Retraction

MEDICAL SCIENCES


The authors wish to note: “We were informed by the Office of Research Integrity of the University of Colorado of the inappropriate duplication of several histological images in Figs. 2 A and D, 3 A–C, S2B, and S3B of our recent PNAS paper. There were no findings of scientific misconduct made against any of the authors. The problems with the histologic slides appear to be due to honest errors. We apologize to our colleagues and the scientific community for any inconvenience this might have caused. Based on these concerns, we request to retract our paper.”

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Signaling through hepatocellular A2B adenosine receptors dampens ischemia and reperfusion injury of the liver

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Ischemia and reperfusion significantly contributes to the morbidity and mortality of liver surgery and transplantation. Based on studies showing a critical role for adenosine signaling in mediating tissue adaptation during hypoxia, we hypothesized that signaling events through adenosine receptors (ADORA1, ADORA2A, ADORA2B, or ADORA3) attenuates hepatic ischemia and reperfusion injury. Initial screening studies of human liver biopsies obtained during hepatic transplantation demonstrated a selective and robust induction of ADORA2B transcript and protein following ischemia and reperfusion. Subsequent exposure of gene-targeted mice for each individual adenosine receptor to liver ischemia and reperfusion revealed a selective role for the Adora2b in liver protection. Moreover, treatment of wild-type mice with an Adora2b−/− selective antagonist resulted in enhanced liver injury, whereas Adora2b−/− agonist treatment was associated with attenuated hepatic injury in wild-type, but not in Adora2b−/− mice. Subsequent studies in mice with Adora2b deletion in different tissues—including vascular endothelia, myeloid cells, and hepatocytes—revealed a surprising role for hepatocellular-specific Adora2b signaling in attenuating nuclear factor NF-κB activation and thereby mediating liver protection from ischemia and reperfusion injury. These studies provide a unique role for hepatocellular-specific Adora2b signaling in liver protection during ischemia and reperfusion injury.

Hepatic ischemia and reperfusion injury is an important source of morbidity and mortality during major liver surgery and hepatic transplantation (1, 2). Extracellular adenosine is a signaling molecule that has been implicated in attenuating inflammation during conditions of hypoxia or ischemia (3–5). Indeed, conditions of limited oxygen availability—such as hepatic ischemia—are associated with increased production of adenosine from precursors, such as ATP, ADP, or AMP (6, 7). Hypoxia and ischemia are associated with the transcriptional induction of the ecto-5′ nucleotidase CD39 (ecto-5′ conversion of ATP to AMP) (8), and cd39−/− mice are more prone to hepatic ischemia and reperfusion, whereas CD39 is also critical for liver regeneration (9). Moreover, hypoxia also drives the transcriptional induction of the ecto-5′ nucleotidase CD73 (conversion of AMP to adenosine), and gene-targeted mice for cd73 experience a more severe phenotype in models of hepatic ischemia and reperfusion (10), thus suggesting a protective role of extracellular adenosine generation during hepatic ischemia and reperfusion injury. Based on these findings, we hypothesized that extracellular adenosine could represent therapeutic targets for attenuating hepatic ischemia and reperfusion injury. At present, the functional role of individual adenosine receptors (ARs), their tissue-specific roles, and the relevance of the adenosine-signaling pathway during human liver ischemia and reperfusion injury remain unknown. To make progress on these issues, we combined studies of liver biopsies in patients undergoing liver transplantation with pharmacologic and genetic studies in mice. Indeed, our findings point toward a protective role of adenosine receptor 2b (Adora2b) signaling on hepatocytes in liver protection.

Methods

Human Liver Tissue. Liver samples were obtained from patients undergoing orthotopic liver transplantation (Fig. 1A, SI Methods, and Table 51).

Murine Model of Partial Liver Ischemia. A murine model of partial liver ischemia was employed using a hanging-weight system as previously described (SI Methods) (11).

