Risk of sudden death during sleep in syringomyelia and syringobulbia

Martin A Nogués, Ricardo Gené, Horacio Encabo

Abstract
Clinical, respiratory, and polysomnographic findings in three patients with syringomyelia and syringobulbia who developed severe respiratory complications are described. Neurological examination showed evidence of IXth and Xth cranial nerve involvement with dysphagia and dysphonia, but there were no complaints of serious sleep difficulties. Two patients died during sleep and the other was resuscitated a day later. All patients showed moderate restrictive ventilatory defects with reduced maximal buccal pressures and one also showed a low ventilatory response to CO₂ rebreathing. Protracted central, obstructive, and mixed apnoeas and hypopnoeas were commonly observed during sleep. There were no changes in heart rate during these events. A combination of respiratory and cardiovascular mechanisms might have been responsible for the severe complications described.

Respiratory abnormalities have been described in patients with syringomyelia and syringobulbia. 6,7 Several nuclei and pathways known to be involved in the voluntary or automatic regulation of breathing in animals may eventually be damaged by a syrinx, such as the vagal afferent fibres from the lungs to the medulla; the nucleus tractus solitarius; 6 the inspiratory and expiratory neurons located at the motor dorsal vagus, ambiguous, and tractus solitarius nuclei; 6 the ascending afferents and descending efferents involved in breathing control at the medulla or cervical spinal cord; and the efferent motor neurons to the diaphragm and intercostal muscles. 6 Despite the recognition of respiratory hazards in syringomyelia-syringobulbia patients, breathing abnormalities have so far received scant attention, particularly regarding their potential implication in sudden death.

The integrity of pathways and centres involved in voluntary and automatic breathing may be assessed by functional volume and pressure studies, together with polygraphic recordings during sleep. We describe the clinical and laboratory findings in three patients with advanced syringomyelia and syringobulbia. Respiratory and polysomnographic studies disclosing abnormalities may explain the severe acute complications reported.

Patients and methods
Case 1
This 54 year old man first noticed dizziness, dysphagia, and dyspnœa at the age of 37. A diagnosis of syringobulbia and syringomyelia was made at that time. Though he was treated with radiotherapy, his condition worsened. He then remained stable for 15 years, after which he noticed progressive weakness of all four limbs. He had been born by an abnormal delivery with forceps, suffering postnatal asphyxia.

On examination in March 1988 his overall appearance was normal, but he had noticeable wasting and weakness of the shoulder girdle. He had an ataxic gait, a left Horner's syndrome, horizontal and vertical nystagmus, definite wasting and weakness of proximal muscles in upper limbs, and areflexia. He had a spastic paraparesis, with brisk jerks and bilateral extensor plantar responses. Deep sensation was impaired in both hands, and there was an area of dissociated sensory loss from the face down to T5 bilaterally.

An MRI showed atrophy of the right cerebellar hemisphere and a cervical syrinx extending upwards to the lower part of the medulla (fig 1). He denied any sleep difficulties.

He had posterior fossa decompression with laminectomy at C1 and C2 in June 1988, and an extensive cervical arachnoiditis was found. The floor of the fourth ventricle was also explored, and a syringocysternal catheter was placed. After operation his swallowing difficulties worsened and the left upper limb became weaker. Nine days after surgery the patient had a sudden cardiorespiratory arrest while asleep in hospital and was successfully resuscitated. He received intermittent positive pressure ven-
Case 2
This 68 year old man developed wasting and weakness of both hands at the age of 30. This was followed by progressive loss of pain and temperature sensation over the right upper limb, and later the left, in a cape distribution down to the waist. Over the past 10 years he developed progressive weakness of the lower limbs, severe oscillopsia, dysphagia, and occasional dyspnoea. The diagnosis of syringomyelia was confirmed by myelography. Born by normal delivery, he had suffered meningitis at the age of 11.

