

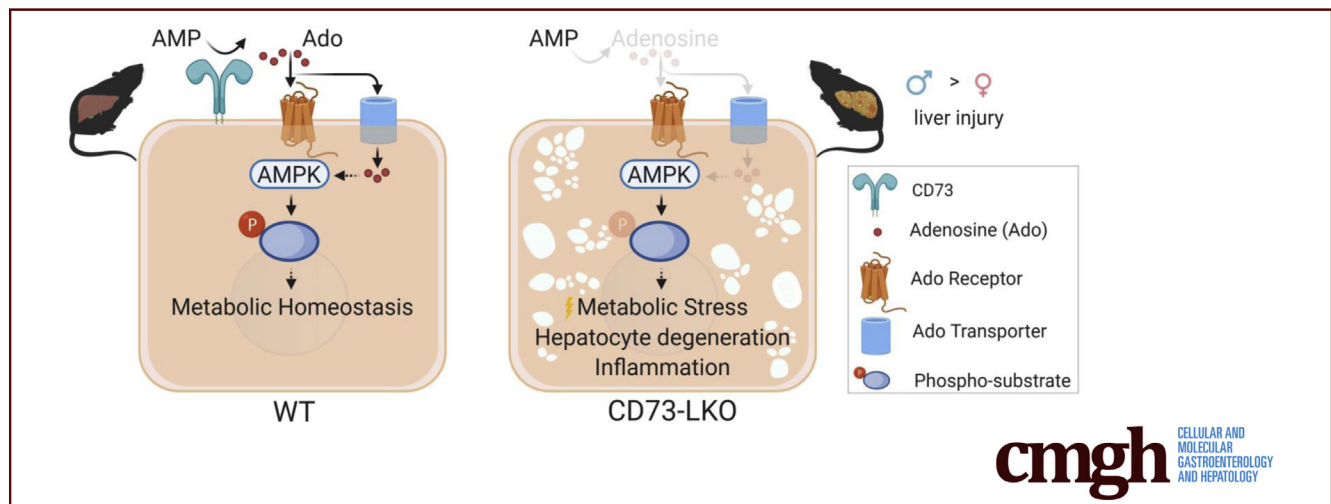
## ORIGINAL RESEARCH

## CD73 Maintains Hepatocyte Metabolic Integrity and Mouse Liver Homeostasis in a Sex-Dependent Manner



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## SUMMARY

We developed a new mouse model that showed that CD73, the major extracellular adenosine-producing enzyme, supports long-term liver homeostasis. These results advance our understanding of human disease and therapeutics because CD73 is down-regulated in chronic liver diseases and CD73 inhibitors are in clinical trials for cancer and coronavirus disease 2019.

**BACKGROUND & AIMS:** Metabolic imbalance and inflammation are common features of chronic liver diseases. Molecular factors controlling these mechanisms represent potential therapeutic targets. CD73 is the major enzyme that dephosphorylates extracellular adenosine monophosphate (AMP) to form the anti-inflammatory adenosine. CD73 is expressed on pericentral hepatocytes, which are important for long-term liver homeostasis. We aimed to determine if CD73 has nonredundant hepatoprotective functions.

**METHODS:** Liver-specific CD73 knockout (CD73-LKO) mice were generated by targeting the *Nt5e* gene in hepatocytes. The CD73-LKO mice and hepatocytes were characterized using multiple approaches.

**RESULTS:** Deletion of hepatocyte *Nt5e* resulted in an approximately 70% reduction in total liver CD73 protein ( $P < .0001$ ). Male and

female CD73-LKO mice developed normally during the first 21 weeks without significant liver phenotypes. Between 21 and 42 weeks, the CD73-LKO mice developed spontaneous-onset liver disease, with significant severity in male mice. Middle-aged male CD73-LKO mice showed hepatocyte swelling and ballooning ( $P < .05$ ), inflammation ( $P < .01$ ), and variable steatosis. Female CD73-LKO mice had lower serum albumin levels ( $P < .05$ ) and increased inflammatory genes ( $P < .01$ ), but did not show the spectrum of histopathologic changes in male mice, potentially owing to compensatory induction of adenosine receptors. Serum analysis and proteomic profiling of hepatocytes from male CD73-LKO mice showed significant metabolic imbalance, with increased blood urea nitrogen ( $P < .0001$ ) and impairments in major metabolic pathways, including oxidative phosphorylation and AMP-activated protein kinase (AMPK) signaling. There was significant hypophosphorylation of AMPK substrates in CD73-LKO livers ( $P < .0001$ ), while in isolated hepatocytes treated with AMP, soluble CD73 induced AMPK activation ( $P < .001$ ).

**CONCLUSIONS:** Hepatocyte CD73 supports long-term metabolic liver homeostasis through AMPK in a sex-dependent manner. These findings have implications for human liver diseases marked by CD73 dysregulation. (*Cell Mol Gastroenterol Hepatol* 2021;12:141–157; <https://doi.org/10.1016/j.jcmgh.2021.01.016>)

**Keywords:** Adenosine; Inflammation; AMP-Activated Protein Kinase; Ecto-5'-Nucleotidase; Hepatic Zonation.

The highly integrated metabolic activities of hepatocytes control physiological homeostasis and are perturbed in chronic liver diseases. Nonalcoholic fatty liver disease (NAFLD) is the most prevalent chronic liver disease in children and adults, and a major risk factor for the development of cirrhosis and hepatocellular carcinoma (HCC).<sup>1,2</sup> A lack of approved therapies renders NAFLD a major health problem, estimated to affect 25% of the global population, including 80 million people in the United States.<sup>3</sup> In recognition that NAFLD is principally a metabolic disease, a new nomenclature was proposed: metabolic-associated fatty liver disease.<sup>4</sup> From that standpoint, understanding how hepatocytes maintain long-term energy homeostasis will be critical for addressing the key mechanisms behind this complex and heterogeneous disorder.

It was shown more than 50 years ago that adenosine triphosphate (ATP) injections increase hepatic ATP content and can prevent the development of fatty liver in rodents.<sup>5</sup> Moreover, patients with nonalcoholic steatohepatitis, the most severe form of NAFLD, have significantly impaired capacity for replenishing hepatic ATP stores after transient depletion.<sup>6</sup> Numerous biochemical reactions control ATP metabolism and utilization by hepatocytes, including the enzymatic hydrolysis of extracellular ATP to form adenosine monophosphate (AMP), which is metabolized further to adenosine by ecto-5'-nucleotidase (CD73).<sup>7</sup> Adenosine regulates many physiological responses via activation of adenosine receptors,<sup>8</sup> and it also can be taken up inside the cell via specific transporters<sup>9</sup> and phosphorylated back to AMP by adenosine kinase.<sup>10</sup> A major function of adenosine is to reduce metabolic demand and conserve energy, but it is not known what role CD73 plays in this process, despite being the major extracellular adenosine regulator across different tissues and cell types.<sup>11,12</sup> Insights into CD73 mechanisms have particular relevance to liver biology and disease because the messenger RNA (mRNA) expression and enzymatic activity of CD73 are down-regulated significantly in human NAFLD, cirrhosis, and HCC via transcriptional,<sup>13</sup> post-transcriptional,<sup>14</sup> and post-translational<sup>15</sup> mechanisms. The latter 2 mechanisms involve production of a catalytically deficient splice variant (CD73S) and catalytically impaired high-mannose glycosylation variant.<sup>12</sup>

Although CD73 has ubiquitous expression,<sup>12</sup> recent single-cell profiling experiments have shown that its mRNA is zonally distributed in epithelial tissues, including the liver,<sup>16</sup> intestine,<sup>17</sup> and kidney.<sup>18</sup> CD73 is induced transcriptionally by hypoxia<sup>19</sup> and CD73-generated adenosine directly protects multiple epithelial tissues, including the liver, against hypoxic injury.<sup>20–22</sup> Previous studies have shown an allostatic function for CD73 in response to severe oxygen deprivation, but presently it is not known if CD73 functions in the long-term maintenance of liver homeostasis. Addressing this question is important because oxygen tension across the hepatic lobule is a key determinant of physiological metabolic zonation.<sup>23</sup> Metabolic zonation

refers to the heterogeneous distribution of enzyme activities, resulting in periportal predominance of protein secretion, cholesterol synthesis, fatty acid oxidation, and ureagenesis, and pericentral predominance of glycolysis, lipogenesis, and bile acid synthesis. This structured division of labor among healthy hepatocytes is disrupted in chronic liver diseases such as NAFLD.<sup>24</sup>


Given the earlier-described observations, and the known functions of adenosine in regulating hepatic glucose and lipid metabolism,<sup>25</sup> we hypothesized that CD73 has nonredundant homeostatic functions in the liver. To test this hypothesis, we generated mice with a targeted deletion of the CD73-encoding gene *Nt5e* in hepatocytes and characterized their liver phenotypes using multiple approaches. Our findings showed unanticipated age- and sex-dependent functions of CD73 in the long-term maintenance of hepatocyte metabolism and liver homeostasis. As such, these results add a cellular-level understanding of this key enzyme, in particular its relatively underappreciated functions in epithelial tissues. Importantly, these findings have translational implications for human liver diseases, as well as anti-CD73 antibodies and inhibitors, which currently are undergoing clinical development for cancer<sup>26</sup> and COVID-19 immunotherapy.<sup>27</sup>

## Results

### *Pericentral Hepatocytes Account for the Bulk of CD73 Expressed in the Normal Mouse Liver*

Using single-cell RNA transcriptomic data from the Tabula Muris project,<sup>28</sup> we compared *Nt5e* expression across different cell types in the mouse liver. Both fluorescence-activated cell-sorted and microfluidic droplet-captured cells showed that hepatocytes are the primary cell types expressing *Nt5e* (Figure 1A–D). Approximately 30% of hepatocytes, 7% of leukocytes and natural killer cells, and <2% of Kupffer cells and liver sinusoidal endothelial cells express *Nt5e*, while cholangiocytes and B cells lack *Nt5e* expression (Figure 1A–D). To determine if *Nt5e* presence correlates with CD73 protein, we performed coimmunofluorescence staining for CD73 and markers of hepatocytes, cholangiocytes, endothelial cells, and Kupffer cells (Figure 1E). In agreement with the

**Abbreviations used in this paper:** A<sub>2A</sub>R, G<sub>s</sub>-protein-coupled adenosine receptor 2A; AMP, adenosine monophosphate; AMPase, adenosine monophosphatase; AMPK, adenosine monophosphate-activated protein kinase; APCP, adenosine 5'-( $\alpha,\beta$ -methylene)diphosphate; ATP, adenosine triphosphate; CD73, ecto-5'-nucleotidase; CD73-LKO, liver-specific CD73 knockout; CoA, coenzyme A; HCC, hepatocellular carcinoma; HFD, high-fat diet; K, keratin; mTOR, mechanistic target of rapamycin; NAFLD, nonalcoholic fatty liver disease; PBS, phosphate-buffered saline; PCR, polymerase chain reaction; TBST, tris-buffered saline with 0.1% Tween 20; WT, wild-type; loxP, locus of cross-over P1; GPI, glycosylphosphatidylinositol.

