Metabolically Activated Adipose Tissue Macrophages Perform Detrimental and Beneficial Functions during Diet-Induced Obesity

Graphical Abstract

Highlights

- Inflammation and dead adipocyte clearance by MMe macrophages require NOX2

- \(\text{Nox}^2^{-/-}\) improves the metabolic phenotype in early DIO but worsens it in late DIO

- Early improvements associate with suppressed ATM inflammation

- Late worsening associates with lower ATM lysosomal exocytosis to dead adipocytes

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In Brief

During obesity, adipose tissue macrophages are metabolically activated (MMe). Coats et al. show that MMe macrophages perform detrimental (potentiate inflammation) and beneficial (exocytose lysosomes to clear dead adipocytes) functions, controlled by NOX2. \(\text{Nox}^2^{-/-}\) mice exhibit improved or worsened metabolic phenotypes depending on high-fat-diet duration, highlighting the dynamic contributions of MMe macrophages in obesity.

Coats et al., 2017, Cell Reports 20, 3149–3161

September 26, 2017 © 2017 The Authors.

http://dx.doi.org/10.1016/j.celrep.2017.08.096
Metabolically Activated Adipose Tissue Macrophages Perform Detrimental and Beneficial Functions during Diet-Induced Obesity

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SUMMARY

During obesity, adipose tissue macrophages (ATMs) adopt a metabolically activated (MMe) phenotype. However, the functions of MMe macrophages are poorly understood. Here, we combine proteomic and functional methods to demonstrate that, in addition to potentiating inflammation, MMe macrophages promote dead adipocyte clearance through lysosomal exocytosis. We identify NADPH oxidase 2 (NOX2) as a driver of the inflammatory and adipocyte-clearing properties of MMe macrophages and show that, compared to wild-type, Nox2−/− mice exhibit a time-dependent metabolic phenotype during diet-induced obesity. After 8 weeks of high-fat feeding, Nox2−/− mice exhibit attenuated ATM inflammation and mildly improved glucose tolerance. After 16 weeks of high-fat feeding, Nox2−/− mice develop severe insulin resistance, hepatosteatosis, and visceral lipoatrophy characterized by dead adipocyte accumulation and defective ATM lysosomal exocytosis, a phenotype reproduced in myeloid cell-specific Nox2−/− mice. Collectively, our findings suggest that MMe macrophages perform detrimental and beneficial functions whose contribution to metabolic phenotypes during obesity is determined by disease progression.

INTRODUCTION

During obesity, macrophages accumulate in visceral adipose tissue, where they promote chronic low-grade inflammation (Weisberg et al., 2003; Xu et al., 2003). It is well accepted that this inflammation is causally associated with insulin resistance in mice. Inhibiting several pathways that drive inflammatory signaling and/or production in macrophages improves insulin sensitivity during obesity (Han et al., 2013; Patsouris et al., 2008; Saberi et al., 2009; Wei et al., 2016).

In addition to producing inflammatory cytokines, adipose tissue macrophages (ATMs) have also been postulated to perform beneficial functions during diet-induced obesity (DIO) (Fitzgibbons and Czech, 2016). They internalize excess free fatty acids (FFAs) released by insulin-resistant adipocytes, thereby buffering metabolic tissues from the damage caused by ectopic accumulation of saturated FFAs. They also clear dead adipocytes that accumulate during prolonged obesity when adipose tissue expands enough to produce hypoxia (Strissel et al., 2007; Sun et al., 2011). The clearance of these dead adipocytes promotes adipocyte turnover and maintains adipose tissue health during nutrient excess.

The diverse functions of macrophages, such as those described earlier, have often been conceptualized through an M1 and M2 paradigm (Gordon and Taylor, 2005). The M1 phenotype is caused by Th1 mediators such as lipopolysaccharide (LPS) and interferon gamma (IFNg) and is characterized by increased production of pro-inflammatory cytokines, while the M2 phenotype is driven by Th2 mediators (e.g., interleukin-4 [IL-4]), which activates expression of immunosuppressive factors that promote macrophage clearance of dead cells and tissue remodeling (Odegaard et al., 2007). Although this M1/M2 paradigm has been a useful construct for understanding macrophage heterogeneity, studies suggest that it cannot adequately describe the functions of ATMs during obesity.

Studies from Xu et al., (2013) showed that macrophages from obese mice induce lysosomal and lipid metabolism pathways, raising the possibility that these pathways might help macrophages to clear dead adipocytes and their large FFA reservoirs. Moreover, studies from our laboratory showed that saturated FFAs produce a pro-inflammatory, metabolically activated (MMe) macrophage phenotype that is mechanistically distinct from M1 or M2 activation (Kratz et al., 2014). Although we showed that MMe macrophages accumulate in visceral and subcutaneous adipose tissue of obese humans and mice, their roles...
in regulating macrophage inflammation, function, and insulin sensitivity have not been explored.

Here we combine proteomic and functional analyses with genetic models and dietary interventions to investigate the roles of MMe macrophages in DIO. We show that metabolic activation of macrophages induces inflammatory cytokine production (a detrimental function) and lysosomal exocytosis to dead adipocytes (a beneficial function). We identify NADPH oxidase 2 (NOX2) as a key regulator of the inflammatory cytokine expression and lysosomal exocytosis in MMe macrophages and show that ablating Nox2 both improves and worsens the metabolic phenotype during DIO, depending on the duration of high-fat feeding. Collectively, our findings underscore the dynamic functions of ATMs during DIO and demonstrate the importance of the MMe phenotype in regulating these functions and their metabolic consequences.

