Necroptosis contributes to chronic inflammation and fibrosis in aging liver

Sabira Mohammed1, Nidheesh Thadathil2, Ramasamy Selvarani2, Evan H Nicklas2, Dawei Wang2, Benjamin F Miller2,3,4, Arlan Richardson1,2,3,5, and Sathyaseelan S. Deepa1,2,3*

1Stephenson Cancer Center, 2Department of Biochemistry and Molecular Biology, 3Oklahoma Center for Geroscience & Brain Aging, University of Oklahoma Health Sciences Center, 4Aging and Metabolism Research Program, Oklahoma Medical Research Foundation, 5Oklahoma City VA medical Center, Oklahoma City, OK, USA

*Corresponding Author: Sathyaseelan S Deepa, Ph.D.
Stephenson Cancer Center
Department of Biochemistry and Molecular Biology
Oklahoma Center for Geroscience & Brain Aging
University of Oklahoma Health Sciences Center
975 NE 10th Street, BRC-1368B
Oklahoma City, OK 73104, USA
E-mail: Deepa-Sathyaseelan@ouhsc.edu
Phone: 405-271-8001 (Extn:48393)
Fax: 405-271-2298

Running title: Necroptosis, inflammation and aging liver
Abstract

Inflammaging, characterized by an increase in low-grade chronic inflammation with age, is a hallmark of aging and is strongly associated with various age-related diseases, including chronic liver disease (CLD) and hepatocellular carcinoma (HCC). Because necroptosis is a cell death pathway that induces inflammation through the release of DAMPs, we tested the hypothesis that age-associated increase in necroptosis contributes to chronic inflammation in aging liver. Phosphorylation of MLKL and MLKL-oligomers, markers of necroptosis, as well as phosphorylation of RIPK3 and RIPK1 were significantly upregulated in the livers of old mice relative to young mice and this increase occurred in the later half of life (i.e., after 18 months of age). Markers of M1 macrophages, expression of proinflammatory cytokines (TNFα, IL6 and IL-1β), and markers of fibrosis were significantly upregulated in the liver with age and the change in necroptosis paralleled the changes in inflammation and fibrosis. Hepatocytes and liver macrophages isolated from old mice showed elevated levels of necroptosis markers as well as increased expression of proinflammatory cytokines relative to young mice. Short term treatment with the necroptosis inhibitor, necrostatin-1s (Nec-1s), reduced necroptosis, markers of M1 macrophages, expression of proinflammatory cytokines, and markers of fibrosis in the livers of old mice. Thus, our data show for the first time that liver aging is associated with increased necroptosis and necroptosis contributes to chronic inflammation in the liver, which in turn appears to contribute to liver fibrosis and possibly CLD.

KEYWORDS

Necroptosis, Inflammation, Fibrosis, Necrostatin-1s, Aging, Liver
INTRODUCTION

Aging is characterized by an increase in low-grade chronic inflammation, termed inflammaging, that is strongly associated with various age-associated diseases such as type 2 diabetes, cardiovascular disease, cancer, and neurodegenerative diseases such as Alzheimer’s disease. Therefore, inflammaging is considered as an important factor in the etiology of these age-associated diseases (Franceschi & Campisi, 2014). In humans, inflammaging is characterized by an increase in the levels of circulating proinflammatory cytokines interleukin 6 (IL6), tumor necrosis factor-α (TNF-α), and IL-1β (Ferrucci et al., 2005; Hager et al., 1994; Pedersen et al., 2003; Roubenoff et al., 1998), and increased levels of these cytokines are associated with diseases and mortality (De Martinis et al., 2005; Heneka et al., 2013; Youm et al., 2013). Inflammaging is the net effect of a multifactorial and multiorgan involvement where immune cells, especially macrophages, in various tissues such as adipose tissue, liver and kidney contribute to the production of these proinflammatory cytokines. Proinflammatory cytokines produced by tissues can affect its own function as well as other distal tissues. For example, proinflammatory cytokine production by adipose tissue increases with age (Mancuso & Bouchard, 2019) and inhibits the ability of adipose tissue to store fat leading to lipotoxicity in liver and skeletal muscle resulting in hepatic steatosis and muscle dysfunction, respectively (Schwartz et al., 1990).

Aging is associated with an increase in the levels of inflammatory cytokines in the liver (Stahl et al., 2020). Damage-associated molecular patterns (DAMPs, example, HMGB1, mitochondrial DNA, nuclear DNA, ATP etc.) released from damaged or dying hepatocytes are one of the proposed mediators of chronic inflammation in the liver (Brenner et al., 2013). Necroptosis, a form of programmed necrosis, is a regulated cell death pathway that is strongly associated with increased inflammation through the release of DAMPs from the necroptotic cells (Galluzzi et al., 2018; Newton & Manning, 2016). In contrast, cell death by apoptosis is associated with limited release of DAMPs and therefore is less inflammatory in nature. Absence of caspase-8, an initiator of extrinsic apoptosis, switches cell death from apoptosis to necroptosis (Fritsch et al., 2019). Necroptosis most likely evolved as an alternative form of cell death to kill cells infected by viral pathogens and to promote inflammatory and immune responses to limit the spread of the viruses (Dondelinger et al., 2016). Necroptosis is initiated when necroptotic stimuli (e.g. TNFa, oxidative stress or mTOR/Akt pathway) (Royce et al., 2019) sequentially phosphorylate and activate receptor-interacting protein kinase 1(RIPK1) and RIPK3 which in turn phosphorylate the
pseudokinase mixed lineage kinase domain-like (MLKL) protein. Phosphorylation of MLKL leads to its oligomerization, oligomerized MLKL then binds to and disrupts the cell membrane releasing cellular components including the DAMPs, which initiate and exacerbate the inflammatory process (Newton & Manning, 2016; Samson et al., 2020). DAMPs released from necroptotic cells bind to pattern recognition receptors (PRRs) such as toll like receptors (TLRs) on innate immune cells resulting in the induction of pro-inflammatory cytokines (Owens, 2009; Rodríguez-Gómez et al., 2020). Necroptosis has emerged as a novel mode of cell death in various chronic liver diseases such as non-alcoholic fatty liver disease (NAFLD) and non-alcoholic steatohepatitis (NASH), conditions that are also associated with aging liver (Afonso et al., 2015; L. Chen et al., 2020; Gautheron et al., 2014; Majdi et al., 2020). Various liver cell types such as hepatocytes (Zhong et al., 2020; B. Zhang et al., 2019a), Kupffer cells (Blériot et al., 2015) and endothelial cells (Zelic et al., 2018) are shown to undergo necroptosis under pathological conditions or injury.

Blocking necroptosis has been shown to reduce hepatic inflammation in various mouse models of liver diseases (Afonso et al., 2015; Gautheron et al., 2014; Majdi et al., 2020), whereas the role of necroptosis in age-associated hepatic inflammation is unexplored. Previously, we reported that necroptosis increases with age in the white adipose tissue of mice and is reduced by interventions that delay aging (dietary restriction or in Ames dwarf mice) (Deepa et al., 2018; Royce et al., 2019). Recently, we found that necroptosis and inflammation are increased in the livers of a mouse model of accelerated aging (mice deficient in the antioxidant enzyme, Cu/Zn superoxide dismutase, $Sod1^{-/-}$ mice) and blocking necroptosis reduced inflammation in the livers of $Sod1^{-/-}$ mice (Mohammed et al., 2021). Based on these findings, we hypothesized that age-associated increase in necroptosis might contribute to the age-related increase in hepatic inflammation. Because chronic inflammation can lead to liver fibrosis, we also studied the effect of necroptosis on the markers of fibrosis in the liver with age. Our data show that markers of necroptosis, inflammation, and fibrosis increase with age in the livers of mice and blocking necroptosis (using a pharmacological inhibitor of necroptosis, necrostatin-1s, Nec-1s) reduced the expression of inflammatory cytokines and fibrosis markers in the livers of old mice.

2 RESULTS

2.1 Necroptosis markers increase with age in the livers of mice

We first measured necroptosis in the livers of mice ranging in age from 7 to 24 months by two different approaches: (1) flow cytometry using Annexin V/propidium iodide (PI) staining...
(Pietkiewicz et al., 2015), and (2) biochemical analysis by western blotting to identify changes in the expression of proteins involved in necroptosis (Mohammed et al., 2021). Figure 1a shows flow cytometry data where cells positive for Annexin V are early apoptotic and cells double positive for Annexin V and PI are necroptotic/late apoptotic. As shown in Figure 1a, the percent of liver cells undergoing apoptosis and late apoptosis/necroptosis were significantly increased with age. Early apoptotic cell population was significantly higher in the liver starting from 12 months of age (2.5 fold) and remained elevated at 18 (2.7 fold) and 24-months of age (2.2-fold), whereas late apoptotic/necroptotic population was significantly higher at 18- and 24-months relative to 7-month-old mice (3.3- and 4.6-fold). There was no significant increase in the necrotic population at 18- and 24-months relative to 7-month-old mice, however, necrosis was increased at 12 months relative to 7-month-old mice. The percent of cells undergoing necroptosis/late apoptosis was 2 fold higher than the percent of cells undergoing early apoptosis in 24-month-old mice.

As flow cytometry cannot differentiate between necroptotic and late apoptotic cell populations, we also measured the expression of the three proteins involved in necroptosis. As can be seen in Figures 1b and c, the levels of MLKL and RIPK3 were significantly increased in 18- and 24-month-old mice compared to the 7- and 12-month old mice. However, we observed no changes in the level of RIPK1 with age. Figure S1a shows that the changes in the levels of MLKL and RIPK3 were associated with increased transcript levels of these proteins and no change in the transcript levels of RIPK1 was observed. Because the phosphorylation of MLKL, RIPK3 and RIPK1 trigger necroptosis (He et al., 2016) the levels of phospho-MLKL (P-MLKL), P-RIPK3, and P-RIPK1 were measured. As shown in Figures 1b and c, the levels of these three phosphorylated proteins were significantly increased in the 18- and 24-month-old mice compared to the 7- and 12-month-old mice. Even though the levels of P-MLKL and P-RIPK3 showed a slight reduction at 24 months of age relative to 18 months, this reduction was not statistically significant. In contrast, P-RIPK1 showed a significant increase at 24 months relative to 18 months. The increased expression of P-MLKL in the livers of old mice was further confirmed by immunofluorescence staining of young and old mice livers. The specificity of the P-MLKL staining was confirmed by using liver tissues from old Mlkl−/− mice (Figure 1d). RIPK3-mediated phosphorylation of MLKL leads to conformational changes in MLKL leading to the formation of MLKL tetramers or octamers. For mouse MLKL, tetramers fail to translocate to the plasma membrane and octamer formation is required for pore formation in the membrane(Huang et al.,
Therefore, we measured the MLKL-oligomers in the liver tissues of 7-, 12-, 18-, and 24-month-old mice (Figure 1e). Consistent with the age-associated increase in the levels of MLKL, MLKL oligomerization also showed a significant increase with age; levels of MLKL-oligomers were similar in the livers of 7- and 12-month-old mice, whereas MLKL-oligomer levels were significantly increased by 2 fold in 18- and 5 fold in 24-month-old mice, compared to 7- or 12-month-old mice.

Because DAMPs are released from cells undergoing necroptosis, we measured the levels of circulating high-mobility group box-1 (HMGB1), a DAMP that has been shown to be released by hepatic necroptosis (Wen et al., 2020) and is associated with acute liver injury and chronic liver disease (Gaskell et al., 2018). As shown in Figure 1f, circulating levels of HMGB1 are significantly elevated at 18-months (4-fold) and 24-months (7-fold) of age relative to 7- or 12-month-old mice. The changes in circulating HMGB1 levels parallel the changes in the necroptosis markers in the liver, providing additional data supporting that necroptosis increases with age in liver.

Hepatocytes constitute 80% of the liver mass and are reported to undergo cell death in CLD (Shojaie et al., 2020), therefore, we tested whether hepatocytes were undergoing increased necroptosis. Hepatocytes were isolated from young (7-month) and old (24-month) mice and purity of the isolated hepatocytes were confirmed by western blotting using albumin, the marker for hepatocytes (Liu et al., 2017; Malato et al., 2011); CD31, the marker for endothelial cells (Katz et al., 2004) and desmin, marker for hepatic stellate cells (An et al., 2020) and also mRNA levels of albumin, Clec4f (Kupffer cell marker) (Scott et al., 2016), F4/80 (macrophage marker) (Delire et al., 2016; Stahl et al., 2020) and CD31 (Figure S1 b,c). Figure 2a shows flow cytometry data of isolated hepatocytes from young and old mice. There was significant increase in apoptotic (3-fold) and necroptosis/late apoptosis (4.4-fold) populations from old mice relative to young mice, whereas the percentage of necrotic cell population was significantly reduced in hepatocytes from old mice. Necroptosis in hepatocytes was confirmed by assessing changes in the levels of necroptosis marker, P-MLKL. Levels of P-MLKL and MLKL were both significantly elevated (2-fold) in the hepatocytes isolated from old mice relative to young mice (Figure 2b). This was further confirmed by co-immunostaining of liver sections from old mice with P-MLKL and albumin, which showed that P-MLKL colocalizes with albumin (Figure 2c). Similarly, levels of MLKL-oligomers were significantly elevated (4-fold) in hepatocytes isolated from old mice (Figure 2d).
Next, we assessed the levels of P-RIPK3, RIPK3, P-RIPK1, and RIPK1 in isolated hepatocytes from young and old mice. As shown in Figure 2e, levels of P-RIPK3 and RIPK3 are significantly elevated in the hepatocytes from old mice (2-fold and 1.4-fold respectively), whereas levels of P-RIPK1 and RIPK1 were similar in hepatocytes from young and old mice. *Mlkl* and *Ripk3* transcript levels are also significantly elevated in the hepatocytes of old mice relative to young mice (7.5-fold and 8-fold, respectively), whereas *Ripk1* transcript levels were similar in hepatocytes from young and old mice (Figure S1d). Because Kupffer cells are reported to undergo necroptosis during bacterial infection (Blériot et al., 2015), we tested whether liver macrophages undergo necroptosis in aging. Double immunostaining using P-MLKL and F4/80 (macrophage marker) showed colocalization of these two proteins in the livers of old mice (Figure 2f). Similar to the increased transcript levels of *Mlkl* in hepatocytes from old mice, macrophage population isolated from old mice also showed a significant increase in *Mlkl* transcripts (2-fold) relative to young (Figure S1e). Moreover, we also checked annexin/propidium iodide staining in LSEC and Kupffer cell fractions isolated by Magnetic activated cell sorting (Figure S1f). There was significant increase (1.7 fold) in the late apoptotic/necroptotic population in the Kupffer cell fraction isolated from old liver. The purity of the isolated fractions was checked by real time PCR using cell specific primers (Figure S1g).