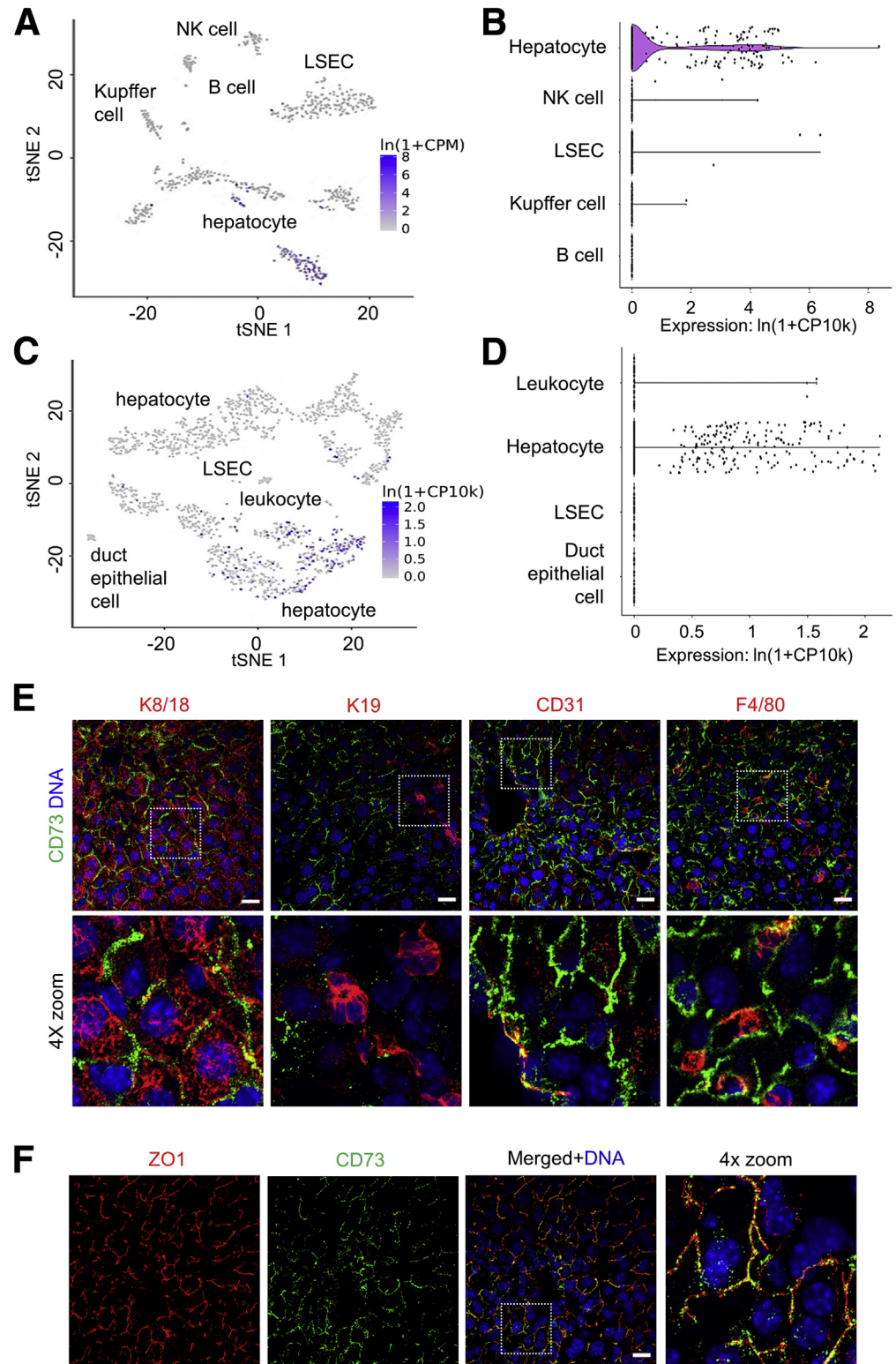
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**Figure 1. CD73 is expressed primarily on hepatocytes in normal mouse liver.** t-distributed stochastic neighbor embedding (tSNE) and violin plots showing RNA sequencing analysis of the mouse liver from (A and B) fluorescence-activated cell sorter (FACS)-sorted and (C and D) microfluidic droplet captured cells showing the highest expression of the mouse CD73-encoding gene *Nt5e* in hepatocytes. Data were obtained from Tabula Muris. (E) Fresh-frozen liver sections were stained with antibodies against CD73 (green), and the cell-specific markers keratins K8/18 (hepatocyte), K19 (cholangiocyte), CD31 (endothelium), and F4/80 (Kupffer cells) in red. *Bottom row*: Magnified views of the boxed areas in the top row. Blue, 4',6-diamidino-2-phenylindole (DAPI)-stained nuclei (DNA). (F) Immunofluorescence staining for CD73 (green) in frozen liver sections showing colocalization with the bile-canalicular marker zonula occludens 1 (ZO1) (red). *Right*: 4× magnified view of the boxed area in the merged panel. DAPI-stained nuclei. Blue, DAPI-stained nuclei (DNA). Scale bars: 20 μm. NK, natural killer; LSEC, liver sinusoidal endothelial cell.



transcriptomic data, CD73 is present on hepatocytes, but absent from cholangiocytes, which are marked by keratin (K) K8/K18 and K19 staining, respectively (Figure 1E). In addition, subsets of endothelial cells, expressing CD31, and Kupffer cells, expressing F4/80,

co-stained for CD73 (Figure 1E). However, the most abundant expression of CD73 was detected on the bile canalicular membranes of hepatocytes, as shown by co-staining with the tight junction marker protein zonula occludens 1 (Figure 1F).



### Liver-Specific CD73 Knockout Mice Develop Normally and Do Not Show Major Liver Abnormalities at a Young Age

To address the function of CD73 in the mammalian liver, we generated mice with a targeted deletion of the *Nt5e* gene in hepatocytes. The critical exon 2 of *Nt5e* was flanked by loxP sites and deleted in the presence of Cre recombinase driven by the albumin (*Alb*) promoter. As expected, *Nt5e* was selectively targeted in the liver (Figure 2A) and CD73 protein was absent from primary hepatocytes isolated from the CD73-LKO mice (Figure 2B). Immunoblotting of total liver lysates showed that CD73 levels were reduced by 75% and 67% in male and female CD73-LKO mice, respectively, compared with age-matched wild-type (WT) mice (Figure 2C and D). The expression of a functionally related GPI-anchored protein, tissue nonspecific alkaline phosphatase, was unaffected (Figure 2C). Immunofluorescence analysis of liver tissues showed the absence of CD73 from the apical membrane of hepatocytes in CD73-LKO mice (Figure 2E), and this corresponded to significantly diminished *ecto*-adenosine monophosphatase (AMPase) activity in the pericentral region of CD73-LKO livers (Figure 2F). Therefore, the CD73-LKO mice lack hepatocyte CD73 expression and are deficient in CD73-associated *ecto*-AMPase function.

To determine how this genetic manipulation affected liver function, we analyzed mice at different ages. Histologic examination of liver tissues from 7-week-old WT and CD73-LKO male mice did not show major differences (Figure 3A). In agreement with previous observations using constitutive *Nt5e*-knockout mice,<sup>22</sup> the CD73-LKO mice did not show major defects in viability, growth, and development up to the mature adult stage at 21 weeks (Figure 3B). Both male and female CD73-LKO mice gained weight at the same rate (Figure 3B) and had similar liver weight-to-body weight ratios compared with their WT counterparts (Figure 3C). Taken together, these data show that hepatocyte CD73 is not required for normal mouse liver development and maturation.

### CD73-LKO Mice Develop Spontaneous Hepatocyte Degeneration in a Sex- and Age-Dependent Manner

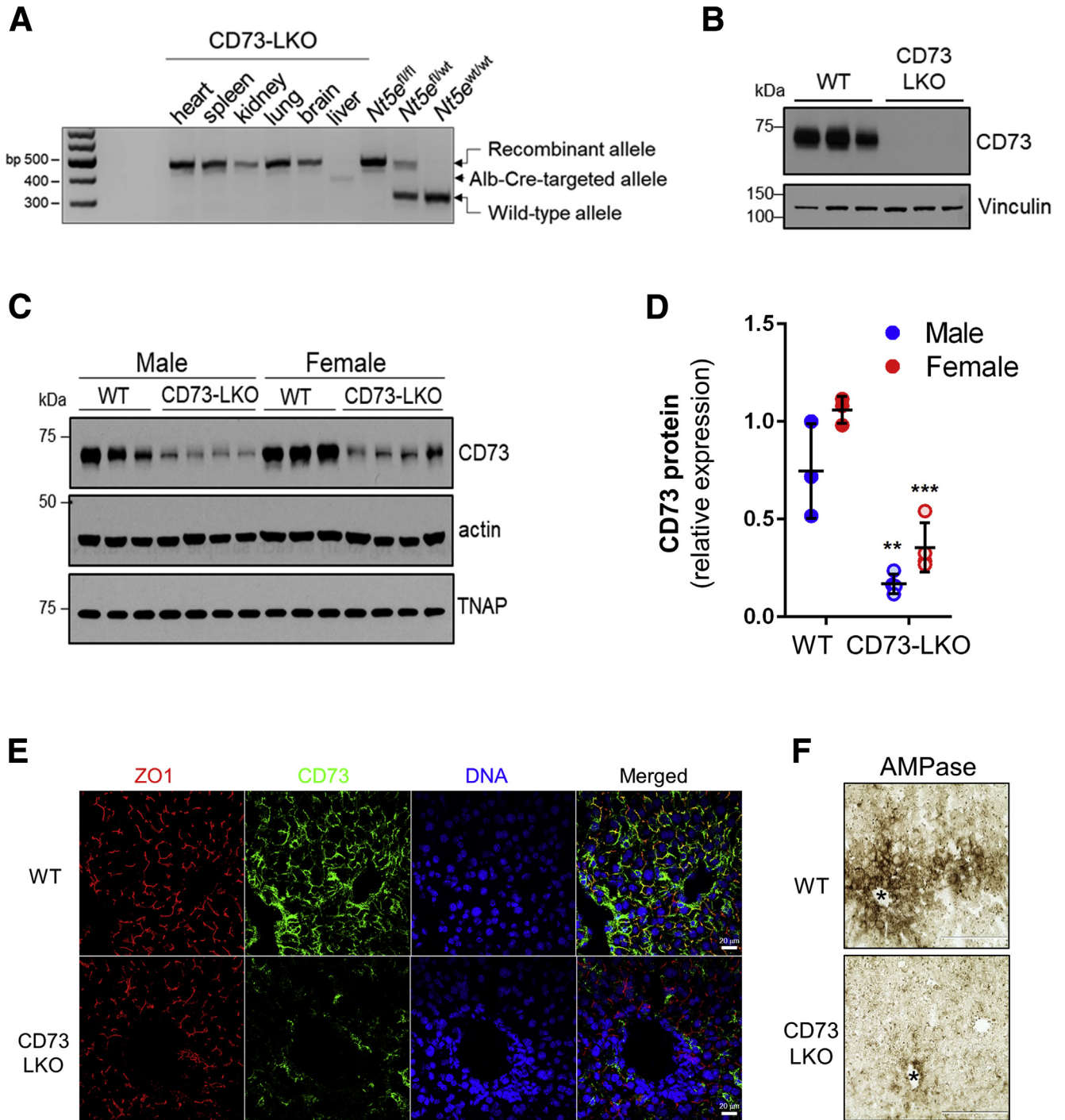
The zonal expression pattern of CD73 on pericentral hepatocytes prompted us to investigate the long-term consequences of CD73 deficiency because pericentral hepatocytes are important for the homeostatic renewal in the mouse liver.<sup>29,30</sup> Therefore, we studied the liver phenotypes of standard chow-fed WT and CD73-LKO mice between 22 and 42 weeks (5–10 mo) of age, which represents the period between the mature adult and middle-aged stages. Serum analysis showed a trend ( $P < .08$ ) toward increased alanine aminotransferase levels in male mice, but not in female mice ( $P < .3$ ) at 21–22 weeks of age (Figure 4A). Histologically, we