RESULTS

Metabolic Activation Induces Macrophage Inflammation and Lysosomal Exocytosis In Vitro and In Vivo

We previously showed that saturated FFAs produce macrophage inflammation via an MMe pathway that is mechanistically distinct from the classically activated M1 phenotype induced by bacterial challenge (Kratz et al., 2014). Although we showed that these MMe macrophages are present in adipose tissue of obese humans and mice, their functions in DIO and insulin resistance are poorly understood.

To interrogate how metabolic activation might alter macrophage function, we produced pro-inflammatory MMe macrophages by treating murine bone marrow-derived macrophages (BMDMs) with glucose, insulin, and palmitate (Figure 1A) and performed plasma membrane proteomics. We focused on plasma membrane proteins because they mediate a variety of important macrophage functions, including phagocytosis, cell migration, and antigen presentation.

Mass spectrometric analysis of the plasma membrane identified 664 proteins in M0 (unstimulated) and MMe macrophages. Statistical analysis with the t test and G test identified 176 proteins that were differentially abundant (false discovery rate [FDR] < 0.05) in MMe versus M0 macrophages (Figure 1B). Gene ontology analysis of proteins induced in MMe macrophages identified enrichments in several annotations expected for a cell surface proteome (i.e., plasma membrane, membrane raft, and extracellular matrix) (Figure 1C), suggesting that our methods for isolating plasma membrane proteins were robust. Moreover, we observed increased levels of MMe-associated proteins (PLIN2 and P62), but not M1-associated proteins (CD40 and STAT1), that we previously identified in proteomic analyses of human MMe and M1 macrophages (Figure S1) (Kratz et al., 2014).

Unexpectedly, plasma membrane proteins induced in MMe macrophages were enriched in lysosomal membrane proteins (p = 10−6) (Figure 1C; Table S1), and this enrichment was specific to the MMe phenotype, because they were not observed in M1 macrophages (Figure 1C). To confirm these observations using an alternative approach, we quantified cell surface LAMP1 and LAMP2 levels on M0, MMe, and M1 macrophages by flow cytometry. Consistent with our proteomics data, we found that LAMP1 and LAMP2 were induced on the cell surface of MMe, but not M1, macrophages (Figure 1D; Figure S1). Because lysosomal membrane proteins appear on the cell surface during lysosomal exocytosis, when lysosomal and plasma membranes fuse (Rodríguez et al., 1997), these findings suggest that metabolic activation induces macrophages to exocytose their lysosomes.

Previous studies showed that lysosomal exocytosis by macrophages is required for them to clear dead adipocytes (Haka et al., 2016). Because macrophages (~20 μm diameter) are much smaller than adipocytes (~120 μm diameter), they cannot clear dead adipocytes through a traditional phagocytosis mechanism. Instead, the macrophages form a tight attachment (a lysosomal synapse) on the dead adipocyte. They acidify the contact region using the plasma membrane proton pumping V-ATPase and secrete lysosomal contents into the lysosomal synapse. The lysosomal lipases in the acidic milieu liberate FFAs that are subsequently internalized by macrophages, leading to foam cell formation.

Because FFAs are required for metabolic activation, and dead adipocytes are an abundant reservoir of FFA, we hypothesized that dead adipocytes could induce the inflammation and lysosomal exocytosis associated with the MMe phenotype. To test this hypothesis, we treated unstimulated BMDMs with media conditioned by live or apoptotic 3T3-L1 adipocytes and measured inflammatory cytokine expression by qRT-PCR and quantified lysosomal exocytosis by staining for cell surface LAMP1 and LAMP2.

We found that media from apoptotic adipocytes coordinately induced the expression of Il1β and Il6 (Figure 1E) and the appearance of cell surface LAMP1 and LAMP2 on macrophages (Figure 1F). The effects on cytokines and lysosomal exocytosis were specific to apoptotic adipocytes, because media from live 3T3-L1 adipocytes or apoptotic neutrophils (a dead cell without FFA) were incapable of inducing these changes (Figures 1E and 1F). Thus, dead adipocytes release factors that induce the inflammation and lysosomal exocytosis characteristic of MMe macrophages.

Our previous studies showed that ATMs from obese mice adopt an MMe phenotype (Kratz et al., 2014). We therefore reasoned that ATMs from obese mice would also show coordinated induction of cytokine expression and lysosomal exocytosis. To test this, we fed C57BL/6 mice a low-fat diet (LFD) or 45% high-fat diet (HFD) for 16 weeks, isolated ATMs (Figure 1G), confirmed their purity based on CD11c and F4/80 staining (98.1%) (Figure 1H), and assessed cytokine expression by qRT-PCR and cell surface lysosomal membrane protein levels by mass spectrometry.

Consistent with previous findings, ATMs from obese mice were characterized by elevated Tnfa, Il6, and Il1β expression (Figure 1I). Moreover, plasma membrane proteins elevated in ATMs from obese mice (relative to lean mice) were significantly enriched for lysosomal membrane proteins (p = 10−7). Indeed, 10 lysosomal membrane proteins were induced on the cell surface of ATMs from obese mice (Figures 1J and 1K; Table S1), 6 of which (LAMP2, ATP6V1B2, ATP6V1A, AHNAK, NPC1, and ANXA2) were also induced on the cell surface of MMe macrophages in vitro.
TLR2, NOX2, and MYD88 Coordinately Regulate Inflammation and Lysosomal Exocytosis in MMe Macrophages

What signaling pathways regulate cytokine expression and lysosomal exocytosis in MMe macrophages? To begin to answer this question, we searched the literature for proteins known to mediate macrophage responses to FFAs, the driver of the MMe phenotype. We focused on toll-like receptor (TLR) 2, NOX2, MYD88, and TLR4 because previous studies showed that these proteins are required for palmitate-induced inflammation and/or endoplasmic reticulum (ER) stress in macrophages (Kratz et al., 2014; Robblee et al., 2016; Seimon et al., 2010).

