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IDENTIFICATION OF ENLARGED CONAL ARTERY IN TETRALOGY OF FALLOT WITH SECTOR SCAN ECHOCARDIOGRAPHY

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Abstract — In a 9-month-old boy with tetralogy of Fallot, prominent conal branch of the right coronary artery traversing the right ventricular outflow tract was visualized with sector scan echocardiography. Angiography in the aortic root confirmed the echocardiographic diagnosis. The advantage of noninvasive diagnosis of these anomalies is in postponing angicardiography and surgery in cases of tatralogy of Follot when major branch of the coronary artery crosses the right ventricular outflow tract. An alternative palliative operation may be necessary in such patients.

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Case report

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Introduction — In 5—9% of patients with tetralogy of Fallot, anterior descending coronary artery or another major branch crosses the right ventricular outflow tract (1, 2). When total correction is planned, the right ventricular outflow tract or transanular patch must be frequently used especially in infants. The anomalous artery can be severed at the time of right ventricular outflow tract incision and fatalities may occur. Thus preoperative delineation of coronary arteries is essential. Routinely employed aortic root angiography with a venous catheter passed through ventricular septal defect, or if this fails, with a retrograde catheter usually satisfactorily shows the coronary arteries.

Sector scan echocardiography has been used to assess the coronary arteries in transposition of great arteries (3).

In this report we describe the sector scan echocardiographic imaging of prominent conal artery branch of the right coronary artery crossing the right ventricular outflow tract in tetralogy of Fallot.

Case report — A 9-month-old boy with diagnosis of tetralogy of Fallot was presented to us at the age of 4 months with the episodes of cyanotic spells. At that time he was started on pro-



Fig. 1a — Short axis parasternal view of the left anterior descending coronary artery (lad) and circumflex branch of the left coronary artery (cir). AO = aorta, PA = pulmonary artery.

pranolol and received the transfusion of packed red cells because of anemia.

Cyanotic spells reappeared a few days before his second admission. Sector scan echocardio-



Fig. 1b — Short axis parasternal view of the conal branch (cb) of the right coronary artery crossing the right ventricular outflow tract. AO=aorta, PA=pulmonary artery.



Fig. 2 — Aortic root angiography showing the conal branch (arrow) of the right coronary artery (rca) which crosses the right ventricular outflow tract.

graphy revealed typical features of tetralogy of Fallot with pronounced infundibular stenosis. Special attention was paid to the examination of the coronary arteries. We used 5 MHz medium focus transducer from high left parasternal short axis view just above the aortic valve (1). To see the left coronary artery the transducer was oriented inferiorly so that the pulmonary valve annulus was shown on cross section. Anticlockwise rotation of the transducer revealed prominent conal branch of the right coronary artery (Fig. 1 a, b). Cardiac catheterization and angiocardiography confirmed echocardiographic diagnosis of tetralogy of Fallot. Injection of the contrast medium into the aortic root in left anterior oblique projection of 45 degrees and cranial angulation of 20 degrees showed the right coronary artery and the conal branch crossing the right ventricular outflow tract (Fig. 2).

Discussion — Recent report on the echocardiographic examination of the anomalies of the coronary arteries in tetralogy of Fallot has identified coronary arteries in 78 % of studied children. Major obstacles for adequate imaging of coronary arteries in this defect have been poor penetration of ultrasound beam in large patients, concomitant pulmonary disease with persistent overlaying of lung tissue, and pulmonary artery atresia with severe hypoplasia of the right ventricular outflow tract. The anomalies have been found in 9 of 29 studied children.

Only in one patient a small conal branch has been visualized. It has not been considered abnormal (4). However, conal branch of the right coronary artery in tetralogy of Fallot tends to be long and swings lower over the right ventricular outflow tract instead of being short and high as is the case in normal situation. The enlarged conal artery is the response to right ventricular hypertrophy. It may render right ventriculotomy undesirable (1, 5). Surgeon can usually identify anomalies of the coronary arteries during surgery but sometimes these may be obscured by adhesions of previous operations or may run intramurally.

Thus the advantage of sector scan echocardiographic examination of the coronary arteries is in postponing the invasive investigation and surgery in cases of tetralogy of Fallot where major coronary artery crosses the right ventricular outflow tract. An alternative palliative operation may be indicated in such patients.

Povzetek

PRIKAZ POVEČANE KONUSNE ARTERIJE PRI TETRALOGIJI FALLOT Z DVODIMENZIONALNO EHOKARDIOGRAFIJO

Pri 9 mesečnem dečku s tetralogijo Fallot sva prikazala povečano konusno arterijo, ki je prečkala iztočni trakt desnega prekata z dvodimenzionalno ehokardiografijo. Diagnozo sva potrdila z angiografijo v aortni koren. Prednost neinvazivne ehokardiografske diagnostike prikaza koronarnih arterij je v preložitvi invazivne diagnostike in kirurškega posega pri bolnikih s teralogijo Fallot, kjer večja koronarna arterija prečka iztočni trakt desnega prekata. Včasih je pri teh pacientih potrebno napraviti alternativno paliativno operacijo.

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27th Congress of the European Society of Paediatric Radiology

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