A single-center experience of magnetic resonance imaging findings of fetal sacrococcygeal teratomas

MESUDE TOSUN
İSA ÇAM
HANDE USLU
YASEMİN DOĞAN
YONCA ANIK

Follow this and additional works at: https://journals.tubitak.gov.tr/medical

Part of the Medical Sciences Commons

Recommended Citation
TOSUN, MESUDE; ÇAM, İSA; USLU, HANDE; DOĞAN, YASEMİN; and ANIK, YONCA (2022) "A single-center experience of magnetic resonance imaging findings of fetal sacrococcygeal teratomas," Turkish Journal of Medical Sciences: Vol. 52: No. 4, Article 38. https://doi.org/10.55730/1300-0144.5423
Available at: https://journals.tubitak.gov.tr/medical/vol52/iss4/38

This Article is brought to you for free and open access by TÜBİTAK Academic Journals. It has been accepted for inclusion in Turkish Journal of Medical Sciences by an authorized editor of TÜBİTAK Academic Journals. For more information, please contact academic.publications@tubitak.gov.tr.
A single-center experience of magnetic resonance imaging findings of fetal sacrococcygeal teratomas

Mesude TOSUN1*, İsa ÇAM1, Hande USLU1, Yasemin DOĞAN2, Yonca ANIK1

1Department of Radiology, Faculty of Medicine, Kocaeli University, Kocaeli, Turkey
2Department of Obstetrics and Gynecology, Faculty of Medicine, Kocaeli University, Kocaeli, Turkey

1. Introduction
Sacroccygeal teratomas (SCT) are known as rare tumors, but they are the most common tumor in fetuses and newborns. This study aims to present fetal magnetic resonance imaging (MRI) findings of SCT diagnosed prenatally and compare them with that of the prenatal Ultrasound (US) findings.

Background/aim: Sacrococcygeal teratomas (SCT) are known as rare tumors, but they are the most common tumor in fetuses and newborns. This study aims to present fetal magnetic resonance imaging (MRI) findings of SCT diagnosed prenatally and compare them with that of the prenatal Ultrasound (US) findings.

Materials and methods: Eleven patients diagnosed as SCT prenatally by US and further assessed by MRI are included. MRI was performed via a 1.5-T magnet with a body coil. The presence, size, content extension, and compressive effects of each mass were determined and correlated with US findings and with postnatal examinations, including surgery and pathology. As complications, the presence of ascites and skin edema or pleural or pericardial effusion was diagnosed as hydrops. The amniotic fluid index was calculated for the assessment of oligo- or polyhydramnios.

Results: US findings are found strongly correlated with MRI findings. An agreement on the extent of each mass was observed in eight patients, disagreement in one fetus was an extension of the tumor within the spinal canal recognized only at MR and assessment of intrapelvic-abdominal extension was superior in MRI. There were n = 6 type I, n = 2 type II, n = 1 type III and, n = 2 type IV tumors. MRI was superior to US for detecting displacement of the colon (n = 3), intrapelvic-intraabdominal extension (n = 1), urinary tract complication (n = 2), and intraspinal extension (n = 1).

Conclusion: MRI is found to be superior to US especially in the assessment of intrapelvic and intraspinal extension of the tumor, colonic displacement, and complications.

Key words: Fetal, magnetic resonance imaging, sacrococcygeal teratoma, Altmand type, prenatal, diagnosis

Received: 05.12.2021 • Accepted/Published Online: 07.05.2022 • Final Version: 10.08.2022

This work is licensed under a Creative Commons Attribution 4.0 International License.
for early detection and diagnosis of fetal SCT and is the primary screening method [15,16]. If an SCT is suspected on prenatal US screening, in utero fetal magnetic resonance imaging (MRI) is recommended whenever possible. MRI offers a superior anatomic resolution. Our aim in this study is to present fetal MRI findings as a single-center experience of SCT diagnosed prenatally.

2. Material and methods

2.1. Patients

The University Hospital research ethics committee approved this study (ID 2021-340). Written informed consent was waived for the retrospective analyses by the Institutional Review Board. We conducted a retrospective single-center study at our University Hospital to review all fetal SCT cases suspected before birth and confirmed postnatally between January 2011–2021. Data were collected from the digital patient archiving system (Sectra, PACS). Most cases were submitted with a presumed diagnosis of tumor or fetal hydrops. All pregnant women underwent a detailed fetal anomaly scan using high-resolution US equipment.

2.2. USG and MR examinations

Fetal ultrasonographic evaluations were performed by a perinatologist (at least 10 years of experience) and a radiologist (at least 15 years of experience) experienced in the obstetric US, separately. Fetal growth, exact tumor dimensions, morphology, and location, the presence or absence of additional malformations, and assessment of amniotic fluid volume were evaluated. There was no conflict between the experts in the interpretation of the images. It was agreed that MRI should be performed as advanced imaging to evaluate the extent of SCT and additional anomalies. MRI was performed via a 1.5-T magnet (Gyrosan Intera; Philips medical systems, Eindhoven, The Netherlands) with a body coil. In our center, fetal MRI (1.5 Tesla) is standard for additional imaging (MRI) is recommended whenever possible. MRI offers a superior anatomic resolution. Our aim in this study is to present fetal MRI findings as a single-center experience of SCT diagnosed prenatally.