On examination in August 1986 the patient had a short neck and low hair implantation. There was a left Horner’s syndrome, marked rotatory nystagmus to the left, wasting of the left sternocleidomastoid muscle, and wasting and fasciculations of the left side of the tongue. There was noticeable bilateral wasting and weakness of proximal and distal upper limb muscles, and reflexes were absent. He had spastic paraparesis, and bilateral extensor plantar responses were elicited. Dissociated sensory loss down to T4 was found. Vibration sense was impaired in all four limbs, but position sense was normal throughout. Although he never complained of sleep-wake difficulties, over the past 10 years relatives had noticed irregular breathing during sleep, such as respiratory pauses followed by strenuous snoring. In December 1988, the patient’s wife reported he had stopped breathing altogether in the course of an afternoon nap, and became cyanotic. She aroused him violently and started cardiopulmonary resuscitation. After several minutes he began to breathe normally, showing a decerebrate posture for several more minutes. He was admitted to hospital and recovered fully. The patient refused to be investigated by MRI.

Case 3
This 38 year old man first noticed progressive wasting and weakness of both arms at the age of 25. A year later his legs were also weak and walking had become difficult. His medical history was unremarkable apart from his birth after a prolonged and traumatic labour.

In 1975 a myelogram showed widening of the cervical and thoracic spinal cord. A CT scan showed moderate hydrocephalus, considerable dilatation of the fourth ventricle; a ventriculoperitoneal shunt was placed in December 1979, resulting in a slight improvement. In 1986, after a fall, his neurological condition deteriorated progressively. An MRI showed a normal brainstem and cerebellum with no evidence of cerebellar tonsillar herniation. The fourth ventricle was of normal size and location. The cervical spinal cord was noticeably atrophic from the level of the foramen magnum downwards, and a collapsed syrinx extended from the 4th ventricle down to T5. There was no radiological evidence of a lesion in the medulla. The patient had a T5 laminecotomy, syringotomy, and placement of a syringoperitoneal shunt. Postoperatively, he recovered gradually, with some improvement of the spastic tone.

On examination the patient had bilateral Horner’s syndrome and considerable nystagmus in all directions. There was weakness of the muscles innervated by the IXth, Xth, and X1th cranial nerves. Both upper limbs and shoulder girdles showed definite muscle wasting with reduced muscle tone and claw-like deformities of both hands. His upper limbs were areflexic. There was severe bilateral spasticity with hyperreflexia of the lower limbs, ankle clonus, and bilateral extensor plantar responses. Sensory testing disclosed an appreciable dissociated loss, mainly in a right hemi-cape distribution. Overall, he was severely disabled with hardly any useful function of his hands and he could walk only with assistance.

Over the past two years he had had occasional respiratory difficulties at night, waking up suddenly with breathlessness. Relatives were well aware of these episodes, which were worse after heavy meals or alcohol intake. They turned him over and woke him up. In November 1988 the patient went to sleep after a large meal, waking up with shortness of breath. He sat up in bed, recovered within a few minutes, fell asleep again, and was found dead one hour later.

Pulmonary function tests
The following variables were determined: flow-volume curves using a Fleisch pneumotach; forced vital capacity (FVC); forced expiratory volume at one second (FEV1); FEV1/FVC ratio; flow expiratory peak; and instantaneous flows at 25%, 50%, and 75% FVC. Expiratory muscle endurance was estimated from the values of maximal voluntary ventilation during 12 seconds. Respiratory muscle force was estimated from maximal inspiratory and expiratory pressures (PiMax measured at total lung capacity and maximal expiratory buccal pressure (PeMax) measured from residual volume. Values obtained were compared with normal control values from our own laboratory.

Arterial blood gases were determined from a sample obtained by brachial puncture while the patients were breathing room air. Respiratory centre output was then evaluated by means of the response to CO2 rebreathing, with a Collins spirometer containing a volume equal to the subject’s vital capacity plus one litre of the mixed gases containing 7% CO2 and 93% O2 (hyperoxygen rebreathing). The end tidal PCO2 (PECO2) was measured with an IL infrared CO2 analyser. A CO2 response curve was plotted over successive intervals of half a minute against PECO2 at their midpoints. Slopes of response curves for ventilation/PECO2 were calculated with a linear regression analysis. The mean value for normal subjects was 2-5 l/min Hg. Values below 1-0 l/min Hg were considered abnormal. During the rebreathing test in patient 1, the occlusion pressure at 0-1 seconds was evaluated by using a balloon inflated during expiration (without the subject’s knowledge) and measured 0-1
seconds after the start of the next inspiration. All data were recorded by a 2600 Gould polygraph at a paper speed of 50 mm/s. The day after the pulmonary function studies were performed the subjects were investigated with polysomnography.

