those of controls, p<0.05. The expression of IL-23 protein were also higher than those of controls, which in concordance with the changes of mRNA.

**Conclusions** Our data show that local significantly increased levels of IL-23p19mRNA in myocardium and IL-23 may play a role in the pathogenesis of mice virus myocarditis.

**e0217** STUDY ON THE PROTECTIVE EFFECT OF THE MIXTURE OF SHENMAI PULVIS AND DANSHEN DECOCCTION ON THE MYOCARDIUM OF TYPE 2 DIABETIC CARDIOMYOPATHY IN RATS MODELS
doi:10.1136/hrt.2010.208967.217

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**Objective** To study the effect of the Mixture of Shengmai Pulvis and Danshen Decoction in protecting rats of the type 2 diabetic cardiomyopathy (DCM) model.

**Methods** 42 SD rats models of DCM, established by combination of insulin resistance by a high-fat diet with intraperitoneal injection of high dose streptozotocin (50 mg/kg), were evaluated in the damage of the myocardium by ECG at the twelfth week after modelling, and the serum were analysed for blood glucose (GLU), cholesterol and triglyceride (TG); the content of the left cardiac ventricle myocardial collagen was quantified by Masson staining test; the level of myocardial cell apoptosis was tested with Tunel apoptosis kit; the damage extent of the myocardial subcellular structure was observed by electron microscopy; the expression levels of cardiac TSP-1, TGF-β1 and TRB-3 proteins were detected by immunohistochemistry, the changes of the expression levels of the cardiac TSP-1, A-TGFβ1 and L-TGF-β1 protein were detected by Western blotting; and the changes of the mRNA expression levels of TSP-1 and TRB-3 were detected by real-time quantitative PCR.

**Results** Compared with the control group, the rat blood glucose, cholesterol, triglycerides were significantly decreased; the myocardial tissue was less damaged and the collagen fibre content was reduced in the group of the Mixture of Shengmai Pulvis and Danshen Decoction; The myocardial sub-cellular structural damage in electron microscopy was to a lesser extent, the expression levels of the myocardial TSP-1, TGF-β1 and TRB-3 by immunohistochemical detection and the average expression levels of the myocardial TSP-1, A-TGFβ1 and L-TGF-β1 by Western blotting were decreased; and the expression levels of TSP-1mRNA and TRB-3 mRNA by PCR detection were decreased than those of the control group.

**Conclusion** The Mixture of Shengmai Pulvis and Danshen Decoction can inhibit through multiple pathways the process of myocardial fibrosis in the rat myocardium of diabetic cardiomypathy, and significantly delay the formation course of diabetic cardiomyopathy in hyperglycemia rats.

**e0218** BIOLOGICAL CHARACTERISTICS RESEARCH OF STENT COATING WITH ZEDORAY CONSTITUENTS IN A PORCINE MODEL
doi:10.1136/hrt.2010.208967.218

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**Background** In-stent restenosis is caused by neointimal hyperplasia, which involves abnormal growth of vascular smooth muscle cells (VSMC). Zedoryy constituents is known to inhibit smooth muscle cell hyperplasia and migration, while inhibit ADP induced platelet aggregation.

**Objective** To evaluate the biological characteristics of stents coating with zedoryy constituents in porcine coronary model.

**Methods** Bare metal stents (BMS, n=56), Sirolimus eluting stents (SES, n=56) and Zedoryy eluting stents (ZES, n=56) were implanted in the proximal segment of three different epicardial coronary arteries in 36 swines randomly. Coronary angiography, optical coherence tomography (OCT) and histomorphologic analysis were performed at 30 days and 90 days after the procedure.

**Results** The 30 day (n=24) OCT examination showed ZES arm has larger lumen diameter (LD), acceptable mean lumen stenosis of area (MSA) compared with BMS (LD: ZES 1.9±0.51 mm, SES 1.85±0.41 mm, BMS 1.1±0.3 mm, p<0.05; MSA: ZES 21.7±19.3, SES 25.2±18.9, BMS 41.7±21.3, p<0.001). By histomorphometric analysis, similar injury scores were observed at the three arms (p>0.05). However, significant inflammation score reduction was seen in ZES group (ZES: 0.65±0.54, SES: 1.05±0.44, BMS: 0.94±0.75, p<0.001) compared to other two groups at 30 day, no differences in three groups at 90 day. Either at 30 day or 90 day, by qualitative analysis, well developed endothelium was seen in ZES arm, while impaired endothelium was observed with part of stent strut naked at vessel lumen at SES arm.

