CASE REPORT

Successful treatment of atelectasis with Dornase alpha in a patient with congenital muscular dystrophy

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KEYWORDS
Dornase alpha; Neuromuscular diseases; Atelectasis

Abstract A 28-year-old neuromuscular patient chronically treated with nocturnal noninvasive ventilation developed pulmonary lobar atelectasis and daytime hypoxemia. Twenty-four-hour 5L/min oxygen was begun, while mechanical cough assist aids were applied for seven days. In the following three days, treatment with nebulized Dornase alpha (rhDNase) b.i.d. was tested, without any significant improvement. On 11 and 13th days rhDNase was instilled by flexible bronchoscopy. A rapid resolution of the atelectasis was observed with relief of hypoxemia, without significant side effects. On day 16 the patient was discharged without oxygen requirements. In non-intubated neuromuscular patients with atelectasis who do not respond successfully to non-invasive treatments intrabronchial instillation of rhDNase may safely help to improve airway clearance.

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PALAVRAS-CHAVE
Dornase alfa; Doenças neuromusculares; Atelectasia

Tratamento bem sucedido de atelectasia com Dornase alfa num doente com distrofia muscular congênita

Resumo Um doente neuromuscular crónico de 28 anos de idade, tratado com ventilação noturna não invasiva, desenvolveu atelectasia lobar pulmonar e hipoxemia diurna. Foi iniciado suporte de oxigénio durante 24 horas, enquanto uma ajuda mecânica para a tosse foi aplicada por 7 dias. Nos 3 dias seguintes o tratamento com Dornase alfa nebulizado (rgDNase) b.i.d. foi testado, sem qualquer melhoria significativa. No 11.º e 13.º dias rhDNase foi introduzido por broncoscopia flexível. Um restabelecimento rápido da atelectasia foi observado com alívio da hipoxemia, sem efeitos secundários significativos. No 16.º dia o doente teve alta sem necessidade de oxigénio. Em doentes neuromusculares não intubados, com atelectasia,
Introduction

In neuromuscular disease, impairment of cough mechanisms due to expiratory muscle weakness can favor the development of atelectasis after pulmonary infections. This can lead to hypoxia with a life-threatening clinical situation. There is a lack of evidence-based studies on the management of infectious atelectasis.

Although chest physiotherapy, mechanical in-exsufflation (Mi-E) and high-frequency chest wall oscillation (HFCWO) improve airway clearance, they may not be sufficient, particularly when secretions become highly viscous due to accumulation of significant amounts of extracellular DNA.

Recombinant human Dornase alpha (rhDNase) may cleave and depolymerize extracellular DNA, and separate it from proteins: this allows endogenous proteolytic enzymes to break proteins and decrease viscoelasticity and surface tension of purulent sputum. In patients with cystic fibrosis rhDNase has proved extremely effective, with both aerosol administration and bronchoscopic instillation. Studies in newborns or in children have reported beneficial effects of rhDNase on atelectasis in non-cystic fibrosis patients, and anecdotal reports have suggested a beneficial effect on respiratory syncytial virus bronchiolitis and in atelectasis due to mucus plugs in newborns and children.

Most of these studies were performed by instilling the drug directly into the trachea in intubated patients. Although most patients showed an improvement within 24 h, in some of them direct instillation resulted in clinical deterioration, presumably due to a mucus mobilization that was too rapid. This effect may be dangerous in neuromuscular patients if means for removal of the mucus are not immediately available, or if the rate of mobilization exceeds capacity of elimination. However, no studies reporting effects of rhDNase in neuromuscular patients have been published. In this article we describe the favorable clinical course of a non-intubated neuromuscular patient with infectious atelectasis treated with bronchoscopic instillation of rhDNase.