**Polysonomography**

Patients were studied for one night in a quiet, darkened, sound attenuated room. Routine polysonomnographic techniques were used. Conventional leads were used for EEG, EOG, EMG (chin and tibialis anterior muscles), and EKG recordings.

Respiratory movements were transduced to pressure changes with rubber bellows placed at the maximal excursion abdominal site (Phipps and Bird pneumograph and Grass PTT5 transducer). EMG intercostal activity was also recorded with surface electrodes. Airflow was monitored by means of nasal and oral thermistors and snoring by a microphone.

The study was repeated in patients 2 and 3 two years later. On the second occasion, and in patient 1, oxygen saturation was monitored by means of an ear oximeter (Ohmeda Biox 3700), while a Model D G D polygraph was used throughout. Recordings were made at a chart speed of 15 mm/s, which allowed a one page scoring epoch of 20 seconds.

**Results**

**Pulmonary function studies**

As shown in table 1, patients 1 and 3 had a reduced respiratory rate, ranging from 7 to 9 breaths per minute. Patient 1 had a prolonged inspiratory time as well. Overall, patients showed moderate restrictive ventilatory incapacity with reduced maximal buccal pressures. Arterial blood gases at rest showed no evidence of CO₂ retention, and two of the three patients had mild hypoxaemia. Ventilatory response to CO₂ rebreathing was normal in patient 3, abnormally reduced in patient 2, and at the lower normal limit in patient 1 (fig 1).

Patient 1, who had an abnormally low respiratory rate at rest, showed only a slight rate increase during CO₂ rebreathing, ranging from eight to 12 respirations per minute. There were no important changes in heart rate during CO₂ rebreathing in any of the patients.

**Polysonomography**

All three patients showed low sleep efficiency (table 2). Sleep was frequently interrupted by

### Table 1 Results of pulmonary function tests

<table>
<thead>
<tr>
<th>Test or variable</th>
<th>Case 1</th>
<th>Case 2</th>
<th>Case 3</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>A</td>
<td>N</td>
<td>%</td>
</tr>
<tr>
<td>Respiratory rate (breaths/min)</td>
<td>7-0</td>
<td>18-0</td>
<td>9-0</td>
</tr>
<tr>
<td>FEV₁ (%)</td>
<td>2-0</td>
<td>62-0</td>
<td>4-0</td>
</tr>
<tr>
<td>FEV₁/FVC (%)</td>
<td>0-8</td>
<td>0-8</td>
<td>0-8</td>
</tr>
<tr>
<td>Pmax (%)</td>
<td>88-0</td>
<td>71-0</td>
<td>123-0</td>
</tr>
<tr>
<td>Pmax (%)</td>
<td>157-0</td>
<td>50-0</td>
<td>123-0</td>
</tr>
<tr>
<td>PaO₂ (mmHg)</td>
<td>77-7</td>
<td>68-0</td>
<td>77-7</td>
</tr>
<tr>
<td>PaCO₂ (mmHg)</td>
<td>43-1</td>
<td>36-7</td>
<td>39-5</td>
</tr>
<tr>
<td>CO₂ rebreathing:</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>VE/PECO₂</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PO₂/PECO₂</td>
<td>1-89</td>
<td>3-20</td>
<td>1-78</td>
</tr>
</tbody>
</table>

A = Absolute value  
N = Normal value  
% = Percentage of predicted value  
FEV₁ = Forced vital capacity; FVC = Forced expiratory volume at one second; Pmax = Maximal buccal inspiratory pressure as percentage of predicted value; PaO₂ = Maximal buccal inspiratory pressure as percentage of predicted value; VE/PECO₂ = Ventilatory response: end tidal CO₂ ratio (mmHg CO₂); PO₂/PECO₂ = Occlusion pressure: end tidal CO₂ ratio (cm H₂O/mmHg CO₂).

