Characterization of a Murine Model System to Study MicroRNA-147 During Inflammatory Organ Injury

Boyun Kim,1 Victor Guaregua,1 Xuebo Chen,1 Chad Zhao,2 Wanyi Yeow,1 Nathaniel K. Berg,1 Holger K. Eltzschig,1 and Xiaoyi Yuan1,3

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Abstract—Inflammatory organ injury and sepsis have profound impacts on the morbidity and mortality of surgical and critical care patients. MicroRNAs are small RNAs composed of 20–25 nucleotides that have a significant contribution to gene regulation. MicroRNA-147 (miR-147), in particular, has been shown to have an emerging role in different physiological functions such as cell cycle regulation and inflammatory responses. However, animal model systems to study tissue-specific functions of miR-147 during inflammatory conditions in vivo are lacking. In the present study, we characterize miR-147 expression in different organs and cell types. Next, we generated a transgenic mouse line with a floxed miR-147 gene. Subsequently, we used this mouse line to generate mice with whole-body deletion of miR-147 (miR-147−/−) by crossing “floxed” miR-147 mice with transgenic mice expressing Cre recombinase in all tissues (CMVcre mice). Systematic analysis of miR-147−/− mice demonstrates normal growth, development, and off-spring. In addition, deletion of the target gene in different organs was successful at baseline or during inflammation, including the heart, intestine, stomach, liver, spleen, bone marrow, lungs, kidneys, or stomach. Moreover, miR-147−/− mice have identical baseline inflammatory gene expression compared to C57BL/6 mice, except elevated IL-6 expression in the spleen (7.5 fold, p < 0.05). Taken together, our data show the successful development of a transgenic animal model for tissue and cell-specific deletion of miR-147 that can be used to study the functional roles of miR-147 during inflammatory organ injury.

KEY WORDS: Inflammation; Organ injury; MicroRNA; Transgenic mouse model; MicroRNA-147.

INTRODUCTION

Inflammation is commonly observed in organ injuries including acute respiratory distress syndrome (ARDS) [1–3], myocardial infarction [4], acute kidney injury [5–7], and in sepsis and septic shock [8, 9], which contribute significantly to the mortality and morbidity of surgical or critical care patients. Sepsis and septic shock alone affect more than 31.5 million patients globally each year [8].
Different cell populations play a distinct role in the pathogenesis of inflammatory organ injuries. For example, innate immune cells such as macrophages and neutrophils are involved in the acute phase of inflammation in response to the pathogen-associated molecular patterns (PAMP) and danger-associated molecular patterns (DAMP) and contribute to the production of early inflammatory cytokines including tissue necrosis factor (TNF) and interleukin 1 (IL-1) [10]. Endothelial cells and epithelial cells could further respond to DAMP to promote tissue injury and organ dysfunction [10]. So far, the treatment option of inflammatory organ injury involves correcting the underlying causes and supportive care [11]. Thus, understanding the pathogenesis and the search for novel therapeutic targets for inflammatory organ injury is at the center of attention. To this front, several animal models have been developed to study pathophysiology and therapeutic targets of systemic inflammation and organ dysfunction [12]. For example, cecal ligation and puncture (CLP) is one of the most commonly used models to induce systemic inflammation and organ injury mainly caused by microbial infection [13].

MicroRNAs are small RNAs ranging from 20 to 25 nucleotides that are crucial for post-transcriptional gene regulation. So far, there are more than 2000 microRNAs identified in the human genome according to the most recent miRBase database (http://www.mirbase.org/). The main role of microRNAs is to regulate target gene expression through interaction with the untranslated region (UTR) [14]. It is estimated that microRNAs could target 60% of the human genes [15], indicating the critical role of microRNAs in many physiological and pathological conditions. Previous studies have suggested that microRNAs play crucial roles in development, cell cycle regulation, inflammatory responses, and many other physiological processes [14, 16–18]. For example, miR-223 functions as a regulator of macrophage and neutrophil differentiation and activation [19]. A recent study identified the shunting of miR-223 from neutrophil to alveolar epithelial cells to provide tissue protection during acute lung injury [20], suggesting diverse mechanisms of action for microRNAs. In the clinic, microRNAs could be therapeutically targeted via several approaches [21]. Specifically, microRNA overexpression is achieved by the delivery of microRNA mimetics, while microRNA inhibition is achieved by inhibitors such as the locked nucleic acid (LNA) [21]. For example, miR-122 has been identified as an enhancer for hepatitis C virus (HCV) replication via binding to the 5′ UTR of the viral genome [22]. MiR-122 LNA has been developed and studied by phase III clinical trials as HCV therapy showing promising results [23, 24]. Stemming from the functional diversity and the ease of therapeutic targeting, investigations on microRNAs have been intensive. Thus, developing mouse models to pinpoint the specific contribution of microRNAs in a particular tissue or cell type is of great importance.

