ORIGINAL RESEARCH COMMUNICATION

Nuclear Factor E2-Related Factor-2 Negatively Regulates NLRP3 Inflammasome Activity by Inhibiting Reactive Oxygen Species-Induced NLRP3 Priming

Xiuting Liu, Xin Zhang, Yang Ding, Wei Zhou, Lei Tao, Ping Lu, Yajing Wang, and Rong Hu

Abstract

Aims: The NLRP3 inflammasome is a multiprotein complex that protects hosts against a variety of pathogens. However, the molecular mechanisms of modulating NLRP3 inflammasome activation, especially at the priming step, are still poorly understood. This study was designed to elucidate the negative regulation of nuclear factor E2-related factor-2 (Nrf2) on the activation of NLRP3 inflammasome.

Results: We reported that Nrf2 activation inhibited NLRP3 expression, caspase-1 cleavage, and subsequent IL-1 β generation. Compared with normal cells, Nrf2-deficient cells showed upregulated cleaved caspase-1, which were attributed to the increased transcription of NLRP3 caused by excess reactive oxygen species (ROS). Furthermore, priming of the NLRP3 inflammasome was sensitive to the exogenous ROS levels induced by H₂O₂ or rotenone. Combined with adenosine triphosphate, rotenone triggered higher activity of the NLRP3 inflammasome compared with lipopolysaccharide, suggesting that ROS promoted the priming step. In addition, Nrf2-induced NQO1 was involved in the inhibition of the NLRP3 inflammasome. In an *in vivo* alum-induced peritonitis mouse model, Nrf2 activation suppressed typical IL-1 signaling-dependent inflammation, whereas Nrf2^{-/-} mice exhibited a significant increase in the recruitment of immune cell and the generation of IL-1 β compared with wild-type mice.

Innovation: We elucidated the effects and possible mechanisms of Nrf2 activation-induced NQO1 expression on NLRP3 inflammasome inactivation and established a novel regulatory role of the Nrf2 pathway in ROS-induced NLRP3 priming.

Conclusions: We demonstrated Nrf2 negatively regulating NLRP3 inflammasome activity by inhibiting the priming step and suggested that Nrf2 could be a potential target for some uncontrolled inflammasome activation-associated diseases. *Antioxid. Redox Signal.* 26, 28–43.

Keywords: Nrf2, NLRP3 inflammasome, ROS, priming

Introduction

THE NLRP3 INFLAMMASOME is a multiprotein complex composed of the NLRP3 scaffold, ASC adaptor, and procaspase-1 (38). Recent studies indicate that the NLRP3 inflammasome can be activated when stimulated by both endogenous “danger signals” and compounds associated with pathogens (39). The assembly of the inflammasome leads to the maturation and secretion of some proinflammatory cytokines, such as IL-1 β and IL-18 (24, 25). Moreover, the NLRP3 inflammasome has been regarded as a key component of innate immunity. However, excessive and chronic activation of the inflammasome results in many human diseases, such as fa-

miliar cold autoinflammatory syndrome (FCAS), cancer, type II diabetes (14), gout, atherosclerosis (22), and Alzheimer’s disease (5). Thus, activation of the NLRP3 inflammasome should be effectively controlled (13).

Many studies have shown that activation of the NLRP3 inflammasome requires two steps. At first, pattern recognition or cytokine receptor-induced priming signal leads to the synthesis of NLRP3 and pro-IL-1 β proteins. Then, a second signal promotes the NLRP3 inflammasome assembly-mediated cleaved caspase-1 and IL-1 β secretion (3). The protein level of NLRP3 is critical for inflammasome assembly and regarded as a rate-limiting element for the inflammasome activation (6, 46). In resting macrophages,

Innovation

Nuclear factor erythroid 2-related factor 2 (Nrf2) activation-induced NQO1 expression contributed to the inhibition of NLR family, pyrin domain containing 3 (NLRP3) inflammasome activation in macrophages. In addition, tert-butylhydroquinone negatively regulated NLRP3 transcription by enhancing the antioxidant capacity in an Nrf2-antioxidant response element (ARE)-dependent manner. As a reactive oxygen species scavenger, Nrf2-mediated NQO1 expression blocked NLRP3 priming, followed by attenuated activation of NLRP3. These results established a link between the Nrf2-ARE pathway and the NLRP3 inflammasome. These results also suggested that pharmacological activation of Nrf2 could be a potential therapeutic strategy for NLRP3-associated diseases.

NLRP3 expression is relatively low and the assembly of the inflammasome complex is poorly induced. Many exogenous and endogenous factors can promote NLRP3 expression, including TLR agonists, proinflammatory cytokines, and reactive oxygen species (ROS) (37). It is interesting to note that almost all NLRP3 agonists can trigger ROS generation and that oxidative stress plays a vital role in NLRP3 inflammasome activation.

Redox signaling is a critical mechanism that mediates NLRP3 inflammasome activation. Changes in the redox microenvironment regulate the NLRP3 inflammasome activation resulting in cleaved caspase-1 and mature IL-1 β secretion (35). In addition, *in vitro* suppressing of ROS production by some scavengers or knockdown of NADPH oxidase subunits revealed that when ROS production was absent, NLRP3-induced IL-1 β generation was impaired, which indicated that ROS played crucial roles in inflammasome activation (10).

Notably, as a redox-sensitive transcription factor, Nrf2 regulates cellular antioxidant responses through modulating many phase II antioxidants, detoxifying enzymes, and some stress-responsive proteins (12, 20, 42). Under basal conditions, Kelch ECH associating protein 1 (Keap1), a repressor protein, tethers Nrf2 to the actin cytoskeleton within the cytosol. The interaction between Nrf2 and Keap1 leads to continual ubiquitination and degradation of Nrf2. Under cellular stress conditions, Nrf2 is accumulated and translocated to the nucleus, binding to the antioxidant response element (ARE) to initiate the antioxidant response, which protects against a variety of reactive toxicants, including mitochondrial and NADPH oxidase-induced ROS (28, 23, 45). The protective roles of Nrf2 have been reported in asthma (34), sepsis (44), and cigarette smoke-induced emphysema (43). Furthermore, sulforaphane has been demonstrated to ameliorate DSS-induced colitis (48), to which Nrf2^{-/-} mice are more susceptible (19). Although the effects of Nrf2 on the regulation of many inflammatory diseases are emerging, the mechanisms of which remain largely unknown.

