ARVC or sudden death (OR=7.300, 95% CI 1.606 to 33.177, p=0.013) and number of regions with myocardial fibrosis (OR=2.204, 95% CI 1.116 to 4.354, p=0.023) were independent predictors for life-threatening ventricular arrhythmia in ARVC.

Conclusions MRI is the optimal imaging approach for detecting ARVC. Familial history of ARVC or sudden death, the accordion sign and number of regions with myocardial fibrosis were associated with an increased risk of life-threatening ventricular arrhythmia in patients with ARVC.

**e0605** "ONE-STOP SHOP" EXAMINATION OF COMPLICATED AND COMPLEX CONGENITAL HEART DISEASE WITH MRI

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Objective To evaluate the diagnostic value of MRI in complicated congenital heart disease, comparing with echocardiography and x-ray cardio-angiography.

Methods 20 patients with complicated or complex congenital heart disease (male 12, female 8, mean age 17.8 y, ranging from 3–46 y) underwent MR examination. A wireless vector ECG-gating was used for triggering, all the sequences were applied with breath hold or free breathing. Turbo spin echo and multiple gradient echo sequences including FLASH (fast low angle shot) and TrueFISP (true fast imaging with steady-state precession.) with TSENSE (adaptive sensitivity encoding incorporating temporal filtering) and TREAT (time resolved echo-shared technique) were used to evaluate the morphology, function, flow status of the heart. MR findings were compared with the results of echocardiography and x-ray cardio-angiography.

Results All the examination were accomplished successfully and diagnostic MR images were obtained satisfactorily. The mean total scan time was 35 min ranging from 29–55 min. According to the main malformation diagnosed by MRI, double-outlet of right ventricle (DORV) were 11 cases, single ventricle were 3 cases, function-corrected transposition of great arteries were 2 cases, endocardial cushion defect, interruption of aortic arch, coarctation of the aorta and pulmonary atresia was 1 case, respectively. The total coincidence was 95% comparing with echocardiography, of the aorta and pulmonary atresia was 1 case, respectively. The function-corrected transposition of great arteries were 2 cases, main malformation diagnosed by MRI, double-outlet of right ventricle.

Conclusion The "One-stop shop" examination of complicated and complex congenital heart disease with MRI offers a comprehensive evaluation of complicated congenital heart disease, comparing with echocardiography and x-ray cardio-angiography.

**e0607** ANALYSIS ON EARLY DIAGNOSIS GRADING MODEL OF ACUTE AORTIC DISSECTION

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Objective To investigate the features of the clinical manifestation, laboratory data and imageology information in acute Aortic Dissection (AD) patients, to find the early, right and easy grading model of diagnosing AD.

Method Analyze the clinical manifestation, Laboratory data and imageology information of 182 AD patients, who were our patients at Emergency Department in last three years, and meanwhile, compared them with 184 chest and back ache patients, to search the early diagnosis grading model of AD.

Results If 5 score was the standard of the grading system to early diagnose AD on the basis of logistic regression equation and clinical practice, the sensitivity of AD forecast is 96.7%, specificity is 81.0%.

Conclusions We can improve the emergency diagnose level of AD through establishing the early grading model which contains the stabling and severe pain, distinct rise of the blood pressure, asymmetry of the blood pressure and/or the pulse, wide arteriae aorta and/or mediastinum on chest x-ray, obvious rise of D-dimmer.

**e0608** A NITINOL OCCLUDER AND A SPECIAL DELIVERY DEVICE FOR PATENT DUCTUS ARTERIOSUS (PDA) CLOSURE

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Background Nitrol occluders used for patent ductus arteriosus (PDA) may result in the stenosis of aortic and pulmonary artery when applied for smaller-sized children. The present study sought to design and develop an improved nitinol occluder and delivery system, and to verify the reliability and safety of this device on canine model of PDA.

Methods A canine FDA model was established by anastomosing the internal jugular vein with the left pulmonary artery and the descending aorta in an end-to-side fashion. The effects of the novel nitinol occluder and its delivery system were followed-up for 6 months after the closure.

Results FDA was successfully established in 10 out of 18 dogs and was treated by transcatheter occlusion with the novel nitinol occluder. Postoperative echocardiography showed that the location and shape of the occluder were normal without any regurgitation. Morphological examination revealed that the surface of the occluder was covered by a grey membrane-like tissue, which was identified as endothelial tissue by histological and electron microscopy examinations. There was no corrosion or severe inflammation on the occluder.

**e0609** TRANSCATHETER CLOSURE OF A GIGANTIC RIGHT SINUS OF VALSALVA ANEURYSM TO RIGHT VENTRICULAR FISTULA: A CASE REPORT

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Ruptured aneurysms of the sinus of Valsalva with a gigantic fistulous tract are extremely rare. A single origin of the coronary arteries is a rare coronary anomaly. We describe the case of a 6-year-old girl who had a ruptured aneurysm of the right coronary sinus with a gigantic fistulous tract and an abnormal origin of the right coronary artery. Transthoracic echocardiography showed the right coronary artery (RCA) aneurysm and RCA fistula connecting to the right ventricular (RV). Aortography revealed the presence of a right coronary sinus of Valsalva aneurysm (SVA)-RV fistula and the originating of right coronary artery (RCA) from left circumflex artery (LCX). She was diagnosed as having a gigantic SVA to right ventricle fistula and a single origin of the coronary arteries. Successful transcatheter closure of the fistulous ostium was performed.