Hydroxychloroquine attenuates autoimmune hepatitis by suppressing the interaction of GRK2 with PI3K in T lymphocytes

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Hydroxychloroquine (HCQ) is a derivative of the heterocyclic aromatic compound quinoline, which has been used for the treatment of autoimmune diseases. The central purpose of this study was to investigate therapeutic effects and inflammatory immunological molecular mechanism of HCQ in experimental autoimmune hepatitis (AIH). Treatment with HCQ ameliorated hepatic pathologic damage, inflammatory infiltration, while promoting regulatory T cell (T reg) and down-regulated CD8+T cell differentiation in AIH mice induced by S-100 antigen. In vitro, HCQ also suppressed pro-inflammatory cytokine (IFN-γ, TNF-α, and IL-12) secretion, promoted anti-inflammatory cytokine (TGF-β1) secretion. HCQ mainly impaired T cell lipid metabolism but not glycolysis to promote T reg differentiation and function. Mechanistically, HCQ down-regulated GRK2 membrane translocation in T cells, inhibited GRK2-PI3K interaction to reduce the PI3K recruiting to the membrane, followed by suppressing the phosphorylation of PI3K-AKT-mTOR signal. Pretreating T cells with paroxetine, a GRK2 inhibitor, disturbed HCQ effect to T cells. HCQ also reversed the activation of the PI3K-AKT axis by 740 Y-P (PI3K agonist). Meanwhile, HCQ inhibited the PI3K-AKT-mTOR, JAK2-STAT3-SOCS3 and increased the AMPK signals in the liver and T cells of AIH mice. In conclusion, HCQ exhibited specific and potent therapeutic effects on AIH and attendant liver injury, which was attributed to HCQ acted on GRK2 translocation, inhibited metabolism-related PI3K-AKT and inflammation-related JAK2-STAT3 signal in T lymphocytes, thereby modulating lipid metabolism of T cell function to regulate T reg differentiation and function.

KEYWORDS
hydroxychloroquine, autoimmune hepatitis, regulatory T cells, glycolipid metabolism, G protein-coupled receptor kinase 2, PI3K-AKT axis
Introduction

Autoimmune hepatitis (AIH) is a severe inflammatory liver disease, characterized by lymphocytic infiltration and immune cells imbalance, which induce the destruction of liver parenchyma and the elevated levels of transaminase (Yang et al., 2018). The current standard therapy of AIH is glucocorticoid alone or combinating with azathioprine, which would usually produce severe adverse effects (Christen, 2019). It is necessary to develop specific and safer therapeutic agents. Hydroxychloroquine (HCQ) is derivative of the heterocyclic aromatic compound quinoline, which has been used as antimalarial agent for a long time. Currently, it is an excellent candidate for the treatment of autoimmune diseases, such as systemic lupus erythematosus (SLE) and rheumatoid arthritis (RA) (Martinez et al., 2020). It was reported that the liver injury of COVID-19 patients could be alleviated with the treatment of HCQ (Sima et al., 2021). Other study found that HCQ probably have inhibitory effects on immune cellular inflammation and activation underlying joint inflammation (Ben-Zvi et al., 2012; Rainsford et al., 2015).

Defective immunoregulation in AIH might result from $T_{reg}$ number and function reducing (Zhu et al., 2021). It was reported that HCQ rebalances Th1/$T_{reg}$-mediated immunity and ameliorates SLE (An et al., 2017). Whether HCQ play a therapeutic role by regulating $T_{reg}$ function in AIH has been attracted. Cell glucolipid metabolism is one of the key factors affecting $T_{reg}$ differentiation (Matias et al., 2021). PI3K-AKT-mTOR pathway is reported to involve in immune cells metabolism (Xiang et al., 2016). mTOR inhibition could induce AMP-activated protein kinase (AMPK) activity and lipid oxidation in $T_{reg}$, which promoted $T_{reg}$ differentiation (Pompura and Dominguez-Villar, 2018).