We first investigated the effects of these knockouts on inflammatory cytokine expression. We found that ablating Nox2 (gp91 subunit), Tlr2, Myd88, or Tlr4 attenuated Il6 and Il1β levels in MMe macrophages (Figure 2A; Figure S2). The anti-inflammatory effect of ablating Nox2 or Tlr2 was specific to MMe macrophages, because cytokine expression in M1 macrophages was unaffected (Figure 2A). In contrast, ablating Tlr4 or Myd88, an adaptor protein used by most TLRs to activate nuclear factor-kB (NF-kB) (Akira et al., 2001), lowered inflammatory cytokine expression in both MMe and M1 macrophages (Figure 2A; Figure S2). Altogether, these studies identified TLR2, NOX2, MYD88, and TLR4 as regulators of inflammatory cytokine expression in MMe macrophages and further demonstrated that distinct signaling pathways regulate inflammatory cytokine expression in MMe and M1 macrophages.
We further investigated the effects of these knockouts on lysosomal exocytosis. To this end, we used a more physiologically relevant co-culture assay (Haka et al., 2016) that quantifies (1) the delivery of lysosomal contents from macrophages to apoptotic 3T3-L1 adipocytes and (2) the subsequent accumulation of adipocyte-derived lipids in the macrophage (Figure 2B). As a control, macrophages were co-incubated with live adipocytes, demonstrating the specificity of these processes to apoptotic adipocytes (Figure 2B).

We found that lysosomal exocytosis to apoptotic adipocytes was significantly impaired in Nox2−/−, Tir2−/−, and Myd88−/− macrophages (Figure 2C). Moreover, Nox2−/−, Tir2−/−, and Myd88−/− macrophages accumulated fewer lipids from apoptotic adipocytes (Figure 2C), perhaps due to their inability to deliver lysosomal lipases to mobilize FFAs from the adipocyte lipid droplet. Consistent with this interpretation, the extent of macrophage lipid accumulation was strongly and positively correlated (R² = 0.94, p < 0.0001) to the amount of lysosomal exocytosis (Figure 2D). Moreover, Nox2−/−, Tir2−/−, and Myd88−/− macrophages were not defective in their ability to internalize palmitate (Figure 2E), one of the major FFAs liberated during lipolysis from adipocytes.

Lysosomal exocytosis to apoptotic adipocytes was not impaired in Tlr4−/− macrophages (Figure S2), even though this knockout suppressed inflammatory cytokine expression. These findings suggest that the defective lysosomal exocytosis observed in Tlr2−/−, Nox2−/−, and Myd88−/− macrophages was not reliant upon the suppression of inflammatory cytokines. Consistent with this hypothesis, supplementing Nox2−/− macrophages with tumor necrosis factor alpha (TNF-α), a key cytokine produced by ATMs, could not rescue the defect in lysosomal exocytosis (Figure 2F).

ATM Inflammation Precedes Visceral Fat Adipocyte Death during DIO

Our in vitro findings suggest that MMe macrophages might play complex roles during DIO. However, MMe macrophages overexpress inflammatory cytokines, which could promote insulin resistance (Chawla et al., 2011; Olefsky and Glass, 2010). However, MMe macrophages exocytose their lysosomes and clear
dead adipocytes, which could limit insulin resistance by main-
taining adipose tissue health and protecting against ectopic lipid
deposition in the liver (Shulman, 2014; Strissel et al., 2007; Sun
et al., 2011). Because these potentially detrimental and benefi-
cial properties of MMe macrophages are coordinately regulated
by TLR2, NOX2, and MYD88, we reasoned that interpreting the
metabolic phenotypes of mice deficient in one or more of these
proteins might be challenging.

To help facilitate interpretation, we first determined whether
ATM inflammation and adipocyte death (and hence the need
for dead adipocyte clearance) occurred at different time points
during DIO. Previous studies showed that ATM inflammation
might be an early and persistent event, while adipocyte death
may predominantly occur after prolonged high-fat feeding
when adipose tissue becomes hypoxic (Figure 3A) (Strissel
et al., 2007). However, these studies were performed in mice
fed a 60% HFD, and our studies used the 45% HFD. To better
understand the timing of these events and to establish reproduc-
ible conditions to investigate the effects of our knockouts on
these processes in vivo, we studied adipose tissue biology in
wild-type C57BL/6 mice fed an LFD or 45% for 8 and 16 weeks
(Figure 3B).