observed significant pericentral hepatocyte injury in 22-week-old male CD73-LKO mice, which was not present in the female mice, or the corresponding age- and sex-matched WT controls (Figure 4B). The hepatocyte injury and tissue damage were more extensive in the older (age, 42 wk) compared with the younger (age, 22 wk) male mice (Figure 4C), pointing to a progressive and age-dependent liver injury. Blinded quantitative histologic analysis further unmasked male-predominant hepatocellular damage ( $P < .05$ ), which was defined by swelling and ballooning hepatocyte degeneration (Figure 4D). Another notable sex difference was that hepatocytes in female CD73-LKO appeared smaller (Figure 4B), which corresponded with decreased serum albumin levels (Figure 4E), whereas serum albumin was unaffected in the male CD73-LKO mice (Figure 4E). Furthermore, we observed a significant increase in serum blood urea nitrogen levels in the male, but not female, CD73-LKO mice (Figure 4F).

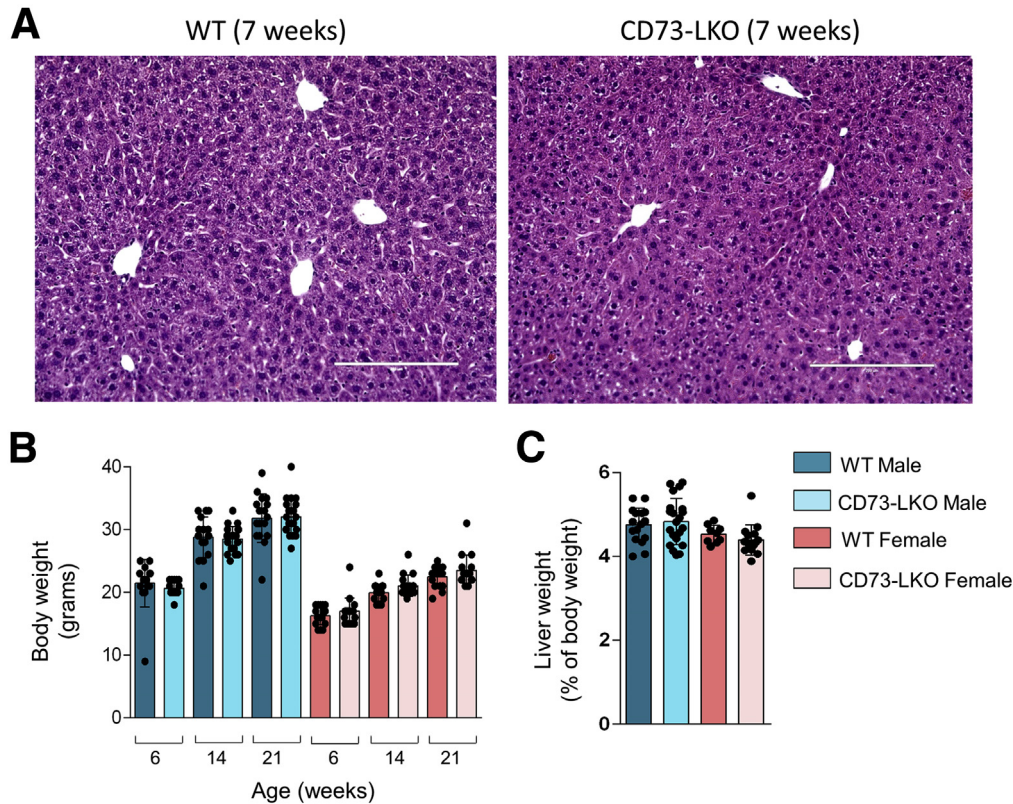
### Deficiency of Hepatocyte CD73 Leads to AMP-Activated Protein Kinase Hypoactivation and Perturbs Metabolic Homeostasis

To understand the cellular basis for the male-predominant histopathologic changes, we performed proteomic profiling on freshly isolated hepatocytes from 5-month-old male WT and CD73-LKO mice (Figure 5A). This analysis showed up-regulation in protein synthesis pathways associated with the transcriptional regulator eukaryotic initiation factor 2 (EIF2), apoptosis, and nucleotide excision repair. Pathways involved in fatty acid oxidation (peroxisome proliferator-activated receptor  $\alpha$ /retinoid X receptor  $\alpha$ , PPAR $\alpha$ /RXR $\alpha$ ), cholesterol, and stearate biosynthesis also were up-regulated (Figure 5A). On the other hand, hormone signaling (estrogen and aldosterone), amino acid metabolism (transfer RNA charging), nutrient sensing (mechanistic target of rapamycin, mTOR; p70S6K), and overall cellular energy homeostasis (oxidative phosphorylation and AMP-activated protein kinase [AMPK]) pathways were down-regulated in the CD73-null hepatocytes (Figure 5A). Consistent with abnormal lipid metabolism, as suggested by the proteomic analysis, male CD73-LKO mice developed significantly more microvesicular and macrovesicular steatosis, compared with female mice (Figure 5B and C). We profiled a set of lipid metabolism genes to determine if transcriptional compensation may account for the sex differences we observed. Although most genes remained unchanged, the gene *Hmgcs1*, which encodes hydroxymethylglutaryl-coenzyme A synthase, was significantly higher in female WT mice and further induced in the CD73-LKO mice (Figure 5D). Hydroxymethylglutaryl-CoA synthase converts acetyl-CoA to hydroxymethylglutaryl-CoA, which is used in the mevalonate pathway.

To further examine the link between CD73 activity and hepatic lipid accumulation, we treated WT hepatocytes



**Figure 2. Generation of liver-specific CD73-LKO mice.** (A) PCR analysis of *Nt5e* generated a 349-bp fragment from the Cre-targeted allele in the CD73-LKO mouse liver. *Nt5e<sup>(wt/wt)</sup>*, *Nt5e<sup>(fl/wt)</sup>*, and *Nt5e<sup>(fl/fl)</sup>* mice are controls. Representative immunoblots of CD73 in (B) primary hepatocytes, and (C) in total liver lysates from male and female WT and CD73-LKO mice. Actin and tissue nonspecific alkaline phosphatase (TNAP) immunoblots serve as controls. (D) Semiquantitative analysis of CD73 protein expression based on immunoblot band intensities in panel C. \*\**P* < .01, \*\*\**P* < .001, 2-way analysis of variance. Error bars represent SD. (E) Immunofluorescence staining for CD73 (green), tight junction protein zonula occludens 1 (ZO1) (red), and 4',6-diamidino-2-phenylindole (DAPI)-stained DNA (blue) on frozen liver tissue sections from WT (top) and CD73-LKO (bottom) mice. Scale bar: 20  $\mu$ m. (F) AMPase activity in WT and CD73-LKO mice using formalin-fixed liver tissue sections. The brown deposits indicate ecto-AMPase activity in the presence of AMP. Stars indicate the central vein. Scale bar: 400  $\mu$ m.



**Figure 3. CD73-LKO mice develop normally and do not show major liver abnormalities up to 21 weeks of age.** (A) Formalin-fixed liver sections of 7-week-old male mice stained with H&E. Scale bar: 200  $\mu$ m. (B) Comparison of body weight from WT and CD73-LKO mice at 6–21 weeks of age. n = 15–21 mice/group. (C) Analysis of liver-to-body weight ratios of 21-week-old mice. Males: n = 18 WT, n = 21 CD73-LKO. Females: n = 9 WT, n = 15 CD73-LKO. Error bars represent SD.

with the CD73 inhibitor adenosine 5'-( $\alpha,\beta$ -methylene) diphosphate (APCP) in the absence and presence of oleic acid and monitored neutral lipid accumulation using a fluorescence imaging method. Treatment with APCP alone led to significant accumulation of lipid droplets in hepatocytes (Figure 6A and B). Co-treatment of hepatocytes with oleic acid and APCP together led to significantly more accumulation of lipid droplets than either treatment alone (Figure 6A), suggesting that CD73 enzymatic activity regulates hepatocyte lipid metabolism. To determine if CD73 enzymatic activity is altered during hepatic steatosis, we fed mice a high-fat diet (HFD) for 14 weeks and assessed CD73 ecto-AMPase activity using enzyme histochemistry. As shown in Figure 6C, hepatic steatosis leads to a dramatic decrease in CD73 activity in vivo. Together with the results in isolated hepatocytes, this suggests that CD73 enzymatic activity and hepatic lipid metabolism are closely linked.

