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***Basic Study***

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

Zhou YK *et al.* SHED ameliorates ConA-induced autoimmune hepatitis

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**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 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

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**Core Tip:** Autoimmune hepatitis poses an enhancing global burden, underscoring the need to identify novel treatments. To determine the effects of stem cells from human exfoliated deciduous teeth (SHED) on concanavalin A (ConA)-induced autoimmune hepatitis, we pretreated ConA-induced hepatitis mice with SHED. SHED blocked increased expression of alanine aminotransferase and aspartate aminotransferase, and prevented aberrant histopathological liver architecture due to infiltration of CD3+, CD4+, tumor necrosis-alpha+, and interferon-gamma+ inflammatory cells. ConA upregulated nuclear factor-kappa B-activated tumor necrosis-alpha and interferon-gamma expression, inducing hepatocyte apoptosis and acute liver injury. SHED administration protected hepatocytes from apoptosis, providing a potential treatment for hepatitis and acute hepatic injury.

**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[1]. 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[2,3]. 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[4]. 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[5–7]. However, the limited sources of BMMSCs 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 BMMSCs[8,9]. Yamaza *et al*[10] 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[11]. 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[12,13]. 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[14–16]. 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[17–19]. 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*[20]. 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 × 106) 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 *in vivo* 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 × 106/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](https://www.sciencedirect.com/topics/medicine-and-dentistry/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, according to the manufacturer's instructions (eBioscience, Inc., San Diego, CA, United States)[17].

***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-κB (1:400) antibodies were added and incubated overnight at 4 °C. Nuclear staining was performed with [DAPI](https://www.sciencedirect.com/topics/medicine-and-dentistry/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](https://www.sciencedirect.com/topics/medicine-and-dentistry/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 × 105 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-κB (1:200) overnight at 4 °C, followed by incubation with goat anti-rabbit [fluorescein isothiocyanate](https://www.sciencedirect.com/topics/medicine-and-dentistry/fluorescein-isothiocyanate) antibody. Nuclear staining was performed with [DAPI](https://www.sciencedirect.com/topics/medicine-and-dentistry/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-κB (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 × 106) 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[21]. 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 (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 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[23–25]. SHED therapy has been used to treat refractory liver disease, which involves severe dysfunction and fibrosis, through hepatocyte transdifferentiation and paracrine mechanisms[10,26]. 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[27]. 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[28]. Previous studies have shown that mice pretreated with TNF-α inhibitors or anti-TNF-α antibodies can reduce ConA-induced hepatitis[29]. TNF-α and IFN-γ knockout mice could not induce liver damage following ConA stimulation[28–30]. It has been reported that SHED has superior anti-inflammatory effects compared to BMMSCs[31]. 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[32–34]. 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[34–37]. Since apoptosis may also be regulated by death receptors or the mitochondrial pathway, the caspase family needs to be activated in both cases[38,39]. 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.

**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-gammaexpression, 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-gammaexpression 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](http://www.baidu.com/Link?url=5NjF4qYatiRHNxG4so4pW6gzW34mHqrh0gXxd8KfPVgK6962SOfM1BXRuKQAXdXkwzRbnoWmQ-MDxtNICnpx3VIUX1D0NSK-8hhOU34wq8Qv_03S6ZoDpjCsOV7J52HI).

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**Footnotes**

**Institutional animal care and use committee statement:** This study was approved by the Animal Care and Use Committee of the Health Science Center, Peking University (No. 2015-186).

