ABSTRACT

Purpose: The Korean Association of Pediatric Surgeons (KAPS) performed a nationwide survey on sacrococcygeal teratoma in 2018.

Methods: The authors reviewed and analyzed the clinical data of patients who had been treated for sacrococcygeal teratoma by KAPS members from 2008 to 2017.

Results: A total of 189 patients from 18 institutes were registered for the study, which was the first national survey of this disease dealing with a large number of patients in Korea. The results were discussed at the 34th annual meeting of KAPS, which was held in Jeonju on June 21–22, 2018.

Conclusions: We believe that this study could be utilized as a guideline for the treatment of sacrococcygeal teratoma to diminish pediatric surgeons' difficulties in treating this disease and thus lead to better outcomes.

Keywords: Teratoma; Sacrococcygeal region; Surveys and questionnaires

INTRODUCTION

Since 1991, the Korean Association of Pediatric Surgeons (KAPS) has performed annual nationwide studies, each year addressing a different topic relating to pediatric diseases, and the results of these studies are discussed at each respective annual meeting of KAPS. They are also summarized and published in Advances in Pediatric Surgery, the official journal of the KAPS. The list of study topics is summarized in Table 1. The 34th annual meeting of KAPS was held in Jeonju on June 21–22, 2018, and the topic was sacrococcygeal teratoma, which was discussed for the first time at an annual meeting of KAPS.
METHODS

The authors reviewed and analyzed the clinical data of patients who had been treated for sacrococcygeal teratoma by KAPS members from 2008 to 2017. We used Microsoft Access 2016® (Microsoft, Redmond, WA, USA) for the patient registry and data collection. All the data were analyzed using IBM SPSS version 23 statistical software (IBM Co., Armonk, NY, USA), and p-values <0.05 were considered statistically significant.

RESULTS

1. Demographics

A total of 189 patients with sacrococcygeal teratoma from 18 institutes were registered for the study (Fig. 1). The top 4 institutes treated 73.5% of all the patients during the study period. The patients’ demographics are summarized in Table 2. The disease was found to have occurred predominantly in females (M:F=1:2.71). A total of 37 accompanying malformations were present in 29 patients (15.3%), and 15 of these patients were diagnosed with Currarino syndrome. Most patients (n=137, 72.5%) were diagnosed and underwent surgery during the neonatal period, but 27 (14.3%) patients underwent surgery beyond the age of 1 year. In this

---

Table 1. The list of topics addressed at each annual meeting of the Korean Association of Pediatric Surgeons since 1991

| Year | Topic                                                                 | Topic                                                                 |
|------|----------------------------------------------------------------------|----------------------------------------------------------------------|
| 1991 | Current situation in Korean pediatric surgery                        | 2005<sup>5</sup> Necrotizing enterocolitis                            |
| 1992 | Inguinal hernia                                                      | 2006<sup>6</sup> Acute appendicitis                                   |
| 1993 | Hirschsprung disease                                                | 2007 Prospect of pediatric surgery                                   |
| 1994 | Anorectal malformation                                              | 2008 Inguinal hernia                                                 |
| 1995<sup>a</sup> | Esophageal atresia and tracheoesophageal fistula                      | 2009 Hirschsprung disease                                             |
| 1996<sup>a</sup> | Branchial anomalies                                                | 2010<sup>4</sup> Intestinal atresia                                |
| 1997<sup>a</sup> | Infantile hypertrophic pyloric stenosis                              | 2011<sup>4</sup> Biliary atresia                                    |
| 1998<sup>a</sup> | Intestinal atresia                                                 | 2012 Statistics of pediatric surgery disease                        |
| 1999<sup>a</sup> | Anorectal malformations                                             | 2013<sup>4</sup> Minimally invasive surgery                          |
| 2000<sup>a</sup> | Index cases in pediatric surgery                                    | 2014<sup>4</sup> Newborns surgery with congenital anomalies          |
| 2001<sup>a</sup> | Biliary atresia                                                     | 2015<sup>4</sup> Neonate congenital Bochdalek hernia                 |
| 2002<sup>a</sup> | Choledochal cyst                                                    | 2016 Esophageal atresia with tracheoesophageal fistula               |
| 2003<sup>a</sup> | Congenital posterolateral diaphragmatic hernia                      | 2017 Choledochal cyst                                                |
| 2004 | Trend of pediatric surgery disease                                  | 2018<sup>8</sup> Sacrococcygeal teratoma                             |

<sup>a</sup>These studies were published in Advances in Pediatric Surgery.
study, we compared the results of the neonates who were younger than 29 days with those of the “old-age” group, which comprised patients who were older than 1 year.