### 2.2 Age-associated increase in chronic inflammation in the livers of mice

DAMPs released from cells undergoing necroptosis bind to cell surface receptors on innate immune cells (e.g. macrophages) to induce an inflammatory response through the production of pro-inflammatory cytokines (Chen & Nuñez, 2010; Rock et al., 2011). Therefore, we measured changes in liver macrophages with age as well as levels of proinflammatory cytokines in the liver, isolated hepatocytes and isolated macrophage population. Flow cytometry analysis showed that the total immune cell population, as measured by the percentage of CD45^+^ cells (An et al., 2020; Blom et al., 2009), was significantly increased in the livers of old mice (67.9±6.84%) relative to young mice (32.86±0.92%) (Figure 3a). Next, total macrophage population was sorted by staining the cells with F4/80, a cell surface marker expressed by liver macrophages, i.e. resident Kupffer cells and infiltrating monocyte derived macrophages (Ramachandran et al., 2012; Stahl et al., 2020). As shown in Figure 3b, a significant increase in the F4/80^+^ cell population was observed in the livers of old mice (27.9±1.8%) relative to young mice (16.6±1.6%). We also measured transcript levels of F4/80 and monocyte chemoattractant protein-1 (MCP-1/CCL2), which is
produced predominantly by Kupffer cells, across different age groups. As shown in Figure S2a, transcript levels of F4/80 were similar in 7- and 12-month-old mice, whereas it was significantly increased (2-fold) at 18- and 24-months of age (left panel). The expression of MCP-1 is also significantly elevated in the livers of 18- and 24-month-old mice (5- and 6-fold) compared to 7- or 12-month-old mice (Figure S2a, right panel). These data are consistent with macrophage number in the liver increasing with age.

Macrophages are categorized into M1 or M2 phenotypes. In general, M1 macrophages play a more proinflammatory role in liver injury and M2 macrophages exert an anti-inflammatory effect (Alisi et al., 2017). Expression of cell surface receptors such as CD68, CD86, CD80, TLR4 and CD11c are the common markers for M1 macrophages, and Arg1, CD206 and Fizz1 are used as markers of M2 macrophages. To quantify the changes in M1/M2 macrophage population, we performed flow cytometric analysis with the liver lysates from young and old mice. M1 macrophage population assessed by CD45+CD11b+Ly6G−CD80+ F4/80+ cells (Davies et al., 2013; Lynch et al., 2018; Ramachandran et al., 2012) were significantly increased in the livers of old mice (9.03±0.59%) in comparison to young mice (2.45±0.18%) (Figure 3c). M2 macrophage population assessed by CD45+CD11b+Ly6G−F4/80+ CD206+ cells (Bai et al., 2017; Beljaars et al., 2014; C. Lee et al., 2019) was reduced in the livers of old mice (5.06±0.12) relative to young mice (6.48±0.56), however, this reduction did not reach statistical significance (Figure 3d). We also analyzed the transcript levels of M1 and M2 macrophage markers in the livers of mice in the four different age groups. M1 macrophage markers (CD68, CD86, TLR4, and CD11c) were significantly upregulated at 18-months compared to their expression at 7 or 12-months of age. Expression of M1 macrophage markers at 24-months were similar to those at 18-months, except for CD11c, which was increased between 18 and 24 months of age (Figure S2b). The markers of M2 macrophages (Arg1 and Fizz1) are significantly reduced at 12-, 18-, and 24-months compared to their expression at 7 months of age (Figure S2c).

Consistent with the increase in M1 macrophage markers, expression of proinflammatory cytokines TNFα, IL6 and IL-1β were significantly up-regulated (2- to 3-fold) in the livers of 18- and 24-month-old mice compared to 7- or 12-month-old mice (Figure 3e). The changes in protein levels of TNFα and IL-1β also were significantly increased in the livers of 18- and 24-month-old mice relative to 7- or 12-month old mice (Figure 3f). Analysis of the isolated macrophage population (F4/80+ fraction) showed a significant increase in the transcript levels of IL-1β (3.5-
fold) and MCP-1 (5.6-fold) in old mice relative to young mice (Figure 3g). However, transcript levels of TNFα or IL6 in the F4/80+ fraction of cells was similar in young and old mice. Because hepatocytes are also reported to produce proinflammatory cytokines (Stahl et al., 2020), we measured the expression of proinflammatory cytokines in hepatocytes isolated from young and old mice. Transcript levels were significantly increased for TNFα (4-fold), IL6 (4-fold) and IL-1β (40-fold) and MCP1/CCL2 (4-fold) in the hepatocytes isolated from old mice relative to hepatocytes from young mice (Figure 3h). Next we tested whether liver sinusoidal endothelial cells (LSEC), which are known to produce inflammatory cytokines in chronic liver diseases (Wang & Peng, 2021), also contributes to inflammation in aging liver. As shown in Figure S2d expression of TNFα, IL6, IL-1β, and MCP1 were similar in LSEC isolated from young and old mice.

### 2.3 Age-associated increase in fibrosis in the livers of wild type mice

Chronic hepatic inflammation is a major driver of liver fibrosis (Koyama & Brenner, 2017), and the transformation of quiescent hepatic stellate cells (HSCs) to trans differentiated myofibroblasts, which produce extracellular matrix components (ECM), drives fibrogenesis (U. E. Lee & Friedman, 2011). Transforming growth factor beta (TGF β) is a major player in the induction of fibrosis and is produced by several liver cells like LSEC, Kupffer cells, platelets, hepatocytes and macrophages (Dewidar et al., 2019; Ghafoory et al., 2018; Schon & Weiskirchen, 2014). TGFβ expression, as measured by transcript levels, was significantly increased (1.5- and 2-fold) in the livers of 18- and 24-month-old mice relative to its expression in the livers of 7- or 12-month old mice (Figure 4a). Desmin, a protein produced by activated HSCs, which is strongly upregulated in liver fibrosis (Puche et al., 2013), also showed a significant increase with age. At 18- and 24-months of age, desmin protein levels in the liver were significantly higher (3- to 4-fold) than in 7- or 12-month-old mice (Figure 4b). Consistent with the activation of HSCs, the expression of ECM components Col1α1 and Col3α1 were significantly increased (2.5-fold) in the livers of 18- and 24-month-old mice relative to 7- or 12-month-old mice (Figure 4c). Next, we assessed total collagen content by measuring concentration of hydroxyproline (OHP) because total collagen content is an indicator of the severity of fibrosis (Wei et al., 2020). At 18-and 24-months of age, OHP content in the liver was significantly higher (1.5- and 1.9-fold) relative to 7- or 12-month-old mice (Figure 4d). We also performed picrosirius red staining (PSR), a commonly used histological technique for visualizing collagen in paraffin-embedded tissue sections (An et al., 2020; Lankadasari et al., 2018). PSR staining in the old mice liver was significantly higher (2.7-
fold) relative to young mice, suggesting increased levels of collagen in old mice liver (Fig. 4e). Thus, markers of fibrosis show an age-associated increase in livers of mice that paralleled the changes in inflammation and necroptosis.

2.4 Blocking necroptosis reduces hepatic inflammation and fibrosis

To determine if necroptosis was responsible for the increased inflammation and fibrosis seen in the livers of old mice, we tested the effect of blocking necroptosis on hepatic inflammation and fibrosis in old mice. Mice were treated with Nec-1s (RIPK1 inhibitor) for 30 days, which has been shown to effectively block necroptosis and reduce inflammation in the liver (Mohammed et al., 2021) as well as other tissues (Liang et al., 2018; Lin et al., 2020; Shen et al., 2019). Three groups of mice were used for these experiments: (1) young wild type (WT) mice (8-month-old, vehicle treated), (2) old WT mice (24-month-old, vehicle treated), and (3) old WT mice (24-month-old, Nec-1s treated). Nec-1s treatment had no effect on the body weight or liver weight of old WT mice (Figure S3a and S3b), indicating no obvious negative effect of Nec-1s on old mice. To test whether Nec-1s treatment blocked necroptosis in the livers of old WT mice, levels of P-MLKL and MLKL-oligomers were measured. The data in Figures 5a and 5b show that Nec-1s treatment significantly reduced necroptosis as predicted, i.e., the levels of P-MLKL and MLKL-oligomers were reduced to the levels seen in young mice. Since apoptosis is increased in liver of old mice and Nec-1s inhibits RIPK1, which is known to be involved in apoptosis, we checked whether Nec-1s has an effect on apoptosis. Towards this, the expression of Cleaved Caspase-3 was checked (Figure S3c). As seen in the figure, Nec-1s did not exert an effect on apoptosis. To test whether Nec-1s had any effect on other pathways, we assessed changes in cell senescence. To our surprise, we found that transcript levels of p16 and p21 expression, markers of cell senescence, which increased with age, (Y. Zhang et al., 2017) was significantly reduced after Nec-1s treatment (Figure 5c).

Next, we measured the effect Nec-1s treatment on the age-related increase in macrophage number, macrophage activation, and hepatic inflammation. Transcript levels of F4/80, MCP1 and M1 macrophage markers (CD68, CD86, CD11c and TLR4) that were elevated in the livers of old WT mice were significantly reduced by Nec-1s to levels present in the young WT mice (Figures S3d and S3e), whereas expression of M2 macrophage markers (Arg1 and Fizz1) were unaffected by Nec-1s treatment (Figure S3f). Consistent with this, protein expression of expression of F4/80 (4-fold) and CD68 (5-fold) was significantly increased in the livers of old mice, which were
significantly reduced by Nec-1s treatment to levels similar to levels in young liver as shown by immunofluorescence staining (Figure 5d). In contrast, protein expression of CD206 (M2 macrophage marker) was significantly reduced (80%) in the livers of old mice relative to young mice and Nec-1s had no effect on CD206 expression in the livers of old mice (Figure 5d).

Similarly, levels of transcripts for proinflammatory cytokines TNFα, IL6 and IL-1β, which were elevated in the livers of old WT mice, were significantly reduced by Nec-1s treatment and were comparable to the levels in young mice (Figure S3g). Consistent with the reduction in transcript levels, protein levels of TNFα in the livers of old WT mice were also significantly reduced by Nec-1s treatment (Figure 5e).

Finally, we determined the effect of blocking necroptosis on liver damage and fibrosis. Liver damage was assessed by measuring circulating levels of alanine aminotransferase (ALT). We found a significant increase in the serum levels of ALT in the old mice (2.2-fold) relative to young mice, and Nec-1s treatment significantly reduced serum ALT levels in old mice to levels comparable to young mice (Figure 5f). Fibrosis was assessed by measuring the levels of Col1α1 and Col3α1 as well as OHP content. As shown in Figures S3h and Figure 5g the higher levels of these markers of fibrosis in the livers of old WT mice were significantly reduced by Nec-1s. To check for the effect of Nec-1s on deposition of collagen fibers, Picrosirius red staining was done (Figure 5h). Consistent with the effect of Nec-1s on the markers of fibrosis, we saw that Nec-1s reduced the increased collagen deposition in old mice (2.5 fold) to levels similar to that of young mice. Thus, our data shows that short term Nec-1s treatment can exert its effect on the M1 macrophages and reduce age-associated hepatic inflammation and thereby liver damage and fibrosis.

3 DISCUSSION

The main findings of the study are that liver aging is associated with increased necroptosis, inflammation, and fibrosis and inhibiting necroptosis reduced age-associated inflammation and fibrosis in the liver. This suggests that necroptotic cell death contributes to chronic inflammation and subsequent development of fibrosis in liver aging.

The age-associated increase in chronic, sterile inflammation, termed inflammaging, is strongly associated with aging and age-associated diseases (Franceschi & Campisi, 2014). For example, studies have shown that an age-related increase in hepatic inflammation (Stahl et al.,
2020) is associated with the onset and progression of CLD, e.g., fibrosis and HCC (Franceschi et al., 2000; Franceschi et al., 2007; Morsiani et al., 2019). However, the pathway(s) that contribute to the age-related increase in hepatic inflammation is not clearly understood. Necroptosis is an inflammatory mode of cell death resulting in the release of DAMPs, which initiate an inflammatory response by binding to the cell surface receptors on innate immune cells (Newton & Manning, 2016). An age-associated increase in DAMPs has been proposed to be a potential driver of inflammaging (Goldberg & Dixit, 2015), and studies have shown that necroptosis-mediated inflammation may play an important role in a variety of age-related diseases such as Alzheimer's disease, Parkinson's disease, and atherosclerosis (Royce et al., 2019). We have reported that necroptosis increases with age in the adipose tissue of mice, and this increase was attenuated by dietary restriction (Deepa et al., 2018), suggesting for the first time that necroptosis plays a role in aging. We also observed that necroptosis was increased in the livers of Sod1KO mice, which show accelerated aging and develop CLD and spontaneous HCC (Mohammed et al., 2021). In this study, we determined if necroptosis played a role in the age-related increase in inflammation in liver of mice. Our findings show that necroptosis, as measured by the increase in the expression of P-MLKL and MLKL-oligomers, increases with age in the livers of mice. While the expression of P-MLKL was similar at 18- and 24-months of age, MLKL oligomer level was significantly increased from 18 to 24 months in the livers of old mice. Recent studies have identified TAM kinases as positive regulator of oligomer formation and HSP70 as a negative regulator of oligomer formation (Johnston et al., 2020; Meng et al., 2021; Najafov et al., 2019). Therefore, it is possible that levels/activities of HSP70 or TAM kinases might be regulating MLKL oligomerization at 24 months. We also found that P-RIPK1 levels were significantly higher in the liver at 24-months relative to 18-months. In addition to necroptosis, RIPK1 is also involved in apoptosis as well as cell survival pathways such as NF-kB, Akt, and JNK (Dara et al., 2015; Kondylis et al., 2017; McNamara et al., 2013). Therefore, it is possible that an increase in P-RIPK1 at 24 months might reflects the involvement of P-RIPK1 in the other cellular pathways.

