Management of respiratory failure with ventilation via intranasal stents in cystic fibrosis

F P Edenborough, M Wildman, D W Morgan

Abstract
The case history is presented of a patient with acute respiratory failure complicated by nasal obstruction resulting in intolerance of nasal ventilation. Urgent insertion of nasal stents permitted restoration of ventilation with resolution of breathlessness and stabilisation of arterial blood gases.

Keywords: cystic fibrosis; respiratory failure; nasal stents; nasal intermittent positive pressure ventilation

Type II respiratory failure is common in patients with cystic fibrosis with advanced disease and may respond to long term nasal ventilation. We report a case of acute respiratory failure complicated by nasal obstruction resulting in intolerance of nasal ventilation which required urgent insertion of nasal stents.

Case history
The 28 year old man with cystic fibrosis had diabetes mellitus, mild liver disease, aminoglycoside induced renal tubulopathy, sputum colonised with Pseudomonas aeruginosa, and type I (hypoxic) respiratory failure requiring long term oxygen therapy. His spirometric test results when well gave forced expiratory volume in one second (FEV1)/forced vital capacity (FVC) of 0.95/2.67 l with stable capillary blood gases as in table 1 (column A).

He was admitted with an FEV1/FVC of 0.55/1.80 l and an oxygen saturation of 84% breathing room air. His nebulised medication included DNase (Pulmozyme®), β2 agonists, and colistin sulphate with inhaled fluticasone. He received supplemental nasogastric feeding and required Creon 8000, cisapride, ranitidine, and metoclopramide, and Humulin insulin.

Investigations revealed hypokalaemia (K+ = 2.7 (normal range 3.5–5.3) mmol/l), hypomagnesaemia (Mg2+ = 0.68 (normal range 0.7–1.0) mmol/l), and neutrophilia of 11 700/ml. A full blood count and renal and liver function results were otherwise normal. Treatment with intravenous aztreonam and tobramycin and K+ and Mg2+ supplementation was commenced and he initially improved, but after 14 days his antibiotics were changed to Tazocin, colistin sulphate, and ciprofloxacin for extended cover of a resistant Pseudomonas and metronidazole for possible anaerobic infection.

He continued to improve but on the 40th day of admission he again deteriorated with a fever of 39.6°C with arterial blood gas tensions as shown in table 1 (column B). Nasal intermittent positive pressure ventilation (NIPPV) was commenced at settings of 3/12/12 on 4 l oxygen (expiratory positive airway pressure (EPAP) 3 cm H2O/inspiratory positive airway pressure (IPAP) 12 cm H2O/ default rate = 12 breaths/min). He deteriorated overnight (table 1 (column C)) and was treated with intravenous hydrocortisone, increased ventilation (3/14/12 on 6 l oxygen), and physiotherapy for presumed sputum retention. The antibiotics were changed to ceftazidime, tobramycin, and Tazocin and by the following morning he had improved (table 1 (column D)). The following night his nose blocked and, despite fluticasone 50 µg nasal spray, ephedrine 0.5% drops, and saline washes, he was unable to tolerate NIPPV either nasally or via a full face mask. Despite maximal medical treatment including intensive physiotherapy and maximal oxygen by mask his gases deteriorated (table 1 (column E)) and he began to tire. The changes in blood gas tensions are shown in fig 1.

Nasendoscopy performed to determine the cause of the nasal blockage revealed minimal crusting and debris but markedly oedematous nasal mucosa unresponsive to 1:80 000 adrenaline and 5% cocaine spray. An attempt to insert size 6 nasopharyngeal tubes failed due

Table 1 Blood gas tensions during nasal ventilation in relation to placement of stents

