Circular RNA-0007059 protects cell viability and reduces inflammation in a nephritis cell model by inhibiting microRNA-1278/SHP-1/STAT3 signaling

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Abstract

Background: Increasing evidence has indicated that circular RNAs (circRNAs) play a role in various diseases. However, the influence of circRNAs in nephritis remains unknown.

Methods: Microarray analysis and RT-qPCR were used to detect the expression of circRNA. Type I IFN were administrated to RMC and HEK293 cells to establish a nephritis cell model. CCK-8, MTT assay, and flow cytometry were used to assess cell proliferation, viability, and apoptosis of cells. Bioinformatics analysis and dual luciferase reporter assay detect the interaction of circ_0007059, miRNA-1278, and SHP-1. Glomerulonephritis was performed in a mouse model by administration of IFNα-expressing adenovirus. IHC staining showed the pathogenic changes.

Results: In the present study, the expression of circ_0007059 in type I interferon (IFN)-treated renal mesangial cells (RMCs), lupus nephritis (LN) specimens, and HEK293 cells was downregulated compared with that in normal healthy samples and untreated cells. Circ_0007059 overexpression resulted in increased cell proliferation, cell viability, apoptosis, and inflammation-associated factors (CXCL10, IFIT1, ISG15, and MX1) in RMCs and HEK293 cells. In addition, circ_0007059 overexpression significantly restored cell proliferation and viability and inhibited IFN-induced apoptosis. Further, the increased expression resulted in reduced inflammation and the downregulation of CXCL10, IFIT1, ISG15, and MX1 in RMCs and HEK293 cells. Circ_0007059 serves as a sponge for miR-1278 so that the latter can target the 3′-untranslated region of SHP-1. Overexpressed circ_0007059 inhibited miR-1278 expression and elevated SHP-1 expression, subsequently reducing STAT3 phosphorylation. Meanwhile, miR-1278 was upregulated and SHP-1 was downregulated in LN samples and IFN-treated cells. The restoration of miR-1278 counteracted the effect of circ_0007059 on viability, apoptosis, and inflammation as well as on SHP-1/STAT3 signaling in RMCs and HEK293 cells. We also investigated the role of SHP-1 overexpression in IFN-treated RMCs and HEK293 cells; SHP-1 overexpression resulted in a similar phenotype as that observed with circ_0007059 expression.

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Background
Systemic lupus erythematosus (SLE) is a heterogeneous systemic autoimmune disease characterized by extensive clinical features (Tsokos et al. 2016). Lupus nephritis (LN) is one of the worst complications of SLE and a significant indicator of poor prognosis (Zubiria Salgado and Herrera-Diaz 2012; Hahn et al. 2012). Inflammation of the glomeruli and tubules is the most common symptom of LN, which is believed to result from the sediment of immune complexes in the kidney (Davidson and Aranow 2010; Zhen et al. 2005). The subsequent activation of a battery of inflammatory reactions and generation of type I interferon (IFN) are instrumental for studying LN (Rönnblom et al. 2006; Tang et al. 2009; Baechler et al. 2004). Unfortunately, the development of LN in lupus-prone mice requires a long time, and the condition is unstable. This is an impediment for investigating the pathogenesis of LN and development of effective treatments for this disease (Davidson and Liu 2013).

The deposition of immune complexes in renal tissue activates immune cells, including plasmacytoid dendritic cells, via STAT signaling, which attenuates the downstream expression of genes in the type I IFN pathway (TI-IFN-P) (Banchereau and Pascual 2006). Recently, several studies have demonstrated that resident nephrocytes are also directly activated by immune complexes to produce inflammatory mediators that promote the accumulation of immune cell types (Lema et al. 2001). Resident renal cells (RRCs), including renal mesangial cells (RMCs), are essential for the pathogenesis of LN (Ka et al. 2007; Krötz 2011). RMCs represent a type of RRC that can release type I IFN and aggravate autoimmune renal injury (Fairhurst et al. 2009). Nevertheless, the regulatory mechanism of type I IFN signals in RMCs remains unclear.

