRNA steady-state ΔC_T values of each group. C_T, threshold cycle at which an increase in reporter fluorescence above a baseline signal can first be detected; ΔC_T, C_T of mRNA of interest minus C_T of the housekeeping gene GAPDH. Lower ΔC_T values denote higher mRNA levels. a, p = 0.001 comparing TGF-β1 ΔC_T values of sham-operated and BDL CTR rats; b, p < 0.001 comparing CTGF ΔC_T values between all three groups; c, p < 0.01 comparing PDGF-β-chain ΔC_T values of sham-operated and BDL CTR rats and p < 0.001 comparing PDGF-β-chain ΔC_T values of BDL SIR and BDL CTR rats and p < 0.001 comparing sham-operated rats with BDL CTR or BDL SIR rats.

**TABLE 5**

Results of morphometric analysis of liver sections of BDL rats with and without rapamycin treatment

<table>
<thead>
<tr>
<th>Group</th>
<th>Volume Fraction (%)</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>BDL CTR (n = 7)</td>
</tr>
<tr>
<td>Hepatocytes</td>
<td>55.4 ± 12.2</td>
</tr>
<tr>
<td>Connective tissue</td>
<td>10.4 ± 4.4</td>
</tr>
<tr>
<td>Bile duct epithelial cells</td>
<td>18.3 ± 7.5</td>
</tr>
<tr>
<td>Desmin-positive cells</td>
<td>14.5 ± 3.4</td>
</tr>
<tr>
<td>Actin-positive cells</td>
<td>11.1 ± 7.6</td>
</tr>
<tr>
<td>Actin/desmin-positive cells</td>
<td>0.71 ± 0.37</td>
</tr>
</tbody>
</table>

* Statistically significant differences between BDL rapamycin-treated and non-treated rats (p < 0.05).

**Fig. 1.** Histological sections of livers from BDL rats with and without rapamycin treatment. CAB stain of BDL animals without treatment (a) and BDL animals with rapamycin treatment (b). Arrows indicate bile duct proliferations and connective tissue. Immunohistochemistry for desmin-positive cells (arrows) in BDL animals without treatment (c) and BDL animals with rapamycin treatment (d). Immunohistochemistry for α-smooth muscle actin (arrows) in BDL animals without treatment (e) and with rapamycin treatment (f). Original magnification, 20×.

**Fig. 2.** Quantitative real-time PCR of total liver TGF-β1 (a), CTGF (b), and PDGF-β-chain (c) mRNA in sham-operated, BDL CTR, and BDL SIR rats. Circles represent ΔC_T values of single animals; horizontal lines represent the mean mRNA steady-state ΔC_T values of each group. C_T, threshold cycle at which an increase in reporter fluorescence above a baseline signal can first be detected; ΔC_T, C_T of mRNA of interest minus C_T of the housekeeping gene GAPDH. Lower ΔC_T values denote higher mRNA levels. a, p = 0.001 comparing TGF-β1 ΔC_T values of sham-operated and BDL CTR rats; b, p < 0.001 comparing CTGF ΔC_T values between all three groups; c, p < 0.01 comparing PDGF-β-chain ΔC_T values of sham-operated and BDL CTR rats and p < 0.001 comparing sham-operated rats with BDL CTR or BDL SIR rats.

Liver PDGF-β-chain mRNA steady-state levels (Fig. 2c) were increased by BDL by a factor of 29.2 (p < 0.001). BDL SIR rats showed an increase by a factor of 8.8 compared with sham-operated rats (p < 0.001; p < 0.01 comparing BDL CTR with BDL SIR animals). These results clearly show that rapamycin treatment inhibits the gene expression of the most important profibrogenic and promitogenic cytokines. However, a complete normalization to the levels of the sham-operated animals was not achieved.

