An ionic control of metabolic-epigenetic reprogramming links inflammation to membrane potential-mediated nutrient acquisition

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Abstract

Metabolic and epigenetic reprogramming play pivotal roles in driving inflammation, but the precise regulatory mechanisms remain minimally understood. Here we show an ionic control mediated by macrophage Kir2.1, an inwardly-rectifying K+ channel, promoting lipopolysaccharide (LPS)-induced inflammation. Kir2.1 blockade by the selective inhibitor ML133 or its specific deletion in macrophages suppressed the production of LPS-induced inflammatory factors, such as interleukin-1β, and protected mice from LPS-induced sepsis in vivo. Kir2.1 loss-of-function led to a nutrient starvation phenotype, with impaired glucose and serine-glycine-one-carbon metabolism, whose synergy promotes the generation of S-adenosylmethionine (SAM). Accordingly, reduced SAM availability by Kir2.1 blockade decreased histone methylation at key inflammatory effector loci, such as Il1b. Although the immunomodulatory effect of Kir2.1 was independent of modulation by Ca2+ flux and general signaling pathways, its loss-of-function led to a depolarized membrane potential (Vm) which decreased the surface expression of nutrient transporters, including GLUT1 and CD98. We thus identify an ionic control of metabolic-epigenetic reprogramming that links inflammation to Vm-mediated nutrient acquisition and identifies potential new strategies for anti-inflammatory therapy.

Introduction

Macrophage activation by Toll-like receptors, notably Toll-like receptor 4 (TLR4), is intrinsically linked to metabolic reprogramming. Upon lipopolysaccharide (LPS) stimulation, inflammatory macrophages require a profound reprogramming of cellular metabolism to meet their metabolic demands for performing diverse inflammatory functions including the production and secretion of inflammatory cytokines. For example, LPS stimulation results in an increase in glycolytic activity and enhanced cellular anabolism to support inflammatory activity (O’Neill et al., 2016). The glycolytic enzyme pyruvate kinase M2 and the tricarboxylic acid (TCA) cycle metabolite succinate have been shown to induce pro-IL-1β expression through hypoxia-inducible factor 1-alpha activity (Palsson-McDermott et al., 2015a; Tannahill et al., 2013). Another TCA cycle metabolite, itaconate, acts as an anti-inflammatory metabolite via nuclear factor erythroid 2-related factor 2 (Mills et al., 2018). In addition, we have previously demonstrated that LPS activates the pentose phosphate pathway, the serine synthesis pathway, and one-carbon metabolism, the synergism of which drives epigenetic reprogramming for interleukin-1β (IL-1β) expression (Yu et al., 2019). However, how inflammatory macrophage metabolic phenotypes reflect biochemical dependencies and how the nutrient availability orchestrates inflammatory responses are still elusive.

In contrast to rapidly-proliferating cells, macrophages are not prone to rapid proliferation after LPS stimulation, but instead produce diverse inflammatory cytokines (O’Neill et al., 2016). However, macrophages still require a constant supply of extracellular nutrients that fuel both anabolic metabolism and the synthesis of diverse inflammatory factors. To acquire extracellular nutrients such as glucose and amino acids, cells including macrophages rely on transporter proteins in the plasma membrane (McCracken and Edinger, 2013). For example, glucose transporter 1 (GLUT1)-mediated glucose
metabolism plays critical roles in driving inflammatory macrophages (Freemerman et al., 2014). Although the alterations of cellular metabolic landscape upon LPS stimulation has been gradually illustrated, the regulatory mechanisms that maintain the nutrient acquisition to meet the metabolic demands for inflammation are minimally understood, particularly how macrophages coordinate different environmental changes (such as changes in ions and pH) to dynamically acquire and exploit extracellular nutrients for driving inflammation.

Many lines of investigation have demonstrated that intact ion transport is required for both innate and adaptive immunity. In immune cells, divalent cations such as Ca\(^{2+}\), Mg\(^{2+}\), and Zn\(^{2+}\) are important as second messengers to regulate immune signaling pathways. By contrast, monovalent cations such as K\(^{+}\) and Na\(^{+}\) are mainly required for the regulation of membrane potential (V\(_m\)), which indirectly controls Ca\(^{2+}\) flux and immune cell signaling (Feske et al., 2015). Studies using patch-clamp recording have revealed that macrophages express several types of K\(^{+}\) channel required for K\(^{+}\) homeostasis, V\(_m\), and Ca\(^{2+}\)-dependent signaling pathways during inflammation (Feske et al., 2015). For example, the voltage-gated K\(_V\)1.3 channel is critical for setting the resting V\(_m\) in macrophages (Mackenzie et al., 2003), and promotes the proliferation and inducible nitric oxide synthase expression stimulated by macrophage colony-stimulating factor (Vicente et al., 2003). Another K\(^{+}\) channel, K\(_{Ca}\)3.1, is activated by Ca\(^{2+}\) influx and is required for MCP-1-mediated macrophage migration into inflamed tissues (Toyama et al., 2008). Recently, the two-pore-domain K\(^{+}\) channels TWIK2 and THIK1 have been demonstrated to activate NLRP3 inflammasome by mediating K\(^{+}\) efflux (Di et al., 2018; Madry et al., 2018). However, despite the surging interest in the role of metabolic reprogramming in driving inflammatory macrophages, little attention has been paid to the ionic control of nutrient access and utilization during inflammation.

Here we found the inward rectier K\(^{+}\) channel, Kir2.1, was essential for the production of LPS-induced inflammatory effectors such as IL-1\(\beta\) by promoting nutrient import and utilization. Blockade of Kir2.1 impaired glucose and serine-glycine-one-carbon (SGOC) metabolism, whose synergy fuels histone methylation for the expression of LPS-induced inflammatory genes by supporting the generation of S-adenosylmethionine (SAM). We also uncovered an unexpected mechanism by which Kir2.1-mediated regulation of V\(_m\) maintains surface nutrient transporters levels to meet the metabolic demands of inflammatory macrophages. Thus, we identify an ionic-metabolic-epigenetic axis that links inflammation to nutrient acquisition and provide potential new strategies of manipulating V\(_m\) for anti-inflammatory therapy.

**Results**

**Kir2.1 is highly expressed and regulates the membrane potential in primary macrophages**
Using RNA-sequencing (RNA-seq) analysis, we found that K\textsubscript{Ca}3.1 (encoded by \textit{Kcnn4}), THIK1 (encoded by \textit{Kcnk13}), TWIK2 (encoded by \textit{Kcnk6}), and Kir2.1 (encoded by \textit{Kcnj2}) were most strongly expressed in bone-marrow-derived macrophages (BMDMs) (Fig. 1A). \textit{K}_{Ca}3.1 plays a role in macrophage activation and migration (Feske et al., 2015), while TWIK2 and THIK1 have recently been shown to be required for NLRP3 inammasome activation by promoting K\textsuperscript{+} efflux (Di et al., 2018; Madry et al., 2018). We thus focused on the inwardly-rectifying K\textsuperscript{+} channel Kir2.1. Under physiological conditions, Kir channels generate a large K\textsuperscript{+} conductance at potentials negative to K\textsuperscript{+} equilibrium potential (E\textsubscript{K}) but also permit a relatively small current flow at potentials slightly more positive to E\textsubscript{K} which is essential to stabilize the resting membrane potential (V\textsubscript{m}) (Hibino et al., 2010; Miyazaki et al., 1974; Sakmann and Trube, 1984). Among different tissues and cell types, the expression of Kir2.1 was relatively high in macrophages and other specialized macrophages including osteoclasts and microglia (Figure S1A). To examine whether primary macrophages exhibit functional Kir2.1 channels, we performed whole-cell patch-clamp recordings from peritoneal macrophages subjected to voltage-ramps from -120 to +60 mV. Ba\textsuperscript{2+} (BaCl\textsubscript{2}) is usually used to block the Kir channels and we found a robust Ba\textsuperscript{2+}-sensitive Kir current in the current-voltage relationship (Figure S1B). ML133 is a selective Kir2 blocker, with an IC\textsubscript{50} of 1.9 µM for Kir2.1 (Wang et al., 2011; Wu et al., 2010). In HEK293 cells transfected with Kir2.1, ML133 potently inhibited the Kir2.1 current of both a large inward current component and a slight outward current component (Edwards and Hirst, 1988; Sakmann and Trube, 1984) (Figure S1C). Consistently, ML133 inhibited the endogenous Kir2.1 currents in both resting and LPS-stimulated peritoneal macrophages (Figs. 1B, 1C, 1F, and S1D).