Circular RNAs (circRNAs) are unique noncoding RNAs (ncRNAs) expressed widely in human cells and have attracted great interest in recent years (Meng et al. 2017). As an endogenous ncRNA, circRNA is considered to be a key regulator of various biological processes (Ebbesen et al. 2017). In addition, circRNAs have been shown to be associated with cancer generation and development. Based on a previous study on lung cancer, the epithelial–mesenchymal transition and proliferation were shown to be inhibited by hsa_circ_0007059 via microRNA (miRNA)-378 inhibition (Gao et al. 2019). Overexpressed circ_0007059 can inhibit cell proliferation, repress CyclinD1 expression, and elevate p53 expression in A549 and H1975 cell lines. This is accompanied by apoptosis and the increased expression of active caspase-3 and Bax. In oral squamous cell carcinoma, circ_0007059 is associated with prognosis and affected malignant behavior via AKT/mTOR signaling (Su et al. 2019). Circ_0007059 upregulation can inhibit cell growth, invasion, and accelerate apoptosis in SCC15 and CAL27 cell lines. Furthermore, tumor formation studies in nude mice indicated that circ_0007059 has a tumor suppressive effect in vivo. However, a role for circ_0007059 in RMC behavior during nephritis has not been reported. Therefore, the current study aimed to evaluate the expression and function of circ_0007059 in IFN-induced RMCs and human embryonic kidney 293 (HEK293) cells. This will provide further understanding of the molecular mechanisms and pathogenesis of LN and lead to the development of new treatment strategies for lupus.

Methods
Patients
We enrolled 30 patients (female 30; age range 19.6–45.0 years; mean age 37.5 years) with SLE from the Affiliated Hospital of Youjiang Medical University for Nationalities from Dec 2015 to Dec 2018. The study was approved by the Ethics Committee of the Affiliated Hospital of Youjiang Medical University for Nationalities (Approval number: YYFY-LL-2020-14). All patients provided written informed consent. Renal biopsy was performed in 18 patients with continuous SLE and active nephritis; 12 were diagnosed with inactive SLE. Partial nephrectomy was performed for renal biopsy, and the tissue was immersed in 4% formaldehyde at room temperature overnight prior to sectioning for in situ hybridization. The patients were diagnosed in accordance with the diagnostic criteria of SLE established by the American College of Rheumatology (Hochberg 1997). We measured the Systemic Lupus Erythematosus Disease Activity Index (SLEDAI) score in patients’ blood. Patients with a score index of < 4 were classified into the inactive disease group, whereas patients with a score index of ≥ 4 were placed into the active disease group. The renal

Conclusions: The study indicates that circ_0007059 protects RMCs against apoptosis and inflammation during nephritis by attenuating miR-1278/SHP-1/STAT3 signaling.

Keywords: Nephritis, circ_0007059, miR-1278, SHP-1, STAT3
SLEDAI scores were measured as described previously (Pitashny et al. 2007). The proliferation of mesangial cells is considered the primary pathological change in SLE. For this study, we used ten pairs of kidney tissue samples as controls. For pathological examination, parts of the kidney were surgically removed.

**LncRNA microarray**

Total RNA from LN samples (n=7) and the adjacent normal tissue (n=7) was extracted using TRIzol. We evaluated the RNA quality using Agilent 2100 bioanalyzer (Agilent Technologies), followed by microarray analysis. We used miRNA QC tool software to analyze rRNA (Affymetrix). The rRNA ratio (28S/18S) of all samples in the experiment exceeded 1.8, and the RNA Integrity Number exceeded 8.0. For the LN tissue, we considered a 1.5-fold difference and P<0.05 as significantly differentially expressed miRNAs.

**Cell culture and stimulation**

Human primary RMCs were obtained from ScienCell Research Laboratories, and HEK293 cells were supplied by the American Type Culture Collection (Manassas, VA, USA). The phenotypes of both cell lines were checked daily. Type I IFN (1000 units/ml; PBL Interferon Source) was used for mouse and primary human RMCs.

**Cell transfection**

Circ_0007059 (pLCDH-hsa_circ_0007059) and SHP-1 (pcDNA3.1-SHP-1) overexpressing plasmids and the miR-1278 mimic (mimic sense 5' GCA UAU CAU CUA U-3') and its related control (NC mimic, 5'-UUUG UAC UAC ACA AAA GUA CUG-3') were obtained from Genecopexy Co., Ltd. (Beijing, China). Lipofectamine 3000 (Invitrogen, Carlsbad, CA, USA) was used to transfet RMCs and HEK293 cells based on the manufacturer's instructions. The transfected RMCs and HEK293 cells were collected for subsequent experiments 40 h after transfection.