Eleven patients diagnosed as SCT prenatally by US and further defined by MRI are included. Diagnostic quality MR images were obtained in all patients. All images taken in MR examination were evaluated. US findings are strongly correlated with MRI findings. An agreement on the extent of each mass was observed in eight patients, disagreement in one fetus was an extension of the tumor within the spinal canal recognized only at MR, and assessment of intrapelvic-abdominal extension was superior in MRI. On MRI, sacrococcygeal teratomas (n = 11) were completely cystic in four fetuses, mixed and solid in seven (Table). According to Altman classification, there were n = 6 type I (Figures 1a, 1b, 2a, 2b, and 3), n = 2 type II (Figures 4a, 4b), n = 1 type III (Figures 5a, 5b, 5c) and, n = 2 type IV tumors (Figures 6a, 6b). MRI was superior to US for detecting displacement of the colon (n = 3), intrapelvic-intraabdominal extension (n = 1), urinary tract complication (n = 2), and intraspinal extension (n = 1). Three patients (bilateral n = 2, unilateral n = 1) had pelvicaliectasis in the kidneys on MRI. However, one of these three patients had unilateral renal ectasia, the other kidney was agenetic (Figures 7a, 7b) and there was severe oligohydramnios. In MRI, tumor compression of adjacent pelvic structures could be evaluated. However, pelvic bones and vertebral column were a limitation in evaluating the extent of SCT on sonography. Chiari type 2 malformation was present in one case (9%) of 11 fetuses and type 3 congenital pulmonary airway malformation in other cases (Figures 8a, 8b). These findings observed in MRI were also detected in the sonographic examination.

SCT causes displacement of the bladder by the effect of pelvic mass, hydronephrosis, and large convoluted ureters. The bladder displacement was observed in 3 of 11 fetuses (27.2%), and bilateral hydronephrosis in 2 fetuses...
and unilateral hydronephrosis in 1 fetus on MRI, but only hydronephrosis was detected by sonography. In the patient with unilateral hydronephrosis, no other kidney was observed (unilateral renal agenesis), and oligohydramnios was present. In the pathological examination, the cases were documented as mature (n = 8), immature (n = 2) and mixed teratomas (n = 1).

4. Discussion
In this study, we aimed to evaluate imaging findings and analyze which imaging approaches are most valuable in fetuses diagnosed with prenatal SCT. SCTs are the most common tumors of the fetus and the neonate, carrying a variable prognosis. The majority tend to be benign (~80%). Alpha-fetoprotein (AFP) and beta HCG may be elevated. A pathology-based classification is as: benign (mature) 70%–80% or malignant (immature). Those presenting in older infants tend to have a higher malignant potential and those presenting in utero have a poor prognosis due to complications. It has been reported that fetal mortality is higher if the gestational week at the time of diagnosis is early in SCT [18]. Mass size, the solid component, and the vascularity of the mass are reported to be more important on prognosis than the gestational age at the time of diagnosis [18]. The prognosis of SCTs detected in the prenatal period is to be worse than those detected in the neonatal period which may be due to the fact that larger tumors are more likely to be detected in fetal life.

Table. Clinical characteristics and outcomes in 11 fetuses with SCT.

<table>
<thead>
<tr>
<th>Patient</th>
<th>Gestational week (GW)</th>
<th>Tumor type</th>
<th>Tumor size (cm³)</th>
<th>US evidence for decompensation</th>
<th>Additional findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>21+2</td>
<td>1</td>
<td>15.9</td>
<td>No</td>
<td></td>
</tr>
<tr>
<td>II</td>
<td>23+1</td>
<td>2</td>
<td>33.3</td>
<td>Increased AF</td>
<td></td>
</tr>
<tr>
<td>III</td>
<td>22+1</td>
<td>2</td>
<td>27.7</td>
<td>No</td>
<td></td>
</tr>
<tr>
<td>IV</td>
<td>21+4</td>
<td>4</td>
<td>25</td>
<td>Increased AF</td>
<td>CPAM</td>
</tr>
<tr>
<td>V</td>
<td>26+8</td>
<td>1</td>
<td>14.5</td>
<td>Decreased AF</td>
<td>Renal Agenesis</td>
</tr>
<tr>
<td>VI</td>
<td>19</td>
<td>1</td>
<td>52.8</td>
<td>No</td>
<td></td>
</tr>
<tr>
<td>VII</td>
<td>22+5</td>
<td>1</td>
<td>150</td>
<td>No</td>
<td></td>
</tr>
<tr>
<td>VIII</td>
<td>22+3</td>
<td>1</td>
<td>8</td>
<td>No</td>
<td></td>
</tr>
<tr>
<td>IX</td>
<td>21+5</td>
<td>1</td>
<td>9.5</td>
<td>Increased AF</td>
<td></td>
</tr>
<tr>
<td>X</td>
<td>32</td>
<td>3</td>
<td>66.5</td>
<td>No</td>
<td></td>
</tr>
<tr>
<td>XI</td>
<td>18+5</td>
<td>4</td>
<td>23.5</td>
<td>No</td>
<td>Chiari Type 2</td>
</tr>
</tbody>
</table>