Because knockout of Kir2.1 is lethal in the neonate (Zaritsky et al., 2000), its role in immune cells has not been reported using Kir2.1-deficient immune cells. We deleted Kir2.1 selectively in myeloid cells by generating \textit{Lyz2-cre-Kcnj2\textsuperscript{f/f}} mice and found the ability of bone marrow cells to differentiate \textit{ex vivo} into macrophages in the absence of Kir2.1 was normal in light of the expression of the macrophage surface markers CD11b and F4/80 (Figure S1F). The proliferation of cultured Kir2.1-deficient BMDMs was also normal (Figure S1G). However, Kir2.1 deficiency greatly impaired the Kir2.1 currents in both resting and LPS-stimulated peritoneal macrophages (Figs. 1D, 1E, 1F, and S1E), and ML133 showed no additional effect (Figs. 1D, 1E, and 1F).

Kir2.1 was reported to stabilize the resting V\textsubscript{m} of many cell types including cardiac myocytes (Sakmann and Trube, 1984), vascular smooth muscle cells (Karkanis et al., 2003; Park et al., 2008; Quayle et al., 1993), endothelial progenitor cells (Quayle et al., 1993; Zhang et al., 2019) and microglial cells (Gattlen et al., 2020). We first adopted a real-time dynamic detection of peritoneal macrophage V\textsubscript{m} for 3 min by patch clamp experiments because of a relatively slow inhibitory effect of ML133 on Kir2.1 (Figs. 1B and 1C). The V\textsubscript{m} of peritoneal macrophages (-38.4 ± 2.8 mV) was not changed when Kir2.1 was blocked by ML133 (-39.4 ± 3.4 mV) in the resting state. However, the V\textsubscript{m} of LPS-stimulated macrophages was dramatically changed from -42.8 ± 2.9 mV to a much more depolarized V\textsubscript{m} of +12.3 ± 1.9 mV by ML133, and to a lesser extent, to -29.7 ± 2.3 mV due to Kir2.1 deficiency (Figs. 1G and S1H). Moreover, the effect of ML133 was abolished in Kir2.1-deficient macrophages (Figs. 1G and S1H), further indicating the
specicity of ML133 on Kir2.1. We also performed patch clamp recording for a short period of time of 30 seconds and found the V_m of LPS-stimulated macrophages (-31.9 ± 1.6 mV) was depolarized to -17.3 ± 1.1 mV due to Kir2.1 deficiency (Fig. 1H). Of note, the V_m of resting macrophages was also depolarized by Kir2.1 deficiency under this condition (Fig. 1H), suggesting a discrepancy in the depolarization of resting macrophages between ML133 (acute blockade) and Kir2.1 deficiency (long-term absence). Together, we conclude that Kir2.1 plays a critical role in regulating the membrane potential of inflammatory macrophages.

**Kir2.1 loss-of-function suppresses LPS- and infection-induced inflammatory cytokines and pathological inflammation**

To explore the role of Kir2.1 in LPS-induced inflammation, we first treated LPS-stimulated peritoneal macrophages with ML133 and found little cytotoxicity of ML133 (Figure S2A). Strikingly, ML133 dose-dependently inhibited the LPS-induced IL-1β, but not the TNF-α (Figs. 2A and 2B). Blockade of Kir2.1 by Ba^{2+} gave similar results (Figure S2B). To more broadly explore the effect of ML133, we performed RNA-seq analysis in BMDMs and found that, while Il1b was the gene most decreased by ML133, a cluster of LPS-induced inflammatory genes including Il1a, Il18, Il12a, and Il6 were also decreased, but Tnf was still not affected (Figs. 2C and S2C). Gene set enrichment analysis (GSEA) of the RNA-seq data showed a striking enrichment of ‘inflammatory response’ after ML133 treatment (Fig. 2D). Next, we used several genetic strategies to further investigate the role of Kir2.1 in LPS-induced inflammation. Silencing of Kcnj2 by small-interfering RNAs (siRNAs) in BMDMs or stably expressing shRNAs in immortalized BMDMs (iBMDMs) suppressed the LPS-induced IL-1β, but not the TNF-α (Figures S2D-S2G). Peritoneal macrophages from Lyz2-cre-Kcnj2^{f/f} mice similarly showed a reduction in LPS-induced IL-1β compared to wild-type mice (Fig. 2E). We also treated Kir2.1-depleted peritoneal macrophages with ML133 and found the inhibitory effect of ML133 on LPS-induced IL-1α and IL-1β was greatly impaired, further suggesting the specificity of ML133 through Kir2.1 (Figures S2H). To address the function of Kir2.1 in vivo, we used an LPS-induced sepsis model which largely reflects the inflammatory functions of monocytes/macrophages. ML133 or Kir2.1 deficiency significantly decreased the serum levels of IL-1β and increased the survival of mice (Figs. 2F-2I). The decreased TNF-α levels in Lyz2-cre-Kcnj2^{f/f} mice are probably due to the contribution of IL-1β to TNF-α production in vivo (Fig. 2H) (Tannahill et al., 2013). When infected with Gram-negative bacteria, such as *Escherichia coli* and *S. typhimurium* (strain SL1344), ML133 or Kir2.1 deficiency similarly led to decreased IL-1β and IL-1α production both in vitro and in vivo, with less effect on the TNF-α (Figs. 2J, 2K, S2I, and S2J). Last, to strengthen the evidence for an anti-inflammatory effect of Kir2.1 blockade in samples from patients, we used synovial fluid cells from gouty patients. Gout is an inflammatory form of arthritis and IL-1 inhibitors are effective in treating patients with acute and chronic gout (Jiang et al., 2017). We found that ML133 prevented IL-1β production in freshly-isolated synovial fluid cells from gouty patients (Fig. 2L). These data together indicate that Kir2.1 stimulates LPS-induced inflammation and reveal a potential anti-inflammatory strategy by targeting Kir2.1.
Kir2.1 promotes LPS-induced glucose uptake and consumption in inflammatory macrophages

Given the inconsistent inhibition of LPS-induced IL-1β and TNF-α, we predicted that Kir2.1 blockade would not affect the general signaling pathways mediated by TLR4, including the NF-κB and MAPK activation essential for both cytokines upon LPS stimulation. As predicted, ML133 or Kir2.1 deficiency had little effect on LPS-induced NF-κB and MAPK activation (Figures S3A-S3C). Accumulating evidence indicates that the Warburg effect of aerobic glycolysis plays critical roles in driving inflammatory macrophages, in particular the LPS-induced IL-1β production (Adamik et al., 2013; O'Neill et al., 2016). We thus further investigated whether Kir2.1 drives inflammatory macrophages by modulating LPS-induced metabolic reprogramming. We first measured real-time changes in the extracellular acidification rate (ECAR) and found ML133 or Kir2.1 deficiency led to a decrease in the LPS-induced long-term commitment to glycolysis (Fig. 3A). Moreover, both unbiased metabolomics profiling and a targeted metabolomics approach revealed that ML133 decreased the LPS-induced accumulation of metabolites representing aerobic glycolysis (Figs. 3B and 3C), suggesting a role of Kir2.1 in supporting a constant glycolysis during inflammation. Akt-mTORC1 signaling and HIF-1α activation were reported to modulate LPS-induced aerobic glycolysis and IL-1β production (Everts et al., 2014; Mills et al., 2016; Mills et al., 2018; Palsson-McDermott et al., 2015b; Tannahill et al., 2013). We found ML133 or Kir2.1 depletion had little effect on the early LPS induction of Akt-mTORC1 signaling in light of the phosphorylation of Akt and S6 (Figures S3A, S3B, and S3D). Moreover, GSEA of RNA-seq data showed that ‘mTORC1 signaling’ and ‘HIF-1α targets’ were not particularly enriched by ML133 (Figures S3E and S3F). Given that macrophages require a constant supply of extracellular glucose to support their intracellular metabolic reprogramming (Freemerman et al., 2014; Gamelli et al., 1996), we next considered the possibility that Kir2.1 promotes glucose import during LPS stimulation. Strikingly, LPS-induced glucose uptake in peritoneal macrophages was dose-dependently inhibited by ML133 in vitro (Fig. 3D). Similar results were obtained in peritoneal macrophages from Lyz2-cre-Kcnj2<sup>f/f</sup> mice compared to wild-type macrophages (Fig. 3E). Furthermore, we assessed this effect of Kir2.1 in vivo by measuring glucose uptake of macrophages in peritoneal exudate cells (PECs) and inflammatory monocytes in peripheral mononuclear cells (PBMCs) after intraperitoneal LPS challenge. Consistently, LPS-induced glucose uptake by these cells was significantly lower in Lyz2-cre-Kcnj2<sup>f/f</sup> mice than that in wild-type mice (Figs. 3F and 3G). Together, these data suggest that Kir2.1 promotes glucose uptake and consumption in inflammatory macrophages.