### Table 2 Polysomnographic findings

<table>
<thead>
<tr>
<th>Case 1</th>
<th>Case 2</th>
<th>Case 3</th>
</tr>
</thead>
<tbody>
<tr>
<td>1st study(7 Mar 88)</td>
<td>2nd study (18 May 88)</td>
<td>1st study (1 Aug 86)</td>
</tr>
<tr>
<td>Total recording time (min)</td>
<td>482</td>
<td>352</td>
</tr>
<tr>
<td>Total sleep time (min)</td>
<td>262</td>
<td>220</td>
</tr>
<tr>
<td>Sleep efficiency*</td>
<td>0-54</td>
<td>0-73</td>
</tr>
<tr>
<td>Sleep latency (min)</td>
<td>17</td>
<td>1</td>
</tr>
<tr>
<td>Waking after onset of sleep (min)</td>
<td>203</td>
<td>75</td>
</tr>
<tr>
<td>Stage 1 (%)</td>
<td>9-2</td>
<td>1-3</td>
</tr>
<tr>
<td>Stage 2 (%)</td>
<td>32-6</td>
<td>46-5</td>
</tr>
<tr>
<td>Stage 3+4 (%)</td>
<td>4-4</td>
<td>4-4</td>
</tr>
<tr>
<td>REM stage (%)</td>
<td>14-6</td>
<td>22-4</td>
</tr>
<tr>
<td>REM latency (min)</td>
<td>32-4</td>
<td>57</td>
</tr>
<tr>
<td>No of wakefulness</td>
<td>38</td>
<td>25</td>
</tr>
<tr>
<td>No of central apnoeas (maximal duration in secs)</td>
<td>34 (25)</td>
<td>18 (30)</td>
</tr>
<tr>
<td>No of obstructive and mixed apnoeas (maximal duration in secs)</td>
<td>36 (35)</td>
<td>46 (35)</td>
</tr>
<tr>
<td>No of hypopnoeas (maximal duration in secs)</td>
<td>13 (10)</td>
<td>4 (35)</td>
</tr>
<tr>
<td>Total no of apnoeas and hypopnoeas (mean duration in secs)</td>
<td>47 (13)</td>
<td>64 (19)</td>
</tr>
<tr>
<td>Apnoea + hypopnoea index*</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Minimum oxygen saturation (%)</td>
<td>83</td>
<td>77</td>
</tr>
<tr>
<td>Bradycardia during apnoeas</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Periodic leg movements</td>
<td>+</td>
<td>+</td>
</tr>
</tbody>
</table>

*Sleep efficiency: total sleep time / total recording time.  
†Apnoea + hypopnoea index = total number of apnoeas + hypopnoeas / total sleep time in minutes × 60.
respiratory events or periodic leg movements, or both. The number of awakenings (arousals lasting at least 10 seconds) and the time to waking after onset of sleep were increased in all studies. Sleep latency was prolonged only in patient 3. Delta sleep (stages 3 and 4) was reduced or absent in the three patients and the REM stage was of brief duration in patients 1 and 3.

Respiration during sleep was disturbed, as shown by the apnoea plus hypopnoea index, which proved higher in subjects 2 and 3. There were frequent central apnoeas, obstructive apnoeas, and mixed apnoeas, as well as hypopnoeas. On occasion, apnoeas and hypopnoeas were extremely long lasting and associated with low oxygen saturation values, but there was no evidence of concomitant bradycardia.

The second study in patients 2 and 3 showed an increase in the absolute number, mean duration, and maximal duration of apnoeas and hypopnoeas. The longest event recorded lasted 170 seconds (in the second study in patient 3). In the second study central and obstructive apnoeas decreased in patient 2, though the number of hypopnoeas increased considerably. Central apnoeas lasting 30–40 seconds were observed in both studies while patient 3 was awake. Patients 1 and 3 also showed a very low respiratory rate (7–9 breaths/minute), and patient 1 had a long inspiratory time (6–10 seconds).

Discussion
All three patients who had syringomyelia-syringobulbia and severe cardiorespiratory complications had clinical involvement of the cervical spinal cord and medulla, the critical structures for respiratory controls. Abnormalities of respiration in syringomyelia-syringobulbia patients have occasionally been described previously. Ventilatory centre insensitivity to CO₂ has been reported in patients with syringomyelia or Chiari type 1 malformation who developed respiratory symptoms. Insensitivity to hypoaxia with a relatively normal response to CO₂ was also observed in a patient with a Chiari malformation and syringomyelia. Bilateral weakness of the diaphragm and isolated central respiratory failure have been described in syringomyelia and Chiari type 1 malformation. Cases of obstructive sleep apnoea have also been reported. Ours is the first study in which respiratory abnormalities were documented by functional respiratory studies and polysomnography before the occurrence of respiratory arrest or sudden death. Furthermore, the recordings have detected abnormalities of heart rate during sleep and low respiratory rates and central apnoeas during wakefulness.