To get an idea of the cell type involved in necroptosis, we measured necroptosis in hepatocytes and macrophages isolated from livers of young and old mice. We found that hepatocytes isolated from old mice exhibit increased necroptosis relative to hepatocytes from young mice, suggesting that necrototic hepatocytes might be a major source of DAMPs in aging liver. Hepatocyte necroptosis has previously been reported in bacterial hepatitis and TNF challenge
(Koschel et al., 2021), ischemia and reperfusion injury (Zhong et al., 2020), chronic alcoholic liver disease (Lu et al., 2016), nonalcoholic fatty liver disease (Afonso et al., 2015; Wu & Nagy, 2020) and from hepatic O-GlcNAc transferase deficiency in mice (Zhang et al., 2019). Similar to hepatocytes, liver macrophages in old mice also expressed necroptosis markers. Macrophages have been reported to undergo necroptosis and contribute to pathological conditions like atherosclerosis and bacterial pneumonia (González-Juarbe et al., 2015; Karunakaran et al., 2016). However, this is the first report showing that macrophage necroptosis occurs in the aging liver. Other cell types such as endothelial cells (Gong et al., 2019) and hepatic stellate cells (Jia et al., 2018) also have been reported to undergo necroptosis in pathological conditions. Whether cell types other than hepatocytes and macrophages undergo necroptosis in the liver during aging and the proportions of these cell types undergoing necroptosis in aging need to characterized.

Because necroptosis has been associated with increased inflammation, we determined if an increase in inflammation in liver was observed that paralleled the increase in necroptosis. We first focused on liver macrophages, because they are the major producers of proinflammatory cytokines in the liver (Wen et al., 2021). Our data show that total macrophage content as well as proinflammatory M1 macrophages increase with age, whereas anti-inflammatory M2 macrophages decline with age in the liver. Consistent with the increase in macrophages, expression of key proinflammatory cytokines associated with inflammaging, TNFα, IL6 and IL-1β, also showed an age-related increase in the liver. Thus, our data show that age-related increase in necroptosis is associated with an age-related increase in inflammation in the liver and the increase in inflammation is seen in a time frame similar to changes in necroptosis, i.e. occurring after 18 months of age. Previous studies have reported that primary hepatocytes can also secrete proinflammatory cytokines (Rowell et al., 1997; Stahl et al., 2020), so we measured the expression of inflammatory cytokines in isolated hepatocytes from young and old mice. Our data show that the expression of proinflammatory cytokines TNFα, IL6 and IL1β and the chemokine CCL2 were increased in hepatocytes isolated from old mice. Thus, both liver macrophages and hepatocytes are potential sources of inflammatory cytokines seen in the liver with age.

Our data show that the age-related increase in inflammation in liver is associated with increased necroptosis. To determine if necroptosis is causative, we inhibited necroptosis by Nec-1s to determine if reducing necroptosis altered hepatic inflammation. Previous studies have shown that Nec-1s, which inhibits necroptosis by targeting RIPK1 (Takahashi et al., 2012), reduces
inflammation associated with necroptosis in multiple sclerosis and amyotrophic lateral sclerosis (Yuan et al., 2019). No off-target effect for Nec-1s has been reported so far and is considered as a very specific and potent inhibitor of RIPK1 (Takahashi et al., 2012). In the present study, we found that short-term treatment of old mice with Nec-1s reduced M1 macrophages and the expression of proinflammatory cytokines. Interestingly, Nec-1s treatment had no effect on the levels of anti-inflammatory M2 macrophages, suggesting that reduction of M1 macrophages could be the possible mechanism for reduced inflammation observed with Nec-1s treatment. DAMPs are one of the factors that are known initiate macrophages activation and polarization (Lee et al., 2020). Therefore, a reduction in DAMPs release due to necroptosis inhibition might contribute to the reduction in M1 macrophage markers. It is also possible that reduction in CCL2 with Nec-1s treatment could reduce monocyte infiltration, thereby reducing the M1 macrophage pool. In support of this, RIPA-56, a necroptosis inhibitor, has been shown to reduce monocyte infiltration to spinal cord in a mouse model of multiple sclerosis (S. Zhang et al., 2019). Even though, we did not see an induction of apoptosis in the livers of mice treated with Nec-1s, we cannot rule out the possibility that M1 macrophage population undergoes selective cell death through apoptosis or other cell death pathway(s) in response to Nec-1s. Future in vitro characterization of liver macrophages regarding their responses to danger signals will be required to identify the nature of inflammatory cytokines produced by different DAMPs. We recently found that short term Nec-1s treatment could effectively reduce necroptosis and inflammation in the livers of Sod1KO mice (Mohammed et al., 2021), which show accelerated aging (Deepa et al., 2018). Even though RIPK1 has been known to have a role in apoptosis, we found that use of Nec-1s that targets RIPK1 had no effect on apoptosis in the liver of old mice. Therefore, it appears that necroptosis plays a role in inflammation associated with aging in liver.

Chronic inflammation is one of the proposed mediators of CLD, especially fibrosis, which has been reported to increase with age in the liver of mice (Abdelmegeed et al., 2016; Jin et al., 2020; Koyama & Brenner, 2017) and humans (Kim et al., 2015). However, the factor(s) contributing to the age-related increase in hepatic fibrosis is not known. Therefore, we were interested in determining if necroptosis-induced inflammation played a role in the age-related increase in fibrosis in the livers of mice. We found that blocking necroptosis and reducing inflammation in the livers of old mice reduced markers of fibrosis. In addition, Nec-1s treatment reduced serum ALT levels in old mice, suggesting that blocking necroptosis reduced
hepatocellular injury and CLD (Kim et al., 2008). Our findings are supported by previous reports showing that blocking necroptosis genetically (Ripk3−/−) in mice fed a methionine-deficient diet or pharmacologically (RIPA56) in mice fed a high-fat diet reduced liver fibrosis (Gautheron et al., 2014; Majdi et al., 2020). Thus, it appears that necroptosis plays an important role in fibrosis in liver under a variety of conditions, including aging.

In our study, we looked at a range of ages in order to delineate the progression of necroptosis and inflammation. Our data demonstrate that necroptosis and inflammation do not increase linearly with age. Rather, the increase in necroptosis and inflammation in the liver occurs in the latter half of life, around 18 months of age when mice began to exhibit some age-associated pathologies (Stahl et al., 2020). We believe that several factors could be involved in the age-related increase in necroptosis. For example, circulating levels of TNFα, a well-known inducer of necroptosis (Degterev et al., 2005) have been reported to increase with age in humans (Bruunsgaard et al., 2003; Kirwan et al., 2001). In addition, oxidative stress, which has been shown to increase with age in a wide variety of tissues including liver (Bokov et al., 2004; Lozada-Delgado et al., 2020) and occurs later in life (Hamilton et al., 2001), has also been shown to induce necroptosis in both cell culture and animal studies. For example, hydrogen peroxide induces necroptosis in retinal pigment epithelial cells (Hanus et al., 2015), and paraquat-induced oxidative stress leads to necroptosis in cardiomyocytes (Zhang et al., 2018). Similarly, increased reactive oxidative species production due to the deletion of the antioxidant enzyme glutathione peroxidase 4 in hematopoietic cells causes necroptosis in erythroid precursor cells (Canli et al., 2016). We also found that Sod1−/− mice, which show a dramatic increase in oxidative stress have increased expression of necroptosis markers in the liver (Mohammed et al., 2021) and white adipose tissue (Royce et al., 2019). Finally, mammalian target of rapamycin (mTOR) signaling has also been reported to induce necroptosis. mTOR signaling has been shown to increase with age in various tissues including liver (Baar et al., 2016) and increased mTORC1 signaling is associated with various age-related diseases, including HCC (Caccamo et al., 2010; Ferrin et al., 2020; Grabiner et al., 2014; Völkers et al., 2014). In support of the role of mTOR pathway in necroptosis activation, studies have shown that induction of necroptosis is blocked by the combined treatment of Akt and mTOR inhibitors in the hippocampal neuronal cell line HT22 (Liu et al., 2014), and aberrant activation of mTOR by genetic deletion of TSC1 in intestinal epithelial cells resulted in the overexpression of RIPK3, epithelial necrosis, and subsequent colitis (Xiao, 2018). Factors
known to induce necroptosis (TNFα, oxidative stress, and mTOR) also increases with age (Royce et al., 2019). Therefore, an age-dependent increase in these pathways in the liver might contribute to increased necroptosis in aging liver. It is possible that age-associated increase in TNFα/oxidative stress/mTOR activation in liver makes cells more prone to undergo necroptosis leading to the release of DAMPs and activation of hepatic macrophages increasing hepatic inflammation leading to liver fibrosis. Our finding that a short term Nec-1s treatment late in life can effectively block hepatic inflammation and reduce liver fibrosis suggest that blocking necroptosis would be an effective approach to reverse fibrosis associated with aging and CLD. The long-term effects of Nec-1s on age associated hepatic inflammation and fibrosis in aging need to be addressed.

This is the first study to demonstrate that aging is associated with elevated necroptotic cell death in the liver (hepatocytes and macrophages). In addition, the study identified that necroptosis is a contributor to hepatic inflammation and fibrosis in liver aging. While the current study was performed using male mice, future studies will address the effect of sex on necroptosis and inflammation in the liver using female mice. Although Nec-1s is specific inhibitor of RIPK1, we cannot rule out off target effects. Therefore, future studies will include other pharmacological compounds that inhibit necroptosis or using genetic knockout models of Ripk3 or Mlkl. Therapeutic applications of RIPK1 inhibitors for the treatment of a variety of human diseases are being tested in clinical trials: For example, GSK′772 is being developed for peripheral autoimmune diseases, including psoriasis, rheumatoid arthritis and ulcerative colitis, whereas the brain-penetrant DNL747 is in human clinical trial phase Ib/IIa for amyotrophic lateral sclerosis (ALS) (Mifflin et al., 2020). Moreover, several of the FDA approved anti-cancer drugs such as Sorafenib and Dabrafenib have been identified as anti-necroptotic agents (Fulda, 2018). While the adverse effect of necroptosis inhibition is not clear, it is highly unlikely that inhibiting necroptosis by targeting RIPK1 will increase the susceptibility to viral infection. This is because, virus-induced necroptosis is through activation of RIPK3-MLKL pathway, independent of RIPK1. Therefore, we predict that blocking RIPK1 will not affect virus-induced necroptosis, however, this need to tested. Similarly, it will be difficult to predict whether necroptosis inhibition will have a pro-tumorigenic effect. Necroptosis has been reported to exhibit dual effects in cancer, i.e. necroptosis can either promote or reduce tumor growth depending on the type of cancer. For example, reduced expression of necroptotic markers promote tumor growth in breast cancer (Stoll et al., 2017) and
acute myeloid leukemia (Höckendorf et al., 2016), whereas in pancreatic ductal adenocarcinoma (Seifert et al., 2016) and cholangiocarcinoma (Lomphithak et al., 2021) increased expression of necroptosis markers are associated with tumor growth. Because aging is a slow and gradual process, further studies with long-term Nec-1s treatment will be needed to address the feasibility of using necroptosis inhibitor(s) for inflammaging and aging.

In conclusion, our study show that necroptosis is a potential contributor to age-associated chronic inflammation and fibrosis in the liver. Thus, necroptosis is a potential therapeutic target for treating chronic liver diseases associated with aging.

4 EXPERIMENTAL PROCEDURES

4.1 Animals

All procedures were approved by the Institutional Animal Care and Use Committee at the University of Oklahoma Health Sciences Center (OUHSC). C57BL/6 male wild-type (WT) mice of different age groups (7-, 12-, 18-, and 22 to 24-month-old) (n= 5-10/group) were received from National Institute on Aging. After receiving the mice, they were group housed as received in ventilated cages 20 ± 2 °C, 12-h/12-h dark/light cycle and were fed rodent chow (5053 Pico Lab, Purina Mills, Richmond, IN) ad libitum at the OUHSC animal care facility for 2 weeks before euthanizing the mice for tissue collection.

4.2 Annexin V/Propidium iodide (PI) staining of whole liver tissue

Mice were euthanized and in vivo perfusion was performed as described by (Liu et al., 2017) and annexin PI staining was performed as described in Mohammed et al., 2019 and analyzed using FACS Calibur flow cytometer (BD Biosciences, San Jose, CA). The data was analyzed using FlowJo (BD Biosciences, NJ, USA) software.

4.3 Isolation of hepatocytes

Hepatocyte isolation was performed as described by (Liu et al., 2017). The purity of the isolated hepatocytes was checked by real time PCR and western blotting for the expression of liver cell specific markers – albumin (hepatocytes), F4/80 (macrophages), Clec4f (Kupffer cells), CD31 (endothelial cells) and desmin (hepatic stellate cells).

4.4 Quantitative real-time PCR (RT-PCR)
RT-PCR was performed using 20mg frozen liver tissues as described previously (Mohammed et al., 2021). The list of primers used are given in Table S1.