**Conclusion** Zedoryy eluting stents can reduce neointimal hyperplasia with good endothelia coverage in porcine coronary model.

**e0219** STUDY ON THE ROLE OF CD4+CD25+ TREG ON Atherosclerosis in ApoE-/- MICE
doi:10.1136/hrt.2010.208967.219

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**Objective** To investigate the mechanism of CD4+CD25+ treg in atherosclerosis in ApoE-/- mice.

**Methods** 28 weeks-old male ApoE-/- mice were randomly divided into two groups feeding high-fat diet (AH) or normal diet (AN). Ten C57BL/6J male mice feeding normal-diet (BN). After 12 weeks, the whole aorta from the root to crotch of iliac artery was separated and the whole blood were centrifugated to get the serum. Paraf sections of aorta were stained with H&E and morphometric analysis were performed at 30 days and 90 days after the procedure. Well developed endothelium was seen in ZES arm, while impaired endothelium was observed with part of stent strut naked at vessel lumen at SES arm.

**Results** The 30 day (n=24) OCT examination showed ZES arm has larger lumen diameter (LD), acceptable mean lumen stenosis of area (MSA) compared with BMS (LD: ZES 1.9±0.51 mm, SES 1.85±0.41 mm, BMS 1.1±0.3 mm, p<0.05; MSA: ZES 21.7±19.3, SES 25.2±18.9, BMS 41.7±21.3, p<0.001). So did in AH group than that in AN group (all p<0.05). There were advanced atherosclerotic plaques in ApoE-/- mice. The thickness of intima (m), plaque area (mm²), plaque/lumen ratio in AN group were significantly higher than in BN group (12.24±1.34 vs 7.58±2.25, 600265±462567625 vs 0, 15.9±5.45 vs 0, all p<0.01). So did in AH group than that in AN group (all p<0.05). The percentage of splenic CD4+CD25+ treg (%) and the serum concentration of TGF-β1, IL-10(pg/ml) in AN group were significantly decreased than in BN group (9.4±4.00 vs 13.2±5.97, 116.05±52.27 vs 191.27±95.27, 41.53±16.15 vs 61.84±23.05, all p<0.05). So did in AH group than that in AN group (all p<0.05). The expression of Foxp3+ and CD25+ cells in plaques-free intima in BN group. The expression of Foxp3, CD25+ cells were significantly decreased in AH group than in AN group (1.28±1.20 vs 3.04±1.92, 2.00±1.59 vs 3.98±1.67, all p<0.05).
**Conclusions** The splenic CD4^{+}CD25^{+}treg and serum concentration of TGF-β1, IL-10 of mice were decreased and there were lower frequency of Foxp3^{+} and CD25^{+} cells in severe atherosclerotic plaques than in mild. It meant that CD4^{+}CD25^{+}treg may has anti-atherosclerotic role on atherosclerotic progression.

**e0220** THE INVOLVEMENT OF IL-23/TH17 PATHWAY IN MURIN MODEL OF COXSACKIE VIRUS B3-INDUCED VIRAL MYOCARDITIS
doi:10.1136/hrt.2010.208967.220
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**Background** The IL-23/Th17 pathway plays an important role in the development of chronic inflammatory diseases and autoimmune diseases. However, the role of the IL-23/Th17 axis in the regulation of virus myocarditis (VMC) is still largely unknown.

**Methods** VMC was induced in male Balb/c mice by CVB3 peritoneal injection. Mice injected with PBS were taken as the controls. IL-23, IL-17 and RORγt mRNA in the myocardium of VMC were assessed by semi-quantitative RT-PCR on the time of 0, 1, 2, 3, 4 and 6 weeks after injection. IL-23, IL-17 protein from blood plasma was evaluated by ELISA. Flow cytometric analysis was used to evaluate the frequencies of Th17 subsets in CD4^{+}Tcell. CD4^{+} T cells were isolated from VMC mice and cultured with rIL-23 in vitro to investigate the function of IL-23 in the IL-23/Th-17 pathway.

