Basic Study

Stem cells from human exfoliated deciduous teeth ameliorate concanavalin A-induced autoimmune hepatitis by protecting hepatocytes from apoptosis

Yi-Kun Zhou, Ling-Su Zhu, Hua-Ming Huang, Sheng-Jie Cui, Ting Zhang, Yan-Heng Zhou, Rui-Li Yang

Abstract

BACKGROUND
Autoimmune hepatitis is a serious autoimmune liver disease that threatens human health worldwide, which emphasizes the urgent need to identify novel treatments. Stem cells from human exfoliated deciduous teeth (SHED), which are easy to obtain in a non-invasive manner, show pronounced proliferative and immunomodulatory capacities.

AIM
To investigate the protective effects of SHED on concanavalin A (ConA)-induced hepatitis in mice, and to elucidate the associated regulatory mechanisms.

METHODS
We used a ConA-induced acute hepatitis mouse model and an in vitro co-culture system to study the protective effects of SHED on ConA-induced autoimmune hepatitis, as well as the associated underlying mechanisms.

RESULTS
SHED infusion could prevent aberrant histopathological liver architecture caused by ConA-induced infiltration of CD3+, CD4+, tumor necrosis-alpha+, and interferon-gamma+ inflammatory cells. Alanine aminotransferase and aspartate aminotransferase were significantly elevated in hepatitis mice. SHED infusion
INTRODUCTION

Autoimmune hepatitis (AIH) is an immune-mediated liver inflammatory disease that has a variety of clinical manifestations, including occult onset without symptoms, acute onset, and even acute liver failure[4]. The course of AIH can undulate, which may begin as acute hepatitis and ultimately lead to fibrosis, cirrhosis, and liver transplantation, eventually leading to death in a short period of time. The therapeutic aim of AIH is to alleviate the state of liver inflammation and prevent fibrosis. Generally, AIH treatment requires long-term or even lifelong use of immunosuppressants, which may cause serious side effects. Ten to twenty percent of AIH patients are insensitive to immunosuppressant treatment, cannot be well controlled even after treatment, and will ultimately develop end-stage cirrhosis[5,6]. AIH is becoming a worldwide health problem, which highlights the need to explore alternative treatment strategies.

As mesenchymal stem cells (MSCs) have multilineage differentiation and immunomodulatory properties, stem cell therapies for inflammation-related diseases have recently attracted more attention. It has been reported that MSC infusion could alleviate a variety of diseases, such as autoimmune disease, nerve injury, spinal cord injury, and diabetes[8]. Previous studies have shown that injection of bone marrow (BM) MSCs, adipose MSCs, and tonsil-derived stem cells can alleviate concanavalin A (ConA)-induced AIH in mice[9]. However, the limited sources of BM MSCs have greatly narrowed their therapeutic applications. Stem cells were found in the pulp of human exfoliated deciduous teeth (SHED), which showed more pronounced proliferative and immunomodulatory capacity compared to BM MSCs[9], Yamaza et al. SHED ameliorates ConA-induced autoimmune hepatitis could therefore block ConA-induced alanine aminotransferase and aspartate aminotransferase elevations. Mechanistically, ConA upregulated tumor necrosis-alpha and interferon-gamma expression, which was activated by the nuclear factor-kappa B pathway to induce hepatocyte apoptosis, resulting in acute liver injury. SHED administration protected hepatocytes from ConA-induced apoptosis.

CONCLUSION

SHED alleviates ConA-induced acute liver injury via inhibition of hepatocyte apoptosis mediated by the nuclear factor-kappa B pathway. Our findings could provide a potential treatment strategy for hepatitis.

Key Words: Autoimmune hepatitis; Stem cells from human exfoliated deciduous teeth; Concanavalin A; Apoptosis; Nuclear factor-kappa B
et al. also reported that spleen transplantation of SHED significantly improved hepatic dysfunction, and that these stem cells directly transformed into hepatocytes in carbon tetrachloride-treated mice without cell fusion, thus suggesting that SHED may provide a feasible treatment for liver regeneration. SHED was also reported to differentiate into hepatocytes in vitro, indicating that SHED may represent the appropriate stem cells to treat hepatitis. In light of the above effects related to SHED, we questioned whether these non-traumatically collected stem cells could block the occurrence of AIH.

ConA, from the jack bean Canavalia ensiformis, binds to sugar residues on the surface of multiple cell types. It has been reported that ConA-induced AIH has the same pathogenicity and histopathological characteristics as AIH. ConA can induce hepatocyte damage by activating inflammatory cytokines produced by macrophages and T cells. It has been reported that inflammatory cytokines, such as interleukin (IL)-2, IL-6, tumor necrosis-alpha (TNF-α), and interferon-gamma (IFN-γ), play important roles in the development of acute hepatitis. Therefore, previous studies used drugs to downregulate the relevant pro-inflammatory factors through different pathways (TRADD/TRAF2, JAK2/STAT, reactive oxygen species [ROS], PI3K, and JNK), so as to reduce liver damage and treat AIH. However, the effects of these drugs are limited by some phenomena, and its exact mechanism requires further study. SHED have demonstrated profound immunomodulatory and anti-inflammatory abilities that inhibit T lymphocyte proliferation both in vitro and in vivo. Whether SHED infusion could block ConA-induced acute inflammation needs to be further investigated.

Here, we explored the effects of SHED on ConA-induced AIH in mice, as well as the potential underlying mechanisms. Our results revealed that SHED infusion alleviated ConA-induced acute liver injury via hepatocyte apoptotic inhibition mediated by the nuclear factor-kappa B (NF-κB) pathway.

