Resveratrol protects against L-arginine-induced acute necrotizing pancreatitis in mice by enhancing SIRT1-mediated deacetylation of p53 and heat shock factor 1

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Abstract. Acute necrotizing pancreatitis (ANP) is a common severe critical illness with a high mortality rate. Resveratrol, a polyphenol compound derived from various plants such as grape skin, peanut, berry and veratrum, exhibits multiple biological activities, especially potent anti-inflammatory activity, but its effect on ANP has not yet been fully elucidated. The present study aimed to investigate the effects of resveratrol on L-arginine-induced ANP and the possible mechanisms. A mouse model of ANP was established by 2 hourly intraperitoneal injections of 8% L-arginine (4 g/kg). Then the mice were treated by intragastric administration of resveratrol (80 mg/kg) every 12 h immediately after the second injection of L-arginine. Mice with ANP showed increased apoptosis of pancreatic acinar cells, pancreatic myeloperoxidase activity, serum lactate dehydrogenase activity, amylase, tumor necrosis factor-α (TNF-α), interleukin-6 (IL-6) levels as well as decreased serum IL-10 level, pancreatic expression of heat shock factor 1 (HSF1), sirtuin 1 (SIRT1) and p53, but the ratio of acetylated HSF1 and p53 was markedly increased. Resveratrol enhanced the survival rate of mice with ANP from 47.8 to 71.4% and obviously restored the changes in mice with ANP as mentioned above. Additionally, interactions between SIRT1 and p53 and between SIRT1 and HSF1 in the pancreas of the mice were confirmed by co-immunoprecipitation. These data suggest that resveratrol protects against L-arginine-induced ANP, which may be related to the enhancement of SIRT1-mediated deacetylation of p53 and HSF1.

Introduction

Acute pancreatitis (AP) is a trypsin-activated sudden inflammatory response in the pancreas and autodigestion of the pancreas caused by various etiological factors such as gallstones, craptulence, hyperlipemia, virus, sepsis and shock (1,2). Currently, AP is divided the mild AP, moderately severe AP (MSAP) and severe AP (SAP) in the clinic according to disease severity and prognosis (3). SAP is also called acute necrotizing pancreatitis (ANP) and generally manifests as diffuse pancreatic necrosis and hemorrhage, adiponecrosis, neutrophil and monocyte infiltration, necrosis and apoptosis of pancreatic acinar cells. It can rapidly develop into systemic inflammation and even multiple organ dysfunction syndrome (MODS), accompanied by an unacceptably high mortality rate. However, the therapy of ANP is now limited to symptomatic and supportive treatments such as analgesia, spasmolysis, improvement in microcirculation, antishock, anti-inflammation and antiemetic. Yet up to 30% of ANP patients die from various complications (4). Therefore, searching for effective therapeutic targets and drugs are of great importance.

Today the pathogenesis of ANP has not been fully elucidated. It has been confirmed that the autodigestion of the pancreas by intrapancreatic trypsin, excessive inflammation, necrosis and apoptosis of pancreatic acinar cells may all be involved in this process (5-7). However, the use of trypsin inhibitors and anti-inflammatory therapy directed at a certain inflammatory factor are usually noneffective in the treatment of ANP. The occurrence and development of ANP can be described as progression from an initial injury of pancreatic acinar cells to local and systemic inflammation (5). Uncontrolled release of a large amount of various inflammatory factors including tumor necrosis factor-α (TNF-α), interleukin-6 (IL-6), IL-1β and IL-10, are not only the initial factors of ANP, but also promote the necrosis and apoptosis of pancreatic acinar cells and eventually aggravate the impairment of the pancreas in ANP (6). Hence, the decrease in the uncontrolled release of various inflammatory factors may be helpful to improve the prognosis of AP.