Figure 3. ATM Inflammation Precedes Visceral Fat Adipocyte Death during DIO
Wild-type C57BL/6 mice were fed a low-fat diet (LFD) or 45% high-fat diet (HFD) for 8 or 16 weeks.
(A) A model for the timing of ATM inflammation and epididymal fat death during DIO.
(B) Body weight gain.
(C) mRNA expression of inflammatory cytokines and lipid metabolism genes in purified ATMs.
(D) Hif1α mRNA levels in epididymal fat (a marker of hypoxia).
(E) Caspase-3 (CASP3) activity levels in epididymal fat.
(F) Epididymal fat was stained with antibodies against PLIN2 (red, identifies adipocytes), MAC2 (green, identifies macrophages), and DAPI (blue, identifies nuclei). Macrophages in crown-like structures (CLSs, arrow) were quantified by microscopy and standardized per unit area.
(G) H&E staining of liver and quantification of liver triglyceride (TG) levels.
Results are mean ± SEM; n = 5–15, *p < 0.05 Student’s t test. See also Figures S3 and S4.
Eight weeks of high-fat feeding was sufficient to induce the MMe phenotype in ATMs characterized by increased inflammatory cytokines (Tnfα, Il6, and Il1β) and genes involved in lipid metabolism (Plin2, Abca1, and Cds2) (Figure 3C). Despite the presence of ATM inflammation, several lines of evidence indicated that epididymal fat health in obese mice was not significantly compromised at this time point. First, adipose tissue levels of Hif1α were not elevated (Figure 3D), suggesting that hypoxia, a key driver of adipocyte death (Wang et al., 1995), had not yet occurred. Second, adipose tissue caspase-3 activity, a marker of apoptosis (Nicholson et al., 1995), was not elevated (Figure 3E). Third, the number of crown-like structures, an indicator of adipocyte death (Murano et al., 2008), was unchanged (Figure 3F; Figure S3). Finally, ectopic fat deposition in the liver, which has been associated with adipocyte death in visceral fat (Strissel et al., 2007), was undetectable (Figure 3G; Figure S4).

Although ATMs still overexpressed the cytokines and lipid metabolism genes characteristic of MMe macrophages following 16 weeks of high-fat feeding (Figure 3C), the state of epididymal fat health was substantially different at this time point. Adipose tissue Hif1α levels were elevated (Figure 3D), suggesting that the tissue was hypoxic. Moreover, adipose tissue caspase-3 activity and crown-like structures were markedly increased (Figures 3E and 3F; Figure S3), indicating the presence of apoptotic adipocytes. Consistent with the deterioration of epididymal fat health and increased FFA release, the MMe phenotype was more pronounced at this time point, particularly with respect to the lipid metabolism genes (Figure 3C). Moreover, ectopic fat deposition in the liver was detectable in obese mice (Figure 3G; Figure S4), and this deposition could not be explained by changes in genes involved in triglyceride (TG) synthesis (Fas1, Acc1, and Sreb1) or fatty acid metabolism (Ppara and Cpt1α) in the liver (Figure S4).

These findings suggest that while MMe ATMs are present at both 8 and 16 weeks of 45% high-fat feeding, adipocyte death, and hence the requirement for macrophages to clear dead adipocytes, may only be an important consideration after 16 weeks of high-fat feeding.

**Nox2** - **Mice Have Improved ATM Inflammation and Glucose Tolerance after 8 Weeks of High-Fat Feeding**

Our identification of NOX2, TLR2, and MYD88 as drivers of cytokine expression and lysosomal exocytosis in MMe macrophages provided an opportunity to inhibit these processes in vivo and assess effects on obesity-associated insulin resistance. We focused our initial studies on Nox2−/− mice because NOX2 is predominantly expressed by phagocytic cells, including macrophages and neutrophils (Bedard and Krause, 2007).

We began by interrogating the metabolic phenotypes of Nox2−/− mice following 8 weeks of high-fat feeding. Because ATM inflammation was present at this time point, but adipocyte apoptosis had not yet occurred (Figure 3), we hypothesized that the anti-inflammatory effect of Nox2−/− would predominate and protection from metabolic derailments during DIO would be observed at this early time point. This hypothesis makes several predictions.

First, ablating Nox2 should attenuate ATM inflammation. Although Nox2−/− mice fed the HFD were heavier than wild-type mice (Figure 4A), ATMs isolated from visceral fat were substantially less inflamed, as evidenced by decreased expression of Tnfα, Il6, and Il1β (Figure 4B). Because NOX2 drives inflammatory cytokine expression in MMe, but not M1, macrophages (Figure 2), these findings reinforce the notion that ATMs are inflamed via an MMe pathway in vivo (Kratz et al., 2014).

Second, ablating Nox2 should not increase the number of dead adipocytes. Nox2−/− mice showed no differences in adipocyte death indicators; Hif1α levels, caspase-3 activity, and the number crown-like structures were unchanged, and ectopic fat accumulation in the liver was only minimally and insignificantly increased in Nox2−/− mice relative to wild-type mice fed the HFD (Figures 4C–4F; Figures S3 and S4). Moreover, none of these adipocyte death indicators were elevated in obese relative to lean mice, irrespective of genotype (Figures 4C–4F), which is consistent with our findings that 8 weeks of 45% HFD is insufficient to induce adipocyte death (Figure 3).

Third, this hypothesis predicts that Nox2−/− mice should demonstrate improved metabolic parameters. Nox2−/− mice fed the HFD had lower fasting glucose levels (Figure 4G) and improved glucose tolerance (Figure 4H). However, fasting insulin levels and insulin tolerance remained unchanged (Figures 4G and 4I), perhaps due to the increased body weight, the mild hepatosteatosis, and the increased number of ATMs in Nox2−/− mice (Figure S5), all of which would limit the anti-inflammatory effect of the knockout.

Altogether, these findings provide further evidence that ATMs are inflamed via an MMe pathway in vivo and further suggest that inhibiting inflammatory cytokine expression in MMe macrophages (by ablating Nox2) protects against metabolic dysfunction following 8 weeks of high-fat feeding.