**Cell Cycle Regulatory Proteins.** In addition to the effect on growth factors, rapamycin is known to have a direct inhibitory effect on the cell cycle machinery. Therefore, we
studied the effect of rapamycin treatment on the most important cell cycle regulatory proteins that are known to be affected by rapamycin. Figure 4a shows the mRNA steady-state levels of the cyclin-dependent kinase inhibitor p27 in the liver. Neither BDL alone nor BDL and rapamycin treatment caused significant changes in the p27 mRNA levels. The protein levels of p27 are given in Fig. 4b. In contrast to the findings on the mRNA level, significant differences were obtained on the protein level. Lowest levels were found in BDL CTR rats. Rapamycin treatment induced a significant increase in the p27 level but did not reach the p27 level of sham-operated animals (p < 0.01), but p4-E-BP1 (Ser65/Thr70) levels of BDL SIR rats did not differ significantly from BDL CTR animals or sham-operated rats.

Next, liver mRNA and protein levels of the cyclin-dependent kinase inhibitor p21 were studied (Fig. 5). Similar to the findings for p27, neither BDL nor BDL and rapamycin treatment significantly altered the mRNA steady-state levels (Fig. 5a). On the protein level (Fig. 5b), significant differences were obtained (p < 0.01 between BDL CTR and BDL SIR and sham-operated animals), with the highest levels in BDL CTR animals. Rapamycin treatment caused a significant decrease of p21 protein levels in BDL SIR animals compared with BDL CTR animals but did not reach the p21 level of sham-operated animals.

The activity of 4E-BP1 is regulated by phosphorylation and dephosphorylation of the phosphorylation sites Thr37/46 and Ser65/Thr70. Figure 6 shows the liver protein levels of nonphosphorylated 4E-BP1, p4-E-BP1 (Thr37/46), and p4-E-BP1 (Ser65/Thr70). Protein levels of total, nonphosphorylated 4E-BP1 did not differ between the treatment groups. Similarly, protein levels of p4-E-BP1 (Thr37/46) did not vary significantly between groups. In contrast, we found significantly lower protein levels of p4-E-BP1 (Ser65/Thr70) in BDL CTR rats compared with sham-operated animals (p < 0.01), but p4-E-BP1 (Ser65/Thr70) levels of BDL SIR rats did not differ significantly from BDL CTR animals or sham-operated rats. These results differ from the documented effect of rapamycin on 4E-BP1 phosphorylation in cell culture models, since we were not able to show an effect on 4E-BP1 phosphorylation in vivo.

As is the case for 4E-BP1, the activity of p70^sk^ is regulated by phosphorylation and dephosphorylation. It is shown in Fig. 7 that rapamycin treatment of BDL rats lowered p-p70^sk^ significantly (p < 0.001). Furthermore, p-p70^sk^ levels in BDL CTR rats were significantly lower than in sham-operated animals (p < 0.01; p < 0.01 comparing sham-operated and BDL CTR rats). To prove that the increase in p-p70^sk^ is not due to a relative increased expression of total, nonphosphorylated p70^sk^, we used an antibody against total, nonphosphorylated p70^sk^ and did not find a difference between groups.

We were not able to detect pRb in the livers of BDL CTR, BDL SIR, or sham-operated animals by Western blot analysis (Fig. 8). However, we found a strong signal in cultured HEK cells and cultured normal rat cholangiocytes (NRC), which were harvested in the exponential growing phase. Serum deprived (24 h) NRC cells showed a weaker band, whereas HEK cells, which were kept without serum for 96 h, displayed no band.

![Fig. 3. TGF-β plasma levels obtained with an immunoassay. Bars represent the mean values ± S.D. TGF-β plasma levels in the BDL CTR animals were higher (p < 0.02) than in the BDL SIR and sham-operated animals (p < 0.005) (n = 6 in all groups).](attachment:image3)

![Fig. 4. Quantitative real-time PCR (a), densitometric analysis of Western blots (b), and Western blot (c) of sham-operated, BDL CTR, and BDL SIR rat total liver p27 mRNA and protein levels. a, circles represent ΔCT values of single animals; horizontal lines represent the mean in p27 mRNA steady-state ΔCT values of each group. Differences are not statistically significant. b, bars represent the mean values ± S.D. of the Western blot band densities × band areas (arbitrary units) of the different treatment groups (p < 0.01 between all groups). c, bands show two representative animals of each treatment group. BDL CTR and BDL SIR rats (n = 8); sham-operated animals (n = 6).](attachment:image4)
Discussion

Prevention of liver fibrosis by rapamycin was first demonstrated in rats exposed to long-term treatment with CCl4 by Zhu et al. (1999). We expand this observation in another model and also demonstrate amelioration of hemodynamic changes, inhibition of bile duct proliferation, and, finally, the signal transduction pathway of the antifibrotic effects of rapamycin.