Resveratrol, a polyphenol compound derived from various plants such as grape skin, peanut, berry and veratrum has multiple biological activities, including potent anti-inflammatory
activity, antioxidant, anti-aging, insulin sensitization and cardiac protection. It has been shown that resveratrol has certain protective effect on cerulein-induced acute edema pancreatitis and 4% sodium taurocholate-induced SAP as well as related complications in rats, but its effect on L-arginine-induced ANP remains unknown to date (8,9). Resveratrol has been found to be an activator of sirtuin 1 (SIRT1). SIRT1 is one of the seven mammalian orthologs of the yeast protein silent information regulator 2 (Sir2), a conserved NAD-dependent protein deacetylase that plays an important role in the cell proliferation, differentiation, apoptosis, metabolism and inflammatory response through deacetylating certain lysine residues in a variety of downstream substrates including histone H3, p53, heat shock factor 1 (HSF1), and further enhancing their transcription activity (10-12). However, the role of SIRT1 in the development of ANP is largely unknown.

In this study, for the first time, to the best of our knowledge, the effect of resveratrol on L-arginine-induce ANP in mice was investigated, and whether the activation of SIRT1 and its deacetylation of downstream substrates are involved in this process was determined.

Materials and methods

Animals. Male Balb/c mice weighed 20-25 g were purchased from the Center of Experimental Animals in Central South University of China. All mice were kept on a 12-h light/dark cycle at a room temperature of 23±2˚C and a relative humidity of 50±5% and housed individually with free access to food and water throughout the experiment. Mice were randomly assigned to the experimental procedures described as follows. Animal use procedures were approved by the Animal Welfare Ethics Committee of Central South University.

Establishment of a mouse model of acute pancreatitis and treatment. A mouse model of acute pancreatitis was established by 2 hourly intraperitoneal injections of 8% L-arginine hydrochloride (pH 7.0, 4 g/kg body weight). The mice were randomly divided into three groups: control group (Con), L-arginine exposure group (AP), L-arginine and resveratrol treatment group (AP+RSV). Mice in the AP+RSV group were treated with intragastric administration of 80 mg/kg resveratrol every 12 h immediately after the second injection of L-arginine. Mice in the Con group and AP group were treated with the same volume of vehicle. Resveratrol was dissolved in 0.5% sodium carboxymethyl cellulose away from light and was used immediately after it was ready. Before administration of L-arginine, the mice were deprived of food and received only water for 10-12 h. After the first injection of L-arginine and clotted at room temperature of 48 and 72 h immediately after the second injection of L-arginine. Mice were given food and water ad libitum immediately and then sacrificed at 48 and 72 h after the second injection of L-arginine.

Reagents. The antibodies to HSF1 (1:2,000 dilution; 12972S), acetylated-lysine (1:1,000 dilution; 9441S), SIRT1 (1:2,000 dilution; 9475S) and p53 (1:1,000 dilution; 2524S) were purchased from Cell Signaling Technology, Inc. (Danvers, MA, USA). The antibody to glyceraldehyde 3-phosphate dehydrogenase (GAPDH; 1:3,000 dilution; G9545), resveratrol, L-arginine hydrochloride, sodium carboxymethyl cellulose, protease inhibitor cocktail were purchased from Sigma-Adrich (St. Louis, MO, USA). Mouse TNF-α, IL-6 and IL-10 enzyme-linked immunosorbent assay (ELISA) kits and IgG horseradish peroxidase-conjugated secondary antibody (1:10,000 dilution; BA1050, BA1054) were purchased from Boster Biological Technology, Ltd. (Wuhan, China). Lactate dehydrogenase and myeloperoxidase (MPO) assay kit were purchased from Nanjing Jiancheng Biology Engineering Institute. In situ cell death detection kit was purchased from Roche Applied Sciences (Mannheim, Germany). PureProteome™ Protein A and Protein G magnetic beads were purchased from Merck Millipore (Billerica, MA, USA). Other reagents were analytically pure.