In this study, we show that Nrf2 activation induced NQO1 expression to scavenge ROS, which then inhibited NLRP3 expression and subsequent caspase-1 cleavage. We identified Nrf2 as an endogenous regulator of the NLRP3 inflammasome and illustrated the fine regulation of inflammasome activation under physiological conditions.

Results

Nrf2 inhibited NLRP3 inflammasome activation

To investigate the possible roles of Nrf2 in NLRP3 inflammasome activation, lipopolysaccharide (LPS)-stimulated THP-Ms were pretreated with tert-butylhydroquinone (tBHQ) before challenging them with adenosine triphosphate (ATP). As shown in Figure 1A and Supplementary Figure S1A and B (Supplementary Data are available online at www.liebertpub.com/ars), tBHQ dose dependently increased the protein level and nuclear translocation of Nrf2. NLRP3-mediated caspase-1 activation (Fig. 1B and Supplementary Fig. S1C) and IL-1 β maturation (Fig. 1C) were inhibited by tBHQ in a dose-dependent manner. Consistent with this, tBHQ treatment significantly attenuated NLRP3 expressions at both the mRNA and protein levels (Fig. 1B–E).

To test whether the inhibitory effect of tBHQ on the activation of NLRP3 inflammasome was mediated by Nrf2, siRNA targeting Nrf2 was used to suppress endogenous Nrf2 expression (Fig. 1F). Knockdown of Nrf2 in THP-Ms significantly increased LPS+ATP-induced NLRP3 inflammasome activation (Fig. 1H, J, and Supplementary Fig. S1E). On the contrary, the NLRP3 inflammasome was poorly activated when Nrf2 was over-expressed (Fig. 1I and Supplementary Fig. S1F). Combined with tBHQ stimulation, transfection in THP-Ms with an Nrf2 over-expression plasmid resulted in a sharp decrease of IL-1 β secretion (Fig. 1I, K and Supplementary Fig. S1F), while transfection with siRNA or plasmid alone had no effect on inflammasome activation (Fig. 1G and Supplementary Fig. S1F). Together, these observations suggested that tBHQ-induced Nrf2 activation negatively regulated the NLRP3 inflammasome activation.

Nrf2 inhibited the priming step of NLRP3 inflammasome activation

A sufficient level of NLRP3 protein is necessary to form and activate the NLRP3 inflammasome. Given that Nrf2 inhibited NLRP3 activation, we tested whether Nrf2 suppressed NLRP3 expression. THP-Ms were pretreated with tBHQ, which was then followed by LPS stimulation. tBHQ treatment remarkably attenuated LPS-induced NLRP3 expression at both the protein (Fig. 2A and Supplementary Fig. S2A) and mRNA levels (Fig. 2B, C). In addition, stimulation of THP-Ms with tBHQ alone decreased NLRP3 expression (Fig. 2A). As shown in Figure 2D and G, Nrf2 knockdown greatly enhanced NLRP3 expression at both the mRNA and protein levels, while transfection with the Nrf2 plasmid had the opposite effect (Fig. 2E, H). Taken together, these results indicated that Nrf2 activation inhibited LPS-induced NLRP3 production. It was previously reported that NF- κ B was inhibited by Nrf2 activation in certain inflammatory diseases (47). However, we were unable to observe any suppressive effects of tBHQ on NF- κ B nuclear translocation (Fig. 2F and Supplementary Fig. S2C). Traditional inflammasome studies adopt sequential stimulation methods, such as stimulating TLR ligands followed by a second stimulus several hours later to trigger NLRP3 inflammasome assembly. However, recent studies have indicated that co-treatment of LPS and ATP can induce the rapid activation of the NLRP3 inflammasome independent of the NLRP3 protein level (29). In this case, NLRP3 synthesis is not necessary for rapid caspase-1 activation. As shown in Figure 2I and Supplementary Figure S2D, the stimulation of THP-Ms with

FIG. 1. tBHQ inhibits NLRP3 inflammasome activation. THP-Ms were treated with tBHQ for 12 h, and the protein level of Nrf2 and its nuclear translocation were determined by Western blot (A). tBHQ-pretreated THP-Ms were stimulated with LPS for 4 h and ATP for 1 h. The panel shows a Western blot of NLRP3, cleaved caspase-1 (p10), and cleaved IL-1 β (p17) in the supernatants or cell lysates (B). IL-1 β was measured by ELISA (C). The mRNA levels of NLRP3 and pro-IL-1 β were detected by real-time RT-PCR (D, E). Western blot analysis of Nrf2 expression in THP-Ms transfected with different concentrations of Nrf2 siRNA or Nrf2 plasmid for 12 h (F). NLRP3, cleaved caspase-1, and IL-1 β were detected by Western blot (G). Nrf2 siRNA-transfected THP-Ms were stimulated with LPS and ATP for the indicated periods. NLRP3, cleaved caspase-1, and IL-1 β were detected by Western blot (H). IL-1 β was measured by ELISA (J). Nrf2 plasmid-transfected THP-Ms were stimulated with LPS and ATP for the indicated periods. NLRP3, cleaved caspase-1, and IL-1 β were detected by Western blot (I). IL-1 β was detected by ELISA (K). The results are representative of three independent experiments and are expressed as the mean $\pm$ SD. * p < 0.05, ** p < 0.01 compared with the control, # p < 0.05, ## p < 0.01 compared with LPS + ATP. ATP, adenosine triphosphate; LPS, lipopolysaccharide; NLRP3, NLR family, pyrin domain containing 3; Nrf2, nuclear factor erythroid 2-related factor 2; tBHQ, tert-butylhydroquinone; THP-Ms, THP-1 cell-derived macrophages.

