Prenatal diagnosis and follow-up of fetal cardiac tumors: case series and review of the literature

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ABSTRACT

Background: Cardiac tumors are frequent in the fetal and infant stages, with rhabdomyoma being the most prevalent in fetal life. Its diagnosis has increased due to ultrasound screening. Objective: To report the first Peruvian series of cases with prenatal diagnosis of fetal cardiac tumors in a maternal perinatal institute. Methods: Descriptive study of case series. The database of all fetuses with prenatal diagnosis of cardiac tumors from January 2009 to January 2021 at the Instituto Nacional Materno Perinatal (INMP), Lima, Peru, was reviewed. These patients were followed up by telephone and in two cases echocardiographic control was performed. Results: We found 6 cases of patients diagnosed with cardiac tumors in the last 12 years in the INMP, with reduction in size in the follow-up of all cases and association with tuberous sclerosis in half of them. Conclusions: Cardiac rhabdomyomas represent the most frequent tumors in fetal life. Most of them have in common the partial remission of the tumor. However, the clinical evolution depends on the location of the tumor, size, and its association or not with tuberous sclerosis. For this reason, close follow-up is recommended, especially of the central nervous system.

Key words: Rhabdomyoma, Tuberous sclerosis, Prenatal diagnosis

RESUMEN

Antecedentes. Los tumores cardiacos son frecuentes en la etapa fetal e infantil, siendo el rabdomioma el más prevalente en la vida fetal. Su diagnóstico se ha incrementado debido al cribado por ecografía. Objetivo. Comunicar la primera serie peruana de casos con diagnóstico prenatal de tumores cardiacos fetales en un instituto materno perinatal. Métodos. Estudio descriptivo de serie de casos. Se revisó la base de datos de todos los fetos con diagnóstico prenatal de tumores cardiacos a partir de enero de 2009 hasta enero de 2021 en el Instituto Nacional Materno Perinatal (INMP), Lima, Perú. Estos pacientes fueron seguidos por vía telefónica y en dos casos se les realizó control ecocardiográfico. Resultados. Se halló 6 casos de pacientes diagnosticados con tumores cardiacos en los últimos 12 años en el INMP, con reducción de tamaño en el seguimiento de todos los casos y asociación con esclerosis tuberosa en la mitad de ellos. Conclusions. Los rabdomiomas cardiacos representan los tumores más frecuentes en la vida fetal. La mayoría tienen en común la remisión parcial del tumor. Sin embargo, la evolución clínica depende de la ubicación, tamaño y su asociación o no de esclerosis tuberosa. Por ello es recomendable un seguimiento estricto, especialmente del sistema nervioso central.

Palabras clave. Rabdomioma, Esclerosis tuberosa, Diagnóstico prenatal

INTRODUCTION

Fetal cardiac tumors are infrequent, more so primary cardiac tumors\(^1,2\). According to autopsy findings in children, an incidence of 0.027%-0.08% is described\(^3\) and in fetuses it is 0.14%\(^4\). This incidence has increased tenfold in recent years due to improved prenatal ultrasound diagnosis\(^5\).

Histologically, the most common cardiac tumors are rhabdomyomas, fibromas, teratomas, myxomas, and hemangiomas, which vary in incidence across studies\(^6,7\) and according to life stage of fetuses, neonates and children.
Benign cardiac tumors are the most common in fetuses and children, with rhabdomyoma being the most frequent, representing 60% of primary cardiac tumors\(^{(4)}\). Although it is classified as benign, it can cause adverse outcomes depending on its location, size and presence of arrhythmias\(^{(5)}\). Malignant cardiac tumors represent only 10% of primary tumors\(^{(4)}\).

Cardiac rhabdomyomas were first described in 1862 by Von Recklinghausen. Histologically they are composed of the pathognomonic 'spider cells'\(^{(6)}\). They are most frequently located in the ventricles and ultrasonographically present as oval, nodular, hyperechogenic, well-defined and frequently multiple lesions\(^{(7)}\). In prenatal life and depending on their location, they can be asymptomatic, cause arrhythmias, obstruct the outflow tract, give rise to pericardial effusion, hydrops and death. At birth they are usually asymptomatic. However, there may also be cyanosis, decreased peripheral pulse and auscultation of heart murmurs. Rarely require surgical management\(^{(4,8)}\).