US: Ultrasonography, AF: Amniotic fluid, CPAM: Congenital pulmonary airway malformation

Figure 1. Type I sacrococcygeal teratoma. a. Sagittal T2-weighted, and b. Ultrasound images show mixed cystic and solid lesions originating from the coccyx. An intrapelvic extension is not seen.
Figure 2. Type I sacrococcygeal teratoma. a. Sagittal T2 weighted, and b. Ultrasound images demonstrate a large, mostly cystic, mass lesion arising from the coccyx.

Figure 3. Type I sacrococcygeal teratoma. Sagittal T2 weighted image shows dichorionic-diamniotic twins. There is a mixed solid and cystic lesion originating from the coccyx. The other fetus that had been imaged partially does not have any tumor or any anomaly.

Figure 4. Type II sacrococcygeal teratoma. a. Ultrasound, and b. Coronal T2 weighted images show large septate cystic mass arising from coccyx with a small intrapelvic component.
Figure 5. Type III sacrococcygeal teratoma. a. Sagittal, and b. Axial T2-weighted, c. Ultrasound images show large mixed signal intensity tumor extending into abdomen.

Figure 6. Type IV sacrococcygeal teratoma. a. Sagittal, and b. Axial T2-weighted images show tumor developed entirely in the fetal pelvis. The urinary bladder is displaced and vertical dimension is increased and contours are lobulated with increased thickness. There is also polyhydramnios.

Figure 7. Fetus with type 1 SCT. a. Coronal and, b. axial T2 weighted images show right renal agenesis (blue arrow), left renal hydronephrosis (yellow arrow).
and tumors detected in early pregnancy have greater growth potential [19]. Besides the prognosis of prenatally detected SCT is also related to its content. Predominantly solid and highly vascularized masses demonstrate a worse prognosis than tumors that are mainly avascular and cystic in appearance [20].

Complications include urologic complications (the most common cause of morbidity), high output cardiac failure from AV shunting: which in turn can cause hydrops fetalis, gastrointestinal tract obstruction, compression of underlying nerves leading to urinary/fecal incontinence, anemia, dystocia, tumor rupture. Tumor compression of the bladder outlet caused retention in the urinary system, followed by renal impairment, oligohydramnios, and pulmonary hypoplasia [12], also tumor compression or infiltration of the sacral nerves and intraspinal extension of the tumor.

Differential diagnoses include sacral chordoma, sacral meningocele, terminal myelocystocele, and enteric (tailgut) cyst for cystic types and for type IV lesions also low-lying neuroblastoma, rhabdomyosarcoma, small round blue cell tumor in the sacral region. Treatment is with surgical excision inclusive of coccygectomy with additional chemotherapy for malignant ones.

MRI is widely used in these fetuses, although there are few studies in the literature that it is effective [14,21–23]. Majorly, the US operator-dependent technique is affected by the thickness of the abdominal fat tissue, fetal position, and amniotic fluid volume. In addition, it has limited soft-tissue contrast and decreased visual field compared with that of MRI. On the other hand, MRI has a wide window of vision unaffected by fetal position or insufficient amniotic. It can also provide a better resolution of the tumor tissue and the relationship with the surrounding tissues. After a prenatal lesion was suspected by US, fetal magnetic imaging was used to confirm the diagnosis and evaluate additional anomalies. In terms of fetal imaging, we found that MRI was superior to US in predicting tumor morphology in teratomas. More data are needed to support this finding, but preliminary results from our series show a high correlation between fetal MRI findings, outcome, and histopathology in SCTs. Two parents decided to terminate their pregnancies. Like US, MRI examination is a safe method and does not expose the patient to unnecessary risks [16,24]. The possibility of multiplanar examination and soft-tissue resolution for multisystemic evaluation in the diagnosis of fetal malformations are very helpful.

The limitation of our study is that it was designed retrospectively, US were performed by a perinatologist and a radiologist, separately. Although there was no conflict among our researchers, perhaps consensus would be a better alternative.

5. Conclusion
MRI is found to be superior to US, especially in the assessment of intrapelvic and intraspinal extension of the tumor, colonic displacement, and complications. While MRI contributes to the diagnosis, it also serves as a guide for treatment, birth planning, and counseling. In order to optimize pre and postnatal management for the evaluation of fetal sacrococcygeal teratoma, fetuses with SCT detected by US should be referred to MRI to evaluate tumor size, content, and extent.
References