**Conflict-of-interest statement:** The authors declare no conflicts of interest.

**Data sharing statement:** Data are available upon reasonable request from R L Yang at ruiliyangabc@163.com

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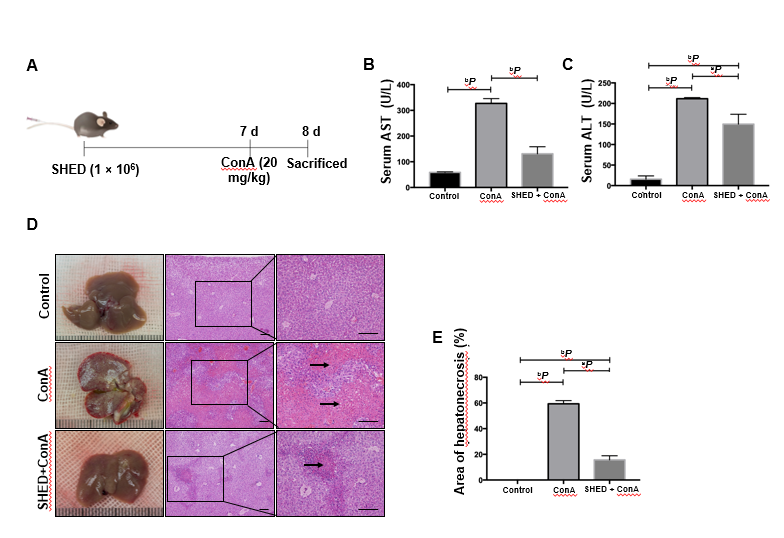
Grade C (Good): 0