MiR-147 (hsa-miR-147b or mmu-miR-147-3p) is located at chromosome 15 in humans and chromosome 2 in mouse. It has been continuously gaining attention as a key regulator of cell cycle progression and inflammatory responses by in vitro and pharmacological studies in vivo [25–32]. For example, miR-147 has been identified as the top upregulated microRNA in lung cancer cells that are tolerant to epidermal growth factor receptor inhibitor, and it orchestrates the metabolic shift of cancer cells for drug tolerance [25]. Another earlier study indicated that miR-147 is induced by toll-like receptor stimulation in macrophages and it is involved in the regulation of inflammatory responses [26]. These studies, along with many others, support the emerging role of miR-147 in the control of many biological processes both in homeostasis and in pathological conditions. Here, we developed a transgenic mouse line using the Cre-flox system for germline and conditional targeting of miR-147 in vivo. The generation of this mouse line will facilitate the study of tissue-/cell-specific contribution of miR-147 in inflammatory organ injury.

**MATERIALS AND METHODS**

**Animals.** Animal procedures were approved by the Institutional Animal Care and Use Committee at the University of Texas Health Center (UTHealth) at Houston. C57BL/6J (wild-type), miR-147^loxP/loxP^, CMVcre mice were purchased from Jackson Laboratories (Bar Harbor, ME). All mice were housed and bred in a specific pathogen-free facility with a 12-h:12-h light:dark cycle at the Center for Laboratory Animal Medicine and Care at the UTHealth. Both gender mice with age between 8 and 12 weeks were used in this experiment. To produce whole-body deficiency of miR-147, miR-147^loxP/loxP^ mice were crossed with CMVcre^+^ mice to generate miR-147^loxP/loxP^CMVcre^+^ mice (miR-147^−/−^).

**CLP Model.** To establish endotoxin-induced organ injury models, we performed cecal ligation and puncture (CLP) procedure. Mice were anesthetized with 3–5% inhaled isoflurane for induction and 1–3% for maintenance. The cecum was ligated with 4–0 sterile suture at 1 cm from the end, and was punctured twice with a 20G needle. A
small amount of feces was extracted with around 2 mm diameter. Sham operation was conducted in the same way without ligation and puncture. Buprenorphine SR was administrated via subcutaneous injection after surgery to reduce pain, and 500 μl of sterile and prewarmed saline was injected to protect from dehydration. All procedures are performed under sterile conditions. The organs were collected 24 h after the procedure.

**Cell Lines.** HEK293, HMEC1, T84, A539, Calu3, HK-2, Caco, HL60, and THP1 cells were purchased from ATCC and cultured according to ATCC recommended conditions. Human cardiomyocytes (HCM) and human pulmonary alveolar epithelial cells (HPAEPIC) were purchased from ScienCell (Catalog #6200 and #3200, respectively) and cultured according to the manufacturer’s instruction. Human monocyte-derived macrophages (MDM) were differentiated from monocytes as previously described [33], using monocytes isolated from peripheral blood collected from healthy volunteers.

**T Cell Differentiation.** Naïve CD4 T cells were isolated from spleens dissected from 8- to 12-week-old C57BL/6J mice using the STEMCell Naïve CD4 T cell isolation kit according to the manufacturer’s instruction. Isolated cells were cultured in a concentration of 2–2.5 million/ml in complete RPMI with L-glutamine supplemented with 10% heat inactivated FBS and antimicrobial reagents. T cell differentiation to Th0/Th1/Th2/Th17/Treg condition was achieved using Th1/Th2/Th17/Treg condition as previously described [34]. Cells were cultured for 72 h and washed with PBS. After centrifugation, cell pellet was lysed by Trizol reagent for RNA isolation.