Kir2.1 supports glycolysis offshoots and SGOC metabolism in inflammatory macrophages and its loss-of-function leads to an amino acid starvation phenotype

A key mechanism for LPS-induced glycolysis is the induction of dimeric PKM2, which enters into a complex with HIF-1α to drive IL-1β expression (Palsson-McDermott et al., 2015a). As HIF-1α activation was minimally affected by ML133 (Figure S3F), we considered other mechanisms underlying the
impaired glycolysis by Kir2.1 blockade. While often represented as a linear metabolic flux, glycolysis-derived carbons also feed into several offshoots and integrate into different biosynthetic pathways (Chaneton et al., 2012; Keller et al., 2012). The pentose phosphate pathway (PPP) generates pentose sugars for nucleotide synthesis, and another three-step offshoot, the serine synthesis pathway (SSP) diverts glucose-derived carbons into serine, which is further integrated into the serine, glycine, one-carbon (SGOC) metabolic network that includes the folate and methionine cycles (Newman and Maddocks, 2017; Yang and Vousden, 2016) (Fig. 4A). Strikingly, unbiased metabolomics profiling revealed that the key metabolites 3-phosphoserine (3PS, representing the SSP) and ribose 5-phosphate (R5P, representing the PPP), increased in response to LPS but were decreased by ML133 (Fig. 4B). In addition, the enzymes mediating the three-step SSP – Phgdh, Psat1, and Psph – (Fig. 4A), and genes previously described as master regulators of the SSP (Yang and Vousden, 2016), such as Atf4 and Mdm2, were all upregulated by ML133 (Fig. 4C), showing a phenotype similar to a compensatory increase in the SSP upon serine starvation (Maddocks et al., 2013; Ye et al., 2012). Many enzymes involved in SGOC metabolism were also upregulated in ML133-treated inflammatory macrophages (Figure S4A). Moreover, GSEA of RNA-seq data showed that ML133-treated inflammatory macrophages were enriched in ‘amino acid transport’ and ‘SGOC metabolism’ (Fig. 4D). Using strategies of pharmacological inhibitor (acute blockade) and genetic depletion (long-term absence) may lead to certain discrepancies in downstream cellular and molecular mechanisms. To reveal the common mechanisms of both ML133 and Kir2.1 deficiency, we analyzed together the RNA-seq data from both ML133-treated and Kir2.1-depleted macrophages upon LPS stimulation. We found 236 overlapped downregulated and 163 overlapped upregulated genes (Fig. 4E). Pathway analysis revealed that the downregulated pathways affected by both ML133 and Kir2.1-depletion were mostly related to ‘inflammatory response’ and ‘response to LPS or IL-1’ (Fig. 4F). Strikingly, the upregulated pathways were related to ‘response to amino acid starvation’ and ‘regulation of translation’ (Fig. 4F). Among these upregulated genes, we found several master sensors and regulators in response to amino acid starvation, including GCN2, PERK, IMPACT, and SLC38A2 (Figure S4B) (Broer and Broer, 2017).

We next used U-[13C]-glucose tracers to confirm the changes in the SSP and SGOC metabolism when Kir2.1 was blocked during LPS stimulation. ML133 significantly decreased intracellular m + 6 glucose, m + 3 serine, and m + 2 glycine, suggesting that Kir2.1 supports glucose uptake and the channeling of glucose-derived carbons into the SSP upon LPS stimulation (Figs. 4G and 4H). To assess the contribution of this suppressed SSP to the anti-inflammatory phenotype of ML133, we supplemented cell penetrable 3-PG (Finder and Hardin, 1999; Yang et al., 2014), which is more channeled into the SSP from glycolysis under serine-depleted conditions (Chaneton et al., 2012; Maddocks et al., 2013; Zhang et al., 2012), and found the suppressed IL-1β production by ML133 was partially restored (Figure S4C). Of note, most of the accumulation of intracellular serine and methionine was unlabeled (m + 0) (Figures S4D and S4E), indicating their accumulation is largely fueled by exogenous import during inflammation. Strikingly, the LPS-induced accumulation of unlabeled serine, glycine, and methionine (key amino acids fueling SGOC metabolism (Locasale, 2013)) was also greatly blocked by ML133 (Fig. 4I). Similarly, we used U-[13C]-serine tracers and found a decrease of m + 3 serine, m + 2 glycine, and m + 0 methionine due to Kir2.1
deficiency, recapitulating the impeded serine and methionine uptake (Figure S4F). These results together highlight a role of Kir2.1 in the nutrient supply fueling glycolysis offshoots and SGOC metabolism during LPS-induced inflammation.

**Kir2.1 supports S-adenosylmethionine generation and configures histone methylation in inflammatory macrophages**

Through coupling with the methionine cycle, the SGOC metabolic network acts as an integrator of nutrient status to generate diverse outputs, including the primary methyl donor S-adenosylmethionine (SAM) (Locasale, 2013; Yang and Vousden, 2016; Yu et al., 2019). We and others have previously showed that glucose can provide carbons to feed SAM generation through de novo ATP synthesis (Maddocks et al., 2016; Newman and Maddocks, 2017; Yu et al., 2019) (Fig. 5A). Using U-[^13]C-glucose, we found ML133 led to a decrease in LPS-induced m + 5 to 9 SAM (via both the PPP and SSP) and its demethylation product S-adenosylhomocysteine (SAH) (Figs. 5A and 5B), as well as m + 5 to 9 ATP (Figure S5A). The total amount of SAM was also reduced by ML133 (Figure S5B). Using U-[^13]C-serine tracers, we obtained similar results that ML133 or Kir2.1 deficiency decreased the LPS-induced incorporation of U-[^13]C-serine-derived carbons into m + 1 to 4 SAM, as well as the total amount of SAM (Figs. 5C, 5D, and S5C). Given that Kir2.1 blockade led to impaired nutrient uptake, exogenous glucose or amino acids (serine, glycine, or methionine) that donate carbons into SGOC metabolism only had mild rescue effects on the suppressed IL-1β production by ML133 (Figures S5D-S5F). However, addition of SAM dose-dependently restored this suppressed IL-1β production (Figs. 5E and 5F), suggesting a role of Kir2.1 in supporting LPS-induced SAM generation by providing the supply of extracellular nutrients.

