

Ependymoma

- 32 year old nurse referred to spinal clinic
- Currently off sick for the last month
- 1 year history of back and right leg pain
- 20 sessions with chiropractor- no help
- NSAIDs no help

- No CES symptoms
- No inflammatory pattern
- No PMH Ca
- Tip toe walker as child with corrective surgery aged 10
- LBP and right leg pain episodic since aged 16
- X-rays aged 16 NAD- no other investigations
- 2-5 episodes per year lasting 1-2 weeks

- Decreased lordosis
- End range pain all movements in Lx
- Catch of pain rising from flexion
- Poor muscle control around trunk
- Tender locally L5/S1 on palpation
- SIJ tests negative
- SLR 30 degrees bilaterally
- Positive slump

- Diagnosis of L5/S1 disc- R sided/central with NRI L5/S1
- Referred for Physio
- Review appt at 6/52
- No improvement at review
- Referred for MRI scan

3-10/100
population

Young persons
tumour- mostly
people in 20s and
30s

Male>female

Main spinal cord
tumour (60%). Glial
cell tumour

40% from the filum
terminale

Symptoms
longstanding before
diagnosis
(82%>12/12)

10-40% recurrence
rate



Slow growing tumour

Intradural mostly- occasional extradural spread

Symptoms dependent on site and size of tumour

May have back pain, leg pain, back and leg pain

May have neurological deficit of not

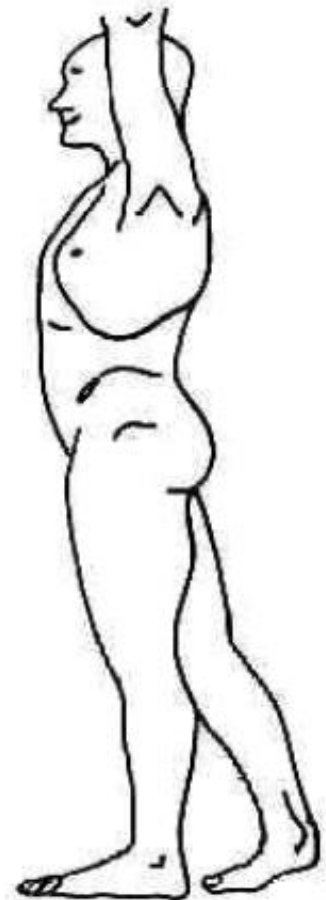
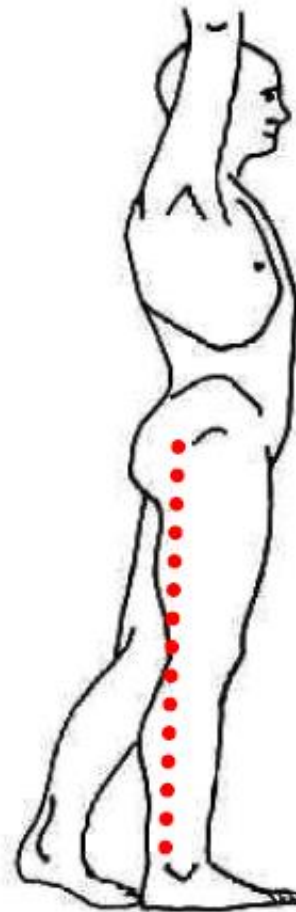
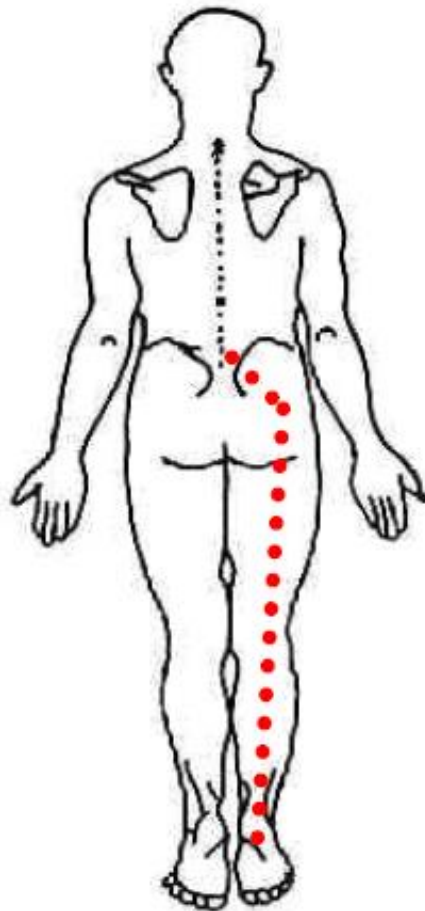
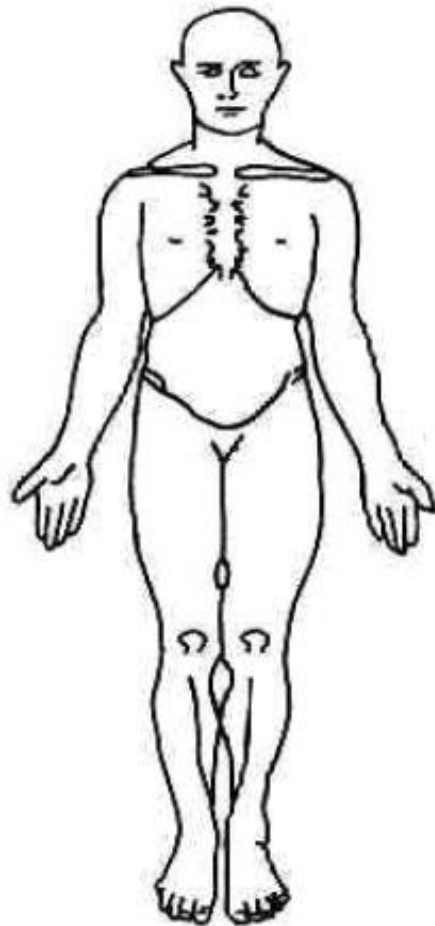
May have stenotic symptoms