G protein-coupled receptor kinases 2 (GRK2) is a key participant to modulate phosphorylation-dependent G protein coupled receptors (GPCRs) desensitization, endocytosis, intracellular trafficking and re-sensitization as well as the subsequent intracellular signaling cascades. Recent data indicated that GRK2 could interact with non-GPCR substrates such as PI3K, AKT, MEK and so on (Ribas et al., 2007), which involved in inflammatory, cardiovascular disease, and tumor treatment (Cheng et al., 2021). The composition of the C-terminal domain of approximately 230 amino acids allows GRK2 to combine with PI3K, AKT, PIP2, Gp$_{0}$ and so on (Han et al., 2016). In RA, inhibiting the expression of GRK2 in membrane and increasing its expression in cytoplasm improved the abnormal proliferation of fibroblast like synovial cells. In addition, inhibition of GRK2 expression in rat spleen T cells can regulate T cell function to alleviate RA (Wang et al., 2017a; Wang et al., 2018). In the fibroblast-like synoviocytes of RA patients, the interaction of GRK2 with PI3K promoted PI3K to recruit to the membrane, which contributed to the signal transduction (Wang et al., 2020). Other study showed that GRK2 also play a crucial role in function and differentiation of $T_{reg}$ due to the interaction of GRK2 with PI3K-AKT pathway (Han et al., 2020).

Based on these observations, we hypothesize that HCQ ameliorates AIH by suppressing the inflammatory T-cell activity and promoting $T_{reg}$ differentiation. To explore this, we detected the role of HCQ in AIH therapy and mechanism in vivo and in vitro. The results suggested that HCQ attenuated inflammation by regulating T cells lipid metabolism and $T_{reg}$ differentiation, which was attributed to promoting the interaction of GRK2 with PI3K in the cytoplasm of T lymphocyte and inhibiting PI3K-AKT axis.

Materials and methods

Experimental autoimmune hepatitis model and treatment

Six-week-old male C57BL/6 mice were provided by Laboratory animal center of Anhui Medical University. All animal experimental procedures were approved by the Laboratory Animal Ethics Committee of Anhui Medical University (No. LLSC20190534). Mice were randomly divided into six groups ($n = 6$ per group) including control, AIH model, AIH+HCQ 10 mg/kg, AIH+HCQ 20 mg/kg, AIH+Prednisone (PRE) 8 mg/kg (positive control), AIH+HCQ 20 mg/kg+ PRE 8 mg/kg (drug combination) and control+HCQ 20 mg/kg (biosafety of HCQ). Hepatic syngeneic liver antigen (S-100) preparation: under non-sterile culture conditions, the livers of four female C57BL/6 mice were removed and cut into pieces on the surface of ice, then grinded on glass slides. After PBS washing, the protein in the liver cells was fully released and supernatant fluid was collected by ultracentrifugation. The experimental group was administrated by intraperitoneal injection S-100 after fully emulsified on 1st day and 7th day with 0.5 ml of 0.5–2.0 g/L and an equal volume of complete Freund’s adjuvant (CFA) (Beyotime, China). On the 14th day of modeling, the drug was administrated by gavage for 2 weeks until all mice were sacrificed. The control mice were orally administered with normal saline.

Histopathology and immunohistochemistry staining

Liver tissues were fixed in 4% paraformaldehyde for more than 24 h. Then the fixed liver tissues were stained for hematoxylin and eosin (H&E). Sections were incubated overnight at 4°C with CD3 and F4/80 primary antibodies (Proteintech, China) and incubated with biotin-labeled secondary antibody for 30 min after washing with phosphate-buffered saline. Slices were stained with chromogen diaminobenzidine and hematoxylin, dehydrated, and then treated with xylene.
ALT/AST/MDA/SOD assay

The levels of alanine aminotransferase (ALT), aspartate aminotransferase (AST), malondialdehyde (MDA) and superoxide dismutase (SOD) in the serum of mice were assessed using Kit (#C009-1-1, #C010-1-1, #A003-1 and #A001-1, Nanjing Jiancheng Bioengineering Institute, China) according to manufacturer’s instructions.

Cell culture and drug treatment

Spleen T lymphocytes were collected from the spleen of mice and activated with concanavalin A (ConA, 0.1 μM) and incubated with HCQ using doses in our concentration screening tests (100, 50, and 25 μM) for 24 h. Bone marrow derived dendritic cells (BMDCs) were collected from the tibia and femur of AIH mice. DCs were cultured with cytokines GM-CSF (20 ng/ml) and IL-4 (10 ng/ml) (Liu et al., 2015). All cells were cultured in RPMI 1640 medium (#BC-M-017, Biochannel, China) with 10% fetal bovine serum (FBS; Biological Industries, Israel) and maintained at 37°C in an incubator with 5% CO2 (Liu et al., 2014; Fang et al., 2020).