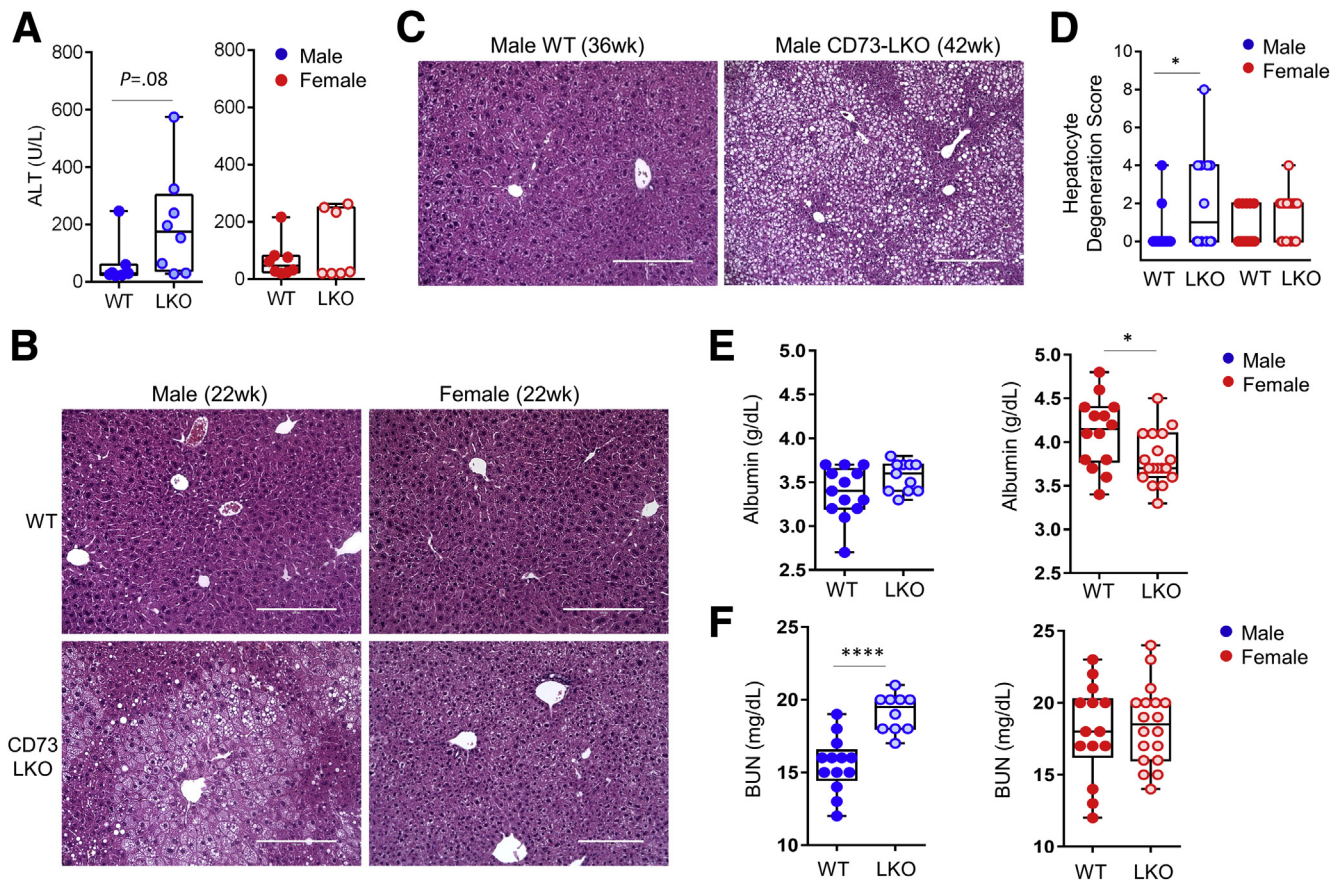
The proteomic profiling showed that AMPK, which is an ATP sensor and master metabolic regulator of lipid and glucose homeostasis, was down-regulated in CD73-LKO hepatocytes. To determine if the metabolic dysregulation in the absence of CD73 was linked to altered AMPK signaling, we probed for AMPK substrate phosphorylation using a specific phospho-motif antibody targeting LXRXX [pS/pT].<sup>31</sup> Despite equal expression of total AMPK $\alpha$ , male CD73-LKO mice had a dramatic reduction in the phosphorylation of AMPK substrates (Figure 7A and B). On the other hand, baseline AMPK substrate phosphorylation was lower in WT female mice and

was not changed significantly by CD73 deletion. The down-regulation of AMPK substrate phosphorylation was not related to major changes in the abundance or subcellular distribution of the kinase (Figure 7C), leading us to hypothesize that adenosine generated by CD73 promotes AMPK activity. In support of this hypothesis, extracellular addition of AMP significantly, and dose-dependently, induced AMPK activation (phosphorylation at Thr-172) in primary hepatocytes co-treated with soluble, enzymatically active CD73 (Figure 7D and E). To determine if this effect was dependent on the uptake of extracellular adenosine, we tested AMPK activation in the absence and presence of nitrobenzylthioinosine, an inhibitor of the equilibrative nucleoside transporter 1 adenosine re-uptake transporter.<sup>32</sup> Although AMPK phosphorylation appeared blunted in the presence of nitrobenzylthioinosine, this was not statistically significant (Figure 7F–G), suggesting multiple potential mechanisms. Combined, these data show that CD73, in the presence of extracellular AMP, activates AMPK in hepatocytes potentially via transport-dependent and independent mechanisms.

### *Mature Adult CD73-LKO Mice Develop Liver Inflammation, With a Male Predominance in Severity*

Because CD73 and adenosine play a major role in immune suppression, we next asked whether CD73-LKO mice develop hepatic inflammation. Blinded histologic analysis showed the presence of portal and lobular inflammation





**Figure 4. CD73-LKO mice develop spontaneous liver injury after 21 weeks of age in a sex-dependent manner.** (A) Serum analysis of alanine aminotransferase (ALT) in 21- to 22-week-old male (blue) and female (red) WT (filled circles) and CD73-LKO mice (open circles). Males:  $n = 8$  WT;  $n = 8$  CD73-LKO;  $P = .08$  (unpaired  $t$  test); females:  $n = 7$  WT;  $n = 7$  CD73-LKO.  $P = .3$  (unpaired  $t$  test). (B) Representative H&E images of formalin-fixed liver sections from 22-week-old male and female WT and CD73-LKO mice. Scale bars: 200  $\mu\text{m}$ . (C) Representative H&E images of formalin-fixed liver sections from 36- to 42-week-old male WT and CD73-LKO mice. Scale bars: 200  $\mu\text{m}$ . (D) Quantification of blinded histologic scoring of hepatocyte degeneration (defined as swelling and ballooning) in 21- to 42-week-old mice. Scoring: 0, none; 4, mild swelling/focal ballooning; 8, severe swelling/extensive ballooning (scores were averaged for animals with ballooning + swelling). \* $P < .05$ ; 1-way analysis of variance. Box-and-whisker plots of (E) albumin and (F) blood urea nitrogen (BUN) in male (blue) and female (red) WT (filled circles) and CD73-LKO mice (open circles). Same animals as in panel D. Males:  $n = 13$  WT;  $n = 12$  CD73-LKO; females:  $n = 14$  WT;  $n = 18$  CD73-LKO mice. \* $P < .05$ , \*\*\*\* $P < .0001$ ; unpaired  $t$  test.

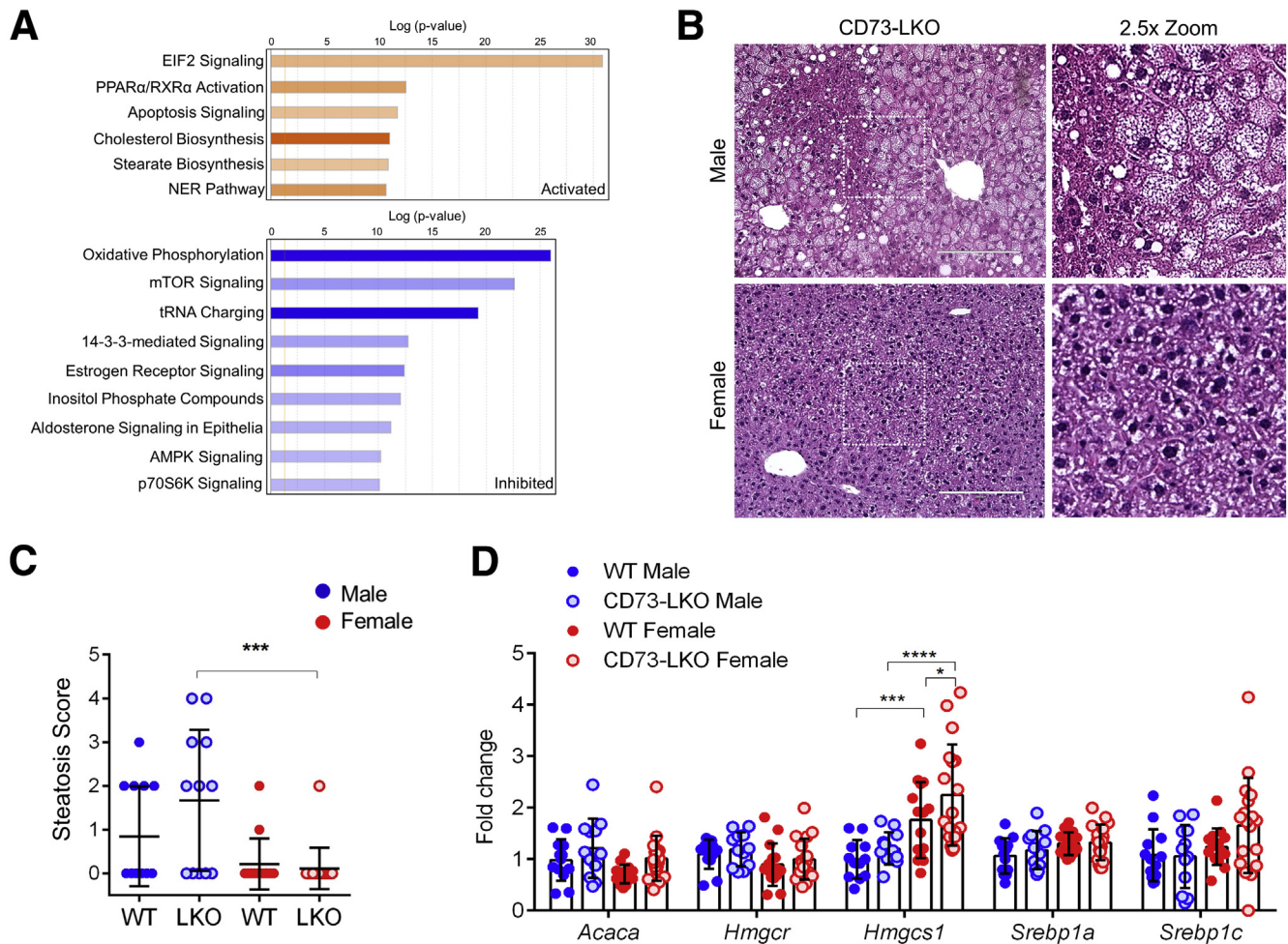
in both male and female CD73-LKO mice, but the male mice were affected more severely (Figure 8A and B). Consistent with the known roles of adenosine<sup>33</sup> and AMPK<sup>34</sup> as anti-inflammatory mediators, we observed induction in the proinflammatory cytokines interleukin 1 $\beta$  and tumor necrosis factor  $\alpha$  in the CD73-LKO livers, with the latter reaching statistical significance only in female mice (Figure 8C). Despite the similar transcriptional changes of proinflammatory genes, male mice showed significantly more neutrophil infiltration, as assessed by lymphocyte antigen 6 complex locus G6D (Ly6G) staining (Figure 8D and E). Profiling of CD73-associated adenosine pathway components, including the AMP substrate-generating enzyme CD39 (*Entpd1*) and adenosine receptors A1, A2A, A2B, and A3 (*Adora 1*, *2A*, *2B*, and *3*), showed female-specific compensatory induction of these targets in the absence of hepatocyte CD73 (Figure 8F). These results show that loss of hepatocyte