**MATERIALS AND METHODS**

**Animals and treatment**

Male Balb/c mice (6-8 wk old; 23 ± 2 g) were purchased from Vital River Laboratory Animal Technology (Beijing, China). Mice were randomly assigned to a control group (n = 10; intravenous injections with saline solution), a ConA group (n = 10; intravenous injections with 20 mg/kg ConA at 20 mg/kg body weight; Sigma-Aldrich, Street Louis, MO, United States), and an SHED + ConA group (n = 10; intravenous injections with SHED cells (1 × 10^6) 7 d prior to ConA challenge). After injections of ConA for 24 h, the mice were sacrificed for further analysis (Figure 1A). Protocols were approved by the Animal Care and Use Committee of the Health Science Center, Peking University (No. 2015-186).

**Optical imaging**

Mice (n = 3) were intravenously injected with prepared DiR-labeled SHED cells in phosphate-buffered saline (PBS) and scanned at 1 d, 3 d, and 7 d post-injection using an Interactive Video Information System Lumina Series III Imaging System (Caliper Life Sciences, Perkin Elmer, United States). Ex vivo imaging was carried out immediately afterward by imaging excised major organs (heart, lung, liver, spleen, and kidney).

**Frozen sections**

SHED cells were labeled with 5-(and-6)-carboxy fluorescein diacetate succinimidyl ester (CFSE, Molecular Probes Biotec. Co) and diluted to 1 × 10^6/mL for transplantation. The liver and spleen were flash-frozen at optimum cutting temperature at 1 d, 3 d, and 7 d post-injection. Nuclear staining was performed with diamidine phenyl indoles (DAPI) mounting medium (Vector Lab, Burlingame, CA, United States). The positions of CFSE-labeled SHED cells could be traced using a confocal laser scanning microscope (Zeiss, Thornwood, NY, United States).

**Biochemical assay**

Mouse serum samples were collected for the analysis of inflammatory cytokines, alanine aminotransferase (ALT), and aspartate aminotransferase (AST). ALT and AST levels were evaluated with commercial kits, according to the manufacturer’s instructions (Nanjing Jiancheng Bioengineering Institute, Nanjing, China). TNF-α and IFN-γ levels were measured using enzyme-linked immunosorbent assay kits.
Figure 1 Pretreatment with stem cells from human exfoliated deciduous teeth alleviates concanavalin A-induced acute liver injury in mice. A: Mice were subjected to intravenous injections with stem cells from human exfoliated deciduous teeth cells (1 × 10^6) 7 d prior to a concanavalin A (20 mg/kg) challenge. Mice were sacrificed 1 d after injection; B: Levels of serum alanine aminotransferase; C: Levels of serum aspartate aminotransferase; D: Anatomical examination and hematoxylin-eosin staining showing necrotic area in the three groups. Arrows indicate massive cell death in the liver section; E: Percentage of liver tissue necrotic areas. Scale bars = 200 μm. *P < 0.05; **P < 0.01; n ≥ 3.

Flow cytometric analysis
To detect the levels of CD3+/CD4+ T cells and Th1 cells in mouse blood, anti-mouse CD3, CD4, and IFN-γ antibodies were included according to the manufacturer’s instructions. Antibodies against cell surface markers were added for 15 min at room temperature. After lysing red blood cells, leukocytes were fixed and permeabilized by using the Cytofix/Cytoperm Fixation/Permeabilization Kit as described above. The cell pellets were resuspended, fluorescein isothiocyanate-conjugated anti-mouse IFN-γ was added and incubated at 4 °C for 30 min, and flow cytometry (FACS) analysis was performed according to the manufacturer’s instructions.

Immunohistochemistry
For immunohistochemistry, sections were dewaxed with xylene, treated with gradient concentrations of ethanol, washed with deionized water, and heated in a microwave in 10 mmol/L sodium citrate (pH 6). Then, 0.05% Tween 20 was added and samples were blocked for 1 h with 5% bovine serum albumin and 0.1% Triton X-100 in PBS. Anti-CD3 (1:100), anti-CD4 (1:100), anti-total caspase3 (1:100), anti-TNF-α (1:100), and anti-IFN-γ (1:100) antibodies were added and incubated overnight at 4 °C. After washing three times, sections were incubated with secondary antibodies at room temperature for 1 h, and then observed under a microscope (Leica, Wetzlar, Germany). Positive areas were measured with Image-Pro Plus software 6.0 (Media Cybernetics, Rockville, MD, United States).
**Immunofluorescence**

After washing in PBS for 5 min, frozen tissue sections were treated with 0.1% Triton at room temperature for 30 min to promote membrane rupture, and non-specific antigen binding sites were blocked with 5% bovine serum albumin. Rabbit anti-total caspase3 (1:100), anti-phosphorylated (p) NF-κB (1:400), and anti-NF-kB (1:400) antibodies were added and incubated overnight at 4 °C. Nuclear staining was performed with DAPI mounting medium after incubation with secondary antibody for 1 h. All sections were observed with a Zeiss fluorescence microscope (Zeiss, Thornwood, NY, United States).