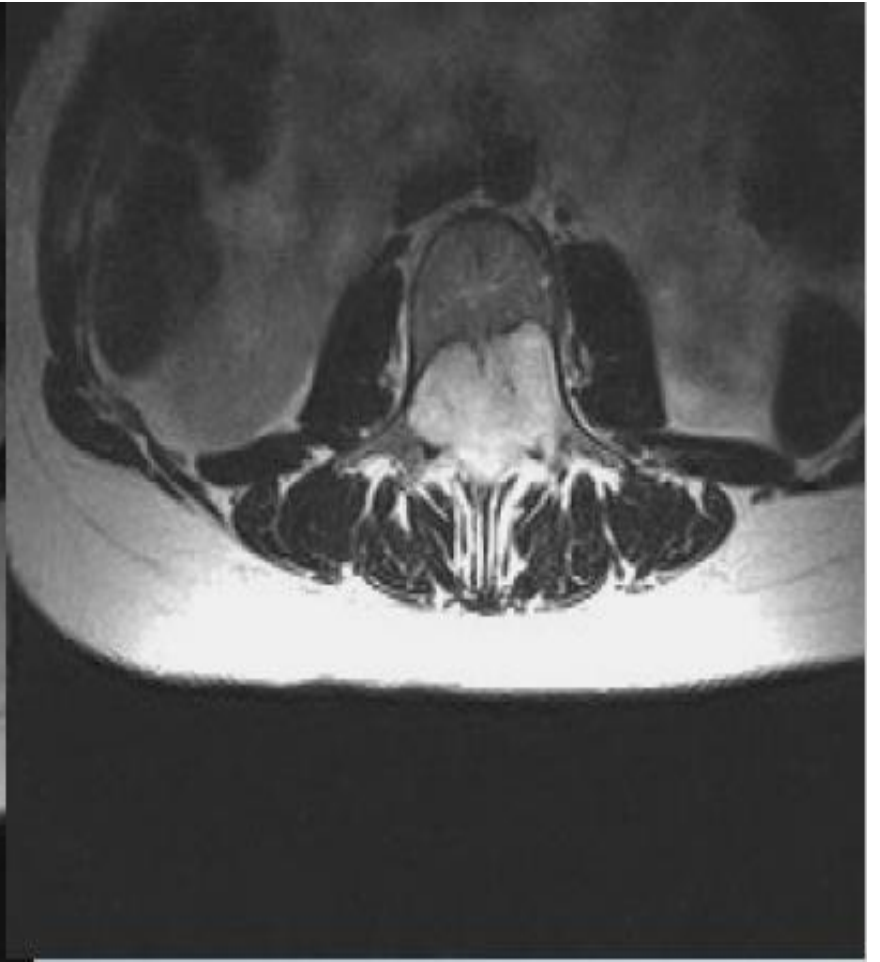
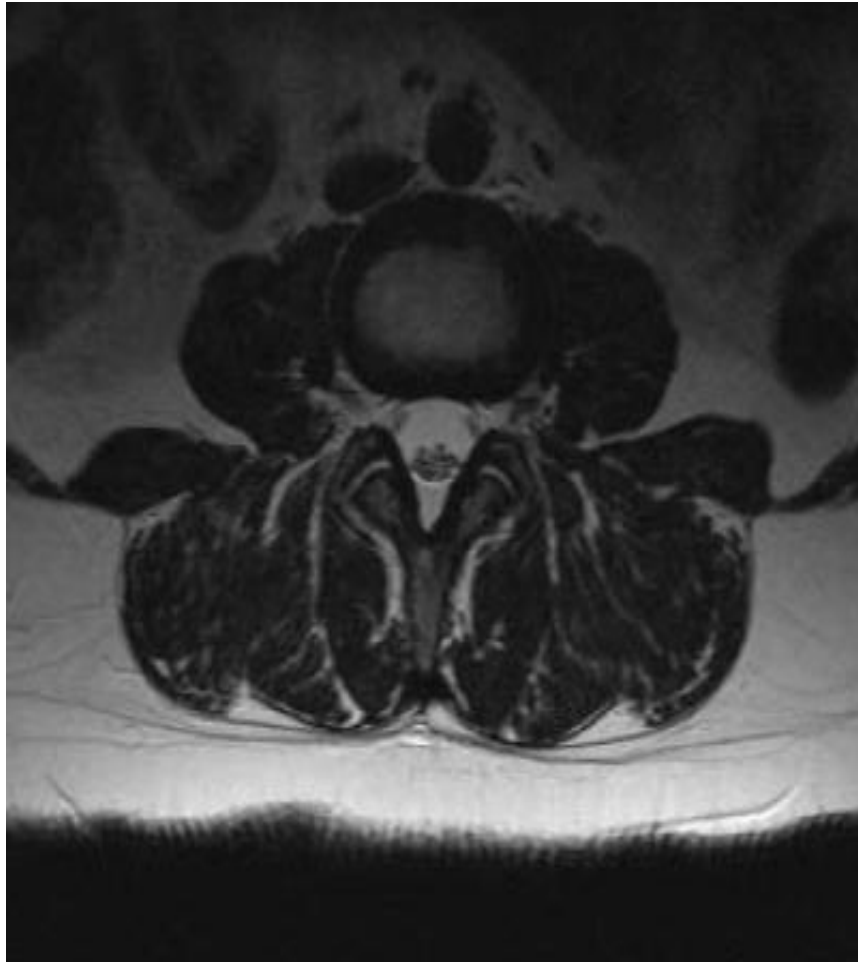
May have CES symptoms