CD73 leads to spontaneous liver inflammation, and that in female mice this may be partly opposed by compensatory up-regulation of anti-inflammatory adenosine signaling mediators to limit tissue damage.

## Discussion

### Physiological Hepatoprotection of CD73 and Adenosine

Our study shows the importance of CD73, a major extracellular AMPase, for the long-term metabolic integrity of the mammalian liver. Previous seminal work using constitutive CD73 knockout mice established CD73 as a key factor in maintaining epithelial integrity via adenosine-mediated protection of the tissue barrier function, especially during hypoxia,<sup>19,22,35,36</sup> while our present findings provide new insights into the homeostatic mechanisms that are attributed specifically to CD73 on hepatocytes.



**Figure 5. Metabolic imbalance and increased steatosis in male CD73-LKO mouse hepatocytes and livers.** (A) Ingenuity pathway analysis of proteomic results showing activated (orange) and inhibited (blue) pathways in freshly isolated hepatocytes from male CD73-LKO mice relative to WT controls. (B) H&E staining of formalin-fixed liver sections. Magnified view of boxed areas in the left panel comparing the presence of steatosis in male and female CD73-LKO mice. Scale bar: 200  $\mu$ m. (C) Quantification of blinded histologic scoring of microvesicular and macrovesicular steatosis. Scoring: 0, none; 1, mild; 2, moderate; 3, severe (scores were averaged for animals with microsteatosis and macrosteatosis). Males: n = 13 WT; n = 12 CD73-LKO; females: n = 14 WT; n = 18 CD73-LKO. \*\*\* $P$  < .001, 1-way analysis of variance. (D) Gene expression analysis of lipid metabolism genes *Acaca*, *Hmgcr*, *Hmgcs1*, *Srebp1a*, and *Srebp1c* using total liver messenger RNA in male and female WT and CD73-LKO mice. n = 13 WT male, n = 12 LKO male, n = 14 WT female, n = 18 LKO female. \* $P$  < .05, \*\*\* $P$  < .001, \*\*\*\* $P$  < .0001, 2-way analysis of variance. Error bars represent SD. EIF2, eukaryotic initiation factor 2; NER, nucleotide excision repair; PPAR $\alpha$ /RXR $\alpha$ , peroxisome proliferator-activated receptor  $\alpha$ /retinoid X receptor  $\alpha$ ; tRNA, transfer RNA.

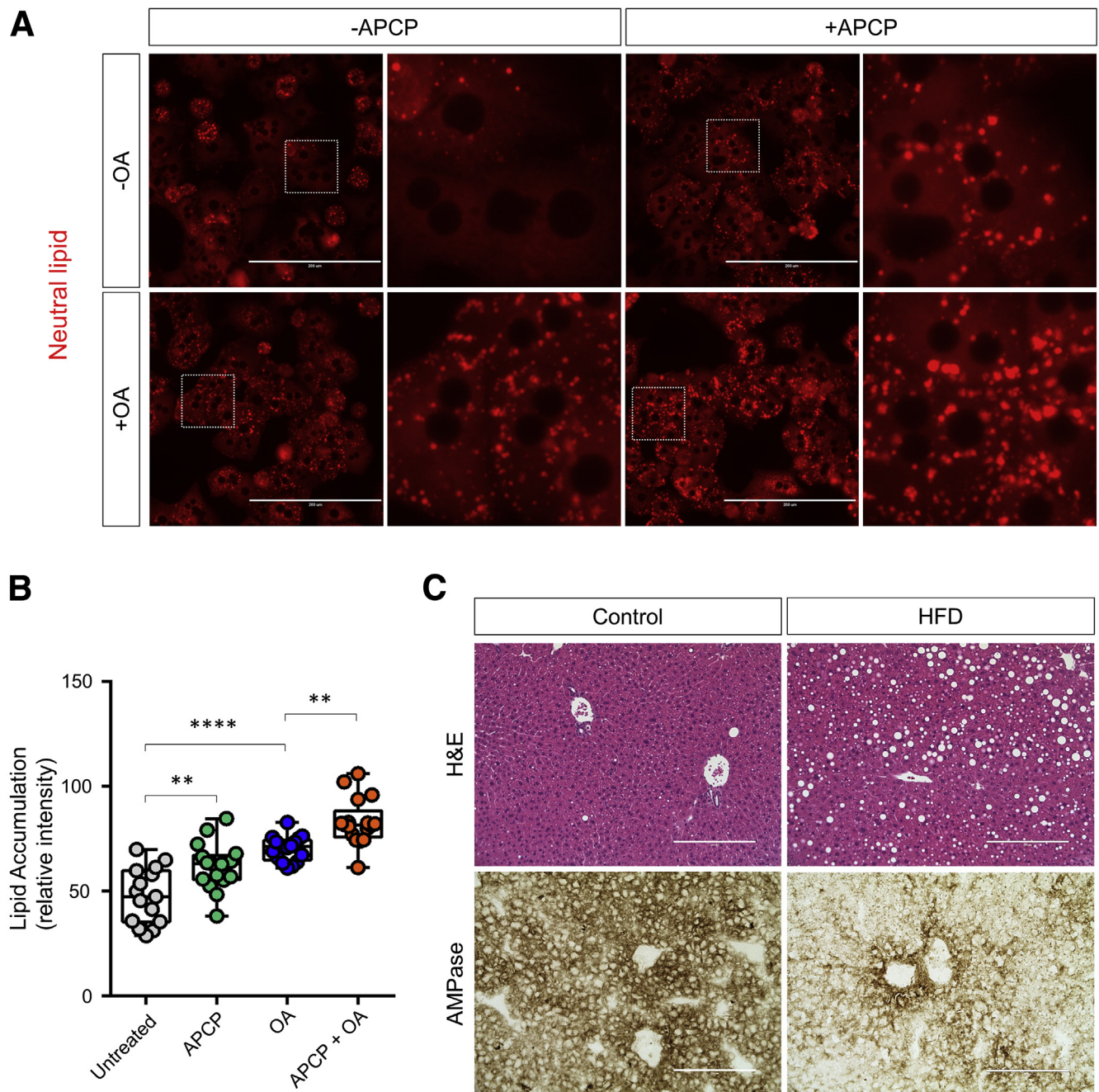
The male-predominant, spontaneous development of liver disease that we observed in CD73-LKO mice points to a physiological protective mechanism of CD73 that shows a sexual dimorphism. We conclude that AMPK hypoactivation, likely secondary to extracellular adenosine deficiency, contributes to hepatocellular injury, metabolic stress, and inflammation in male CD73-LKO mice. CD73 is down-regulated significantly under conditions of severe and persistent chronic stress in the mouse liver,<sup>13</sup> and in chronic human liver diseases of different etiologies.<sup>14,15</sup> Future studies addressing the detailed mechanisms of how CD73 supports metabolic homeostasis in hepatocytes throughout the lifespan, and in a sex-dependent manner, will have important implications for understanding mammalian liver biology as well as disease

development and progression. To that end, the *Nt5e<sup>fl/fl</sup>* and CD73-LKO mice represent tools for studying the cell-specific mechanisms of human liver diseases that are driven by perturbed metabolism and inflammation, such as NAFLD.<sup>37</sup>

### Sex Differences and Hormonal Regulation of the CD73-Adenosine Axis Across the Lifespan

At present, we do not fully understand the genetic and metabolic factors contributing to the sex differences. Our findings here highlight the importance for future work to consider both biological sex and age when examining the role of CD73-adenosine in hepatocyte biology and liver disease. Previous reports using whole-body knockout mice to address

