Hepatocyte mitochondrial DNA drives nonalcoholic steatohepatitis by activation of TLR9

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Nonalcoholic steatohepatitis (NASH) is the most common liver disease in industrialized countries. NASH is a progressive disease that can lead to cirrhosis, cancer, and death, and there are currently no approved therapies. The development of NASH in animal models requires intact TLR9, but how the TLR9 pathway is activated in NASH is not clear. Our objectives in this study were to identify NASH-associated ligands for TLR9, establish the cellular requirement for TLR9, and evaluate the role of obesity-induced changes in TLR9 pathway activation. We demonstrated that plasma from mice and patients with NASH contains high levels of mitochondrial DNA (mtDNA) and intact mitochondria and has the ability to activate TLR9. Most of the plasma mtDNA was contained in microparticles (MPs) of hepatocyte origin, and removal of these MPs from plasma resulted in a substantial decrease in TLR9 activation capacity. In mice, NASH development in response to a high-fat diet required TLR9 on lysozyme-expressing cells, and a clinically applicable TLR9 antagonist blocked the development of NASH when given prophylactically and therapeutically. These data demonstrate that activation of the TLR9 pathway provides a link between the key metabolic and inflammatory phenotypes in NASH.

Introduction

Hepatocyte steatosis and ballooning and inflammation of the liver constitute steatohepatitis, which occurs in response to a diverse range of stressors ranging from excess alcohol to drug-induced liver injury. Obesity-driven metabolic syndrome is the most common reason for steatohepatitis and is termed nonalcoholic steatohepatitis (NASH) (1). Overnutrition and obesity uniformly result in hepatic steatosis, which can progress to steatohepatitis, fibrosis, and cirrhosis. The biological processes promoting progression from steatosis to steatohepatitis are of great interest but are currently poorly understood.

TLR9 is an endosomal pattern recognition receptor (PRR) for which CpG-rich bacterial DNA and mammalian self-DNA are ligands (2). We have identified a requirement for TLR9 in toxic liver injury. For hepatocyte mtDNA to activate TLR9, whether a 12-week HFD model of NASH resulted in greater activation of a TLR9 reporter than did mtDNA from mice fed a chow diet (CD). This was confirmed in mouse Kupffer cells (KCs), which demonstrate upregulation of Tnfa transcripts in response to hepatocyte mtDNA from HFD-fed mice (Figure 1B).

Hepatocyte mtDNA is increased in NASH and is known to be proinflammatory (6). For hepatocyte mtDNA to activate TLR9, it would require release from hepatocytes and entry into TLR9-containing endosomes of cells capable of mounting an inflammatory response. Plasma extracellular mtDNA levels increase in tissue injuries in mice and humans including toxic liver injury (8). We tested whether a 12-week HFD model of NASH resulted in an increase in plasma DNA. There was an increase in total DNA and mtDNA, but not nDNA in plasma from HFD-fed compared with CD-fed mice (Supplemental Figure 1, A–C; supplemental material available online with this article; doi:10.1172/JCI83885DS1). Plasma from HFD-fed mice gave a small but significantly greater signal than did plasma from CD-fed mice from a TLR9 reporter cell line (Supplemental Figure 1D). To confirm whether these changes were present in humans, plasma from 3 previously well-characterized groups was analyzed: group 1 consisted of lean subjects without

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liver disease and normal alanine transaminase (ALT) levels; group 2 consisted of obese subjects with normal serum ALT levels; and group 3 consisted of obese subjects with elevated ALT levels (Table 1 and ref. 9). We used ALT as a marker of ongoing liver injury. The human data mirrored the mouse HFD model data, with group 3 subjects having increased plasma levels of total DNA and mtDNA, but not nDNA compared with groups 1 and 2 (Figure 1, C–E). In addition, plasma from group 3 subjects produced a stronger signal from the TLR9 reporter cell line (Figure 1F). These data show an increase in plasma mtDNA and greater TLR9 ligand activity in HFD-induced NASH.