Transcriptional Analysis and Immunoblotting. Transcript levels and protein content were measured as previously described (SI Methods and Tables S2 and S3) (12–14).

ELISA and Assays. IL-6 and TNFα (R&D Systems), neutrophil sequestration, transcriptional activity of NF-xB and CAMP was quantified as previously described (see SI Methods) (14, 15).

Liver Histology. Liver tissue was stained as previously described (SI Methods) (10).

Isolation of Hepatocytes, Endothelial Cells, and Macrophages. Specific cell isolation was performed as described previously (SI Methods) (16–18).

Adora2b Genetic Deletion in Cell Cultures. Human hepatocytes (HepG2) were obtained from ATCC and cells were cultured and transfected as previously described (SI Methods) (10).

Statistical Analysis. Liver injury score data are given as median and range. All other data are presented as mean ±SD from three to eight animals per condition. We performed statistical analysis using the Student’s t test. A value of P <0.05 was considered statistically significant. For Western blot analysis three repeats were performed. For all statistical analysis GraphPad Prism 5.0 software for Windows XP was used.

Study Approval. Collection and use of patient samples were approved by the Colorado Multiple Institutional Review Board (COMIRB) at University of Colorado, Denver (COMIRB Protocol 10–0100) and informed consent was obtained. All animal protocols were in accordance with the US guidelines by the institutional animal care and use committee for use of live animals and were approved by the Institutional Animal Care and Use Committee of the University of Colorado guidelines for animal care.

Results

Selective Induction of Human Adora2b Transcript and Protein Levels During Hepatic Ischemia and Reperfusion Injury. Previous studies had indicated that the enzymatic control of extracellular adenosine contributes to liver protection from ischemia and reperfusion (9, 10).
induction of the ADORA2B (over 10-fold induction; \( P < 0.001 \)), whereas transcript levels of other ARs were unchanged (Fig. 1B). Moreover, we found dramatic increases of ADORA2B protein levels following warm ischemia and reperfusion for each individual patient (Fig. 1C). Taken together, these studies uniquely indicate that hepatic ischemia and reperfusion is associated with a selective induction of the ADORA2B during human liver transplantation.

Global Deletion of Adora2b Aggravates Murine Ischemia and Reperfusion Injury of the Liver. Consistent with the above studies during human liver transplantation, we observed a selective induction of Adora2b transcript and protein levels following hepatic ischemia (Fig. 1 D and E). As next step, we examined previously described mice with genetic deletion for individual ARs during hepatic ischemia and reperfusion (12). Indeed, mice gene targeted for Adora1 (Fig. 2A), Adora2A (Fig. 2B), or Adora3 (Fig. 2D) had similar degrees of hepatic tissue injury as corresponding littermate controls matched in weight, age, and sex exposed to 45 min of partial hepatic ischemia followed by 2 h of reperfusion. In contrast, Adora2b+/- mice exhibited significantly higher levels of tissue injury as examined by elevations of the transaminases aspartate transaminase (AST) and alanine transaminase (ALT) and liver histology (Fig. 2C). To correlate Adora2b expression levels with hepatic injury, we measured cAMP levels and hepatic neutrophil accumulation assessed by myeloperoxidase (MPO) measurements; Fig. S1C) were increased in Adora2b+/- mice compared with controls following liver ischemia and reperfusion injury. Moreover, second organ injury as examined by increases of IL-6 or myeloperoxidase in the lungs was significantly enhanced in gene-targeted mice for the Adora2b (Fig. S1D). In addition we performed experiments with prolonged reperfusion times. In these experiments, we followed 45 min of liver ischemia with 24 h of reperfusion. Indeed, Adora2b+/- mice exhibited significantly higher levels of tissue injury as examined by elevations of the transaminases AST and ALT (Fig. S1E) and liver histology (Fig. S1F) after 24 h of reperfusion time. Taken together, these studies in gene-targeted mice for individual ARs implicate a selective role for the Adora2b in liver protection from ischemia and reperfusion injury. Furthermore these findings highlight that Adora2b receptor expression levels correlate with Adora2b-mediated tissue protection from ischemia and reperfusion injury.