FCM analysis

Immune cells were analyzed by FCM. Briefly, the cells underwent staining with fluorochrome-antibodies (Multi Sciences Lianke Bio, China) targeting cell antigens at 4°C for 60 min. Tissue abrasive fluid and T cell suspension were stained with FITC-CD4, PE-CD25, APC-Foxp3 to identify Treg and APC-CD3, FITC-CD4, PE-CD8 to identify CD4+ or CD8+T cells. The expressions of Foxp3 were observed in CD4+CD25+ cell gate, and the expressions of CD4 and CD8 were observed in CD3+ cell gate. BMDCs were stained with FITC-CD11c, APC-CD86 and PE-MHC-II (Liu et al., 2015), the expressions of MHC-II and CD86 were calculated in CD11c+ cell gate. The gate was chosen by the compare between negative cells (without fluorescent dye) and single fluorescent dye or mixed fluorescent dye cells. Furthermore, mitochondrial membrane potential (MMP) of cells was evaluated using JC-1 molecular probes and analyzed by flow cytometry. In cells with a high MMP (∆Ψm >80–100 mV), JC-1 forms aggregates that emit red-orange fluorescence (wavelength, 590 nm), whereas in cells with low mitochondrial potential (∆Ψm <80–100 mV), JC-1 forms monomers that emit green fluorescence (wavelength, 525–530 nm). The high MMP exerted high ratio of JC-1 Red/JC-1 Green. T lymphocytes were stained with JC-1 and tested in FITC and PE channels. The gate was chosen by the clustering of cells.

Membrane and cytoplasm protein extraction

Membrane and cytoplasm protein of T lymphocytes was extracted by membrane and cytosol protein extraction kit (#P0033, Beyotime, China). The total proteins of liver tissues and spleen T lymphocytes were extracted by RIPA (#P0013C, Beyotime, China) with phosphatase inhibitor and PMSF (#P0012S, Beyotime, China). Immunoblotting was performed as previously described (Xiang et al., 2022). Primary antibodies were listed in Supplementary Table S1. Secondary antibodies were as follows: HRP-conjugated affinipure goat anti-rabbit IgG (H+L) (#SA00001-2, Proteintech, China) and HRP-conjugated affinipure goat anti-rabbit IgG (H+L) (#SA00001-1, Proteintech, China). The protein samples were visualized using the ECL-chemiluminescent kit (#WBKLS0100, Millipore, United States) and analysed by ImageJ software.

Quantitative real-time-PCR

Specifically, total RNA was extracted from liver tissues and spleen T lymphocytes by TRizol reagent (#257401, Invitrogen, United States). RNA was reverse transcribed in to cDNA by reverse transcription system kit (#R222-01, Vazyme, China). The mRNA levels were measured by qPCR using SYBR Green qPCR Master Mix (#Q111-02/03, Vazyme, China) according to the manufacturer protocol. The primers were listed in Supplementary Table S2. Gene expression was normalized to expression of β-actin.

Enzyme-linked immunosorbent assay, free fatty acid secretion and glucose uptake assay

Levels of transforming growth factor-β1 (TGF-β1), interferon-γ (IFN-γ), interleukin-12 (IL-12), and tumor necrosis factor-α (TNF-α) in cell culture supernatant and serum were assessed with the ELISA assay (Proteintech, China). Free fatty acid was examined with the free fatty acid assay kit (#A042-1-1, Nanjing Jiancheng Bioengineering Institute, China).
Institute, China), glucose uptake was examined by the glucose oxidase-peroxidase method with test kit (#361510, Rongsheng, China).

Cell counting Kit-8 assay

The proliferation of T cell was assessed using CCK-8 kit (#GK10001, GLPBIO, China). After stimulation, the reagent was added into the culture medium, and the mixture was maintained in an incubator comprising 5% CO2 + 95% air at 37°C for 2 h. The absorbance at 450 nm was detected utilizing a Microplate Reader.