**Real-time quantitative polymerase chain reaction (RT-qPCR)**

The RNeasy Mini Kit (Qiagen, Hilden, Germany) was used to isolate total RNA as per the manufacturer’s instructions. Quality of total RNA was checked by Nanodrop® ND-1000 UV–Vis Spectrophotometer. RNA was treated with 3 U/mg RNase R (Epicenter, Madison, WI) for 15 min at 37 °C to digest linear RNA. The Prime Script RT Master Mix (Takara Bio Inc., Kusatsu, Japan) with oligo (dT) or random primers was used to reverse-transcribe the treated RNA (500 ng). Quantitative PCR was performed using 2 × PCR Master Mix (Thermo Fisher Scientific, Waltham, MA). The relative expression of the different genes was calculated by the ΔΔCt method using GAPDH and U6 snRNA expression as internal controls for mRNA and miRNA, respectively. The following primers were used for qPCR: circ_0007059 F, 5'-GAG ACA GTA GCC ATC CAG C-3'; hsa_circ_0007059 R, 5'-TGA TCT GAG TCC AGG TGT T-3'; GAPDH F, 5'-TCA AGG CTG AGA ACG GGA AG-3'; GAPDH R, 5'-TCC CCC CAC TTG ATT TTG GA-3'; and U6 F, 5'-CTC GCT TCG GCA GCA CA-3'; U6 R, 5'-AAC GCT TCA CGA ATT TGC GT-3'.

**Cell Counting Kit-8 (CCK-8) assay**

Cells were plated into 96-well plates at a density of 2 × 10³ cells/well. At 0, 24, 48, and 72 h, CCK-8 (10 µl) (Biyuntian Biotechnology Co. Ltd., Shanghai, China) was added with fresh medium (90 µl) to each well and incubated the plate for 1 h. Optical density (OD) was measured with a microplate reader at 450 nm and 630 nm. The analysis of OD values and graph preparation was performed using GraphPad 5.0 (GraphPad Software, La Jolla, CA).

**MTT assay**

The MTT assay was used to assess cell viability. MTT (20 µl of 0.5 mg/mL) was added to the cells after the supernatant was discarded to dissolve the formazan dye. DMSO (150 µl) was added to each well and incubated for 10 min. OD was measured at 540 nm using Infinite M200 microplate reader (Tecan, Männedorf, Switzerland).

**Flow cytometry**

Trypsin was used to digest cells after culturing for 48 h. The apoptotic rate was assessed using Annexin V FITC/propidium iodide (PI) Apoptosis Assay Kit (Biyuntian Biotechnology Co. Ltd.) as per the manufacturer's instructions. The cells were suspended in 1 x Annexin V binding buffer and Annexin V (5 µl) and PI (1 µl) were added to the cell suspension (100 µl) and mixed. The mixture was incubated in darkness at room temperature for 15 min, and then 1 x Annexin V binding buffer (400 µl) was added to each sample to stop the reaction. The apoptosis rate was measured using FACSCalibur flow cytometer (BD Biosciences, Franklin Lakes, NJ).

**Western blot (WB) analysis**

Extracts of cells and specimens were prepared in RIPA buffer (Beyotime Biotechnology) at 4 °C. WB was performed with the cell extracts using commercially available primary antibodies. Primary antibodies used in this study includes: Antibodies to GAPDH (1:5000, ab8245, Abcam), Caspase-3 (1:1000, ab4051, Abcam), Caspase-9 (1:1000, ab25758, Abcam), cleaved Caspase-3 (1:1000, ab2302, Abcam), cleaved Caspase-9 (1:1000, ab2324, Abcam), SHP-1 (1:2000, AB227503, Abcam), STAT3...
Circ_0007059 expression is downregulated in LN specimens and IFN-treated RMCs

To study the role of circRNAs in the progression of LN, abnormal circRNA expression was evaluated in renal biopsy samples from patients with SLE using microarray analysis. The expression of several miRNAs in the kidneys of patients with LN was significantly lower than that of healthy controls (Fig. 1B). Because type I IFN is essential for LN in humans and mice, we treated RMCs and HEK293 cells with IFN to induce the LN phenotype in these models (Wolf et al. 2018). Of note, IFN treatment reduced the expression of circ_0007059 (Fig. 1C, D). Hence, our results suggest that circ_0007059 is involved in LN progression.