**Isolation of Blood Neutrophils, Lymphocytes, and Monocytes.** Neutrophil was isolated from blood from 8- to 12-week-old C57BL/6J mice as previously described [20]. Lymphocytes were isolated from blood from 8- to 12-week-old C57BL/6J mice as previously described [35]. Monocytes were isolated from blood from 8- to 12-week-old C57BL/6J mice using EasySep™ Mouse Monocyte Isolation Kit from STEMCELL Technologies according to the manufacturer’s recommended protocol. After centrifugation, cell pellet was lysed by Trizol reagent for RNA isolation.

**Isolation of Alveolar Epithelial Cells.** Alveolar epithelial cells were isolated as previously described [36]. In brief, 8- to 12-week-old C57BL/6J mice were euthanized by overdose of pentobarbital. After opening up the chest cavity, lungs were perfused with 10 ml of PBS and a small incision was made at the trachea for the insertion of 20G blunt ended catheter. 1.5 ml of 5 unit/ml dispase in DMEM/F12 media was instilled intratracheally via the catheter and followed by 300 μl of 1% low melting point agarose in PBS. Lungs were removed and incubated in 0.5 ml of dispase for 45 min at room temperature. After the incubation, lung tissues were cut into small pieces and rotate for 15 min at 4°C. Digested tissues run through a 70-micron cell strainer and biotinylated antibodies for CD16/32, TER119, CD 45, and CD90 were added. Alveolar epithelial cells were negatively selected using streptavidin labeled magnetic beads. The resulting cells were incubated for 2 h to remove fibroblast. The cell pellet was lysed by Trizol reagent for RNA isolation.

**Isolation of Colon Epithelial Cells.** Colon epithelium was isolated from 8- to 12-week-old C57BL/6J mice as previously described [37]. Cell pellet was lysed by Trizol reagent for RNA isolation.

**Isolation of Renal Tubule Cells.** Kidney tubules were isolated according to a modified protocol described previously [38]. Mice were euthanized by high dose of pentobarbital sodium and the kidneys were reperfused with ice-cold PBS. The kidneys were washed with ice-cold PBS twice and were chopped into small pieces on ice. The chopped tissues were enzymatically dissociated with collagenase type II (0.25 mg/ml; Worthington) using a gentleMACS tube (Miltenyi Biotec). The tissue was incubated and dissociated in gentleMACS™ Octo Dissociator (Miltenyi Biotec) at 37°C for 30 min. After enzymatic reaction, collagenase activity was inhibited by adding one volume of Renal Epithelial Growth Medium 2 (PromoCell). To collect tubular cells, the dissociated kidney was centrifuged at 50×g for 5 min. First pellet was resuspended with Renal Epithelial Growth Medium 2 and the supernatant was centrifuged again at 50×g for 5 min. Second pellet was resuspended in the same medium. First and second pellets were combined and used for analyses.

**Quantitative Real-time Polymerase Chain Reaction.** Total RNA was isolated from cultured cells and mouse tissues using a Trizol reagent (Ambion, Life Technologies) according to the manufacturer’s instructions. Reverse transcription was carried out from 50 ng of total RNA using a High-Capacity cDNA RT kit (Applied Biosystems, ThermoFisher Scientific). TaqMan real-time PCR assay was performed to detect Il-6, Cxcl1, and 18s (internal control). TaqMan™ Gene Expression Assay (FAM): 18s (catalog number: 4351368, Assay ID: Hs99999901_s1); Il-6 (catalog number: 4351370, Assay ID: Mm00446190_m1); Cxcl1 (catalog number: 4351370, Assay ID: Mm04207460_m1). For miRNA detection, quantitative PCR was conducted in two-step PCR using TaqMan MicroRNA Assay. First, reverse transcription (RT) was performed from 10 ng total RNA using miR-147 and U6 snoRNA (internal control) primers on a Bio-Rad T100 Thermal Cycler. Second, the RT-PCR product was amplified using TaqMan MicroRNA Assay
plus the TaqMan Universal PCR Master Mix on Bio-Rad CFX384 real-time system. The relative expressions of target genes were calculated using $2^{-\Delta \Delta Ct}$ method after normalizing by 18s or U6 snoRNA. TaqMan™ MicroRNA Assay: miR-147 (Catalog #: 4440887, Assay ID: 002262); U6 snoRNA (Catalog #: 4440888, Assay ID: 001973).