Histological examination. Pancreatic tissues were rinsed and fixed in 4% paraformaldehyde (PFA) at 4˚C overnight. The PFA-treated pancreatic tissues were then processed with sequential clearing and dehydrating steps, and embedded in paraffin blocks. Samples were sectioned into 5-µm slices and subjected to standard hematoxylin and eosin (H&E) staining for the evaluation of pancreatic tissue injury. Images were taken under an optical microscope (Olympus, Tokyo, Japan). The severity of pancreatic tissue lesions was evaluated and scored by two experienced pathologists in a random and double-blinded manner according to a modified Grewal method as follows (13): edema was scored from 0-4 points (0, none; 1, patchy interlobular septum; 2, diffuse interlobular septum; 3, diffuse interlobular and intraacinar septum; 4, diffuse intercellular space); leukocytic infiltration was scored from 0-4 points (0.5, per 5 leukocytes, 4, >30 leukocytes); the percentage of acinar cell necrosis was evaluated by ImageJ software and scored from 0-4 points (0, none; 1, 1-10% necrosis area; 2, 11-20% necrosis area; 3, 21-30% necrosis area; 4, >30% necrosis area) and hemorrhage was scored from 0-1 points (0, none; 1, hemorrhage).

Measurement of serological index. Blood was obtained by direct cardiac puncture under deep anesthesia at 48 h and 72 h after the second injection of L-arginine and clotted at room temperature for 4 h. To obtain serum samples, the blood was centrifuged at 1,500 x g for 10 min. The serum amylase level was further detected at the clinical laboratory of Xiangya Hospital. ELISA assays were performed to detect the TNF-α, IL-1β, IL-6 and IL-10 contents in the serum according to the manufacturer’s protocol. Serum lactate dehydrogenase (LDH) activity was measured by a colorimetric method according to the manufacturer’s protocol.

MPO activity assay. The pancreatic MPO activity was assessed by an MPO colorimetric activity assay kit according to the manufacturer’s protocol and presented as units per milligram of pancreatic tissue.

Terminal deoxynucleotidyltransferase-mediated dUTP nick end labeling (TUNEL) assay. TUNEL staining was performed to detect the apoptosis of pancreatic acinar cells through in situ cell apoptosis detection kits according to the manufacturer’s instructions. Pancreatic tissues were rinsed and fixed in 4% PFA at 4˚C overnight followed by paraffin embedding. Paraffin-embedded sections of the samples were deparaffinated and hydrated using graded ethanol, and then incubated.
in 20 g/ml protease K at 37°C for 30 min. After being washed with phosphate-buffered saline (PBS) for 5 min three times, the samples were incubated with 0.1% Triton X-100 for 20 min at room temperature and then washed with PBS for 5 min three times. Terminal deoxynucleotidyl transferase (TdT) and dUTP were mixed at a 2.29 ratio and then added to the sample and incubated at 37°C for 2 h. After washing with PBS for 5 min three times, the sections were immersed into 3% H2O2 prepared by methanol for 15 min away from light, and then covered with converter-POD at 37°C for 30 min in a humidified box. After rinsing, the samples were incubated with DAB substrate kit under a light microscope. The slides were lightly counterstained with haematoxylin and then dehydrated and mounted. For each pancreas specimen, tissue sections were examined under a light microscope at x100 and x200 magnifications. Ten random fields per section were counted and analyzed by Image Pro-Plus 7.0 software and the percentage of apoptotic cells was finally calculated as the apoptotic index.

