SHORT REPORT

Ipecac myopathy and cardiomypathy

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Abstract
Two cases of ipecac myopathy, one with associated cardiomypathy are reported. Both patients were young women with eating disorders who came to medical attention because of diffuse muscle weakness. Clinical and electromyographic data suggested ipecac myopathy and muscle biopsies confirmed this diagnosis. One patient had associated clinical and echocardiographic evidence of significant cardiomypathy. The myopathy resolved and the echocardiogram returned to normal after discontinuing the use of ipecac.

Case reports
Patient 1
This was a 26 year old female admitted for evaluation of progressive muscle weakness. At the time of presentation, she had a two month history of progressive difficulty climbing stairs, carrying a laundry basket, and holding her head up while seated at work. She admitted to mild myalgia and had noted mild atrophy of her thigh and arm musculature. She had occasional palpitations, increased sweating, and mild pedal oedema. She denied chest pain or dyspnoea on exertion. She had a history of asthma, mild hypertension, depression and anorexia nervosa. At the time of her father’s death, four years before evaluation, she had lost 18 kilograms, dropping from 61 kilograms to 43 kilograms. Since then, her weight had increased to 57 kilograms and remained stable (height 165 cm). For the six month period before her evaluation, she had ingested several 30 ml doses of syrup of ipecac (containing approximately 21 mgs of emetine) per week to induce emesis after eating. She claimed only occasional alcohol use.

Her general physical examination revealed a well nourished, young, white woman. Her resting heart rate was 104 with S3 and S4 heart sounds present. Lungs were clear. Icthyosis was present on the extensor surface of her arms and legs. Her neurological examination was remarkable for diffuse mild extremity atrophy and moderate diffuse weakness, greater proximally. Her sensory examination was normal. Laboratory evaluation revealed blood counts, serum electrolytes, LDH, SGOT, SGPT, and alkaline phosphatase to all be within normal limits. A rheumatoid factor and antinuclear antibody were negative. A serum creatinine kinase (CK) level was elevated, with a peak at 1027 units/l. An ECG showed sinus tachycardia with a prolonged QT interval and T-wave inversion in the lateral leads. An echocardiogram showed left ventricular dilatation and diffuse hypocontractility with an ejection fraction of 36-5% (see table). Electromyography showed low-amplitude, polyphasic units in proximal muscles, without increased spontaneous activity.

A left deltoid biopsy showed slight variation in fibre size without evidence of inflammation. ATPase and NADH stains showed targetoid changes in both type 1 and 2 fibres with

Table 1 Echocardogram results (patient 1)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Study 1 (15 months after discontinuing ipecac)</th>
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</thead>
<tbody>
<tr>
<td>LVIDd (short axis)</td>
<td>5-1 cm</td>
</tr>
<tr>
<td>LVIDd (short axis)</td>
<td>4-2 cm</td>
</tr>
<tr>
<td>EDV</td>
<td>109-4 ml</td>
</tr>
<tr>
<td>ESD (apical view)</td>
<td>69-5 ml</td>
</tr>
<tr>
<td>EF (apical view)</td>
<td>36-5%</td>
</tr>
<tr>
<td>FS</td>
<td>17-6%</td>
</tr>
<tr>
<td></td>
<td>26-1%</td>
</tr>
</tbody>
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Abbreviations: LVIDd (left ventricular internal dimension, diastole), LVIDs (left ventricular internal dimension, systole), EDV (end diastolic volume), ESV (end systolic volume), EF (ejection fraction), FS (fractional shortening).
Ipecac myopathy and cardiomyopathy

Patient 2
This was a 29 year old, white female who was evaluated for muscle weakness during a psychiatric admission for an eating disorder. She had a several month history of progressive generalised weakness, exercise intolerance and amenorrhea. For the six months before evaluation, she drank approximately one bottle of wine per day. During this period, she had a pattern of minimal food intake alternating with binge eating, always followed by ipecac-induced emesis. General physical examination showed a thin white female with dry skin. Heart and lungs examination were normal. Diffuse moderate muscle atrophy and weakness were present, more prominent proximally. Sensory examination was normal.

Laboratory examination showed blood counts, serum chemistries, thyroid function test, and serum CK to be within normal limits. An ECG showed non-specific T-wave abnormalities. Electromyography showed long, polyphasic motor units with no increased insertional or spontaneous activity.