**Nox2** - **Mice Develop Severe Visceral Lipoatrophy, Hepatosteatosis, and Insulin Resistance following 16 Weeks of HFD**

Next, we investigated the metabolic phenotypes of Nox2−/− mice fed a HFD for 16 weeks. Because ATM inflammation and adipocyte death were both present at this time point (Figure 3), and ablating Nox2 attenuated metabolic activation and lysosome secretion in response to dead adipocytes (Figure 2), we hypothesized that Nox2−/− mice might exhibit a more complex phenotype at this later time point. To explore this, we characterized the adipose tissue, liver, and metabolic parameters of Nox2−/− mice fed the HFD for 16 weeks.

The protection observed in Nox2−/− mice following 8 weeks of HFD was reversed after prolonged high-fat feeding. Nox2−/− mice fed a HFD for 16 weeks were more obese, hyperglycemic, hyperinsulinemic, and insulin resistant than wild-type controls (Figures 5A–5D). Nox2−/− mice retained glucose tolerance, but they required higher levels of insulin production to maintain glucose homeostasis (Figure 5C).

An examination of the epididymal fat in Nox2−/− mice fed the HFD for 16 weeks revealed marked increases in the number of crown-like structures (Figure 5E; Figure S3) and caspase-3 activity (Figure 5F) relative to control mice, suggesting a higher accumulation of dead adipocytes. The extent of this adipocyte death was substantial given that Nox2−/− mice fed the HFD had less epididymal fat mass than wild-type mice (Figure 5G), despite
being more obese (Figure 5A). In contrast, Nox2<sup>−/−</sup> mice had more epididymal fat than wild-type mice following 8 weeks of HFD (Figure 5G). These findings suggest that Nox2<sup>−/−</sup> mice are capable of expanding their epididymal fat mass during early DIO but that they cannot sustain it during prolonged exposure to the HFD.

Despite the increased number of dead adipocytes in Nox2<sup>−/−</sup> mice, which should induce macrophage inflammation (Figure 1), ATM expression of Tnfα, Il6, and Il1β was similar to that of wild-type mice (Figure 5H). Thus, it is likely that the anti-inflammatory effect of ablating Nox2 was still present at 16 weeks but masked by the hyper-inflammatory environment produced by the dead adipocytes.

Consistent with the deterioration of epididymal fat health and mass at 16 weeks of HFD, Nox2<sup>−/−</sup> mice displayed severe hepatosteatosis, characterized by increased hepatic TG mass and liver weight (Figures S1–S5; Figure 5I). This hepatosteatosis could not be explained by changes in the expression of liver genes involved in TG synthesis (Fas1, Acc1, and Srebp1) or fatty acid metabolism (Pparα and Cpt1α) (Figure 5L).

Altogether, these studies showed that Nox2<sup>−/−</sup> mice develop severe late-onset adipocyte death, hepatosteatosis, and insulin resistance. Because Nox2<sup>−/−</sup> mice are more obese than their wild-type counterparts, we explored the possibility that these late-onset effects were due to excess body mass. To test this, we fed wild-type mice a HFD until their average body weight
matched Nox2−/− mice at 16 weeks (wild-type match) and quantified key aspects of the metabolic phenotype.

We found that wild-type-match mice could not mimic the late-onset dead adipocyte accumulation, hepatosteatosis, and insulin resistance observed in Nox2−/− mice (Figures S3, S4, and S6), suggesting that these metabolic abnormalities were not driven by the increased body weight of Nox2−/− mice fed the HFD. Liver TGs and fasting glucose levels were not correlated to body weight in wild-type and Nox2−/− mice fed the HFD (Figures 5M and 5N). In contrast, these parameters were significantly and positively correlated to the extent of adipocyte death in epididymal fat, as determined by caspase-3 activity (Figures 5M and 5N). Thus, the inability of Nox2−/− mice to sustain their epididymal fat may help to explain the late-onset hepatosteatosis and metabolic defects observed in these mice.

Dead Adipocyte Accumulation Is Associated with Defective Lysosomal Exocytosis by Nox2−/− ATMs

In principle, the number of dead adipocytes observed in vivo is determined both by the rate of death and the rate of clearance (Figure 6A); the latter process, which is mediated by ATMs, has not been the subject of intensive study. Because Nox2−/− macrophages exhibited defective lysosomal exocytosis and clearance of apoptotic adipocytes in vitro (Figures 1 and 2), we reasoned that defective dead adipocyte clearance by ATMs might help to explain the late-onset accumulation of dead adipocytes. We used three approaches to test this hypothesis.

First, we investigated whether drivers of adipocyte death were elevated in Nox2−/− mice. Adipocyte apoptosis can be caused by hypoxia and/or inflammation (Sun et al., 2011), both of which increase as adipose tissue expands during obesity. Neither of
these mechanisms are a likely cause of dead adipocyte accumulation in Nox2\(^{-/-}\) mice, because adipose tissue levels of Hif1\(\alpha\) (hypoxia marker) and Tnf\(\alpha\) and Ili1\(\beta\) (pro-inflammatory markers) were unchanged after 16 weeks of high-fat feeding compared to wild-type mice (Figures 6B and 6C). Moreover, ATM inflammation (Figure 5) and ATM number (Figure S5) were not significantly different between wild-type and Nox2\(^{-/-}\) mice fed the HFD.