SAM is the universal methyl donor for all methylation reactions in cells, which plays critical roles in the chromatin state and gene transcription (Goll and Bestor, 2005). SAM availability can directly modulate several epigenetic methylation marks required for the maintenance of downstream gene transcription (Mentch et al., 2015; Mews et al., 2014; Shiraki et al., 2014; Shyh-Chang et al., 2013). RNA-seq analysis of the expression of 183 annotated SAM-dependent methyltransferase enzymes (Kottakis et al., 2016) revealed that Kir2.1 deficiency upregulated all methyltransferases for histone H3 methylation at lysine 36 (H3K36me) (Wagner and Carpenter, 2012) compared to those for other histone methylation marks (Figs. 5G, 5H, and S5G). Among the most upregulated methyltransferase genes, ASH1L and NSD1 have specific mono- and dimethylase activity for H3K36, and SETD2 is the only reported trimethylase catalyzing the trimethylation of H3K36 (Fig. 5G) (Chen et al., 2017; Wagner and Carpenter, 2012). H3K36me3 is one of the most dynamic histone methylation marks and its main distribution appears in a wide range of gene body regions as a SAM ‘sink’ (Ye et al., 2017), making it more sensitive to SAM availability for downstream gene expression (Wagner and Carpenter, 2012). To test whether Kir2.1-induced SAM generation fuels histone methylation such as H3K36me3 upon LPS stimulation, we coupled chromatin immunoprecipitation with quantitative PCR (ChIP-qPCR) and found that LPS-induced
occupancy of H3K36me3 in \textit{Il1b} gene-body regions farther from the 5' end (Adamik et al., 2013) was significantly decreased by ML133 or Kir2.1 deficiency (Figs. 5I and S5H). In addition, LPS-induced H3K36me3 enrichment in the gene-body regions of other inflammatory factors including \textit{Il1a}, \textit{Il18}, and \textit{Cxcl10} were also decreased (Figs. 5J and S5H). In contrast, H3K36me3 enrichment in the gene-body regions of \textit{Tnf} was unaffected in these experiments (Figure S5I). We thus conclude that deregulation of histone methylation marks, at least H3K36me3, contributes to the anti-inflammatory outcome of Kir2.1 loss-of-function.

**Kir2.1-mediated regulation of membrane potential orchestrates metabolic-epigenetic reprogramming in inflammatory macrophages**

Given that monovalent cations such as K\textsuperscript{+} regulate the membrane potential, which indirectly controls the flux of Ca\textsuperscript{2+} and intracellular signaling pathways (Feske et al., 2015; Franchini et al., 2004; Lam and Schlichter, 2015), we initially investigated whether Kir2.1 stimulates inflammatory macrophages by modulating Ca\textsuperscript{2+} homeostasis. In LPS-stimulated BMDMs loaded with Ca\textsuperscript{2+}-sensitive fluorescent dye Fluo-4-AM, we found an elevation of cytosolic Ca\textsuperscript{2+} level in response to ML133 (Figure S6A). However, when we used BAPTA or EGTA to chelate the extracellular Ca\textsuperscript{2+} or BAPTA-AM to chelate the intracellular Ca\textsuperscript{2+}, the inhibitory effect of ML133 on LPS-induced IL-1\textbeta production was not affected (Figures S6B-S6F). Notably, BAPTA-AM showed an additional inhibitory effect on the production of IL-1\textbeta (Figures S6E and 6F), suggesting a separate contribution of intracellular Ca\textsuperscript{2+} in parallel to ML133. Consistent with idea, when we depleted ER Ca\textsuperscript{2+} store by thapsigargin (TG), although TG alone was able to reduce the production of IL-1\textbeta, ML133 still had a similar inhibitory efficiency on IL-1\textbeta production (Figure S6G). Moreover, the effect of ML133 was independent of the concentrations of extracellular Ca\textsuperscript{2+} (0, 1.2, 2.4, and 3.6 mM) or the other divalent cations Mg\textsuperscript{2+} and Mn\textsuperscript{2+} (Figures S6H and S6I). We thus conclude that the changes in Ca\textsuperscript{2+} homeostasis is not required for the anti-inflammatory outcome of Kir2.1 blockade.

Given the critical role of Kir2.1 in maintaining V\textsubscript{m} of inflammatory macrophages (Figs. 1E and 1F), we considered the possibility that Kir2.1 promotes LPS-induced inflammation by its regulation of V\textsubscript{m}. We first examine whether V\textsubscript{m} depolarization by other means results in an analogous suppression of inflammatory macrophages. Increased extracellular K\textsuperscript{+} ([K\textsuperscript{+}]\textsubscript{e}) is widely used to depolarize V\textsubscript{m}. We found the V\textsubscript{m} of both resting and LPS-stimulated macrophages was depolarized by elevated [K\textsuperscript{+}]\textsubscript{e} as determined by either patch clamp experiments or a membrane sensitive fluorochrome DiBAC4(3) (Eil et al., 2016) (Figures S6J and S6K). Strikingly, elevation of [K\textsuperscript{+}]\textsubscript{e} similarly decreased the LPS-induced IL-1\textbeta without affecting the TNF-\alpha (Fig. 6A), as well as glucose uptake detected by intracellular m+6 glucose using U-[13C]-glucose tracers (Fig. 6B). The effect of elevated [K\textsuperscript{+}]\textsubscript{e} was not due to an osmotic effect, as choline chloride or mannitol did not induce a similar suppression of IL-1\textbeta production (Figure S6L). Gramicidin is another widely used strategy to depolarize V\textsubscript{m} by forming pores permeable to both K\textsuperscript{+} and Na\textsuperscript{+} in the plasma membrane (Munoz-Planillo et al., 2013). As expected, gramicidin depolarized the V\textsubscript{m} of inflammatory
macrophages (Figure S6K), and it recapitulated the suppressed LPS-induced IL-1β production and glucose uptake, but not the TNF-α (Figs. 6C and 6D). Furthermore, using U-[13C]-glucose, we found that elevated [K+]_e and gramicidin led to a decrease in LPS-induced m + 3 serine, m + 5 to 9 ATP, and m + 5 to 9 SAM, as well as unlabeled m + 0 serine and methionine (Figs. 6E and 6F). Elevated [K+]_e also reduced the enrichment of H3K36me3 in the gene-body regions of Il1b, Il1a, Il18, and Cxcl10 loci (Figs. 6G and 6H). Together, these data suggest that the maintenance of membrane potential may be a common mechanism that orchestrates metabolic-epigenetic reprogramming in inflammatory macrophages.

**Kir2.1 supports nutrient supply by promoting the surface expression of nutrient transporters during LPS-induced inflammation**

Nutrient import is the proximal step at which intracellular metabolism can be tightly regulated. Cells require adaptations in nutrient transport mechanisms to meet different metabolic demands (McCracken and Edinger, 2013). Consistent with this idea, glucose transporter expression levels are elevated in proliferating cells and in a wide variety of tumor types (Adekola et al., 2012). To investigate the mechanisms underlying the regulation of nutrient import by Kir2.1 in inflammatory macrophages, we first analyzed the RNA-seq data with the KEGG (Kyoto Encyclopedia of Genes and Genomes) and Reactome pathway databases. Strikingly, apart from pathways related to inflammation, Kir2.1 blockade led to a strong enrichment for the pathways related to endocytosis, vesicle transport, and membrane trafficking (Figs. 7A and 7B). Given the critical roles of surface transporters in the regulation of nutrient import, we next evaluated the potential contribution of Kir2.1 to the surface expression of nutrient transporters fueling glucose and amino acid metabolism. Glucose transporter 1 (GLUT1) is the primary transporter that rewires glucose metabolism in LPS-stimulated macrophages (Freemerman et al., 2014). While the mRNA expression was not affected (Figure S7A), ML133 dose-dependently decreased the membrane GLUT1 expression in LPS-stimulated macrophages examined by flow cytometry (Fig. 7C). This effect was also confirmed by immunoblotting for biotinylated cell-surface proteins in ML133-treated or Kir2.1-depleted inflammatory macrophages (Figs. 7C and S7B). Consistently, this decreased membrane GLUT1 expression was recapitulated by elevated [K+]_e (Fig. 7D). Among the membrane amino acid transporters, L-type amino acid transporter (LAT1) together with 4F2 cell-surface antigen heavy chain (4F2hc, also known as CD98) transport large neutral amino acids including methionine (McCracken and Edinger, 2013; Yanagida et al., 2001). Both ML133 and elevated [K+]_e decreased the surface expression of CD98 in LPS-stimulated macrophages (Fig. 7E), and depolarization by gramicidin gave similar results including GLUT1 and CD98 (Figs. 7F and 7G). These results further indicated the contribution of membrane potential to the nutrient import in inflammatory macrophages.