### 4.5 Western Blotting

Western blotting was performed as described previously (Mohammed et al., 2021). Images were taken using a Chemidoc imager (Bio-Rad) and quantified using ImageJ software (U.S. National Institutes of Health, Bethesda, MD, USA). Primary antibodies against the following proteins were used: phospho(S345) MLKL, phospho(T231+ S232)-RIPK3, HMGB1 and TNFα from Abcam (Cambridge, MA); RIPK1 and RIPK3 from Novus Biologicals (Centennial, CO); MLKL from Millipore Sigma (Burlington, MA); Phospho(Ser166)-RIPK1, Cleaved Caspase-3, Caspase 3 and Cleaved IL-1β from Cell Signaling Technology (Danvers, MA); desmin from ThermoFisher Scientific (Waltham, MA); GAPDH, β-tubulin and β-actin were from Sigma-Aldrich (St. Louis, MO). HRP-linked anti-rabbit IgG, HRP-linked anti-mouse IgG and HRP-linked anti-rat IgG from Cell Signaling Technology (Danvers, MA) were used as secondary antibodies.

### 4.6 Detection of MLKL-oligomers by western blotting

MLKL oligomerization in liver was detected by western blotting under non-reducing conditions following the protocol described by (Cai & Liu, 2018).

### 4.7 Western blotting for HMGB1

Plasma concentration of HMGB1 was analyzed by western blotting as described by Higgins et al. (Higgins et al., 2013).

### 4.8 Magnetic activated cell sorting (MACS) for liver macrophages and LSEC

The different hepatic cell populations were isolated by MACS as described by Liu et al, 2017. After isolation of hepatocyte fraction, magnetically labeled CD146 antibody (Miltenyi, CA) was added to non-parenchymal fraction and LSEC fraction was collected by passing through a magnetic column as per manufacturer’s instructions. The CD146+ fraction was incubated with biotin labeled F4/80 antibody (Miltenyi, Auburn, CA) and anti-biotin microbeads, and magnetic separation was performed to isolate the F4/80+ cell fraction. The purity of the isolated fractions was checked by real time PCR and western blotting for cell specific markers.

### 4.9 Characterization of M1/M2 macrophages by flow cytometry
Immune cell population was analyzed as described by Issac Mohar et al. (2015) with some modifications. Hepatocytes and non-parenchymal cells (NPC) were separated as described above. The pelleted NPC was purified by density gradient centrifugation using OptiPrep density gradient media (Cosmo Bio USA, Carlsbad, CA) and interphase was used for flow cytometric analysis. Live cell gating was done with Live/Dead fixable violet dead cell stain kit from Thermo Fisher Scientific (Waltham, MA). The following antibodies were used for staining (Biolegend, San Diego, CA): CD45-APC/Cy7, CD11b-PE/Cy7, Ly6G-FITC, F4/80-PE, CD80-PE/Cy5, CD206-PE Dazzle, CD16/32. Data was collected using Stratagem 4-Laser flowcytometer (San Jose, CA), and analyzed using Flow Jo software (BD Biosciences, NJ, USA) software. The gating strategy that was followed for the analysis is included in Figure S4.

4.10 Hydroxy proline (OHP) assay

OHP was performed as described by Smith et al. (2016). The OD values were converted into µg units using the 4-parameter standard curve generated using the standards and expressed as µg/g of tissue.

4.11 Picrosirius red staining

Picrosirius red staining was done using standardized protocol at the Imaging Core facility at the Oklahoma Medical Research Foundation. The images were taken using a Nikon Ti Eclipse microscope (Nikon, Melville, NY) for 3 random fields per sample and quantified using Image J software.

4.12 Immunofluorescence staining.

Immunofluorescence staining was performed as described in Thadathil et al., (2021) with modifications. Briefly, liver cryosections (10µm in thickness) were permeabilized, blocked and incubated with primary antibodies against P-MLKL, albumin (Novus Biologicals), F4/80 (Novus Biologicals) overnight at 4°C. This was followed by staining with the corresponding fluorescent tagged secondary antibodies for 1 hour at room temperature: goat anti-rabbit Alexa Fluor 647 (Thermo Fisher Scientific), donkey anti-goat Alexa Fluor 488 (Thermo Fisher Scientific), goat anti-rat Alexa Fluor 488 (Abcam). The nuclei were counter stained with DAPI (Thermo Fisher Scientific) and mounted with ProLong™ Diamond antifade mountant (Thermofisher scientific). All the imaging was acquired with a confocal microscope (Zeiss LSM 710) at 200X and 630X magnifications in five non-overlapping fields per mouse.
Paraffin sections of liver were used for immunofluorescent staining for F4/80, CD68 and CD206 as described by Lankadasari et al (PMID: 30083262). The primary antibodies used are F4/80 (Novus Biologicals), CD68 (Bio-Rad) or CD206 (Abcam) and the secondary antibodies used are goat anti-rat Alexa Fluor 488 (Abcam), donkey anti-rabbit Alexa Fluor 488 (Abcam). Nuclei were counter stained with DAPI and the sections mounted with VectaMount AQ Aqueous Mounting Medium (Vector laboratories, Burlingame, CA). Images were acquired with Nikon TE2000-E fluorescent microscope. Fluorescent intensity (percentage area of green fluorescence) was calculated using Image J software (U.S. National Institutes of Health) from 3 random fields per sample.

4.13. Administration of Nec-1s
Nec-1s administration was performed as described previously (Mohammed et al, 2021). Daily consumption of Nec-1s based on this protocol is reported to be 2.5–5 mg/day (Ito et al., 2016).

4.14. Serum alanine aminotransferase (ALT) measurement
Serum levels of ALT was measured using alanine transaminase colorimetric activity assay kit from Cayman Chemical Company (Ann Arbor, MI) following manufacturer’s instructions.

4.15. Statistical analyses
Ordinary one-way ANOVA with Tukey's post hoc test and Student’s t test was used to analyze data as indicated. P<0.05 is considered statistically significant.

ACKNOWLEDGMENTS
The authors would like to thank Claire Abbott, Aging and Metabolism Research Program, OMRF for her help in performing OHP assay; Laboratory for Molecular Biology and Cytometry Research at the University of Oklahoma Health Sciences Center for providing the facilities for the flow cytometry experiments and immunofluorescence imaging and the Imaging Core facility at the Oklahoma Medical Research Foundation for performing Picrosirius red staining and providing facilities for confocal imaging; The efforts of authors were supported by NIH grants R01AG059718 (SD), R01AG057424 (AR), Oklahoma Center for the Advancement of Science and Technology research grant (HR18-053) (SD), Presbyterian Health Foundation (OUHSC) Seed grant (SD), a Senior Career Research Award (AR) and a Merit grant I01BX004538 (AR) from the Department of Veterans Affairs, and NIH funding (R01 AG064951, R56 AG067754, R21 AR077387) to BFM.
CONFLICT OF INTEREST

The authors declare no competing financial interests.

AUTHORS’ CONTRIBUTIONS

S.M. performed the experiments, analyzed data, and prepared figures, N.T. performed immunofluorescence experiments and edited the manuscript, R.S. performed western blots for Nec-1s study, E.H.N. performed all the real-time qPCR, D.W. helped with animal studies, collaborated with B.F.M. for hydroxyproline assay, B.F.M. and A.R. gave critical comments and suggestion for the manuscript; and S.S.D designed the experiments, wrote and edited the manuscript.

DATA AVAILABILITY STATEMENT

The data that supports the findings of this study are available in the manuscript and supplementary material of this article. Correspondence and requests for information should be addressed to S.S.D.

FIGURE LEGENDS

FIGURE 1 Necroptosis markers increase with age in the livers of mice. (a) Left: Annexin V-FITC/PI staining of liver cells from 7, 12, 18, and 22 to 24-month-old mice. The upper right quadrant (FITC+/PI+) represents late apoptotic/necroptotic population, the lower right quadrant (FITC+/PI-) represents early apoptotic population and the upper left quadrant (FITC-/PI+) represents the necrotic population. Right: Graphical representation of the early apoptotic, late apoptotic/necroptotic and necrotic populations. (b) Left: Immunoblots of liver extracts prepared from 7 (white bars), 12 (grey bars), 18 (blue bars), and 22 to 24-month-old (red bars) mice for P-MLKL, MLKL and β-tubulin. Right: Graphical representation of quantified blots normalized to β-tubulin. (c) Left: Immunoblots of liver extracts for P-RIPK3, RIPK3, P-RIPK1, RIPK1 and β-tubulin. Right: Graphical representation of quantified blots normalized to β-tubulin. (d) Left: Immunostaining of liver sections from young (white bar), old (red bar) and Mlkl knockout (yellow bar) mice for P-MLKL. Scale bar: 50µM. Right: Graphical representation of the percentage of red fluorescent area. (e) Top: Immunoblots probed using anti-MLKL antibody for oligomers (>250kDa). β-tubulin was used as loading control. Bottom: Graphical representation of quantified oligomer normalized to β-tubulin. (f) Top: Immunoblots of plasma samples for HMGB1. Coomassie stained gel is used as loading control. Bottom: Graphical representation of quantified
blot normalized to Coomassie stained gel. Data represented as mean ± SEM, * p< 0.05, ** p< 0.005, *** p<0.0005, n = 5-7 per group.

FIGURE 2 Increased expression of necroptosis markers in hepatocytes isolated from old mice. (a) Left: Representative image of Annexin V/PI staining of hepatocytes isolated from young (white bars) and old mice (red bars). The upper right quadrant (FITC+/PI+) represents late apoptotic/necroptotic population, the lower right quadrant (FITC+/PI-) represents early apoptotic population and the upper left quadrant represents necrotic population. Right: Graphical representation of the early apoptotic, late apoptotic/necroptotic and necrotic populations. (b) Left: Immunoblots of hepatocytes isolated from young and old mice for P-MLKL, MLKL and GAPDH. Right: Graphical representation of quantified blot normalized to GAPDH. (c) Immunostaining for the albumin (green) and P-MLKL (red) in the livers from old mice. Arrows indicate co-localization (yellow). Scale bar: 50µM. (d) Left: Immunoblots of isolated hepatocytes from young and old mice for MLKL-oligomer. Right: Graphical representation of quantified oligomer band normalized to GAPDH. (e) Top: Immunoblots of isolated hepatocytes from young and old mice for P-RIPK3, RIPK3, P-RIPK1, RIPK1 and GAPDH. Bottom: Graphical representation of quantified blots normalized to GAPDH. (f) Immunostaining for F4/80 (green) and P-MLKL (red) in livers from old mice. Arrows indicate co-localization (yellow). Scale bar: 50µM. Data represented as mean ± SEM, * p< 0.05, ** p< 0.005, *** p<0.0005, n = 5-7/group.

FIGURE 3 Markers of inflammation increase with age in the livers of mice. (a) Left: Representative flow cytometric analysis of CD45+ cells. Right: Graphical representation of the percentage population of CD45+ cells in the liver of young (white bar) and old mice (red bar), gated on live cell population. (b) Left: Representative flow cytometric analysis of F4/80+ cells. Right: Graphical representation of the percentage population of F4/80+ cells in the livers of young and old mice, gated on live, CD45+ cells. (c) Left: Representative flow cytometric analysis of M1 macrophages (CD80+ F4/80+ cells). Right: Graphical representation of the percentage population of M1 macrophages (CD80+ F4/80+ cells) in the livers of young and old mice, gated on live, CD45+ CD11b+ Ly6G- cells. (d) Left: Representative flow cytometric analysis of M2 macrophages (CD206+ F4/80+ cells). Right panel: Graphical representation of the percentage population of M1 macrophages (CD206+ F4/80+ cells) in the livers of young and old mice, gated on live, CD45+ CD11b+ Ly6G- cells. (e) Transcript levels of TNFα, IL6, IL-1β in the livers of 7 (white bars), 12
(grey bars), 18 (blue bars), and 22 to 24-month-old (red bars) mice normalized to β-microglobulin and expressed as fold change. (f) Left: Immunoblots of liver extracts for TNFα, cleaved IL-1β and β-tubulin. Right: Graphical representation of quantified blots normalized to β-tubulin. Transcript levels of TNFα, IL6, IL-1β and MCP-1 in isolated F4/80+ cells (g) hepatocytes (h) from young and old mice normalized to β-microglobulin and expressed as fold change. Data represented as mean ± SEM, * p< 0.05, ** p< 0.005, *** p<0.0005, n = 5-7/group.

FIGURE 4 Markers of fibrosis increase with age in the livers of mice. (a) Transcript levels of TGF-β in liver of 7 (white bars), 12 (grey bars), 18 (blue bars), and 22 to 24-month-old mice (red bars) normalized to β-microglobulin and expressed as fold change. (b) Left: Immunoblots of liver extracts for desmin and β-tubulin. Right: Graphical representation of quantified blots normalized to β-tubulin. (c) Transcript levels of Col1α1, Col3α1 in livers normalized to β-microglobulin and expressed as fold change. (d) Levels of hydroxyproline in the livers, expressed as microgram of hydroxyproline/g of liver tissue. (e) Left: Picrosirius red staining in young and old mice. Scale bar: 200µm. Right: Quantification of fibrotic area in young (white bar) and old (red bar) mice. Data represented as mean ± SEM, * p< 0.05, ** p< 0.005, *** p<0.0005, n = 7-10/group.