**Cell culture and treatment**

Murine liver NCTC-1469 cells were purchased from the Institute of Biochemistry and Cell Biology, Chinese Academy of Science (Shanghai, China). SHED were provided by the Oral Stem Cell Bank (Beijing, China) and isolated as previously reported[8]. NCTC-1469 cells were cultured in Dulbecco’s modified Eagle’s medium supplemented with 10% fetal bovine serum, 100 U/mL penicillin, and 100 U/mL streptomycin. SHED cells were cultured in α-MEM supplemented with 15% fetal bovine serum, 100 U/mL penicillin, and 100 U/mL streptomycin. NCTC cells (2 × 10⁵ cells/well) were cultivated in 6-well plates and incubated with ConA of different concentrations (10, 20, 50, 80 and 100 μg/mL), which were analyzed for cell proliferative activity by cell counting. We then selected the appropriate treatment concentration (50 μg/mL) to study whether SHED cells inhibit NCTC cell death caused by ConA in vitro. PKH26-labeled NCTC-1469 cells were inoculated into 6-well plates at a density of 2 × 10⁵ cells/well for 12 h. Cells were then cultured with 50 μg/mL ConA and SHED cells (1 × 10⁵ cells/well) for 24 h. Determination of cell seeding density was based on our preliminary experiments by which we found that this ratio is more suitable for the study of cell co-cultures. Cells on the slide were fixed and observed with a fluorescence microscope (Zeiss).

**Determination of apoptosis and DNA damage**

Apoptosis in liver tissues was detected by terminal dUTP nick-end labeling (TUNEL) staining. Paraffin-embedded sections were deparaffinized and digested with 20 μg/mL proteinase K for 30 min at room temperature. After washing with PBS, slices were incubated with the TUNEL reaction overnight at 4 °C, followed by incubation with rabbit anti-mouse fluorescein isothiocyanate, and DAPI for nuclear labeling. TUNEL+ cells were observed and counted in at least three different fields per section. For the in vitro study, NCTC 1469 cells were inoculated in 6-well plates at a density of 2 × 10⁵ cells/well, followed by the addition of ConA (50 μg/mL) and SHED (1 × 10⁶ cells/well). After 24 h of culture, cells were collected, washed with PBS, and stained using an Annexin V apoptosis detection kit. FACS was used to observe cell fluorescence. Cells on the slide were fixed, and immunofluorescence staining was then used to determine apoptotic levels. After incubation in blocking buffer, cells were incubated with anti–caspase3 (1:400), anti-phosphorylated (p) NF-κB (1:200), and anti-NF-kB (1:200) overnight at 4 °C, followed by incubation with goat anti-rabbit fluorescein isothiocyanate antibody. Nuclear staining was performed with DAPI. Sections were observed with a fluorescence microscope (Zeiss).

**Western blot analysis**

Mouse normal liver cells (NCTC 1469) were treated with ConA (50 mg/mL) or ConA + SHED cells (1 × 10⁵ cells/well, transwell). When the cells grew to 80%, proteins were extracted after 24 h of treatment. The proteins were separated by sodium dodecyl sulfate polyacrylamide-gel electrophoresis and transferred to membranes, which were then blocked for 1 h. The membranes were incubated with the following primary antibodies: Anti–pNF-κB (1:1000; Cell Signaling) and anti–NF-kB (1:1000; Cell Signaling). After washing with TBS containing Tween-20, the membranes were incubated with a secondary antibody for 1 h, and chemiluminescence was used to visualize protein bands.

**RESULTS**

**SHED moderates ConA-induced acute liver injury in mice**

To analyze the effects of SHED infusion on ConA-induced AIH, we infused the SHED cells (1 × 10⁷) 7 d prior to ConA injection (Figure 1A). The results showed that pretreatment of SHED infusion significantly inhibited the transaminase (ALT and AST) release in AIH (Figure 1B and C). Anatomical and histological examinations,
which were used to estimate liver injury, showed that the structure of liver lobule and the arrangement of liver cells were clear in the control group, while the structure and arrangement of liver lobules were aberrant, with inflammatory cell infiltration and massive necrotic areas in the ConA group. Moreover, SHED infusion could significantly restore liver structure and reduce inflammatory cell infiltration (Figure 1D and E). DiR-labeled SHED cells were detected in mouse liver and spleen 1 d, 3 d, and 7 d after the transplantation (Supplementary Figure 1A). With the extension of time after SHED injection, the intensity of dye decreased gradually. Moreover, CFSE labeled cells were detected in mouse liver and spleen, which also decreased in a time dependent manner. However, no obvious recruitment of SHED cells was observed in the lung, kidney, and heart (Supplementary Figure 1B and C). These results suggested that SHED cells recruited in the liver of mice may play protective effects on acute liver injury.

**SHED pretreatment reduces inflammation in ConA-induced hepatitis**

T cell-mediated immune responses play a crucial role in the occurrence and development of AIH. We further explored the underlying mechanism by analyzing T cell infiltration and inflammation. The results showed that the number of CD3+ and CD4+ T cells in the liver tissues of mice in the ConA group was significantly higher than that of the control group, and that SHED pretreatment markedly reduced the number of CD3+ and CD4+ T cells (Figure 2A-C). Similarly, we detected the level of CD3+ and CD4+ T cells in whole blood, and the results were consistent with previous results (Supplementary Figure 2). Hepatitis is associated with changes in inflammatory cytokine levels. TNF-α and IFN-γ were significantly increased in the ConA-treated group, while pretreatment with SHED cells significantly reduced the ConA-induced increase of TNF-α and IFN-γ in liver tissues (Figure 2D-F). In addition, biochemical assays were used to detect the levels of TNF-α and IFN-γ in mouse sera, and the findings were consistent with the results of those in the liver tissues (Figure 2G and H). Besides, we found that the number of CD4+ IFN-γ+ T cells in ConA group increased, and SHED injection could reverse this situation (Figure 2I and J). These results showed that SHED played an immunomodulatory role in AIH mice, which alleviated liver damage by inhibiting the activation of T helper 1 cell-mediated inflammation.