The surface expression and activity of glucose transporters were regulated by lipid rafts-mediated trafficking and internalization (Kumar et al., 2004; Michel and Bakovic, 2007; Yan et al., 2018). Nearly 80% of GLUT4 is internalized by a lipid rafts-dependent mechanism which can be blocked by the cholesterol-chelating drug nystatin (Blot and McGraw, 2006; Ros-Baro et al., 2001). Without affecting TLR4-mediated
downstream signaling upon LPS stimulation (Figure S7C), nystatin partially restored the decreased level of surface GLUT1, the suppressed glucose uptake, and IL-1β production but not TNF-α when Kir2.1 was blocked by ML133 or depleted (Figs. 7H-7K and S7D). Nystatin also rescued the enrichment of H3K36me3 in the gene-body regions of Il1a, Il1b, Il18, and Cxcl10 loci in Kir2.1-depleted inflammatory macrophages (Figure S7E). Last, nutrient transporters including GLUT1, CD98, and LAT1 are ARF6/GRP1 cargo proteins that can be recycled back to the plasma membrane via the tubular recycling endosome (Eyster et al., 2009; Finicle et al., 2018; Maldonado-Baez et al., 2013; McCracken and Edinger, 2013). Strikingly, a constitutively active mutant of GRP1 (S155D/T280D, GRP1 DD mutant), which forces the recycling of these transporters back to the plasma membrane and prevents their loss (Finicle et al., 2018; Li et al., 2012), largely restored the ML133-induced suppression of IL-1β (Fig. 7K), further strengthening the contribution of surface nutrient transporters loss to the anti-inflammatory outcome of Kir2.1 blockade.

Discussion

The coordination between metabolism and epigenetics has been proposed as a key mechanism in immunity (Chisolm and Weinmann, 2018; Phan et al., 2017), but how they are supported by nutrient availability remains poorly understood. In contrast to tumor cells or activated T cells, LPS stimulation does not induce rapid proliferation of macrophages. However, it also creates increasing metabolic demands that would be supported by extracellular nutrients. The present work provides evidence for fundamental membrane potential control of optimizing nutrient acquisition and utilization for intracellular metabolic and epigenetic reprogramming in inflammatory macrophages (Figure S7F). We propose a model whereby Kir2.1-mediated adaptation in nutrient import (including glucose and amino acids) fueling SAM generation and epigenetically driving LPS-induced inflammation. Mechanistically, we reveal an unexpected role of Kir2.1 in the adaptive regulation of surface nutrient transporters levels in LPS-stimulated macrophages.

Recently, an ionic checkpoint of the elevated [K+]e that blocks T cell function has been found within the tumor microenvironment, and this elevated [K+]e impairs T cell effector functions but promotes T cell stemness (Eil et al., 2016; Vodnala et al., 2019). Augmenting K+ efflux in tumor-specific T cells improves effector functions and enhances tumor clearance and survival (Eil et al., 2016), while exposure to increased [K+]e enhances the persistence and anti-tumor function of T cells (Vodnala et al., 2019). Changes in environmental K+ concentrations have also been reported in different inflammation-related pathophysiological situations (Billman, 1994; Gido et al., 1997; Levite et al., 2000). Disturbances in the concentrations of different ions are closely associated with critical inflammatory disorders, including sepsis and severe burns (Lee, 2010). Accordingly, modulating the ionic regulation of immune cells may represent a potential therapeutic approach for treating inflammatory diseases. Consistent with this idea, Kir2.1 loss-of-function impaired LPS- and infection-induced inflammatory gene expression and rendered mice resistant to sepsis. Moreover, ML133 suppressed IL-1β production in freshly-isolated synovial fluid cells from gouty patients, suggesting Kir2.1 as a potential target for treating inflammatory diseases.
However, given the expression of Kir2.1 also in other cell types such as endothelial cells, much work remains to determine the systemic effect of targeting Kir2.1 during inflammatory diseases.

Besides the suppressed LPS-induced inflammation, another key phenotype induced by both ML133 and Kir2.1 depletion is nutrient starvation. Several master regulators in response to amino acid starvation such as GCN2, PERK, IMPACT, and SLC38A2 (Broer and Broer, 2017) were also upregulated. There is growing evidence that local nutrient limitations at immune effector sites can be obstacles to both antimicrobial and anti-tumor immunity (Olenchock et al., 2017). Understanding how the interactions among microenvironment factors, immune cell nutrient demands, and cellular metabolic state shape the “metabolic phenotyping” is critical to obtain a more complete understanding of immune cell metabolism. Moreover, instead of acting alone, different types of nutrients would synergistically feed the generation of specific immunometabolites to modulate immune functions. Although other metabolic pathways may also be regulated by Kir2.1 as the low specificity and selectivity of amino acid transporters, our study unveils an ionic control on the synergism of glucose and SGOC metabolism to epigenetically drive LPS-induced inflammation by promoting SAM generation. As a universal methyl donor, the dynamic production and utilization of SAM is critical for the regulation of gene expression by methylation reactions. As a SAM 'sink', H3K36me is one of the more dynamic histone methylation marks required for transcription elongation and splicing (Wagner and Carpenter, 2012; Ye et al., 2017). We found a different sensitivity to the SAM supply and H3K36me regulation between IL-1β (also a set of inflammatory genes) and TNF-α, which may provide an explanation for the distinct sensitivity to the metabolic control of these two important inflammatory cytokines during inflammation.

Our study additionally highlights a Kir2.1 control on the adaptations in nutrient uptake of inflammatory macrophages by dynamically regulating the surface expression of nutrient transporters, including GLUT1 (glucose) and CD98 (amino acids). Given the apical position of the nutrient transporters in metabolic pathways, these proteins are intriguing pharmacologic targets for cancer treatment. The strategy of starving cancer cells of required amino acids including serine and methionine has been proved to be effective both in mice and humans (Gao et al., 2019; Locasale, 2013; Maddocks et al., 2013). More recently, efforts have been directed at blocking nutrient import rather than availability. Several compounds that inhibit nutrient transport in tumor cells can prevent proliferation and induce cell death (McCracken and Edinger, 2013). Given that $V_m$ regulation on the surface retention of nutrient transporters may be a common mechanism during inflammation and cancer, it will be important to further investigate the roles of other ionic channels on nutrient uptake, as well as the nutrients acquired through those less well characterized transporters.

In sum, this current study unveils a Kir2.1 control acting on macrophage activation and shows that targeting ion channels may have implications for treating inflammatory diseases.

### Experimental Model And Subject Details

#### Mice
C57BL/6 mice were purchased from the Model Animal Research Center of Nanjing University, Lyz2-cre mice were purchased from the Jackson Laboratory, and the \textit{Kcnj2}^{ff} mice were kind gifts from Professor Mark T. Nelson of the University of Vermont. \textit{Kcnj2}^{ff} mice were crossed with Lyz2-cre mice to obtain Lyz2-cre-\textit{Kcnj2}^{ff} mice. Animals were housed in a specific pathogen-free facility in the University Laboratory Animal Center. The animal experimental protocols were approved by the Review Committee of Zhejiang University School of Medicine and were in compliance with institutional guidelines.

**Cells**

HEK293T cells were from ATCC and cultured in Dulbecco's modified Eagle's medium (DMEM).

To obtain Mouse peritoneal macrophages, in day 0, mice were injected peritoneally with 2.5 ml of 4% thioglycolate (Merck) medium, and in day 3 to 5, peritoneal macrophages were obtained by flush the peritoneal cavity with PBS or DMEM medium. Each mouse usually yields approx. $2 \times 10^7$ cells, and the non-adherent cells were discarded after macrophages adhere.

Bone marrow cells were flushed from tibias and femurs with cold DMEM and cultured in DMEM supplemented with 10% fetal bovine serum (FBS), 1% penicillin/streptomycin, and 10 ng/ml macrophage colony-stimulating factor (PeproTech) to generate BMDMs.

The iBMDMs were a kind gift from Prof. Shao (National Institute of Biological Sciences, China). J774.1 (from ATCC) and iBMDMs were cultured in DMEM supplemented with 10% FBS and 1% penicillin/streptomycin.

Synovial fluid cells ($5 \times 10^5$/well) were seeded in 12-well plates in RPMI 1640 supplemented with 10% FBS. They were stimulated with 100 ng/ml LPS and inhibitors as indicated for 12 h. Then sample supernatants were used for IL-1β measurements using ELISA.

