A 1-day-old boy presented with a protruding non-tender soft mass (8 × 7 cm) in the posterior sacral area at birth, with intact overlying skin. Sonography revealed a cystic dilation of the distal central canal surrounded by dilated subarachnoid space (figure 1). MRI confirmed the diagnosis of terminal myelocystocele by the typical finding of a

trumpetlike flaring syringocele and meningoceles (figure 2).

Terminal myelocystocele, a rare form of occult spinal dysraphisms, is characterized by hydromyelic dilation of the distal central canal (syringocele), bulging through a posterior spina bifida and surrounded by an expanded dural sheath (meningocele). Even though uncommon, it should be included in differential diagnosis of congenital sacral mass. Although ultrasound is an excellent initial screening modality, MRI is necessary to confirm the diagnosis and delineate the anatomy.

REFERENCES


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