Plasma mtDNA could be present in solution, as intact free mitochondria, or as mitochondria enclosed within extracellular microparticles (MPs) (10). Of relevance, in a choline deficiency model of NASH, there is a dramatic increase in MPs of hepatocytic origin in plasma, with hepatocyte MPs visible in the space of Disse (11). To identify whether intact mitochondria were present in plasma from our subjects with NASH and whether they were free or within MPs, we performed flow cytometric analysis of plasma from subjects in each of the 3 groups. Mitochondria were readily detectable in plasma by flow cytometry after staining with MitoTracker Deep Red, which stains mitochondria with an intact membrane potential, and MPs were identified by a combination of size gating (0.2–1.0 μm) and positivity for the plasma membrane dye PKH67 green. Dual staining with MitoTracker Deep Red and PKH67 allows the identification of mitochondria enclosed within a plasma membrane (Figure 1G, upper right quadrant). Obese patients with elevated serum ALT levels had a greater percentage of mitochondria staining for oxDNA, and the cellular origin of these free mitochondria is yet to be determined. The hepatocyte origin of plasma MPs containing mitochondria with increased oxDNA is consistent with the findings of increased oxidation of hepatocyte DNA in NASH (6). Finally, we confirmed that MPs were responsible for the ability of plasma to activate a TLR9 reporter cell line, as the majority of this activity was lost in MP-free plasma (Figure 1L).

TLR9 ligands can preferentially activate 2 downstream signaling pathways, resulting in differing degrees of production of NF-κB-dependent proinflammatory cytokines and IRF7-dependent type 1 IFN (13). The IRF7-dependent pathway can upregulate the IL-1 receptor antagonist (IL-1RA), which has antiinflammatory effects in the liver (14). To test whether mtDNA from hepatocytes of HFD-fed mice can activate type 1 IFN and IL-1RA, primary murine macrophages were stimulated with nDNA and mtDNA from hepatocytes of HFD-fed and CD-fed mice, and mRNA levels of Ifna, Ifnb, and Il1ra were quantified (Supplemental Figure 2, A–C). In contrast to the upregulation of TNF-α, hepatocyte mtDNA from HFD-fed mice, when compared with mtDNA from CD-fed mice, resulted in no change in Ifna, Ifnb, or Il1ra levels (15). This demonstrates that mtDNA from hepatocytes of HFD-fed mice results in the selective upregulation of proinflammatory cytokines, but not type 1 IFN.

Total body TLR9 deletion results in loss of NASH in experimental models, but the cellular requirement for TLR9 is not known (5). Liver macrophages, which include resident KCs and infiltrating monocytes, are the main proinflammatory cell populations in NASH. We tested the role of TLR9 signaling in these cells by generating mice in which TLR9 was removed from hepatocytes of HFD-fed mice (Tlr9fl/fl) and along with total TLR9-deficient mice, were placed on a HFD. The IRF7-dependent pathway can upregulate the IL-1 receptor antagonist (IL-1RA), which has antiinflammatory effects in the liver (14). To test whether mtDNA from hepatocytes of HFD-fed mice can activate type 1 IFN and IL-1RA, primary murine macrophages were stimulated with nDNA and mtDNA from hepatocytes of HFD-fed and CD-fed mice, and mRNA levels of Ifna, Ifnb, and Il1ra were quantified (Supplemental Figure 2, A–C). In contrast to the upregulation of TNF-α, hepatocyte mtDNA from HFD-fed mice, when compared with mtDNA from CD-fed mice, resulted in no change in Ifna, Ifnb, or Il1ra levels (15). This demonstrates that mtDNA from hepatocytes of HFD-fed mice results in the selective upregulation of proinflammatory cytokines, but not type 1 IFN.