Grade D (Fair): D

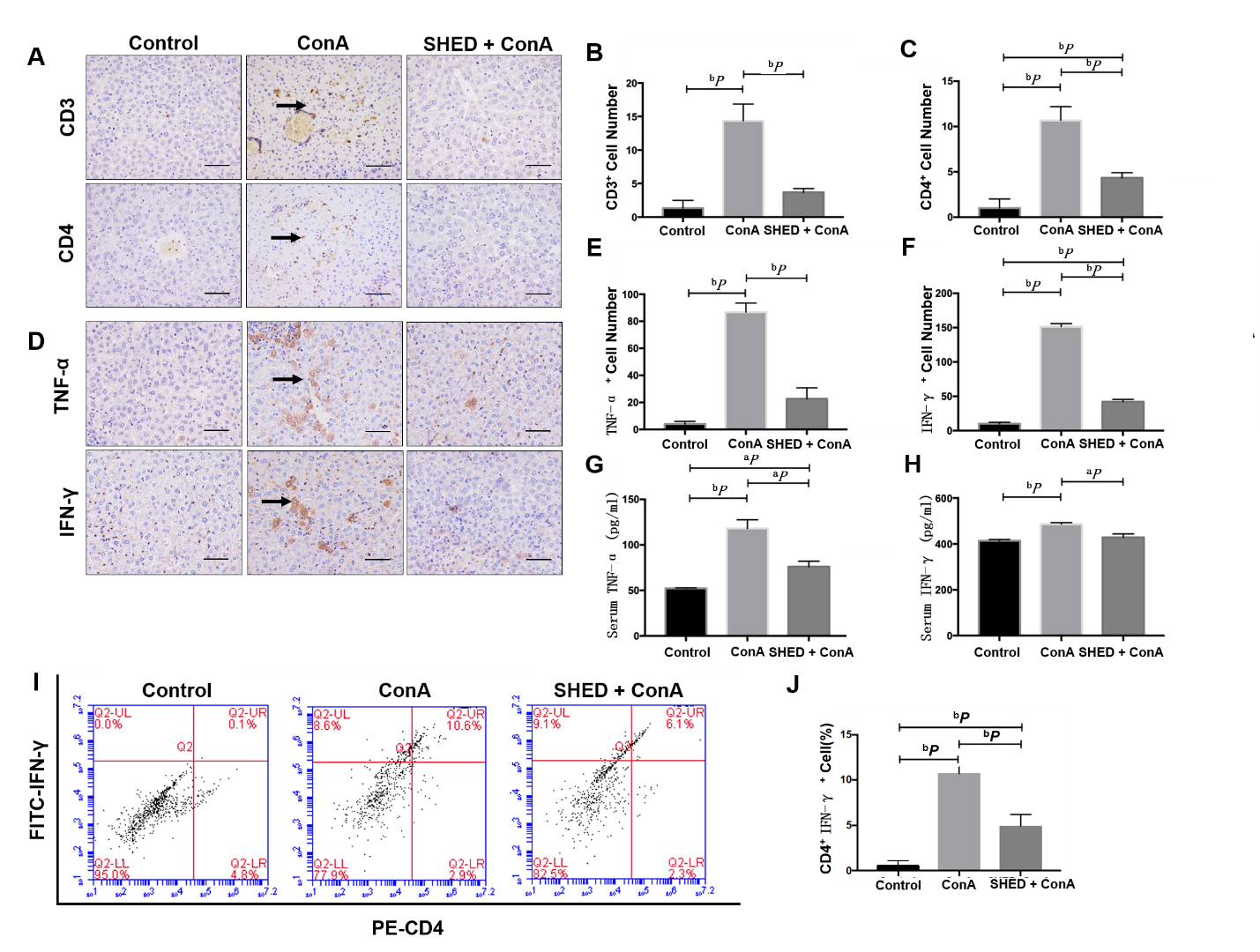
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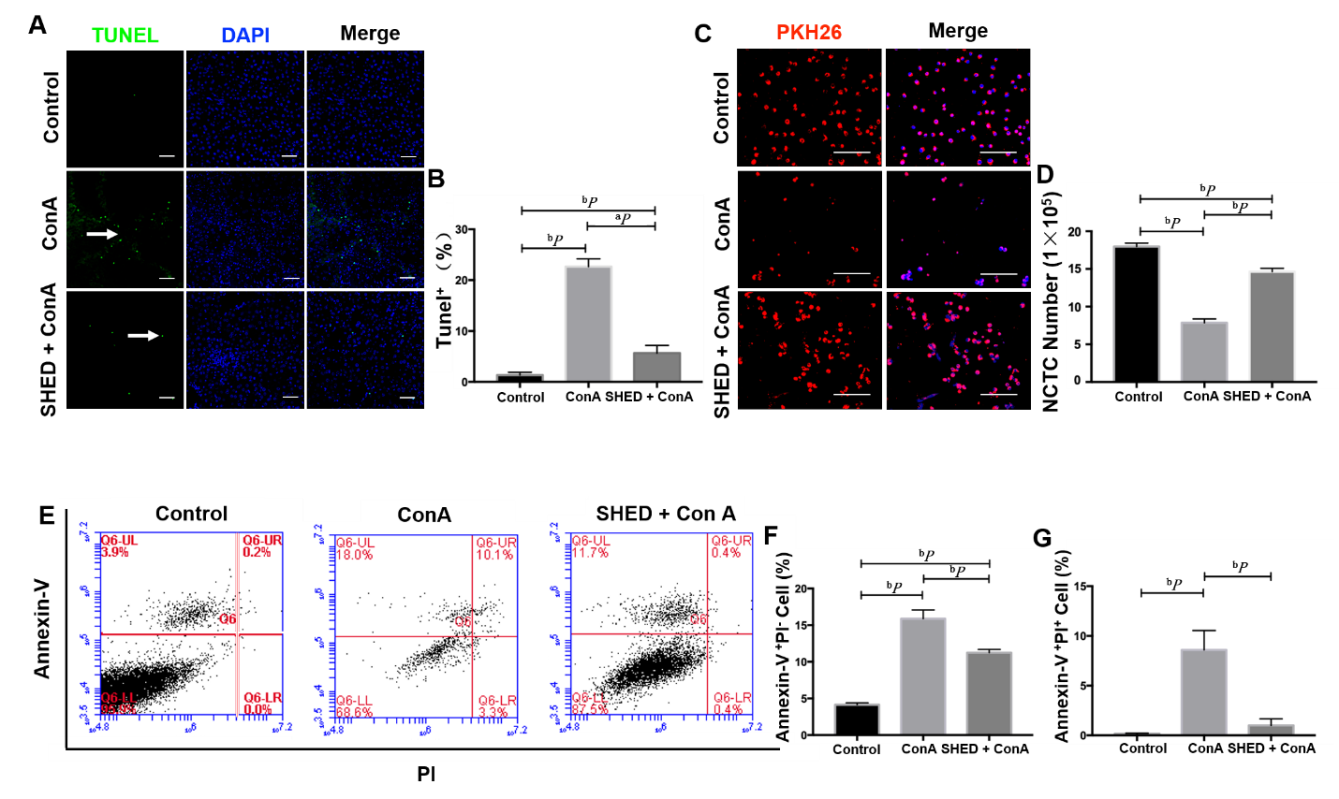
**Figure Legends**