FIG. 2. Nrf2 inhibited the priming step of NLRP3 inflammasome activation. THP-Ms were pretreated with tBHQ for 12 h and then stimulated with or without LPS for the indicated periods. NLRP3 was detected by Western blot (A). The mRNA levels of NLRP3 and pro-IL-1 β were measured by real-time RT-PCR (B, C). THP-Ms were transfected with Nrf2 siRNA or Nrf2 plasmid and then stimulated with LPS or ATP. The protein levels of NLRP3 were detected by Western blot (D, E). The mRNA levels of NLRP3 and pro-IL-1 β were measured by real-time RT-PCR (G, H). tBHQ-pretreated THP-Ms were stimulated with LPS for 4 h, and the protein level of NF- κ B and its nuclear translocation were determined by Western blot (F). tBHQ-pretreated THP-Ms were stimulated with LPS together with ATP for 1 h. Cleaved caspase-1 (p10), IL-1 β , and NLRP3 were detected by Western blot (I). tBHQ-stimulated THP-Ms were pretreated with actidione for 1 h and then stimulated with LPS and ATP. Cleaved caspase-1 (p10), IL-1 β , and NLRP3 were detected by Western blot (J). The results are representative of three independent experiments and are expressed as the mean $\pm$ SD. * p < 0.05, compared with the control, # p < 0.05, ## p < 0.01 compared with LPS + ATP.

LPS and ATP for 1 h led to rapid NLRP3 inflammasome activation, while tBHQ treatment had no significant effect on NLRP3 inflammasome activation. Furthermore, when actidione was given before LPS stimulation to inhibit NLRP3 protein synthesis, the inhibitory effect of tBHQ on inflammasome activation was abolished (Fig. 2J and Supplementary Fig. S2E), indicating that Nrf2 inhibited NLRP3 inflammasome activation by suppressing NLRP3 transcription.

tBHQ decreased the production of ROS induced by LPS and ATP

Nrf2 plays a vital role in protection against oxidative stress. Previous reports identified that ROS were critically involved in NLRP3 inflammasome activation (35). Therefore, we mea-

sured the production of ROS in THP-Ms under different stimulation conditions. tBHQ decreased total ROS in a time- and dose-dependent manner (Fig. 3A, B). ROS generation induced by LPS and ATP was also inhibited by tBHQ stimulation (Fig. 3C, D). Furthermore, Nrf2 siRNA caused higher ROS release (Fig. 3E), while a lower ROS level was observed in the context of Nrf2 overexpression (Fig. 3F). Similar results were also found in bone marrow-derived macrophages (BMDMs), where Nrf2^{-/-} BMDMs exhibited higher levels of ROS that could not be downregulated by tBHQ (Fig. 3G-I).

Superoxide is a component of mitochondrial ROS. Using flow cytometry, we examined the effect of tBHQ on the intracellular superoxide level. It was found that most of the intracellular superoxide elevation caused by LPS and ATP was decreased when Nrf2 was activated in advance (Fig. 4A).

FIG. 3. tBHQ decreased the production of ROS induced by LPS and ATP. THP-Ms were stimulated by tBHQ with different doses for the indicated time, and the intracellular level of total ROS was measured by flow cytometry (A, B). Cells pretreated with different concentrations of tBHQ were stimulated by LPS and ATP; then, ROS were measured (C, D). Nrf2 siRNA and Nrf2 plasmid-transfected THP-Ms were stimulated by LPS and ATP at the indicated time and dose; then, the intracellular level of total ROS was measured by flow cytometry (E, F). WT BMDMs and Nrf2^{-/-} BMDMs were stimulated by tBHQ, and the intracellular level of total ROS was measured by flow cytometry (G–I). The results are representative of three independent experiments and are expressed as the mean $\pm$ SD. * p < 0.05, ** p < 0.01 compared with the control, # p < 0.05, ## p < 0.01 compared with LPS + ATP. N.S. means no significant difference. BMDM, bone marrow-derived macrophage; ROS, reactive oxygen species. To see this illustration in color, the reader is referred to the web version of this article at www.liebertpub.com/ars

Moreover, superoxide was markedly enhanced in Nrf2-deficient cells and inhibited by Nrf2 overexpression (Fig. 4B). The activity of superoxide dismutase (SOD) was also measured after different stimulation, which was found to be closely related to Nrf2 activation (Fig. 4C, D). Similar phenomena were observed in the levels of peroxide (Fig. 4E, F) and catalase (Fig. 4G, H). Taken together, these results strongly suggested that Nrf2 activation enhanced the activities of SOD and catalase to scavenge different sources of ROS.

ROS promoted NLRP3 activation by affecting the priming step

The above results suggested that tBHQ suppressed ROS production and NLRP3 inflammasome activation. To investigate the role of ROS in the activation of the NLRP3 inflammasome, H₂O₂ and rotenone were used to upregulate total intracellular ROS (Fig. 5A). H₂O₂ represented an exogenous way to promote ROS production, while rotenone, a respiratory chain inhibitor, increased mitochondrial ROS and enhanced total ROS. We found that ROS can both induce NLRP3 transcription and promote LPS-induced NLRP3 expression at the mRNA and protein levels (Fig. 5B, C). When NAC or rotenone

was given to LPS-treated THP-Ms for 3 h followed by ATP stimulation, the production of both IL-1 β (Fig. 5F) and cleaved caspase-1 (Fig. 5D) was decreased by NAC, whereas the activation of the NLRP3 inflammasome was increased by rotenone.

Based on the finding that ROS promoted the transcription of NLRP3 (Fig. 5B, C), we explored whether ROS primarily regulated the priming step. Rotenone was used as a substitute for either LPS or ATP to stimulate THP-Ms. As expected, both LPS and rotenone were able to prime the NLRP3 inflammasome, and when combined with ATP, they all induced cleaved caspase-1 activation. Hence, we speculated that ROS may play the same role as LPS in the activation of the NLRP3 inflammasome (Fig. 4E, G). Indeed, IL-1 β secretion was not increased in response to LPS plus rotenone. That is, rotenone had little effect on the assembly of the NLRP3 inflammasome. Taken together, these results demonstrated that ROS promoted the priming step of NLRP3 inflammasome activation.