Circ_0007059 overexpression increases viability and suppresses apoptosis and IFN signaling in RMCs and HEK293 cells

To analyze the effect of circ_0007059 on the viability of IFN-treated RMCs and HEK293 cells, we transfected cells with a circ_0007059-overexpressing vector or control vector. RT-qPCR data confirmed that the transfection of circ_0007059 resulted in a significant increase in circ_0007059 expression levels (Fig. 2A, B). The CCK-8 assay revealed that RMC and HEK293 cell proliferation were noticeably inhibited at 24, 48 and 72 h after IFN induction, but circ_0007059 expression completely restored proliferation (Fig. 2C, D). Furthermore, as determined by the MTT assay, circ_0007059 overexpression led to a noticeable restoration in cell viability, which was suppressed by IFN-treated RMCs and HEK293 cells (Fig. 2E, F). These results suggested a beneficial role of circ_0007059 on the viability of IFN-treated RMCs and HEK293 cells.

Considering that circ_0007059 restored the IFN-impaired proliferation of RMCs and HEK293 cells, we
hypothesized that circ_0007059 may also inhibit cell apoptosis in IFN-induced RMCs and HEK293 cells. First, to test its effects on cell apoptosis, Annexin V/PI flow cytometry indicated significant apoptosis in cells treated with IFN, whereas apoptotic cell proportion was significantly reduced after circ_0007059 overexpression (Fig. 3A, B). In addition, the reduced cleavage of caspase-3 and caspase-9 indicated that circ_0007059 inhibited IFN-induced cell apoptosis (Fig. 3C, D). We also used apoptosis inhibitor QVD to clearly incriminate apoptosis. Caspase inhibitor QVD (100 nM) was administrated in IFN-treated RMC and HEK293 cells, and the expression and cleavage of caspase-3, as well as flow cytometry were performed to determine apoptosis. Expectedly, QVD also ameliorated IFN-caused apoptosis (Additional file 1: Fig. 1A). Our data also showed that QVD treatment contributed to the inhibition of Caspase-3 cleavage (Additional file 1: Fig. 1B), suggesting that circ_0007059 possessed a similar inhibitory effect with QVD on IFN-induced apoptosis.

(See figure on next page.)

**Fig. 1** Expression of circ_0007059 in kidneys, IFN-induced RMCs and HEK293 cells, and patients with lupus nephritis (LN). A microarray analysis revealed differentially expressed genes between kidney samples from patients with LN (n = 7) and normal healthy tissue (n = 7) (biological replicates, 7; technical replicates, 1; repeat time, 3). B the expression of circ_0007059 in renal biopsy samples from patients with LN (n = 30) and surrounding normal tissue samples (n = 10) by RT-qPCR (biological replicates, as indicated; technical replicates, 3; repeat time, 3). C, D treatment of RMCs and HEK293 cells with IFN (1,000 units/mL) for 24 h. RT-qPCR was used to measure the expression of circ_0007059 (biological replicates, 3; technical replicates, 2; repeat time, 3). Results are showed as the mean±SEM for biological replicates. *P < 0.05 vs. the indicated group

**Fig. 2** Influence of circ_0007059 overexpression on the cell viability of IFN-induced RMC and HEK293 cells. RMC and HEK293 cells were transfected with a circ_0007059-overexpressing vector or NC vector for 24 h, followed by treatment with 1,000 units/mL IFN for 24 h. A, B Expression level of circ_0007059 in RMC and HEK293 cells was detected by RT-qPCR (biological replicates, 3; technical replicates, 2; repeat time, 3). C, D A CCK-8 assay was used to measure the cell proliferation rate at 24, 48, and 72 h after transfection (biological replicates, 1; technical replicates, 3; repeat time 3). E, F An MTT assay was used to determine cell viability, whereas untreated cells served as controls (biological replicates, 1; technical replicates, 3; repeat time, 3). Results are expressed as the mean±SEM for technical replicates. *P < 0.05, **P < 0.01 vs. the indicated group
Fig. 2 (See legend on previous page.)
We then tested whether circ_0007059 regulates TI-IFN-P in HEK293 and RMC cells. The expression of several genes induced by IFN was evident following the treatment of RMCs with IFN, including CXCL10, IFIT1, ISG15 and MX1, whereas the overexpression of circ_0007059 consistently decreased the expression of these genes (Fig. 4A, C, E, G). Circ_0007059 exerted the same effect on IFN-treated HEK293 cells (Fig. 4B, D, F, H).

