

Multifocal nodular hypertrophy of trigeminal nerve in multifocal acquired demyelinating sensory and motor neuropathy

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A 60-year-old woman presented with asymmetric distal limb weakness and muscle atrophy. Electrophysiologic study revealed demyelinating polyneuropathy, and she was diagnosed with multifocal acquired demyelinating sensory and motor (MADSAM) neuropathy as a subtype of chronic inflammatory demyelinating polyneuropathy (CIDP) based on the established diagnostic criteria.¹ Anti-neurofascin155 antibody was negative. Cervical and lumbosacral nerve roots showed asymmetric multifocal hypertrophy (figure 1), which were distinct from those in typical CIDP.² Brain MRI revealed multifocal nodular hypertrophy of trigeminal nerves (figure 2 and video 1). These hypertrophic patterns in trigeminal nerves as well as nerve roots might be characteristics of MADSAM neuropathy.

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Author contributions

M. Kuwahara has contributed to the acquisition, analysis, and interpretation of data and drafted the article. I. Numoto has analyzed and interpreted data. S. Kusunoki has made substantial contributions to the conception and design of the study and also revised the paper critically for important intellectual content. S. Kusunoki provided the final approval to the paper.

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Disclosure

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Figure 1 Cervical and lumbosacral nerve roots hypertrophy

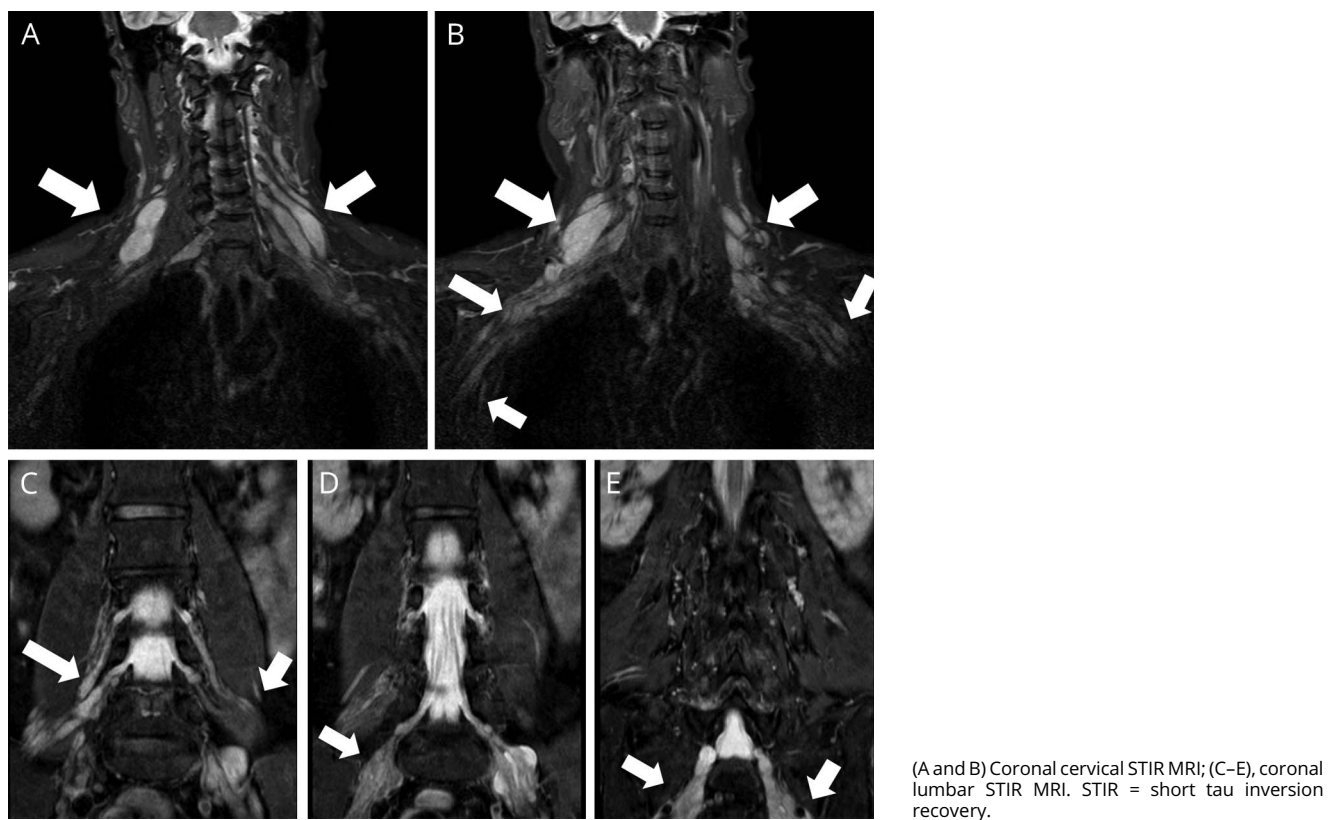


Figure 2 Trigeminal nerves hypertrophy