Histopathological Analysis. Histological comparison of major organs including the brain, heart, liver, lung, stomach, intestine, spleen, and kidney was performed in 8- to 12-week-old, sex- and age-matched C57BL/6J mice and miR-147−/− mice as previously described [39]. After harvesting, tissues were fixed in 10% formaldehyde for 24 – 48 h and paraffin-embedded. Five-micrometer sections were cut and stained with hematoxylin and eosin. Pictures were taken from the slides using a Leica microscope.

Statistical Analysis. All data included were shown as mean ± standard error (SEM) of the mean. Data following normal distribution and have equal variances were compared using parametric two-sample unpaired t tests. Data that are not normally distributed were analyzed using non-parametric two-sample unpaired t tests using Mann-Whitney rank-sum tests. Comparison of three or more groups was achieved by one-way ANOVA and corrected for multiple comparisons. Detailed information of statistical analysis for each experiment shown was included in the figure legend. Statistical analyses were performed using GraphPad Prism software.

RESULTS

MiR-147 Expression in Different Cell Lines, Primary Cells, and Organs

Previous studies have suggested the functional role of miR-147 in many physiological processes including cell cycle regulation and metabolism [25–32]. Here, the expression pattern of miR-147 across different cells and organs is investigated by RT-qPCR. Firstly, we measured the expression level of miR-147 by RT-qPCR in different human cell lines (Fig. 1a) and found relatively high expression levels of miR-147 in T84, A549, and Calu 3 cells. The miR-147 levels in primary cells including human cardiomyocytes (HCM), human monocyte-derived macrophages (MDM), and human pulmonary alveolar epithelial cells (HPAEpiC) are further investigated (Fig. 1b). To study the miR-147 level across different organs, we harvested different organ compartments including the bone marrow, brain, heart, intestine, kidney, liver, lung, spleen, and stomach from C57BL/6J (WT) mice, and measured the miR-147 expression by RT-qPCR. MiR-147 is mostly enriched in the intestine, stomach, and heart tissue (Fig. 1c). Subsequently, different types of epithelial cells were isolated from C57BL/6J mice and higher miR-147 levels were observed in the colon epithelial cells compared to the alveolar epithelial cells and kidney tubule epithelial cells (Fig. 1d). Furthermore, no significant differences in miR-147 expression were observed in several types of blood immune cells including neutrophils, lymphocytes, and monocytes from the peripheral blood of C57BL/6J mice (Fig. 1e). Finally, there are no significant changes in the miR-147 levels across the different T helper differentiation conditions when naïve CD4 T cells isolated from the spleen of C57BL/6J mice are differentiated into T helper 0 (Th0), T helper 1 (Th1), T helper 2 (Th2), T helper17 (Th17), and regulatory T (Treg) cells in vitro (Fig. 1f).

The Expression Level of MiR-147 After CLP

After having shown the expression level of miR-147 under baseline conditions across different organs, we next investigated the expression of miR-147 under systemic inflammatory conditions. To achieve systemic organ inflammation, CLP or sham procedure was performed in 8- to 10-week-old C57BL/6J mice and major organs including the brain, heart, intestine, kidney, liver, lung, spleen, and stomach were harvest 24 h after CLP. Firstly, to understand the inflammatory conditions in each organ, the expression level of Cxcl1 in these organs was assessed by RT-qPCR in the CLP and sham group and significant induction of Cxcl1 was found in all of the organs (Fig. 2a). In addition, previous studies have indicated an upregulation of Il6 in the CLP models. Thus, we compared the expression level of Il6 in these organs by RT-qPCR in the CLP and sham group and found significant induction of these Il6 in the majority of the organs except the brain and intestine (Fig. 2b). These results indicate profound multi-organ inflammation in the CLP group. Finally, the expression pattern of miR-147 in inflamed organs was further measured by RT-qPCR. Surprisingly, miR-147 is induced in the lung, kidney, and stomach while decreased in the brain following the CLP procedure when compared to the sham group (Fig. 2c). Taken together, these studies indicated strong systemic inflammation in all major organs following CLP, and miR-147 is selectively upregulated in the lung, kidney, and stomach while down-regulated in the brain.
Strategy for the Generation of MiR-147 Floxed Mice