The negative regulation of Nrf2 on the NLRP3 inflammasome was dependent on ROS

Next, we study the underlying mechanism of the inhibitory effects of Nrf2 on NLRP3 inflammasome activation. Nrf2

FIG. 4. tBHQ enhanced the activities of SOD and catalase to scavenge superoxide and peroxide. Cells pretreated with different concentrations of tBHQ were stimulated by LPS and ATP, and the intracellular levels of superoxide and peroxide were measured (A, E). Nrf2 siRNA or Nrf2 plasmid-transfected THP-Ms were stimulated by LPS and ATP at the indicated time and dose, and then, superoxide and peroxide were measured (B, F). THP-Ms were subjected to the above stimulation, and the activities of SOD (C, D) and catalase (G, H) were detected. The results are representative of three independent experiments and are expressed as the mean $\pm$ SD. * $p < 0.05$, compared with the control, # $p < 0.05$, ## $p < 0.01$ compared with LPS + ATP. SOD, superoxide dismutase.

signaling suppressed ROS generation and caspase-1 activation. Hence, we tested the role of ROS in Nrf2-induced NLRP3 inflammasome inhibition, and it was found that the inhibitory effect of tBHQ on NLRP3 inflammasome activation was rescued by rotenone (Fig. 6A–C). Combined with ATP, H₂O₂ or rotenone (instead of LPS) could trigger NLRP3 inflammasome activation, which was inhibited by tBHQ pretreatment (Fig. 6D–I).

Moreover, transfection with Nrf2 siRNA increased intracellular ROS levels and promoted the transcription of NLRP3, cleaved caspase-1 production and IL-1 β secretion. This enhancement was significantly suppressed by NAC (Fig. 6J–L), suggesting that Nrf2-induced NLRP3 inflammasome inhibition was dependent on ROS. When transfected with the Nrf2

plasmid, neither rotenone nor NAC had any significant effect on NLRP3 inflammasome activation (Fig. 6M–O). We speculated that when Nrf2 was overexpressed, exogenous ROS were scavenged before activating the NLRP3 inflammasome. These data suggested that Nrf2 inhibited the activation of NLRP3 inflammasome by scavenging ROS.

The negative regulation of Nrf2 on NLRP3 inflammasome activation involved NQO1

As shown in Figure 1H, Nrf2 knockdown significantly increased LPS- and ATP-induced NLRP3 inflammasome activation. To further confirm this, the effect of tBHQ was tested in Nrf2^{-/-} BMDMs. The results showed that

FIG. 5. ROS promoted NLRP3 activation by affecting the priming step. LPS-stimulated THP-Ms were exposed to NAC, H₂O₂, and rotenone and then treated with ATP. The intracellular level of ROS was measured by flow cytometry (A). The mRNA levels of NLRP3 and pro-IL-1β were measured by real-time RT-PCR (B, C). NLRP3, cleaved caspase-1, and IL-1β were detected by Western blot (D). IL-1β was detected by ELISA (F). Rotenone was substituted with LPS or ATP, and then, cleaved caspase-1, IL-1β, and NLRP3 were detected by Western blot (E). IL-1β was detected by ELISA (G). The results are representative of three independent experiments and are expressed as the mean ± SD. **p* < 0.05, ***p* < 0.01 compared with the control, #*p* < 0.05, compared with LPS + ATP. To see this illustration in color, the reader is referred to the web version of this article at www.liebertpub.com/ars

FIG. 6. The negative regulation of Nrf2 on the NLRP3 inflammasome was dependent on ROS. THP-Ms were pretreated with tBHQ for 12 h and then stimulated with LPS for 4 h and ATP for 1 h. NAC or rotenone was added for the indicated time before ATP treatment. NLRP3, cleaved caspase-1, and IL-1 β were detected by Western blot (A). IL-1 β was detected by ELISA (B). The mRNA level of NLRP3 was detected by real-time RT-PCR (C). Western blot analysis of NLRP3, cleaved caspase-1, and IL-1 β in tBHQ-pretreated cells stimulated with rotenone (D) or H₂O₂ (G) plus ATP. IL-1 β was detected by ELISA (E, H). The mRNA level of NLRP3 was measured by real-time RT-PCR (F, I). Nrf2 siRNA-transfected THP-Ms were treated with the same stimulation. NLRP3, cleaved caspase-1, and IL-1 β were detected by Western blot (J) IL-1 β was detected by ELISA (K). The mRNA level of NLRP3 was measured by real-time RT-PCR (L). Nrf2 plasmid-transfected cells were shown (M–O). The results are representative of three independent experiments and are expressed as the mean $\pm$ SD. * p < 0.05, ** p < 0.01.

the negative regulation of tBHQ on LPS+ATP-induced caspase-1 activation in BMDMs was abolished almost completely when Nrf2 was absent (Fig. 7A, B and Supplementary Fig. S3F, G). On the contrary, the Nrf2 overexpression plasmid significantly rescued the inhibitory function of tBHQ in Nrf2^{-/-} BMDMs (Fig. 7C, D, and Supplementary Fig. S3H). We found that treating BMDMs with different doses of tBHQ for 12 h increased the transcription of ARE genes, including GCLC, GCLM, GST, HO-1, and NQO1 (Fig. 7E). Notably, the transcription of NQO1 induced by tBHQ was significantly increased by 20-fold. By contrast, in Nrf2^{-/-} BMDMs, the NQO1 mRNA level was not affected by tBHQ, while the mRNA levels of other ARE genes were increased by approximately fourfold when exposed to 50 μ M tBHQ (Fig. 7F). In addition, in THP-Ms, the effect of tBHQ on the expression of NQO1, but not other ARE genes, was more profound than in Nrf2-silenced cells (Fig. 7G). A previous study reported that glutathione (GSH) may be the dominant factor in the cytoprotective actions of Nrf2 (11). However, we did not detect any remarkable effects of tBHQ on GSH in either THP-Ms or BMDMs (Supplementary Fig. S3I), suggesting that GSH may not be involved in the regulation of the NLRP3 inflammasome. Taken together, these data indicated that NQO1 was involved in the inhibitory effects of Nrf2 on NLRP3 inflammasome activation.