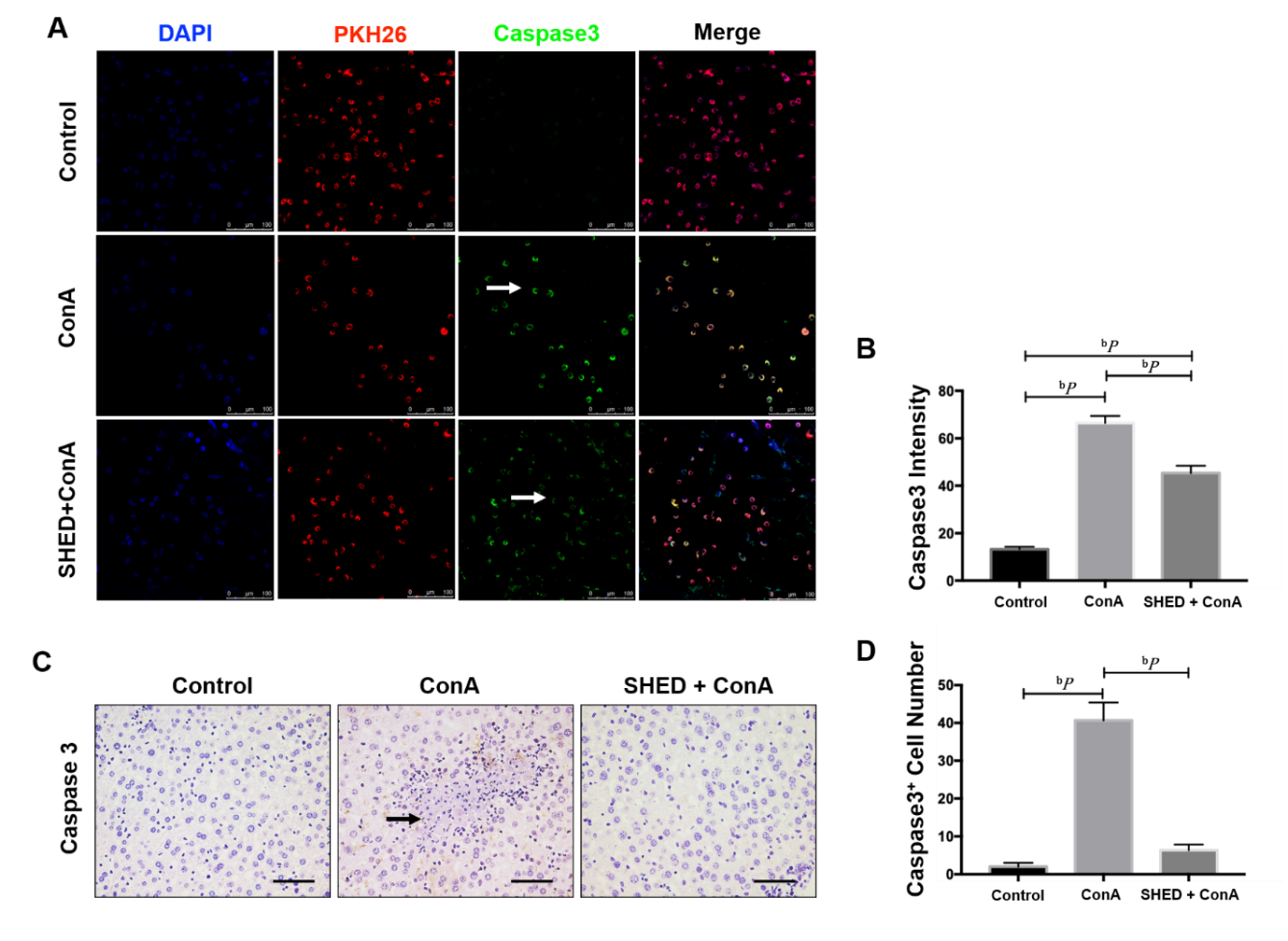
**Figure 1 Pretreatment with stem cells from human exfoliated deciduous teeth alleviates concanavalin A-induced acute liver injury in mice.** A: Mice were subjected to intravenous injections with stem cells from human exfoliated deciduous teeth cells (1 × 106) 7 d prior to a concanavalin A (20 mg/kg) 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. a*P* < 0.05; b*P* < 0.01; *n* ≥ 3.