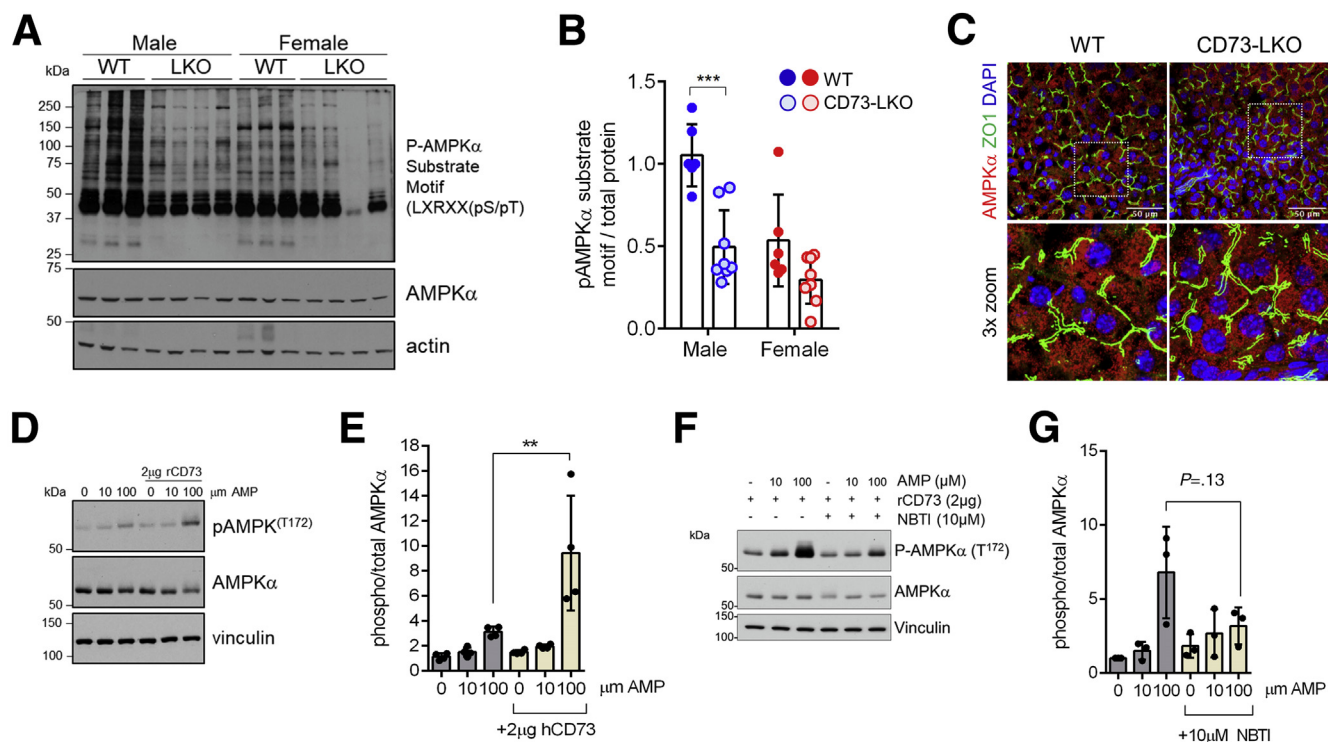




**Figure 6. Inhibition of CD73 enzymatic activity promotes lipid accumulation in hepatocytes.** (A) Isolated WT C57BL/6 male hepatocytes were treated for 24 hours with 1  $\mu\text{mol/L}$  oleic acid in the presence or absence of the CD73 inhibitor, APCP (10  $\mu\text{mol/L}$ ). Intracellular lipid accumulation was analyzed using LipidTOX neutral lipid stain (red). (B) Quantification of lipid accumulation in panel A.  $**P < .01$ ,  $****P < .0001$ ; 1-way analysis of variance. Error bars represent SD. (C) WT mice were fed normal chow (Control) or a HFD for 14 weeks. *Top panels:* Representative H&E images of Control- and HFD-fed mice, showing the development of steatosis in the HFD condition. *Bottom panels:* Representative enzyme histochemistry images showing a decrease in CD73 AMPase activity in the HFD condition. Scale bars: 200  $\mu\text{m}$ .

the role of CD73 and adenosine in hepatic steatosis<sup>38</sup> and fibrosis<sup>39</sup> examined only young (age, 6–8 wk) male mice. We found that 7-week-old CD73-LKO mice are indistinguishable from their WT littermates, regardless of sex, whereas CD73 loss is detrimental at older ages, but that female mice are relatively protected. Our study shows sex-dependent liver injury resulting from CD73 deficiency, and the CD73-

adenosine axis has been linked recently to the protective action of female sex hormones in other organ systems and diseases. For example, estrogen-receptor signaling maintains CD73 expression and adenosine production by osteoprogenitors, while the loss of CD73-generated adenosine promotes postmenopausal bone loss in mice.<sup>40</sup> In addition, several studies have shown complex and region-specific modulation



**Figure 7. CD73-generated extracellular adenosine activates AMPK in hepatocytes and livers from CD73-LKO mice and show impaired AMPK signaling.** (A) Representative immunoblots of AMPK substrate phosphorylation in liver lysates from male and female mice. (B) Semiquantitative analysis of AMPK substrate phosphorylation relative to total protein based on immunoblots in panel A.  $***P < .0001$ , 2-way analysis of variance. Error bars represent SD. (C) Fresh-frozen liver sections were stained with antibodies against AMPK (red) and zonula occludens 1 (ZO1) (green) show similar AMPK distribution in WT and CD73-LKO mice. *Bottom panels:* Magnified view of the boxed areas. Scale bar: 50  $\mu$ m. (D) Isolated WT hepatocytes were treated with AMP and recombinant soluble CD73 (rCD73) at the indicated concentrations. Representative immunoblots of total and phospho-AMPK $\alpha$ . (E) Quantification of phosphorylated/total AMPK.  $n = 4$  replicates.  $**P < .001$ ; 2-way analysis of variance. (F) Representative immunoblot analysis showing AMPK phosphorylation in WT hepatocytes treated with AMP, rCD73, and the adenosine transport inhibitor nitrobenzylthioinosine (NBTI). (G) Quantification of phosphorylated/total AMPK in rCD73-treated hepatocytes, in the absence/presence of AMP and NBTI.  $n = 3$  replicates. 2-way analysis of variance. Error bars represent SD. DAPI, 4',6-diamidino-2-phenylindole.

of spontaneous adenosine production by CD73 in the central nervous system that is mediated by both estrogen receptors  $\alpha$  and  $\beta$ ,<sup>41-43</sup> accounting for sex differences underlying adenosine-dependent neuromodulation. Although the exact significance of increased Hmgcs1 in the female mice is not presently clear, it is possible that it serves a protective mechanism against lipid accumulation because deficiency of the closely related mitochondrial Hmgcs2 gene promotes mouse fatty liver pathology via ketogenic insufficiency.<sup>44</sup> Moreover, it was shown recently that, in regulatory T cells, perturbation of the mevalonate pathway leads to metabolic stress and a dramatic reduction in cell surface CD73 expression,<sup>45</sup> showing a link between these 2 pathways.

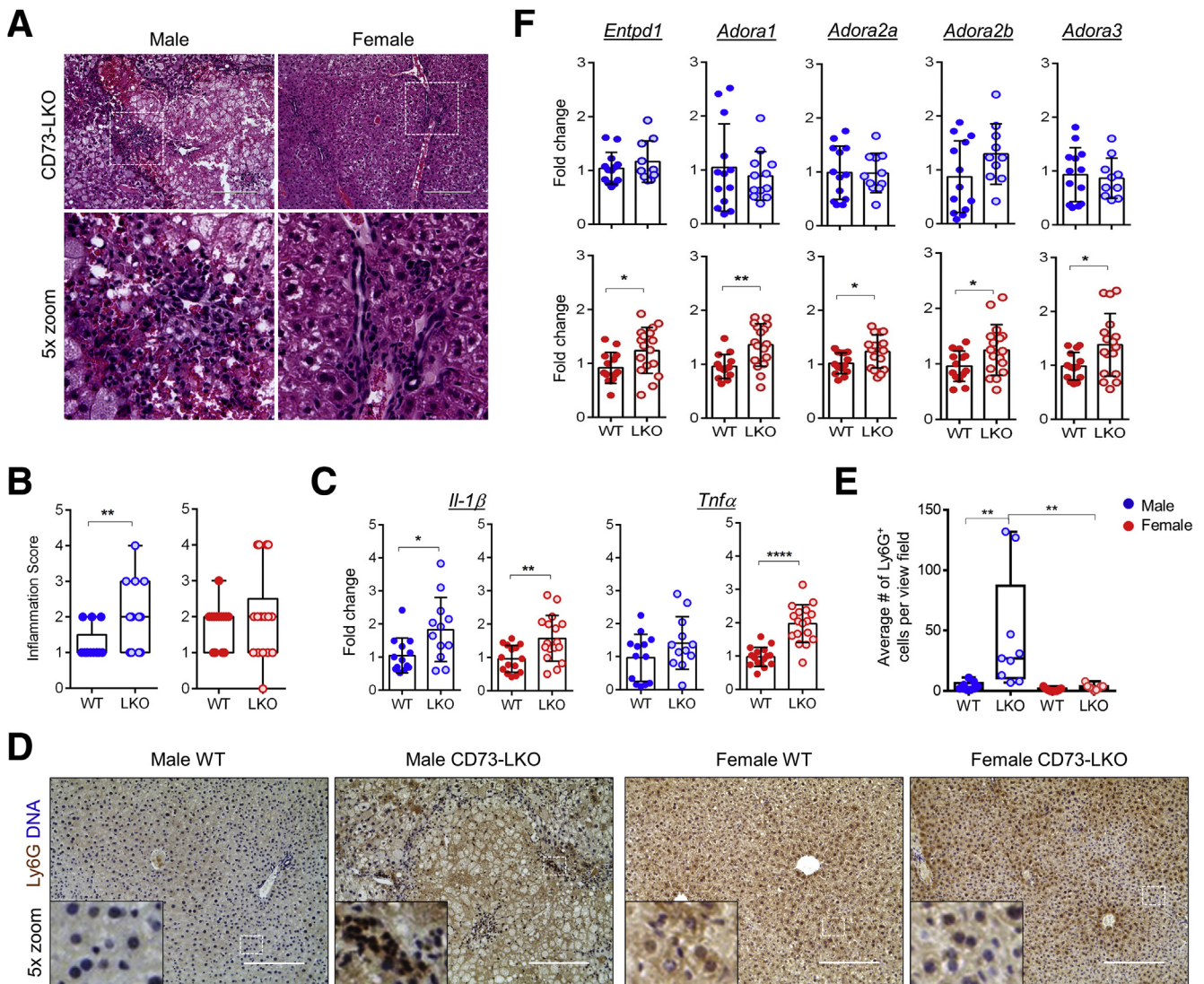
Biological sex and reproductive hormones exert a significant effect over individual variability to human disease development and progression.<sup>46</sup> This is especially evident in chronic liver disease, which currently ranks among the top 10 causes of death in men, but not in women.<sup>47</sup> Both male sex and older age are associated with worse survival and greater incidence of HCC among patients with biopsy-confirmed NAFLD.<sup>48</sup> However, the basic biological mechanisms of how age and biological sex affect patient susceptibility to disease development and progression have

been difficult to understand, especially in the case of NAFLD.<sup>49</sup> Recent studies in mice have shown that the genetic background, body fat distribution, plasma high-density lipoprotein, ceramide, and metabolic gene (eg, *Pklr*, *Lcn2*) levels underlie sex differences in hepatic steatosis.<sup>50-54</sup> An extensive analysis examined 50 metabolic and cardiovascular traits in 100 inbred mouse strains and showed significant sex differences at the molecular and physiological levels.<sup>54</sup> In general, sex hormones partially accounted for the metabolic traits, but the major conclusion from this study was that sex differences cannot be reconciled by examining single molecules or even pathways, given the multitude of variables that exert effects on the organism (eg, environment, age, genetic background) and their interactions. Therefore, the CD73-adenosine axis is one of many pathways that can influence sex differences in liver biology and disease.