To address the function of miR-147 during organ injury, we generated a novel transgenic mouse line with a floxed miR-147 gene. Utilizing the Cre-loxP system, these mice allow us to generate mice with germline deletion of miR-147 (miR-147−/−) mice. Gene targeting of miR-147 conditional knockout was first established in Bruce4 C57BL/6 embryonic stem (ES) cells. To generate miR-147 conditional mice, a 114-bp region encompassing the miR-147 gene was floxed with loxP sites (Fig. 3a). An FRT-flanked neo cassette was also inserted within the floxed region for targeted ES cell selection by neomycin. Correctly targeted ES cells were confirmed by Southern blotting with Neo and 3′ probes (Fig. 3b). Subsequently, one of the clones with correct targeting (I-IC7) was selected for the generation of chimera mice. Breeding of chimera mice to flipase (flp) mice resulted in flp-mediated recombination at the FRT sites to delete the neo cassette. Litter from the mating of chimera and flipase mice was confirmed by Southern blot (Fig. 3c). These mice have been crossbred with CMVcre for the generation of miR-147−/− mice. miR-147−/− allele was confirmed by genotyping on DNA isolated from tail snip. The primer sequence and PCR protocol are shown in Fig. 3 da n de. From the representative gel picture shown in Fig. 3f, the deletion band was successfully detected in the animals. Heterozygous mice carrying miR-147 deletion allele were further bred to generate homozygous miR-147−/− mice.

Confirmation of MiR-147 Knockout and Breeding Characteristics of miR-147−/− Mice

After the identification of miR-147−/− mice, we pursue to further confirm the knockout of miR-147 under inflammatory conditions and define the breeding
characteristics. Firstly, to confirm the knockout of miR-147, several major organs including the brain, heart, intestine, kidney, liver, lung, spleen, and stomach were harvested from WT or miR-147−/− mice under baseline condition. The expression level of miR-147 was assessed by RT-qPCR and the result indicated that miR-147 is successfully deleted in all major organs (Fig. 4a). To investigate the knockout efficiency of miR-147 under inflammatory conditions, WT or miR-147−/− mice were exposed to CLP. Tissues showing significant upregulation of miR-147 during CLP, including the lung, kidney, and stomach, were harvest 24 h later for the analysis of miR-147 expression. miR-147−/− mice showed complete abolishment of miR-147 levels compared to WT mice (Fig. 4b), indicating successful knockout of miR-147 during inflammatory conditions. Next, the reproductive performance and growth curves were assessed in miR-147−/− mice. A total of 30 mice in the WT group and 29 mice in the miR-147−/− were born in the study period. There are no significant differences in litter size and gender ratio (Fig. 4c). The weight of each mouse from the WT and miR-147−/− groups was recorded for the generation of weight curve. There is no significant difference in weight gain comparing WT and miR-147 as indicated in the growth curve (Fig. 4d). Furthermore, when separated based on gender, the weight curves are similar between WT and miR-147−/− mice (Fig. 4e, f). Overall, the miR-147−/− mice show a successful knockout of miR-147 under inflammatory conditions and the breeding or growth of miR-147−/− mice has similar characteristics compared to WT animals.

**Baseline Inflammation in miR-147−/− Mice**

After confirming the successful knockout of miR-147 in several organs following CLP, we next investigated the baseline inflammation in major organs and tissues to further characterize miR-147−/− mice. For this purpose, we harvested organs and tissues including the bone marrow, brain, heart, intestine, kidney, liver, lung, spleen, and stomach, from age- and gender-matched WT or miR-147−/− mice and measured the expression level of Cxcl1 and Il6 by RT-qPCR. Statistically significant differences were not detected in the expression level of Cxcl1 in different organs or tissues when comparing WT and miR-147−/− mice (Fig. 5a). However, the Il6 level is upregulated in the spleen from miR-147−/− mice compared to that from WT mice, suggesting potential increases in the baseline inflammation in the spleen resulted from miR-147 knockout.

**Fig. 2.** Organ inflammation and miR-147 levels after CLP. CLP was conducted on 8--12-week-old C57BL/6J mice, and the indicating organs were collected in 24 h. a, b Induction of inflammatory genes including Cxcl1 and Il6 in CLP group. c Relative transcript level of miR-147 in CLP group compared to sham group. All graphs are presented as mean ± SEM. *P < 0.05, **P < 0.01, ***P < 0.0001 relative to sham, by Mann-Whitney test.
147 deletion (Fig. 5b). Furthermore, we performed histological analysis on major organs including the brain, heart, intestine, kidney, liver, lung, spleen, and stomach, from age- and gender-matched WT or miR-147−/− mice by H&E staining. From the histological analysis, we did not observe significant abnormality in miR-147−/− mice when compared to WT animals (Fig. 5c). Taken together, these studies suggest that miR-147−/− mice have comparable baseline inflammation and histological characteristics in major organs with WT animals, except for an elevated Il6 expression in the spleen.