**SHED pretreatment inhibits ConA-induced hepatocyte apoptosis**

Since ConA could induce hepatocyte damage, we analyzed whether ConA treatment affected liver cell apoptosis. The results revealed that the ConA group showed more apoptotic liver cells, and SHED significantly reduced this apoptotic trend (Figure 3A and B). In order to test whether SHED infusion could protect liver cells from apoptosis, we used mouse liver NCTC 1469 cells co-cultured with SHED in vitro. ConA treatment decreased NCTC cell viability in a dose-dependent manner (Supplementary Figure 3A and B). NCTC cells were stimulated with ConA at a concentration of 50 μg/mL for 24 h, then co-cultured with SHED to analyze its protective effect. The data showed that pretreatment with SHED considerably increased NCTC cell viability, which was inhibited by ConA treatment (Supplementary Figure 3C and D). To verify the effects of SHED, NCTC cells were labeled with PKH26 and co-cultured with SHED, and the results showed that the number of PKH26-labeled NCTC cells significantly decreased after ConA treatment, while SHED co-culture blocked this decrease in live PKH26-labeled NCTC cells (Figure 3C and D). Furthermore, FACS showed that ConA treatment induced remarkable (Annexin-V·PI and Annexin-V·PI') cell apoptosis, which could be blocked by SHED pretreatment (Figure 3E-G). Caspase3 is one of the most important end shear enzymes in the process of apoptosis. We found that caspase3 expression was upregulated in ConA-treated NCTC cells, while SHED co-culture decreased this upregulation, as assessed by immunofluorescence staining (Figure 4A and B). Furthermore, we found that the number of caspase3+ cells was also increased in the liver of ConA-induced hepatitis mice. SHED pretreatment could block the increase of caspase3+ cells in the liver (Figure 4C and D). These results indicated that SHED infusion could protect hepatocytes from ConA-induced apoptosis.

**SHED infusion inhibits the NF-κB signaling pathway in ConA-induced hepatitis**

Inflammatory cytokines TNF-α and IFN-γ were elevated in ConA-induced hepatitis, as the NF-κB pathway plays a crucial role in this cytokine-mediated inflammation[23]. We analyzed the NF-κB pathway and found that pNF-κB expression was upregulated in ConA-treated NCTC cells, while SHED co-culture decreased this upregulation (Figure 5A and B). NF-κB also showed the same trend (Figure 5C and D). In addition, we found that the numbers of cells positive for pNF-κB (Figure 5E and F) and NF-κB...
Zhou YK et al. SHED ameliorates ConA-induced autoimmune hepatitis
Figure 2 Pretreatment with stem cells from human exfoliated deciduous teeth reduces inflammation in concanavalin A-induced hepatitis. A: CD3+ and CD4+ T cell expression in liver tissues; B: Quantitative analysis of the number of CD3+ T cells; C: Quantitative analysis of the number of CD4+ T cells; D: Tumor necrosis factor-alpha (TNF-α) and interferon-gamma (IFN-γ) expression in liver tissues; E: Quantitative analysis of the number of TNF-α+ cells; F: Quantitative analysis of the number of IFN-γ+ cells; G: Serum TNF-α levels; H: Serum IFN-γ levels. Arrows indicate positive cells in the liver section; I: Th1 cells detected by flow cytometry; J: Quantitative analysis of the percentage of CD4+ IFN-γ+ T cells; Scale bars = 100 μm. *P < 0.05; **P < 0.01; n ≥ 3.

(Figure 5G and H) were significantly increased in the ConA-treated group and significantly decreased in the SHED cell pretreated groups. These results were verified by Western blot analysis (Figure 5I), suggesting that SHED cell treatment may ameliorate liver injury in ConA-induced hepatitis, in part, through the NF-κB pathway.

DISCUSSION

AIH is an immune-mediated inflammatory disease of the liver that can exist in acute or chronic forms, and has an increasing incidence and mortality rate. It is characterized by the elevation of serum transaminase, immunoglobulin G levels, and positive circulating autoantibody[22]. Here, we reveal that pretreatment with SHED for 7 d has a protective effect against ConA-induced hepatitis by restoring liver structure and reducing serum transaminase levels. Studies have reported that several MSCs could partially alleviate ConA-induced hepatitis in mice when MSCs and ConA were co-transfused[6,7].

SHED are a brilliant resource solution for tissue engineering and immunotherapy.
Zhou YK et al. SHED ameliorates ConA-induced autoimmune hepatitis

Figure 3 Pretreatment with stem cells from human exfoliated deciduous teeth pretreatment inhibits concanavalin A-induced hepatocyte apoptosis. A: Terminal dUTP nick-end labeling staining of apoptotic cells; B: Semi-quantitative analysis of the percentage of terminal dUTP nick-end labeling+ area in liver tissues. Arrows indicate positive cell death in the liver section; C: NCTC-1469 liver cells in the indicated treatment groups. NCTC cells are labeled with PKH26; D: Quantitative analysis of NCTC-1469 liver cell number; E: NCTC cell apoptosis detected by flow cytometry; F: Quantitative analysis of Annexin V+PI- rate; G: Quantitative analysis of Annexin V+PI+ rate. Scale bars = 100 μm. *P < 0.05, **P < 0.01, n ≥ 3.
because of its wide range of sources, safety, convenience, low immune rejection, high proliferation ability, strong multidirectional differentiation potential, and significant immunomodulatory properties. Their therapeutic potential has been extensively studied in various animal disease models, and previous research has achieved promising insights for clinical treatment. SHED therapy has been used to treat refractory liver disease, which involves severe dysfunction and fibrosis, through hepatocyte transdifferentiation and paracrine mechanisms. Here, we first report that pretreatment with SHED demonstrates protective effects for acute hepatitis. ConA-induced hepatitis progresses rapidly, and causes death in more than 50% of mice in 24 h without intervention (data not shown). This phenomenon suggests that earlier SHED infusion may benefit acute hepatitis treated with stem cells. According to our results, SHED cells can be specifically recruited to the liver and spleen after tail vein injection, so as to play the role of pretreatment through blood circulation into the lesion site. Spleen and liver accumulation can be related to phagocytic cell uptake by the mononuclear phagocyte system. SHED are derived from the pulp of deciduous teeth, which are easily obtained in a non-invasive manner.