Rapamycin treatment of rats started immediately after bile duct ligation significantly inhibits the development of liver fibrosis, as evidenced by a decrease in the volume fraction of connective tissue and activated hepatic stellate cells. Activation of HSC is mediated predominantly by PDGF, TGF-β1, and its downstream mediator CTGF. All three cytokines were markedly up-regulated in BDL and reduced by a factor of 2 to 8 by rapamycin treatment. PDGF, the most potent proliferative factor for HSC, is synthesized not only by HSC but also by cholangiocytes during cholestasis (Grappone et al., 1999). The cytokine is a dimer composed of PDGF-A or -B with the three possible isoforms AA, BB, or AB. PDGF-BB is the most potent form in stimulating HSC proliferation; mRNA steady-state levels of the B-chain in the liver was decreased by half by rapamycin treatment. Further studies will be needed to delineate which cell type is most responsible for the decrease in fibrogenic cytokines.

The expression of TGF-β1 mRNA—thought to be the major stimulus for fibrogenesis in HSC (Friedman, 1999)—was decreased by half by rapamycin treatment. The reduced expression of TGF-β1 mRNA in rapamycin-treated animals was paralleled by only a minor decrease in TGF-β1 plasma levels; this is due to the predominantly auto- and paracrine action of TGF-β1. Our results are in line with a suppression of TGF-β1 secretion in an immortalized human HSC line (Shibata et al., 2003) and the decreased TGF-β1 mRNA levels in the liver of rapamycin-treated rats (Zhu et al., 1999). However, our results differ from the findings of other authors who described an increase in TGF-β1 mRNA and/or plasma levels by rapamycin in the plasma of human kidney transplant recipients (Khanna et al., 2002), human lymphocytes (Khanna, 2000), transformed and nontransformed cultured cells (Law et al., 2002), rat kidney (Shihab et al., 2004), and prostate cancer cells (van der Poel, 2004). Because these experiments were not conducted using either hepatocytes or HSC, we conclude that the effect of rapamycin in the liver might differ from the effect in other tissues.

The most dramatic effect was seen in the message of CTGF, a downstream mediator of TGF-β (Moussad and Briggerstock, 2000) involved in experimental and human liver cirrhosis (Paradis et al., 1999; Williams et al., 2000). CTGF is produced in HSC (Paradis et al., 1999; Williams et al., 2000) and proliferating cholangiocytes (Sedlaczek et al., 2001). Both cholangiocytes and activated HSC were markedly reduced by rapamycin treatment. Further studies will be needed to delineate which cell type is most responsible for the decrease in fibrogenic cytokines.

The hemodynamic changes induced by bile duct ligation in rats are favorably influenced by rapamycin treatment, as shown by a decrease in portal pressure and increase in portal blood flow. Interestingly, hepatic artery blood flow was significantly lower in BDL SIR compared with both BDL CTR and sham rats. An increase in hepatic arterial blood flow is seen in different models of portal hypertension, in particular in BDL owing to the marked ductular proliferation (Gross et al., 1987; Van de Casteele et al., 2001). Thus, a reduction in arterial flow in rapamycin treated animals is not unexpected in light of the reduction of ductular proliferation. However, hepatic arterial flow was decreased even compared with the sham-operated rats, suggesting some specific effect of rapamycin on hepatic arterial flow.

Stereological analysis has demonstrated inhibition of two expanding cell lines induced by BDL: cholangiocytes and HSC. The antiproliferative activity of rapamycin is well established (Hidalgo and Rowinsky, 2000). Rapamycin is thought to affect the cell cycle through inhibition of mTOR by the FKBP12-rapamycin complex, thereby inhibiting the phosphorylation of p70S6K and 4E-BP1, impairment of pRB hyperphosphorylation, and prevention of p27 down- and p21 up-regulation. p27 and p21 belong to the kinase inhibitor
Fig. 6. Densitometric analysis of Western blots (a) and representative Western blots of two animals of each treatment group (b) of total rat liver nonphosphorylated 4E-BP1, phosphorylated 4E-BP1 at Thr37/46, and phosphorylated 4E-BP1 at Ser65/Thr70. Bars represent the mean values ± S.D. of the Western blot band densities × band areas (arbitrary units) of the different treatment groups. There are no statistically significant differences between the treatment groups for nonphosphorylated 4E-BP1 and phosphorylated 4E-BP1 (Thr37/46). Only the difference of phosphorylated 4E-BP1 (Ser65/Thr70) in sham versus BDL CTR rats is significant (p < 0.01). BDL CTR and BDL SIR rats (n = 8); sham-operated animals (n = 6).