**DISCUSSION**

The present study aimed at establishing transgenic animal models to study miR-147 in organ injury. Previous studies had indicated a crucial role of miR-147 in key biological processes such as cell cycle regulation and inflammatory response. Thus, we investigated miR-147 expression levels in different cell lines, different organs, and different immune cell populations under baseline conditions. Following organ injury achieved by CLP, we observed systemic inflammation across all examined organs...
Fig. 4. Confirmation of miR-147 KO in various tissues and breeding characteristics and growth curve of miR-147<sup>−/−</sup> (KO) mice. a Basal transcript level of miR-147 in the brain, heart, intestine, liver, lung, kidney, spleen, and stomach from C57BL/6J (WT) and miR-147<sup>−/−</sup> mice (KO) (<i>n</i> = 6/group). b Transcript level of miR-147 in the lung, stomach, and kidney from WT and miR-147<sup>−/−</sup> mice following CLP procedure (<i>n</i> = 6 for WT; <i>n</i> = 5 for KO mice). c Comparison of gender ratio and average litter size from WT and KO mice that were used for the present study. <i>P</i> value from Fisher’s exact test and unpaired <i>t</i> test, respectively. d Growth curve based on body weight of all animals from WT and KO mice over 42 days. e Growth curve based on body weight of females from WT and KO mice. f Growth curve based on body weight of males from WT and KO mice. Graphs for mRNA expression represent mean ± SEM and growth curves represent mean ± SD. *<i>P</i> < 0.05, **<i>P</i> < 0.01 relative to WT, by Mann-Whitney test.
and miR-147 upregulation in the lung, stomach, and kidney. Furthermore, to address the function of miR-147 during organ injury, we generated a novel transgenic mouse line with a floxed miR-147 gene and crossed the mice with CMVcre for germline deletion of miR-147. The successful deletion of miR-147 was confirmed as miR-147−/− mice showed completely abolished miR-147 expression under inflammatory conditions. Next, we assessed the reproductive performance and growth curves of the miR-147−/− mice and observed no significant difference in weight gain between WT and miR-147−/− mice. After confirming the successful knockout of miR-147 in several organs following CLP, we next investigated the baseline inflammation in major organs and tissues using RT-qPCR and histological analysis. Results from these analyses suggest that miR-147−/− mice have baseline inflammation and histological characteristics in major organs comparable with WT animals, except an elevated Il6 expression in the spleen. Taken together, these studies confirmed the successful generation of transgenic animals to study the role of miR-147 in organ injury.

MicroRNAs are studied in vivo by several different strategies. First and foremost, pharmacological overexpression and inhibition of microRNAs are essential tools for the
therapeutic targeting of microRNAs [21]. For instance, delivery of microRNA mimic results in the overexpression of target microRNA and downregulation of the target genes. On the other hand, microRNA inhibition could be achieved by treatment of LNA or antagonomers. Expression of microRNAs in vivo could be achieved by microRNAscope, which will localize and visualize mature microRNAs on tissue slides. Transgenic animal models are instrumental for studying the functional role of microRNAs in vivo, especially models with conditional potential for cell- or tissue-specific manipulation of microRNAs [40]. Transgenic mice could be generated with the injection of a transgene into the fertilized eggs and including an exogenous promoter would facilitate the constitutive or tissue-specific overexpression [41]. Constitutive or conditional knockout of microRNAs could be achieved by gene-targeting vectors using several systems including Cre-loxP system [42] and flp-FRT system [43]. Moreover, several systems have been established for inducible modification of microRNAs that could also have the potential for inducible knockdown or overexpression, including the tetracycline (Tet)-inducible system [44] and the Cre-ER(T) system [45]. Combining the inducible system with the conditional system results in powerful tools to study the temporal and tissue-specific functional role of microRNAs.