May have none of these

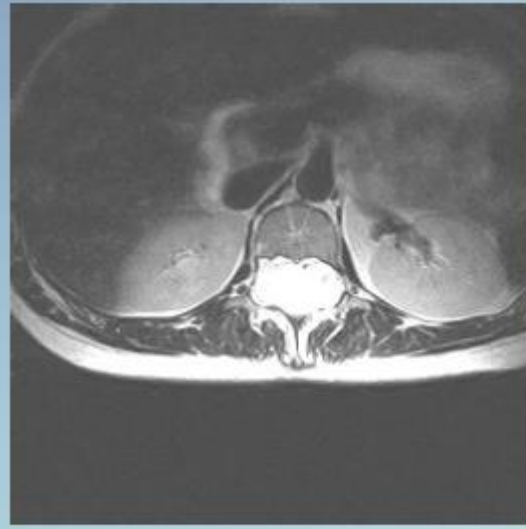
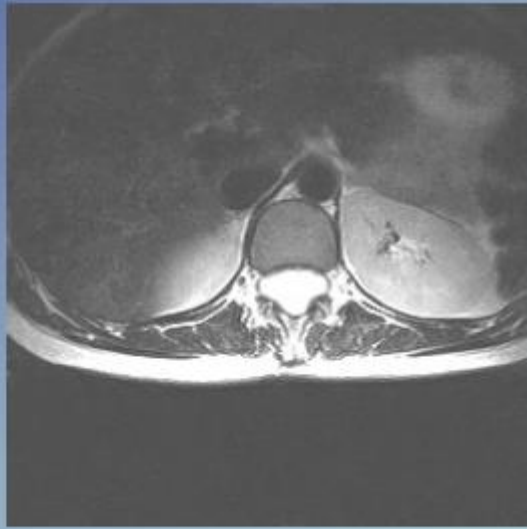
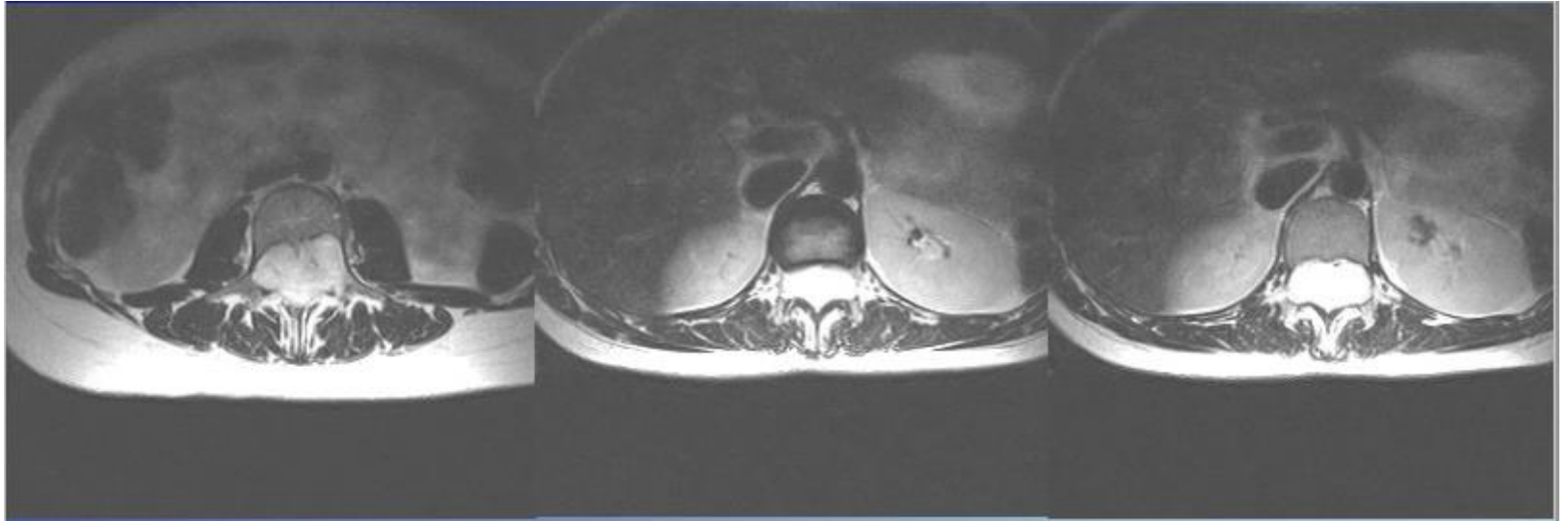
Treatment