**Human Subjects**

Synovial fluid (approximate 4–5 ml) was obtained from two gouty patients (a 36-year-old man and a 56-year-old man) with serum uric acid levels > 500 mmol/l and knee effusion. The patients were not involved in previous procedures or drug tests. To use these clinical materials for research purposes, prior patient written informed consent and approval from the Institutional Research Ethics Committee of The Second Affiliated Hospital of Zhejiang University School of Medicine were obtained (approval no. 2018-064). Synovial fluid cell ($5 \times 10^5$) were seeded in 12-well plates in RPMI 1640 supplemented with 10% FBS. They were stimulated with 100 ng/ml LPS and inhibitors as indicated for 12 h. Then sample supernatants were used for IL-1β detection using ELISA.

**Reagents**

LPS from \textit{Escherichia coli} O111:B4, U-$^{13}$C]-serine, serine, glycine, methionine, SAM, 3-PG and BaCl$_2$ were from Sigma; ML133 was from Selleck; amino-acid- and KCl- deficient DMEM was from Shanghai
BasalMedia Technologies; dialyzed FBS was from Biological Industries; and U-[\textsuperscript{13}C]-glucose was from Cambridge Isotope Laboratories; 2NBDG was from Cayman; Nystatin was from Sigma.

The following Biolegend antibodies were used: pro-IL-1\(\beta\) (AF-401-NA,) was from R&D systems; anti-Kir2.1 (19965-1-AP) was from Proteintech; anti-phospho-p65 (#3033), anti-p65 (#8242), anti-phospho-I\(\kappa\)B\(\alpha\) (#2859), anti-I\(\kappa\)B\(\alpha\) (#4814), anti-phospho-Erk (#4370), anti-Erk (#4695), anti-phospho-JNK (#9255), anti-JNK (#9252), anti-phospho-P38 (#4511), anti-P38 (#8690), anti-phospho-AKT (#4060), anti-AKT (#2920), anti-phospho-P38 (#4511), anti-P38 (#8690), anti-phospho-AKT (#4060), anti-AKT (#2920), anti-phospho-JNK (#9252), anti-phospho-JNK (#9255), anti-phospho-P38 (#4511), anti-P38 (#8690), anti-phospho-AKT (#4060), anti-AKT (#2920), anti-H3K36me3 (#4909) were from Cell Signaling Technology; anti-GLUT1 (ab115730) was from Abcam, PE anti-mouse CD98 (#128207) was from Biolegend, anti-\(\beta\)-actin (M1210-2), anti-rabbit IgG-HRP (HA1001), anti-mouse IgG-HRP (HA1006), and anti-goat IgG-HRP (HA1005) were from HuaBio, DyLight549 goat anti-rabbit IgG [H+L] (70-GAR5492) was from MultiSciences, and the ChIP kit (#9005) was from Cell Signaling Technology.

**Method Details**

**Immunoblot Analysis**

Cells were lysed in 2 \(\times\) SDS buffer (100 mM Tris-HCl, 4% SDS, 20% glycerol, 2% 2-mercaptoethanol, and 0.05% bromophenol blue) followed by boiling for 10 min. Then samples were separated by SDS-PAGE on 12% gels, after which the proteins were transferred to nitrocellulose membranes (#28637358, Pall). The membranes were blocked for 1 h in blocking buffer (5% skimmed milk and 0.1% Tween 20 in TBS), and then incubated with primary antibodies in 5% BSA overnight. The membranes were incubated with secondary antibodies in 0.1% Tween 20 in TBS at room temperature for 1 h. To detect proteins, we used ECL blotting reagents (Thermo Fisher).

**In vivo LPS challenge**

Mice were injected intraperitoneally with LPS (25 mg/kg body weight) alone or along with ML133 (30 mg/kg body weight). In the sepsis model, mice were sacrificed 4 h after LPS challenge, and the serum levels of IL-1\(\beta\) and TNF-\(\alpha\) were measured by ELISA (Thermo Fisher) according to the manufacturer’s instructions. For mouse survival rate analysis, mice were injected intraperitoneally with LPS (20 mg/kg body weight) alone or along with ML133 (30 mg/kg body weight), then survival rates were analyzed.

**Bacterial infection**

For \textit{in vitro} bacterial infection assay, mouse peritoneal macrophages were seeded in 12 well plate (5 \(\times\) \(10^5\)/well) and infected with 5 \(\times\) \(10^6\) \textit{E.coli} or \textit{salmonella} SL1344 for 6 h in the presence or absence of ML133 (25 \(\mu\)M) followed by qPCR analysis of inflammatory genes transcription. For \textit{in vivo} bacterial infection assay, 8-week-old mice were injected with 1 \(\times\) \(10^7\) bacteria peritoneally in the presence or absence of ML133 (30 mg/kg), 6 hours later, the mice were sacrificed and the serum levels of IL-1\(\beta\) and TNF-\(\alpha\) were measured by ELISA (Thermo Fisher) according to the manufacturer’s instructions.
Quantitative PCR (qPCR)

RNA was extracted using RNAiso Plus reagent (Takara). Complementary DNA was synthesized using HiScript® II Reverse Transcriptase (Vazyme Biotech) according to the manufacturer’s protocol. qPCR was performed using SYBR Green (Vazyme Biotech) on a CFX96 Touch Real Time PCR (BioRad). The PCR program was initial denaturation at 95 °C for 2 min, then cDNA amplification for 40 cycles at 95 °C for 30 s and 60 °C for 30 s. The samples were individually normalized to Gapdh.

The primers are listed in the table.

| Gene | Forward 5’to 3’       | Reverse 5’to 3’       |
|------|-----------------------|-----------------------|
| Gapdh | AGGTCGGTGTGAACGGATTTG | TGTAGACCATGTAGTTGAGGTCA |
| Il1b  | ATCAACAAACAGTATATTCTCAT | GGGTGCCCTTTTCATTAC |
| Tnf   | CCTGTAGCCACGTCTGTA    | GGGAGTAGACAAGGTACAACCC |
| Il1a  | GGAGAAGACCAGCCGTGGTTGCT | CCGTGCAGGTCACCGACTT |

Measurement of extracellular acidification rate (ECAR)

The Seahorse XF96 analyzer (Agilent Technologies) was used to measure extracellular ECAR. BMDMs were seeded on XF96 plates at 8 × 10^4 cells/well one day prior to the XF assay. On the day of assay, the medium was replaced with assay medium composed of XF Base Medium without phenol red (Agilent Technologies, 103335-100) supplemented with 10 mM glucose, 2 mM L-glutamine, 1 mM sodium pyruvate, adjusted to pH 7.4 and incubated at 37 °C without CO₂ 45 min prior to XF assay. The assay protocol was as follows: baseline measurement with 5 cycles (mix 3 min, wait 0 min, measure 3 min); then LPS and/or ML133 was injected with the final concentration of 500 ng/ml or 25 µM respectively and measurements continued with 10–99 cycles (mix 3 min, wait 0 min, measure 3 min). Data shown are the mean ± SD, n = 6 technical replicates.