Quantitative (real-time) polymerase chain reaction. Total RNA of the pancreatic tissues was extracted by TRizol and reverse transcribed to cDNA with Primerscript™ RT reagent kit with gDNA Eraser according to the manufacturer’s instructions (Takara Shuzo Co., Kyoto, Japan). The concentration and purity of total RNA were determined by measuring the OD260 and OD260/OD280 ratio, respectively. The mRNAs of TNF-α, IL-6 and IL-10 were measured by SYBR Premix Ex Taq™ (Takara Shuzo Co.) through an ABI 7500 real-time PCR system (Life Technology Corp., Carsbad, CA, USA). Each cDNA sample was analyzed in triplicate. The relative quantitation of mRNA was analyzed using the equation: ratio=2^{ΔΔCt} and normalized to GAPDH. The following primers synthesized by Sangon Biotech Co., Ltd. (Shanghai, China) were used: TNF-α forward, 5'-ACCCCTACACTCTACAAA CCA-3' and reverse, 5'-ACAAGGTACAACCCATGCGG-3'; IL-6 forward, 5'-CCCCAATTTCATTCAATCTCC-3' and reverse, 5'-CGCATTAGTTTGGCGGATA-3'; IL-10 forward, 5'-GCTTCTTGACATCCAAGGCC-3' and reverse, 5'-CTGCG TGAATCCATGCAGT-3'; GAPDH forward, 5'-GGGTC CAGCTTTAGGTTCAT-3' and reverse, 5'-TACGGCCAAATC CGTTCACA-3'.

Western blotting. Mouse pancreatic tissue (100 mg) was homogenized in 0.5 ml lysate buffer containing 20 mM Tris-HCl (pH 7.4), 5 mM EDTA, 100 mM Na3P04, 2 mM Na2VO4, 100 mM NaF and 1% Nonidet P-40 and protease inhibitors were added at a fixed volume ratio [lysate buffer:phenylmethylsulfonyl fluoride (PMSF):protease inhibitor cocktail, 100:1:1]. Homogenized samples were centrifuged at 4°C, at 14,000 rpm for 15 min. The concentrations of protein from the supernatant were determined using BCA protein assay reagent. Protein (25 μg) was loaded for 10% sodium dodecyl sulfate-polyacrylamide-gel electrophoresis (SDS-PAGE) and then transferred to polyvinylidene difluoride (PVDF) membranes (Merck Millipore). After blocking with 5% bovine serum albumin, immunoblotting was carried out with various primary antibodies at 4°C overnight or at 25°C for 2 h (anti-GAPDH antibody was used as an internal control). After incubation with IgG horseradish peroxidase-conjugated secondary antibody at room temperature for 1 h, the membranes were washed three times successively. Then the specific proteins were detected by enhanced chemiluminescence (Invitrogen, Carlsbad, CA, USA) according to the manufacturer’s instructions. The relative band intensity was quantified by Quantity One software (Bio-Rad Laboratories, Inc., Hercules, CA, USA).

Co-immunoprecipitation. Fresh mouse pancreatic tissue was rapidly homogenized in lysate buffer as mentioned above. The protein concentration was then adjusted to 3 μg/μl and 800 μg total protein and 4 μg capture antibody (SIRT1, HSF1 or p53) were incubated at 2-8°C with continuous mixing overnight. Fifty microliters of protein A and protein G magnetic beads was resuspended and washed with 500 μl PBS containing 0.1% Tween-20. The reaction containing the pre-formed antibody-antigen complex mentioned above was subsequently added to the magnetic beads and incubated for 8 h at 2-8°C with continuous mixing to capture the immune complex. The magnetic beads were collected by putting the tube into the magnetic stand, allowing the beads to migrate to the magnet, and then removing the sample with a pipette. After washing the beads with 500 μl PBS containing 0.1% Tween-20 for 3 times, 60 μl of 5X sample loading buffer suitable for electrophoresis was added and mixed to resuspend the beads. The beads were heated at 70-90°C for 10 min and stored at -80°C until being used for western blotting.

Acetylation assay. Fresh mouse pancreatic tissue was subjected to immunoprecipitation with mouse HSF1 and p53 antibodies, while acetylated HSF1 and p53 were detected by western blotting with an acetylated lysine antibody. Then the acetylation of HSF1 and p53 was evaluated by calculating the ratio of acetylated HSF1 and p53 to total HSF1 and p53, respectively.

