Prenatal diagnosis of major aortopulmonary collateral arteries

Tomasz Moszura1, Katarzyna Janiak2, Maria Respondek-Liberska2, Anna Mazurek-Kula1, Paweł Dryżek1, Jacek Moll3, Andrzej Sysa1

1Cardiology Department, Polish Mothers’ Memorial Hospital, Research Institute, Lodz, Poland
2Department for Diagnosis and Prophylaxis of Foetal Malformations, Polish Mothers’ Memorial Hospital, Research Institute and Medical University, Lodz, Poland
3Cardiosurgery Department, Polish Mothers’ Memorial Hospital, Research Institute, Lodz, Poland

Abstract

Background: The presence of foetal major aortopulmonary collateral arteries (MAPCAs) is associated with adverse outcome, therefore early diagnosis is essential.

Aim: To evaluate the usefulness of foetal echocardiography in the diagnosis and evaluation of MAPCAs in foetuses with pulmonary atresia, as well as to assess the effects of prenatal diagnosis on the management of neonates with pulmonary atresia.

Methods: From 11,678 examined foetuses, we retrieved 15 cases of patients with MAPCAs and congenital heart defects which had been diagnosed by foetal echocardiography (1994–2008), using 2D echocardiography + color-Doppler (CD) + pulsed Doppler (2DD) and spatio-temporal image correlation (STIC) techniques. In 13 patients, MAPCAs were confirmed after birth based on angiography.

Results: In all cases, vessels corresponding to MAPCAs were visible in longitudinal view with CD, and in three cases were additionally confirmed by STIC technique. In nine cases one, in four cases two, and in two cases three MAPCAs were suspected. In two cases, MAPCAs were not confirmed after birth; one due to misdiagnosis secondary to aberrant right subclavian artery, and one because of abnormal ductus arteriosus course coexistent with right aortic arch.

Conclusions: In foetuses with pulmonary atresia, it is possible to find MAPCAs with current technology (both 2D + CD, power angiography and real time-3D echocardiography [4D]). The differential diagnosis (MAPCAs or other vessels) should be included. Although prenatal diagnosis does not change the obstetrical management, it is important information for a paediatric cardiologist. Early neonatal angiography might be of great value not only in confirming MAPCAs, but also in performing cardiac intervention and in some cases preventing future heart failure.

Key words: foetal echocardiography, MAPCAs, pulmonary atresia
The presence of diverse pulmonary vascularisation of the lung by aortopulmonary collaterals has serious clinical implications [1]. Different sources of pulmonary blood flow can result in abnormal vascular development, with the risk of pulmonary hypertension developing in different pulmonary segments [2, 3]. In those with poorly developed pulmonary arteries and restrictive pulmonary blood flow, cyanosis is the main clinical manifestation, while in those with a large number of wide aortopulmonary collaterals, early heart failure is manifest [4]. Current ultrasonographic diagnostics should allow for the identification of aortopulmonary collaterals before birth [5]. The aim of our study was to report on the utility of foetal echocardiography in the diagnosis and evaluation of major aortopulmonary collateral arteries (MAPCAs) in the foetus with PA. Prenatal diagnosis will allow for determination of prognosis and may speed correct therapeutic interventions directly after birth.

**METHODS**

**Study group**

From 11,678 foetuses evaluated between 1994 and 2008, we identified 15 cases with pulmonary outflow obstruction and suspicion of MAPCAs diagnosed by foetal echocardiography: 13 of them presented with PA and two with critical pulmonary stenosis. Nine foetuses had tetralogy of Fallot and PA; three had complex single ventricle; two had tricuspid atresia with PA; and one had atrio-ventricular septal defect with PA. All had post-natal evaluation in our institution. In 11 patients, MAPCAs were confirmed by post-natal cardiac catheterisation (Fig. 1). In cases where there were wide collaterals possibly leading to heart failure and pulmonary hypertension, we interventionaly closed collaterals where dual supply to lung segments was identified (blood inflow into pulmonary segments from a minimum of two different sources). Collateral embolisation was performed using a Jackson-coil PDA. In one patient with critical pulmonary valve stenosis, pulmonary artery hypoplasia and the presence of numerous aortopulmonary collaterals, we simultaneously conducted percutaneous balloon pulmonary valvuloplasty (balloon catheter TYSHEAK II (6 × 20 mm) and collateral embolisations.

**Foetal echocardiography**

The ultrasound systems used were an ATL HDI-5000 and GE 730 Voluson Expert with transabdominal probes convex type C5 — 2 MHz and C7 — 4 MHz, cardiologic probe P4 — 2 MHz. A consistent standard technical approach to cardiac anatomy and function was applied in all cases: short- and long-axis scans of the intracardiac anatomy, aorta and pulmonary arteries were obtained. Colour Doppler (CD), pulsed-wave, and continuous wave were used as well. Echocardiographic recordings were retrospectively analysed from videotapes or digital media. Gestational age at the time of initial diagnosis and gestational age at the time of MAPCAs visualisation were recorded.