[^13]C-glucose and[^13]C-serine tracing

Mouse peritoneal macrophages (5 × 10^6/dish) were seeded in 60-mm dishes in complete DMEM medium (supplemented with 10% fetal bovine serum (FBS), 1% penicillin/streptomycin) to adhere. In metabolite tracing experiments, complete DMEM medium was replaced with glucose- or serine-deficient DMEM supplemented with 25 mM U[^13]C-glucose (Cambridge Isotope Laboratories) or 0.4 mM U[^13]C-serine (Sigma) (supplemented with 10% dialyzed fetal bovine serum (FBS), 1% penicillin/streptomycin). Cells were treated with 500 ng/ml LPS in the presence or absence of compounds as indicated for 6 hours. For metabolite extraction, cells were washed twice with PBS and once more with 0.9% NaCl. After completely aspirating the liquid, the plates were put on dry ice, 1 ml of 80% (v/v) methanol (pre-chilled to −80 °C) was added, and the plates were kept at −80 °C for 2 h. The plates were scraped on dry ice, the cell lysate/methanol mixture was transferred to a 2-ml tube on dry ice, then another 0.8 ml of 80% methanol...
was added to wash the plate and transfer the mixture to a tube. Each tube was centrifuged at 14,000 g for 20 min at 4 °C and the metabolite-containing supernatant was transferred to a new tube and lyophilized. Metabolites were analyzed using a TSQ Quantiva Ultra triple-quadrupole mass spectrometer coupled with an Ultimate 3000 UPLC system (Thermo Fisher, CA) equipped with a heated electrospray ionization probe. Chromatographic separation was done by gradient elution on a reversed-phase UPLC HSS T3 column (2.1 × 100 mm, 1.7 µm, Waters). Mobile phase A consisted of 2 mM perfluoroheptanoic acid in 100% H$_2$O, and mobile phase B of 100% acetonitrile. A 10-min gradient with a flow rate of 300 µL/min was used as follows: 0-1.5 min at 2% B; 1.5-5 min, 2–98% B; 5–7 min, 98% B; 7-7.1 min, 2% B; 7.1–10 min 2% B. The column chamber was held at 45 °C and the sample tray at 10 °C. Data were acquired in Selected Reaction Monitoring in positive/negative switch ion mode and optimal transitions are reported in the table as indicated below. Both the precursor and fragment ions were separately collected at a resolution of 0.7 FWHM. The source parameters were as follows: spray voltage, 3000 V; ion transfer tube temperature, 350 °C; vaporizer temperature, 300 °C; sheath gas flow rate, 35 Arb; auxiliary gas flow rate, 12 Arb. CID gas, 1.5 mTorr. Data analysis and quantitation were performed using Xcalibur 3.0.63 (Thermo Fisher, Carlsbad, CA). The liquid chromatography-mass spectrometry was done at the Metabolomics Facility of Technology Center for Protein Sciences, Tsinghua University, Beijing.

**Metabolite quantification and metabolomics**

Mouse peritoneal macrophages (5 × 10$^6$ /dish) were seeded in 60-mm dishes in complete DMEM medium (supplemented with 10% fetal bovine serum (FBS), 1% penicillin/streptomycin) to adhere. Cells were treated with 500 ng/ml LPS in the presence or absence of compounds as indicated for 6 hours. Then metabolites were extracted as above. For unbiased and targeted metabolomics, samples were analyzed at Applied Protein Technology (Shanghai, China).

For unbiased metabolomics, LC-MS/MS Analysis (HILIC/MS) were performed using an UHPLC (1290 Infinity LC, Agilent Technologies) coupled to a quadrupole time-of-flight (AB Sciex TripleTOF 6600). For HILIC separation, samples were analyzed using a 2.1 mm × 100 mm ACQUIY UPLC BEH 1.7 µm column (waters, Ireland). In both ESI positive and negative modes, the mobile phase contained A = 25 mM ammonium acetate and 25 mM ammonium hydroxide in water and B = acetonitrile. The gradient was 85% B for 1 min and was linearly reduced to 65% in 11 min, and then was reduced to 40% in 0.1 min and kept for 4 min, and then increased to 85% in 0.1 min, with a 5 min re-equilibration period employed. The ESI source conditions were set as follows: Ion Source Gas1 (Gas1) as 60, Ion Source Gas2 (Gas2) as 60, curtain gas (CUR) as 30, source temperature: 600°C, IonSpray Voltage Floating (ISVF) ± 5500 V. In MS only acquisition, the instrument was set to acquire over the m/z range 60-1000 Da, and the accumulation time for TOF MS scan was set at 0.20 s/spectra. In auto MS/MS acquisition, the instrument was set to acquire over the m/z range 25-1000 Da, and the accumulation time for product ion scan was set at 0.05 s/spectra. The product ion scan is acquired using information dependent acquisition (IDA) with high sensitivity mode selected. The collision energy (CE) was fixed at 35 V with ± 15 eV. Declustering potential (DP) was set as ± 60 V.
A targeted metabolic analysis was performed using an LC-MS/MS system. The dried metabolites were dissolved in 100 µL of acetonitrile/ H₂O (1:1, v/v) and centrifuged (13000 rpm) for 15 min. Electrospray ionization was conducted with an Agilent 1290 Infinity chromatography system and AB Sciex QTRAP 5500 mass spectrometer. NH₄COOH (15 mM) and acetonitrile were used as mobile phases A and B, respectively. A binary solvent gradient was used as follows: A, NH₄COOH; B, 0–18 min at 90–40% acetonitrile; 18-18.1 min at 40–90% acetonitrile; and 18.1–23 min at 90% acetonitrile. The LCMS/MS was operated in the negative mode under the following conditions: source temperature, 450 °C; ion source gas 1, 45; ion source gas 2, 45; curtain gas, 30; and ion spray voltage floating (ISVF), -4500 V.

RNA-sequencing and differential expressed genes analysis

Library preparation for Transcriptome sequencing. A total amount of 3 µg RNA per sample was used as input material for the RNA sample preparations. Sequencing libraries were generated using NEBNext® UltraTM RNA Library Prep Kit for Illumina® (NEB, USA) following manufacturer's recommendations and index codes were added to attribute sequences to each sample. Briefly, mRNA was purified from total RNA using poly-T oligo-attached magnetic beads. Fragmentation was carried out using divalent cations under elevated temperature in NEB. Next First Strand Synthesis Reaction Buffer (5×). First strand cDNA was synthesized using random hexamer primer and M-MuLV Reverse Transcriptase (RNase H⁻). Second strand cDNA synthesis was subsequently performed using DNA Polymerase I and RNase H. Remaining overhangs were converted into blunt ends via exonuclease/polymerase activities. After adenylation of 3' ends of DNA fragments, NEBNext Adaptor with hairpin loop structure were ligated to prepare for hybridization. In order to select cDNA fragments of preferentially 250 ~ 300 bp in length, the library fragments were purified with AMPure XP system (Beckman Coulter, Beverly, USA). Then 3 µl USER Enzyme (NEB, USA) was used with size-selected, adaptor-ligated cDNA at 37 °C for 15 min followed by 5 min at 95 °C before PCR. Then PCR was performed with Phusion High-Fidelity DNA polymerase, Universal PCR primers and Index (X) Primer. At last, PCR products were purified (AMPure XP system) and library quality was assessed on the Agilent Bioanalyzer 2100 system.

Clustering and sequencing. The clustering of the index-coded samples was performed on a cBot Cluster Generation System using TruSeq PE Cluster Kit v3-cBot-HS (Illumia) according to the manufacturer's instructions. After cluster generation, the library preparations were sequenced on an Illumina Hiseq platform and 125 bp/150 bp paired-end reads were generated.

Quantification of gene expression level. featureCounts v1.5.0-p3 was used to count the reads numbers mapped to each gene. And then FPKM of each gene was calculated based on the length of the gene and reads count mapped to this gene. FPKM, expected number of Fragments Per Kilobase of transcript sequence per Millions base pairs sequenced, considers the effect of sequencing depth and gene length for the reads count at the same time, and is currently the most commonly used method for estimating gene expression levels.

Differential expression analysis. Differential expression analysis of two conditions/groups (two biological replicates per condition) was performed using the DESeq2 R package (1.16.1). DESeq2 provide statistical
routines for determining differential expression in digital gene expression data using a model based on the negative binomial distribution. The resulting P-values were adjusted using the Benjamin and Hochberg's approach for controlling the false discovery rate. Genes with an adjusted P-value < 0.05 found by DESeq2 were assigned as differentially expressed.

**GO and KEGG enrichment analysis of differentially expressed genes.** Gene Ontology (GO) enrichment analysis of differentially expressed genes was implemented by the cluster Profiler R package, in which gene length bias was corrected. GO terms with corrected P-value less than 0.05 were considered significantly enriched by differential expressed genes.

Gene Set Enrichment Analysis (GSEA) (http://www.broadinstitute.org/gsea/index.jsp) of the expression data was used to assess enrichment of the KEGG as well as the SGOC genesets.