2. Preoperative evaluation and treatment
The most common symptom was sacral mass, and 85.6% of the neonates were diagnosed prenatally. Ultrasonography was the most common diagnostic tool in the prenatal period, but magnetic resonance imaging (MRI) was the most popular study after birth. Eleven patients underwent an in-utero procedure. Preoperative serum levels of α-fetoprotein (AFP) were evaluated in 138 patients, and it was the most common tumor marker, followed by β-human chorionic gonadotropin and carcinoembryonic antigen. Almost all the patients underwent surgical treatment without preoperative treatment, with only 8 patients requiring preoperative chemotherapy or embolization. Anterior displacement of the rectum and obstructive hydronephrosis were the most common preoperative complications because of the mass effect of the tumor (Table 3).

3. Operative treatment
Almost all the neonatal patients underwent surgical treatment without delay. Their median age at the time of surgery was 4 days after birth, and the median body weight was 3.0 kg. Most of the patients underwent surgery once, but 22.2% required an operation more than one time. The perineal approach as a surgical method was so common that 93.7% of the operations were performed using only a perineal approach. A total of 84.1% of the patients underwent complete excision of the tumor, and 21 patients had 24 intraoperative complications among them. Of these, intraoperative bleeding was the most common complication, followed by cerebrospinal fluid leakage. Postoperatively, 15.9% had complications. Problems with the wound constituted the most common postoperative complication and the most common cause of reoperation during the postoperative hospitalization period (Table 4).
4. Tumor characteristics

The maximal diameters of the tumors in the neonate group were significantly larger than those in the old-age group (7.9±5.1 vs. 4.2±4.8 cm, p<0.001). Using the Altman classification, the most common tumor type was type I in the neonatal group, but type IV was the most common type in the old-age group (p<0.001). Mature and cystic or predominantly cystic mixed type were the main characteristics of the tumors. In this study, we found that the pathologic reports of 12 patients did not belong to teratoma or germ cell tumors, but we included them in this study because their clinical features were more compatible with sacrococcygeal teratoma. A total of 11 patients required postoperative chemotherapy (Table 5).

5. Postoperative treatment and follow-up

Postoperative follow-up was available at a median age of 41 months (Table 6). Most of the patients underwent follow-ups at less than a one-year interval. Ultrasonography and MRI were the most common evaluation methods for follow-up. During the follow-up period, 39 patients had tumor recurrence. The pathological characteristics of the recurrent tumors are described in Table 7. Thirty-three patients underwent excision or excision with chemotherapy. Long-term complications were found in 44 patients, and most of them correlated with the function of defecation.

---

### Table 3. Preoperative evaluation and treatment

| Characteristic                              | Total  | Neonate | Old-age |
|--------------------------------------------|--------|---------|---------|
| Clinical presentation<sup>a</sup>          | (n=182)| (n=132) | (n=26)  |
| Abdominal pain/distension                  | 13 (7.1)| 5 (3.8) | 2 (7.7) |
| Mass                                       | 145 (79.7)| 120 (90.9)| 11 (42.3)|
| Constipation                               | 10 (5.5)| 0       | 5 (19.2)|
| No symptoms                                | 9 (4.9)| 7 (5.3) | 1 (3.8) |
| Other                                      | 14 (7.7)| 1 (0.8) | 9 (34.6)|
| Prenatal diagnosis in neonate<sup>b</sup>  | 113/135 (85.6) |
| Prenatal US                                | 113    |
| Prenatal MRI                               | 3 (2.2)|
| In-utero procedure                         | 11/135 (8.1) |
| RFA                                        | 8      |
| Aspiration                                 | 2      |
| Cystic-amniotic shunt                      | 1      |
| Diagnostic work-up after delivery<sup>b</sup>| (n=182)| (n=132) | (n=26)  |
| US                                         | 105 (57.7)| 83 (62.9)| 10 (38.5)|
| CT                                         | 16 (8.8)| 9 (6.8) | 5 (19.2)|
| MRI                                        | 159 (87.4)| 112 (84.8)| 24 (92.3)|
| Preoperative biopsy                         | 4 (2.2)| 0       | 4 (15.4)|
| Other                                      | 3 (1.6)| 2 (1.5) | 1 (3.8) |
| Pre-op AFP (ng/mL)                         | (n=138)| (n=111) | (n=14)  |
| Mean±SD                                    | 87,658±77,076 | 20,287±42,396 | 64,745 (0.8–600,000) | 2.1 (0.8–150,730) |
| Preoperative treatment                     | (n=189)| (n=137) | (n=27)  |
| Preoperative chemotherapy                  | 6      | 6       |
| Preoperative embolization                   | 1      | 1       |
| Preoperative chemotherapy and ASCT<sup>b</sup>| 1      | 1       |
| Preoperative complications                  | (n=189)| (n=137) | (n=27)  |
| Yes                                        | 30 (15.9)| 26 (19.0)| 1 (3.7) |
| No                                         | 159 (84.1)| 111 (81.0)| 26 (96.3)|