tBHQ attenuated alum-induced peritonitis in vivo

To further test whether the *in vitro* biological effects of Nrf2 activation on NLRP3 inflammasome were also observable *in vivo*, an IL-1-dependent mouse peritonitis model was adopted. Mice received a gavage of tBHQ (300 mg/kg and 600 mg/kg) for 1 week followed by an intraperitoneal (i.p.) injection of alum. After that, we collected peritoneal exudate cells (PECs), and inflammatory cells recruited by alum were analyzed by flow cytometry. As shown in Figure 8A–D, alum-induced recruitment of Ly6G⁺ neutrophils and Ly6C⁺ monocytes was significantly blocked by different doses of tBHQ. In addition, less caspase-1 cleavage and mature IL-1 β secretion were observed in the lavage fluid from mice that received a gavage of tBHQ (Fig. 8E, F). Notably, we observed that alum-recruited PECs had a fusiform shape, rather than a circular shape, as observed in normal and tBHQ pretreatment groups (Fig. 8G).

However, Nrf2^{-/-} mice showed a remarkable increased recruitment of inflammatory cells, including Ly-6G⁺ neutrophils and Ly-6C⁺ inflammatory monocytes, after intraperitoneal (i.p.) injection of alum compared with wild-type mice (Fig. 9A–D), supporting a critical anti-inflammatory role of Nrf2 in alum-induced peritonitis. In addition, higher NLRP3 and cleaved caspase-1 expression, as well as IL-1 β secretion, were also observed in the lavage fluid from Nrf2^{-/-}

FIG. 7. The negative regulation of Nrf2 on NLRP3 inflammasome activation involved NQO1. tBHQ-pretreated WT BMDMs, Nrf2^{-/-} BMDMs, and Nrf2 plasmid-transfected Nrf2^{-/-} BMDMs were stimulated with LPS for 4 h and ATP for 1 h, and then, NLRP3, cleaved caspase-1, and IL-1β were detected by Western blot (A–C). IL-1β was detected by ELISA (D). WT BMDMs and Nrf2^{-/-} BMDMs were treated with tBHQ for 12 h, and the mRNA levels of GCLM, GCLC, GST, HO-1, and NQO1 were measured by real-time RT-PCR (E, F). The expressions of the above mRNA were detected in THP-Ms treated with different stimulations (G). The results are representative of three independent experiments and are expressed as the mean ± SD. **p* < 0.05, ***p* < 0.01, #*p* < 0.05, ##*p* < 0.01 compared with LPS + ATP, N.S. means no significant difference.

mice (Fig. 9E, F). Nrf2 deficiency may abolish its protective effect against alum-induced peritonitis, as immune cells had a more mature phenotype and a fusiform shape (Fig. 9G). Overall, these findings further confirmed that Nrf2 activation inhibited NLRP3 inflammasome activity *in vivo*.

Discussion

Recently, Nrf2 and the NLRP3 inflammasome have been identified as regulators of many inflammatory diseases. They have also become useful targets of anti-inflammatory interventions for the treatment of liver injury (32), lupus nephritis (18), and colitis (49). More recently, Pei-Yi Tsai demon-

strated that epigallocatechin-3-gallate had prophylactic effects on lupus nephritis by increasing the Nrf2 antioxidant signaling pathway and inhibiting renal NLRP3 inflammasome activation (11), which could also be applied to the treatment of glomerulonephritis (50). In addition, schisandrin B was reported to induce Nrf2 expression and attenuate ROS-dependent peritonitis (26). Although studies have revealed that enhancing Nrf2 and suppressing NLRP3 can be used to treat inflammation, it is still unknown whether there is a crosslink between the Nrf2-ARE pathway and NLRP3 inflammasome and whether Nrf2 inducers could be adopted to protect against NLRP3-associated diseases. Further in-depth biochemical analysis about the interaction between Nrf2 and

FIG. 8. tBHQ attenuated alum-induced peritonitis *in vivo*. Different groups of mice ($n=8$) were intraperitoneally injected with alum ($700 \mu\text{g}$ /per mouse) diluted in sterile PBS. Peritoneal lavage was performed for 12 h after injection. The ratio of CD11b⁺Ly-6C⁺ inflammatory monocytes (**A**, **C**) and CD11b⁺Ly-6G⁺ neutrophils (**B**, **D**) was evaluated by FACS analysis. PECs were seeded in 12-well plates and HE stained (**G**). The protein levels of NLRP3 and cleaved caspase-1 were detected by Western blot (**E**). IL-1 β was detected by ELISA (**F**). The results are representative of at least three independent experiments and are expressed as the mean $\pm$ SD. ** $p < 0.01$, *** $p < 0.001$ compared with the control, # $p < 0.05$, ## $p < 0.001$ compared with alum. PEC, peritoneal exudate cell. To see this illustration in color, the reader is referred to the web version of this article at www.liebertpub.com/ars

the NLRP3 inflammasome should provide mechanistic insights. In this study, we found that Nrf2 activation inhibited NLRP3 inflammasome activation by decreasing ROS-induced NLRP3 transcription.

