



Misdiagnosis is an important factor for diagnostic delay in McArdle disease

Renata Siciliani Scalco^{a,b,*}, Jasper M. Morrow^a, Suzanne Booth^a, Sherryl Chatfield^a,
Richard Godfrey^{a,c}, Ros Quinlivan^{a,d}

^a MRC Centre for Neuromuscular Diseases and Department of Molecular Neuroscience, University College London Institute of Neurology and National Hospital for Neurology and Neurosurgery, Queen Square, London WC1N 3BG, UK

^b CAPES Foundation, Ministry of Education of Brazil, Brasilia, DF, Brazil

^c Centre for Human Performance, Exercise and Rehabilitation, Brunel University London, London UB8 3PH, UK

^d Dubowitz Neuromuscular Centre, Great Ormond Street Hospital, London WC1N 3JH, UK

Received 17 January 2017; received in revised form 29 March 2017; accepted 28 April 2017

Abstract

Diagnosis of McArdle disease is frequently delayed by many years following the first presentation of symptoms to a health professional. The aim of this study was to investigate the importance of misdiagnosis in delaying diagnosis of McArdle disease. The frequency of misdiagnosis, duration of diagnostic delay, categories of misdiagnoses and inappropriate medical interventions were assessed in 50 genetically confirmed patients. The results demonstrated a high frequency of misdiagnosis (90%, $n = 45/50$) most commonly during childhood years (67%; $n = 30/45$) compared with teenage years and adulthood (teenage: $n = 7/45$; adult $n = 5/45$; not known $n = 3/45$). The correct diagnosis of McArdle disease was rarely made before adulthood (median age of diagnosis 33 years). Thirty-one patients (62%) reported having received more than one misdiagnosis; the most common were “growing pains” (40%, $n = 20$) and “laziness/being unfit” (46%, $n = 23$). A psychiatric/psychological misdiagnosis was significantly more common in females than males (females 6/20; males 1/30; $p < 0.01$). Of the 45 patients who were misdiagnosed, 21 (47%) received incorrect management. This study shows that most patients with McArdle disease received an incorrect explanation of their symptoms providing evidence that misdiagnosis plays an important part in delaying implementation of appropriate medical advice and management to this group of patients.

© 2017 The Authors. Published by Elsevier B.V. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

Keywords: Glycogen storage disease type V; McArdle disease; Growing pains; Exercise intolerance; Rhabdomyolysis; Myoglobinuria

1. Introduction

McArdle disease (GSDV) is an autosomal recessive disorder characterised by the absence of muscle glycogen phosphorylase. The enzyme deficit results in impaired muscle metabolism with symptoms such as exercise intolerance and muscle pain beginning in childhood. Muscle pain occurs within a few minutes of starting physical activity and can lead to muscle contracture and rhabdomyolysis (RM) if that activity persists or is more vigorous. Muscle contracture and RM in McArdle disease do not just follow exercise and can also be triggered by sustained isometric muscle contraction in everyday activities or ‘unusual’

circumstances [1,2]. RM may result in potential life-threatening complications requiring urgent hospital admission such as compartment syndrome and acute kidney failure (Table 1).

Early diagnosis, ideally when the individual is still a child, is important to facilitate learning the life skills required to manage the condition and prevent RM [3]. Timely diagnosis facilitates appropriate screening, management and prevention of known comorbidities associated with the condition such as sedentariness and obesity [4]. Currently, a correct diagnosis frequently occurs years after first presentation of symptoms [5–8]. This could, in part, be due to its rarity, as doctors might not be familiar with the clinical hallmarks of the condition such as the *second wind* phenomenon, which occurs after about 8–10 minutes of aerobic activity when the symptoms of exercise intolerance (tachycardia, myalgia and fatigue) disappear and the patient can exercise more freely.

* Corresponding author. MRC Centre for Neuromuscular Diseases, Queen Square, London WC1N 3BG, UK. Fax: +0203 448 4725.

E-mail address: r.scalco@ucl.ac.uk (R.S. Scalco).

Table 1
McArdle disease diagnostic features.

