Prenatal Diagnosis of Persistent Left Superior Vena Cava and its Clinical Significance

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Background: Persistent left superior vena cava (PLSVC) is a variant of systemic venous return which is observed in 0.3% of autopsies in the general population and in 4-8% of patients with congenital heart disease. Aims: To evaluate associated cardiac, extracardiac and chromosomal anomalies in prenatally diagnosed cases of PLSVC and to review their outcome.

Study Design: Retrospective comparative study.

Methods: The data of patients with a prenatal diagnosis of PLSVC between May 2008 and January 2013 were reviewed retrospectively.

Results: Data of 31 cases were reviewed. Fifteen (48.4%) cases were associated with cardiac defects and 17 (54.8%) cases had associated extracardiac sonographic or postpartum findings. Two fetuses had karyotype anomalies. Outcome was significantly more favorable in cases not associated with cardiac defects in comparison to those associated with cardiac anomalies (84.6% vs. 33.3%, p=0.009). All cases with isolated PLSVC survived, while among the cases associated with extracardiac anomalies, with cardiac anomalies and with both extracardiac and cardiac anomalies, the survival rate was 75%, 50% and 22.2%, respectively. The most frequent group of cardiac anomalies associated with PLSVC was septal defects and VSD was the most common heart defect individually, being observed in nine fetuses.

Conclusion: Prenatally diagnosed PLSVC is associated with cardiac and extracardiac anomalies in the majority of cases. Outcome is significantly worse if PLSVC is associated with a cardiac defect, and the prognosis is excellent in isolated cases.

Key Words: Cardiac anomaly, congenital anomaly, persistent left superior vena cava, prenatal diagnosis, ultrasound

Persistent left superior vena cava (PLSVC) is a variant of systemic venous return which is observed in 0.3% of autopsies in the general population and in 4-8% of patients with congenital heart disease (1). In the embryonic period, the right and left anterior cardinal veins constitute the main venous drainage of the cephalic portion, and as a result of the development of the left innominate vein that bridges the anterior cardinal veins at 8 weeks gestation, the proximal part of the left cardinal vein regresses and only a small portion of it remains as the left superior intercostal vein (LSVC) (2-4). A defect in the regression of this vein is thought to be the cause of the persistence of the LSVC. PLSVC drains mostly into the right atrium via the coronary sinus, and into the left atrium in 10% of cases (4). In general, the diagnosis of PLSVC has no clinical impact, because it is asymptomatic, and the systemic venous blood continues to return to the right atrium (5, 6). However, a prenatal diagnosis of PLSVC plays an important role in prenatal counseling and management, since it is associated with cardiac and extra-cardiac diseases with an incidence as high as 83% and 48%, respectively (2, 6).

There are only a few series of prenatally diagnosed PLSVC in the literature (2, 6, 7). In this study, we aimed to identify prenatally diagnosed cases of PLSVC in our clinic, to evaluate the associated cardiac, extracardiac and chromosomal anomalies, and to review their outcome.

MATERIAL AND METHODS

In this study, the data of patients with prenatal diagnosis of PLSVC between May 2008 and January 2013 were reviewed retrospectively. All patients were examined in a tertiary referral center for prenatal diagnosis where an anatomic survey and fetal echocardiography were performed in a standardized fashion with high resolution equipment (Voluson 730 Expert, GE Healthcare, Milwaukee, WI, USA and Xario SSA-660A, Toshiba Medical Systems). Fetal echocardiography was carried out by visualizing standard anatomical planes including the observation of the situs, the position of the heart, the four-chamber view, the outflow tracts, the three-vessel view, the aortic and ductal arches and both systemic and pulmonary venous...
RESULTS

Over the study period, a total of 35 fetuses with PLSVC were examined. Four of them were excluded due to incomplete data and loss to follow up. In all of the 31 cases, a normal right superior vena cava (RSVC) was also present. Seven cases were referred to our center with a suspicion of PLSVC. Fourteen cases were referred due to other cardiac and extracardiac defects, and PLSVC was detected during echocardiography. In 10 cases, PLSVC was detected in our clinic during routine ultrasound scanning. The demographic characteristics, associated extracardiac findings, karyotype and outcome of the fetuses are shown in Table 1. The cases were categorized into two groups: Group 1 constituted cases without associated sonographic or postnatal cardiac defects, while group 2 included those with associated sonographic and/or postnatal cardiac defects. Group 1 included 16 (51.6%) fetuses, and in half of them additional extracardiac findings were observed. Thirteen (81.25%) fetuses had a normal karyotype. In a case with choroid plexus cyst and thickened nuchal fold, trisomy 21 was diagnosed postnatally, because the parents did not accept an invasive procedure during pregnancy. Group 2 involved 15 (48.4%) fetuses and in 9 (60%) of them additional extracardiac findings were observed. Eleven (73.3%) fetuses had a normal karyotype, and one fetus with overlapping fingers, fetal growth restriction, choroid plexus cyst, double outlet right ventricle, ventricular septal defect (VSD) and aberrant right subclavian artery was found to have a 47,XX+18 karyotype. The rate of aneuploidy in the total cohort was 7.7% (2/26).