Co-immunoprecipitation

To examine the association between GRK2 and PI3K or AKT in spleen T lymphocytes under different conditions, total cell lysates were immunoprecipitated using anti-GRK2 antibody and analyzed by western blot with an anti-GRK2, anti-PI3K or anti-AKT. Briefly, the preparation of lyse is as described above for western blots, and then pre-cleared using IgG. The total protein (1,000 µg of each lysate sample) was incubated with GRK2 antibody (4 µl) on a rotating shaker at 4°C overnight, the Protein A/G was added with agarose and then the small ball (20 µl) was added to each tube and placed in the rotating shaker at 4°C for 2 h. Then the tube was washed three times with cracking buffer. Resin-bound immune complexes were boiled for 5 min after protein loading buffer was added. Finally, repeating steps as Western blot.

Statistical analyses

All data were statistically analyzed using GraphPad Prism 8.0. The data were normally distributed and expressed by the mean ± standard deviation (SD). Multiple comparisons were carried out by one-way (ANOVA). Comparisons between two groups were performed using independent-sample t-tests. All experiments were performed at least three times, \( p < 0.05 \) was considered statistically significant.
Results

Hydroxychloroquine ameliorated hepatic pathologic damage and inflammatory infiltration of autoimmune hepatitis mice

In S-100-treated AIH mice, oral administration of 10 or 20 mg/kg HCQ, which referred to the report (Zheng et al., 2021), for 2 weeks markedly diminished the extent of AIH with the loss of liver architecture, congestion, lymphocytic infiltration and large area necrosis (arrows represent inflammatory infiltrations) (Figure 1A). We focused on inflammatory cell related markers, such as mature macrophages cell marker F4/80 and lymphocyte marker CD3 to investigate the alleviating effect of HCQ on the inflammatory infiltration of AIH. The results showed that the expression of CD3 and F4/80 significantly increased in the liver of AIH, which was slipped to basal levels by HCQ treatment (Figure 1B). The liver index and spleen index decreased in HCQ treatment compared with AIH group to a certain extent (Figure 2A). Serum levels of ALT and AST were decreased after HCQ treatment (Figure 2B). HCQ memorably attenuated hepatic oxidative stress with lower MDA and higher SOD compared with S-100 treatment (Figure 2C). Furthermore, the decrease of anti-inflammatory factor, TGF-β1 and the increase of proinflammatory factor, IFN-γ, TNF-α and IL-12 in serum caused by S-100 could be reversed by HCQ (Figure 2D). These effects were better or comparable to those of PRE, which was used in clinical settings against AIH. In addition, there was synergistic effect in treating AIH by combination of HCQ and PRE. Furthermore, HCQ treatment did not alter levels of ALT, AST, or inflammatory factors in healthy mice (Figures 1, 2). The results demonstrated that HCQ exerted liver-protective and anti-inflammatory effects on S-100-induced AIH and appeared to have no serious adverse effects.

Hydroxychloroquine regulated T cells function and Treg differentiation in autoimmune hepatitis

Numerous immune cells regulate the development of hepatic inflammation (Koyama and Brenner, 2017). We quantitatively determined the proportions of various immune cells of multiple organs in AIH. The results showed that, after S-100 administration, Treg, reduced in the spleen and liver, CD8+ T cells differentiated in the spleen, and bone marrow derived dendritic cells (BMDCs) matured. HCQ reversed the effect of S-100 (Figures 3A–C), but had little effect on other subtypes of immune cells in the liver and spleen of AIH mice (Date not shown). Here, we also noted that HCQ increased the fork head box protein 3 (Foxp3) mRNA expressions in the liver of AIH mice, which facilitated Treg immune tolerance. The serine protease granzyme B (GzmB), markers of CD8+ T cell-induced cytotoxicity, was reduced by HCQ in the liver of AIH mice. HCQ showed a potential immunosuppressive effect on DC-CD8+ T cell communication by upregulating the negative regulatory molecules CTLA-4 and PD-1 in the liver. The effect of HCQ to other costimulatory molecules were indistinctive (Figure 3D).

In vitro, HCQ significantly suppressed spleen T-lymphocyte proliferation in a dose dependent manner (Figure 4A). It also decreased the pro-inflammatory cytokines, and increased the anti-inflammatory cytokines production in the supernatant of T cells (Figure 4B). These effects were comparable to those of MTX, which is used to inhibit T cells proliferation and function. Of note, in vitro, HCQ strongly increased Treg and slightly retarded CD8+ T cells differentiation (Figure 4C). HCQ represented similar effects on Foxp3 and GzmB mRNA expressions in vitro compared to in vivo experiment (Figure 4D). The apparent modulation involving Treg of HCQ might be important to maintain immune homeostasis during progression of AIH.

