A patient with unilateral upper medial medullary infarction presented with contralateral paralysis of the pharyngeal constrictor muscle in association with lemniscal sensory loss, pyramidal insufficiency, and central facial palsy on the same side. Individual differences in supranuclear control of the pharyngeal muscles may explain this rare occurrence. The combination of these signs is described as a syndrome of upper medial medullary lesion.

**CASE REPORT**

A 53 year old right handed man experienced heaviness of the right hand, then minimal speech and deglutition difficulties of acute onset. By the next day, the heaviness involved both the entire right upper and lower limbs, and clumsiness of the right hand and unstable gait followed. He had diabetes mellitus, which had been controlled medically for 4 years. On examination, he was conscious and fully oriented. His pupils were equal and reacted to light. Eye movement was normal, and there was no nystagmus. Trigeminal sensory and motor functions were normal. There was minimal weakness of the

**Abbreviations:** LMI, lateral medullary infarction; MMI, medial medullary infarction; MRI, magnetic resonance imaging; TMS, transcranial magnetic stimulation; MEP, motor evoked potential
lower half of the face on the right, as evidenced by impaired elevation of the upper lip when he opened his mouth. His head inclined slightly posterolaterally towards the left, but vestibular function otherwise was normal. While attempting to phonate, the posterior pharyngeal wall moved to the left, indicative of paralysis of the right side of the upper pharyngeal constrictor muscle (signe de rideau of Vernet\textsuperscript{10}) (fig 1A). The soft palate was normal in position and motion, but there was no soft palate reflex on either side. There was no tongue paralysis, and laryngeal examination showed normal vocal cord motion. In the limbs, muscle strength was normal and symmetrical. On the right side, muscle stretch reflexes were exaggerated, and there was extensor plantar response. Thermal and pinprick sensations in the limbs were unimpaired. Perception of joint movement and vibration was impaired in the right fingers and toes. Limb ataxia, present on the right, worsened when visual guidance was deprived. Magnetic resonance imaging of the brain showed a left side medial medullary infarction just below the pontomedullary junction (fig 1B). The rest of the brain stem and cerebellum, the hemispheres, and the cranial base otherwise were normal. Magnetic resonance angiography of the brain showed significant stenosis of the left vertebral artery.

Routine laboratory tests and echocardiography were normal except for elevated blood sugar and haemoglobin A1C levels. After a course of medical treatment, stable gait was restored. Minimal dysarthria and dysphagia lasted only a few days, whereas the right sided, central type facial and pharyngeal weakness still persisted at the follow up examination 2 months after stroke onset.

**DISCUSSION**

This patient had persistent paralysis of the right side of the upper posterior pharyngeal constrictor muscle due to a well demarcated, left sided medial medullary infarct. Unilateral motor weakness of the pharyngeal muscle, which is innervated by the nucleus ambiguus, usually indicates a nuclear or infranuclear lesion, but unilateral paralysis of the palatal, pharyngeal, and laryngeal muscles caused by a supranuclear lesion occasionally has been reported. Unilateral faciolingual-massereter-pharyngeal-laryngeal weakness from a contralateral oculopercular infarction was reported in the 1960s,\textsuperscript{11} but no additional cases have been reported since brain imaging techniques have come into use. In 1990, Bogousslavsky and Regli\textsuperscript{12} reported six patients with facial and lingual hemiparesis due to a contralateral lesion limited to the genu of the internal capsule, originally described by D\text{\textregistered}jeiner and D\text{\textregistered}jeiner-Klumpke.\textsuperscript{13} On the same side as their faciolingual hemiparesis, three patients showed unilateral weakness of the masseter muscle associated with palatal weakness (two), pharyngeal hypotonia and vocal cord paralysis (one), and decreased pharyngeal reflex (one). A review of the literature turned up two previous reports of unilateral paralysis of muscle(s) innervated by the nucleus ambiguus contralateral to a lesion in the corona radiata,\textsuperscript{14} and in the genu of the internal capsule.\textsuperscript{15} In some individuals therefore, the nucleus ambiguus seems to be governed by crossed corticobulbar innervation. A transcranial magnetic stimulation study of humans by Ertekin \textit{et al} suggested that in the nucleus ambiguous there is a crossed corticofugal connection.\textsuperscript{16} Motor evoked potentials were elicited in the cricothyroid muscle after contralateral motor cortex stimulation.

Our patient had unilateral, probably supranuclear, paralysis of the upper pharyngeal constrictor muscle, which has not been reported in medial medullary syndrome. The case of our patient further suggests that in humans supranuclear control of the nucleus ambiguous may have contralateral predominance.

**REFERENCES**