<table>
<thead>
<tr>
<th>A</th>
<th>B(–3)</th>
<th>C(–2)</th>
<th>D(–1)</th>
<th>E(0)</th>
<th>F(+1)</th>
<th>G(+2)</th>
<th>H(+4)</th>
<th>I(+6)</th>
<th>J(+7)</th>
<th>K(+17)</th>
<th>L(+20)</th>
</tr>
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<tbody>
<tr>
<td>FIO2 (l)</td>
<td>Air</td>
<td>4</td>
<td>4</td>
<td>4</td>
<td>6</td>
<td>8</td>
<td>12</td>
<td>10</td>
<td>8</td>
<td>8</td>
<td>4</td>
</tr>
<tr>
<td>HCO3</td>
<td>33.5</td>
<td>40.8</td>
<td>55.1</td>
<td>37.7</td>
<td>40.5</td>
<td>34.8</td>
<td>42.0</td>
<td>37.7</td>
<td>36.4</td>
<td>39.8</td>
<td>34.7</td>
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<tr>
<td>FPAO2 (%)</td>
<td>92</td>
<td>84</td>
<td>91</td>
<td>94</td>
<td>75</td>
<td>82</td>
<td>95</td>
<td>93</td>
<td>95</td>
<td>99</td>
<td>94</td>
</tr>
</tbody>
</table>
to the extent of the obstruction airway. Portex Blue Line size 4.5 siliconised endotracheal tubes (internal diameter 4.5 mm, external diameter 6.5 mm, Portex Ltd, Hythe, Kent, UK) were therefore placed under direct vision using a nasendoscope to lie in the postnasal space on each side, trimmed at approximately 14 cm with 2 cm projecting externally allowing the tubes to be sutured together anterior to the columella (fig 2A and B). Nasal toilet was then performed with saline douches. Advancing the tubes into the nasopharynx (fig 2C and D) by extending the patient’s neck enabled pulmo-

tory toilet via a size 10 suction catheter after which the tubes were withdrawn back into the postnasal space to avoid excessive gagging. After two hours on NIPPV set at 3/11/12 with 8 l oxygen the gases were as shown in table 1 (column F) and fig 1, improving overnight (table 1 (column G)) on increased ventilation (3/14/12 and 12 litres oxygen).

Continued maximal medical therapy including physiotherapy five times daily resulted in improved gas tensions despite reducing inspired oxygen. Nasogastric feeding was recom-

menced via a size 8 Portex infant feeding tube passed through the right sided stent. The left tube was removed on day 18 and by day 20, when no longer reliant on overnight ventilation (table 1 (column L)) the second stent was removed. Fluticasone 50 µg and xylometazolone 0.1% nasal sprays were commenced but the patient reported no nasal stuffiness or blockage and was later able to recommence NIPPV.

No organism was found to account for his pyrexial illness and deterioration. Multiple central and peripheral blood cultures, urine and stool cultures were negative. Tests for Legionella and respiratory viruses including CMV and EBV were also negative. His Pseudomonas remained sensitive to tobramycin and colistin sulphate throughout with intermit-
tent sensitivity to Tobramycin. At the time of writing he remains in hospital on intravenous anti-
biotics and on the active transplant list.

**Discussion**

Nasal polyposis and chronic rhinosinusitis is not uncommon in cystic fibrosis. Nasal ventil-

ation has previously been described in cystic fibrosis.

![Blood gas tensions during nasal ventilation in relation to placement of stents.](http://thorax.bmj.com/)
fibrosis and, in this centre, is used to treat type II respiratory failure in patients awaiting transplantation. Despite routinely prescribed nasal steroids, dryness and crusting is common though recently introduced custom designed warm air humidifiers such as HumidAire (ResMed (UK) Ltd, Abingdon, Oxon, UK) appear to lessen the incidence of nasal problems.

Alternative patient interfaces with NIPPV are available including full face masks, unsuccessful in this case, and mouth only or nose and mouthpieces which have been used extensively elsewhere but which were not available in this case. Standard nasopharyngeal airways were found to be too soft to bypass the obstruction and hence an alternative was sought. Silicone tubes have been used as stents following surgery for acquired choanal atresia in infants and rarely in adults. Endotracheal tubes have similarly been used in infants but we are unaware of the use of this technique acutely in adults.

In this case, having been unable to oxygenate the patient, urgent insertion of nasal stents permitted restoration of NIPPV which successfully treated the acute hypoxia and improved hypercapnia over time. The procedure was well tolerated; nasal complications such as columellar or septal ulceration, nasal cavity or sinus infection were not seen and the nares remained patent after removal of the tubes. We believe the use of urgently inserted nasal stents was life saving in this case.

We thank Dr D E Stableforth, consultant physician, for permission to publish on this patient and Dave Thomas, senior ODA, ENT theatres for modelling the nasal stents. Dr Wildman is sponsored by the UK CF Trust.

Funding: none.

Conflicts of interest: none.


Recurrent laryngeal nerve palsy associated with mediastinal amyloidosis

Philip Conaghan, Darryl Chung, Roger Vaughan

Abstract

Amyloidosis affecting peripheral nerves causing isolated nerve palsies is uncommon. Localised amyloidosis occurs less frequently than the reactive or immune related systemic forms, and mediastinal localisation is virtually unknown. We present a case of recurrent laryngeal nerve palsy associated with mediastinal AL amyloidosis in a middle aged man.