Statistical analysis. All data were analyzed by SPSS 18.0 software. Measurement data are shown as mean ± SEM of three different experiments and analyzed by unpaired two-tailed
student's t-tests. Kaplan-Meier analysis was performed to compare the differences in survival rate between different groups. P<0.05 was considered statistically significant.

Results

Effect of resveratrol on the survival of mice with L-arginine-induced ANP. To assess the effect of resveratrol on the prognosis of ANP, a mouse model of ANP was established by intraperitoneal injection of L-arginine as reported previously (14,15).

No mouse died in the control group; the 7-day survival rate was 100%, while the 7-day survival rate of mice with ANP was obviously decreased to 47.8%, which was significantly lower than that of the control group (p<0.05). In contrast, resveratrol enhanced the 7-day survival rate of mice with ANP to 71.4% (p<0.05) (Fig. 1).

Effect of resveratrol on the pancreatic morphology of mice with L-arginine-induced ANP. The mice treated with 2 hourly intraperitoneal injections of 8% L-arginine hydrochloride...
developed ANP at 48 h after the second injection of L-arginine. As shown in Fig. 2A, the pancreas of the mice in the Con group was colored grey-white with normal structure, while the pancreas of the mice with ANP was colored gray-red with obvious roughness, necrosis and diffuse hemorrhage at 48 h and 72 h after the second injection of L-arginine. The pancreas in the resveratrol-treated mice with ANP showed a lighter color than that of the mice with ANP and minimal necrosis. The structure of the pancreas was close to that of the mice in the Con group.

H&E staining showed wider pancreatic interlobular and acini septum, leukocyte infiltration, and up to a 20% necrotic area as well as hemorrhage in the mice with ANP at 48 h after the second injection of L-arginine, while pancreatic interlobular and acini septum, and intercellular space were all widened. A large amount of leukocyte infiltration and a >30% necrotic area as well as hemorrhage were observed in the mice with ANP at 72 h after the second injection of L-arginine. However, the resveratrol-treated mice with ANP showed narrower pancreatic interlobular and acini septum, less leukocyte infiltration and a <10% necrotic area (Fig. 2B). According to the modified Grewal method, the pancreatic damage was scored. The pathological scores of AP mice at 48 h and 72 h after the second injection of L-arginine were both significantly higher than those of the control group, while resveratrol obviously decreased the pathological scores of the mice with ANP (p<0.05) (Fig. 2C).

Effect of resveratrol on the serum amylase level, LDH activity, and pancreatic MPO activity of mice with L-arginine-induced ANP. As the serum amylase level is the most commonly used biochemical marker of AP, the serum LDH activity and pancreatic MPO activity are related to neutrophil infiltration. Thus, in this study, the serum amylase level, LDH activity and pancreatic MPO activity were detected at 48 and 72 h after the second injection of L-arginine. The results showed that the serum amylase level, LDH activity and pancreatic MPO activity were significantly higher in the mice with ANP compared with those parameters of the mice in the Con group (p<0.05). Resveratrol obviously decreased the serum amylase level, LDH activity and pancreatic MPO activity of the mice with ANP (p<0.05) (Fig. 3).

Effect of resveratrol on pancreatic IL-6, TNF-α, IL-10 mRNA expression and serum IL-6, TNF-α, IL-10 levels of the mice with L-arginine-induced ANP. The development of ANP is considered as progression from the initial injury of exocrine pancreas to local and systemic inflammatory response (5). Therefore, the pancreatic IL-6, TNF-α, IL-10 mRNA expression and serum IL-6, TNF-α, IL-10 levels were detected. It was found that the pancreatic IL-6, TNF-α mRNA expression and serum IL-6, TNF-α levels of mice with ANP were significantly increased compared with the mice in the Con group (p<0.05), while the pancreatic IL-10 mRNA expression and serum IL-10 level were obviously decreased (p<0.05). Resveratrol did not only markedly decrease the pancreatic IL-6, TNF-α mRNA expression and serum IL-6, TNF-α levels of mice with ANP but also enhanced the pancreatic IL-10 mRNA expression and serum IL-10 level (p<0.05) (Fig. 5).