Fig. 7. Densitometric analysis of Western blots (a) and representative Western blots of two animals of each treatment group (b) of rat total liver nonphosphorylated p70^S6K and phosphorylated p70^S6K. Bars represent the mean values ± S.D. of the Western blot band densities × band areas (arbitrary units) of the different treatment groups. There are no statistically significant differences between the treatment groups for nonphosphorylated p70^S6K. p-p70^S6K, p < 0.001 comparing BDL CTR rats with BDL SIR rats and p < 0.01 comparing sham-operated animals with BDL CTR rats and BDL SIR rats. BDL CTR and BDL SIR rats (n = 8); sham-operated animals (n = 6).
protein family and are instrumental in the transition from G1 to S phase. p27 mRNA levels are constant throughout the cell cycle, whereas p27 protein levels are high in nonproliferating cells and low in S phase (Hengst and Reed, 1996). Rapamycin increases p27 protein levels in different cell types (Law et al., 2002; Woltman et al., 2002), thereby inhibiting proliferation. Moreover, fibroblasts and T-lymphocytes from p27 knockout mice exhibited resistance to the growth-inhibitory effect of rapamycin (Luo et al., 1996). Unchanged p21 mRNA levels in quiescent and proliferating cells have been shown by others (Gaben et al., 2004). p21 protein levels are low in proliferating and high in nonproliferating cells, as shown in hepatocytes (Ilyin et al., 2003) and transformed mouse fibroblasts (Gaben et al., 2004). Our observations of p27 and p21 regulation by rapamycin are consistent with these findings: We showed that liver p27 and p21 mRNA steady-state levels are not influenced by BDL and that rapamycin treatment significantly increased p27 and decreased p21 protein levels. This might contribute to the inhibition of bile duct proliferation and HSC activation.

The complex of rapamycin and FKBP12 inhibits mTOR (Schmelze and Hall, 2000). mTOR controls translation via activation through phosphorylation of p70<sup>60K</sup> (Shama and Meyuhas, 1996) and inhibition of the eIF4E inhibitor 4E-BP1 (Hara et al., 1998). p70<sup>60K</sup> is phosphorylated not only by mTOR, but also by the phosphoinositide-dependent kinase PDK1 that mediates phosphorylation at Thr252 (Alessi et al., 1998). We used an antibody specific for the Ser411 phosphorylation site, a target of mTOR (Isotani et al., 1999). The suppression of p70<sup>60K</sup> phosphorylation by rapamycin and, hence, the growth-inhibitory effect has been shown in a variety of different cell types (Jiang et al., 2001). With our findings, we add further evidence that rapamycin inhibits the phosphorylation of p70<sup>60K</sup> in vivo.

mTOR does not only phosphorylate p70<sup>60K</sup>, it also phosphorylates 4E-BP1. In a first step, phosphorylation sites Thr37/46 are phosphorylated by mTOR. This is the priming step for the resultant phosphorylation of Ser65 and Thr70 in response to the PI3K or mitogen-activated protein kinase pathways (Gingras et al., 1998, 1999). Nonphosphorylated 4E-BP1 inhibits the initiation of translation through the association with eIF4E. As expected, we did not find a difference in the protein levels of total, nonphosphorylated 4E-BP1 between the different animal groups. To our surprise, neither p4E-BP1 (Thr37/46) nor p4E-BP1 (Ser65/Thr70) protein levels were significantly influenced by rapamycin. Moreover, we found significantly more p4E-BP1 (Ser65/Thr70) in sham-operated rats than in BDL CTR animals. Most of the studies demonstrating inhibition of 4E-BP1 phosphorylation by rapamycin were done in cell culture. Few studies investig-

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