Values are presented as number (%).
US, ultrasound; MRI, magnetic resonance imaging; RFA, radiofrequency ablation; CT, computed tomography; AFP, α-fetoprotein; SD, standard deviation; ASCT, autologous stem cell transplantation.
<sup>a</sup>Including multiple selection; <sup>b</sup>Autologous stem cell transplantation.
Currently, among the 189 patients, 1 death had occurred in a case of incomplete excision, 136 patients are living without tumors, 19 are living with tumors, and 33 patients have been lost to follow-up (Fig. 2).

6. Questionnaire for sacrococcygeal teratoma

The questionnaire for sacrococcygeal teratoma consisted of 7 questions, and 22 regular members of KAPS answered them. The following are the questions and the number of answers for each item (Table 8).
Studies on the clinical characteristics of and strategies used to treat sacrococcygeal teratoma are not rare, but its low incidence usually makes it challenging for pediatric surgeons to accumulate experience in this disease. Previous studies about this disease in Korea have been limited, and only a few studies have been published [1-4]. The significance of this study is that it is the first nationwide survey on sacrococcygeal teratoma in Korea and includes a significant number of patients. Our study showed excellent results after surgical treatment. There was only 1 reported case of death after surgery, and the number of long-term functional complications was not high.

Table 5. Tumor characteristics

| Characteristic                      | Total (n=189) | Neonate (n=137) | Old-age (n=27) |
|------------------------------------|---------------|-----------------|----------------|
| Pathology of tumor                 |               |                 |                |
| Mature teratoma                    | 138 (73.0)    | 103 (75.2)      | 16 (59.3)      |
| Immature teratoma                  | 33 (17.5)     | 30 (21.9)       | 2 (7.4)        |
| Grade 1                            | 4 (13.8)      | 4 (14.8)        | 0 (0)          |
| Grade 2                            | 10 (34.5)     | 8 (29.6)        | 1 (100)        |
| Grade 3                            | 15 (51.7)     | 15 (55.6)       | 0 (0)          |
| Grade unknown                      | 4 (13.8)      | 3               | 1              |
| Mixed                              | 4 (2.1)       | 2 (1.5)         | 0              |
| Malignant (yolk sac)               | 2 (1.1)       | 1 (0.7)         | 1 (3.7)        |
| Other*                             | 12 (6.3)      | 1 (0.7)         | 8 (29.6)       |
| Largest tumor length (cm)          | 6.8±4.8       | 7.9±5.1         | 4.2±4.8        |
| Type of tumor component            |               |                 |                |
| Cystic type                        | 63 (34.1)     | 43 (31.6)       | 9 (37.5)       |
| Predominantly cystic mixed type    | 61 (33.0)     | 48 (35.3)       | 6 (25.0)       |
| Predominantly solid mixed type     | 32 (17.3)     | 29 (21.3)       | 1 (4.2)        |
| Solid type                         | 29 (15.7)     | 16 (11.8)       | 8 (33.3)       |
| Unknown                            | 4             | 1               | 2              |
| Altman classification              |               |                 |                |
| I                                  | 68 (36.0)     | 53 (38.7)       | 8 (29.6)       |
| II                                 | 57 (30.0)     | 47 (34.3)       | 2 (7.4)        |
| III                                | 25 (13.2)     | 21 (15.3)       | 1 (3.7)        |
| IV                                 | 45 (23.8)     | 16 (11.7)       | 16 (59.3)      |

Values are presented as number (%) or mean±standard deviation.