FIGURE 5 Nec-1s treatment reduces markers of necroptosis, inflammation, fibrosis, and hepatic damage in the livers of old mice. (a) Left: Immunoblots of liver extracts from young (7-month, white bars), old (24-month, red bars), and old mice treated with Nec-1s (24 months, old-Nec-1s, green bars ) for P-MLKL, MLKL and β-tubulin. Right: Graphical representation of quantified blots normalized to β-tubulin. (b) Left: Immunoblots of non-reduced samples for MLKL-oligomer and β-tubulin. Right: Graphical representation of quantified oligomer normalized to β-tubulin. (c) Transcript levels of p16 and p21 normalized to β-microglobulin and expressed as fold change. (d) Left: Immunostaining for F4/80, CD68 and CD206 (green). Nucleus counterstained with DAPI (blue). Scale bar: 75µM. Right: Graphical representation of the percentage area of green fluorescence. (e) Left: Immunoblots of liver extracts for TNFα and β-tubulin. Right: Graphical representation of quantified blots normalized to β-tubulin. (f) ALT activity measured in the serum expressed as IU/L. (g) Levels of OHP expressed as microgram of hydroxyproline/g of liver tissue. (h) Left: Picrosirius red staining. Scale bar: 200µm. Right: Quantification of staining. Data represented as mean ± SEM, * p< 0.05, ** p< 0.005, *** p<0.0005, n = 5-7/group.
REFERENCES:

Abdelmegeed, M. A., Choi, Y., Ha, S.-K., & Song, B.-J. (2016). Cytochrome P450-2E1 promotes aging-related hepatic steatosis, apoptosis and fibrosis through increased nitroxidative stress. *Free Radical Biology & Medicine, 91*, 188–202. https://doi.org/10.1016/j.freeradbiomed.2015.12.016

Afonso, M. B., Rodrigues, P. M., Carvalho, T., Caridade, M., Borralho, P., Cortez-Pinto, H., Castro, R. E., & Rodrigues, C. M. P. (2015). Necroptosis is a key pathogenic event in human and experimental murine models of non-alcoholic steatohepatitis. *Clinical Science (London, England : 1979), 129*(8), 721–739. https://doi.org/10.1042/CS20140732

Alisi, A., Carpino, G., Oliveira, F. L., Panera, N., Nobili, V., & Gaudio, E. (2017). The Role of Tissue Macrophage-Mediated Inflammation on NAFLD Pathogenesis and Its Clinical Implications. *Mediators of Inflammation, 2017*, 8162421. https://doi.org/10.1155/2017/8162421

An, P., Wei, L.-L., Zhao, S., Sverdlov, D. Y., Vaid, K. A., Miyamoto, M., Kuramitsu, K., Lai, M., & Popov, Y. V. (2020). Hepatocyte mitochondria-derived danger signals directly activate hepatic stellate cells and drive progression of liver fibrosis. *Nature Communications, 11*(1), 2362. https://doi.org/10.1038/s41467-020-16092-0

Baar, E. L., Carbajal, K. A., Ong, I. M., & Lamming, D. W. (2016). Sex- and tissue-specific changes in mTOR signaling with age in C57BL/6J mice. *Aging Cell, 15*(1), 155–166. https://doi.org/10.1111/acel.12425

Bai, L., Liu, X., Zheng, Q., Kong, M., Zhang, X., Hu, R., Lou, J., Ren, F., Chen, Y., Zheng, S., Liu, S., Han, Y.-P., Duan, Z., & Pandol, S. J. (2017). M2-like macrophages in the fibrotic liver protect mice against lethal insults through conferring apoptosis resistance to hepatocytes. *Scientific Reports, 7*(1), 10518. https://doi.org/10.1038/s41598-017-11303-z

Beljaars, L., Schippers, M., Reker-Smit, C., Martinez, F. O., Helming, L., Poelstra, K., & Melgert, B. N. (2014). Hepatic Localization of Macrophage Phenotypes during Fibrogenesis and Resolution of Fibrosis in Mice and Humans. *Frontiers in Immunology, 5*, 430. https://doi.org/10.3389/fimmu.2014.00430

Blériot, C., Dupuis, T., Jouvion, G., Eberl, G., Disson, O., & Lecuit, M. (2015). Liver-Resident
Macrophage Necroptosis Orchestrates Type 1 Microbicidal Inflammation and Type-2-Mediated Tissue Repair during Bacterial Infection. *Immunity*, 42(1), 145–158. https://doi.org/10.1016/j.immuni.2014.12.020

Blom, K. G., Qazi, M. R., Matos, J. B. N., Nelson, B. D., DePierre, J. W., & Abedi-Valugerdi, M. (2009). Isolation of murine intrahepatic immune cells employing a modified procedure for mechanical disruption and functional characterization of the B, T and natural killer T cells obtained. *Clinical and Experimental Immunology*, 155(2), 320–329. https://doi.org/10.1111/j.1365-2249.2008.03815.x

Bokov, A., Chaudhuri, A., & Richardson, A. (2004). The role of oxidative damage and stress in aging. *Mechanisms of Ageing and Development*, 125(10–11), 811–826. https://doi.org/10.1016/j.mad.2004.07.009

Brenner, C., Galluzzi, L., Kepp, O., & Kroemer, G. (2013). Decoding cell death signals in liver inflammation. *Journal of Hepatology*, 59(3), 583–594. https://doi.org/10.1016/j.jhep.2013.03.033

Bruunsgaard, H., Andersen-Ranberg, K., Hjelmborg, J. v B., Pedersen, B. K., & Jeune, B. (2003). Elevated levels of tumor necrosis factor alpha and mortality in centenarians. *The American Journal of Medicine*, 115(4), 278–283. https://doi.org/10.1016/s0002-9343(03)00329-2

Caccamo, A., Majumder, S., Richardson, A., Strong, R., & Oddo, S. (2010). Molecular interplay between mammalian target of rapamycin (mTOR), amyloid-beta, and Tau: effects on cognitive impairments. *The Journal of Biological Chemistry*, 285(17), 13107–13120. https://doi.org/10.1074/jbc.M110.100420

Cai, Z., & Liu, Z.-G. (2018). Detection of MLKL Oligomerization During Programmed Necrosis. *Methods in Molecular Biology (Clifton, N.J.*), 1857, 85–92. https://doi.org/10.1007/978-1-4939-8754-2_8

Canli, Ö., Alanküş, Y. B., Grootjans, S., Vegi, N., Hültner, L., Hoppe, P. S., Schroeder, T., Vandenabeele, P., Bornkamm, G. W., & Greten, F. R. (2016). Glutathione peroxidase 4 prevents necroptosis in mouse erythroid precursors. *Blood*, 127(1), 139–148. https://doi.org/10.1182/blood-2015-06-654194

Chen, G. Y., & Nuñez, G. (2010). Sterile inflammation: sensing and reacting to damage. *Nature*
Chen, L., Cao, Z., Yan, L., Ding, Y., Shen, X., Liu, K., Xiang, X., Xie, Q., Zhu, C., Bao, S., & Wang, H. (2020). Circulating Receptor-Interacting Protein Kinase 3 Are Increased in HBV Patients With Acute-on-Chronic Liver Failure and Are Associated With Clinical Outcome. *Frontiers in Physiology, 11*, 526. https://doi.org/10.3389/fphys.2020.00526

Dara, L., Johnson, H., Suda, J., Win, S., Gaarde, W., Han, D., & Kaplowitz, N. (2015). Receptor interacting protein kinase 1 mediates murine acetaminophen toxicity independent of the necrosome and not through necroptosis. *Hepatology (Baltimore, Md.), 62*(6), 1847–1857. https://doi.org/10.1002/hep.27939

Davies, L. C., Jenkins, S. J., Allen, J. E., & Taylor, P. R. (2013). Tissue-resident macrophages. *Nature Immunology, 14*(10), 986–995. https://doi.org/10.1038/ni.2705

De Martinis, M., Franceschi, C., Monti, D., & Ginaldi, L. (2005). Inflamm-ageing and lifelong antigenic load as major determinants of ageing rate and longevity. *FEBS Letters, 579*(10), 2035–2039. https://doi.org/10.1016/j.febslet.2005.02.055

Deepa, S. S., Unnikrishnan, A., Matyi, S., Hadad, N., & Richardson, A. (2018). Necroptosis increases with age and is reduced by dietary restriction. *Aging Cell, 17*(4), e12770. https://doi.org/10.1111/acel.12770

Degterev, A., Huang, Z., Boyce, M., Li, Y., Jagtap, P., Mizushima, N., Cuny, G. D., Mitchison, T. J., Moskowitz, M. A., & Yuan, J. (2005). Chemical inhibitor of nonapoptotic cell death with therapeutic potential for ischemic brain injury. *Nature Chemical Biology, 1*(2), 112–119. https://doi.org/10.1038/nchembio711

Delire, B., Lebrun, V., Selvais, C., Henriet, P., Bertrand, A., Horsmans, Y., & Leclercq, I. A. (2016). Aging enhances liver fibrotic response in mice through hampering extracellular matrix remodeling. *Aging, 9*(1), 98–113. https://doi.org/10.18632/aging.101124

Dewidar, B., Meyer, C., Dooley, S., & Meindl-Beinker, A. N. (2019). TGF-β in Hepatic Stellate Cell Activation and Liver Fibrogenesis-Updated 2019. *Cells, 8*(11). https://doi.org/10.3390/cells8111419

Dondelinger, Y., Hulpiau, P., Saeys, Y., Bertrand, M. J. M., & Vandenabeele, P. (2016). An evolutionary perspective on the necroptotic pathway. *Trends in Cell Biology, 26*(10), 721–
Ferrín, G., Guerrero, M., Amado, V., Rodríguez-Perálvarez, M., & De la Mata, M. (2020). Activation of mTOR Signaling Pathway in Hepatocellular Carcinoma. *International Journal of Molecular Sciences, 21*(4). https://doi.org/10.3390/ijms21041266

Ferrucci, L., Corsi, A., Lauretani, F., Bandinelli, S., Bartali, B., Taub, D. D., Guralnik, J. M., & Longo, D. L. (2005). The origins of age-related proinflammatory state. *Blood, 105*(6), 2294–2299. https://doi.org/10.1182/blood-2004-07-2599

Franceschi, C, Bonafè, M., Valensin, S., Olivieri, F., De Luca, M., Ottaviani, E., & De Benedictis, G. (2000). Inflamm-aging. An evolutionary perspective on immunosenescence. *Annals of the New York Academy of Sciences, 908*, 244–254. https://doi.org/10.1111/j.1749-6632.2000.tb06651.x

Franceschi, Claudio, & Campisi, J. (2014). Chronic inflammation (inflammaging) and its potential contribution to age-associated diseases. *The Journals of Gerontology. Series A, Biological Sciences and Medical Sciences, 69 Suppl 1*, S4-9. https://doi.org/10.1093/gerona/glu057

Franceschi, Claudio, Capri, M., Monti, D., Giunta, S., Olivieri, F., Sevini, F., Panourgia, M. P., Invidia, L., Celani, L., Scurti, M., Cevenini, E., Castellani, G. C., & Salvioli, S. (2007). Inflammaging and anti-inflammaging: a systemic perspective on aging and longevity emerged from studies in humans. *Mechanisms of Ageing and Development, 128*(1), 92–105. https://doi.org/10.1016/j.mad.2006.11.016

Fritsch, M., Günther, S. D., Schwarzer, R., Albert, M.-C., Schorn, F., Werthenbach, J. P., Schiffmann, L. M., Stair, N., Stocks, H., Seeger, J. M., Lamkanfi, M., Krönke, M., Pasparakis, M., & Kashkar, H. (2019). Caspase-8 is the molecular switch for apoptosis, necroptosis and pyroptosis. *Nature, 575*(7784), 683–687. https://doi.org/10.1038/s41586-019-1770-6

Fulda, S. (2018). Repurposing anticancer drugs for targeting necroptosis. *Cell Cycle (Georgetown, Tex.), 17*(7), 829–832. https://doi.org/10.1080/15384101.2018.1442626

Galluzzi, L., Vitale, I., Aaronson, S. A., Abrams, J. M., Adam, D., Agostinis, P., Alnemri, E. S., Altucci, L., Amelio, I., Andrews, D. W., Annicchiarico-Petruzzelli, M., Antonov, A. V, Arama, E., Baehrecke, E. H., Barlev, N. A., Bazan, N. G., Bernassola, F., Bertrand, M. J. M., Bianchi, K., … Kroemer, G. (2018). Molecular mechanisms of cell death: recommendations
of the Nomenclature Committee on Cell Death 2018. *Cell Death and Differentiation*, 25(3), 486–541. https://doi.org/10.1038/s41418-017-0012-4

Gaskell, H., Ge, X., & Nieto, N. (2018). High-Mobility Group Box-1 and Liver Disease. *Hepatology Communications*, 2(9), 1005–1020. https://doi.org/10.1002/hep4.1223

Gautheron, J., Vucur, M., Reisinger, F., Cardenas, D. V., Roderburg, C., Koppe, C., Kreggenwinkel, K., Schneider, A. T., Bartneck, M., Neumann, U. P., Canbay, A., Reeves, H. L., Luedde, M., Tacke, F., Trautwein, C., Heikenwalder, M., & Luedde, T. (2014). A positive feedback loop between RIP3 and JNK controls non-alcoholic steatohepatitis. *EMBO Molecular Medicine*, 6(8), 1062–1074. https://doi.org/10.15252/emmm.201403856

Ghafoory, S., Varshney, R., Robison, T., Kouzbari, K., Woolington, S., Murphy, B., Xia, L., & Ahamed, J. (2018). Platelet TGF-β1 deficiency decreases liver fibrosis in a mouse model of liver injury. *Blood Advances*, 2(5), 470–480. https://doi.org/10.1182/bloodadvances.2017010868

Goldberg, E. L., & Dixit, V. D. (2015). Drivers of age-related inflammation and strategies for healthspan extension. *Immunological Reviews*, 265(1), 63–74. https://doi.org/10.1111/imr.12295

Gong, Y., Fan, Z., Luo, G., Yang, C., Huang, Q., Fan, K., Cheng, H., Jin, K., Ni, Q., Yu, X., & Liu, C. (2019). The role of necroptosis in cancer biology and therapy. *Molecular Cancer*, 18(1), 100. https://doi.org/10.1186/s12943-019-1029-8