Pharmacologic Inhibition of Adora2b Aggravates Hepatic Ischemia and Reperfusion Injury. After having shown that genetic deletion of the Adora2b is associated with more severe hepatic injury and liver inflammation following ischemia and reperfusion, we next pursued pharmacologic studies with a previously described Adora2b antagonist 1-propyl-8-(p-sulfophenyl) xanthine (PSB1115). For the purpose of these studies, we administered PSB1115 30 min before the onset of ischemia via retroorbital injection (0.5 mg/25 g mouse i.v.). Consistent with the previous studies in gene-targeted mice for Adora2b, we observed elevated liver injury as assessed by elevations of transaminases (Fig. S2A), histologic tissue injury (Fig. S2B), liver inflammation (Fig. S2C), and second organ injury (Fig. S2D). Moreover, the observed alterations of liver injury following PSB1115 treatment were abolished in gene-targeted mice for Adora2b (Fig. S2E), thereby providing evidence for the specificity of PSB1115 for Adora2b. Together with the studies in Adora2b+/- mice.
mice, these findings indicate that inactivation of Adora2b via genetic deletion or pharmacologic inhibition significantly aggravates liver injury following hepatic ischemia and reperfusion.

**Treatment with Adora2b Agonist Provides Potent Protection from Hepatic Ischemia and Reperfusion Injury.** After having shown that targeted deletion or specific inhibition of the Adora2b is associated with enhanced tissue injury, we next examined a potential therapeutic role for Adora2b agonist treatment during liver ischemia and reperfusion. Indeed, we recently described a specific and effective Adora2b agonist [2-[6-amino-3,5-dicyano-4-(4-cyclopropylmethoxy)phenyl]pyridin-2-ylsulfanyl]acetamide (BAY 60–6583)] (12). For the purpose of these studies, BAY 60–6583 (0.25 mg/25 g mouse i.v.) was administered via retro-orbital injection 30 min before the induction of liver ischemia for 45 min and 2 h reperfusion. Indeed, Adora2b agonist treatment was associated with a marked reduction in liver injury and inflammation as assessed by plasma levels of transaminases AST, ALT (Fig. S3A), hepatic histology (Fig. S3B), liver inflammation (Fig. S3C), and second organ injury of the lungs (Fig. S3D). Additional studies in gene-targeted mice for Adora2b revealed that the protective effects of BAY 60–6583 are completely abolished in Adora2b<sup>−/−</sup> mice, thereby providing evidence for the specificity of BAY 60–6583 for Adora2b (Fig. S3E). To test the long-term effect of liver protection via Adora2b agonist (BAY 60–6583) treatment, we performed experiments with 24 h reperfusion following 45 min of liver ischemia. Indeed, Adora2b agonist treatment provided similar protection after 24 h of reperfusion as indicated by a robust reduction in plasma levels of transaminases AST, ALT, and hepatic histology (Fig. S4 A and B). Together, these studies provide evidence that Adora2b agonist treatment is therapeutic during liver ischemia and reperfusion injury.