*Epidermal cyst (2), lipoma (2), lipoblastoma (2), duplication cyst (1), solitary fibrous tumor (1), infantile fibrosarcoma (1), lipomyelomenigocele (1), lymphangioma (1), epithelioid hemangioendothelioma (1).

Fig. 2. Summary of treatments and prognoses.
SCT, sacrococcygeal teratoma; F/U, follow-up.
Although not all the patients were followed-up for a long time, it is likely that the mortality of those who were lost to follow-up was not affected because the pediatric surgeons would probably have treated them if they had experienced any problems. These results are similar to or better than those of previously published studies [5-10].

| Table 6. Postoperative treatment and follow-up |
|-----------------------------------------------|
| Characteristic                                | Total (n=189) | Neonate (n=137) | Old-age (n=27) |
| Postoperative follow-up (mo)                  |              |                 |                |
| Age at last follow-up                         | 47.9±38.9 (median 41, range 0–243) |
| Postoperative follow-up                       | 37.9±27.6 (median 34, range 0–112) |
| Follow-up method(5)                           |              |                 |                |
| US                                            | 89 (47.1)    | 66 (48.2)       | 13 (48.1)      |
| CT                                            | 23 (12.2)    | 17 (12.4)       | 4 (14.8)       |
| MRI                                           | 98 (51.9)    | 77 (56.2)       | 10 (37.3)      |
| PET                                           | 5 (2.6)      | 3 (2.2)         | 2 (7.4)        |
| Other                                         | 4 (2.1)      | 3 (2.2)         | 0 (0)          |
| Follow-up interval (mo)                       |              |                 |                |
| 1–3                                           | 25           |                 |                |
| 4–6                                           | 30           |                 |                |
| 7–12                                         | 37           |                 |                |
| >12                                          | 14           |                 |                |
| Tumor recurrence                              | 39 (20.6)    | 28              | 5              |
| Detection of tumor recurrence                 |              |                 |                |
| Physical examination                          | 1 (2.6)      | 1 (3.6)         | 0 (0)          |
| Elevated tumor marker                         | 3 (7.7)      | 3 (10.7)        | 0 (0)          |
| MRI                                           | 27 (69.2)    | 20 (71.4)       | 5 (100)        |
| U/S                                           | 7 (17.9)     | 4 (14.3)        | 0 (0)          |
| Other                                         | 1 (2.6)      | 0 (0)           | 0 (0)          |
| Treatment of tumor recurrence                 |              |                 |                |
| Excision                                      | 21 (53.8)    | 14 (50.0)       | 3 (60.0)       |
| Excision+CTx                                  | 12 (30.8)    | 9 (30.3)        | 1 (20.0)       |
| CTx only                                      | 2 (5.1)      | 1 (3.6)         | 1 (20.0)       |
| Observation                                   | 2 (5.1)      | 2 (7.1)         | 0 (0)          |
| Other                                         | 2 (5.1)      | 2 (7.1)         | 0 (0)          |
| Long-term functional complications(5)         | 44           | 21              | 7              |
| Constipation                                  | 21           |                 |                |
| Soiling                                       | 14           |                 |                |
| Urinary incontinence                          | 7            |                 |                |
| Lower extremity weakness                      | 5            |                 |                |
| Other                                         | 7            |                 |                |

Values are presented as mean±standard deviation or number (%).
US, ultrasound; CT, computed tomography; MRI, magnetic resonance imaging; PET, positron emission tomography; CTx, chemotherapy.
(5)Including multiple selection.