González-Juarbe, N., Gilley, R. P., Hinojosa, C. A., Bradley, K. M., Kamei, A., Gao, G., Dube, P. H., Bergman, M. A., & Orihuela, C. J. (2015). Pore-Forming Toxins Induce Macrophage Necroptosis during Acute Bacterial Pneumonia. *PLoS Pathogens*, 11(12), e1005337. https://doi.org/10.1371/journal.ppat.1005337

Grabiner, B. C., Nardi, V., Birsoy, K., Possemato, R., Shen, K., Sinha, S., Jordan, A., Beck, A. H., & Sabatini, D. M. (2014). A diverse array of cancer-associated MTOR mutations are hyperactivating and can predict rapamycin sensitivity. *Cancer Discovery*, 4(5), 554–563. https://doi.org/10.1158/2159-8290.CD-13-0929

Hager, K., Machein, U., Krieger, S., Platt, D., Seefried, G., & Bauer, J. (1994). Interleukin-6 and selected plasma proteins in healthy persons of different ages. *Neurobiology of Aging*, 15(6),
771–772. https://doi.org/10.1016/0197-4580(94)90066-3

Hamilton, M. L., Van Remmen, H., Drake, J. A., Yang, H., Guo, Z. M., Kewitt, K., Walter, C. A., & Richardson, A. (2001). Does oxidative damage to DNA increase with age? *Proceedings of the National Academy of Sciences of the United States of America, 98*(18), 10469–10474. https://doi.org/10.1073/pnas.171202698

Hanus, J., Anderson, C., & Wang, S. (2015). RPE necroptosis in response to oxidative stress and in AMD. *Ageing Research Reviews, 24*(Pt B), 286–298. https://doi.org/10.1016/j.arr.2015.09.002

He, S., Huang, S., & Shen, Z. (2016). Biomarkers for the detection of necroptosis. *Cellular and Molecular Life Sciences : CMLS, 73*(11–12), 2177–2181. https://doi.org/10.1007/s00018-016-2192-3

Heneka, M. T., Kummer, M. P., Stutz, A., Del ekate, A., Schwartz, S., Vieira-Saecker, A., Gripp, A., Axt, D., Remus, A., Tzeng, T.-C., Gelpi, E., Halle, A., Korte, M., Latz, E., & Golenbock, D. T. (2013). NLRP3 is activated in Alzheimer’s disease and contributes to pathology in APP/PS1 mice. *Nature, 493*(7434), 674–678. https://doi.org/10.1038/nature11729

Higgins, S. J., Xing, K., Kim, H., Kain, D. C., Wang, F., Dhabangi, A., Musoke, C., Cserti-Gazdewich, C. M., Tracey, K. J., Kain, K. C., & Liles, W. C. (2013). Systemic release of high mobility group box 1 (HMGB1) protein is associated with severe and fatal Plasmodium falciparum malaria. *Malaria Journal, 12*, 105. https://doi.org/10.1186/1475-2875-12-105

Höckendorf, U., Yabal, M., Herold, T., Munkhbaatar, E., Rott, S., Jilg, S., Kauschinger, J., Magnani, G., Reisinger, F., Heuser, M., Kreipe, H., Sotlar, K., Englertner, T., Rad, R., Weichert, W., Peschel, C., Ruland, J., Heikenwalder, M., Spiikermann, K., … Jost, P. J. (2016). RIPK3 Restricts Myeloid Leukemogenesis by Promoting Cell Death and Differentiation of Leukemia Initiating Cells. *Cancer Cell, 30*(1), 75–91. https://doi.org/10.1016/j.ccell.2016.06.002

Huang, D., Zheng, X., Wang, Z.-A., Chen, X., He, W.-T., Zhang, Y., Xu, J.-G., Zhao, H., Shi, W., Wang, X., Zhu, Y., & Han, J. (2017). The MLKL Channel in Necroptosis Is an Octamer Formed by Tetramers in a Dyadic Process. *Molecular and Cellular Biology, 37*(5). https://doi.org/10.1128/MCB.00497-16
Ito, Y., Ofengeim, D., Najafov, A., Das, S., Saberi, S., Li, Y., Hitomi, J., Zhu, H., Chen, H., Mayo, L., Geng, J., Amin, P., DeWitt, J. P., Mookhtiar, A. K., Florez, M., Ouchida, A. T., Fan, J., Pasparakis, M., Kelliher, M. A., … Yuan, J. (2016). RIPK1 mediates axonal degeneration by promoting inflammation and necroptosis in ALS. *Science (New York, N.Y.)*, 353(6299), 603–608. https://doi.org/10.1126/science.aaf6803

Jia, Y., Wang, F., Guo, Q., Li, M., Wang, L., Zhang, Z., Jiang, S., Jin, H., Chen, A., Tan, S., Zhang, F., Shao, J., & Zheng, S. (2018). Curcumol induces RIPK1/RIPK3 complex-dependent necroptosis via JNK1/2-ROS signaling in hepatic stellate cells. *Redox Biology*, 19, 375–387. https://doi.org/10.1016/j.redox.2018.09.007

Jin, C. J., Baumann, A., Brandt, A., Engstler, A. J., Nier, A., Hege, M., Schmeer, C., Kehm, R., Höhn, A., Grune, T., Witte, O. W., & Bergheim, I. (2020). Aging-related liver degeneration is associated with increased bacterial endotoxin and lipopolysaccharide binding protein levels. *American Journal of Physiology. Gastrointestinal and Liver Physiology*, 318(4), G736–G747. https://doi.org/10.1152/ajpgi.00345.2018

Johnston, A. N., Ma, Y., Liu, H., Liu, S., Hanna-Addams, S., Chen, S., Chen, C., & Wang, Z. (2020). Necroptosis-blocking compound NBC1 targets heat shock protein 70 to inhibit MLKL polymerization and necroptosis. *Proceedings of the National Academy of Sciences of the United States of America*, 117(12), 6521–6530. https://doi.org/10.1073/pnas.1916503117

Karunakaran, D., Geoffrion, M., Wei, L., Gan, W., Richards, L., Shangari, P., DeKemp, E. M., Beanlands, R. A., Perisic, L., Maegdefessel, L., Hedin, U., Sad, S., Guo, L., Kolodgie, F. D., Virmani, R., Ruddy, T., & Rayner, K. J. (2016). Targeting macrophage necroptosis for therapeutic and diagnostic interventions in atherosclerosis. *Science Advances*, 2(7), e1600224. https://doi.org/10.1126/sciadv.1600224

Katz, S. C., Pillarisetty, V. G., Bleier, J. I., Shah, A. B., & DeMatteo, R. P. (2004). Liver sinusoidal endothelial cells are insufficient to activate T cells. *Journal of Immunology (Baltimore, Md. : 1950)*, 173(1), 230–235. https://doi.org/10.4049/jimmunol.173.1.230

Kim, I. H., Kisseleva, T., & Brenner, D. A. (2015). Aging and liver disease. *Current Opinion in Gastroenterology*, 31(3), 184–191. https://doi.org/10.1097/MOG.0000000000000176

Kim, W. R., Flamm, S. L., Di Bisceglie, A. M., & Bodenheimer, H. C. (2008). Serum activity of
alanine aminotransferase (ALT) as an indicator of health and disease. *Hepatology (Baltimore, Md.),* 47(4), 1363–1370. https://doi.org/10.1002/hep.22109

Kirwan, J. P., Krishnan, R. K., Weaver, J. A., Del Aguila, L. F., & Evans, W. J. (2001). Human aging is associated with altered TNF-alpha production during hyperglycemia and hyperinsulinemia. *American Journal of Physiology. Endocrinology and Metabolism,* 281(6), E1137-43. https://doi.org/10.1152/ajpendo.2001.281.6.E1137

Kondylis, V., Kumari, S., Vlantis, K., & Pasparakis, M. (2017). The interplay of IKK, NF-κB and RIPK1 signaling in the regulation of cell death, tissue homeostasis and inflammation. *Immunological Reviews,* 277(1), 113–127. https://doi.org/10.1111/imr.12550

Koschel, J., Nishanth, G., Just, S., Harit, K., Kröger, A., Deckert, M., Naumann, M., & Schlüter, D. (2021). OTUB1 prevents lethal hepatocyte necroptosis through stabilization of c-IAP1 during murine liver inflammation. *Cell Death and Differentiation.* https://doi.org/10.1038/s41418-021-00752-9

Koyama, Y., & Brenner, D. A. (2017). Liver inflammation and fibrosis. *The Journal of Clinical Investigation,* 127(1), 55–64. https://doi.org/10.1172/JCI88881

Lankadasari, M. B., Aparna, J. S., Mohammed, S., James, S., Aoki, K., Binu, V. S., Nair, S., & Harikumar, K. B. (2018). Targeting S1PR1/STAT3 loop abrogates desmoplasia and chemosensitizes pancreatic cancer to gemcitabine. *Theranostics,* 8(14), 3824–3840. https://doi.org/10.7150/thno.25308

Lee, C., Jeong, H., Bae, Y., Shin, K., Kang, S., Kim, H., Oh, J., & Bae, H. (2019). Targeting of M2-like tumor-associated macrophages with a melittin-based pro-apoptotic peptide. *Journal for Immunotherapy of Cancer,* 7(1), 147. https://doi.org/10.1186/s40425-019-0610-4

Lee, H., Fessler, M. B., Qu, P., Heymann, J., & Kopp, J. B. (2020). Macrophage polarization in innate immune responses contributing to pathogenesis of chronic kidney disease. *BMC Nephrology,* 21(1), 270. https://doi.org/10.1186/s12882-020-01921-7

Lee, U. E., & Friedman, S. L. (2011). Mechanisms of hepatic fibrogenesis. *Best Practice & Research. Clinical Gastroenterology,* 25(2), 195–206. https://doi.org/10.1016/j.bpg.2011.02.005

Liang, S., Lv, Z.-T., Zhang, J.-M., Wang, Y.-T., Dong, Y.-H., Wang, Z.-G., Chen, K., Cheng, P.,
Yang, Q., Guo, F.-J., Lu, W.-W., Zhu, W.-T., & Chen, A.-M. (2018). Necrostatin-1 Attenuates Trauma-Induced Mouse Osteoarthritis and IL-1β Induced Apoptosis via HMGB1/TLR4/SDF-1 in Primary Mouse Chondrocytes. *Frontiers in Pharmacology*, 9, 1378. https://doi.org/10.3389/fphar.2018.01378

Lin, B., Jin, Z., Chen, X., Zhao, L., Weng, C., Chen, B., Tang, Y., & Lin, L. (2020). Necrostatin-1 protects mice from acute lung injury by suppressing necroptosis and reactive oxygen species. *Molecular Medicine Reports*, 21(5), 2171–2181. https://doi.org/10.3892/mmr.2020.11010

Liu, J., Huang, X., Werner, M., Broering, R., Yang, D., & Lu, M. (2017). Advanced Method for Isolation of Mouse Hepatocytes, Liver Sinusoidal Endothelial Cells, and Kupffer Cells. *Methods in Molecular Biology (Clifton, N.J.)*, 1540, 249–258. https://doi.org/10.1007/978-1-4939-6700-1_21

Liu, Q., Qiu, J., Liang, M., Golinski, J., van Leyen, K., Jung, J. E., You, Z., Lo, E. H., Degterev, A., & Whalen, M. J. (2014). Akt and mTOR mediate programmed necrosis in neurons. *Cell Death & Disease*, 5(2), e1084. https://doi.org/10.1038/cddis.2014.69

Lomphithak, T., Akara-Amornthum, P., Murakami, K., Hashimoto, M., Usubuchi, H., Iwabuchi, E., Unno, M., Cai, Z., Sasano, H., & Jitkaew, S. (2021). Tumor necroptosis is correlated with a favorable immune cell signature and programmed death-ligand 1 expression in cholangiocarcinoma. *Scientific Reports*, 11(1), 11743. https://doi.org/10.1038/s41598-021-89977-9

Lozada-Delgado, J. G., Torres-Ramos, C. A., & Ayala-Peña, S. (2020). *Chapter 4 - Aging, oxidative stress, mitochondrial dysfunction, and the liver* (V. R. Preedy & V. B. B. T.-A. (Second E. Patel (eds.); pp. 37–46). Academic Press. https://doi.org/10.1016/B978-0-12-818698-5.00004-3

Lu, C., Xu, W., Zhang, F., Shao, J., & Zheng, S. (2016). Nrf2 Knockdown Disrupts the Protective Effect of Curcumin on Alcohol-Induced Hepatocyte Necroptosis. *Molecular Pharmaceutics*, 13(12), 4043–4053. https://doi.org/10.1021/acs.molpharmaceut.6b00562

Lynch, R. W., Hawley, C. A., Pellicoro, A., Bain, C. C., Iredale, J. P., & Jenkins, S. J. (2018). An efficient method to isolate Kupffer cells eliminating endothelial cell contamination and selective bias. *Journal of Leukocyte Biology*, 104(3), 579–586.
Majdi, A., Aoudjehane, L., Ratziu, V., Islam, T., Afonso, M. B., Conti, F., Mestiri, T., Lagouge, M., Foufelle, F., Ballenghien, F., Ledent, T., Moldes, M., Cadorret, A., Fouassier, L., Delaunay, J.-L., Aït-Slimane, T., Courtois, G., Fève, B., Scatton, O., … Gautheron, J. (2020). Inhibition of receptor-interacting protein kinase 1 improves experimental non-alcoholic fatty liver disease. *Journal of Hepatology, 72*(4), 627–635. https://doi.org/10.1016/j.jhep.2019.11.008

Malato, Y., Naqvi, S., Schürmann, N., Ng, R., Wang, B., Zape, J., Kay, M. A., Grimm, D., & Willenbring, H. (2011). Fate tracing of mature hepatocytes in mouse liver homeostasis and regeneration. *The Journal of Clinical Investigation, 121*(12), 4850–4860. https://doi.org/10.1172/JCI59261