Impaired lipid metabolism in hydroxychloroquine-modulated T lymphocyte

Manipulating the metabolism of immune cells may alter immune homeostasis (Fox et al., 2005). Induced Treg differentiate from conventional naïve CD4+ T cells under a variety of conditions that range from inflammatory environments, in the presence of particular cytokines, mainly TGF-β1, to suboptimal glycolysis and/or fatty acid oxidation signals (Murphy et al., 1990). Previous reports indicated that HCQ exerted substantial metabolic regulation effects including lipid and insulin metabolism (Hu et al., 2017). Thus, HCQ might ameliorate AIH partly by suppressing T cells metabolism, which promoted Treg development. Our results showed that HCQ inhibited NEFA secretion of T cells, but had little effect on T cell glucose uptake (Figures 5A,B). In the presence of HCQ, glucose transporter (GLUT)-1, the predominant glucose transporter in T cells, were lower compared to that of activated cells without treatment (Figure 5C). We found that HCQ treatment resulted in a significant reduction in MMP of T cells (Figure 5D). We also tested the marker genes of glycolysis and lipid metabolism [lipid synthesis and fatty acid oxidation (FAO)]. The result showed that there was no...
FIGURE 2
HCQ ameliorated hepatic pathologic damage and inflammatory infiltration of AIH mice. (A) Liver index (liver wet weight [mg]/mouse body weight [g] ×100%) and spleen index (spleen wet weight [10 mg]/mouse body weight [g] × 100%). (B) Levels of ALT and AST in serum. (C) Levels of MDA and SOD in serum. (D) Serum inflammatory cytokine concentrations. Data expressed as mean ± SD (n = 5 or 6). *p < 0.05, **p < 0.01, ***p < 0.001 relative to controls; *p < 0.05, **p < 0.01, ***p < 0.001 relative to S-100-induced AIH mice; ns, not significant.
significant effect of HCQ to glycolysis in T cells, however, HCQ substantially downregulated mRNAs encoding lipid utilization components in T cells (Srebp2 and Acaca), upregulated the fatty acid oxidation related genes expression (SirT1, SirT2, and SirT3), which might promote $T_{reg}$ differentiation (Figure 5C). Given that mitochondria were central to the metabolism of lipid (Seenappa et al., 2016), we speculated that suppression of fuel metabolism and promotion of FAO might contribute importantly to the mechanism of $T_{reg}$ development induced by HCQ.
HCQ regulated T cell function and differentiation in vitro. (A) Proliferation of T cells ($n = 5$). (B) Inflammatory cytokine concentrations in T cell culture supernatant ($n = 5$). (C) The percentage of Treg, CD4$^+$ T cells, CD8$^+$ T cells in T lymphocytes ($n = 3$). (D) Expression of Foxp3, GzmB, CTLA-4, PD-1, CD28 mRNA in T lymphocytes ($n = 5$). Data expressed as mean ± SD. $^* p < 0.05$, $^** p < 0.01$, $^*** p < 0.001$ relative to controls; $^# p < 0.05$, $^## p < 0.01$, $^### p < 0.001$ relative to ConA group; ns, not significant.
Hydroxychloroquine acted on G protein-coupled receptor kinases 2 translocation and reduced the membrane recruitment of PI3K in spleen T lymphocytes

