

Surgical Outcomes of Pediatric Patients with Asymptomatic Tethered Cord Syndrome

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Study Design: A retrospective cohort study.

Purpose: To examine the validity of prophylactic surgery for children with tethered cord syndrome (TCS).

Overview of Literature: Prophylactic surgery for pediatric patients with TCS remains controversial.

Methods: We retrospectively analyzed the surgical outcomes of 14 children (nine boys and five girls) with asymptomatic TCS who were surgically treated at Hokkaido University Hospital between 1989 and 2015.

Results: The median age at the time of initial surgery for asymptomatic TCS was 28.6 months (range, 0–66 months). The median final follow-up period was 142 months (range, 7–232 months). Of the 14 children with asymptomatic TCS, 12 had lumbosacral lipoma and two had meningocele. According to the classification of spinal lipoma, two children had dorsal type, four had caudal type, two had transitional type, and four had filar type. There were no children with lipomyelomeningocele. All children were free of neurological symptoms until 94 months after the initial surgery. Subsequently, one child exhibited delayed neurological deficits and underwent a second surgery because of motor and sensory disturbances; slight sensory disturbance was noted at the final follow-up examination. Another child later showed bowel and bladder dysfunction. However, a second surgery was not performed for this child because his motor and sensory functions were normal; hence, we chose to avoid nerve injury in the case of dissecting adhesion.

Conclusions: All 14 children with asymptomatic TCS were free of neurological symptoms until 94 months after the initial surgery. However, two children exhibited delayed neurological deficits at 94 months and 177 months. We believe that prophylactic surgery for asymptomatic TCS is effective for a certain period. However, because the natural history of TCS is poorly understood, strict follow-up after surgery is necessary.

Keywords: Child; Lipoma; Spinal dysraphism; Prophylactic surgery; Outcome

Introduction

Tethered cord syndrome (TCS) is the clinical manifestation of an abnormal stretch of the spinal cord and is presumed to cause mechanical injury, a compromised blood supply, and altered spinal cord metabolism. TCS is associated with neurological deficits in the lower limbs, leg

numbness, lower back pain, foot deformities, and bowel and bladder disturbances [1,2]. It is unlikely that symptoms will sufficiently improve if surgery is performed after the onset of neurological symptoms associated with TCS. Hence, prophylactic surgery for asymptomatic TCS is advocated and performed. In the present study, the surgical outcomes of children with asymptomatic TCS were

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retrospectively analyzed, and the validity of prophylactic surgery for asymptomatic TCS was examined.

Materials and Methods

Between 1989 and 2015, 14 children with asymptomatic TCS underwent surgical treatment at Hokkaido University Hospital. Patients with presacral lipomas, other spinal dysraphisms (such as split cord malformations), or severe neuro-orthopedic deformities (such as clubfoot or worsening scoliosis) were excluded from this study. We explained the surgical procedure and obtained prior in-

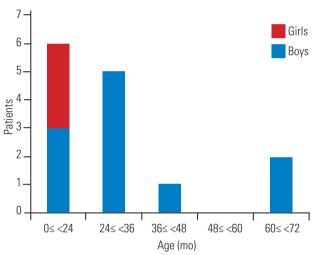


Fig. 1. Age (in months) of patients with asymptomatic tethered cord syndrome at the initial surgery.

formed consent from all patients. The surgical outcomes of these children were retrospectively reviewed.

Results

All 14 children (nine boys and five girls) with asymptomatic TCS underwent prophylactic surgery to release the tethered conus. The median duration of follow-up was 142 months (range, 7-232 months).

The median age at the time of the initial surgery was 28.5 months (range, 0-66 months) (Fig. 1). Six children underwent the initial surgery when aged <24 months, which was the most common age period. Subsequently,

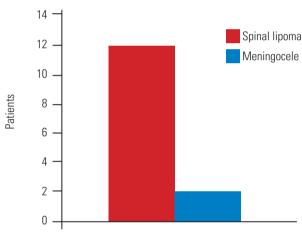


Fig. 2. Lumbosacral spine lesion in patients with asymptomatic tethered cord syndrome. Twelve patients had lumbosacral lipoma and two had meningocele.