Our *in vivo* study showed that Nrf2^{-/-} mice are more susceptible to peritonitis than wild-type mice, which does not agree with a recent study by Zhao *et al.* indicating that Nrf2^{-/-} mice have significantly less recruitment of immune cells and are less susceptible to peritonitis (51). Alum-induced peritonitis is regarded as a typical NLRP3-IL-1 β axis-dependent inflammation animal model, and notably, the dose of alum and its stimulation time influence immune cell recruitment in the peritoneal cavity. The omentum is a major site of entry for leukocytes during peritoneal inflammation (8), and a sufficient dose of alum is required to instigate leukocyte recruitment. Generally, a dose of $700 \mu\text{g}$ per mouse

has been adopted in most experiments (9, 16). In addition, monocyte-derived macrophages are known to migrate from the peritoneum in the resolution phase of acute inflammation (4), and the longer time course would increase the migration of immune cells. In our study, we used an alum dose of $700 \mu\text{g}$ per mouse for 12 h to induce acute peritonitis, which is different from Zhao's methodology ($400 \mu\text{g}$ of alum per mouse for 16 h). We speculate that the dose of alum used and the duration of the exposure may explain the contradiction between our study and Zhao's.

We hypothesized that Nrf2 preactivation leads to the inhibition of NLRP3 inflammasome activation *via* a ROS-dependent mechanism. Indeed, tBHQ decreased the production of superoxide and peroxide, while the activities of SOD and catalase were significantly enhanced. Larry *et al.* demonstrated that Nrf2 played a critical role in controlling

FIG. 9. Enhanced alum-induced peritonitis in Nrf2^{-/-} mice. WT or Nrf2^{-/-} mice ($n = 8$) were intraperitoneally injected with alum (700 μg /per mouse) diluted in sterile PBS. Peritoneal lavage was performed 12 h after injection. The ratio of CD11b⁺Ly-6C⁺ inflammatory monocytes (A, B) and CD11b⁺Ly-6G⁺ neutrophils (C, D) was evaluated by FACS analysis. Cytospin preparation followed by HE staining was performed (G). The protein levels of NLRP3 and cleaved caspase-1 were detected by Western blot (E). IL-1 β was measured by ELISA (F). The results are representative of at least three independent experiments and are expressed as the mean $\pm$ SD. * $p < 0.05$, ** $p < 0.01$. To see this illustration in color, the reader is referred to the web version of this article at www.liebertpub.com/ars

basal GSH levels by modulating the rate of GSH synthesis in mouse embryonic fibroblasts. Moreover, GSH was entirely responsible for the protection against a soft and hard electrophile, an organic hydroperoxide, and a redox-cycling agent, indicating that increasing the abundance of GSH is more important than the induction of enzymes, such as heme oxygenase-1 and NAD(P)H:quinone oxidoreductase-1, in scavenging ROS (11). In the present study, we measured the mRNA levels of Nrf2-targeted ARE genes, including GCLC, GCLM, GST, HO-1, and NQO1, and found that the effect of

tBHQ on the mRNA level of NQO1, but not other ARE genes, was more profound than that in Nrf2-deficient cells. Notably, the expression of NQO1 was closely related to Nrf2-mediated inhibition of the NLRP3 inflammasome, which indicated that Nrf2-induced NQO1 expression played a critical role in scavenging ROS. In addition, NQO1 was identified as a highly inducible cytoprotective gene that regulated ROS generation (21). tBHQ could ameliorate arsenic-induced cytotoxicity and apoptosis by inducing Nrf2-dependent antioxidant responses in which NQO1 was

FIG. 10. Proposed model illustrating how Nrf2 negatively regulated NLRP3 inflammasome activation. ROS induced by LPS promoted NLRP3 transcription. Then, ATP stimulation activated the NLRP3 inflammasome, which led to mature IL-1 β secretion. tBHQ induced the nuclear translocation of Nrf2, NQO1 expression, and scavenged intracellular ROS, which resulted in the decrease of NLRP3 transcription and subsequent inflammasome activation. To see this illustration in color, the reader is referred to the web version of this article at www.liebertpub.com/ars

involved (7). In addition, the NQO1 inhibitor diminutol (Dim) abolished the suppressive function of Nrf2 on hepatotoxicity (33). Therefore, NQO1 may account for the inhibitory effects of Nrf2 on the regulation of NLRP3 inflammasome activation.

Oxidative stress is considered as a common feature of inflammatory responses, and ROS act as an upstream signal mediating the NLRP3 inflammasome activation (36, 54). In this study, our results showed that ROS exerted their effects by promoting the transcription of NLRP3. NF- κ B was reported to be an ROS-sensitive transcription factor that mediated NLRP3 inflammasome activation (27), but we did not find any change in the NF- κ B protein level or its nuclear translocation after exposure to tBHQ. Bauernfeind *et al.* (2) demonstrated that the priming step of NLRP3 inflammasome strictly requires ROS production, which is not necessary for inflammasome activation (1). Recently, a novel mechanism for ROS regulation of the post-translational activation of NLRP3 by deubiquitination has been put forward (17). In addition, ROS induced oxidized mitochondrial DNA release into the cytosol (40), where it binds to and activates the NLRP3 inflammasome (41). It was also reported that ROS-mediated TXNIP detachment from thioredoxin triggered IL-1 β expression (53). However, Masters *et al.* (30) identified no difference in NLRP3 inflammasome activation in response to inflammasome activators between WT and TXNIP^{-/-} macrophages, thereby excluding a role of TXNIP in modulating the NLRP3 inflammasome. Further studies need to be carried out to clarify the precise mechanism of ROS-mediated NLRP3 transcription.

However, there are studies indicating a negative regulatory role of ROS on NLRP3 inflammasome activation. It was found that excessive ROS could inhibit NLRP3-mediated caspase-1 activation and that NLRP3 inflammasome activation and IL-1 β secretion were suppressed by HO-1 inhibitor zinc (II) protoporphyrin IX (ZnPP) and the knockdown of Nrf2 and with the consumption of GSH-induced superoxide anion production (15). Studies also showed that in SOD1-deficient mice, persistent high ROS levels in macrophages were able to inhibit the activity of caspase-1 and the maturation of IL-1 β (31). ROS overproduction specifically inhibited caspase-1 by reversible oxidation and glutathionylation of the redox-sensitive cysteine residues Cys397 and Cys362. A

similar phenomenon may occur in Nrf2^{-/-} macrophages under chronic oxidative stress conditions, as Zhao suggested a positive role for Nrf2 in NLRP3 inflammasome activation (52). Further experiments are worth to dissect the precise mechanisms.