**Identification of Hepatocyte-Specific Adora2b Signaling in Liver Protection from Ischemia and Reperfusion Injury.** As the next step, we pursued studies to address the tissue-specific contributions of...
Adora2b signaling in hepatic ischemia and reperfusion injury. Indeed, we used a recently described transgenic mouse line with a “flxed” Adora2b gene (12). We used this mouse line to generate mice with tissue-specific deletions of Adora2b in different cellular compartments of the liver. As such we generated mice with endothelial Adora2b deletion (Adora2bfloxP/loxP VE-cadherin Creκ), deletion of Adora2b on macrophages, monocytes, and granulocytes (Adora2bfloxP/loxP Lysm Creκ), and deletion of Adora2b on hepatocytes (Adora2bfloxP/loxP albumin Creκ). As shown in Western blots from isolated endothelial cells, Adora2b expression was significantly attenuated in Adora2bfloxP/loxP VE-cadherin Creκ mice compared with VE-cadherin Creκ controls (Fig. S5A). Similarly, Western blot analysis for Adora2b expression from isolated macrophages demonstrates that Adora2b expression was significantly attenuated in Adora2bfloxP/loxP Lysm Creκ mice compared with Lysm Creκ controls (Fig. S5B). Finally, Adora2b expression was essentially undetectable in Western blots from isolated hepatocytes derived from Adora2bfloxP/loxP albumin Creκ compared with controls (Fig. S5C). Based on previous reports indicating an important role of Adora2b in vascular leakage during hypoxic conditions (19) or during ischemic injury of the kidneys (12, 16), we first attempted studies in mice with endothelial Adora2b deletion. Adora2bfloxP/loxP VE-cadherin Creκ mice exposed to 45 min of ischemia followed by 2 h of reperfusion showed no difference in liver injury as assessed by plasma transaminase levels and liver histology (Fig. 3A). Because Adora2b signaling has been implicated in macrophage functions (20, 21), we subsequently exposed Adora2bfloxP/loxP Lysm Creκ mice to 45 min of ischemia followed by 2 h of reperfusion. However, we observed no difference in liver injury between the experimental groups and liver histology (Fig. 3B). As the next step, we examined Adora2bfloxP/loxP albumin Creκ mice with hepatocellular deletion of Adora2b to similar experimental conditions. Interestingly, these mice demonstrated a similar phenotype as we had previously observed in mice with global deletion of Adora2b. Indeed, we found dramatic increases in liver injury as assessed by plasma transaminase levels and liver histology (Fig. 3C) determined following 45 min of liver ischemia and 24 h of reperfusion. Together, these studies indicate a hepatocyte-specific role of Adora2b signaling in liver protection from ischemia.

**Hepatocyte Adora2b Signaling Attenuates Hypoxic Nuclear Factor NF-κB Activation in Vitro.** After having shown that Adora2b-dependent liver protection from ischemia and reperfusion predominantly involves hepatocyte-specific Adora2b signaling, we next attempted studies to gain additional insight into the mechanism of Adora2b-dependent liver protection. Previous studies had shown that hypoxic conditions such as ischemia and reperfusion are associated with the posttranslational stabilization of NF-κB, leading to a subsequent activation of proinflammatory gene programs (22). A very elegant study from the laboratory of Sean P. Colgan demonstrated that Adora2b signaling in hypoxic lungs represents a means of preventing the activation of NF-κB during hypoxic conditions (14). Based on these findings, we hypothesized that hepatocyte-specific Adora2b signaling could function as a break for hypoxic NF-κB activation during hepatic ischemia and reperfusion. To address this hypothesis, we used a lentiviral approach to generate a hepatocyte cell line (HepG2) with siRNA-mediated permanent repression of ADORA2B (Fig. S6A and B). When we exposed control cells to conditions of ambient hypoxia (1%
oxygen, 24 h) we observed robust increases in NF-κB reporter activity (Fig. S6C) or its regulatory component IkBα (Fig. S6D). This response was significantly increased in cells with siRNA-mediated repression of ADORA2B. Moreover, increases of hypoxic NF-κB or IkBα stabilization were significantly followed by Adora2b inhibition with PSB115, whereas treatment with Adora2b agonist BAY 60-6583 dampened this response. Together, these in vitro studies indicate that hepatocyte-specific ADORA2B signaling attenuates hypoxia-driven activation of proinflammatory programs involving NF-κB.