**Results** Comparing with the controls, IL-23, IL-17 and RORγt mRNA were steadily expressed in the myocardium of infected mice from 1 week after virus infection (p<0.01). IL-23 and IL17 protein level increased from 1st week to 6th week. The frequencies of Th17 cells were obviously increased in VMC mice 1 week after infection (p<0.01), the maximum level of Th17 cells was reached at 4th week. The ratio of Th17 cells in the spleen lymphocyte significantly improved after rIL-23 stimulation, the IL-17 and RORγt mRNA expression of the cultured cells and the IL-17 protein in the culture supernatants increased after rIL-23 stimulation (p<0.05).

**Conclusions** IL-23/Th-17 pathway may play an essential role in VMC.

**e0221** EFFECTS OF EXTRACORPOREAL CARDIAC SHOCK WAVE THERAPY ON EXPRESSION OF ENDOTHELIAL NITRIC OXIDE SYNTHASE AND BASIC FIBROBLAST GROWTH FACTOR IN SWINE WITH ACUTE MYOCARDIAL INFARCTION
doi:10.1136/hrt.2010.208967.221
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**Objective** To observe the effects of extracorporeal cardiac shock wave therapy (CSWT) on cintent of endothelial nitric oxide synthase (eNOS) and basic fibroblast growth factor (bFGF) in serum and myocardial tissue in swine with acute myocardial infarction (AMI).

**Methods** 12 Model swines with acute myocardial infarction was made, and were randomly divided into two groups with six in each group: simplex myocardial infarction as the control group and the experimental group which received CSWT treatment. (three times on the first, third, fifth days after operation. 200 hit/point. Total: 12 points. Energy: 0.09 mJ/mm^{2}). Peripheral blood was extracted at eight different time points before and after operation (immediate, the first, third, fifth days, 1, 2, 3, 4 weeks after operation) to detect serum eNOS content with enzymelinked immunoassorbent assay.

Materials at myocardial tissue from the swines killed 1 month later were obtained to detect expression amount of eNOS and bFGF with semi quantitative RT-PCR method.

**Results** eNOS rose up in the experimental group 1 day after CSWT and a reached the peak on the fifth day, whereas eNOS gradually reduced in the control group, which showed significant difference between the two groups (p<0.01). Detection with semi quantities RT-PCR of myocardial tissues of infarction border area showed that eNOS and bFGF expression in the experimental group were obviously higher than those in the control group (eNOS 27.705±4.13) vs (16.445±3.51) (bFGF 52.571±4.23) vs (17.853±4.17 p<0.01).

**Conclusions** Extracorporeal cardiac shock wave therapy of acute myocardial infarction can effectively promote rise of eNOS and bFGF, which may be a new way to cure AMI.

**e0222** EFFECTS AND FUNCTION MECHANISM OF HYDROGEN SULFIDE ON MYOCARDIAL ISCHAEMIA REPERFUSION ARRAHYTHMIA IN RATS
doi:10.1136/hrt.2010.208967.222
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**Objective** To explore the Effects and Function Mechanism of hydrogen sulfide on Myocardial Ischaemia reperfusion Arrhythmia in Rats.

**Methods** We used sodium hydrosulfide (NaHS) as the donor of H_{2}S, SD rats were randomly divided into sham group, Myocardial Ischaemia reperfusion group (IR group), IR+NaHS group, and IR+NaHS+glibenclamide group. We monitor the Haemodynamics of rats, including heart rate, arterial pressure, left ventricular pressure et al. We also observe the rate of ventrical arrhythmia in each group.

**Result** H_{2}S can significantly reduces rats’ heart rate, arterial pressure and left ventricular pressure. It also reduces the rate of ventrical arrhythmia in Myocardial Ischaemia reperfusion Rats. The K_{ATP} Channel Blocker glibenclamide can weaken the H_{2}S Antiarrhythmic effects (p<0.01).

**Conclusions** H_{2}S can reduces the rate of ventrical arrhythmia in Myocardial Ischaemia reperfusion Rats. The Function Mechanism may be associated with the K_{ATP} signal transduction pathway in cells.