In conclusion, a model was proposed to illustrate how Nrf2 negatively regulated NLRP3 expression and subsequent caspase-1 activation (Fig. 10). NLRP3 is the rate-limiting element for the activation of the inflammasome. It has been demonstrated that the activation status of the NLRP3 inflammasome, which resulted from high level of NLRP3 expression, is closely related to many diseases, including metabolic diseases, autoinflammatory syndromes, and cancers. Hence, it is critical to tightly control NLRP3 expression to prevent and treat these diseases. Our work identified Nrf2 as an inhibitor of NLRP3 expression at the transcriptional level and provided a molecular pathway through an endogenous regulator affecting the immune response and even controlling the development of immune-mediated diseases. Furthermore, our results provided a strategy to downregulate the activity of inflammasome and suggested that Nrf2 could be a potential target for the therapeutic modulation of NLRP3-associated diseases, such as autoinflammatory diseases, gout, and diabetes.

Materials and Methods

Reagents

tBHQ purchased from Sigma-Aldrich was dissolved in DMSO at 20 mM. Primary antibodies for Nrf2, β -actin, lamin A, IL-1 β , horseradish peroxidase (HRP)-conjugated secondary antibodies were purchased from Bioworld Technology, Inc. NLRP3, NQO1, and caspase-1 antibody were purchased from CST Technology, Inc. Antibodies for flow cytometry, including CD11b, Ly-6C, and Ly-6G, were purchased from eBioscience (eBioscience, Inc.). Nrf2 siRNA and control siRNA were purchased from Santa Cruz Biotechnology, Inc. Phorbol myristate acetate (PMA), lipopolysaccharide (LPS), adenosine triphosphate (ATP), and rotenone were purchased from Sigma-Aldrich. N-acetyl-cysteine (NAC) and ROS Assay kit were purchased from Beyotime Institute of Biotechnology. ELISA kits for

human and murine IL-1 β were purchased from Boster Biotech Co. Ltd.

Cell culture

Human THP-1 cells were obtained from the Cell Bank of Shanghai, Institute of Biochemistry and Cell Biology, Chinese Academy of Sciences and maintained in RPMI 1640 medium (Gibco) supplemented with 10% heat-inactivated fetal bovine serum (Gibco) and 100 U/ml benzylpenicillin and 100 mg/ml streptomycin. THP-1 cells were stimulated by PMA (100 ng/ml) for 12 h to differentiate into macrophages. The peritoneal macrophages were isolated according to the following procedures. They were isolated from C57/BL6 mice and cultured with RPMI 1640 medium supplemented with 10% heat-inactivated fetal bovine serum. After 8 h, culture fluid was exchanged with a fresh culture medium. The adherent macrophages were used for experiments. L929 cells were also purchased from the Institute of Biochemistry and Cell Biology and cultured with DMEM (Gibco) supplemented with 10% heat-inactivated fetal bovine serum. BMDM cells were isolated according to the following procedures. Bone marrow cells were isolated from C57/BL6 mice and cultured with DMEM supplemented with 10% heat-inactivated fetal bovine serum and 30% L929 cell-conditioned medium. Culture fluid was exchanged with a fresh culture medium every 3 days. Under these conditions, adherent macrophages were obtained within 7 days. Non-adherent cells were removed by washing the plate twice with PBS. The adherent macrophages were used for experiments. All cell lines were cultured under a humidified 5% (v/v) CO₂ atmosphere at 37°C.

Mice

Six- to eight-week-old female C57BL/6 mice were purchased from Model Animal Genetics Research Center of Nanjing University (Nanjing, China). Six- to eight-week-old Nrf2^{-/-} mice were on C57BL/6 background. Animal welfare and experimental procedures were carried out strictly in accordance with the Guide for the Care and Use of Laboratory Animals (National Institutes of Health, the United States) and the related ethical regulations of our university. All efforts were made to minimize animals' suffering and to reduce the number of animals used.

Western blot assay

Total proteins were extracted by adding RIPA lysis buffer with 1 mM PMSF for 1 h on the ice and centrifuging at 13,000 rpm for 30 min at 4°C. Protein concentration in the supernatants was measured by BCA protein assay (Thermo). Cell culture supernatants were concentrated with methanol/chloroform and the pellets were lysed in the SDS sample buffer. Then, equal amount of sample was run on 12% SDS-PAGE. The proteins were transferred to polyvinylidene difluoride (PVDF) membranes (Millipore) using a semidry transfer system (Bio-rad). Proteins were detected using specific antibodies of Nrf2, NQO1, NLRP3, caspase-1, β -actin, and lamin A overnight at 4°C. The membranes were then washed three times with PBST followed by HRP-conjugated secondary antibodies for 1 h at 37°C. All of the antibodies were diluted in PBST containing 1% BSA. The signals were

analyzed using the ECL chemiluminescence detection system (Tanon).

Nuclear and cytoplasmic extraction

Following treatments, cells were harvested by centrifugation and washed twice with PBS. Nuclear and cytosol lysates were isolated using a Nuclear/Cytosol Fractionation Kit (BioVision) according to the manufacturer's instruction. The protein concentration of the nuclear and cytoplasmic extracts was measured with the BCA protein assay reagent. Extracts were stored at -20°C until further experimentation.

Measurement of cytokine production

Peritoneal macrophages and THP-1 cells were seeded in 12-well plates and cultured overnight. After different stimulation, the supernatants were collected, and the concentrations of IL-1 β were measured using mouse IL-1 β and human IL-1 β ELISA kits (Boster) according to the manufacturer's instruction.