Case report

A female patient with a congenital muscular dystrophy had been treated with nocturnal nasal noninvasive positive pressure ventilation (NIV) since she was 20. At the age of 26 she was hospitalised for a complete atelectasis of the right lower lobe, and had recovered after more than a month with the application of an intensive combined protocol of HFCWO (The Vest Airway Clearance System, Hill-Rom St. Paul, MN, USA) plus manual and mechanical assist cough (In-Exsufflator, Cough-Assist®, Philips Respironics, Murrysville, PA, USA). Her clinical condition remained good until the age of 28, when her lung function tests showed the following values: vital capacity 0.55 L (16% of the predicted value), maximal inspiratory pressure 10 cmH2O (11% of predicted), maximal expiratory pressure 14 cmH2O (13% of predicted), and peak cough expiratory flow 80 L/min. A few months later, due to the appearance of fever and copious mucus production, she was treated with manual and mechanical chest physiotherapy plus antibiotics (ceftriaxone) initially. However, she still complained of dyspnea and a feeling of retained secretions, and was then admitted to hospital. She had a severe left convex scoliosis with a mean Cobb angle of over 70°. After a chest X-ray, a CT scan was performed (Fig. 1) which showed atelectasis of the right lower lobe. Her diurnal arterial oxygen saturation (SpO2) fluctuated between 82 and 85% in room air, while PaCO2 was normal. The patients showed minimal clinical signs of dehydration, namely dry mouth. She had good skin turgor and normal urine output. Haematocrit and electrolytes were normal except for potassium which was lower than normal. We began nutritional support and hydration because the patient was not able to eat and drink enough. Negative results were obtained from sputum cultures; however, intravenous antibiotics were administered. Twenty-four-hour NIV was begun, with the addition of oxygen (5 L/min), as NIV alone was not enough to maintain SpO2 above 90%. Fifteen-twenty minutes sessions of HFCWO at a pressure of 5 cm H2O and a frequency of 12 Hz were performed; each session was followed by five or six sessions of mechanical assist cough with an In-Exsufflator at pressures of 10/—+5 cm H2O, delivered respectively over 3 and over 2 s, with an abdominal thrust timed to the exsufflation cycle. This protocol was applied 4 times/day; additionally, In-Exsufflator

Figure 1 Chest CT scan at admission: atelectasis of the lower right lobe.
was used on demand. Once 24-h NIV had begun, we asked the patient to frequently change her decubitus. However, as she had a severe scoliosis, she hardly changed her body position tending to remain on her right side. As the patient demonstrated a modest clinical improvement, we hypothesized that bronchial secretions had a high concentration of DNA due to accumulation of degenerated leukocytes. Therefore, one week after admission we tested rhDNase (Pulmozyme®; Roche, Basel, Switzerland). For three days, 2.5 mg were delivered twice daily with a jet nebulizer, using an in-line nebulizer with NIV, without success. The next day flexible fiberoptic bronchoscopy (FOB) was performed, during NIV plus oxygen, to get a better evaluation of the cause of the obstruction and to possibly instill rhDNase bronchoscopically. FOB showed a lot of very thick mucus in the lower right lobe, but, due to its high viscosity, only a small amount could be removed. Then, a single dose of 2.5 mg rhDNase was instilled directly over the affected area. As the procedure was performed in a clinical ward and the patient was at risk of intubation, this was carried out together with an intensivist. Anyway, we had no complications except for a mild transient decrease in SpO₂. The patient was closely monitored by trained nurses. Copious but thinner secretions were removed with the help of the In-Exsufflater. It had to be used six times in the first hour and three times in the second hour for periods lasting from 1 to 2 min to 10 or more consecutive minutes. In the following hours the In-Exsufflater was applied only according to the protocol and within 24 h a significant clinical improvement was evident, with a reduced need of oxygen. A new CT scan demonstrated a significant reduction of the atelectatic area (Fig. 2). The treatment was repeated two days after the first instillation with the same dose of the drug, with a further clinical improvement that allowed us to withdraw oxygen. Even after the second instillation, Ml-E was often required, four times in the first hour and once in the second hour, unlike in the following days when it was applied only according the protocol. Two days later the patient was discharged (Table 1). A month later the clinical and radiological recovery was confirmed (Fig. 3).

Discussion

This experience illustrates the safety and efficacy of intrabronchial rhDNase treatment in the management of infectious atelectasis in a non-invasively ventilated neuromuscular patient in whom physiotherapy techniques alone did not produce a rapid significant improvement.