These data indicate that pretreatment with SHED may be one alternative strategy to prevent and treat severe acute hepatitis and liver injury.

T cell-secreted TNF-α and IFN-γ play important roles in inflammatory responses. Previous studies have shown that mice pretreated with TNF-α inhibitors or anti-TNF-α antibodies can reduce ConA-induced hepatitis. TNF-α and IFN-γ knockout mice could not induce liver damage following ConA stimulation. It has been reported that SHED has superior anti-inflammatory effects compared to BM-MSCs. In this study, the number of CD3 and CD4 T lymphocytes in the liver decreased following SHED injection, so as to inhibit T helper 1 cell inflammatory factors and downregulate TNF-α and IFN-γ expression in liver tissues and peripheral blood. In vitro, we found that SHED can reverse this decrease in ConA-induced liver cells, and we thus studied its effect on liver cell apoptosis. The results indicated that SHED could reverse ConA-induced liver cell apoptosis.

Several pathways, such as NF-κB, Fas/FasL, JNK, MAPK, and ROS, play crucial roles in cell apoptosis. Based on the significant protective effects of SHED, we explored the potential underlying mechanisms of SHED against ConA-induced apoptosis. The results showed that pretreatment with SHED could effectively inhibit p65 phosphorylation, suggesting that SHED may inhibit NF-κB activation by reducing TNF-α and IFN-γ expression, thus protecting mice from AIH invasion. Since apoptosis may also be regulated by death receptors or the mitochondrial pathway, the caspase family needs to be activated in both cases. The results showed that SHED may block caspase3 activation to inhibit external apoptosis (Figure 6). However, details of the mechanism and whether this signaling contributes to the protective effects of SHED on ConA-induced hepatitis require further investigation.

CONCLUSION

In summary, our results reveal that SHED exhibits protective effects in a ConA-induced AIH mouse model. SHED treatment could inhibit the activation of T helper 1 cells and reduce the secretion of TNF-α and IFN-γ to inhibit NF-κB pathway activation induced by ConA, so as to reduce hepatocyte apoptosis and inflammatory liver damage. Our results suggest that SHED could be used as a potential preventive and therapeutic strategy for acute hepatitis and liver injury.
Figure 4 Pretreatment with stem cells from human exfoliated deciduous teeth reduces Caspase3 in concanavalin A-induced hepatitis. 

**A:** Caspase3 protein expression in NCTC cells; 
**B:** Semi-quantitative analysis of caspase3 protein fluorescence intensity; 
**C:** Caspase3 protein expression in liver tissues; 
**D:** Quantitative analysis of the number of caspase3+ cells. Arrows indicate positive cells in the liver section. Scale bars = 100 μm. *P < 0.05, **P < 0.01, n ≥ 3.
Zhou YK et al. SHED ameliorates ConA-induced autoimmune hepatitis

A

B

C

D

pNF-κB

DAPI

Merge

Control

ConA

SHED + ConA

NF-κB

DAPI

Merge

Control

ConA

SHED + ConA

pNF-κB intensity

NF-κB intensity

Control

ConA

SHED + ConA

Control

ConA

SHED + ConA

\(^{\gamma}p\)
Figure 5 Pretreatment with stem cells from human exfoliated deciduous teeth reduces nuclear factor-kappa B activation in concanavalin A-induced hepatitis. A: Phosphorylated nuclear factor-kappa B (pNF-κB) protein expression in NCTC cells; B: Semi-quantitative analysis of (p)NF-κB protein fluorescence intensity; C: NF-κB protein expression NCTC cells; D: Semi-quantitative analysis of NF-κB protein fluorescence intensity; E: Phosphorylated (p)nuclear factor-kappa B (NF-κB) protein expression in liver tissues; F: Quantitative analysis of the number of NF-κB+ cells; G: NF-κB protein expression in liver tissues; H: Quantitative analysis of the number of pNF-κB+ cells; I: Immunoprecipitation and immunoblotting analysis of (p)NF-κB and NF-κB levels in different groups. Arrows indicate positive cells in the liver section. Scale bars = 100 μm. *P < 0.05, **P < 0.01, n ≥ 3.
In concanavalin A-induced acute liver injury, tumor necrosis factor-alpha and interferon-gamma activate nuclear factor-kappa B and caspase3 to induce hepatocyte apoptosis. Stem cells from human exfoliated deciduous teeth alleviate concanavalin A-induced acute liver injury via hepatocyte apoptotic inhibition, which is mediated by the nuclear factor-kappa B pathway.

**ARTICLE HIGHLIGHTS**

**Research background**
Autoimmune hepatitis (AIH) is a serious autoimmune liver disease that threatens human health globally, thus emphasizing the need to identify novel treatments. Stem cells from human exfoliated deciduous teeth (SHED), which are easy to non-invasively obtain, showed pronounced proliferative and immunomodulatory capacities.