- Surgery treatment of choice
- Surgery only possible in 43-59% of conus medullaris tumours
- Radiotherapy post partial resection
- Complete resection - 100% % year survival
- Partial resection - 50% survival















S



T1



T2



STIR



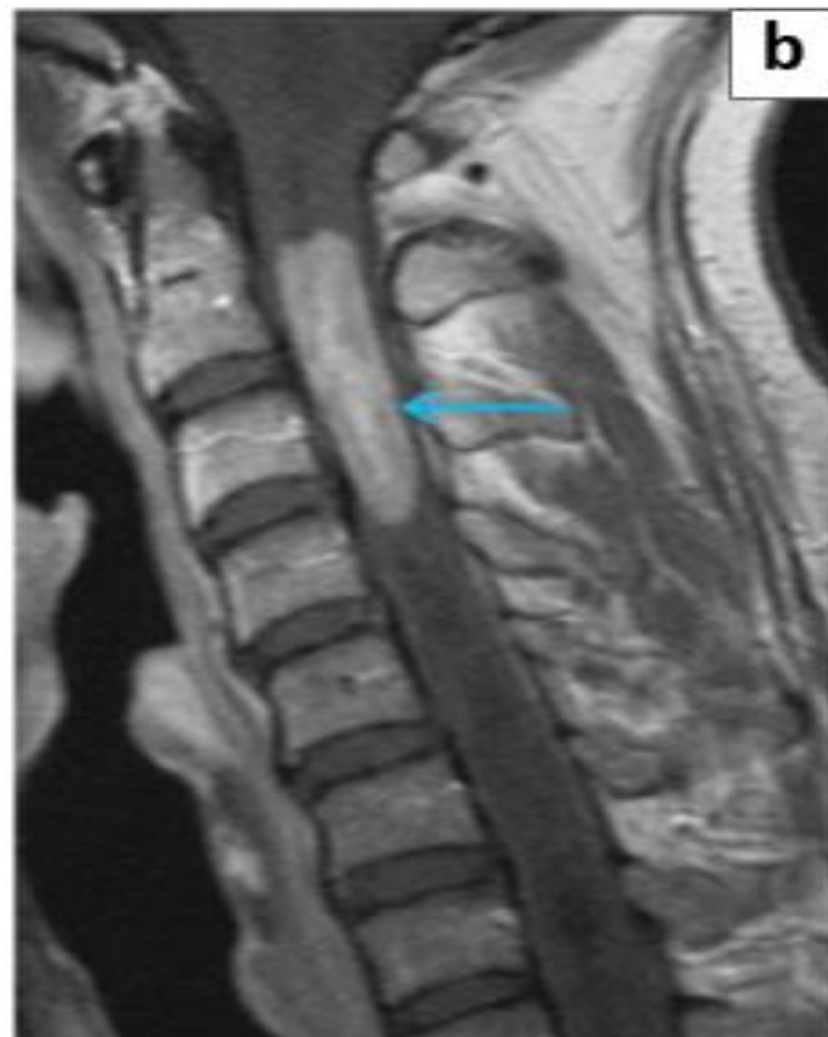


FIGURE 1: Intramedullary ependymoma in a 41-year-old man. (a) Sagittal T2-weighted image demonstrates a heterogeneous, well-circumscribed mass (white arrow) with extensive syringomyelia (yellow arrows); (b) Contrast-enhanced T1-weighted image demonstrates intense contrast enhancement of the tumour (blue arrow).