Second, we determined whether ATMs from Nox2\(^{-/-}\) mice had lower cell surface levels of lysosomal membrane proteins, the hallmark of lysosomal exocytosis (Rodrı́guez et al., 1997). We isolated ATMs from HFD-fed wild-type and Nox2\(^{-/-}\) mice and interrogated them by plasma membrane proteomics. This analysis identified 8 lysosomal membrane proteins on the cell surface of wild-type ATMs, all of which were lower on the cell surface of ATMs from Nox2\(^{-/-}\) mice (Figures 6D and 6E; Table S1). The decreased levels of lysosomal membrane proteins on Nox2\(^{-/-}\) ATMs were likely underestimated given that Nox2\(^{-/-}\) mice had significantly more dead adipocytes, which should stimulate lysosomal exocytosis (Figures 1 and 2). These findings suggest that ATMs from Nox2\(^{-/-}\) mice exhibit impaired lysosomal exocytosis in vivo.

Third, we determined whether ablating Nox2 in myeloid cells specifically could reproduce the late-onset accumulation of dead adipocytes and associated metabolic derailing. We crossed Nox2\(^{-/-}\) mice with LysM-Cre\(^{-/-}\) mice to generate myeloid cell-specific, Nox2-deficient mice (Nox2\(^{-/-}\)LysM-Cre\(^{-/-}\)) and littermate controls (Nox2\(^{-/-}\)LysM-Cre\(^{-/-}\)). After validating Nox2 knockout in ATMs (Figure S7), we examined the metabolic phenotypes of these mice following prolonged high-fat feeding.

Myeloid cell-specific ablation of Nox2 mimicked the late-onset accumulation of dead adipocytes in the absence of increased hypoxia or adipose tissue inflammation (Figures 7A–7D; Figure S3), hepatosteatosis and increased liver weight (Figures 7E and 7F; Figure S4), obesity (Figure 7G), hyperglycemia (Figure 7H), and hyperinsulinemia (Figure 7I) phenotypes of the whole-animal knockout. The stronger increase in the number of crown-like structures in the LysM-Cre model relative to the global knockout was due to the presence of fewer crown-like structures in LysM-Cre\(^{-/-}\) control mice, which likely reflects the variability in the appearance of crown-like structures in vivo. These findings suggest that the accumulation of dead adipocytes and the associated metabolic dysfunction are driven by the loss of Nox2 in myeloid cells, not an unanticipated effect in adipocytes.

**DISCUSSION**

Accumulating evidence suggests that ATMs perform both detrimental and beneficial functions during obesity. They produce inflammatory cytokines that promote insulin resistance (Chawla et al., 2011; Olefsky and Glass, 2010). They also might protect metabolic tissues from the deleterious effects of excess FFAs and contribute to adipose tissue homeostasis by clearing dead adipocytes. Traditionally, these diverse functions have been attributed to distinct ATM populations; the detrimental functions have been associated with M1-like macrophages, which may predominate during early DIO (Lumeng et al., 2007), while the beneficial functions have been ascribed to M2-like macrophages, which may accumulate during prolonged DIO (Shaul et al., 2010).

Here we provide evidence that inflammatory cytokine production and dead adipocyte clearance are functional properties of a single MMe macrophage phenotype that is present during early and late DIO. We further show that inflammatory signaling through NOX2, TLR2, and MYD88 coordinately regulates both the detrimental and the beneficial functions of MMe macrophages. Accordingly, ablating Nox2 produces a complex metabolic phenotype determined by the duration of high-fat feeding, which in turn highlights the relative importance of macrophage...
inflammation and dead adipocyte clearance during the progression of obesity.

After 8 weeks of HFD, wild-type mice exhibited ATM inflammation in the absence of adipose tissue hypoxia and adipocyte death, indicating a low need for dead adipocyte clearance at this time point. Accordingly, the anti-inflammatory benefit of ablating Nox2 was observed without exposing the costs associated with impaired dead adipocyte clearance. Nox2−/− mice had attenuated ATM inflammation and exhibited improvements in fasting glucose and glucose tolerance, even though mice were more obese, had more ATMs in epididymal fat, and were trending to hepatosteatosis—all of which should exacerbate metabolic dysfunction during DIO. Because ablating Nox2 inhibits inflammation in MMe macrophages, but not M1 macrophages, our findings further underscore the importance of metabolic activation in producing ATM inflammation in obesity.

In contrast, after 16 weeks of HFD, we found that dead adipocytes accumulated in epididymal fat of wild-type mice, and this accumulation was significantly exacerbated in Nox2−/− mice, suggesting that the costs associated with impaired dead adipocyte clearance were evident at this time point.

Three lines of evidence suggest that this late-onset dead adipocyte accumulation in Nox2−/− mice was due to impaired clearance by ATMs. First, we showed that inflammation and hypoxia, the main drivers of adipocyte death, were not increased in Nox2−/− mice relative to wild-type mice. Second, we showed that ablating Nox2 interfered with dead adipocyte clearance through lysosomal exocytosis in vitro and further showed that Nox2−/− ATMs had impaired lysosomal exocytosis in vivo. Third, we showed that mice lacking Nox2 in myeloid cells specifically reproduced this late-onset phenotype. Although neutrophils are also targeted by the LysM-Cre system (Clausen et al., 1999), ATMs are the most abundant cell type in adipose tissue and form the adipocyte-clearing crown-like structures during DIO. It is therefore unlikely that neutrophils significantly contribute to the impaired adipocyte clearance observed in vivo. However, because neutrophils promote insulin resistance during DIO (Talukdar et al., 2012), we cannot ignore their potential contribution to the metabolic phenotypes observed in the Nox2−/− mice. Collectively, these findings implicate MMe ATMs and their ability to exocytose their lysosomes to dead adipocytes as a key determinant of adipose tissue remodeling and health during prolonged obesity.