The cases were followed up for a period of 3 months to 6 years. The outcome was significantly more favorable in Group 1 in comparison to Group 2 (84.6% vs. 33.3%, p=0.009) (Table 1). Table 2 shows the outcome of cases (except for the unborn fetuses) in detail. All cases with isolated PLSVC survived and were doing well at the time of writing. On the other hand, among the cases associated with extracardiac defects, with other cardiac defects and with both extracardiac and cardiac defects, the survival rate was 75%, 50% and 22.2%, respectively. Eight cases were operated on for extracardiac defects or cardiac defects other than PLSVC, and three of them died postoperatively. Five of them were good in health at the time of writing. The features of all cases in Group 1 and Group 2 are summarized in Table 3 and Table 4, respectively.

The most frequent group of cardiac anomalies was septal defects (eleven cases) followed by conotruncal anomalies (four cases), left ventricular outflow obstructive diseases (four cases) and heterotaxy syndromes (three cases). VSD was the most common heart defect individually and was observed in nine fetuses.

DISCUSSION

In this study, we reviewed the prenatal and postnatal cardiac and extracardiac findings, associated chromosomal anomalies and outcome in 31 PLSVC cases. Prenatally, PLSVC is easily and accurately diagnosed in the three-vessel view, where four vessels are visualized instead of three, with an extra vessel lo-
cated to the left of the pulmonary trunk (3). In rare instances of the absence of the RSVC, there are three abnormally arranged vessels aligned, i.e. the aorta, pulmonary artery and PLSVC from right to left. Significant dilatation of the coronary sinus (Figure 2), which is best visualized in a transverse transthoracic cross-sectional plane slightly caudal to the apical four-chamber view, is an indirect sign of PL SVC, as well (8). However, the presence of PL SVC must be confirmed in the three-vessel view, because anomalous pulmonary venous drainage may also cause significant dilatation of the coronary sinus (8). Also, a dilated coronary sinus may be misdiagnosed as an ostium primum atrial septal defect at the level of the opening of the coronary sinus into the right atrium or as mitral atresia since the mitral valve is pushed anteriorly by the coronary sinus and may not be seen in the four-chamber view (7).

The most significant clinical implication of prenatally diagnosed PL SVC is the association with cardiac and extracardiac defects. Previously, Galindo et al. (6) reported that 44 of 54 cases (81.4%) with PL SVC were associated with a cardiac defect; the corresponding rate in the series of Berg et al. (2) was 83% (68/82). Unlike these results, more than half (16/31) of the cases in our series were not associated with additional cardiac malformation. Furthermore, seven cases (9%) in the series of Berg et al. (2) and three cases (5.5%) in the series of Galindo et al. (6), had isolated PL SVC with no associated cardiac or extracardiac findings, in contrast to our series in which the corresponding rate was as high as 25.8% (8/31). The inconsistency of our findings with the previous studies may be a result of more common incorporation of three-vessel view in routine systematic ultrasound examination of fetal heart in the last few years, since these previous series were published in 2006 and 2007.

Nshah et al. (9) postulated that for the regression of the left anterior cardinal vein, compression by the growing lungs and the increasing size of the atria are necessary. In their postnatal series, a significantly more frequent association was recorded with anomalies, such as atrioventricular canal malformation, mitral atresia and cor triatratum, which reduce compression of the developing LSVC (9). On the other hand, PL SVC is present in 50-70% of the cases with heterotaxy, suggesting that the presence of PL SVC is more likely to be associated with the underlying defect of lateralization rather than mechanical factors (2). In our series, septal defects were the most common cardiac defect, followed by left outflow tract obstruction and conotruncal anomalies. Heterotaxy was observed in only 3 of the 15 cases with cardiac defects being much lower than the other previous prenatal series in which it was reported as 41-54% (2, 6). Atrioventricular septal defects, right outflow tract obstruction and double outlet right ventricle were the most common cardiac anomalies in the cases with heterotaxy in previous series, while the spectrum of cardiac malformations differed in the

![FIG. 2. Transverse view of the fetal chest in a plane slightly caudal to the apical four-chamber view showing a dilated coronary sinus (arrow).](image)

**TABLE 1.** Demographic and clinical characteristics of cases. (Group 1: cases not associated with cardiac defects, Group 2: cases associated with cardiac defects).