### Stable gene-knockdown with short-hairpin RNAs

We constructed pLVX-shRNAs plasmids targeting \( knj2 \). iBMDMs were spinfected with retrovirus encoding short-hairpin RNA for 90 min at 2500 rpm and 32 °C. Forty-eight hours after infection, the cells were selected by culture with 5 mg/mL puromycin (Sigma).

shRNAs used in the study.

| \( knj2 \)-1  | CCCATCACTATCTGTTGCACGAA |
| \( knj2 \)-2  | GCTCTTTGAAGAGAAACACTA |

### Gene-knockdown by siRNAs

For gene-knockdown experiments, cells were transfected with siRNAs (GenePharma, Shanghai) using RNAiMAX Transfection Reagent (Thermo Fisher Scientific). Briefly, cells were seeded in 12-well plates in complete DMEM to adhere, then washed with PBS. Cells were transfected with 20 µM siRNAs in 1 ml Opti-MEM (Thermo Fisher Scientific). After 6 h, the medium was changed to DMEM and the cells were cultured for another 2 days.

siRNAs used in the study:

| siRNA          | sense(5'-3') | antisense(5'-3') |
|---------------|-------------|-----------------|
| siRNA-control | UUCUCCGAACGUGUCACGUTT | ACGUGACACGUUCGGGAAATT |
| siRNA-knj2-1  | CCCAUCACUAUCGUUCACGAAATT | UUCGUGAACGGAUGUGGGTT |
| siRNA-knj2-2  | GCUCUUGAAGAGAAACACUATT | UAGUGUUUCUCUUCAAGAGGCTT |

### Electrophysiology
Whole-cell current recordings were performed using a HEKA EPC10 amplifier controlled with PatchMaster software (HEKA) at room temperature with a voltage ramp of 500 ms duration from −120 mV to +60 mV applied every 5 s. ML133 and BaCl₂ were performed when the currents reached to the steady state. Specifically, the extracellular solution (ECS) contained 135 mM NaCl, 5 mM KCl, 2 mM CaCl₂, 1 mM MgCl₂ and 10 mM HEPES (adjusted to pH 7.4 with NaOH). For high potassium induced depolarization, the external solution was changed into high potassium ECS containing 50 mM potassium L-aspartate, 90 mM NaCl, 2 mM CaCl₂, 1 mM MgCl₂ and 10 mM HEPES (adjusted to pH 7.4 with NaOH). The pipette solution contained 147 mM potassium L-aspartate, 2 mM MgCl₂ and 10 mM HEPES (adjusted to pH 7.3 with KOH). The membrane potential was held at -40 mV. Glass pipettes with a resistance of 3–5 MΩ were used. Data were acquired at 10 kHz and filtered offline during data analysis. The series resistance (Ra) was less than 15 MΩ and monitored after experiments. Change of the extracellular solution was performed using an RSC-200 system (Bio-Logic Science Instruments).

For continuously monitoring changes of membrane potential (Vm), whole-cell configuration was adopted using current clamp (I = 0). we recorded a total of 3 min with or without ML133 and 5 s for High-potassium (50 mM). The membrane potential was the average Vm within 30 s of the initiation or the average Vm of the last 30 s of a total 3 minutes record.

Membrane potential detection by FACS

Mouse peritoneal macrophages were seeded into 12-well plate (6 × 10⁵/well). Cellular membrane potential sensitive probe, DiBAC₄(3), was mixed with indicated stimuli and the peritoneal macrophages were incubated for 1 h. Then the cells were washed with cold PBS for 3 times and FACS analysis were executed.

Chromatin immunoprecipitation (ChIP) and ChIP-qPCR

Cells were seeded at 1 × 10⁷/dish in 100-mm dishes in complete DMEM medium (supplemented with 10% fetal bovine serum (FBS), 1% penicillin/streptomycin) to adhere. Cells were treated with 500 ng/ml LPS in the presence or absence of compounds as indicated for 6 hours. Then experiments were performed according to the kit protocol (SimpleChIP® Plus Enzymatic Chromatin IP Kit, CST#9005) and the final DNA samples were used for qPCR. DNA amplification for 40 cycles at 95 °C for 30 s and 60 °C for 30 s. ChIP primers used in the study.
Glucose uptake assay

For in vitro glucose uptake assays, mouse peritoneal macrophages were seeded at 5 × 10^6/well in 12-well plates and treated as indicated. Cells were starved for 30 min at 37°C in Krebs-Ringer bicarbonate solution (KRBH, in mM: NaCl 135, KCl 3.6, NaH₂PO₄ 0.5, CaCl₂ 1.5, NaHCO₃ 2, HEPES 10 and 0.1% BSA) after washing 3 times with cold KRBH. Then the cells were incubated in KRBH supplemented with 2NBDG (60 µM) at 37°C for 15 min. After washing with cold KRBH for 3 times, the cells were scraped for FACS analysis. For in vivo glucose uptake assays, 8-week-old mice were intraperitoneally injected with LPS (1 mg/kg), 30 min later 2NBDG (500 nmol/mouse) was injected, and after another one hour, blood and peritoneal exudate cells were sequentially collected. Peripheral blood mononuclear cell (PBMC) were isolated from blood sample by Ficoll-Paque™ PLUS (GE Healthcare) according to the manufacturer’s instructions. Peritoneal exudate cell (PEC) were flushed with 5 ml cold PBS. Macrophages (CD45⁺ CD11b⁺ F4/80⁺ cells) in PECs and monocyte (CD45⁺ CD11b⁺ Ly6C<sub>high</sub> cells) in the blood were stained with corresponding antibody for 30 min on ice. After PBS washing, FACS analysis of 2NBDG MFI was executed by BD Fortessa Multicolor flow cytometer.

Calcium Flux Analysis

Intracellular calcium flux was detected by Fluo4-AM following the manufacturer’s instructions. In brief, Mouse peritoneal macrophages were seeded in 6-well plate (3 × 10^6/well). Fluo4-AM and Pluronic F127 were dissolved in HBSS (with Ca²⁺ and Mg²⁺), cells were incubated at 37°C for 2 h, then changed into HBSS (1% BSA) without Fluo4 for further 30 min. After that, cells were scraped and stained with CD45 and F4/80 in 4°C for 30 min for next FACS analysis. CD45⁺ F4/80⁺ macrophages were gated first and Fluo4 intensity of macrophages were recorded by BD Fortessa. Raw data was analyzed by GraphPad Prism.

Membrane protein biotinylation assay

Mouse peritoneal macrophages were seeded at 4 × 10^6/dish in 60-mm dishes and treated as indicated. Cells were washed three times with ice-cold PBS to remove any contaminating proteins. Then cell surface proteins were biotinylated by incubating cells with Sulfo-NHS-SS-Biotin solution on ice with a concentration of 1 mg/ml in PBS. 2 h later, cells were washed three times with ice-cold PBS to remove non-reacted biotinylation reagent. Alternatively, 25–50 mM Tris (pH 8.0) may be used for the initial wash.
to quench any non-reacted biotinylation reagent. After being lysed and centrifuged at 12000 rpm, 10 min, the supernatant was then transferred to a 1.5 ml microcentrifuge tube containing pre-washed streptavidin magnetic beads and incubate at 4°C overnights with rotation. Streptavidin magnetic beads were washed and lysed in SDS-PAGE Reducing Sample Buffer and the samples were used for western blot.

**Flow cytometry**

$6 \times 10^5$ mouse peritoneal macrophages were seeded and stimulated under the indicated conditions. Then cells were collected by cell scraper, followed by antibody staining for 30 min at 4°C. After washing with PBS solution to exclude nonspecific staining, the surface expression of CD98 or GLUT1 was detected by flow cytometry (ACEA NovoCyte).

**Statistical Analysis**

The results for q-PCR are expressed as the mean ± SD. The mouse sepsis model and the LC/MS experiments are expressed as the mean ± SEM and analyzed using two-tailed Student’s t-test for two groups. The q-PCR results are representative of at least three independent experiments. For mouse survival rate analysis, GraphPad Prism7 was used to plot Kaplan-Meier survival curves and to compare survival using log-rank tests.

**Declarations**

**CONTACT FOR REAGENT AND RESOURCE SHARING**

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Di Wang (diwang@zju.edu.cn).

**DATA AND CODE AVAILABILITY**

The RNA-seq data in this paper have been deposited in the Gene Expression Omnibus (GEO) with accession number GEO: GSE146158.

**AUTHOR CONTRIBUTIONS**

W.Y., Z.W., X.Y., Z.C., K.Z., S.C., T.X., D.J., X.G., M.L., J.Z., H.F., and Q.Y. performed the experiments; X.Y., X.Z., Y.W., W.Y., and D.W. designed the research; W.Y., Z.W., and D.W. wrote the manuscript; and D.W. supervised the project.

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