### Cross-Talk Between CD73 and AMPK in Hepatocytes

CD73 is strongly induced in epithelial tissues under hypoxic conditions via hypoxia-inducible factor-1.<sup>19</sup> Rapid and sustained CD73 up-regulation during hepatic ischemia





**Figure 8. Loss of hepatocyte CD73 results in spontaneous liver inflammation.** (A) Representative H&E images of inflammatory lesions in livers from 22-week-old CD73-LKO mice. *Boxed areas in the top panels* are magnified in the *bottom panels*. (B) Quantification of blinded histologic analysis of portal and lobular inflammation (scores were averaged for animals with portal + lobular inflammation). Scoring: 0, none; 2, minimal; 4, mild; 6, moderate; 8, severe. (C) Total liver messenger RNA analysis of proinflammatory markers interleukin  $1\beta$  and tumor necrosis factor  $\alpha$  in male (blue) and female (red) mice aged 21–42 weeks. (D) Representative images of immunohistochemical-stained paraffin-embedded liver tissues for neutrophil marker Ly6G (brown). Harris hematoxylin-stained nuclei (blue). *Inset*: magnified view of the *boxed area* showing signal distribution. Scale bar: 200  $\mu\text{m}$ . (E) Quantification of Ly6G<sup>+</sup> cells per view field in panel D. (F) Gene expression analysis of ectonucleotidase CD39 (*Entpd1*) and adenosine receptors A1R, 2AR, 2BR, and 3R (*Adora1*, *2a*, *2b*, *3*) using total liver messenger RNA in male and female WT and LKO mice.  $n = 13$  WT male,  $n = 12$  LKO male,  $n = 14$  WT female,  $n = 18$  LKO female. \* $P < .05$ , \*\* $P < .01$ , \*\*\*\* $P < .0001$ , unpaired  $t$  test. Error bars represent SD.

confers adenosine-dependent hepatoprotection.<sup>21</sup> The latter is very similar to the reported hepatoprotective effect of AMPK during ischemic preconditioning,<sup>55</sup> although a functional link between these pathways has not been established. At 5 months of age, AMPK activity in mouse liver is augmented significantly by hypoxia, but this regulation is lost at older ages.<sup>56</sup> The age-dependent activation of AMPK by hypoxia in the liver coincides with the age of onset of spontaneous liver injury in the CD73-LKO

mice, which became apparent at 5 months in our study. Based on this, we propose that the detrimental effects stemming from CD73 deficiency in hepatocytes may be associated with impaired capacity of the mature liver to calibrate oxygen responses properly via AMPK. In vitro work in cell lines, and in vivo studies using lower organisms, have linked AMPK and hypoxia-inducible factor-1 cross-talk with cellular homeostasis and organismal longevity through their roles in

balancing intracellular reactive oxygen species.<sup>57,58</sup> The purinergic signaling pathway also plays important roles in oxidative stress responses in the liver and is disrupted in liver disease.<sup>59</sup> As a key regulator of purinergic signaling, and the major extracellular enzymatic source of the immunosuppressive and anti-inflammatory mediator adenosine, CD73 represents a promising target for metabolic inflammation, which is considered a major driver of chronic human diseases and associated comorbidities.<sup>39,60</sup>

### Anti-inflammatory Effects of Adenosine in the Liver

In various liver injury models, adenosine exerts protective anti-inflammatory effects through the activation of the G<sub>s</sub>-protein-coupled adenosine receptor 2A (A<sub>2A</sub>R) on hepatocytes and immune cells.<sup>35,61,62</sup> For example, genetic deletion or pharmacologic inhibition of the A<sub>2A</sub>R causes severe inflammation and liver damage after concanavalin A-induced liver injury in a mixed cohort of male and female mice.<sup>61</sup> Whole-body hypoxia treatment provided protection against concanavalin A-induced liver inflammation.<sup>62</sup> Although the detailed mechanism behind the hepatoprotective effects of hypoxia in liver inflammation are not clear, it is plausible that hypoxia-mediated CD73 induction, followed by increased extracellular adenosine production, may be important during immune-mediated liver damage, which remains to be examined in detail in future studies. CD73 protein was shown to be up-regulated in a HFD model of liver injury performed in male mice, in which the hepatic protein expression of CD39 and A<sub>2A</sub>R also were increased.<sup>35</sup> The up-regulation of these purinergic factors was likely a protective mechanism because the absence of A<sub>2A</sub>R led to more severe inflammation, steatosis, and impaired insulin sensitivity after HFD feeding.<sup>35</sup>

The hypoactivation of AMPK and liver injury in the CD73-LKO mice that we observed were similar to the findings from another study using a mixed cohort of older (age, 12–16 mo) male and female mice, which found that the absence of the A<sub>2A</sub>R receptor leads to hypoactivation of AMPK and metabolic dysregulation, specifically obesity, hyperglycemia, and glucose intolerance.<sup>63</sup> The relative protection that we observed in the female CD73-LKO mice in our study could be attributed to compensatory induction of adenosine receptors, in particular A<sub>2A</sub>R and A<sub>2B</sub>R. Paradoxically, our proteomic analysis showed that the mTOR pathway is decreased in CD73-LKO mice despite the AMPK hypoactivation. It is plausible that the decrease in mTOR activity as predicted by the proteomic analysis is secondary to nucleotide imbalance because it recently was shown that mTOR is inhibited by purine depletion (adenylate and its derivatives) irrespective of AMPK activation.<sup>64</sup> Specifically, genetic and pharmacologic blockade of de novo purine biosynthesis or purine salvage significantly decreased the activation of the key mTOR downstream targets S6K and 4EBP1. This inhibitory effect was rescued by exogenous purines and their intracellular conversion to adenylylated nucleotides.

Taken together, our study provides broad insight into potential novel and sex-dependent functions of CD73 in promoting hepatocyte metabolic homeostasis under physiological conditions. Future studies will illuminate deeper understanding of the mechanistic networks by which CD73 plays a role in the liver. Ultimately, our findings can improve target identification for drug discovery in chronic liver diseases.

## Materials and Methods

### Chemicals and Reagents

The following chemicals and reagents were purchased from Sigma-Aldrich (St. Louis, MO): bovine serum albumin (A2153), adenosine 5'-( $\alpha,\beta$ -methylene)diphosphate (M3763), insulin (I0516), glucagon (G2044), hydrocortisone (H0888), levamisole (196142), calcium chloride dihydrate (223506), glucose (G8270), Percoll (P1644), and ethylene glycol-bis( $\beta$ -aminoethyl ether)-*N,N,N',N'*-tetraacetic acid (E3889). Laemmli sample buffer (4 $\times$ , 1610747) was purchased from Bio-Rad Laboratories (Hercules, CA).

### Generation of Hepatocyte-Specific CD73 KO Mice

CD73 floxed (CD73<sup>fl/fl</sup>) mice were generated by Cyagen Bioscience, Inc (Santa Clara, CA). Briefly, exon 2 of the CD73-encoding gene, *Nt5e* (GenBank accession number: NM\_011851.4; Ensembl: ENSMUSG00000032420), was targeted to generate a frameshift mutation from downstream exons. Bacterial artificial chromosome (BAC) clone RP23-137M4 from the C57BL/6J library was the template for the targeting vector that contains a Neo cassette flanked by Frt sites, and *Nt5e* exon 2 region flanked by loxP sites. Linearized targeting vector was electroporated into embryonic stem cells, followed by neomycin selection of resistant clones. Targeted embryonic stem clones were injected into blastocysts and transferred into surrogate mothers to obtain chimera lines. The conditional knockout allele was obtained after flippase-mediated recombination. Heterozygote (flox/wt) mice were bred with each other to obtain flox/flox. For this study, CD73<sup>fl/fl</sup> mice were backcrossed to C57BL/6J WT mice 4 times. To generate the CD73 hepatocyte-specific knockout mice (CD73-LKO), CD73<sup>fl/fl</sup> mice were bred with Cre mice driven by the albumin promoter (stock 003574; The Jackson Laboratory).<sup>65</sup> Genotyping was performed at 7–10 days of age and the mice were weaned at 21 days. WT and CD73-LKO littermates were maintained on a normal chow diet or a 14-week high-fat diet (60% calories from fat [Bio-Serv, Flemington, NJ]; as noted in Figure 6 legend) in the pathogen-free animal facility at the University of North Carolina at Chapel Hill. All experiments were approved by the Institutional Animal Care and Use Committee and in accordance with the National Institutes of Health guidelines.

### Genotyping

Polymerase chain reaction (PCR) analysis using DNA extracted from toe clips and DreamTaq Green PCR Master



Mix 2× (K1081; Thermo Fisher, Waltham, MA) was performed to identify WT and CD73-LKO mice. Floxed PCR analysis used primers Floxed-Forward (5'-AGCA-CATTTAGTTTGAATCCC-3'), and Floxed-Reverse (5'-AAACAGACTTCTTGATTGGCAT-3'). To identify the presence of Cre recombinase, we used primers Fabpi-200R (5'-TAGAGCTTTGCCACATCACAGGTCATTCAG-3'), Fabpi-200F (5'-TGGACAGGACTGGACCTCTGCTTTCCTAGA-3'), Cre-281R (5'-TCGCCATCTTCCAGCAGG-3'), and Cre-281F (5'-CCATCTGCCACCAGCCAG-3'). Amplicons were analyzed using 2% agarose gel electrophoresis. The presence of a 426-bp and 286-bp bands indicate a floxed CD73 and WT allele, respectively. A 300-bp and 200-bp band is positive for Cre recombinase.

### Serum Analysis

Blood was collected by cardiac puncture from male and female WT and CD73-LKO mice between 5 and 9 months of age. Serum was analyzed for alanine aminotransferase, albumin, and blood urea nitrogen levels using the VetScan VS2 Analyzer and VetScan mammalian liver profile reagent rotor (Abaxis, Union City, CA).

