Premature coronary artery disease and the Prader-Willi syndrome

S.R. Page1, S.S. Nussey1, G.A. Haywood2 and J.S. Jenkins1

Departments of 1Medicine and 2Cardiology, St George’s Hospital Medical School, Cranmer Terrace, London SW17 0RE, UK.

Summary: A 28 year old woman with the Prader-Willi syndrome developed chest pain and loss of anterior R wave amplitude on the electrocardiogram. Cardiac catheterization demonstrated a severe proximal stenosis of the left anterior descending artery with delayed antegrade flow together with antero-apical akinesia consistent with myocardial infarction. Physicians involved in the management of patients with the Prader-Willi syndrome should be aware of this association with premature coronary artery disease.

Introduction

The Prader-Willi syndrome is a relatively common congenital disorder occurring with an incidence of 1 in 10,000 live births.1 The syndrome is characterized by poor fetal movements and breech presentation; hypotonia and delayed attainment of developmental milestones; hyperphagia and morbid obesity; small hands and feet; hypogonadism and mental retardation.1,2 A deletion from the long arm of chromosome 15 has been described in about 50% of cases.3 Premature coronary artery disease has been reported in only two previous cases of the Prader-Willi syndrome4,5 and in one report the diagnosis was only appreciated at post mortem.6 We report a third case in which coronary angiography demonstrated a severe proximal stenosis of the left anterior descending coronary artery in a 28 year old female with this syndrome.

Case report

A 28 year old female with the Prader-Willi syndrome attended an endocrine follow-up clinic. She complained of a single episode of severe constricting chest pain and breathlessness at rest two weeks previously. This had been followed by intermittent central chest pain of shorter duration, unrelated to exercise, and increasing breathlessness on exertion. There was no history of familial cardiac problems, diabetes or hypertension but she had smoked 5 cigarettes a day for 7 years. Her only medication was triiodothyronine 20 μg three times daily for control of obesity.

The mother’s pregnancy had proceeded to term but delivery was by breech presentation. The neonatal period was complicated by feeding difficulties and generalized hypotonia. Subsequent motor development was considerably delayed. She was unable to sit independently until 18 months and did not walk until 4 years of age. Marked obesity developed from 18 months of age. She had a documented I.Q. of 84 at 5 years of age but required special schooling during adolescence; she remained grossly obese. She presented with delayed puberty and primary amenorrhoea when aged 15 years. Basal gonadotrophin concentrations were low but combined anterior pituitary function tests with luteoliberin (GnRH 100 μg), thyroliberin (TRH 200 μg) and insulin (0.2 U/kg) were normal. The gonadotrophin response to GnRH was considerably enhanced following a 6 week course of clomiphene [pre-clomiphene, peak luteinising hormone (LH) 12.4 mU/l, peak follicle stimulating hormone (FSH) 2.3 mU/l; post-clomiphene, peak LH 40 mU/l, peak FSH 14.0 mU/l]. This, together with the neonatal history, short stature (height 1.37 m), massive obesity (body mass index 55.8 kg/m², normal < 25 kg/m²), small hands and feet and mild mental retardation established the diagnosis of the Prader-Willi syndrome. Oestrogen replacement therapy was commenced when she was 17 years of age but discontinued five years later due to further weight gain. Chromosomal studies using the Trypsin-Giemsa method7 confirmed the presence of a deletion of the q11.1–q11.2 region of the long arm of chromosome 15.

Her pulse was 86 per minute sinus rhythm, blood pressure 130/80 mmHg. The jugular venous pressure was 3 cm and cardiac auscultation was normal. There was no clinical evidence of pul-
monary or peripheral oedema. A chest radiograph revealed a large cardiac silhouette and electrocardiography showed loss of anterior R wave amplitude. Cardiac enzymes and the blood urea and electrolytes were normal. Fasting serum cholesterol and triglycerides were 4.2 mmol/l and 1.1 mmol/l respectively (normal range; cholesterol 3.3–7.3 mmol/l, triglycerides 0.8–2.0 mmol/l) and the fasting blood glucose was 4.6 mmol/l (normal range 3.0–6.0 mmol/l). An exercise electrocardiogram was terminated at stage 2 of the Bruce protocol by chest pain and exhaustion. There was no significant ST segment depression or arrhythmia. In view of the evidence suggesting ischaemic heart disease cardiac catheterization was performed. The left ventriculogram showed antero-apical akinesia consistent with myocardial infarction and angiography revealed a severe stenosis of the left anterior descending coronary artery with delayed antegrade flow to the distal vessel (Figure 1). The remaining coronary arteries were normal. Since starting diltiazem 60 mg three times daily she has remained free of chest pain.

**Discussion**

Morbidity and mortality in the Prader-Willi syndrome usually result from the complications of massive obesity. Impaired pulmonary function and diminished responses to carbon dioxide are associated with the gradual development of the obesity-hypoventilation syndrome characterized by somnolence, hypoxia and carbon dioxide retention with resulting cardiorespiratory failure. Pulmonary embolic death and death following surgical procedures have been reported as other causes of morbidity and mortality though there are few post-mortem reports available in the world literature. In the present case the angiographic findings clearly demonstrate single vessel coronary artery disease associated with myocardial infarction in a female at 28 years of age. Premature coronary artery disease is not acknowledged as a cause of mortality or morbidity in any of the major reviews of this syndrome and we are aware of only two previous cases of premature coronary artery disease in association with the Prader-Willi syndrome in the world literature.

Non-insulin dependent diabetes related to gross obesity has been reported in approximately 7% of cases of the Prader-Willi syndrome. Poorly controlled diabetes together with serum cholesterol concentrations near the upper limit of the normal range were considered risk factors for coronary artery disease in one of the previous case reports. The present case, however, had no evidence of diabetes or hypertension and the fasting blood cholesterol concentration of 4.2 mmol/l was in the range considered optimal for the prevention of ischaemic heart disease. In addition there was no significant family history of heart disease. Obesity is a universal feature of patients with the Prader-Willi syndrome. However, it is generally accepted in the absence of hypertension and increased serum total cholesterol concentrations, that obesity is not an independent risk factor for coronary artery disease. Therefore the only established independent risk factors for the development of premature coronary artery disease in the present case were a history of modest smoking and physical inactivity.

Loss of endogenous oestrogens due to premature menopause is associated with an increased risk of coronary artery disease due in part to adverse effects on the serum lipid profile. In the present case, oestrogen replacement therapy for hypop-
gonadism had been prescribed only between 17 and 22 years of age. However, in view of the normal serum lipids, we believe it is unlikely that the subsequent lack of oestrogen therapy was a significant risk factor for coronary artery disease.

The absence of significant risk factors in this case suggests that premature coronary artery disease may be a direct and largely unrecognized feature of the Prader-Willi syndrome. It is interesting to speculate that the chromosomal deletion from the q11.1–q11.2 region of the long arm of chromosome 15 which is commonly associated with the Prader-Willi syndrome may be also implicated in the development of premature arterial disease.

References

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