**Research motivation**
AIH treatment requires long-term or even lifelong use of immunosuppressants, which may cause serious side effects. Some patients are insensitive to treatment and cannot be well controlled. More appropriate and effective medical technologies to cure diabetes are needed. Mesenchymal stem cell infusion could alleviate a variety of diseases such as AIH. SHED show more pronounced proliferative and immunomodulatory capacities compared to bone marrow mesenchymal stem cells, and can differentiate into hepatocytes *in vitro*. Non-traumatically collected stem cells (SHED) may therefore be a promising option for treating AIH.

**Research objectives**
To investigate the protective effects of SHED on concanavalin A (ConA)-induced hepatitis in mice, and to elucidate the associated regulatory mechanisms.

**Research methods**
We used a ConA-induced acute hepatitis mouse model (6-week-old to 8-week-old male Balb/c mice) and an *in vitro* co-culture system to study the protective effects of SHED on ConA-induced AIH, as well as the underlying mechanisms. Protocols were approved by the Animal Care and Use Committee of the Health Science Center, Peking University (No. 2015-186).

**Research results**
SHED infusion could prevent aberrant histopathological liver architecture caused by
ConA-induced infiltration of CD3+, CD4+, tumor necrosis-alpha”, and interferon-gamma” inflammatory cells. Alanine aminotransferase and aspartate aminotransferase were significantly elevated in hepatitis mice. SHED infusion could block the ConA-induced elevation of alanine aminotransferase and aspartate aminotransferase. Mechanistically, ConA upregulated tumor necrosis-alpha and interferon-gamma expression, which were activated by the nuclear factor-kappa B pathway, to induce hepatocyte apoptosis, resulting in acute liver injury. SHED administration protected hepatocytes from ConA-induced apoptosis.

**Research conclusions**

We demonstrate that SHED exhibits protective effects in a ConA-induced AIH mouse model. SHED treatment could inhibit the activation of Th1 cells, and reduce the secretion of tumor necrosis-alpha and interferon-gamma expression to inhibit ConA-induced nuclear factor-kappa B pathway activation. This reduces both hepatocyte apoptosis and inflammatory liver damage.

**Research perspectives**

These results could provide a potential prevention and therapeutic strategy for hepatitis and acute hepatic injury.

**ACKNOWLEDGEMENTS**

We thank the Oral Stem Cell Bank of China for providing the SHED cells. This article

**REFERENCES**

1. Tansel A, Katz LH, El-Serag HB, Thrift AP, Parepally M, Shakhateh MH, Kanwal F. Incidence and Determinants of Hepatocellular Carcinoma in Autoimmune Hepatitis: A Systematic Review and Meta-analysis. *Clin Gastroenterol Hepatol* 2017; 15: 1207-1217. e4 [PMID: 28215616 DOI: 10.1016/j.cgh.2017.02.006]

2. Czaja AJ. Diagnosis and Management of Autoimmune Hepatitis: Current Status and Future Directions. *Gut Liver* 2016; 10: 177-203 [PMID: 26934884 DOI: 10.5009/gnl15352]

3. Czaja AJ, Bianchi FB, Carpenter HA, Krawitt EL, Lohse AW, Manns MP, McFarlane IG, Mielini-Vergani G, Toda G, Vergani D, Vierling J, Zeniya M. Treatment challenges and investigational opportunities in autoimmune hepatitis. *Hepatology* 2005; 41: 207-215 [PMID: 15690485 DOI: 10.1002/hep.20359]

4. Squillaro T, Peluso G, Galdieri U. Clinical Trials with Mesenchymal Stem Cells: An Update. *Cell Transplant* 2016; 25: 829-848 [PMID: 26423725 DOI: 10.3727/096368915X689622]

5. Zhu X, He B, Zhou X, Ren J. Effects of transplanted bone-marrow-derived mesenchymal stem cells in animal models of acute hepatitis. *Cell Tissue Res* 2013; 351: 477-486 [PMID: 23143676 DOI: 10.1002/cttr.2035]

6. Higashimoto M, Sakai Y, Takamura M, Usai S, Nasti A, Yoshida K, Seki A, Komura T, Honda M, Wada T, Fujiuchi K, Ochiya T, Kaneko S. Adipose tissue derived stromal stem cell therapy in murine ConA-induced hepatitis is dependent on myeloid-lineage and CD4+ T-cell suppression. *Eur J Immunol* 2013; 43: 2956-2968 [PMID: 23934743 DOI: 10.1002/eji.201343531]

7. Ryu KH, Kim SY, Kim YR, Woo SY, Sung SH, Kim HS, Jung SC, Jo I, Park JW. Tonsil-derived mesenchymal stem cells attenuated concanavalin A-induced acute liver injury. *Exp Cell Res* 2014; 326: 143-154 [PMID: 24955408 DOI: 10.1016/j.yexcr.2014.06.007]

8. Miuwa M, Grottlo s H, Zhao M, Lu B, Fisher LW, Robey PG, Shi S. SHED: stem cells from human exfoliated deciduous teeth. *Proc Nail Acad Sci USA* 2003; 100: 5807-5812 [PMID: 12716973 DOI: 10.1073/pnas.0937635100]

9. Nakamura S, Yamada Y, Katagiri W, Sugito T, Ito K, Ueda M. Stem cell proliferation pathways comparison between human exfoliated deciduous teeth and dental pulp stem cells by gene expression profile from promising dental pulp. *J Endod* 2009; 35: 1536-1542 [PMID: 19840643 DOI: 10.1016/j.joen.2009.07.024]