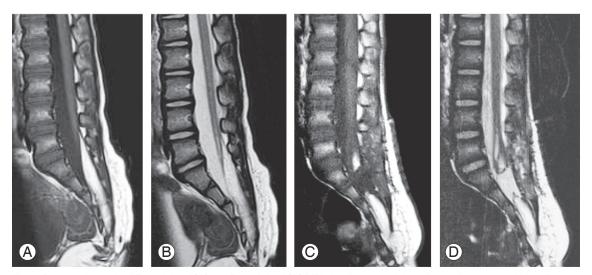


Fig. 3. Preoperative sagittal T1-weighted (A) and T2-weighted (B) MRI showing a transitional lipoma in a 2-year-old boy. Postoperative sagittal T1-weighted (C) and T2-weighted (D) MRI performed 1 month after surgery showing partial removal of the lipoma and forward movement of the spinal cord. MRI, magnetic resonance imaging.

five children underwent prophylactic surgery when aged between 24 and 36 months. Two children underwent surgery when aged >60 months.

Of the 14 children with asymptomatic TCS, 12 had lumbosacral lipoma and two had lumbosacral meningocele (Fig. 2). The classification of spinal lipoma was dorsal type in two children, caudal type in four, transitional type in two, and filar type in four. There were no children with lipomyelomeningocele. The level of conus medullaris determined via magnetic resonance imaging (MRI) was L2-3 in one child, L3-4 in two, L4-5 in one, L5 in two, S1 in three, S2 in two, and S3 in three. Bipolar stimulation and continuous electromyography were performed for all patients during surgery. Although intraoperative evoked potential monitoring became available only in the last part of the present series, we did not use it because it may trigger epileptic seizures. Treatment consisted of surgical untethering of the spinal cord, partial resection of the spinal lipoma (in case of lipoma), and creation of subarachnoid space, where possible. Next, the dural sac was constructed by direct dural repair using GORE-TEX threads (W. L. Gore and Associates Inc., Newark, DE, USA). The extradural space was sealed using fibrin glue (Beriplast P Combi-Set Tissue Adhesion; CSL Behring, King of Prussia, PA, USA). Afterward, the superficial layers were reconstructed in the usual manner. Fig. 3 shows preoperative and postoperative MRIs for a representative patient.

All children were free of neurological symptoms until 94 months after surgery (Fig. 4). Subsequently, two boys exhibited delayed neurological deficits: one had caudal type spinal lipoma at 94 months, and the other had dorsal type spinal lipoma at 177 months. One child underwent a second surgery because of deterioration of neurological function exhibiting as motor and sensory disturbances. Slight sensory disturbance was noted at the final follow-up examination. The other child showed bowel and bladder dysfunctions; however, a second surgery was not

performed because his motor and sensory functions were normal, and we thought it best to avoid nerve injury in the case of dissecting adhesion (Table 1).

Discussion

It is currently unclear whether prophylactic surgery for children with asymptomatic TCS can prevent the development of future neurological deficits. Regarding the natural course of asymptomatic TCS, conversion to symptomatic TCS is reported at a rate of 3%-4% per year [3-5]; hence, it seems natural to consider preventive surgery. However, several studies investigating the success of early preventive surgery have reported conflicting outcomes. Kulkarni et al. [3] performed conservative therapy in 53 patients with lipomas of the spinal cord conus. During the 9-year follow-up period, there was no significant difference in the appearance of neurological symptoms between the conservatively treated and early surgery groups. However, the expression rate of delayed neurological deficits was lower in the conservative therapy group (33% in the conservative therapy group versus 46% in the early surgery group). Wykes et al. [5] performed conservative treatment for 56 patients with asymptomatic lipomas of the spinal

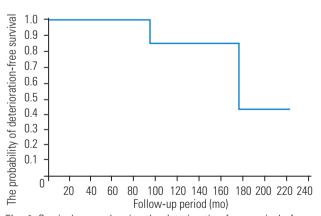


Fig. 4. Survival curve showing the deterioration-free survival of patients who underwent prophylactic surgery.