Figure 7. Ablating Nox2 in Myeloid Cells Is Sufficient to Produce Late-Onset Lipoatrophy, Hepatosteatosis, and Insulin Resistance

A comparison of the late-onset phenotypes in whole-animal, NOX2-deficient (global knockout [KO]), and myeloid cell-deficient (LysM-Cre KO) mice fed the HFD. All parameters are plotted relative to the appropriate wild-type control (CTRL).

(A) Epididymal fat was stained with antibodies against PLIN2 (red, identifies adipocytes), MAC2 (green, identifies macrophages), and DAPI (blue, identifies nuclei). Macrophages in crown-like structures (CLSs, arrow) were quantified by microscopy and standardized per unit area.

(B) Relative caspase-3 (CASP3) activity in epididymal fat.

(C) Relative epididymal fat mass.

(D) Relative levels of Hif1α (a hypoxia marker) and Tnfα and Il1β (inflammatory markers) in epididymal fat.

(E) H&E staining of the liver.

(F) Relative liver weight.

(G) Relative body weight gain.

(H and I) Relative fasting glucose (H) and insulin (I) levels. Data for the global control and KO mice are re-presented in modified form for comparative purposes. Results are mean ± SEM; n = 5–10, *p < 0.05 Student’s t test. See also Figures S3, S4, and S7.
The late-onset dead adipocyte accumulation was so extensive in Nox2\(^{-/-}\) mice that they had less epididymal fat mass than their wild-type counterparts despite being more obese, suggesting that visceral fat could no longer store excess dietary nutrients. This phenotype shares some similarities with lipodystrophy, in which the inability to store nutrients in adipose tissue results in ectopic fat accumulation in the liver and hepatosteatosis, which was likely responsible for the hyperglycemia, hyperinsulinemia, and insulin resistance at this time point (Cohen et al., 2011; Shulman, 2014).

However, unlike lipodystrophy, Nox2\(^{-/-}\) mice did not exhibit inherent defects in adipose tissue development or expansion. Epididymal fat in Nox2\(^{-/-}\) mice fed the LFD was normal at both time points. Furthermore, epididymal fat mass was higher in Nox2\(^{-/-}\) mice fed the HFD for 8 weeks relative to wild-type mice. Instead, Nox2\(^{-/-}\) mice had a diminished capacity to maintain their epididymal fat during prolonged nutrient excess, when adipose tissue hypoxia and adipocyte death were apparent, and ATM-mediated dead adipocyte clearance was required.

The clearance of dead adipocytes poses two unique challenges for macrophages. First, because adipocytes are so large, macrophages cannot phagocytose them. Instead, their clearance relies on the formation of extracellular lysosomal compartments between many macrophages and one adipocyte (Haka et al., 2016), resulting in the formation of crown-like structures. Second, adipocytes are predominantly composed of TGs. Thus, during clearance, macrophages are exposed to FFAs such as palmitate, which induce macrophage inflammation (Kratz et al., 2014). For this reason, clearance of dead adipocytes, unlike clearance of other dead cells (Fadok et al., 1998), is an inflammation-propagating event. Crown-like structures have been positively correlated with inflammation in adipose tissue of obese humans and mice (Lumeng et al., 2007).

From this perspective, the MMe macrophage is well positioned to handle the unique challenges imposed by dead adipocytes. FFAs released by dead adipocytes promote the MMe macrophage phenotype and its associated upregulation of lysosomal exocytosis, lipid metabolism, and inflammation. Lysosomal exocytosis may drive FFA liberation to help clear the TG droplet, which may be further enabled by the induction of lysosomal genes reported in ATMs from morbibly obese mice (Xu et al., 2013). Increased expression of lipid metabolism genes (i.e., ABCA1, CD36, and PLIN2) could help macrophages to productively handle this excess fat, while inflammation could facilitate recruitment of additional macrophages to the crown-like structure to assist in clearance.

Prolonged macrophage inflammation also carries its consequences. Cytokines such as TNF-\(\alpha\) promote damage to metabolic organs and drive insulin resistance during obesity (Hotamisligil et al., 1993) and our findings during early DIO provide further support for this widely accepted paradigm.

However, studies showed that the metabolic consequences of this inflammation might not be as straightforward as this paradigm suggests. For example, thermoneutral housing accelerated the onset of adipose tissue inflammation but did not worsen the metabolic phenotype (Tian et al., 2016). Similarly, ablating TNF-\(\alpha\) expression by adipocytes exacerbated obesity-associated insulin resistance and hepatosteatosis (Wernstedt Asterholm et al., 2014).

Our findings reinforce the emerging complex relationship among inflammation, macrophages, and metabolic phenotypes during DIO. We observed contradictory metabolic phenotypes with the same anti-inflammatory perturbation (Nox2\(^{-/-}\)) over two time points during DIO. Previous studies with Nox2\(^{-/-}\) mice fed the HFD reported similar contradictory findings. One study reported improvements in fasting glucose, glucose tolerance, and brain injury in Nox2\(^{-/-}\) mice fed a 60% HFD for 14 weeks (Pepping et al., 2013). Another study showed increased adiposity due to hyperphagia and worsened insulin resistance and hepatosteatosis in Nox2\(^{-/-}\) fed a 60% HFD for 18 weeks (Costford et al., 2014).