Effect of resveratrol on the SIRT1 protein expression and activity in the pancreas of mice with L-arginine-induced ANP. Resveratrol has been confirmed as a classic SIRT1 activator, but whether SIRT1 is involved in the pathogenesis of L-arginine-induced acute pancreatitis is still unknown. The expression and activity were further detected. It was shown that the expression of SIRT1 in the pancreas of mice with ANP was significantly decreased to ~40% of that in the mice in the control group at 48 and 72 h after the second injection of L-arginine (p<0.05), but obviously increased SIRT1 expression was detected in the resveratrol-treated mice with ANP (p<0.05) (Fig. 6).

p53 is a pro-apoptotic protein, the deacetylation of which indirectly reflects the activity of SIRT1 (5,16,17). Firstly, the expression of p53 was detected. p53 protein expression was significantly downregulated in the pancreas of the mice with ANP at 48 h and 72 h after the second injection of L-arginine (p<0.05), while resveratrol obviously upregulated the p53 protein expression of the mice with ANP (p<0.05). Co-immunoprecipitation assay showed that p53 was
precipitated by the SIRT1 antibody and further acetylation assay showed that the ratio of acetylated p53 was markedly higher in the pancreas of the mice with ANP at 48 h and 72 h after the second injection of L-arginine (p<0.05), but resveratrol markedly attenuated the acetylation of p53 in the mice with ANP (p<0.05) (Fig. 7).

**Effect of resveratrol on the expression and acetylation of HSF1 in the pancreas of mice with L-arginine-induced ANP.** HSF1 is a crucial transcription factor in heat shock response and also plays an important role in the inflammatory response and cell apoptosis. The activity of HSF1 is closely related to its acetylation. To investigate the role of HSF1 in the L-arginine-induced ANP, co-immunoprecipitation assay was firstly performed to detect the interaction between HSF1 and SIRT1. HSF1 was precipitated by the SIRT1 antibody in the different groups, although the total expression of HSF1 was significantly decreased in the pancreas of the mice with ANP at 48 and 72 h after the second injection of L-arginine (p<0.05). Acetylation assay showed that the ratio of acetylated

Figure 4. Effect of resveratrol on the apoptosis of pancreatic acinar cells in mice with L-arginine-induced acute necrotizing pancreatitis (ANP). Apoptotic pancreatic acinar cells were detected by terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL) staining. TUNEL-positive cells are indicated by brown staining. Representative photomicrographs demonstrating TUNEL staining of pancreatic sections (A) and quantitative analysis of TUNEL-positive cells as the apoptotic index (B) are shown. *p<0.05 vs. Con; †p<0.05 vs. acute pancreatitis (AP) (n=5). Con, control group; AP, L-arginine exposure group; AP+RSV, L-arginine and resveratrol treatment group.
HSF1 was significantly upregulated (p<0.05). However, resveratrol obviously enhanced the total expression of HSF1 but decreased the acetylation of HSF1 in the pancreas of the mice with ANP (p<0.05) (Fig. 8).

Discussion

AP, a common inflammatory disorder of the pancreas, has become the leading cause of admission to intensive care units (ICUs) worldwide (18). It has been confirmed that AP is initiated by the activation of various enzymes in pancreatic acinar cells, which leads to the injury of acinar cells, followed by the release of pro-inflammatory factors such as TNF-α, IL-1β and IL-6 (19). The chemokines further released from the injured acinar cells then promote inflammatory cells including neutrophils and macrophages to accumulate in the pancreas, which results in exacerbating the local inflammatory response in the pancreas and progressing rapidly into a systemic inflammatory response (5). Under this condition, the excessive release of various inflammatory factors increases the microvessel density, induces thrombosis and hemorrhage and finally increases the apoptosis and necrosis of acinar cells. In
addition, the gut barrier dysfunction is usually found in AP, which can result in bacterial translocation, further activate the macrophages to release a large quantity of inflammatory factors and induce a secondary attack to the pancreas (20,21). Hence, controlling the rapidly progressive inflammatory response and inhibiting the injury of pancreatic acinar cells may be beneficial to improve the prognosis of ANP.