**Angiography**

After birth, all neonates had left and right-side angiography with detailed delineation of the pulmonary vasculature. Angiographic views were recorded using a uniplanar angiograph (Philips Integris CV). Aortopulmonary collateral estimation required selective angiography with use of a Judkins catheter R 3.0, vertebral and multipurpose. Pressure measurements in the pulmonary arteries and collaterals were obtained.

**RESULTS**

Mean maternal age was 29 ± 4.2 years. Congenital heart defects in our group of foetuses were diagnosed at mean 30.3 weeks of gestation (21–37 weeks). However, a suspicion of foetal MAPCAs was visible on foetal echocardiography later on: at mean 32.6 weeks of gestation (27–38 weeks). There were seven high-risk pregnancies (46.7%): three after in-vitro fertilisation, two based on family history (in one case the first child with common arterial trunk the other with a chest abnormality), one based on obstetric history (miscarriages at first and second pregnancy), and one with diabetes mellitus. Mean neonatal weight at the time of delivery was 2,590 ± 470 g. There was one case of intrauterine growth retardation; the others foetuses were appropriate for gestational age. There was normal mean heart size (heart area/chest area ratio) 0.39 ± 0.07, but in one case there was cardiomegaly (HA/CA 0.55 — case 12) [6]. There was dextrocardia in two cases and situs inversus in one case. A right aortic arch was suspected in two cases. In nine cases, there was identification of an abnormal four-chamber view with overriding aorta, ventricular septal defect and a rudimentary right ventricular outflow tract; and in six cases there was an abnormal four-chamber view due to the
presence of a complex single ventricle (thrice), tricuspid atresia (twice), or atrio-ventricular septal defect (once). There was one false positive diagnosis of d-trasposition of great arteries in the patient number 4.

In all cases, MAPCAs were visible in the longitudinal view (Fig. 2) with CD imaging (Fig. 3). In three cases, it was also confirmed by spatio-temporal image correlation (STIC) and there was no false positive diagnosis. In nine cases, one collateral vessel was seen, in four cases two abnormal vessels, and in two cases three abnormal vessels. The vessels measured 2–4 mm in diameter on 2D echocardiography and CD. Foetal Doppler spectrum was obtained in ten cases and was similar for pulmonary flow, which can be obtained from pulmonary peripheral branches. In two cases, MAPCAs were not confirmed after birth (two cases of false positive prenatal detection). During catheterisation in these cases, an abnormal course of the subclavian artery was seen in one case (aberrant right subclavian artery) and in the other case there was an atypical course for the ductus arteriosus due to the right aortic arch.

In our study group, we noticed specific findings: more than one collateral visualised in foetal echocardiography can be associated with a poor postnatal prognosis (anatomical type II and III of PA). In our referral centre (Foetal Cardiology type C in Poland) we also had several foetuses without prenatal diagnosis of PA and post-natal angiographic diagnoses of MAPCAs. However, these cases were not included in this type of analysis.

In four foetuses, in addition to the main cardiac diagnosis, there was a prenatal suspicion of thymic hypoplasia. Of these, in three neonates CATCH 22 syndrome was confirmed and in one case thyroid hypoplasia was diagnosed post-natally. Cardiac catheterisation confirmed additional sources of pulmonary vascular supply in 13 patients (Fig. 4). Because of the risk of pulmonary hypertension and development of cardiac failure, two neonates had coil embolisation performed; in one case simultaneously with percutaneous pulmonary artery valvuloplasty. All the patients with MAPCAs confirmed post-natally were qualified for cardiological surgical treatment as listed in Table 1.
Table 1. Prenatal and post-natal diagnoses in a series of 15 foetuses and neonates with MAPCA