Although we cannot rule out the contribution of other NOX2-driven mechanisms, the contradictory findings that we and others report with Nox2\(^{-/-}\) mice may be conceptualized by integrating the beneficial and detrimental functions of ATMs through a single MMe phenotype, whose impact on adipose tissue biology and metabolic phenotypes is determined by the duration of high-fat feeding. Future experiments will be required to determine whether this paradigm holds true in other anti-inflammatory settings.

**EXPERIMENTAL PROCEDURES**

**Mice**

All animal studies were approved by the University of Chicago Institutional Animal Care and Use Committee (ACUP 72209). Wild-type, Gp91\(^{-/-}\), Tlr2\(^{-/-}\), Tlr4\(^{-/-}\), and Myd88\(^{-/-}\) male mice on the C57BL/6 background are from Jackson Laboratory. For DIO studies, wild-type, Gp91\(^{-/-}\), Tlr2\(^{-/-}\), and Tlr4\(^{-/-}\) mice were placed on an LFD or HFD (45% fat, Research Diets, D12451) at 8 weeks of age for up to 16 weeks.

**Nox2\(^{-/-}\) Mice**

Nox2\(^{-/-}\) mice were described (Sag et al., 2017) and were crossed with LysM-cre knockin mice (Jackson Laboratory, 004781) to generate LysM-cre\(^{-/-}\) Nox2\(^{-/-}\) mice and littermate control Nox2\(^{+/+}\) mice. See Supplemental Experimental Procedures for genotyping details.

**Differentiation and Activation of BMDMs**

Murine BMDMs were differentiated from bone marrow stem cells as previously described (Kratz et al., 2014). For M1 activation, BMDMs were treated with LPS (5 ng/mL) and IFN\(\gamma\) (12 ng/mL) for 24 hr. For MMe activation, macrophages were treated with a combination of glucose (30 mM), insulin (10 nM), and palmitate (0.4 mM) for 24 hr. Macrophages were also treated with media conditioned by live 3T3-L1 adipocytes, apoptotic 3T3-L1 adipocytes, or apoptotic neutrophils for 24 hr.

**Macrophage Clearance of Apoptotic Adipocytes**

Macrophage clearance of apoptotic adipocytes and subsequent accumulation of lipids were performed as previously described (Haka et al., 2016). See Supplemental Experimental Procedures for additional details.

**ATM Isolation and Analysis**

ATMs were isolated using anti-CD11b antibody coupled to magnetic beads as previously described (Kratz et al., 2014), and purity was assessed by flow cytometry. ATMs were interrogated by qRT-PCR, plasma membrane proteomics, and flow cytometry. A workflow for flow cytometric analysis of ATMs is provided in the Supplemental Information (Figure S5).
Plasma Membrane Proteomics
Plasma membrane proteins of BMDMs or ATMs were isolated and analyzed by mass spectrometry as previously described (Becker et al., 2012). Proteins were quantified by spectral counting and statistical significance was assessed using a combination of the G test (G statistic) and t test (p value) with correction for the false discovery rate as previously described (Heinecke et al., 2010).

Plasma Measurements
Mice were fasted for 3 hr and serum insulin levels were measured by ELISA (Millipore) and blood glucose levels were measured with a One Touch Ultra 2 glucometer (Lifescan). For glucose tolerance test (ITT), mice were fasted for 16 hr, injected with 1 g/kg glucose, and blood was collected at several time points for the measurement of glucose and insulin levels. For insulin tolerance test (ITT), mice were fasted for 6 hr, injected with 0.5 mL/kg insulin, and blood was collected at several time points for the measurement of glucose.

Adipose Tissue Measurements
Epidydmal fat was stained with antibodies against murine MAC2 (Cedarlane) and PLIN2 (Abcam), and fluorescence images were acquired using an Olympus IX81 inverted wide field microscope. The number of crown-like structures in epidydmal fat was by counting the number of adipocyttes surrounded by MAC-2 positive signal per unit area. Between 4 and 6 mice were used per condition and 3 and 5 images were used per mouse. Epidydmal fat caspase-3 activity was measured using the Apo-ONE kit (Promega) according to the manufacturer’s protocol.

Liver Measurements
Livers were perfused with 30 mL of PBS before excision, homogenized, and TGs were measured using a TG quantification kit (Abcam) according to the manufacturer’s protocol.

Statistics
Statistical significance was assessed using an unpaired, two-tailed, Student’s t test. Replicate numbers for each experiment are indicated in the figure legends.

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Experimental Procedures, seven figures, and one table and can be found with this article online at http://dx.doi.org/10.1016/j.celrep.2017.08.096.

AUTHOR CONTRIBUTIONS
Conceptualization, all authors; Investigation, B.R.C., K.Q.S., V.C.B.-L., G.Z., McPherson, S.M., Livers, and B.R.C.; Writing – Original Draft, L.B.; Writing – Reviewing & Editing, all authors; Supervision, L.B. and F.R.M.; Funding Acquisition, L.B. and F.R.M.

ACKNOWLEDGMENTS
This research was supported by grants from the NIH (R01DK102960 to L.B.; R37DK27083 to F.R.M.; R01HL93024 to F.R.M.; R01DK055257 to C.J.R.; P30 DK020593 to the DRTC Cell Biology Core, University of Chicago; T32DK087703 as support for B.R.C. and S.F.; and T32DK007074 as support for A.H.), the Bernice Goldblatt Endowment Fellowship, the University of Chicago (as support for C.C.), and the American Heart Association (10SDG3600027 to L.B.).

Received: June 1, 2017
Revised: July 25, 2017
Accepted: August 29, 2017
Published: September 26, 2017

REFERENCES