10. Yamaza T, Alatas FS, Yuniartha R, Yamaza H, Fujiyoshi JK, Yanagi Y, Yoshimaru K, Hayashida M, Matsura T, Aijima R, Ibara K, Ohga S, Shi S, Nonaka K, Taguchi T. In vivo hepatogenic capacity and therapeutic potential of stem cells from human exfoliated deciduous teeth in liver fibrosis in mice. *Stem Cell Res Ther* 2015; 6: 171 [PMID: 26358689 DOI: 10.1186/s13287-015-0154-6]

11. Ohkoshi S, Hirono H, Nakahara T, Ishikawa H. Dental pulp cell bank as a possible future source of individual hepatocytes. *World J Hepatol* 2018; 10: 702-707 [PMID: 30386463 DOI: 10.4254/wjh.v10.i10.702]
Zhou YK et al. SHED ameliorates ConA-induced autoimmune hepatitis

12 Bies C, Lehr CM, Woodley JF. Lectin-mediated drug targeting: history and applications. Adv Drug Deliv Rev 2004; 56: 425-435 [PMID: 14969751 DOI: 10.1016/j.addr.2003.10.030]

13 Heymann F, Hamesch K, Weiskirchen R, Tacke F. The concanavalin A model of acute hepatitis in mice. Lab Anim 2015; 49: 12-20 [PMID: 25835734 DOI: 10.1177/0023677215572841]

14 Erhardt A, Biburger M, Papadopoulos T, Tiefs G. IL-10, regulatory T cells, and Kupffer cells mediate tolerance in concanavalin A-induced liver injury in mice. Hepatology 2007; 45: 475-485 [PMID: 17226743 DOI: 10.1002/hep.21498]

15 Wang L, Lu L, Zhang J, Xu K, Qian W. Stellate Cell Activation and Imbalanced Expression of TGF-β1/TGF-β3 in Acute Autoimmune Liver Lesions Induced by ConA in Mice. Biomed Res Int 2017; 2017: 2540540 [PMID: 28246592 DOI: 10.1155/2017/2540540]

16 Schumann J, Wolf D, Pahl A, Brune K, Papadopoulos T, van Rooijen N, Tiefs G. Importance of Kupffer cells for T-cell-dependent liver injury in mice. Am J Pathol 2000; 157: 1671-1683 [PMID: 11073826 DOI: 10.1016/S0002-9440(10)64804-3]

17 Li J, Chen K, Li S, Liu T, Wang F, Xia Y, Lu J, Zhou Y, Guo C. Pretreatment with Fucoidan from Fucus vesiculosus Protected against ConA-Induced Acute Liver Injury by Inhibiting Both Intrinsic and Extrinsic Apoptosis. PLoS One 2016; 11: e0152570 [PMID: 27035150 DOI: 10.1371/journal.pone.0152570]

18 Schwabe RF, Brenner DA. Mechanisms of Liver Injury. 1. TNF-alpha-induced liver injury: role of IKK, JNK, and ROS pathways. Am J Physiol Gastrointest Liver Physiol 2006; 290: G583-G589 [PMID: 16537970 DOI: 10.1152/ajpgi.00422.2005]

19 Li J, Wang F, Xia Y, Dai W, Chen K, Li S, Liu T, Zheng Y, Wang J, Lu W, Zhou Y, Yin Q, Lu J, Zhou Y, Guo C. Astaxanthin Pretreatment Attenuates Hepatic Ischemia Reperfusion-Induced Apoptosis and Autophagy via the ROS/MAPK Pathway in Mice. Mar Drugs 2015; 13: 3368-3387 [PMID: 26623842 DOI: 10.3390/md13063368]

20 Martin pastors D, Sasaki RT, Neves AD, da Silva MC. Stem Cells from Human Exfoliated Deciduous Teeth: A Growing Literature. Cells Tissues Organs 2016; 202: 269-280 [PMID: 27544531 DOI: 10.1159/000447555]

21 Mohan RR, Mohan RR, Kim WJ, Wilson SE. Modulation of TNF-alpha-induced apoptosis in corneal fibroblasts by transcription factor NF-kappaB. Invest Ophthalmol Vis Sci 2000; 41: 1327-1336 [PMID: 10798647]

22 Balitser D, Shafizadeh N, Peters MG, Ferrell LD, Alshak N, Kakar S. Autoimmune hepatitis: review of histologic features included in the simplified criteria proposed by the international autoimmune hepatitis group and proposal for new histologic criteria. Mod Pathol 2017; 30: 773-783 [PMID: 28106105 DOI: 10.1038/modpathol.2016.267]

23 Fujii H, Matsubara K, Sakai K, Ito M, Ohno K, Ueda M, Yamamoto A. Dopaminergic differentiation of stem cells from human deciduous teeth and their therapeutic benefits for Parkinsonian rats. Brain Res 2015; 1613: 59-72 [PMID: 25863132 DOI: 10.1016/j.brainres.2015.04.001]

24 Rossato C, Brandão WN, Castro SBR, de Almeida DC, Maranduba CMC, Camara NOS, Peron JPS, Silva FS. Stem cells from human-exfoliated deciduous teeth reduce tissue-infiltrating inflammatory cells improving clinical signs in experimental autoimmune encephalomyelitis. Biologicals 2017; 49: 62-68 [PMID: 28666719 DOI: 10.1016/j.biologicals.2017.06.007]