<table>
<thead>
<tr>
<th>N</th>
<th>Gender</th>
<th>Gest age</th>
<th>Foetal dgn</th>
<th>No. of foetal MAPCA</th>
<th>Neonatal dgn</th>
<th>Angio type</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Female</td>
<td>37</td>
<td>ToF+PA</td>
<td>2</td>
<td>ToF+PA + MAPCA</td>
<td>II</td>
<td>No, MAPCA occlusion with coil, unifocalisation at four months of age</td>
</tr>
<tr>
<td>2</td>
<td>Male</td>
<td>23</td>
<td>SV+PA, situs inversus</td>
<td>1</td>
<td>SV+PA + MAPCA, situs inversus</td>
<td>III</td>
<td>B-T shunt to the main collateral because of low oxygen saturation due to its proximal stenosis, 46th day of life, death after surgery</td>
</tr>
<tr>
<td>3</td>
<td>Female</td>
<td>27</td>
<td>ToF+PA</td>
<td>1</td>
<td>ToF+PA + MAPCA</td>
<td>I</td>
<td>No surgery in four weeks of post-natal life</td>
</tr>
<tr>
<td>4</td>
<td>Female</td>
<td>24</td>
<td>SV/DILV+d-TGA+PA, dextrocardia</td>
<td>1</td>
<td>SV+PA + MAPCA, dextrocardia</td>
<td>I</td>
<td>No surgery in four weeks of post-natal life</td>
</tr>
<tr>
<td>5</td>
<td>Female</td>
<td>29</td>
<td>ToF+PA + Ao arch dxt</td>
<td>3</td>
<td>PA+VSD + MAPCA</td>
<td>III</td>
<td>B-T shunt 21st day of post-natal life (CATCH 22)</td>
</tr>
<tr>
<td>6</td>
<td>Male</td>
<td>33</td>
<td>ToF+PA</td>
<td>1</td>
<td>ToF+PA + MAPCA</td>
<td>II</td>
<td>B-T shunt 28th day of post-natal life (CATCH 22)</td>
</tr>
<tr>
<td>7</td>
<td>Female</td>
<td>36</td>
<td>DORV/SV+VSD+PA</td>
<td>3</td>
<td>DORV+PA + MAPCA</td>
<td>II</td>
<td>B-T shunt 31st day of post-natal life (CATCH 22)</td>
</tr>
<tr>
<td>8</td>
<td>Female</td>
<td>31</td>
<td>ToF+PS</td>
<td>2</td>
<td>ToF+critical PS + MAPCA</td>
<td>II</td>
<td>B-T shunt 35th day of post-natal life</td>
</tr>
<tr>
<td>9</td>
<td>Male</td>
<td>31</td>
<td>TVA+PA</td>
<td>3</td>
<td>TVA + PA + MAPCA</td>
<td>II</td>
<td>No surgery in four weeks of post-natal life</td>
</tr>
<tr>
<td>10</td>
<td>Female</td>
<td>23</td>
<td>ToF+PA</td>
<td>1</td>
<td>ToF+PA + MAPCA</td>
<td>II</td>
<td>B-T shunt 20th day of post-natal life</td>
</tr>
<tr>
<td>11</td>
<td>Female</td>
<td>36</td>
<td>ToF+PA</td>
<td>2</td>
<td>ToF+PA + artery subclavia lusoris</td>
<td>II</td>
<td>No surgery in four weeks of post-natal life</td>
</tr>
<tr>
<td>12</td>
<td>Male</td>
<td>29</td>
<td>TVA+PA+AS+FO, RC, dextrocardia</td>
<td>1</td>
<td>SV+PA + AS, dextrocardia</td>
<td>II</td>
<td>Post-natal death, no autopsy</td>
</tr>
<tr>
<td>13</td>
<td>Female</td>
<td>31</td>
<td>ToF+PA</td>
<td>1</td>
<td>ToF+PA + artery subclavia lusoris</td>
<td>II</td>
<td>Post-natal death, autopsy (+)</td>
</tr>
<tr>
<td>14</td>
<td>Female</td>
<td>31</td>
<td>AVC+PS+Ao arch dxt, DA dxt, Dandy Walker S.</td>
<td>1</td>
<td>AVC + d-TGA+PS</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>15</td>
<td>Male</td>
<td>32</td>
<td>ToF+PA</td>
<td>1</td>
<td>ToF+PA + MAPCA</td>
<td>II</td>
<td>B-T shunt 36th day of post-natal life</td>
</tr>
</tbody>
</table>

MAPCA — major aortopulmonary collateral arteries; ToF — tetralogy of Fallot; PA — pulmonary atresia; SV — single ventricle; B-T — Blalock-Taussig; DILV — double inlet left ventricle; d-TGA — d-transposition of great arteries; Ao arch dxt — aortic arch duplication; VSD — ventricular septal defect; CATCH — cardiac defect, abnormal face, thymic hypoplasia, cleft palate, hypocalcemia; DORV — double outlet right ventricle; PS — pulmonary stenosis; BVP — balloon valvuloplasty; TVA — tricuspid valve atresia; AS — aortic stenosis; FO — foramen ovale; RC — restrictive cardiomegaly; AVC — atrioventricular canal; DA — ductus arteriosus.
CONCLUSIONS

In foetuses with PA, it is possible to find MAPCAs using current technology (both 2D + color Doppler, power angiography and 4D). A differential diagnosis (MAPCAs or other vessels) should be included. Although prenatal diagnosis does not change the obstetric management, it is valuable information for a paediatric cardiologist. Early neonatal angiography may be of great value, not only in confirming MAPCAs, but also in allowing cardiac intervention and in some cases preventing future heart failure.