<table>
<thead>
<tr>
<th></th>
<th>Group 1 (n=16)</th>
<th>Group 2 (n=15)</th>
<th>Total (%) (n=31)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (mean±SD, year)*</td>
<td>30.1±3.5</td>
<td>29.5±6.1</td>
<td>29.8±4.9</td>
</tr>
<tr>
<td>GAD (mean±SD, week)*</td>
<td>24.1±4</td>
<td>25±5.2</td>
<td>24.5±4.6</td>
</tr>
<tr>
<td>Fetal gender*</td>
<td>8 (50)</td>
<td>9 (60)</td>
<td>17 (54.8)</td>
</tr>
<tr>
<td>Male</td>
<td>5 (31.2)</td>
<td>4 (33.3)</td>
<td>9 (29)</td>
</tr>
<tr>
<td>Female</td>
<td>5 (31.2)</td>
<td>5 (40.0)</td>
<td>10 (31)</td>
</tr>
<tr>
<td>Extracardiac anomalies*</td>
<td>8 (50)</td>
<td>9 (60)</td>
<td>17 (54.8)</td>
</tr>
<tr>
<td>Karyotype*</td>
<td>8 (50)</td>
<td>9 (60)</td>
<td>17 (54.8)</td>
</tr>
<tr>
<td>Normal</td>
<td>13 (81.7)</td>
<td>11 (91.7)</td>
<td>24 (92.3)</td>
</tr>
<tr>
<td>Aneuploidy</td>
<td>1 (6.2)</td>
<td>1 (8.3)</td>
<td>2 (6.5)</td>
</tr>
<tr>
<td>Outcome†</td>
<td>11 (68.7)</td>
<td>10 (12.5)</td>
<td>21 (67.7)</td>
</tr>
<tr>
<td>Favorable</td>
<td>13 (81.7)</td>
<td>15 (12.5)</td>
<td>28 (90.3)</td>
</tr>
<tr>
<td>Unfavorable</td>
<td>8 (50)</td>
<td>9 (60)</td>
<td>17 (54.8)</td>
</tr>
</tbody>
</table>

GAD: gestational age at diagnosis

(1) (Unfavorable outcome includes neonatal death, termination of pregnancy and in utero death)

*Group 1 vs. Group 2 is insignificant (p>0.05)

†Group 1 vs. Group 2 is significant (p<0.009)

**TABLE 2.** Outcome of the cases (except for unborn fetuses)

<table>
<thead>
<tr>
<th></th>
<th>Isolated PL SVC (n=5)</th>
<th>PL SVC and extracardiac defects (n=8)</th>
<th>PL SVC and other cardiac defects (n=6)</th>
<th>PL SVC plus both cardiac and extracardiac defect findings (n=9)</th>
<th>Total (n=28)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Well</td>
<td>5 (100%)</td>
<td>6 (75%)</td>
<td>3 (50%)</td>
<td>2 (22.2%)</td>
<td>16 (57.1%)</td>
</tr>
<tr>
<td>NND</td>
<td>-</td>
<td>-</td>
<td>3 (50%)</td>
<td>4 (44.4%)</td>
<td>7 (25%)</td>
</tr>
<tr>
<td>TOP</td>
<td>-</td>
<td>2 (25%)</td>
<td>-</td>
<td>2 (22.2%)</td>
<td>4 (14.3%)</td>
</tr>
<tr>
<td>IUD</td>
<td>-</td>
<td>-</td>
<td>1 (11.1%)</td>
<td>1 (3.6%)</td>
<td></td>
</tr>
</tbody>
</table>

PL SVC: persistent left superior vena cava; NND: neonatal death; TOP: termination of pregnancy; IUD: in utero death

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other cases (2, 6). The most frequent anomalies were reported to be VSD and coarctation of the aorta in the study by Berg et al. (2), and left outflow tract obstructive diseases and conotruncal anomalies in the study by Galindo et al. (6).