25 LEIMDORFER A. Electroencephalographic analysis of action of amidone, morphine and strychnine on the central nervous system. Fed Proc 1948; 7: 70 [PMID: 18935268 DOI: 10.1111/j.1600-0825.2007.01396.x]

26 Ishktiev N, Yaegaki K, Inai T, Tanaka T, Fushimi N, Mitev V, Okada M, Tominaga N, Ono S, Ishikawa H. Novel management of acute or secondary biliary liver conditions using hepatically differentiated human dental pulp cells. Tissue Eng Part A 2015; 21: 586-593 [PMID: 25234861 DOI: 10.1089/tissueengine.2014.0162]

27 Xie F, He J, Chen Y, Hu Z, Qin M, Hui T. Multi-lineage differentiation and clinical application of stem cells from exfoliated deciduous teeth. Hum Cell 2020; 33: 295-302 [PMID: 32006349 DOI: 10.1007/s13577-020-00323-z]

28 Constant SL, Bottomly K. Induction of Th1 and Th2 CD4+ T cell responses: the alternative approaches. Annu Rev Immunol 1997; 15: 297-322 [PMID: 9143600 DOI: 10.1146/annurev.immunol.15.1.297]

29 Wolf D, Hallmann R, Sass G, Srix M, Kästners S, Fregien B, Trustein C, Tiefs G. TNF-alpha-induced expression of adhesion molecules in the liver is under the control of TNFR1--relevance for concanavalin A-induced hepatitis. J Immunol 2001; 166: 1300-1307 [PMID: 11145713 DOI: 10.4049/jimmunol.166.2.1300]

30 Ksontini R, Colagiovanni DB, Josephs MD, Edwards CK 3rd, Tannahill CL, Solorzano CC, Norman J, Denham W, Clare-Salzler M, Mackay SL, Moldawer LL. Disparate roles for TNF-alpha and Fas ligand in concanavalin A-induced hepatitis. J Immunol 1998; 160: 4082-4089 [PMID: 9558119]

31 Otabe K, Munea T, Kawashima N, Suda H, Tsuji K, Sekiya I. Comparison of Gingival, Dental Pulp, and Periodontal Ligament Cells From the Standpoint of Mesenchymal Stem Cell Properties. Cell Med 2012; 4: 13-21 [PMID: 26858852 DOI: 10.3727/215157912X653319]

32 Li J, Xia Y, Liu T, Wang J, Dai W, Wang F, Zheng Y, Chen K, Li S, Abudumijiti H, Zhou Z, Wang J, Lu W, Zhu R, Yang J, Zhang H, Yin Q, Wang C, Zhou Y, Lu J, Zhou Y, Guo C. Protective effects of astaxanthin on ConA-induced autoimmune hepatitis by the JNK/p-JNK pathway-mediated inhibition of autophagy and apoptosis. PLoS One 2015; 10: e0120440 [PMID: 25761053 DOI: 10.1371/journal.pone.0120440]
33 Tao YY, Yan XC, Zhou T, Shen L, Liu ZL, Liu CH. Fuzheng Huayu recipe alleviates hepatic fibrosis via inhibiting TNF-α induced hepatocyte apoptosis. *BMC Complement Altern Med* 2014; 14: 449 [PMID: 25407538 DOI: 10.1186/1472-6882-14-449]

34 Zhang P, Yin Y, Wang T, Li W, Li C, Zeng X, Yang W, Zhang R, Tang Y, Shi L, Li R, Tao K. Maresin 1 mitigates concanavalin A-induced acute liver injury in mice by inhibiting ROS-mediated activation of NF-κB signaling. *Free Radiac Biol Med* 2020; 147: 23-36 [PMID: 31785331 DOI: 10.1016/j.freeradbiomed.2019.11.033]

35 Chen YY, Li RY, Shi MJ, Zhao VX, Yan Y, Xu XX, Zhang M, Zhao XT, Zhang YB. Demethyleberberine alleviates inflammatory bowel disease in mice through regulating NF-κB signaling and T-helper cell homeostasis. *Inflamm Res* 2017; 66: 187-196 [PMID: 27900412 DOI: 10.1007/s00011-016-1005-3]

36 Mehmood T, Maryam A, Zhang H, Li Y, Khan M, Ma T. Deoxyelephantopin induces apoptosis in HepG2 cells via oxidative stress, NF-κB inhibition and mitochondrial dysfunction. *Biofactors* 2017; 43: 63-72 [PMID: 27628030 DOI: 10.1002/biof.1324]

37 ELnahas EM, Zeid MS, Kawy HS, Hendawy N, Baheer W. Celecoxib attenuates depressive-like behavior associated with immunological liver injury in C57BL/6 mice through TNF-α and NF-κB dependent mechanisms. *Life Sci* 2016; 163: 23-37 [PMID: 27580523 DOI: 10.1016/j.lfs.2016.08.027]

38 Liu T, Xia Y, Li J, Li S, Feng J, Wu L, Zhang R, Xu S, Cheng K, Zhou Y, Zhou S, Dai W, Chen K, Wang F, Lu J, Zhou Y, Guo C. Shikonin Attenuates Concanavalin A-Induced Acute Liver Injury in Mice via Inhibition of the JNK Pathway. *Mediators Inflamm* 2016; 2016: 2748367 [PMID: 27293314 DOI: 10.1155/2016/2748367]

39 Mao Y, Wang J, Yu F, Cheng J, Li H, Guo C, Fan X. Gihrelin reduces liver impairment in a model of concanavalin A-induced acute hepatitis in mice. *Drug Des Devel Ther* 2015; 9: 5385-5396 [PMID: 26451093 DOI: 10.2147/DDDT.S89096]