RNA extraction and real-time RT-PCR

Total RNA was isolated using the TRIzol reagent (Invitrogen) according to the manufacturer's protocol. The concentration and purity of the extracted RNA were measured with the optical densities at 260 and 280 nm. RNA samples were reverse transcribed to cDNA and subjected to quantitative PCR, which was performed with the Light-Cycler[®] 96 Real-Time PCR System (Roche) using AceQ qPCR SYBR Green Master Mix (Vazyme). The primer sequences used in this study were as follows:

Nlrp3, 5'-AACATTTCGAGATTGTGGTTGGG-3' (forward)
5'-GTGCGTGAGATTCTGATTAGTGCTG-3' (reverse)
IL-1 β , 5'-AGGCTGCTCTGGGATTC-3' (forward)
5'-GCCACAACAACCTGACGC-3' (reverse)
GAPDH, 5'-AAGGTCGGAGTCAACGGATTT-3' (forward)
5'-AGATGATGACCCTTTTGGCTC-3' (reverse)
Nrf2, 5'-CGAGATATACGCAGGAGAGGTAAGA-3' (forward)
5'-GCTCGACAATGTTCTCCAGCTT-3' (reverse)
GCLC, 5'-GCTGTCTTGCAGGGAATGTT-3' (forward)
5'-ACACACCTTCCTTCCCATTTG-3' (reverse)
GCLM, 5'-TTGGAGTTGCACAGCTGGATT-3' (forward)
5'-TGGTTTTACCTGTGCCCCACTG-3' (reverse)
GST- 5'-CCTGTACCAGTCCAATACCATCCT-3' (forward)
5'-TCCTGTGTCCTTCCCATA-3' (reverse)

Transfection of Nrf2 plasmid and Nrf2 siRNA

Nrf2 plasmid was donated by Professor Siwang Yu from Peking University (Beijing, China) and Nrf2 siRNA was purchased from Santa Cruz Biotechnology, Inc. and transfected according to the manufacturer's instruction of Efect Transfection reagent (Vazyme).

Measurement of intracellular ROS level

ROS Assay kit purchased from Beyotime Institute of Biotechnology was used according to the manufacturer's

instructions. Cells were cultured in a 12-well plate, treated with various stimulations, and then, the cells were harvested and incubated with 100 μ M DCFH-DA attenuated with serum-free medium for 20 min at 37°C in the dark, washed twice with cold PBS. The fluorescence intensity was measured by the FACSCalibur flow cytometer (Becton Dickinson) at Ex./Em. = 488/525 nm.

Measurement of production of cellular superoxide

THP-Ms superoxide levels were measured with dihydroethidium (DHE, Beyotime Institute of Biotechnology), tBHQ-pretreated THP-Ms suffered different stimulations, stained by DHE (10 μ M, 30 min, 37°C) in the dark in a humidified chamber, briefly washed by PBS, and detected by FACS.

Hydrogen peroxide assay

The Hydrogen Peroxide Assay Kit was purchased from Beyotime Institute of Biotechnology. First, the cells were lysed by the schizolysis solution supplied by the kit. Then, the supernatants were collected by centrifuging at 1100 g for 10 min at 4°C. Next, 50 μ l of supernatants and 100 μ l of test solutions were mixed and placed at room temperature for 20 min and measured instantly with a microplate reader at a wavelength of 570 nm. Finally, the concentration of H₂O₂ released was calculated according to a standard concentration curve that originated from standard solutions on identical experiments.

Measurement of SOD activity

The SOD activity kit purchased was used according to the manufacturer's instructions. Cells were cultured in a six-well plate and treated with various stimulations. SOD activity was tested by measuring its inhibition on the process of xanthine oxidase catalyzing xanthine to generate superoxide anion free radicals, and then, the superoxide radicals could oxidize hydroxylamine to generate nitrite, which reacts with a developer to give a purple color. Based on this principle, we determine the SOD1 activity. The detailed procedures were performed as per the manufacturer's instruction.

Measurement of catalase activity

The Catalase Activity Kit purchased from Beyotime Institute of Biotechnology was used according to the manufacturer's instructions.

Measurement of GSH

The GSH kit purchased from Beyotime Institute of Biotechnology was used according to the manufacturer's instructions.

Alum-induced peritonitis

Mice were injected intraperitoneally with 700 μ g alum diluted in sterile PBS. Twelve hours later, peritoneal lavage was collected and PECs were stained with cell surface markers CD11b, Ly6C, and Ly6G. Flow cytometry analysis was performed by the FACSCalibur flow cytometer (Becton Dickinson).

HE staining

Peritoneal macrophages were seeded in six-well plates and cultured for 8 h, and the standard HE staining was performed according to the manufacturer's instructions. Hematoxylin was used to stain nucleic acids and eosin was used to stain the cytoplasm and extracellular matrix.

Statistical analyses

Results are presented as the mean $\pm$ SD. Statistical analyses were performed with the *t*-test for two groups or one-way ANOVA (GraphPad Software) for multiple groups. *p* Values <0.05 were considered significant.

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Author Disclosure Statement

The authors state no conflicts of interest exist.

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Address correspondence to:

Prof. Rong Hu

Department of Physiology

State Key Laboratory of Natural Medicines

China Pharmaceutical University

24 Tongjia Xiang

Nanjing 210009

China

E-mail: ronghu@cpu.edu.cn

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Abbreviations Used

ARE = antioxidant response element

ASC = apoptosis-associated speck-like protein that contains a CARD

ATP = adenosine triphosphate

BMDM = bone marrow-derived macrophage

GSH = glutathione

IL-18 = interleukin-18

IL-1 β = interleukin-1 β

LPS = lipopolysaccharide

NAC = N-acetyl-cysteine

NF- κ B = nuclear factor- κ B

NLR = NOD-like receptor

NLRP3 = NLR family, pyrin domain containing 3

Nrf2 = nuclear factor erythroid 2-related factor 2

PEC = peritoneal exudate cell

PMA = phorbol myristate acetate

ROS = reactive oxygen species

SOD = superoxide dismutase

tBHQ = tert-butylhydroquinone

THP-Ms = THP-1 cell-derived macrophages

TNF- α = tumor necrosis factor α