Table 1. Characteristics of patient groups

<table>
<thead>
<tr>
<th>Group 1: Lean (n = 19)</th>
<th>Group 2: Obese, normal ALT (n = 19)</th>
<th>Group 3: Obese, high ALT (n = 9)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yr)</td>
<td>15.4 (9–16.5)</td>
<td>13.7 (9.5–18.8)</td>
</tr>
<tr>
<td>Sex (M/F)</td>
<td>11/8</td>
<td>9/10</td>
</tr>
<tr>
<td>Race (MED/AA/H)</td>
<td>9/3/7</td>
<td>6/5/8</td>
</tr>
<tr>
<td>GT (NGT/IGT/T2D)</td>
<td>17/2/0</td>
<td>11/3/1</td>
</tr>
<tr>
<td>Z-score BMI</td>
<td>0.41 (–1.70–1.04)</td>
<td>2.16 (1.75–2.67)</td>
</tr>
<tr>
<td>Fasting glucose (mg/dl)</td>
<td>86 (77–89)</td>
<td>87.5 (71–96)</td>
</tr>
<tr>
<td>2-h glucose (mg/dl)</td>
<td>110 (24–138)</td>
<td>117 (92–219)</td>
</tr>
<tr>
<td>Fasting insulin (mcU/ml)</td>
<td>14 (7–45)</td>
<td>16 (4–109)</td>
</tr>
<tr>
<td>Insulin sensitivity index</td>
<td>3.64 (2.52–5.53)</td>
<td>2.91 (0.48–8.1)</td>
</tr>
<tr>
<td>ALT (UI/l)</td>
<td>13 (8–27)</td>
<td>18 (9–21)</td>
</tr>
</tbody>
</table>

Demographic, biometric, and metabolic characteristics of the 3 patient groups. Normal ALT levels were below 35 U/l. M, male; F, female; MED, mixed European descent; AA, African-American; H, Hispanic; GT, glucose tolerance; NGT, normal glucose tolerance; IGT, impaired glucose tolerance; T2D, type 2 diabetes.
in NASH, NF-κB activation was examined in KCs in vivo using an NF-κB reporter mouse and gating on CD11b F4/80 double-positive cells. The HFD induced upregulation of the NF-κB reporter in CD11b F4/80 double-positive cells (Supplemental Figure 3E), and KCs from WT mice on a HFD had greater proinflammatory responsiveness to a TLR9 ligand (Supplemental Figure 3, F–H).

The demonstration of high plasma TLR9 ligand activity and a requirement for TLR9 in the development of HFD-induced NASH have immediate therapeutic implications. TLR9 antagonists have excellent safety profiles and represent a novel therapeutic strategy for NASH. We tested whether the TLR7/9 antagonist IRS954 could block the ability of hepatocyte mtDNA from HFD-fed mice to activate proinflammatory cytokines in primary macrophages. WT mice were placed on a HFD and concurrently administered...
HMGB1, which can enhance TLR9 signaling by DNA, and additional damage-associated molecular patterns (DAMPs), such as formyl peptides (16, 17). The systemic distribution of hepatocyte-derived mitochondria with the ability to activate TLR9 may have consequences in addition to hepatic inflammation. A chronic, low-level increase in plasma TLR9 ligand activity can be expected to activate TLR9 systemically, and this is consistent with the low-level systemic inflammation in NASH that is found in many organs including white adipose tissues and its loss in the absence of TLR9 (5). This leads to the intriguing possibility that hepatocyte-derived mtDNA is responsible for activation of macrophage populations in extrahepatic sites.

Methods

Statistics. Differences between 2 groups were compared with a Mann-Whitney U test using GraphPad Prism 6 software (GraphPad Software). A P value of less than 0.05 was considered significant. See the Supplemental Methods for additional details.

Study approval. This study was approved by the Yale University Animal Care and Use Committee and the Human Investigation Committee.

Author contributions

IGM, NS, YC, RH, and XO performed experimental work, data analysis, and reviewed the manuscript. RLC, AC, and MJS were responsible for experimental design, data analysis, and review of the manuscript. SC handled clinical management and phenotypic characterization of subjects. IGM and WZM were responsible for experimental design, data analysis, and manuscript preparation.
Figure 3. Weekly administration of a TLR9 antagonist protects against NASH. Soluble mtDNA (100 ng/ml) from hepatocytes of CD- and HFD-fed mice was added to KCs in the presence and absence of IRS954 (IRS) for 8 hours, and the upregulation of pro-
Il1b and Tnfa message was quantified (A and B). In a HFD model of NASH, weekly s.c. administration of the TLR9 antagonist IRS954 (5 mg/kg, n = 5) resulted in reduced steatosis, ballooning, and inflammation (C and D), reduced ALT (E), and reduced upregulation of inflammatory cytokines in the liver (F) compared with mice fed a HFD without injection of IRS954 (n = 5). Original magnification, ×100. Data represent the mean ± SEM. *P < 0.05 and **P < 0.01, by Mann-Whitney U test.

Acknowledgments

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