Inflammation is commonly observed in many pathogenic conditions including infection, acute organ injury, and chronic organ dysfunctions [46–54]. Initially intended as a response to help resolve tissue injury, inflammation could become excessive and uncontrolled, which leads to further tissue injury and, occasionally, systemic inflammation [55–59]. For example, uncontrolled alveolar epithelial inflammation results in ARDS [1, 16, 60, 61], which is the common and main cause of death in COVID-19 [62, 63]. Several previous studies have suggested the importance of endogenous anti-inflammatory pathways for the control of organ inflammation. These pathways include the hypoxia-inducible factor signaling pathway [61, 64, 65], purinergic signaling pathway [66–75], microRNAs [19, 20, 76–78], resolvins [79–81], and many others [82–90]. Along these lines, those pathways not only facilitate tissue protection during pulmonary injuries but also contribute to other inflammatory organ conditions such as myocardial infarction [91–98], kidney injury [99–101], and inflammatory bowel diseases [102–108]. Thus far, the majority of the studies on miR-147 focuses on the regulation of different biological processes such as cell proliferation and drug tolerance in cancer cells [29, 31, 109–112]. Additional studies indicated that miR-147 is crucial for the mechanical stretch-induced apoptosis in myoblast [30]. Moreover, a recent study suggested that miR-147 is reduced during rat myocardial infarction models and overexpression of miR-147 provides cardiac protection [113]. Taken together, miR-147loxPloxP mice could facilitate the study of miR-147 in different organs and cell types during inflammatory organ injury by breeding with different Cre recombinase transgenic mice.

The functional role of miR-147 in inflammation has also been indicated by several studies. For instance, a study from Liu et al. demonstrated that miR-147 overexpression in macrophages could dampen TLR activation-induced cytokine production in vitro [26]. Furthermore, a recent study demonstrated that virulent factor from the Mycobacterium marinum downregulated miR-147 levels and overexpression of miR-147 dampened Mycobacterium marinum-induced cytokine production in murine macrophage cell lines in vitro [28]. These studies imply the functional role of miR-147 during infection and inflammatory stimulations in macrophages, and potentially other myeloid cell populations. Our successful generation of floxed miR-147 transgenic mice will facilitate the study of cell-specific function in vivo. Several Cre recombinase mouse lines have been developed to target different populations of macrophages and other myeloid cells [114, 115]. However, most of these Cre recombinase transgenic mice, including LysM-Cre, Csf1r-Cre, Cd11b-Cre, F4/80-Cre, and CX3CR1-Cre, have limited ability in the specific targeting of a certain myeloid cell population. Recent studies suggested that hCD68-rtTA transgenic system could facilitate the selective and inducible targeting of CD11b+ macrophages, including pulmonary recruited and interstitial macrophages [116–118]. Crossbreeding of miR-147loxPloxP mice with hCD68-rtTA and Teto-Cre mice will facilitate the functional study of miR-147 in macrophages in vivo.

CONCLUSION

In this study, we have generated a transgenic mouse line with a floxed miR-147 gene (miR-147loxPloxP) and crossing the miR-147loxPloxP mice with CMVCre mice successfully generated mice with germline deletion of miR-147 (miR-147−/−). Firstly, we demonstrate the successful deletion of the target gene in different organs under baseline or inflammatory conditions in organs that showed induction of miR-147 during inflammation. Furthermore, we show that miR-147−/− mice experience normal growth, development, and off-spring. Moreover, miR-147−/− mice have identical baseline inflammatory gene expression compared to C57BL/6 mice, except elevated IL-6 expression in the spleen (7.5 fold, p < 0.05). Taken together, our data show that we have successfully developed a transgenic animal model for tissue- and cell-
specific deletion of miR-147 that can be used to study the functional role of miR-147 during inflammation.

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AUTHOR CONTRIBUTION

B. Kim designed and performed the experiments, analyzed the data, and drafted the manuscript. V. Guaregua, X. Chen, C. Zhao, W. Yeow, and N. K. Berg performed the experiments and analyzed the data. H. K. Eltzschig edited the manuscript and provided critical advice on the study. X. Yuan designed and performed the experiments, analyzed the data, and drafted and finalized the manuscript.

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DATA AVAILABILITY

The data that support this study are available upon request.

DECLARATIONS

Ethics Approval. All included studies have approval from the IACUC or CPHS committee at the University of Texas Health Science Center at Houston.

Conflict of Interest. The authors declare no competing interests.

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