**Pharmacologic Inhibition of NF-κB Restores a “Normal” Phenotype in Gene-Targeted Mice for Adora2b.** Encouraged by the above in vitro studies indicating that ADORA2B signaling on hepatocytes can dampen the hypoxic activation of proinflammatory gene programs via inhibition of NF-κB, we subsequently examined the functional consequences of pharmacologic NF-κB inhibition during in vivo conditions of hepatic ischemia. For the purpose of these studies, we used a previously described inhibitor of NF-κB (inhibitor of kappa B kinase, 10 mg N-(4-Pyrrolidin-1-yl-piperidin-1-yl)-4-(4-benzoylphenothien-2-yl-pirimidin-2-ylamino)phenylcarboxamide hydrochloride (IKK-16) i.v. 1 h before ischemia) (23). Indeed, IKK-16 treatment of mice with global deletion of the Adora2b was associated with the restoration of a phenotype similar to what we had previously observed in wild-type mice, including attenuation of transaminase plasma levels or histologic liver injury (Fig. S7A). Moreover, the increased susceptibility of mice with hepatocyte-specific deletion of the Adora2b (Adora2blox/lox albumin Cre+) was attenuated (Fig. S7B). Finally, treatment of wild-type mice with IKK-16 was associated with dampened hepatocyte injury as assessed by plasma transaminases and histologic liver injury (Fig. S7C). Taken together, these studies indicate that increased susceptibility of gene-targeted mice for Adora2b to hepatic ischemia and reperfusion injury can be reverted via pharmacologic inhibition of NF-κB. As such, these findings implicate Adora2b signaling in an endogenous feedback loop to dampen ischemia and reperfusion-elicited increases in hepatocyte inflammation and point toward a therapeutic role for Adora2b agonists in treating ischemia and reperfusion injury of the liver.

**Discussion**

Ischemia and reperfusion injury of the liver represents one of the major challenges during liver surgery and liver transplantation. Due to a lack of donor organs, marginal livers are more frequently used for transplantation. This leads to a dramatic increase in the risk for developing severe ischemia and reperfusion injury during liver transplantation. Therefore, the search for novel pharmacologic strategies to dampen hepatic ischemia and reperfusion injury is currently an area of very intense investigation (2). In the present studies, we pursued the hypothesis that signaling events through Adora2b receptors expressed on hepatocytes, and CD8+ T cells play a detrimental role in models of ischemia and reperfusion. We gained insights from these findings in a mouse model of hepatic ischemia and reperfusion injury (Fig. S7A). Moreover, combinations of in vitro and in vivo studies suggest that the observed protection mediated by Adora2b-signaling functions through adenosine-dependent inhibition of hypoxic NF-κB activation. Taken together, the present studies demonstrate what we believe to be a previously unrecognized role for ADORA2B signaling in liver protection from these detrimental effects.

In the present studies, we observed a selective induction of Adora2b in patients undergoing liver transplantation or in mice exposed to in situ conditions of hepatic ischemia and reperfusion injury. Indeed, hepatic ischemia and reperfusion injury is characterized by the interdependent relationship of hypoxia and inflammation (22, 24–26). In this context, previous studies had examined transcriptional pathways that regulate Adora2b expression during conditions of limited oxygen availability. Indeed, first evidence for hypoxia as a transcriptional regulator of the ADORA2B comes from studies in cultured endothelial cells that were exposed to ambient hypoxia that demonstrated a selective and very robust induction of endothelial ADORA2B transcript and protein levels (27). Other studies identified a previously unknown binding site for the hypoxia-inducible factor (HIF) within the promoter region of human Adora2b (28). Moreover, consistent with our findings, a recent study in hypoxic preconditioning of the liver found liver protection via Adora2b signaling (29). Indeed, the finding that adenosine signaling can dampen hypoxic inflammation goes back to several landmark papers from the research laboratory of Michael Silovsky, that provided the first genetic in vivo evidence for adenosine receptor signaling in dampening inflammation (30) or inflammatory hypoxia (31). Together, these studies implicate hypoxia-dependent signaling pathways in the transcriptional control of Adora2b during conditions of limited oxygen availability, and suggest a protective role for such pathways in hepatic ischemia and reperfusion injury.