To the best of our knowledge, to date rhDNase has not been tested in neuromuscular patients. We initially used the drug in a nebulized form, and switched to intrabronchial instillation after that it was unsuccessful. Direct intratracheal instillation of the drug appeared to be far more

![Figure 2](image)  Chest CT scan at day 12: partial regression of the atelectasis.

![Figure 3](image)  Chest CT scan one month after hospital discharge: complete resolution.

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Relationship of therapeutic measures with clinical outcomes and oxygen need.</th>
</tr>
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<tbody>
<tr>
<td>Time</td>
<td>Mechanical therapy</td>
</tr>
<tr>
<td>1–10 h</td>
<td>HFCWO + In-Exsufflater, NIV day and night</td>
</tr>
<tr>
<td>11th</td>
<td>HFCWO + In-Exsufflater NIV day and night, intrabronchial rhDNase</td>
</tr>
<tr>
<td>12th</td>
<td>HFCWO + In-Exsufflater, NIV day and night</td>
</tr>
<tr>
<td>13th</td>
<td>HFCWO + In-Exsufflater, NIV day and night, intrabronchial rhDNase</td>
</tr>
<tr>
<td>14-15th</td>
<td>HFCWO + In-Exsufflater, NIV night only</td>
</tr>
<tr>
<td>16th</td>
<td>NIV night only</td>
</tr>
</tbody>
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<sup>a</sup> Reducing of dyspnea, relief of sense of retained secretions.
effective than its inhalation. Actually in patients on mecha-
nical ventilation nebulized administration of the drug may
result in a significant deposition in the ventilator tubing. As
we continued our usual physiotherapy treatment protocol
after the rhDNase administration, we cannot confirm that
the drug caused the patient’s recovery from atelectasis;
however, a significant clinical improvement rapidly started
only after the drug instillation. Interestingly, the same
temporal relationship between drug administration and
clinical improvement has been observed in the other clinical
series. In common clinical practice FOB itself is considered
a possible way of facilitating mucus mobilization. However,
a small randomized controlled trial showed that it is no bet-
ter than physiotherapy for resolving lung volume loss. In
addition, atelectasis frequently recurs after bronchoscopy.
Conversely, since the drug was instilled our patient easily
expelled her secretions and progressively improved with no
side effects apart from a mild and short-lived decrease in
SpO2. This treatment was well tolerated by the patient who
was not willing to undergo a long hospitalization.

Best clinical practice in neuromuscular patients usu-
ally requires non-invasive means to manage respiratory
complications; however, intrabronchial instillation of rhD-
Nase could be a supplementary method of treatment. As
rhDNase is an expensive drug, a cost-benefit assessment
would be warranted. We cannot accurately estimate how
long the hospital stay would have been for our patient if rhD-
Nase had not been used. However, compared to the previous
episode of atelectasis, we saved 19 days of hospitalization.
One vial of the drug costs 43 euros, and two vials were used
for local instillation. Since one day’s stay in our hospital
costs 350 euros, the drug would have been cost-effective
even if the stay were reduced by just one day.

In conclusion, treatment with rhDNase demonstrated a
safe option for this neuromuscular patient with atelectasis
who had not responded to normal intensive treatment. Bron-
choscopic instillation gave the best results, liquefying secre-
tions that were easily eliminated with the In-Exsufflator. As
persistent atelectasis may be a life-threatening condition in
neuromuscular patients, we suggest that this drug should be
considered for neuromuscular patients when appropriate.
Further studies are needed in order to confirm its efficacy
and the best method of administration.

Ethical disclosures

Protection of human and animal subjects. The authors
declare that no experiments were performed on humans or
animals for this study.

Confidentiality of data. The authors declare that they have
followed the protocols of their work center on the publica-
tion of patient data and that all the patients included in the
study received sufficient information and gave their written
informed consent to participate in the study.

Right to privacy and informed consent. The authors have
obtained the written informed consent of the patients or
subjects mentioned in the article. The corresponding author
is in possession of this document.

Conflicts of interest

The authors have no conflicts of interest to declare.

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