Conflict of interest: none declared

References

Diagnostyka prenatalna kolaterali aortalno-płucnych

Tomasz Moszura¹, Katarzyna Janiak², Maria Respondek-Liberska², Anna Mazurek-Kula¹, Paweł Dryżek¹, Jacek Moll³, Andrzej Sysa¹

¹Klinika Kardiologii, Instytut Centrum Zdrowia Matki Polki, Łódź
²Zakład Diagnostyki i Profilaktyki Wad Wrodzonych, Instytut Centrum Zdrowia Matki Polki, Łódź
³Klinika Kardiochirurgii, Instytut Centrum Zdrowia Matki Polki, Łódź

Streszczenie

Wstęp: Obecność kolaterali aortalno-płucnych u pacjentów z atrezją zastawki płucnej wiąże się z poważnymi skutkami klinicznymi. W grupie osób z dużą liczbą kolaterali szybko dochodzi do rozwoju niewydolności serca. U pacjentów ze zmniejszonym przepływem płucnym i słabo rozwiniętymi naczyniami płucnymi dominującym objawem jest sinica, podczas gdy odaortalne unaczynienie płuc może prowadzić do rozwoju nadciśnienia płucnego w poszczególnych segmentach płuc. Nowoczesna diagnozyka echokardiograficzna pozwala na rozpoznanie odaortalnego unaczynienia płuc już w okresie prenatalnym.

Cel: Celem pracy była ocena przydatności echokardiografii płodowej w rozpoznaniu i ocenie kolaterali aortalno-płucnych (MAPCAs) u płodów z atrezją zastawki płucnej oraz wpływ diagnozyki płodowej na postępowanie z noworodkiem z atrezją zastawki tętnicy płucnej.

Metody: Analizie retrospektywnej poddano 11 678 prenatalnych badań echokardiograficznych [echokardiografia dwuwymiarowa (2D) + technika Dopplera znakowanego korelem (CD) + Doppler pulsacyjny (2DD) oraz czasoprzestrzenne korelacja obrazu (STIC)] wykonanych w latach 1994–2008 w Zakładzie Diagnostyki i Profilaktyki Wad Wrodzonych ICZMP. Wśród płodów z atrezją zastawki płucnej w 15 przypadkach uwidoczniło kolaterale aortalno-płucne w badaniu prenatalnym. U 13 z nich obecność odaortalnego unaczynienia płuc potwierdzono badaniem angiograficznym.

Wyniki: We wszystkich przypadkach naczynia odpowiadające kolateralom aortalno-płucnym uwidoczniło w osi długiej z przepływem potwierdzonym w CD, w 3 przypadkach potwierdzono je także za pomocą techniki STIC. W 9 przypadkach podejrzewano 1 kolateralę, w 4 przypadkach — 2, w 2 przypadkach — 3. W 2 przypadkach obecności kolaterali nie potwierdzono po urodzeniu. W przypadku pierwszym fałszywie dodano rozpoznanie prenatalne było wynikiem obecności tętnicy podobojczykowej błądzącej, w drugim przypadku — nietypowego przebiegu przewodu tętniczego współistniejącego z prawostronnym łukiem aorty.

Wnioski: U płodów z atrezją zastawki płucnej jest możliwa prenatalna diagnoza kolaterali aortalno-płucnych przy użyciu nowoczesnych technik echokardiograficznych [2D + CD + 2DD, Doppler mocy (power-angio), echokardiografia 3-wymiarowa w czasie rzeczywistym (4D)]. Prenatalne rozpoznanie dodatkowego odaortalnego unaczynienia płuc u płodu z atrezją zastawki płucnej nie wpływa na postępowanie położnicze, stanowi natomiast cenną informację dla kardiologa dziecięcego. Wczesna diagnozyka angiograficzna w tej grupie noworodków ma na celu nie tylko potwierdzenie prenatalnego rozpoznania kolaterali aortalno-płucnych, ale pozwala także w niektórych przypadkach na przeprowadzenie leczenia interwencyjnego w celu uniknięcia rozwoju niewydolności serca w przyszłości.

Słowa kluczowe: echokardiografia prenatalna, kolateral aortalno-płucne, atrezja płucna

Kardiol Pol 2011; 69, 2: 146–151

Adres do korespondencji: Adres do korespondencji: Adres do korespondencji: Adres do korespondencji: Adres do korespondencji:
dr n. med. Anna Mazurek-Kula, Klinika Kardiologii, Instytut Centrum Zdrowia Matki Polki, ul. Rzgowska 281/289, 93–338 Łódź, e-mail: qla@op.pl

www.kardiologiapolska.pl