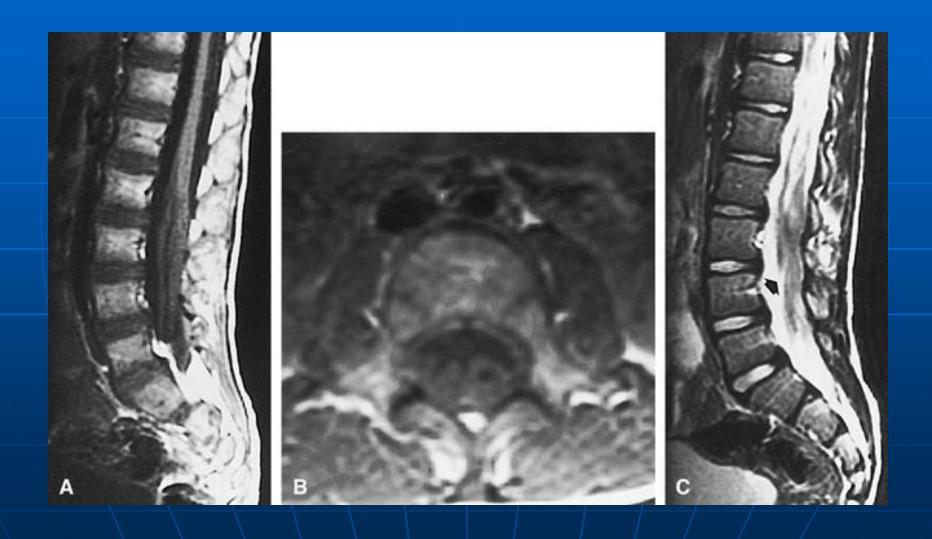
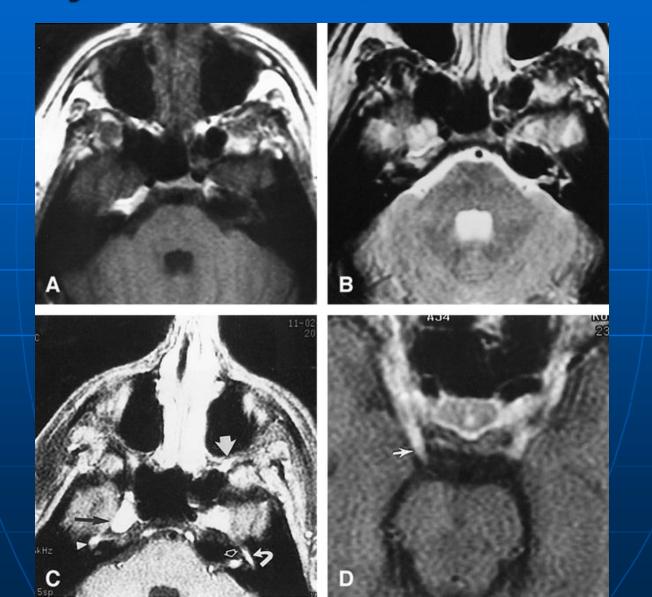
Dejerine-Sottas disease

- Rare hereditary motor and sensory neuropathy that presents with distal extremity motor and sensory symptoms as well as palpable peripheral nerves.
- Cranial nerve involvement is reportedly seen in approximately 15% of cases

Dejerine-Sottas disease



Dejerine-Sottas disease



Differential diagnosis of enlarged nerve roots

Nodular Thickening

- CSF Seeding (carcinomatous meningitis)
- Granulomatous disease
- Arachnoiditis
- Nerve sheath tumors (NF1 >> NF2)

Smooth Thickening

- Charcot-Marie-Tooth (inflammatory dymyelinating polyneuropathy)
- Dejerine-Sottas
- □ Chronic Inflammatory Demyelinating Polyradiculoneuropathy (CIDP)