**Circ_0007059 modulates miR-1278 expression, which, in turn, regulates SHP-1 expression**

Bioinformatics analysis was used to predict the targets of circ_0007059. We discovered that circ_0007059 targets miR-1278, which, in turn, may target the 3′-UTR of SHP-1, a widely reported mediator of inflammation and apoptosis (Chong and Maiese 2007) (Fig. 5A). Next, we investigated the mechanistic connections between the circ_0007059-miR-1278 and miR-1278-SHP-1 interaction (Fig. 5B, C) using DLRA. When RMCs were transfected with an miR-1278 mimic fused to wild-type circ_0007059 and WT SHP-1, luciferase activity decreased by 55% and 50%, respectively, compared with control cells. We then evaluated the expression of SHP-1 and miR-1278 in samples from patients with SLE (n = 30) and normal samples (n = 10). It was found that the level of miR-1278 was elevated and that of SHP-1 mRNA was reduced in LN samples compared with healthy controls (n = 10) (Fig. 5D, E). The IFN treatment of RMCs and HEK293 cells induced miR-1278 expression, and miR-1278 expression was concomitantly downregulated after circ_0007059 overexpression, confirming that circ_0007059 targets miR-1278 (Fig. 5F, G). We also examined SHP-1 protein and mRNA expression in the IFN-treated RMCs and HEK293 cells to explore the association between SHP-1 and circ_0007059. IFN treatment downregulated SHP-1 expression in RMCs and HEK293 cells, and overexpressed circ_0007059 resulted in increased SHP-1 at both the mRNA and protein levels (Fig. 5H–K). In addition, WB data indicated that STAT3 phosphorylation was significantly reduced by circ_0007059 3 h after transfection (Fig. 5H–K). Thus, our findings suggest that circ_0007059 downregulates miR-1278 expression, resulting in increased SHP-1 expression in IFN-treated cells.

**MiR-1278 upregulation counteracts the impact of circ_0007059 on the viability, apoptosis, and IFN signaling pathway of RMCs and HEK293 cells**

We evaluated the effect of miR-1278 on circ_0007059-regulated viability, apoptosis, and IFN-related signaling in IFN-treated RMCs and HEK293 cells. The miR-1278 upregulation also reduced SHP-1 mRNA and protein expression and increased STAT3 phosphorylation (Fig. 6C–F). The MTT assay indicated that transfection with an miR-1278 mimic led to a reduction in viability in IFN-treated RMCs and HEK293 cells (Fig. 6G, H). In addition, Annexin V/PI flow cytometry indicated that miR-1278 upregulation reduced the number of apoptotic RMCs and HEK293 cells (Fig. 6I, J). With respect to IFN-induced signaling, CXCL10, IFIT1, ISG15, and MX1 expression was significantly restored by miR-1278 in RMCs and HEK293 cells with overexpressed circ_0007059 (Fig. 6K, L). These results suggested that circ_0007059 plays a regulatory role by attenuating miR-1278 expression.

**Overexpressed SHP-1 decreases apoptosis and inflammation and promotes proliferation in IFN-treated RMCs and HEK293 cells**

To study the effect of SHP-1 on IFN-treated RMCs and HEK293 cells, STAT3 protein was overexpressed in IFN-treated cells. At first, elevated SHP-1 expression was observed in the SHP-1 overexpression groups (Fig. 7A–D), whereas STAT3 phosphorylation was downregulated in the SHP-1 overexpression groups (Fig. 7C, D). To further evaluate the effect of SHP-1 on cell viability and apoptosis in IFN-treated cells, we assessed these processes by MTT assay and Annexin V/PI flow cytometry. SHP-1 overexpression resulted in a noticeable increase in cell viability and caused a reduction in apoptotic cell proportion compared with the NC group (Fig. 7E–H). Moreover, SHP-1 overexpression resulted
Fig. 3 (See legend on previous page.)
in the downregulation of CXCL10, IFIT1, ISG15, and MX1 expression in IFN-treated RMCs and HEK293 cells (Fig. 7I, J).