A biopsy of the left vastus lateralis muscle showed foci of fibres with increased numbers of internal nuclei with minimal atrophy. Trichrome staining showed focal groups of fibres with blurring of the myofibrillar network and aggregations of dense staining material (fig 2). These aggregates were seen by electron microscopy to represent Z-band streaming. NADH staining also showed loss of the myofibrillar network in many fibres, resulting in a targetoid appearance similar to that seen in patient 1. There was no evidence of inflammation, and phosphorylase and myoadenylate deaminase staining showed normal activity.

Discussion
Ipecac is isolated from the root of Cephalis ipecacuanha. It contains two major alkaloids, emetine and cephaeline. Cephaeline is primarily responsible for producing nausea and emesis, whereas emetine is more toxic to skeletal muscle and the heart. Ipecac has long been used by natives of Brazil to treat diarrhoea and has had widespread use in the treatment of amebiasis. It has also been used in aversive therapy for alcoholism and has been employed experimentally as an antineoplastic agent. In the USA, the only official use of ipecac is as an emetic to treat accidental poisoning. It is available without prescription only in syrup form. Through experiences with its use as a treatment for amebiasis, ipecac has been shown to cause a myopathy.

In one study, nearly 5% of high school age females admitted to bulimic activity. They may resort to ipecac abuse if they cannot effectively induce mechanical emesis. It is estimated that there are one million bulimic women in America and that 35 000 abuse ipecac.

Ipecac myopathy should be considered in all young women with progressive proximal muscle weakness, atrophy, or stiffness, especially when there is any suggestion of an eating disorder. Both of our patients had dry skin,
and several other cases have been reported in which patients were noted to have dry or erythematous skin. The serum CK level may be elevated, but it can also be within normal limits in the presence of significant weakness. The electromyogram shows nonspecific myopathic changes, but a muscle biopsy will help distinguish ipecac myopathy from inflammatory or alcohol-induced myopathy. Nerve conduction velocity is normal.

If ipecac abuse is suspected, an evaluation for cardiomyopathy is warranted. Historical or physical evidence of cardiac failure may be present. The electrocardiogram may show resting tachycardia, prolongation of the QT and PR intervals, inverted T-waves, and ST segment abnormalities. As in the case of patient 1, significant depression of left ventricular function may be detected by echocardiogram or radionuclide tests. The serial echocardiograms performed on our patient demonstrated that ipecac cardiomyopathy is largely reversible.

The lethal dose of emetine has been estimated to be 10 to 25 mgs, or approximately 1-25 grams for an adult. Death is usually caused by ventricular arrhythmias or heart failure. Emetine is so slowly excreted from the body that doses may accumulate over time to approximate the lethal dose, even when emesis is stimulated. Syrup of ipecac is available in 30 ml doses, each dose containing approximately 21 mgs of emetine. It is therefore possible to accumulate a lethal amount of emetine within the body over several months when ipecac is used daily. It has been proposed that emetine inhibits protein synthesis and mitochondrial oxidative phosphorylation in skeletal muscle. These effects may cause decreased ATP formation and a subsequent increased release of calcium into the sarcoplasm that, in turn, activates proteases which cause Z-band lysis, granule formation, and myofilament disorganisation. In the heart, emetine is thought to depress glycolysis and Kreb’s cycle, as well as inhibit synthesis of contractile proteins.

The microscopic pathological changes seen in the muscle biopsies from our patients are very similar to those reported in previous cases of ipecac myopathy and experimental emetine-induced myopathy in rats. Light microscopy reveals variation in fibre size, internal nuclei, granular breakdown of myofilaments, and central core changes or targetoid changes in fibres with ATPase or NADH stains. Electron microscopy shows Z-band streaming and dense bodies. Inflammatory infiltrates are not seen. These findings are characteristic of ipecac myopathy, but are also seen in a variety of other conditions. Although patient 2 was a heavy consumer of alcohol, her muscle biopsy was quite similar to that of patient 1 and demonstrated the characteristic targetoid changes and Z-band streaming of emetine myopathy. The characteristic type II fibre atrophy of chronic alcoholic myopathy was absent and therefore her myopathy was likely predominately due to emetine toxicity. The effects on cardiac muscle are not as well documented, but interstitial oedema, inflammation, and degenerative change of myocardial fibres have been reported.

Therapy consists of discontinuing ipecac use and employing supportive measures. If cardiomyopathy is significant, inotropic and afterload reducing agents may be used. As shown in patient 1, both the cardiomyopathy and myopathy improved after discontinuing ipecac abuse. Electrical and pathological evidence of muscle recovery has also been demonstrated.