Table 1. Patients with delayed neurological deficits

Sex/age at initial surgery (mo)	Type of spinal lipoma	Delayed neurological deficits (mo)	Symptom	Age at 2nd surgery (mo)	Final F/U	Motor or sensory disorder at F/U	BB at F/U
M/16	Caudal	94	BB	None	195	None	Continue
M/7	Dorsal	177	Motor and sensory deficits	184	232	Sensory	None

M, male; BB, bowel and bladder dysfunction; F/U, follow-up.

cord conus with 71% of patients being asymptomatic during the mean follow-up period of 5.9 years. Furthermore, the hazard ratio of becoming symptomatic was 40% in 10 years. Both groups had a negative opinion of prophylactic surgery for asymptomatic TCS. Regarding delayed neurological deficits after prophylactic surgery for asymptomatic TCS, Arai et al. [6] reported that three (8.3%) out of 36 cases exhibited delayed neuropathy 3-6 years after surgery. In these patients, no lesions, such as a re-tethered cord or syringomyelia, were detected on MRI. Hence, the authors considered dysplasia of the lower spinal cord to be involved with the onset of symptoms. In contrast, La Marca et al. [7] reported that 28 patients with asymptomatic filum terminale lipoma who underwent prophylactic surgery exhibited no delayed neurological deficits (mean follow-up period, 3.4 years). However, of 71 patients, nine (12.7%) with asymptomatic spinal cord conus lipomas exhibited delayed neurological deficits after prophylactic surgery. Talamonti et al. [4] performed prophylactic surgery (mean follow-up period, 9.7 years) or conservative therapy (mean follow-up period, 10.4 years) for 56 patients with asymptomatic spinal conus lipomas. During the follow-up period, delayed neurological deficits were observed in 9.7% of patients who underwent prophylactic surgery and in 29.1% of those who underwent conservative therapy. Although there was no statistically significant difference between both groups, the results suggested that prophylactic surgery is useful for asymptomatic spinal cord conus lipomas (p=0.067). In our study, the results of prophylactic surgery for 14 children with asymptomatic TCS revealed that all children remained asymptomatic until 94 months after the initial surgery. However, two children (14.3%) exhibited delayed motor and sensory disturbances or bowel and bladder dysfunctions due to a re-tethered cord. Of these children, one underwent reuntethering surgery.

Although the small patient population is a limitation of our study, the results seem acceptable given the annual incidence of delayed neurological deficits is 3.7%-4.0% as reported by Kulkarni et al. [3] and Wykes et al. [5] (annual deterioration rate in our study, 1.8%). Kanev and Bierbrauer [8] retrospectively analyzed children with lipomeningocele and showed that the incidence of neurological symptoms increased with increasing age. Regarding the surgical timing for asymptomatic TCS, Seki et al. [9] reviewed 31 pediatric patients with TCS and reported that patients with asymptomatic TCS underwent surgery

significantly earlier than patients with symptomatic TCS (p=0.045). In addition, Hoffman et al. [10] examined 97 patients with lipomyelomening ocele and found that 62.5% of patients aged >6 months who underwent prophylactic surgery remained asymptomatic, whereas 70.7% of those aged >6 months who underwent prophylactic surgery became symptomatic. Taken together, these reports suggested that the age of the patient during prophylactic surgery for TCS is a risk factor for the onset of neurological symptoms. Furthermore, the age of the patient should be carefully considered during prophylactic surgery for asymptomatic TCS.

Conclusions

We performed prophylactic surgery for 14 children with asymptomatic TCS. All children remained asymptomatic until 94 months after the initial surgery. However, two patients later exhibited delayed neurological deficits. We believe that prophylactic surgery for asymptomatic TCS is effective for a certain period; however, because the natural history of asymptomatic TCS remains unknown, strict follow-up after surgery is necessary.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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