| Table 7. Recurrent tumor pathology |
|-----------------------------------|
| Characteristic                    | Total (n=33) | Neonate (n=23) | Old-age (n=4) |
| Mature                            | 19 (57.6)    | 15 (78.9)      | 1 (25.0)      |
| Original pathology               |              | Mature (10), immature (8), mixed (1) |
| Immature                         | 2 (6.1)      | 2 (8.7)        | 0 (0)         |
| Original pathology               |              | Mature (1), immature (1) |
| Malignant (yolk sac)             | 6 (18.2)     | 3 (13.0)       | 1 (25.0)      |
| Original pathology               |              | Mature (3), immature (1), yolk sac (2) |
| Mixed                            | 1 (3.0)      | 1 (4.3)        | 0 (0)         |
| Original pathology               |              | Mature (1)     |
| Other(6)                         | 5 (15.2)     | 2 (8.7)        | 2 (50.0)      |
| Original pathology               |              | Mature (3), immature (1), epithelioid hemangioendothelioma (1) |

(6)Epithelioid hemangioendothelioma (1), inflamed granulation tissue (1), lipomeningomyelocele (1), lipoma (1), no evidence of residual teratoma but r/o recurred tumor on follow-up magnetic resonance imaging (1).
This study revealed that this incidence of this disease can be divided into 2 age groups. In the one group, the lesions were detected prenatally, and in the other, the disease was late-onset. Although the incidence of the old-age group was not high, and the prognosis of this group after surgical treatment was also excellent, pediatric surgeons must consider the possibility of this disease in old age.

One of the limitations of this study was that we evaluated only the patients who underwent surgical treatment, and therefore not all patients with sacrococcygeal teratoma were included. The prognosis of sacrococcygeal teratoma would have deteriorated in patients who could not undergo surgical treatment because of their poor general condition or the inoperability of the tumor.

Through the questionnaire, we were able to learn the current status of KAPS members' clinical practices with regard to sacrococcygeal teratoma. Most of the members take care of their patients themselves postoperatively, and they prefer a long-term follow-up of about
5 years. In rare cases, a few members tried to treat patients using laparoscopic surgery or robotic surgery.

We believe that this study could be utilized as a guideline for the treatment of sacrococcygeal teratoma to diminish pediatric surgeons’ difficulties in treating this disease and thus lead to better outcomes.

REFERENCES

1. Choi SH, Hwang EH. Clinical analysis of sacrococcygeal teratoma. J Korean Surg Soc 1985;29:697-702.
2. Gong CS, Kim SC, Kim DY, Kim IK, Namgung JM, Hwang JH, et al. The outcomes of treatment for sacrococcygeal teratoma: the 24-year experiences. J Korean Assoc Pediatr Surg 2013;19:81-9.
3. Jung SE, Bang HY, Mok WK, Lee SC, Park KW, Kim WK. Sacrococcygeal teratoma in infants and children - Problems according to diagnosis and timing of operation -. J Korean Surg Soc 1995;48:127-32.
4. Kim JG, Lee J, Moon IS, Lee MD, Park WB, Chun CS, et al. A clinical analysis of sacrococcygeal teratoma. J Korean Surg Soc 1989;37:224-31.
5. Derikx JP, De Backer A, van de Schoot L, Aronson DC, de Langen ZJ, van den Hoonaard TL, et al. Long-term functional sequelae of sacrococcygeal teratoma: a national study in The Netherlands. J Pediatr Surg 2007;42:1122-6.
6. Draper H, Chitayat D, Ein SH, Langer JC. Long-term functional results following resection of neonatal sacrococcygeal teratoma. Pediatr Surg Int 2009;25:243-6.
7. Ho KO, Soundappan SV, Walker K, Badawi N. Sacrococcygeal teratoma: the 13-year experience of a tertiary paediatric centre. J Paediatr Child Health 2011;47:287-91.
8. Partridge EA, Canning D, Long C, Peranteau WH, Hedrick HL, Adzick NS, et al. Urologic and anorectal complications of sacrococcygeal teratomas: prenatal and postnatal predictors. J Pediatr Surg 2014;49:139-42.
9. Sayed HA, Ali AM, Hamza HM, Mourad AF, Elsayeb AA. Sacrococcygeal tumors: clinical characteristics and outcome of pediatric patients treated at South Egypt Cancer Institute. A retrospective analysis. J Pediatr Surg 2013;48:1604-8.
10. Usui N, Kitano Y, Sago H, Kanamori Y, Yoneda A, Nakamura T, et al. Outcomes of prenatally diagnosed sacrococcygeal teratomas: the results of a Japanese nationwide survey. J Pediatr Surg 2012;47:441-7.