A rare case of type 1 C split cord malformation with single dural sheath

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ABSTRACT
Split cord malformation (SCM) is a rare congenital anomaly in which the cord is split over a portion of its length to form double dural tubes (SCM type I) or two hemicords in a single dural sheath (SCM type II). Dachling Pang classified SCM into 2 types with type I SCM consisting of two hemicords, each contained within its own dural sheath and separated by rigid osseocartilaginous median septum. We report a rare case of SCM type 1 c in which there was a single dural sheath. Only one case of single dural sheath in type 1 SCM has been reported in English literature till now.

Key words: Hemicord, pang’s classification, split cord malformation.

Introduction
Split cord malformation (SCM) is a rare congenital anomaly in which the cord is split over a portion of its length to form double dural tubes (SCM type I) or two hemicords in a single dural sheath (SCM type II). Dachling Pang classified SCM into 2 types with type I SCM consisting of two hemicords, each contained within its own dural sheath and separated by rigid osseocartilaginous median septum. We report a rare case of SCM type 1 c in which there was a single dural sheath. Only one case of single dural sheath in type 1 SCM has been reported in English literature till now.

Case Report
A 10-month-old girl presented to us with a tuft of hair over the lower back since birth. She was moving her lower limbs less as compared to her upper limbs. Bladder and bowel functions were not impaired. Examination revealed tuft of hair in midline in the lumbar region. She also had scoliosis with curvature towards the left side. Her head size was normal for her age. Noncontrast Computed tomography spine revealed bony spur at the level of L3 vertebral body [Figure 1] and it was directed superiorly [Figure 2]. Magnetic resonance imaging scan was done which showed spinal dysraphism in the lumbosacral region with split cord malformation type 1 with two hemicords separated by a spur [Figure 3 and 4]. Cord was ending at L3 [Figure 3].

D 12 to L 4 laminotomy was done. Intraoperatively split cord was extending downwards from L 1. There was bony septum arising from posterior surface of L 3 vertebral body and it was ascending up to L 2 level. There was a single dural sac and the bony septum was indenting the dural sac from anterior aspect, not dividing the posterior dura into two sheaths (contrary to what is seen usually in Type I split). The conus was split into two halves by the septum and there was a single filum. The septum was seen at the lower part of split only, making it type I C split (as per the classification given by Mahapatra and Gupta). The septum was excised and both the dural envelopes were opened and sutured into one sheath. The filum was thickened and detethering was also done. The patient developed weakness in both her lower limbs in the post-operative period. She was given methylprednisolone intravenous for 48 h and at the time of discharge (post-operative day 7), she had recovered power in both lower limbs up to the preoperative level.

Discussion
Ollivier in 1837, first described a case of diplomyelia. He used this term to describe an abnormality of the spinal cord, in which the dura was separated by a bone spur or rigid fibrous band to create 2 sleeves, each containing a portion of spinal cord divided sagittally into 2 parts. Bruce et al. used the term diastematomyelia to describe a spinal cord split by a midline bony spur, reserving the term ‘diplomyelia’ for a true doubling of the spinal cord, without a spur. Feller and Stenberg, first implicated a notochordal cleft, caused by persistence of a

Several authors have proposed theories to explain the genesis of split cord malformation. Gardner[8] proposed the hydromyelic theory, advocating rupture of an intact neural tube by cerebrospinal fluid as the cause of spina bifida defects. Primary mesodermal abnormality theory was advocated by Lichtenstein.[9] Hendrick[10] proposed on the accessory neurenteric canal hypothesis.

Initially, diplomyelia and diastematomyelia were thought to be two different entities. Diplomyelia was thought to be a true duplication of the spinal cord at certain segments[11,12] and the diastematomyelia was thought to be caused by the bony spur, which suggested that it resulted from meso‑dermal invasion of the neural tube.[13,14] This was effectively disproved by Pang et al.,[1] and Pang.[15] They proposed a unified theory of embryogenesis and advocated a new classification, recommending the term ‘split cord malformation’ for all double spinal cords. The unified theory proposes that all split cord malformations originate from one basic ontogenetic error, occurring around the same time, when the primitive neurenteric canal closes. This basic error leads to the formation of an ‘accessory neurenteric canal’ through the midline embryonic disc that maintains communication between yolk sac and amnion, and enables a contact between ectoderm and endoderm within the canal depending on the timing of the formation of the endomesenchymal tract and subsequent mesenchymal infiltration. Thus the neural tube would split into 2 separate components with an intervening fibrocartilaginous or bony septum (Type 1 SCM), or remain a single dural tube with a split cord by fibrous tissues (Type 2 SCM). This would also explain the presence of commonly associated spinal abnormalities such as dermal sinus tracts, spinal lipomas, dermoids, neurenteric cysts, and even a meningocele or myelomeningocele, as these abnormalities can arise from various ectodermal or endodermal remnants.

Pang et al.,[1] and Pang[15] divided SCM into two types. Type I SCM consists of two hemicords, each contained within its own dural sheath and being separated by a rigid osseocartilaginous median septum. A type II SCM consists of two hemicords housed in a single dural sheath being separated by a non‑rigid, fibrous median septum.
Mahapatra and Gupta\textsuperscript{[16]} in 2005, based on their intraoperative findings in regards to the level of the spur and its relation to the split, proposed a subclassification of the existing Pang classification of primary SCM type I (a-d): Ia, a bone spur in the center with an equally duplicated cord above and below the spur; Ib, a bone spur at the superior pole of the split with no space above it and a large duplicated cord lower down; Ic, a bone spur of the lower pole with a large duplicated cord above; and Id, a bone spur straddling the bifurcation with no space above or below the spur. According to these two classification systems, our case conforms to type I c but with single dural tube.

Our case is unique in many respects, firstly there was a single dural sheath containing two hemicords (conus) with a bony septum indenting the single dural sac, contrary to two dural sacs seen in type I split cord malformations. Other unique feature of our case was that the bony septum was attached to the posterior surface of L3 vertebra and was extending superiorly up to L2 level without any attachment to L1 or L2 vertebral body.

Findings of our case are not explained by the classification proposed by Pang \textit{et al.}, as there was a single dural sheath containing two hemicords, with a bony septum indenting the single dural sac, contrary to two dural sacs seen in type I split cord malformations. Only one case of type I SCM with single dural sheath is reported in English literature to the best of our knowledge, that too by the senior author of this paper (Mahapatra).\textsuperscript{[2]} In previous patient, authors reported a long-segment type 1 split cord malformation, with two-level split cord malformation and a single dural sac at the lower split. There was a bony spur arising from the D 11 body level without a separate dural covering but with 2 cords. The split of the cord extended from D 5 to D 11 level.

**Conclusion**

Unified theory of embryogenesis is the most common accepted theory for split cord malformation, however it does not explain type 1 split with single dural sheath. One should be ready to anticipate such findings and avoid injury to the cord during surgery.

**References**


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