Resveratrol is a natural polyphenol compound mainly found in various plants and red wine. It is well known for its anti-inflammatory and antioxidant effects. It has been confirmed that resveratrol has protective effects on sodium taurocholate-induced ANP and cerulean-induced acute edema pancreatitis. L-arginine-induced experimental model of ANP was firstly reported by Tani et al (22), since L-arginine selectively destroys pancreatic acinar cells by inducing amino acid imbalance, decreasing the synthesis of polyamine, nucleic acid and proteinase and resulting in excessive activation of zymogen. In addition, L-arginine could induce the expression of different types of inflammatory factors and apoptosis-related genes and proteins in the pancreatic acinar cells. Compared with other invasive experimental models of ANP, the L-arginine-induced experimental model of ANP has the advantages of non-invasive, easier operation, higher success rate and lower cost, although the death rate is high. It is commonly used as an investigative tool for screening and developing effective therapies for ANP. In this study, the 7-day survival rate of the L-arginine-induced mice with ANP was 47.8%, while administration of 20 and 40 mg/kg resveratrol every 12 h immediately after the second injection of L-arginine had no significant effect on the survival of the mice with ANP (data not shown). However, administration of 80 mg/kg resveratrol significantly enhanced the 7-day survival rate of the mice with ANP to 71.4%. Further studies showed that the L-arginine-induced ANP model was successfully established at 48 h after the second injection of L-arginine, which was mainly manifested as pancreatic and systemic inflammation, apoptosis and necrosis of pancreatic acinar cells. Administration of 80 mg/kg resveratrol every 12 h immediately after the second injection of L-arginine markedly ameliorated L-arginine-induced pancreatic inflammation and injury as well as systemic inflammation. These results proved that resveratrol could protect L-arginine-induced ANP through alleviating the pancreatic inflammatory response.

In recent years, accumulating evidence suggests that the biological activities of resveratrol are closely related to the activation of SIRT1. SIRT1 has been proven to be involved in a variety of cellular functions, such as proliferation, differentiation, apoptosis, aging and metabolism. It also participates in the regulation of inflammation, but its role in ANP is poorly understood to date. Our data showed that the pancreatic expression of SIRT1 in L-arginine-induced mice with ANP was significantly decreased compared with that in the normal mice, which was reversed by resveratrol treatment. The activation of SIRT1 enhances the transcriptional activities...
of its downstream targets mainly through deacetylating the lysine residues at certain sites. p53 has been identified as a downstream target of SIRT1 and the deacetylation level of p53 was proven to be able to indirectly reflect the activity of SIRT1. In this study, the results showed that p53 interacted with SIRT1 in the different experimental groups, the p53 expression was downregulated while the ratio of acetylated p53 was upregulated in the L-arginine-induced mice with ANP, suggesting that the SIRT1 activity was significantly decreased in the mice with ANP, while resveratrol could enhance the SIRT1 activity and subsequent promotion of pancreatic acinar cell apoptosis mediated by p53 deacetylation, while resveratrol eventually decreased the apoptosis of pancreatic acinar cells through other anti-apoptotic mechanisms.

Figure 8. Effect of resveratrol on the expression and acetylation of heat shock factor 1 (HSF1) in the pancreas of mice with L-arginine-induced acute necrotizing pancreatitis (ANP). Interaction between sirtuin 1 (SIRT1) and HSF1 in the pancreas was detected by co-immunoprecipitation at 48 h (A) and 72 h (E) after the second injection of L-arginine. Expression of HSF1 and acetylated HSF1 in the pancreas was detected by immunoblotting at 48 h (B-D) and 72 h (F-H) after the second injection of L-arginine. *p<0.05 vs. Con; #p<0.05 vs. acute pancreatitis (AP) (n=3). Con, control group; AP, L-arginine exposure group; AP+RSV, L-arginine and resveratrol treatment group.