**Circ_0007059 prevents LN development through mediating miR-1278 and SHP-1**

To investigate the potential capacity of circ_0007059, miR-1278, and SHP-1 in preventing or treating LN, Lentiviral-circ_0007059, miR-1278 mimic and SHP-1 were administered to IFNα5 adenovirus-treated (NZB × NZW)F1 mice via tail vein injection. In this study, histologic examination (PAS staining) revealed that IFNα5 adenovirus-treated (NZB × NZW)F1 mice developed severe glomerulonephritis. Notably, Lentiviral-circ_0007059 administrated mice had healthy glomeruli and no significant mesangial hypercellularity or glomerular enlargement. However, upregulation of miR-1278 promoted the glomeruli lesion in the renal. In

![Diagram](image_url)
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contrast, overexpression of SHP-1 further ameliorated the severe glomerulonephritis caused by IFNα5 (Fig. 8A). IHC staining showed that renal circ_0007059 was significantly decreased in the mice with IFNα5 induction, while Lentiviral-circ_0007059 administration obviously upregulated the level of circ_0007059 and SHP-1, and down-regulated miR-1278 in kidney. Furthermore, treatment of Lentiviral-miR-1278 promoted expression of miR-1278 and reduced level of SHP-1. In addition, Lentiviral-SHP-1 induced an upregulation of SHP-1 in the kidney of IFNα5 adenovirus-treated (NZB × NZW)F1 mice (Fig. 8B). These data were consistent with in vitro experiment.

Discussion
Abnormal miRNA expression is related to liver disease severity, for example, in SLE combined with active LN kidneys (Lorenzen et al. 2011; Dai et al. 2009). Therefore, we used miRNA arrays to analyze differential circRNA expression in LN livers and renal biopsy specimens. The expression of circ_0007059 in the renal tissue of patients with LN was significantly lower than that of control patients. Moreover, the levels of circ_0007059 were reduced in IFN-treated RMCs and HEK293 cells. Circ_0007059, a circRNA discovered recently, is associated with lung cancer (Gao et al. 2019), oral squamous cell carcinoma (Su et al. 2019), and other diseases. In the present study, we found that the overexpression of circ_0007059 ameliorated the detrimental effects of IFN in RMCs and HEK293 cells by targeting the miR-1278-SHP-1-STAT3 signaling pathway.

Regarding the pathogenesis of LN, type I IFN is considered a crucial factor. In addition, the expression of IFN-inducible genes (CXCL10, IFIT1, ISG15, and MX1) is increased in patients with SLE, and their expression levels are associated with the activity and severity of LN. Exogenous IFN-alpha can expedite the development and induced IFN-associated gene expression (CXCL10, IFIT1, ISG15, and MX1) indicate that we successfully established the LN cell model.

To the best of our knowledge, this is the first study to demonstrate a novel regulatory loop of circ_0007059-miR-1278-SHP-1-STAT3 in LN development. To date, studies regarding the function of miR-1278 are scant. A previous study indicated that miR-1278 may be differentially expressed in patients with estrogen receptor-positive breast cancer, but its function in breast cancer was not reported. (Sevinc et al. 2015). In this study, compared with control samples, miR-1278 was elevated in the kidneys of the disease group. The ectopic upregulation of miR-1278 impaired cell viability, induced cell apoptosis, and promoted IFN-associated inflammation in IFN-induced RMCs and HEK293 cells with overexpressed circ_0007059. These results suggest that miR-1278 is involved in cell injury due to IFN treatment and in circ_0007059-mediated cell properties and also filled in a gap in our knowledge of the miR-1278’s function during inflammation and apoptosis.