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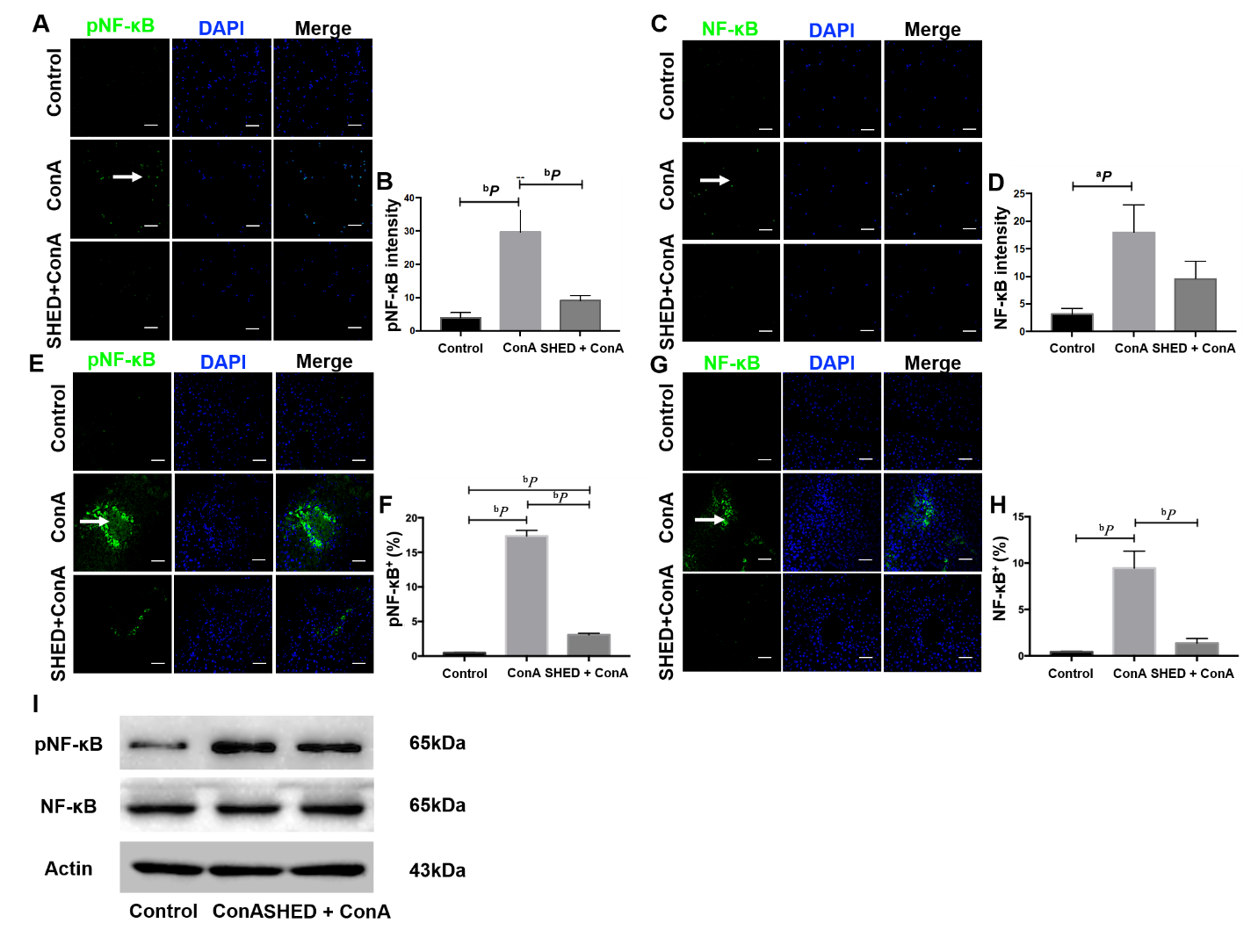
**Figure 2 Pretreatment with stem cells from human exfoliated deciduous teeth** **reduces inflammation in concanavalin A-induced hepatitis.** A: CD3+ and CD4+ T cell expression in liver tissues; B: Quantitative analysis of the number of CD3+ T cells; C: Quantitative analysis of the number of CD4+ T cells; 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. a*P* < 0.05; b*P* < 0.01; *n* ≥ 3.