McArdle disease features	Medical history/Physical exam
Exertion intolerance	Episodes of muscle pain and tachycardia at the beginning of any physical activity and during strenuous activity, isometric muscle contraction and/or resistance training. All skeletal muscles are involved. In children symptoms reported by parents include: <ul style="list-style-type: none"> • Infancy: difficulty crawling more than a few yards • Toddlers: wanting to be carried/or put in a push-chair all of the time, complaining of pain when walking • Children: <ul style="list-style-type: none"> ○ unable to run (maximum running distance 100 m) ○ unable to keep up with peers ○ collapse/vomiting during sporting activities
Muscle contracture	Severe rigidity with associated pain (patients might report it as “muscle seizes up”, “severe cramp”). Muscle contracture may affect any skeletal muscle, for example the forearm, with activities such as opening cans, picking up heavy pots, carrying shopping.
<i>Second wind</i>	During aerobic activity symptoms improve after 8–10 minutes. The <i>second wind</i> can be identified with functional exercise testing with cardiac monitoring such as the 12 minute walk test or cycle ergometry [7,9–11].
Episodes of rhabdomyolysis/myoglobinuria	Severe muscle contracture followed by muscle swelling and pain; flu-like symptoms Discolouration of urine described as: tea, red wine or coca cola With severe episodes there may be collapse and acute renal failure CK is markedly raised (40,000–250,000 IU/L)
Additional investigation	Baseline serum CK is usually raised (10–15× normal) Serum urate is frequently raised Non-ischemic forearm exercise test shows no significant rise in lactate DNA analysis: Initial screen for common mutations in Northern Europeans (p.Arg50X and p.Gly205Ser) Next tier full <i>PYGM</i> sequencing Muscle biopsy rarely required: vacuolar myopathy, sub-sarcolemmal glycogen deposition and absent muscle glycogen phosphorylase activity

CK, creatine kinase.

To investigate the consequences of disease misdiagnosis in McArdle disease patients, a service evaluation was performed to assess the frequency of misdiagnosis, the duration of diagnostic delay, the categories of misdiagnoses and inappropriate medical interventions.

2. Materials and methods

Clinical information from 50 consecutive patients with genetically confirmed GSDV (median age: 48.14; range: 16–73; 30 male, 20 female) was reviewed as part of service evaluation of a ‘Nationally Commissioned Highly Specialized McArdle’s Disease Service’ based in London. The study was registered and approved by the Hospital’s internal review board/audit committee. As this was a service evaluation, informed consent was not required. Detailed data on diagnosis and misdiagnosis are routinely collected as part of patients’ assessment at the UK specialised service. Further information is also available on NHS referral documentation and GP records/referral letter. Data related to onset of symptoms, year of diagnosis and related misdiagnoses, self-perception of GSDV symptoms and incorrect treatment prescription were collected via medical notes review and patients’ personal experience reports using a standardised pre-agreed data extraction form. All data were anonymised, and individual details and precise description of various misdiagnoses and incorrect treatment that could potentially identify an individual were omitted. In patients where more than one misdiagnosis had occurred, we reported data based on the age at the first misdiagnosis. Non-parametric data are

summarised as median (range). Categorical data are summarised as percentages. Gender differences were assessed using Mann–Whitney U tests for continuous variables and chi-squared tests for proportions.

3. Results

The frequency of misdiagnosis in patients with GSDV was 90% (n = 45/50). First misdiagnosis most frequently occurred during childhood years (67%; n = 30/45), and less frequently during teenage years or adulthood (teenage: n = 7/45; adult n = 5/45; not known n = 3/45). However, ongoing or additional misdiagnoses were common through adult years with the median age of correct diagnosis being 33 years (range 6–70). The median delay in correct diagnosis of GSDV was 29 years (range 0–68). The median time from symptom onset to receiving the first misdiagnosis was 3 years (range 0–67). The median time from the misdiagnosis to correct diagnosis was 23 years (range 1–62). There were no significant gender differences in any of these parameters. The diagnostic delay from the first symptoms to the correct diagnosis appeared to decrease over the decades (Fig. 1A). This decrease was associated with an increase in GSDV diagnostic rates with time (Fig. 1B).

Thirty-one patients (62%) reported having received more than one misdiagnosis, with “growing pains” (40%, n = 20) and “laziness/being unfit” (46%, n = 23) representing the most common misdiagnoses (Fig. 2). A psychiatric/psychological misdiagnosis was significantly more common in females than

Download English Version:

<https://daneshyari.com/en/article/5631981>

Download Persian Version:

<https://daneshyari.com/article/5631981>

[Daneshyari.com](https://daneshyari.com)