Another novel observation in this study was that miR-1278 targets the 3'UTR of SHP-1, whereas the latter deactivated STAT3 through dephosphorylation. As non-receptor PTPs, SHP-1 (also named PTPN6) and SHP-2 (also named PTPN11) are the critical regulators of cellular development, hyperplasia, apoptosis, metastasis, differentiation, inflammation, and intermediate metabolism (Chong and Maiese 2007). SHP-1 is primarily expressed in hematopoietic cells and epithelial cells and is considered to have a negative regulatory effect on the inflammatory response (Fish et al. 2008). Previous studies have suggested that multikinase inhibitors, including sorafenib (Tai et al. 2014), dovitinib (Chen et al. 2012a), and the Mcl-1 inhibitor SC-2001 (Chen et al. 2012b), regulate the phosphatase activity of SHP-1 to increase their antitumor effects. It was also reported that the overexpression of SHP-1 can eliminate the phosphorylation of STAT33′UTR by TGFB1 to induce the epithelial–mesenchymal transition, invasion, and migration of HCC cells (Wen et al. 2018). STAT3 is a carcinogenic transcription factor that plays a key role primarily via the phosphorylation of its
Fig. 6 (See legend on previous page.)
tyrosine residues and their subsequent dimerization and translocation to the nucleus (Lim and Cao 2006; Schindler et al. 2007). The STAT3 signal is activated in various human malignant tumors and participates in the progression of many cancer cell types (Lee et al. 2019). A recent study reported that STAT3 phosphorylation is associated with apoptosis and inflammation in cadmium-induced testicular toxicity in rats (Fouad et al. 2019).
Both SHP-1 and STAT3 have been widely investigated in inflammation and apoptosis, but few studies have focused on its role in LN. In this study, the level of SHP-1 in the kidneys of patients with LN was lower than that of control patients. The passive overexpression of SHP-1 by circ_0007059 or overexpression by transfection is accompanied by restored cell viability, inhibited inflammation, and increased apoptosis in IFN-treated RMCs and HEK293 cells. Meanwhile, the passive downregulation of SHP-1 using an miR-1278 mimic resulted in impaired cell viability, induced apoptosis, and IFN-associated inflammation. We also evaluated the phosphorylation of STAT3 along with SHP-1. Our WB data confirmed that phosphorylated STAT3 levels are negatively correlated with SHP-1 expression. The data suggest that SHP-1-STAT3 signaling participates in the IFN-induced nephritis cell model and exerts a cytoprotective role.

However, there are several limitations in the present study: First, the utilization of IFNα to establish a LN cell model. A study has indicated that TLR7-regulated LN is not in a type I IFN-dependent manner (Wolf et al. 2018), suggesting that IFNα is not exclusive inductive factor during LN development; Second, IFNα was also used in the animal model. Although many studies have demonstrated a successful modeling by IFNα induction (Han et al. 2016; Adalid-Peralta et al. 2008; Haselmayer et al. 2014), the difference of pathogenicity between LN animal and human LN samples is not fully elucidated. Finally, to further confirm the role of SHP-1 during LN development, SHP-1 knockdown or knockout animal should be utilized and observed to establish a LN model.

**Conclusions**

In conclusion, the present study is the first attempt to demonstrate a correlation of the circ_0007059-miR-1278-SHP-1-STAT3 regulatory loop with LN development. Circ_0007059, miR-1278, and SHP-1 were indicated as important factors in the pathogenesis of LN. However, the main limitation to our study is that the experiments were performed primarily in an in vitro cell model in response to IFN induction. For the purpose of determining the role of circ_0007059, further studies in knockdown or knockout animal models are required. Our results and conclusions provide novel insights into the pathogenesis of SLE.
Additional file 1: Figure 1. Effect of apoptosis inhibitor QVD on IFN-α-induced apoptosis. Caspase inhibitor QVD (100 nM) was administrated in IFNs-induced treated cells, and the expression and cleavage of caspase-3, as well as flow cytometry were performed to determine apoptosis. A the number of apoptotic cells was evaluated by flow cytometry. (Biological replicates = 1; technical replicates = 1, repeat time = 3). B WB was performed to detect levels of cleaved Caspase-3 and Caspase-3 in RMCs and HEK293 cells. (Biological replicates = 1; technical replicates = 1, repeat time = 2).

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Not applicable.

Authors’ contributions
In this work, PWG, HTH and JM conceived the study and designed the experiments. YZ, DH and LLH contributed to the data collection, ZMW and CC performed the data analysis and interpreted the results. PWG, HTH and JM wrote the manuscript; FY and WYW commented on previous versions of the manuscript. All authors read and approved the final manuscript.

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Availability of data and materials
Not applicable.

Declarations
Ethics approval and consent to participate
The study was approved by the Ethics Committee of Affiliated Hospital of Youjiang Medical University for Nationalities. All patients provided written informed consent.

Consent for publication
Not applicable.

Competing interests
There are no competing financial interests in relation to the work described.

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