Mancuso, P., & Bouchard, B. (2019). The Impact of Aging on Adipose Function and Adipokine Synthesis. *Frontiers in Endocrinology, 10*, 137. https://doi.org/10.3389/fendo.2019.00137

McNamara, C. R., Ahuja, R., Osafo-Addo, A. D., Barrows, D., Kettenbach, A., Skidan, I., Teng, X., Cuny, G. D., Gerber, S., & Degterev, A. (2013). Akt Regulates TNFα synthesis downstream of RIP1 kinase activation during necroptosis. *PloS One, 8*(3), e56576. https://doi.org/10.1371/journal.pone.0056576

Meng, Y., Sandow, J. J., Czabotar, P. E., & Murphy, J. M. (2021). The regulation of necroptosis by post-translational modifications. *Cell Death and Differentiation, 28*(3), 861–883. https://doi.org/10.1038/s41418-020-00722-7

Mifflin, L., Ofengeim, D., & Yuan, J. (2020). Receptor-interacting protein kinase 1 (RIPK1) as a therapeutic target. *Nature Reviews. Drug Discovery, 19*(8), 553–571. https://doi.org/10.1038/s41573-020-0071-y

Mohammed, S., Nicklas, E. H., Thadathil, N., Selvarani, R., Royce, G. H., Kinter, M., Richardson, A., & Deepa, S. S. (2021). Role of necroptosis in chronic hepatic inflammation and fibrosis in a mouse model of increased oxidative stress. *Free Radical Biology & Medicine, 164*, 315–328. https://doi.org/10.1016/j.freeradbiomed.2020.12.449

Mohammed, S., Vineetha, N. S., James, S., Aparna, J. S., Lankadasari, M. B., Allegood, J. C., Li, Q.-Z., Spiegel, S., & Harikumar, K. B. (2019). Examination of the role of sphingosine kinase
2 in a murine model of systemic lupus erythematosus. *FASEB Journal: Official Publication of the Federation of American Societies for Experimental Biology*, 33(6), 7061–7071. https://doi.org/10.1096/fj.201802535R

Mohar, I., Brempelis, K. J., Murray, S. A., Ebrahimkhani, M. R., & Crispe, I. N. (2015). Isolation of Non-parenchymal Cells from the Mouse Liver. *Methods in Molecular Biology (Clifton, N.J.)*, 1325, 3–17. https://doi.org/10.1007/978-1-4939-2815-6_1

Morsiani, C., Bacalini, M. G., Santoro, A., Garagnani, P., Collura, S., D’Errico, A., de Eguileor, M., Grazi, G. L., Cescon, M., Franceschi, C., & Capri, M. (2019). The peculiar aging of human liver: A geroscience perspective within transplant context. *Ageing Research Reviews*, 51, 24–34. https://doi.org/10.1016/j.arr.2019.02.002

Najafov, A., Mookhtiar, A. K., Luu, H. S., Ordureau, A., Pan, H., Amin, P. P., Li, Y., Lu, Q., & Yuan, J. (2019). TAM Kinases Promote Necroptosis by Regulating Oligomerization of MLKL. *Molecular Cell*, 75(3), 457-468.e4. https://doi.org/10.1016/j.molcel.2019.05.022

Newton, K., & Manning, G. (2016). Necroptosis and Inflammation. *Annual Review of Biochemistry*, 85, 743–763. https://doi.org/10.1146/annurev-biochem-060815-014830

Owens, T. (2009). Toll-like receptors in neurodegeneration. *Current Topics in Microbiology and Immunology*, 336, 105–120. https://doi.org/10.1007/978-3-642-00549-7_6

Pedersen, B. K., Steensberg, A., Keller, P., Keller, C., Fischer, C., Hiscock, N., van Hall, G., Plomgaard, P., & Febbraio, M. A. (2003). Muscle-derived interleukin-6: lipolytic, anti-inflammatory and immune regulatory effects. *Pflugers Archiv: European Journal of Physiology*, 446(1), 9–16. https://doi.org/10.1007/s00424-002-0981-z

Pietkiewicz, S., Schmidt, J. H., & Lavrik, I. N. (2015). Quantification of apoptosis and necroptosis at the single cell level by a combination of Imaging Flow Cytometry with classical Annexin V/propidium iodide staining. *Journal of Immunological Methods*, 423, 99–103. https://doi.org/10.1016/j.jim.2015.04.025

Puche, J. E., Saiman, Y., & Friedman, S. L. (2013). Hepatic stellate cells and liver fibrosis. *Comprehensive Physiology*, 3(4), 1473–1492. https://doi.org/10.1002/cphy.c120035

Ramachandran, P., Pellicoro, A., Vernon, M. A., Boulter, L., Aucott, R. L., Ali, A., Hartland, S. N., Snowdon, V. K., Cappon, A., Gordon-Walker, T. T., Williams, M. J., Dunbar, D. R.,
Manning, J. R., Van Rooijen, N., Fallowfield, J. A., Forbes, S. J., & Iredale, J. P. (2012). Differential Ly-6C expression identifies the recruited macrophage phenotype, which orchestrates the regression of murine liver fibrosis. *Proceedings of the National Academy of Sciences of the United States of America, 109*(46). https://doi.org/10.1073/pnas.1119964109

Rock, K. L., Lai, J.-J., & Kono, H. (2011). Innate and adaptive immune responses to cell death. *Immunological Reviews, 243*(1), 191–205. https://doi.org/10.1111/j.1600-065X.2011.01040.x

Rodríguez-Gómez, J. A., Kavanagh, E., Engskog-Vlachos, P., Engskog, M. K. R., Herrera, A. J., Espinosa-Oliva, A. M., Joseph, B., Hajji, N., Venero, J. L., & Burguillos, M. A. (2020). Microglia: Agents of the CNS Pro-Inflammatory Response. *Cells, 9*(7). https://doi.org/10.3390/cells9071717

Roubenoff, R., Harris, T. B., Abad, L. W., Wilson, P. W., Dallal, G. E., & Dinarello, C. A. (1998). Monocyte cytokine production in an elderly population: effect of age and inflammation. *The Journals of Gerontology. Series A, Biological Sciences and Medical Sciences, 53*(1), M20-6. https://doi.org/10.1093/gerona/53a.1.m20

Rowell, D. L., Eckmann, L., Dwinell, M. B., Carpenter, S. P., Raucy, J. L., Yang, S. K., & Kagnoff, M. F. (1997). Human hepatocytes express an array of proinflammatory cytokines after agonist stimulation or bacterial invasion. *The American Journal of Physiology, 273*(2 Pt 1), G322-32. https://doi.org/10.1152/ajpgi.1997.273.2.G322

Royce, G. H., Brown-Borg, H. M., & Deepa, S. S. (2019). The potential role of necroptosis in inflammaging and aging. In *GeroScience* (Vol. 41, Issue 6, pp. 795–811). Springer. https://doi.org/10.1007/s11357-019-00131-w

Samson, A. L., Zhang, Y., Geoghegan, N. D., Gavin, X. J., Davies, K. A., Mlodzianoski, M. J., Whitehead, L. W., Frank, D., Garnish, S. E., Fitzgibbon, C., Hempel, A., Young, S. N., Jacobsen, A. V., Cawthorne, W., Petrie, E. J., Faux, M. C., Shield-Artin, K., Lalaoui, N., Hildebrand, J. M., … Murphy, J. M. (2020). MLKL trafficking and accumulation at the plasma membrane control the kinetics and threshold for necroptosis. *Nature Communications, 11*(1), 3151. https://doi.org/10.1038/s41467-020-16887-1

Schon, H.-T., & Weiskirchen, R. (2014). Immunomodulatory effects of transforming growth
factor-β in the liver. *Hepatobiliary Surgery and Nutrition*, 3(6), 386–406. https://doi.org/10.3978/j.issn.2304-3881.2014.11.06

Schwartz, R. S., Shuman, W. P., Bradbury, V. L., Cain, K. C., Fellingham, G. W., Beard, J. C., Kahn, S. E., Stratton, J. R., Cerqueira, M. D., & Abrass, I. B. (1990). Body fat distribution in healthy young and older men. *Journal of Gerontology, 45*(6), M181-5. https://doi.org/10.1093/geronj/45.6.m181

Scott, C. L., Zheng, F., De Baetselier, P., Martens, L., Saeys, Y., De Prijck, S., Lippens, S., Abels, C., Schoonooghe, S., Raes, G., Devoogdt, N., Lambrecht, B. N., Beschin, A., & Guiliams, M. (2016). Bone marrow-derived monocytes give rise to self-renewing and fully differentiated Kupffer cells. *Nature Communications*, 7, 10321. https://doi.org/10.1038/ncomms10321

Seifert, L., Werba, G., Tiwari, S., Giao Ly, N. N., Alothman, S., Alqunaibit, D., Avanzi, A., Barilla, R., Daley, D., Greco, S. H., Torres-Hernandez, A., Pergamo, M., Ochi, A., Zambirinis, C. P., Pansari, M., Rendon, M., Tippens, D., Hundeyin, M., Mani, V. R., … Miller, G. (2016). The necrosome promotes pancreatic oncogenesis via CXCL1 and Mincle-induced immune suppression. *Nature*, 532(7598), 245–249. https://doi.org/10.1038/nature17403

Shen, B., Mei, M., Pu, Y., Zhang, H., Liu, H., Tang, M., Pan, Q., He, Y., Wu, X., & Zhao, H. (2019). Necrostatin-1 Attenuates Renal Ischemia and Reperfusion Injury via Meditation of HIF-1α/mir-26a/TRPC6/PARP1 Signaling. *Molecular Therapy. Nucleic Acids*, 17, 701–713. https://doi.org/10.1016/j.omtn.2019.06.025

Shojaie, L., Iorga, A., & Dara, L. (2020). Cell Death in Liver Diseases: A Review. *International Journal of Molecular Sciences*, 21(24). https://doi.org/10.3390/ijms21249682

Smith, L. R., Hammers, D. W., Sweeney, H. L., & Barton, E. R. (2016). Increased collagen cross-linking is a signature of dystrophin-deficient muscle. *Muscle & Nerve*, 54(1), 71–78. https://doi.org/10.1002/mus.24998

Stahl, E. C., Delgado, E. R., Alencastro, F., LoPresti, S. T., Wilkinson, P. D., Roy, N., Haschak, M. J., Skillen, C. D., Monga, S. P., Duncan, A. W., & Brown, B. N. (2020). Inflammation and Ectopic Fat Deposition in the Aging Murine Liver Is Influenced by CCR2. *The American
Journal of Pathology, 190(2), 372–387. https://doi.org/10.1016/j.ajpath.2019.10.016

Stoll, G., Ma, Y., Yang, H., Kepp, O., Zitvogel, L., & Kroemer, G. (2017). Pro-necrotic molecules impact local immunosurveillance in human breast cancer. Oncoimmunology, 6(4), e1299302. https://doi.org/10.1080/2162402X.2017.1299302

Takahashi, N., Duprez, L., Grootjans, S., Cauwels, A., Nerinckx, W., Duhadaway, J. B., Goossens, V., Roelandt, R., Van Hauwermeiren, F., Libert, C., Declercq, W., Callewaert, N., Prendergast, G. C., Degterev, A., Yuan, J., & Vandenabeele, P. (2012). Necrostatin-1 analogues: Critical issues on the specificity, activity and in vivo use in experimental disease models. Cell Death and Disease, 3(11), e437-10. https://doi.org/10.1038/cddis.2012.176

Thadathil, N., Nicklas, E. H., Mohammed, S., Lewis, T. L. J., Richardson, A., & Deepa, S. S. (2021). Necroptosis increases with age in the brain and contributes to age-related neuroinflammation. GeroScience. https://doi.org/10.1007/s11357-021-00448-5

Völkers, M., Doroudgar, S., Nguyen, N., Konstandin, M. H., Quijada, P., Din, S., Ornelas, L., Thuerauf, D. J., Gude, N., Friedrich, K., Herzig, S., Glembotski, C. C., & Sussman, M. A. (2014). PRAS40 prevents development of diabetic cardiomyopathy and improves hepatic insulin sensitivity in obesity. EMBO Molecular Medicine, 6(1), 57–65. https://doi.org/10.1002/emmm.201303183

Wang, X.-K., & Peng, Z.-G. (2021). Targeting Liver Sinusoidal Endothelial Cells: An Attractive Therapeutic Strategy to Control Inflammation in Nonalcoholic Fatty Liver Disease. Frontiers in Pharmacology, 12, 655557. https://doi.org/10.3389/fphar.2021.655557

Wei, G., An, P., Vaid, K. A., Nasser, I., Huang, P., Tan, L., Zhao, S., Schuppan, D., & Yury Popov, X. V. (2020). Comparison of murine steatohepatitis models identifies a dietary intervention with robust fibrosis, ductular reaction, and rapid progression to cirrhosis and cancer. Am J Physiol Gastrointest Liver Physiol, 318, 174–188. https://doi.org/10.1152/ajpgi.00441.2019.

Wei, G., An, P., Vaid, K. A., Nasser, I., Huang, P., Tan, L., Zhao, S., Schuppan, D., & Yury Popov, X. V. (2020). Comparison of murine steatohepatitis models identifies a dietary intervention with robust fibrosis, ductular reaction, and rapid progression to cirrhosis and cancer. Am J Physiol Gastrointest Liver Physiol, 318, 174–188. https://doi.org/10.1152/ajpgi.00441.2019.