The prognosis of prenatally detected cardiac rhabdomyomas is favorable, since, having completed their fetal somatic growth, hamartomas lose their mitotic potential and undergo apoptosis\(^{(9)}\). The literature describes that cardiac rhabdomyomas tend to regress with time\(^{(10)}\). However, there is an association between cardiac rhabdomyomas and tuberous sclerosis. Thus, between 60-90% of cardiac rhabdomyomas have a diagnosis of tuberous sclerosis, which worsens the prognosis\(^{(5,8,11)}\). Tuberous sclerosis is a genetic syndrome whose initial manifestation is usually cardiac rhabdomyomas. In addition, they are characterized by the presence of generalized hamartomas in other tissues\(^{(12)}\).

In the present study, we describe the clinical and ultrasound characteristics and the evolution of 6 patients with prenatal diagnosis of cardiac rhabdomyoma over 12 years.

**Methods**

The present is a descriptive case series study, in which the database of the National Maternal Perinatal Institute (INMP), Lima, Peru, of fetuses with prenatal diagnosis of cardiac tumors, from January 2009 to January 2021, was reviewed. Data were extracted from maternal and neonatal medical records. Subsequently, the patients were followed up by telephone and echocardiographic control was performed in two of the patients at the institution. In the remaining four cases, data from other hospitals provided by the parents were considered.

**Results**

Ten cases were found, of which 4 were excluded because they did not have adequate postnatal follow-up. The characteristics of the mothers and the prenatal diagnosis of 6 cases of fetuses with cardiac tumors are summarized in Table 1.

The mean age of the pregnant women at diagnosis was 21.8 years (18-26 years) and the mean gestational age was 29 weeks (18-36 weeks). Four (66.7%) were multiple and two (33.3%) were single. All cases presented tumors in the ventricles, with a similar proportion between the right and left ventricle, and in only one case a cardiac tumor was found in the left atrium. In three cases the size of the tumors was greater than 20 mm (Figure 1). In one case the size was not specified, but it was reported to occupy more than

<table>
<thead>
<tr>
<th>Case</th>
<th>Maternal age (years)</th>
<th>Gestational age (weeks)</th>
<th>Location</th>
<th>Classification</th>
<th>Dimensions</th>
<th>Hemodynamic changes</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>18</td>
<td>18</td>
<td>RV, LV</td>
<td>Multiple</td>
<td>RV: 31x37 mm; LV: 10 x 5.6 mm</td>
<td>Mild tricuspid regurgitation</td>
</tr>
<tr>
<td>2</td>
<td>23</td>
<td>36</td>
<td>RV, LV</td>
<td>Multiple</td>
<td>RV: 34 x 20 mm; LV: 12 x 10 mm</td>
<td>No</td>
</tr>
<tr>
<td>3</td>
<td>22</td>
<td>30</td>
<td>LV</td>
<td>Single</td>
<td>8.4 x 4 mm</td>
<td>No</td>
</tr>
<tr>
<td>4</td>
<td>26</td>
<td>25</td>
<td>RV</td>
<td>Single</td>
<td>Occupied 90% of RV</td>
<td>Tricuspid regurgitation</td>
</tr>
<tr>
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<td>19</td>
<td>36</td>
<td>RV, LV</td>
<td>Multiple</td>
<td>RV: 14 x 9 mm; LV: 9 mm</td>
<td>Mild tricuspid regurgitation</td>
</tr>
<tr>
<td>6</td>
<td>23</td>
<td>31</td>
<td>RV, LV, LA</td>
<td>Multiple</td>
<td>RV: 30 mm; LV: 25 mm; LA: 5 mm</td>
<td>Tricuspid regurgitation Pericardial effusion</td>
</tr>
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RV: right ventricle. LV: left ventricle. LA: left atrium.
90% of the right ventricle (Figure 2). The largest was 31 x 37 mm. In four (66.7%) cases there was mild tricuspid regurgitation. All cardiac tumors were classified as rhabdomyomas due to their echographic characteristics.

Among the perinatal outcomes (Table 2), the mean gestational age at birth was 37.7 weeks (37-38 weeks). Five (83.3%) of the newborns were female. The route of delivery was cesarean section in most cases. Only 16.7% of the newborns had low birth weight. All newborns had Apgar greater than 7 at 5 minutes.

In the follow-up, one of the infants required cardiac surgery due to hemodynamic alterations.

