Eruptive xanthomatosis

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Background
A 33-year-old woman with diet-controlled type 2 diabetes presented with pruritic yellow erythematous papules on her back, arms, legs and buttocks (Figure 1). Glycaemic control had gradually deteriorated over the last one year with a HbA1c reaching 134 mmol/mol. The non-fasting lipid profile had deteriorated acutely from being on-target to a total cholesterol level of 21 mmol/L and triglycerides of 69 mmol/L.

Figure 1 shows severe xanthoma usually seen with a genetic disorder, which may or may not be exacerbated by diabetes or hypothyroidism. Such cases require appropriate investigation, including exploration of family history, alcohol use and ApoE genotyping. Treatment differs according to the underlying diagnosis. Figure 2 demonstrates lipaemia retinalis in the same patient.

Pathology
Eruptive xanthomatosis is a papular cutaneous manifestation of hypertriglyceridaemia which can occur due to a primary genetic defect, or secondary to another disorder such as diabetes, hypothyroidism, liver cirrhosis, nephrotic syndrome or alcohol excess.1 A xanthoma is a dermal deposition of lipid-rich macrophages. Eruptive xanthomas normally resolve rapidly and completely with treatment of the underlying disorder.

In the context of diabetes, hypertriglyceridaemia occurs due to reduced chylomicron clearance leading to an increase in very low density lipoproteins, or lack of insulin (or insulin resistance) resulting in an acquired lipoprotein lipase deficiency.2 Hypertriglyceridaemia needs to be excluded in patients who present with lesions having characteristics of eruptive xanthomas. Management in the context of diabetes should focus on adoption of a low-fat diet, weight reduction and rapid improvement of glycaemic control before considering the use of fibrates or statins.3 Severe hypertriglyceridaemia is associated with increased risk of pancreatitis and requires urgent treatment.

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