Progressive

Wen, S., Li, X., Ling, Y., Chen, S., Deng, Q., Yang, L., Li, Y., Shen, J., Qiu, Y., Zhan, Y., Lai, H., Zhang, X., Ke, Z., & Huang, W. (2020). HMGB1-associated necroptosis and Kupffer cells M1 polarization underlies remote liver injury induced by intestinal ischemia/reperfusion in rats. FASEB Journal : Official Publication of the Federation of American Societies for
Wen, Y., Lambrecht, J., Ju, C., & Tacke, F. (2021). Hepatic macrophages in liver homeostasis and diseases-diversity, plasticity and therapeutic opportunities. *Cellular & Molecular Immunology, 18*(1), 45–56. https://doi.org/10.1038/s41423-020-00558-8

Wu, X., & Nagy, L. E. (2020). MLKL contributes to Western diet-induced liver injury through inhibiting autophagy. *Autophagy, 16*(7), 1351–1352. https://doi.org/10.1080/15548627.2020.1760624

Xiao, H. (2018). Epithelial TSC1-mTOR controls RIPK3-dependent necroptosis and susceptibility to IBD by sensing dietary and microbial metabolites. *The Journal of Immunology, 200*(1 Supplement), 53.12–LP-53.12. www.jimmunol.org/content/200/1_Supplement/53.12.abstract

Youm, Y.-H., Grant, R. W., McCabe, L. R., Albarado, D. C., Nguyen, K. Y., Ravussin, A., Pistell, P., Newman, S., Carter, R., Laque, A., Münzberg, H., Rosen, C. J., Ingram, D. K., Salbaum, J. M., & Dixit, V. D. (2013). Canonical Nlrp3 inflammasome links systemic low-grade inflammation to functional decline in aging. *Cell Metabolism, 18*(4), 519–532. https://doi.org/10.1016/j.cmet.2013.09.010

Yuan, J., Amin, P., & Ofengeim, D. (2019). Necroptosis and RIPK1-mediated neuroinflammation in CNS diseases. *Nature Reviews. Neuroscience, 20*(1), 19–33. https://doi.org/10.1038/s41583-018-0093-1

Zelic, M., Roderick, J. E., O’Donnell, J. A., Lehman, J., Lim, S. E., Janardhan, H. P., Trivedi, C. M., Pasparakis, M., & Kelliher, M. A. (2018). RIP kinase 1-dependent endothelial necroptosis underlies systemic inflammatory response syndrome. *The Journal of Clinical Investigation, 128*(5), 2064–2075. https://doi.org/10.1172/JCI96147

Zhang, B., Li, M.-D., Yin, R., Liu, Y., Yang, Y., Mitchell-Richards, K. A., Nam, J. H., Li, R., Wang, L., Iwakiri, Y., Chung, D., Robert, M. E., Ehrlich, B. E., Bennett, A. M., Yu, J., Nathanson, M. H., & Yang, X. (2019a). O-GlcNAc transferase suppresses necroptosis and liver fibrosis. *JCI Insight, 4*(21). https://doi.org/10.1172/jci.insight.127709

Zhang, B., Li, M. D., Yin, R., Liu, Y., Yang, Y., Mitchell-Richards, K. A., Nam, J. H., Li, R., Wang, L., Iwakiri, Y., Chung, D., Robert, M. E., Ehrlich, B. E., Bennett, A. M., Yu, J.,
Nathanson, M. H., & Yang, X. (2019b). O-GlcNAc transferase suppresses necroptosis and liver fibrosis. *JCI Insight, 4*(21). https://doi.org/10.1172/jci.insight.127709

Zhang, L., Feng, Q., & Wang, T. (2018). Necrostatin-1 Protects Against Paraquat-Induced Cardiac Contractile Dysfunction via RIP1-RIP3-MLKL-Dependent Necroptosis Pathway. *Cardiovascular Toxicology, 18*(4), 346–355. https://doi.org/10.1007/s12012-017-9441-z

Zhang, S., Su, Y., Ying, Z., Guo, D., Pan, C., Guo, J., Zou, Z., Wang, L., Zhang, Z., Jiang, Z., Zhang, Z., & Wang, X. (2019). RIP1 kinase inhibitor halts the progression of an immune-induced demyelination disease at the stage of monocyte elevation. *Proceedings of the National Academy of Sciences of the United States of America, 116*(12), 5675–5680. https://doi.org/10.1073/pnas.1819917116

Zhang, Y., Unnikrishnan, A., Deepa, S. S., Liu, Y., Li, Y., Ikeno, Y., Sosnowska, D., Van Remmen, H., & Richardson, A. (2017). A new role for oxidative stress in aging: The accelerated aging phenotype in Sod1(-/-) mice is correlated to increased cellular senescence. *Redox Biology, 11*, 30–37. https://doi.org/10.1016/j.redox.2016.10.014

Zhong, W., Wang, X., Rao, Z., Pan, X., Sun, Y., Jiang, T., Wang, P., Zhou, H., & Wang, X. (2020). Aging aggravated liver ischemia and reperfusion injury by promoting hepatocyte necroptosis in an endoplasmic reticulum stress-dependent manner. *Annals of Translational Medicine, 8*(14), 869. https://doi.org/10.21037/atm-20-2822

**SUPPLEMENTARY FIGURES:**

FIGURE S1 (a) Transcript levels of *Mlkl, Ripk3, Ripk1* in the livers of 7, 12, 18, and 22 to 24-month-old mice normalized to β-microglobulin and expressed as fold change. (b) Top panel: Immunoblots of whole liver extracts (white bars) and isolated hepatocytes (grey bars) for Albumin, Desmin, CD31 and GAPDH. Bottom panel: Graphical representation of the quantified blots normalized to GAPDH. (c) Transcript levels of Albumin, F4/80, Clec4f and CD31 in the isolated hepatocyte fraction normalized to β-microglobulin and expressed as fold change. (d) Transcript levels of *Mlkl, Ripk3 and Ripk1* in hepatocytes from young (white bars) and old mice (red bars) normalized to β-microglobulin and expressed as fold change. (e) Transcript level of MLKL in the F4/80+ cells isolated from young (white bar) and old mice (red bar). (f) Graphical representation of the early apoptotic and late apoptotic/necroptotic population in the annexin/propidium iodide
staining of LSEC and KC fraction obtained by MACS (g) Transcript levels of albumin (hepatocyte marker), F4/80 (macrophage marker), stabilin and CD31 (endothelial markers) in the cell fractions isolated by MACS. Data represented as mean ± SEM, * p< 0.05, ** p< 0.005, *** p<0.0005, n = 3/group.

FIGURE S2 Transcript levels of (a) F4/80 and MCP1, (b) CD68, CD86, TLR4 and CD11c (c) Arg1 and Fizz1 in the livers of 7 (white bars), 12 (grey bars), 18 (blue bars), and 22 to 24-month-old (red bars) mice normalized to β-microglobulin and expressed as fold change. (d) Transcript levels of TNFα, IL6, IL-1β and MCP-1 in LSEC isolated from young (white bars) and old (red bars). Data represented as mean ± SEM, * p< 0.05, ** p< 0.005, *** p<0.0005, n = 5-7/group.

FIGURE S3 (a) The body weight and (b) percentage liver weight of young (7 months, white bars), old (24 months, red bars) and old mice treated with Nec-1s (24 months, green bars). (c) Left panel: Immunoblots of liver extracts prepared from young, old and old mice treated with Nec-1s for cleaved caspase-3, caspase 3 and β-tubulin. Right panel: Graphical representation of quantified blots normalized to β-tubulin. (d-h) Transcript levels of (d) F4/80 and MCP1, (e) CD68, CD86, TLR4, and CD11C, and (f) Arg1, Fizz1 (g) TNFα, IL6, IL-1β (h) Col1α1, Col3α1 in the livers of young, old, and old-Nec-1s mice normalized to β-microglobulin and expressed as fold change. Data represented as mean ± SEM, * p< 0.05, ** p< 0.005, *** p<0.0005, n = 5-7/group.

Figure S4 Gating strategy that was followed for the flow cytometry experiments.

| Gene                     | Forward sequence         | Reverse sequence         |
|--------------------------|--------------------------|--------------------------|
| Albumin                  | 5'-GCGCAGATGACAGGCGGAA-3' | 5'-GTGCCGTAGCATGCGGAG-3' |
| Arg1                     | 5'-CTCCAAGCCAAAGTCTTAGAG-3' | 5'-AGGAGCTGTCATTGAGGACATC-3' |
| CD11C                    | 5'-CTGGATAGCCTTCTTCTGTG-3' | 5'-GCACACTGTGTCCGAACTC-3' |
| CD31                     | 5'-CTGGTGCTCTATGCAAGC-3'  | 5'-AGTGCTGCCCATTCACTC-3' |
| CD68                     | 5'-CCACAGGCAAGACAGTG-3'  | 5'-TCCACAGCAGACTTGGCCC-3' |
| CD86                     | 5'-ACGATGGACCCCAGATG-3'  | 5'-GCACGTCCAGGAAACAGCA-3' |
| Cdkn1a(p21Cip1)          | 5'-GTCAGGCTGTCTGCCTCCG-3' | 5'-CGGTCCCAGGACAGTGAAG-3' |
| Cdkn2a(p16INK4a)         | 5'-CCACACCGCCCGAACCT-3'  | 5'-GCAGAAGAGCTGCTACGGA-3' |
| Gene         | Forward Primer Sequence | Reverse Primer Sequence |
|--------------|--------------------------|-------------------------|
| Clec4f       | 5′-CTTCGGGGAAGCAACAACTC-3′ | 5′-CAAGCAACTGCACCAGAGAAG-3′ |
| Colla1       | 5′-GCTCCTCTTAGGCGGACT-3′  | 5′-CCACGTCTTACATGGG-3′    |
| Col3a1       | 5′-CTGTAACATGGAACAGGGGGA-3′ | 5′-CCATAGCTGAACTGAAACCACC-3′ |
| F4/80        | 5′-CCCCAGTGTTCTACAGAGTG-3′ | 5′-GTGCCAGATGGATGCT-3′    |
| Fizz1        | 5′-CCAATCAGCTACTATCCATTCC-3′ | 5′-CCAGTCAACAGGAAGACACAG-3′ |
| IL-6         | 5′-TGTTACTCCAGAAGACAGAGG-3′ | 5′-AAGCATGATGCACCTGGAGA-3′ |
| Il-1β        | 5′-AGGTCAAGGTTGGAAGCA-3′  | 5′-TGAAGCAGCTATGGGCAACT-3′ |
| MCP-1        | 5′-TAAACACCTGAGAAGCCAA-3′  | 5′-GCCATAGCTTCAGATTACCGGT-3′ |
| MLKL         | 5′-CTGAGGGAACGTGGATAGAG-3′  | 5′-CGAGAAACTGGAAGCTGCT-3′ |
| RIPK3        | 5′-GAAGACCGGACTCTTGTA-3′   | 5′-CTTGAGGAGCACTTGCT-3′   |
| RIPK1        | 5′-GACTGCTGACCTCTACCTCCG-3′ | 5′-CAGGGATCATCCAGGACGTA-3′ |
| Stabilin     | 5′-GTTCAGACGGTACATCAA-3′   | 5′-CCAGGGATATCCAGGACGTA-3′ |
| TGFβ         | 5′-ACCATGCAAACCTTGCTGGGAC-3′ | 5′-ACAAGTCTCCACCTGGCTTG-3′ |
| TLR4         | 5′-ATGGCATGGCTTACACACC-3′  | 5′-GAGGCAATTTGCTCCACA-3′  |
| TNFα         | 5′-CACAGAAAGCATGATCCGGACGT-3′ | 5′-CGGCAGAGAGGAGGAGTTGACTT-3′ |
| β-actin      | 5′-ATGGATGACGATATCGTG-3′   | 5′-GGGTGCCGTTGATAGATCTTG-3′ |
| β-microglobulin | 5′-CAGGACCAGACTGCTGAT-3′   | 5′-TGAGTACTCATTAGCAGGGCA-3′ |
Figure 1

a) Propidium iodide (PI) and Annexin-FITC staining over different ages (7 months to 24 months).

b) Western blot analysis showing P-MLKL, MLKL, and β-tubulin levels across different ages.

c) Western blot analysis showing P-RIPK3, RIPK3, P-RIPK1, RIPK1, and β-tubulin levels across different ages.

d) fluorescence images and percentage of red fluorescence area of young, old, and Mlkl−/− mice.

e) Fold change of oligomer to β-tubulin levels at different ages.

f) HMGB1 and Coomassie Blue staining over different ages (7 months to 24 months).
Figure 2

(a) Propidium iodide (PI) and Annexin - FITC analysis showing the percentage of population in different quadrants (Q1, Q2, Q3, Q4) for young and old groups.

(b) Western blot analysis showing the protein expression levels of P-MLKL, MLKL, and GAPDH for young and old groups.

(c) Immunofluorescence images of Albumin, P-MLKL, MERGED DAPI, and DAPI showing cellular localization.

(d) Western blot analysis showing the protein expression levels of Monomer, Tetramer, and Oligomer for young and old groups.

(e) Western blot analysis showing the protein expression levels of P-RIPK3, RIPK3, P-RIPK1, RIPK1, and GAPDH for young and old groups.

(f) Immunofluorescence images of F4/80, P-MLKL, MERGED DAPI, and DAPI showing cellular localization.

(Figure 2)
FIGURE 3

(a) CD45+ cells

Young: 32.86 ± 0.92
Old: 67.9 ± 6.84

(b) F4/80+ cells

Young: 16.6 ± 1.6
Old: 27.9 ± 0.92

(c) M1 Macrophages

Young: 2.45 ± 0.18
Old: 9.03 ± 0.59

(d) M2 Macrophages

Young: 6.48 ± 0.56
Old: 5.06 ± 0.12

(e) mRNA levels

TNFα, IL6, IL-1β, MCP1

(f) Protein levels

Active TNFα (52kD), TNFα (25kD), Cleaved IL1β

(g) mRNA level

TNFα, IL6, IL-1β, MCP1

(h) mRNA level

TNFα, IL6, IL-1β, MCP1
Hepatocytes

Kupffer cells

Age

Necroptosis

Nec-1s

M1 Macrophages

Pro-Inflammatory Cytokines

Activated Stellate Cells

Fibrosis

Graphical abstract