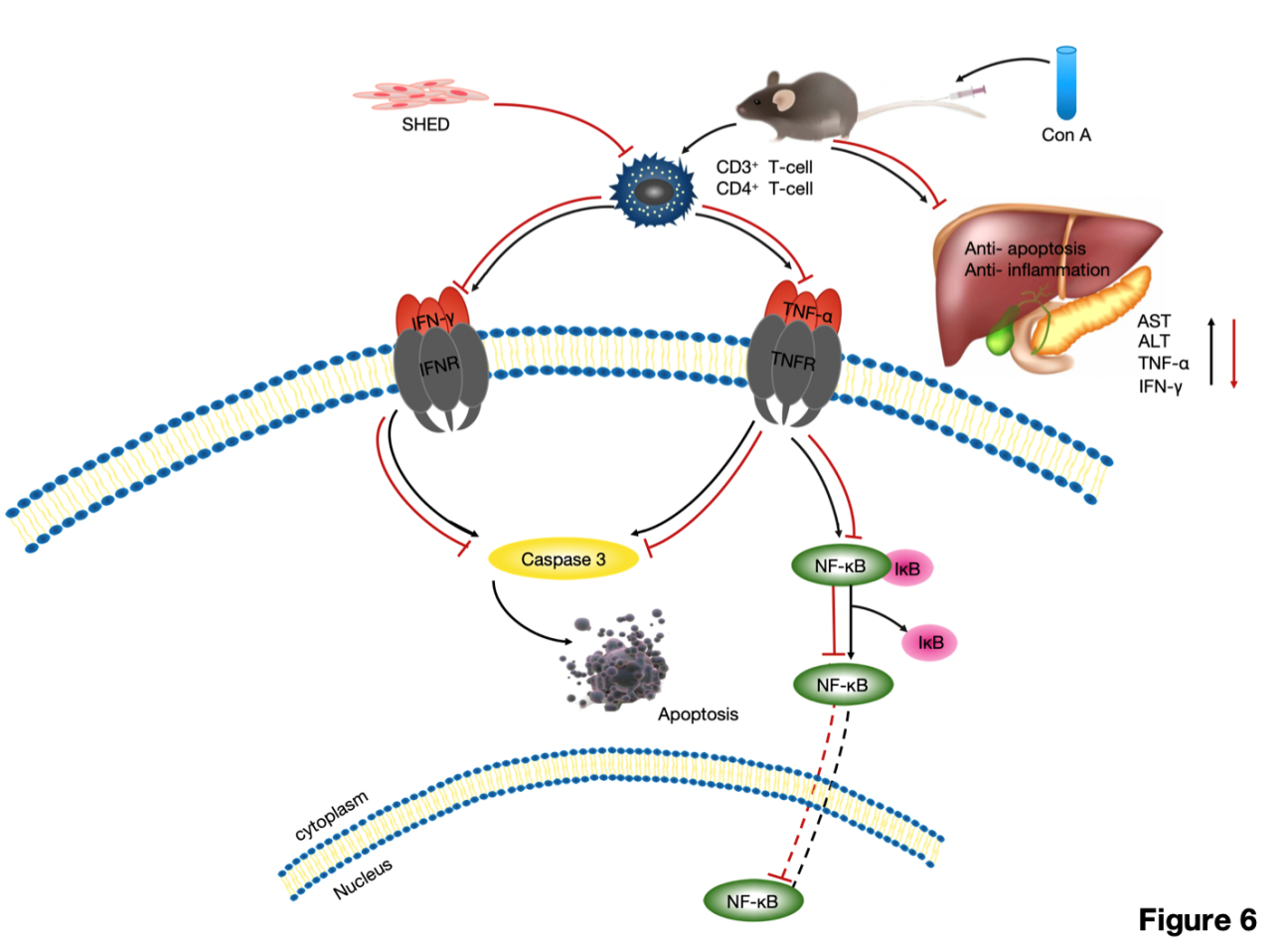
**Figure 3 Pretreatment with stem cells from human exfoliated deciduous teeth pretreatment inhibits concanavalin A-induced hepatocyte apoptosis.** A: Terminal dUTP nick-end labeling staining of apoptotic cells; B: Semi-quantitative analysis of the percentage of terminal dUTP nick-end labeling+ area in 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. a*P* < 0.05, b*P* < 0.01, *n* ≥ 3.



**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. a*P* < 0.05, b*P* < 0.01, *n* ≥ 3.



**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 [(p)NF-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. a*P* < 0.05, b*P* < 0.01, *n* ≥ 3.



**Figure 6 Schematic of treatment with stem cells from human exfoliated deciduous teeth for concanavalin A-induced hepatitis.** 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.