Inflammatory response persists in the occurrence and development of ANP which is described as progression from an initial injury of pancreatic acinar cells to local and systemic inflammation. HSF1, an important heat shock
transcription factor, is widely expressed in various types of cells and plays an important role in inflammation by regulating the transcription of inflammatory factors (29-31). The acetylation of HSF1 decreases its DNA binding activity, and thus promotes the transcription of pro-inflammatory factors and inhibits the transcription of anti-inflammatory factors. In our previous study, HSF1-knockout mice developed more severe acute edema pancreatitis induced by cerulein than the wild-type mice. However, the role of HSF1 in ANP is poorly understood. Our results showed that HSF1 expression was significantly decreased in the pancreatic tissues of mice with ANP, indicating that HSF1 is involved in the pathogenesis of ANP. It is particularly noteworthy that HSF1 is a deacetylase substrate for SIRT1 (10,32). Ghemrawi et al found that a decrease in the cellular availability of B12 led to ER stress activation mediated by decreased SIRT1 expression, which in turn led to both lower HSF1 expression and HSF1 hyper-acetylation (10). Similarly, in our study, although the total HSF1 expression was decreased in the pancreatic tissue of the L-arginine-induced mice with ANP, the ratio of acetylated HSF1 was sharply increased up to 6- to 8-fold of the normal mice, which may consequently increase the transcription of pro-inflammatory factors (TNF-α and IL-6) but decrease the transcription of anti-inflammatory factors (IL-10). However, resveratrol did not only markedly enhance the total expression of HSF1, but also diminished acetylated HSF1. These results suggest that resveratrol enhances the deacetylation of HSF1 in the pancreas of ANP, and subsequently exhibits its potent anti-inflammatory effect and eventually ameliorates ANP by mediating the transcription of inflammatory factors.

In addition to the deacetylation of p53 and HSF1, other intracellular target proteins and signaling pathways may also be involved in the protective effect of SIRT1 activation on ANP. For instance, nuclear factor-κB (NF-κB), forkhead box protein (FoxO1, FoxO3 and FoxO4) are also the deacetylation targets of SIRT1 (33,34). Sirt1 inhibits apoptosis and inflammation via deacetylating these proteins and thereby modulating their activities, which may also contribute to the protective effect of resveratrol on ANP. Mammalian target of rapamycin (mTOR), an important target of SIRT1, inhibits autophagy and has been implicated in the development of pancreatitis (35,36). SIRT1 influences different pathophysiological processes including metabolism, cell apoptosis and aging by downregulating mTOR (37-39). Interestingly, resveratrol exhibited a protective effect on various diseases via the upregulation of mTOR and autophagy. Therefore, the protective effect of resveratrol on ANP may also be attributed to the SIRT1 mTOR axis-enhanced autophagy in the pancreatic acinar cells. Furthermore, the Toll-like receptor 4 (TLR4)/NF-κB pathway mediates the production of various pro-inflammatory factors and has been found to play important roles in the development of pancreatitis, whereas resveratrol was found to relieve the inflammatory injury by inhibiting the TLR4/NF-κB pathway (45,46). Hence, the inhibition of the TLR4/NF-κB pathway may also contribute to the protective effect of resveratrol on ANP.

In conclusion, to the best of our knowledge, our data demonstrated for the first time that resveratrol effectively improved the survival, relieved the inflammatory response and decreased the acinar necrosis and apoptosis in L-arginine-induced ANP in mice, which may be related to the enhancement of SIRT1-mediated deacetylation of p53 and HSF1. Notably, the increase in SIRT1-mediated deacetylation of p53 can promote the apoptosis of pancreatic acinar cells, which might further decrease the necrosis of pancreatic acinar cells and relieve ANP. This may be another important protective effect of resveratrol besides its anti-inflammatory activity.

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