### Immunoblot

Total liver protein lysates were extracted from 25 mg of liver tissue from the left lobe or from cultured primary WT hepatocytes using ice-cold lysis buffer (50 mmol/L n-octyl- $\beta$ -D-glucopyranoside, 1× protease inhibitors [05892970001; Roche, Basel, Switzerland], and 1× phosphatase inhibitors [04906837001; Roche] in 1× phosphate-buffered saline [PBS]). Protein lysates were resolved in 10% sodium dodecyl sulfate–polyacrylamide electrophoresis gels and transferred to a nitrocellulose membrane. After blocking in 5% milk in Tris-buffered saline with 0.1% Tween20 (TBST) for 1 hour at room temperature, the membranes were incubated with anti-CD73 (clone D7F9A, 13160), phospho-AMPK $\alpha$  (T172, clone 40H9, 2535), total AMPK $\alpha$  (clone D5A2, 5831), phospho-AMPK substrate motif (LXRXX [pS/pT], 5759S; Cell Signaling, Beverly, MA); pan actin (ACTN05, MA5-11869; Thermo Fisher), and antivinculin (clone hVIN-1, V9131; Sigma-Aldrich, St. Louis, MO) in 5% bovine serum albumin/TBST overnight at 4°C. Secondary antibodies (A4416, A6154, 1:5000–10,000; Sigma-Aldrich) were incubated for 1 hour at room temperature in 5% milk/TBST. Gels were stained with Coomassie blue, and membranes were stained with Ponceau S stain.

### Primary Hepatocyte Isolation and Treatment

Hepatocyte isolation was performed on 5-month-old CD73<sup>fl/fl</sup> mice weighing 20–35 g. Mice were anesthetized with Avertin (250–300 mg/kg, intraperitoneally) (T48402; Sigma-Aldrich) to achieve deep anesthesia. The portal vein was cannulated and the liver was perfused with 50 mL of 37°C sterile buffer I solution (0.50 mmol/L ethylene glycol-bis[ $\beta$ -aminoethyl ether]-N,N,N',N'-tetraacetic acid, 5.5 mmol/L glucose, 1% penicillin/streptomycin in calcium- and magnesium-free Hank's balanced salt solution) at a rate

of 7 mL/min. Liver digestion was performed with 40 mL of 37°C sterile buffer II solution (1.5 mmol/L CaCl<sub>2</sub>, 5.5 mmol/L glucose, 1% penicillin/streptomycin, and 3600 U of collagenase IV in calcium- and magnesium-free Hank's balanced salt solution) at a rate of 7 mL/min. Livers were surgically removed and hepatocytes were isolated in cold 1× Dulbecco's modified Eagle medium/penicillin/streptomycin media by size exclusion using 100- $\mu$ m and 70- $\mu$ m filters, respectively. Isolated hepatocytes were treated 24 hours after isolation with soluble human recombinant CD73 protein, AMP substrate, and a selective equilibrative nucleoside transporter 1 inhibitor, as specified in the figure legends. For intracellular lipid accumulation experiments, isolated WT C57BL/6 male hepatocytes were treated for 24 hours with 1  $\mu$ mol/L oleic acid in the presence or absence of the CD73 inhibitor, APCP (10  $\mu$ mol/L). Lipid accumulation was visualized using LipidTOX neutral lipid stain (H34477; Thermo Fisher). Fluorescence intensities were quantified in individual hepatocytes using Adobe Photoshop (San Jose, CA) software to measure the signal within individual hepatocytes after selecting equal areas of 20,000 pixels and plotted for each treatment group.

### RNA and Quantitative PCR

Total RNA was isolated from the left lobe of the liver and was extracted according to the manufacturer's protocol (RNeasy Mini Kit, 74104; Qiagen, Hilden, Germany). A total of 2  $\mu$ g RNA was converted to complementary DNA using the high-capacity complementary DNA reverse-transcription kit (4368813; Applied Biosystems, Foster City, CA). PCR was performed using SYBR Green PCR master mix (A25742; Applied Biosystems), and performed in a Quantstudio 6 Flex System (Applied Biosystems). The change in cycle threshold ( $\Delta$ CT) was normalized to 18S, and was expressed in fold change. The primer sequences were as follows: *Adora1* forward: 5'-TGTGACCACCACCCAGAGTA -3'; reverse: 5'-GCAGAGACTGGGACAAGGAG -3'; *Adora2a* forward: 5'-GAAGCAGATGGAGAGCCAAC -3'; reverse: 5'-GAGAGGATGATGGCCAGGTA -3'; *Adora2b* forward: 5'-CCTTTGGCATTG GATTGACT -3'; reverse: 5'-AAAATGCCCACGATCATAGC -3'; *Adora3* forward: 5'-TCATACCGGAAGGAATGAGC -3'; reverse: 5'-AGCTTGACCACCAGATGAC -3'; *Entpd1* forward: 5'-TAC CACCCATCTGGTCATT-3'; reverse: 5'-GGACGTTTTGTTTGGTTGGT -3'; *IL1 $\beta$*  forward: 5'-TCGCTCAGGGTCACAAGAAA-3'; reverse: 5'-CATCAGAGGCAAGGAGGAAAAC-3'; *TNF $\alpha$*  forward: 5'-AGGCTGCCCGACTACGT-3'; reverse: 5'-GAC TTTCTCTGGTATGAGATAGCAAA-3'; *Acaca* forward: 5'-AGC CAGAAGGGACAGTAGAA-3'; reverse: 5'-CTCAGCCAAGCGG ATGTA-3'; *Hmgcr* forward: 5'-CTTGTGGAATGCCTTGT GATTG-3'; reverse: 5'-AGCCGAAGCAGCATGAT-3'; *Hmgcs1* forward: 5'-GCCGTGAACTGGGTGCAA-3'; reverse: 5'-GCAT ATATAGCAATGTCTCCTGCAA-3'; *Srebp1a* forward: 5'-GGC CGAGATGTGCGAACT-3'; reverse: 5'-TTGTTGATGAGCTG GAGCATGT-3'; and *Srebp1c* forward: 5'-GGAGCCATG GATTGCACATT-3'; reverse: 5'-GGCCCGGAAGTCACTGT-3'; *18S* forward: 5'-ACCTGGTTGATCCTGCCAGTAG-3'; reverse: 5'-TTAATGAGCCATTTCGAGTTTC-3'.

### Liver Tissue Staining and CD73 Enzyme Histochemistry

Liver tissues from the left lobe were fixed with paraformaldehyde and paraffin-embedded, or flash-frozen using optimal cutting temperature (OCT) embedding medium. Paraffin-embedded tissues were cut in 10- $\mu$ m sections and stained with hematoxylin and eosin (H&E). For immunohistochemistry, tissue sections were deparaffinized and rehydrated. Antigen was retrieved using HIER buffer pH 6.0 (TA-135-HBL; Thermo Fisher), and endogenous peroxidase was inhibited by 3% hydrogen peroxide. Tissues were blocked with 10% normal goat serum, and the primary antibody for the neutrophil marker Ly6G (2557; Abcam, Cambridge, United Kingdom) was incubated overnight at 4°C in blocking buffer. Secondary biotinylated anti-rat IgG was incubated for 1 hour at room temperature, followed by incubation with tertiary antibody (ABC Elite, PK-6100; Vector, Burlington, Ontario, Canada). Tissue sections were developed using 3,3'-diaminobenzidine substrate (TA-125-QHDX; Thermo Fisher), and counterstained with Harris hematoxylin. Images were analyzed using a 20 $\times$  objective with the EVOS FL auto imaging system (Thermo Fisher). The average number of positive signals was quantified using ImageJ software (National Institutes of Health, Bethesda, MD) and plotted for each animal group. Frozen tissues were cut in 10- $\mu$ m sections and stained by immunofluorescence. Briefly, frozen tissue sections were fixed in cold 10% normal buffered formalin (032-059; Fisher Scientific, Ottawa, Ontario, Canada), followed by cold acetone (A19-1; Fisher Scientific) for 10 minutes each. Tissues were blocked in 2% normal goat serum/2.5% bovine serum albumin/1 $\times$  PBS for 1 hour at room temperature. Primary antibodies CD73 (clone TY/23, 550738; BD Pharmingen, San Jose, CA), K19-AF647 (ab192980; Abcam), K8/18 (clone 8592), and zonula occludens 1 (clone D6L1E, 13663; Cell Signaling) in blocking solution were incubated overnight at 4°C. Tissues were washed in 1 $\times$  PBS, before adding the secondary antibodies anti-rat 488 (a11006; Thermo Fisher) and anti-rabbit 568 (a11079; Thermo Fisher) for 1 hour at room temperature. The nuclei were stained using 4',6-diamidino-2-phenylindole (D1306; Invitrogen, Carlsbad, CA). Images were analyzed at 40 $\times$  oil objective using a Zeiss LSM 880 confocal microscope (Zeiss, Concord, NC). Frozen liver tissues sections of 10- $\mu$ m thickness were fixed in 10% neutral buffered formalin for 5 minutes at 4°C. Enzyme histochemistry was performed using our published protocol.<sup>13</sup> Images were analyzed with a 20 $\times$  objective using the EVOS-FL auto imaging system (Thermo Fisher).

### Proteomic Analysis by Liquid Chromatography With Tandem Mass Spectrometry

Freshly isolated primary hepatocytes from 3 WT and 3 CD73-LKO mice were lysed using 8 mol/L urea. Protein lysates were digested with LysC-protease and trypsin. Peptide samples were cleaned using Pierce C18 spin columns (Thermo Fisher), and then peptide quantitation was conducted. A total of 350 mg per sample was dried and labeled using the TMT 6-plex Isobaric Label Reagent Set

(90061; Thermo Fisher). Label efficiency was evaluated and found to be >99% for all TMT channels. Samples were mixed, then subjected to fractionation into 96 fractions by high-pH, reversed-phase liquid chromatography. Samples were concatenated into 24 samples. Each sample (~1 mg) was analyzed by liquid chromatography–tandem mass spectrometry using the QExactive HF (Thermo Fisher) for a total of 25 analyses. Proteins were identified and quantified using MaxQuant using a reviewed (~18,000 proteins) mouse database appended with a common contaminants database. Core analysis of the liquid chromatography–tandem mass spectrometry data set was analyzed using Ingenuity Pathway Analysis (Qiagen) and filtered based on a log ( $P = 1E^{-10}$ ) and an absolute z-score of 1.

### Statistical Analysis

Data were analyzed using an unpaired *t* test or 2-way analysis of variance built in GraphPad Prism (San Diego, CA). Data are presented relative to WT or untreated controls. Error bars from all graphs indicate the SD for  $n \geq 3$  samples or independent experiments. *P* values are denoted within each respective figure panel. Outliers were tested based on the Grubb's test ( $\alpha = .05$ ). The number of samples or independent experiments is indicated in the figure legends.

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