The association of PLSVC with chromosomal anomalies was previously reported in the literature (4, 7). The rate of aneuploidy was 7.7% in our series, similar to the series of Berg et al. (2) and Galindo et al. (6) in which the corresponding rate
was reported to be 9% in each. Kalache et al. (4) recommend-
ed karyotype analysis when the diagnosis of PLSVC with a
dilated coronary sinus was made, based on the possibility of
missing some cardiac anomalies and obscure extracardiac
anomalies suggesting aneuploidy. The case reported by Ka-
lache et al. (4), as well as 12 fetuses with aneuploidy in the
series of Berg et al. (2) and Galindo et al. (6) had additional
prenatal sonographic cardiac findings. Therefore, it was sug-
gested that cardiac defects themselves, but not PLSVC, were
associated with an abnormal karyotype (2, 6). On the other
hand, chromosomal anomalies were present in five of the 15
fetuses with no additional cardiac defects in a recently pub-
lished study by Barea et al. (7). Moreover, one of these cases
in which neonatal karyotyping revealed trisomy 21 had no
associated extracardiac defect (7). In our series, there were
two aneuploidy cases. In the case with trisomy 18, there were
multiple cardiac and extracardiac anomalies which caused the
suspicion of a chromosomal anomaly. In the other case with
trisomy 21, a choroid plexus cyst and thickened nuchal fold
in the second trimester were clues suggesting the presence of
chromosomal anomaly. Nevertheless, the number of cases re-
ported in the literature is not enough to draw the conclusion
that PLSVC should be accepted as a marker for trisomy 21.

The outcome of PLSVC is associated with the other cardiac
and extracardiac findings. In previous studies, all of the isolated
cases had a favorable prognosis, in contrast to the high neonatal
mortality and TOP rates in those associated with cardiac defects.
Similarly, in our cohort, all fetuses with isolated PLSVC did well
after birth, while the survival rate declined to 50% when an
additional cardiac anomaly was observed, and to 22.2% when an
extracardiac finding was also present. These findings support
the suggestion that isolated PLSVC is a benign anomaly which
should prompt a meticulous examination of the fetus to identify
additional cardiac and extracardiac anomalies.

In contrast to the prenatal diagnosis, the detection of PLS-
VC is difficult postnatally in routine transthoracic echocardi-
ographic studies (6). Usually, a normal R SVC is present in addi-
tion to PLSVC, as with all cases in our series. PLSVC has no
hemodynamic effects and does not need treatment; however, in
rare cases with no R SVC, the diagnosis has significant clinical
implications in adulthood. Firstly, this situation may cause diff-
culties in interventions, such as implantation of a transvenous
pacemaker, systemic venous cannulation for cardiopulmonary
bypass and placement of a pulmonary artery catheter for intra-
operative or intensive care unit monitoring (10, 11). Secondly,
itis may cause dysrhythmias in adulthood due to the deformation
of conduction pathways secondary to coronary sinus dilata-
tion. Prenatal diagnosis of the absence of the R SVC may allow
monitoring of coronary sinus dimensions and help to identify
dysrhythmias using echocardiography and Holter recording, as
was recommended by Guarnieri et al. (11).

This study had some limitations such as its retrospective
nature. Also, our study group involved a selected population
that was referred to a tertiary center, and this might have led
to biased results. Furthermore, not all of the patients under-
went karyotyping, and the karyotypes of a few cases remain
unknown. However, despite these limitations, with this study
we underline the significance of the three-vessel view during
routine fetal screening and draw attention to the increasing
incidence of isolated PLSVC cases in clinical practice.

In conclusion, PLSVC is a vascular variation which is as-
associated with cardiac and extracardiac anomalies in the major-
ority of cases. The outcome is significantly worse if PLSVC is
associated with a cardiac defect, but the prognosis is excellent
in isolated cases. Prenatal diagnosis of PLSVC is significant,
because it paves the way for the diagnosis of structural and
chromosomal anomalies as it leads the physician to perform
fetal echocardiography and a detailed anatomic survey.

Ethics Committee Approval: Ethics committee approval was received for this study from the ethics committee of Istanbul Faculty of Medicine.

Informed Consent: Written informed consent was obtained from patients who participated in this study.

Peer-review: Externally peer-reviewed.

Author contributions: Concept - AÇE, RE, İK; Design - AÇE, AY, RE; Supervision - AY, R.H.; Resource - İK, RE; Materials - AÇE, HÇ, MÖ; Data Collection&or Processing - HÇ, MÖ, RE; Analysis&or Interpretation - AÇE, İK; Literature Search - AÇE; Writing - AÇE; Critical Reviews - AY, R.H.

Conflict of Interest: No conflict of interest was declared by the authors.

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REFERENCES