The network pharmacology studies showed that multi-targets mechanism associated with HCQ treatment in SLE and RA. The related 3,316 proteins’ network data showed that ErbB, HIF-1, NF-κB, FoxO, chemokines, MAPK, JAK-STAT, PI3K-AKT pathways participate in the multi-targets mechanism of HCQ in RA (Lyu et al., 2020; Xie et al., 2021). The efficacy of HCQ against SLE is mainly associated with the targets of cyclin dependent kinase 2 (CDK2), estrogen receptor alpha (ESR1) and CDK1, which regulate PI3K/AKT/GSK3β as well as IFN signaling pathway (Xie et al., 2020). Another study showed that C-C chemokine receptor type 4 (CCR4), a GPCR modulated by GRK2, might an immunomodulatory target of HCQ (Beck et al., 2020). Our studies detected the protein expression of GRK2/PI3K-AKT signal in blood lymphocytes of RA patients and normal people, the results showed that the protein expression of GRK2, p-PI3K and p-AKT were increased in RA patients. The above indicators were reversed after treating by HCQ. In addition, the expression of GzmB was decreased, while the expression of Foxp3 was increased after HCQ treatment in the blood lymphocyte of RA patients (Data not shown). Based on above results and given that the GRK2/PI3K-AKT pathway contributes to the regulation of immune cell metabolism (Yang et al., 2020), we speculated that inhibiting the over activated PI3K-AKT pathway in immune cells might be effective methods for the treatment of RA and other autoimmune diseases such as AIH.
FIGURE 6
HCQ acted on GRK2 translocation and reduced the membrane recruitment of PI3K in spleen T lymphocytes (A) Proliferation of T cells after inhibiting GRK2 (n = 5). (B) Inflammatory cytokine concentrations in T cell culture supernatant after inhibiting GRK2 (n = 5). (C) The percentage of Tregs, CD4+T cells, CD8+T cells in T lymphocytes after inhibiting GRK2 (n = 3). (D) Appearance of NEFA after inhibiting GRK2 (n = 5). (E) The combination of GRK2 and PI3K or AKT (n = 3). (F) Membrane and cytoplasm protein expression of GRK2 and PI3K (n = 3). Data expressed as mean ± SD. *p < 0.05, **p < 0.01 relative to controls; *p < 0.05, **p < 0.01 relative to ConA group; ns, not significant.
We pre-treated ConA induced T cells with paroxetine (GRK2 inhibitor), which retarded the proliferation, proinflammatory factor (IFN-γ, TNF-α, and IL-12) secretion, and NEFA secretion, meanwhile, promoted Treg differentiation and anti-inflammatory factor (TGF-β) secretion. Adding HCQ did not enhance the effect of paroxetine, which suggested that the main target of HCQ might have been blocked (Figures 6A–D). Previous studies showed that the interaction of GRK2 with PI3Kγ promoted PI3K to recruit to the membrane to contribute to the signal transduction. Therefore, we detected the combination of GRK2 and PI3K by CO-IP and the membrane and cytoplasmic expression of GRK2 and PI3K by WB respectively after HCQ treatment. The results found that GRK2 and PI3K co-expression increased by ConA stimulation, and HCQ down-regulated GRK2 and PI3K interaction (Figure 6E). In addition, HCQ down-regulated GRK2 and PI3K translocation, inhibited the expression of GRK2 and PI3K in the cell membrane and increased their expression in the cytoplasm of spleen T lymphocytes (Figure 6F). Based on above results, we proposed that HCQ acted on GRK2, inhibited the interaction of GRK2-PI3K and their translocation, which reduced the recruitment of PI3K to the membrane, inhibited downstream signal transduction to disturb the function of activated T cells.
Hydroxychloroquine decreased the activation of metabolism-related PI3K-AKT-mTOR and inflammation-related JAK2-STAT3-SOCS3 pathways in the T cells and liver tissue of AIH mice

The PI3K-AKT pathway contributes to the regulation of immune cell metabolism (Yang et al., 2020), and JAK2-STAT3 pathway is widely considered to be involved in inflammation related diseases. In addition, it has been reported that HCQ may be involved in the regulation of these two pathways (Lyu et al., 2020; Li et al., 2022). Our results showed that HCQ treatment inhibited the protein and mRNA expression of GRK2 (Figures 7A,B) and reversed the activated PI3K-AKT-mTOR pathway induced by S-100 in the liver (Figure 7C). HCQ also suppressed phosphorylation of JAK2, STAT3, and increased SOCS3, the negative regulatory factor, in the liver of AIH mice (Figure 7D).