In addition to the present studies indicating that hepatocyte-specific adenosine signaling via Adora2b is protective, previous studies from the research laboratory of Joel Linden had implicated hypoxia-dependent adenosine receptor signaling in dampening inflammation (31). Moreover, a very recent study from the laboratory of Mark Okusa revealed that adenosine signaling through dendritic cells expressing Adora2a receptors contributes to attenuating ischemia and reperfusion injury of the kidneys (5). Interestingly, an agonist of ADORA2A (regadenoson) has been approved by the Food and Drug Administration as a coronary vasodilator for patients requiring pharmacologic stress echocardiography (36). Indeed, an ongoing multicenter, dose-finding, and safety trial of infused regadenoson has been initiated to study its safety and efficacy for the treatment of ischemia and reperfusion-related tissue injury in patients with sickle cell disease (37). As such, translational approaches targeting adenosine receptors during liver transplantation might be conceivable in the near future. Whereas there are experimental drugs available as agonists of ADORA2B (12, 38, 39), such compounds have yet to be examined in patients.

The present studies indicate that Adora2b signaling can function as a break for the activation of NK-κB during hepatic ischemia and reperfusion. Indeed, previous studies have shown that conditions of hypoxia or ischemia and reperfusion are associated with the stabilization of NF-κB (22, 26). Mechanistic studies from the laboratory of Cormac Taylor indicate that hypoxia-dependent NK-κB activation is controlled by oxygen sensing enzymes (prolyl hydroxylation domain proteins, PHDs), and thus shares many similarities with the post-translational control of HIF protein levels during hypoxia (26). These studies elegantly revealed that PHD1 negatively regulates NK-κB and hypoxia-associated inhibition of PHD1 subsequently results in the hypoxia-induced activation of NF-κB (26). Subsequent studies from the laboratory of Sean P. Colgan provided exciting insight into how adenosine signaling through Adora2b can interfere with the hypoxia-dependent activation of NK-κB (14). These findings are consistent
with the present studies showing attenuated NF-κB activation during hepatic ischemia and reperfusion via Adora2b signaling. Previous studies have shown that during conditions of hypoxia and ischemia, adenosine is predominantly derived from the breakdown of nucleotides—such as ATP, ADP, and AMP (9, 10, 39–43). However, there may also be alternative pathways of how extracellular adenosine can be enhanced during conditions of inflammation or ischemia. For example, studies from the laboratory of Edwin Jackson implicate a novel molecular pathway in the extracellular accumulation of adenosine. This pathway includes the extracellular generation of adenosine from 2′,3′-cAMP ("extra-cellular 2′,3′-cAMP-adenosine pathway") (44, 45). Other studies implicate adenosine release through equilibrative nucleoside transporters, for example in the brain or the heart—in elevating extracellular adenosine levels (46–48). Whereas some studies implicate CD39- and CD73-dependent nucleotide metabolism as an important source for hepatic adenosine generation during ischemia (9, 10), the contribution of the above discussed pathways in the context of hepatic ischemia and reperfusion injury has to be further examined.

Taken together, the present studies provide unique evidence for a protective role for extracellular adenosine signaling through AdoAR2B expressed on hepatocytes. Therapeutic extensions of these findings suggest that AOR2B agonists could represent a unique group of pharmacologic agents for the treatment or prevention of hepatic ischemia and reperfusion injury, which would be significant for hepatic transplantations and liver surgery in humans. Whereas these findings are very encouraging, future clinical trials should include the translation of these findings into a clinical setting, particularly with regard to the development of AdoAR2B agonists for the treatment of human disease or studies of their safety profile, and their efficiency to prevent or treat ischemia and reperfusion injury during liver transplantation.

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