(Torax 2000;55:436–437)

Keywords: amyloidosis; neuropathy; mediastinum

Case report

A 64 year old white retired bricklayer presented with a three week history of hoarseness and a five week history of dysphagia for solids. Apart from his hoarseness, physical examination was non-contributory. Chest radiography showed a grossly widened mediastinum. Despite a normal Kveim test and bronchial and transbronchial biopsy specimens, he had an unproven diagnosis of sarcoidosis based on a chest radiograph performed 17 years previously showing bilateral hilar lymphadenopathy and bilateral upper lobar nodular shadowing. Computed tomographic scanning showed a large calcified mass in his superior mediastinum surrounding the trachea, the aortic arch, compressing the left brachiocephalic vein, and associated with bilateral calcified upper lobe masses (fig 1). A barium swallow showed a small upper oesophageal posterior diverticulum but no...
Recurrent laryngeal nerve palsy associated with mediastinal amyloidosis

437

rare, with only two cases reported as the AL airways and gastrointestinal tract. However, extramedullary plasmacytomas of the upper tissues. However, proteinaceous, eosinophilic fibrillar deposit in Amyloid is described as an homogeneous, protein fibrillar deposit in various tissues. However, x-ray diffraction accurately identifies the amyloid fibril by its unique β-pleated sheet conformation. Diagnostic characteristics are: (1) apple-green birefringence of Congo Red stained tissue under polarised light, (2) bright yellow-green fluorescence in thioflavine stained preparations, and (3) ultrastructural demonstration of 7–10 nm non-branching fibrils of indeterminate length. Primary immunoglobulin light chain (AL) amyloid occurs in its localised form in extramedullary plasmacytomas of the upper airways and gastrointestinal tract. However, localisation of amyloid to the mediastinum is rare, with only two cases reported as the AL type and one as the reactive (AA) form. Although calcific changes were evident on the chest radiograph, both bronchial and transbronchial biopsy specimens were normal. No histological diagnosis has been obtained of the upper lobe lesions but they are also considered to be amyloid deposits.

It is possible that the earlier chest radiographic findings diagnosed as sarcoidosis could represent early amyloid deposits, although the radiographic changes are typical for sarcoidosis.

Apart from carpal tunnel syndrome associated with β₂-microglobulin deposition in haemodialysis patients, amyloid rarely affects the peripheral nervous system and is usually of the λ type, being of monoclonal origin in 85% of cases. Familial patterns of amyloid mixed peripheral polyneuropathy have been described, commonly associated with variant plasma transthyretin. This case of recurrent laryngeal nerve palsy associated with amyloidosis may be due to perineural infiltration or compression. Infiltration of the vagus nerve itself with amyloid may account for both the vocal cord paralysis and oesophageal dysmotility.

The mediastinal “amyloidoma” may represent an underlying lymphoplasmacytoma or a subtle plasma cell dyscrasia such as a Castleman’s disease tumour of the solitary plasma cell variety, although this patient had no constitutional symptoms or signs of a POEMS syndrome. No steroids or cytotoxic treatment have been considered. Technically challenging surgical resection of this amyloid mass is best reserved for severe compressive and obstructive symptoms of the vascular, upper airway, or oesophageal lumen. Alternatively, intraluminal stenting may be considered.

Discussion

Amyloid is described as an homogeneous, proteinaceous, cosinophilic fibrillar deposit in various tissues. However, x-ray diffraction accurately identifies the amyloid fibril by its unique β-pleated sheet conformation. Diagnostic characteristics are: (1) apple-green birefringence of Congo Red stained tissue under polarised light, (2) bright yellow-green fluorescence in thioflavine stained preparations, and (3) ultrastructural demonstration of 7–10 nm non-branching fibrils of indeterminate length. Primary immunoglobulin light chain (AL) amyloid occurs in its localised form in extramedullary plasmacytomas of the upper airways and gastrointestinal tract. However, localisation of amyloid to the mediastinum is rare, with only two cases reported as the AL type and one as the reactive (AA) form. Although calcific changes were evident on the chest radiograph, both bronchial and transbronchial biopsy specimens were normal. No histological diagnosis has been obtained of the upper lobe lesions but they are also considered to be amyloid deposits.

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Thorax 2000 55: 434-436
doi: 10.1136/thorax.55.5.434

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