Diastematomyelia is a rare form of spinal dysraphism characterized by a sagittal cleft of varying extent that splits the spinal cord, conus medullaris, or filum terminale with splaying of the posterior vertebral elements. This condition is caused by an osseous, cartilaginous, or fibrous septum, producing a complete or incomplete sagittal division of the spinal cord into two hemicords. It may be isolated or associated with other segmental anomalies of the vertebral bodies. Patients may have neurologic deficits in the lower limbs and perineum, causing gait disorders, sphincter disturbances, muscular atrophy, reflex changes, congenital scoliosis, and foot deformities. With modern imaging techniques, spinal dysraphism is being diagnosed in adults with increasing frequency, often as an incidental finding. The figure shows MRIs of a patient with clinically silent diastematomyelia.

Split spinal cord (diastematomyelia)
Hamid Sami, Elliott Ross, Max Walter, et al.
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