In T lymphocytes, HCQ suppressed phosphorylation of PI3K (Tyr458), AKT (Ser473) and mTOR (Ser2448), without affecting their mRNA and total protein expressions (Figures 8A,B). Following the PI3K agonist (740 Y-P, 20 μg/ml, 12 h)-mediated activation of PI3K, T cells exhibited proliferation, with enhanced levels of Treg, elevated secretion of IFN-γ, TNF-α and IL-12, and lowered expression TGF-β1. HCQ, on the other hand, eliminated the impact induced by 740 Y-P (Figures 8C–E). In addition, PI3K inhibitors (LY294002, 10 μM, 12 h) inhibited T cell proliferation and promote Treg differentiation compared with ConA group, adding HCQ did not enhance the effect of LY294002 (Figures 8F,G). GRK2 inhibitor also appeared to inhibit pathway activation in the present of 740 Y-P. And the effect of GRK2 inhibitor was weak after LY294002 treatment in T cells. The results suggested that HCQ effect on PI3K-AKT signal might be mediated by GRK2 (Figures 8C–G).

AMPK is a highly conserved serine/threonine protein kinase, which was the inhibition of mTOR in the lipid oxidation of Tregs (Michalek et al., 2011). Our results showed that HCQ significantly increased the expression of p-AMPK compared with ConA pretreatment T lymphocyte, which indicated there was the cross-talk of GRK2/PI3K-AKT and AMPK for HCQ effect in AIH (Figure 9A). We also examined the protein expressions of JAK2-STAT3-SOCS3 pathway in ConA-treated spleen T lymphocytes. The results showed that HCQ suppressed the phosphorylation of JAK2 (Tyr1007), STAT3 (Tyr705), without affecting their total protein expressions, and increased the expressions of SOCS3 (Figure 9B), which appears to indicate the mechanism of anti-inflammatory effect of HCQ. HCQ might have a potential inhibition to signals cross-talk and modulation for immunologic homeostasis in AIH. The multi-targets mechanism of HCQ in the treatment of AIH was shown in Figure 10.
AIH is an autoimmune liver disease, which progress to cirrhosis and liver failure with untreated (Fan et al., 2019). Although the pathogenesis of AIH is not fully understood, it is generally recognized that immunoregulatory networks play a crucial role (Liberal et al., 2015; Zhu et al., 2021). In a healthy population, circulating autoreactive T cells are suppressed by peripheral tolerance mechanisms to limit autoimmune tissue damage, among which Treg-exerted immune suppression plays a key role. Patients with AIH display a reduced Treg frequency or function compared to healthy subjects (Ferri et al., 2010; Sakaguchi et al., 2010; Liberal et al., 2017; Zhu et al., 2021). In addition, several cytokines are implicated in the pathogenesis of ConA-induced hepatitis, of which TNF-α, IFN-γ, TGF-β are the most important (Sass et al., 2002; Wang et al., 2017b; Zhang et al., 2020). HCQ, as an antimalarial agent, has been used for the treatment of immunological diseases: RA and SLE(3). Recently, HCQ exhibited the probable protective effect in liver injury among the COVID-19 patients (Sima, 2021). It was reported that HCQ restored the Th17/Treg balance in MRL/lpr (mouse model that develops SLE) mice, which resulted in a significant decrease in the expression of IL-17 in Th17 cells and a considerable increase in Foxp3 and TGF-β levels (An et al., 2017).

Based on that observation, we established a model of chronic AIH induced by S-100 combined with CFA, which relatively resembled human AIH. Meanwhile, in vitro, we tested the effect of HCQ in ConA activated murine spleen T-lymphocyte. The results showed that HCQ ameliorated hepatic pathologic damage and inflammatory infiltration. Besides, HCQ promoted Treg differentiation and the mRNA expression of Foxp3, and retarded CD8+T cells differentiation both in vivo and in vitro. The critical role of HCQ was the inhibition of the metabolism-related GRK2/PI3K-AKT pathway and the inflammation-related JAK2-STAT3 pathway in T cells. The sequence appears to be: (1) HCQ acted on GRK2, inhibited the translocation of GRK2, decreased the interaction between GRK2 and PI3K, and reduced the recruitment of PI3K to the membrane (2) the blockade of the PI3K-AKT pathway by HCQ accompanied with the inhibition of the inflammation-related JAK2-STAT3 pathway in T cells (3) the diminished lipid metabolism of T cells impaired its function, followed by increasing Treg differentiation,
disrupting CD8+ T cell responses, upregulating co-inhibition signal and anti-inflammatory cytokine TGF-β1 expression, and decreasing proinflammatory cytokines released from T cells; and consequently, HCQ inhibited inflammatory cell infiltration and immune responses against liver tissue, and thus diminished autoimmune liver injury (Figure 8C).