Respiratory functional studies showed that ventilatory mechanisms were affected to a variable degree in these patients, who had in common a pronounced wasting and weakness of intercostal and neck muscles. Nevertheless, there was no clinical evidence of diaphragm weakness. Dyspnoea in two patients was probably due to muscle weakness since buccal pressures were abnormal in all three patients. Maximal buccal pressures are useful in assessing the strength of respiratory muscles: they are easily measured and more sensitive than clinical examination in detecting muscle weakness. Patients 1 and 2 had mild restriction with abnormal responses to CO₂, whereas patient 3 had severe restriction and normal response to CO₂ rebreathing while awake. Such abnormalities in mechanics may worsen during sleep, leading to more severe ventilatory difficulties, as has already been shown in other conditions, including chronic obstructive pulmonary disease, interstitial lung disease, and scoliosis.

Obstructive, mixed, and central apnoeas were found in all three patients, and the apnoea plus hypopnoea index exceeded the usually accepted normal limit of five events per hour.

Several factors, such as weakness of pharyngeal and laryngeal dilator muscles supplied by pontomedullary cranial nerve nuclei predisposed these patients to obstructive sleep apnoea. Our patients had clinical involvement of the IXth and Xth cranial nerves, as in another case previously reported, in which obstructive sleep apnoea was attributed to damage to the ambiguous and hypoglossal nuclei by a syrinx found at necropsy.

The mechanism of central apnoeas in syringomyelia patients is probably related to compression of descending and ascending affreter pathways in the anterolateral region of the cervical spinal cord by the syrinx. A similar mechanism has been suggested to explain central sleep apnoea after percutaneous cordotomy. Damage to medullary structures may also contribute to this, and explain moreover the low respiratory rate found in patients 1 and 3, whether asleep or awake. In patient 2, basilar impression and a Chiari anomaly may also have had a role in the occurrence of these complications.

The increase in the total number, mean, and maximal duration of apnoeas and hypopnoeas in the second study in patients 2 and 3 suggests deterioration of sleep architecture, most likely due to progression of the disease.

Although patients made no complaints of serious sleep difficulties, relatives were well aware of nocturnal breathing irregularities. Reduced sleep efficiency and increased time to waking after onset of sleep were produced by frequent arousals or awakenings associated with respiratory events or periodic leg movements. The paucity of symptoms of sleep disturbance was in striking contrast to the abnormalities found at polysomnography. Patients denied experiencing excessive daytime sleepiness, repeated nocturnal awakenings, restless sleep, insomnia, or morning headache.

All three patients were at risk of sudden death during sleep. Patients 1 and 3 actually died and patient 2 was resuscitated. These syringobulbia-syringomyelia patients apparently suffered a respiratory arrest while asleep similar to those leading to death in cases of Shy-Drager syndrome, primary cerebral
involvement, or neurosurgical lesions. The concurrence of long lasting apnoeas and hypopnoeas, absence of associated bradycardia or tachycardia, extremely low respiratory rate, and prolonged inspiratory effort point to loss of automatic respiratory regulation during sleep.

In all likelihood, compromise of cardiovascular reflexes contributed to the risk of sudden death in these patients with syringomyelia or syringobulbia. Subclinical involvement of baroreceptor and vagal cardiovascular reflexes are common in syringobulbia patients, particularly in those with clinical evidence of IXth and Xth cranial nerve impairment. Even though autonomic tests were not undertaken, the constant absence of bradycardia during the apnoeas suggests vagal dysfunction.

In conclusion, our findings emphasise the importance of periodic functional respiratory and polysomnographic studies in patients with syringomyelia or syringobulbia. These studies may indicate the need for nocturnal assisted mechanical ventilation in these patients at a greater risk of sudden death.

17 List CF. Neurologic syndromes accompanying developmental anomalies of the occipital bone, atlas and axis. Arch Neurol Psychiatry 1941;45:577–616.
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