We have institutional ultrasound evidence of decrease in tumor volume in two patients. In the rest, indirect information was obtained by telephone follow-up of the parents on the absence of cardiac tumors in the extra-institutional controls.

In 50% of the cases the patients had a diagnosis of tuberous sclerosis; of these, 66.7% had multiple cardiac tumors.

**DISCUSSION**

Cardiac rhabdomyomas are the most frequent fetal tumors, although they are still considered a rare diagnosis. However, their incidence has increased with the more extensive use of ultrasonography and prenatal diagnosis. In our institution 10 cases of cardiac tumors were identified in 12 years of study, but data are available for only 6 of them. All cases had a diagnosis of cardiac rhabdomyoma and 50% were associated with tuberous sclerosis. We have institutional ultrasound evidence of decreased tumor volume in two patients. In the rest of cases, indirect information was obtained through telephone fol-
low-up with the parents regarding the absence of cardiac tumors in extra-institutional controls.

On ultrasonographic evaluation, rhabdomyomas are observed as ovoid, echogenic, unshaded masses that can be located in any of the four cardiac cavities, mainly in the ventricles\(^{12,13}\). In all our cases, rhabdomyomas were found in the ventricles, with equal predominance between both, as indicated in the literature\(^{12}\). In patients with tuberous sclerosis, Jóźwiak et al. found that most were located in the right ventricle (35%), followed by the left ventricle in 22%, interventricular septum in 33%, left and right atrium in 5%\(^{14}\).

The literature reports that 90% of rhabdomyomas are multiple\(^{12}\). Four of our patients (66.7%) had multiple rhabdomyomas. According to what has been reported, 70-90% of rhabdomyomas are associated with tuberous sclerosis. Our case series only found an association in half of the patients, which could mean a lower prevalence of this pathology in our country.

Cardiac rhabdomyomas tend to regress over the years in 90% of cases and do not require surgical management unless they cause severe hemodynamic decompensation. Despite this potential complication, there are currently m-THOR protein inhibitor drugs, such as everolimus, which may help to decrease the size of the tumor\(^{15}\). Dhulipudi et al. performed a prospective observational study in neonates with cardiac rhabdomyomas presenting with clinical symptoms or blood flow obstruction. Five neonates (100%) showed tumor regression during follow-up with everolimus. Therefore, there is potential benefit with the use of this drug in selected symptomatic cases. Multicenter studies are still needed to determine its safety and efficacy in a larger population\(^{16}\).

Cardiac rhabdomyomas are usually asymptomatic. However, depending on their size and location, they may cause obstruction to blood flow, hemodynamic alteration, presence of arrhythmias, pericardial effusion, hydrops and fetal death\(^{8}\). Chao et al. found that a size greater than 20 mm was associated with an increased risk of perinatal death\(^{17}\). Likewise, Okmen F et al. found that cardiac tumor size was associated with perinatal mortality\(^{18}\). In our series, there was no evidence of neonatal death despite the size of the tumor greater than 20 mm. However, 66.7% had mild tricuspid regurgitation on prenatal ultrasound, with only one of them presenting hemodynamic decompensation.

It is important to highlight that there is an association between fetal cardiac rhabdomyomas and tuberous sclerosis in up to 80% of cases. This pathology affects various organs, with the worst prognosis being the involvement of the nervous system, so one of the most frequent symptoms is epilepsy (83%). Early diagnosis and management are important to improve cognitive outcomes\(^{19}\). The fatal complications described are renal angiolioma, sudden unexpected death associated with epilepsy, pulmonary lymphangiolioma, subependymal giant cell astrocytoma and pancreatic cancer\(^{20}\). In our series, 50% of the children had a diagnosis of tuberous sclerosis. Due to this association, a comprehensive evaluation is required when cardiac rhabdomyomas are diagnosed, and magnetic resonance imaging should be performed to look for hamartomas in other locations such as brain, kidney and others, in order to provide counseling to family members and timely multidisciplinary management.

Among the limitations of this presentation, we should point out that 4 patients were lost to follow-up due to migration to another city, genetic study of tuberous sclerosis was not performed on the patients under study due to lack of resources, and some patient data were communicated by telephone interview.

In conclusion, rhabdomyomas represent the most frequent benign cardiac tumors in fetal life; most cases have partial remission of the tumor. However, the clinical course depends on the location, size and its association with tuberous sclerosis. Therefore, a comprehensive and multidisciplinary study is recommended in the fetal stage and close follow-up in childhood.

**References**