Activation of immune cells is associated with a dramatic increase in metabolism. Glucose and lipid metabolism is related to differentiation of Treg (Cluxton et al., 2019; Shan et al., 2020). Besides, it has also been reported that glucose and lipid metabolism is also related to other immune cells, such as CD4+ T, CD8+ T, DC, natural killer T (NKT) cells (Palmer et al., 2016; Giovanelli et al., 2019). Our results showed that HCQ treatment decreased NEFA secretion while increased fatty acid oxidation in T lymphocytes, but HCQ had little effect on T cell glucose uptake and glycolysis. Suppressed lipid metabolism predominantly rather than glycolysis of T cell may contribute to immunosuppressive effects of HCQ in AIH. PI3K-AKT-mTOR has been proved to be a classical pathway linking the activation of the insulin receptor to regulate glucose metabolism, which was also shown to be responsible for the accumulation of intracellular lipids by regulating fatty acid synthesis (Chen et al., 2019; Sun et al., 2022). In addition, the activation of PI3K-AKT-mTOR is accompanied with the inhibition of differentiation and function of Treg (Gao et al., 2020). mTOR is the downstream protein of PI3K-AKT, inhibition of mTOR favors naive CD4+ T-cell conversion to iTreg, and mTOR knockout or treatment with the mTOR-specific inhibitor, rapamycin, favors Treg.
Activated GPCRs, and it also interact with non-GPCRs such as GRK2 specifically recognize and phosphorylate agonist-activated GPCRs, and it also interact with non-GPCRs such as PI3K, AKT (Ribas et al., 2007). The interaction of PI3K with GRK2 mediated PI3K recruitment to the membrane after agonist stimulation. In addition, the inhibition of GRK2-G_{βγ} complex membrane translocation in the fibroblast-like synovial cells might reduce the membrane expression and activation of PI3K, which mediated the inhibition of AKT activity (Wang et al., 2020). AKT had also been reported to associate with GRK2 directly through the GRK2 C-terminus. GRK2 might inhibit agonist-dependent AKT phosphorylation, although the exact mechanisms were not well established (Ribas et al., 2007). The inhibition of GRK2 rescued AKT activity to promote T_{reg} differentiation in mice with diabetic cardiomyopathy (Han et al., 2020). There was also reported that GRK2 knock down prevented AKT activation in cardiac myocytes but the interaction between GRK2 and AKT was not clear (Penela et al., 2010; Pathania et al., 2019). In addition, increasing PI3K produced polyphosphoinositides that interacted with protein kinases, leading to activation of the kinase AKT (Saltiel, 2021). HCQ has been reported to inhibit PI3K-AKT pathway in renal interstitial fibrosis (Li et al., 2022). Therefore, we tested whether HCQ inhibited the PI3K-AKT pathway by regulating the interaction between GRK2 and PI3K or AKT in T cells of AIH. Our results found that GRK2 and PI3K were co expressed in T lymphocytes. HCQ down-regulated GRK2-PI3K interaction to reduce the complex translocation, and suppressed the phosphorylation of PI3K, AKT, and mTOR. The direct interaction of GRK2 with AKT was undefined. After blocking GRK2, the effect of HCQ was diminished.

In conclusion, HCQ exhibits specific and potent therapeutic effects on AIH and attendant liver injury. The critical role appears to be: HCQ acted on GRK2 translocation mediated PI3K-AKT-mTOR signal inhibition in spleen T lymphocyte, thereby modulating lipid metabolism in T lymphocytes and promoting T_{reg} activity. And HCQ might block the cross-talk of JAK2-STAT3-SOCS3, PI3K-AKT-mTOR and AMPK signals.

Data availability statement

Data supporting the results of this study can be obtained from the corresponding author upon reasonable request.

Ethics statement

The animal study was reviewed and approved by the Laboratory Animal Ethics Committee of Anhui Medical University.

Author contributions

T-TL participated in the study design, experimental validation, analyzed the data, material support, coordination, drafted the manuscript, and supervision of the study. CJ designed the experimental validation, performed experiments, analyzed the data, and drafted the manuscript. B-BG performed technical guidance. All authors read and approved the final manuscript.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fphar.2022.972397/full#supplementary-material
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