Case Report

Clinical history and examination

A 19 year old man was referred to the sleep service with a two year history of loud snoring and daytime hypersomnolence. He had lost approximately 20kg in weight over the previous eighteen months with no appreciable symptomatic change. He described short sleep latency, unrefreshing sleep however he did have good sleep hygiene, Epworth score was 14/24. He also described a two year history of hoarseness which initially developed following a three week episode of persistent hiccups. He had sung on a regular basis prior to this episode but was unable to subsequently. There was no past medical history of note and he was not taking any regular medication.

On examination he was of muscular build, weight 84.4Kg, BMI 29.6 KgM⁻². Cardio-respiratory examination was normal as was examination of the oropharynx. He was also noted to have vertical upbeat jerk nystagmus in the primary position persisting in all directions of gaze. Further neurological examination revealed bilateral weakness on finger and thumb abduction (grade 4/5) with brisk knee and ankle reflexes. Sensation mapping demonstrated a deficit in pain and temperature around the hairline of the face, back of neck, both shoulders and along the entire right arm. On subsequent otorhinolaryngology examination there were findings of a right recurrent laryngeal nerve palsy with no identifiable cause.

Chest x-ray demonstrated a scoliosis but was otherwise normal (Figure S1). CT scan of chest and neck showed subtle deviation of the left vocal cord medially, suggestive of a left vocal cord palsy (Figure S2). There were no masses along the expected tract of the left recurrent laryngeal nerve. No significant cervical lymphadenopathy and no abnormal masses were visualized within the neck. MRI of the spine demonstrated a Chiari I malformation (Figure 1). Although it did not appear to cause compression of the medulla or upper cord there was an associated thin but extensive syrinx throughout the cervical and upper thoracic regions (Figure 2). Multichannel polysomnography demonstrated frequent obstructive apnoeas (Figure S3), but a significant number of central or “mixed” apnoeas were also seen (Figure S4). The total arousal Index was 15.2/hr, associated with respiratory events. The Apnoea

Figure S1:
and Hypopnea Index was 28/hr. Flow volume loop (Figure S5) demonstrated flattening of the inspiratory loop in keeping with the vocal cord palsy. The patient was assessed by the neurosurgical service and boarded for a craniocervical decompressive procedure.

**Discussion**

Syringomyelia is the development of a fluid-filled cavity or syrinx within the spinal cord. The most common type of syringomyelia is due to blockage of cerebrospinal fluid (CSF) circulation. Obstruction of CSF circulation from the basal posterior fossa to the caudal space may cause syringomyelia of this type, the most common example being the Arnold–Chiari malformation. Although many mechanisms for syrinx formation have been postulated, the exact pathogenesis is still unknown. Frequently cited theories in cases with Arnold–Chiari malformation are those of William, and Oldfield [1].

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Signs and symptoms of neurological dysfunction that appear with distension of the syrinx are due to compression of long tracts, neurons, and microcirculation. Symptoms referable to raised intramedullary pressure are potentially reversible by syrinx decompression. Syringomyelia usually involves the cervical area, symptomatic presentation depends primarily on the location of the lesion within the neuraxis.

**Examination of the clinical manifestations and relation to their pathogenesis in this case follows;**

Syrinx interrupts the decussating spinothalamic fibers that mediate pain and temperature sensibility, resulting in loss of these sensations (dissociated sensory loss). Pain and temperature sensation may be impaired in either or both arms, or in a shawl-like distribution across the shoulders and upper torso anteriorly and posteriorly.

Syrinx extension into the anterior horns of the spinal cord damages motor neurons (lower motor neuron) and causes diffuse muscle atrophy that begins in the hands and progresses proximally to include the forearms and shoulder girdles. Lower limb spasticity, which may be asymmetrical, appears with other long-tract signs such as paraparesis, hyperreflexia, and extensor plantar responses.

Syringomyelia is a recognized etiological factor in the development of Charcot's joints. Neuropathic joint disease or Charcot joints are a chronic form of a degenerative arthropathy that is associated with decreased sensory innervation of the involved joints [2]. Neurogenic arthropathies may affect the shoulder, elbow, or wrist, scoliosis is seen sometimes in association with this process as in this case [3].

Patients with craniocervical disorders show a wide variety of symptoms and signs suggesting cerebellar and/or high cervical lesion. The anatomic localization of respiratory centers and their possible injury may explain the presence of respiratory disturbances in these diseases [4]. Botelho et al. described the polysomnographic findings in a group of patients with craniocervical disorders. Ninety percent of these patients complained of sleep problems (snoring, choking, and witnessed apneas) and 72% presented hypersomnolence (ESS >9) [4].

Observations of obstructive sleep apnoea in patients with ischaemic pontomedullary lesions, syringobulbia, syringomyelia, olivo-ponto-cerebellar atrophy or disseminated encephalopathy have indicated that it is due to impairment of inspiration–synchronous activation [5].

Brainstem signs are common in syringomyelia associated with Chiari malformations. In addition to the well-recognized sign of downbeat nystagmus, classically associated with foramen magnum abnormalities, a number of other ophthalmic features can be identified [6]. Symptoms include diplopia, oscillopsia, tunnel vision and difficulty in lateral gaze. Signs include nystagmus (downbeat, horizontal, rotatory, and combinations), strabismus, disc pallor, anisocoria, ptosis and field defect. In most cases nystagmus will improve or resolve following treatment [7].

Syringomyelia and Chiari type 1 malformation have previously been described in association with bilateral vocal cord paralysis and obstructive sleep apnea syndrome [8]. A further case report describes bilateral abductor vocal cord paralysis and sleep apnoea developing precipitously following general anaesthesia. Subsequent investigation demonstrated a type-I Chiari malformation with syringomyelia. The authors conclude that brainstem abnormalities such as Chiari malformation with secondary tenth cranial nerve deficits should be considered in previously healthy patients with signs and symptoms of upper airway obstruction and apnoea [9]. Flattening of the inspiratory loop on the flow-volume curve was found on pulmonary function testing, suggesting a variable extra-thoracic obstruction due to a laryngeal lesion [8].

Approximately 20 to 50% of patients with syringomyelia associated with Chiari malformations exhibit cranial nerve or cerebellar symptoms. However, hiccups represent a rare clinical manifestation of this disorder. In this case it is postulated that the initial presentation of intractable hiccups for 3 weeks heralded the early development of syringomyelia secondary to an Arnold–Chiari malformation.

**Conclusion**

Clinicians should consider Chiari malformation as a cause of sleep disturbed breathing and posterior fossa decompression as a potential treatment.

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**References**

3. Spiegel DA; Flynn JM; Staskelis PJ; Dormans, JP; Drummond DS; et al. (2003) Scoliotic curve patterns in patients with Chiari I malformation and/or syringomyelia. Spine 28: 2139-2146. [Link](https://goo.gl/pTMt1T)
