Obstructive sleep apnea in 2 women with familial partial lipodystrophy due to a heterozygous LMNA R482Q mutation

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Although obstructive sleep apnea has many attributes of a complex genetic trait, few genetic mutations have been identified in patients with the condition. Obesity has been shown to be associated with obstructive sleep apnea in the general population, but few rare genetic syndromes affecting total body adipose content or distribution have been evaluated for possible association with obstructive sleep apnea. Familial partial lipodystrophy subtype 2 most often results from mutations in LMNA, the gene encoding nuclear lamin A/C (MIM 151660). Patients with familial partial lipodystrophy subtype 2 are healthy at birth, but around puberty they lose fat stores selectively in the extremities and gluteal region, while visceral, facial and neck fat stores are preserved and may enlarge with caloric excess. Familial partial lipodystrophy subtype 2 has been called “fat neck” syndrome in the past. Using polysomnography, we have documented the presence of obstructive sleep apnea in 2 women with familial partial lipodystrophy subtype 2 who each presented with daytime somnolence.

Patient 1 was a 49-year-old woman of normal weight in whom familial partial lipodystrophy subtype 2 had been diagnosed at age 30 years. She was 161 cm tall, her weight was 72.5 kg, and she had a body mass index of 27.9 kg/m², a waist circumference of 88 cm and blood pressure of 145/88 mm Hg. Patient 2 was a 46-year-old woman of normal weight in whom the lipodystrophy had been diagnosed 1 year earlier. Her height was 155 cm, weight 59.9 kg, body mass index 25.9 kg/m², waist circumference 86 cm and blood pressure 116/60 mm Hg. Both patients had reasonably well-controlled diabetes and dyslipidemia. Both were heterozygous for the LMNA R482Q mutation. The 2 women reported long-standing daytime somnolence and loud snoring at night but no witnessed episodes of obstructive sleep apnea. Neither had a history of allergies, cardiovascular disease or respiratory problems. Patient 1 had 6 female first- and second-degree relatives aged 40–60 years who had the LMNA R482Q mutation; patient 2 had 2 such relatives. None of these relatives had received a diagnosis of obstructive sleep apnea or had symptoms suggestive of it. Both patients had the classic familial partial lipodystrophy fat distribution, with no subcutaneous fat in the limbs and gluteal region, and increased facial, neck and central fat stores (Figure 1). Neither patient had acanthosis nigricans, hirsutism or hepatomegaly, and both had normal dentition, tongue size, pharynx and palate, including uvula and tonsils. In each case, examination of the eyes and of the respiratory, cardiovascular and neurologic systems yielded normal findings, as did resting and exercise electrocardiograms.

Neither patient had thyromegaly or other neck masses felt clinically, and each had normal thyroid function, as assessed by laboratory testing.

Overnight polysomnography was performed according to standard protocols (www.sleeplaboratories.com/physician-5.htm#5-1). Patient 1’s total sleep time was 332 minutes, with 32 apneic and hypopneic episodes. Patient 2’s total sleep time was 391 minutes, with 36 apneic and hypopneic episodes. The apnea–hypopnea index was 5.8 for patient 1 and 5.5 for patient 2, values consistent with mild obstructive sleep apnea. All apneic and hypopneic events were considered to be obstructive rather than central events.

We used a semi-automated algorithm to quantify the subcutaneous fat in a compartment defined as the region of the neck anterior to the rectilinear tangential line at the dorsal most point of the paratracheal region. The proportion of the area of the anterior compartment of the neck that was adipose tissue was 56.8%, 68.7% and 44.1% for patient 1, patient 2 and an age- and sex-matched healthy female control subject who had no lipodystrophy. Semi-automated quantification of subcutaneous fat from T1-weighted magnetic resonance images at vertebral level C5 showed that both patients had more than 1.6 times the adipose tissue content in the anterior compartment of the neck surrounding the trachea as compared with the control subject (Figure 1).

Sleep-disordered breathing in patients 1 and 2 was subsequently controlled with nightly continuous positive airway pressure of 10 and 5 cm H₂O, respectively, administered by nasal mask. Both patients reported marked improvement in their symptoms.

Charcot–Marie–Tooth disease, another monogenic disorder, is also associated with obstructive sleep apnea, but...
the sleep apnea in lipodystrophy is not associated with neuropathy or muscle weakness. Furthermore, obstructive sleep apnea has been reported in patients with acquired lipodystrophy syndromes, specifically partial lipodystrophy associated with HIV infection treated with highly active antiretroviral drugs.  

No clear mechanism links familial partial lipodystrophy subtype 2 with obstructive sleep apnea, although both of the patients we have described were of normal weight but had markedly increased neck fat content, specifically in the region surrounding the trachea. This suggests that the association might be related to the repartitioning of adipose tissue that is characteristic of patients with this type of lipodystrophy.

Although the obstructive sleep apnea in these 2 cases may have been coincidental, it is important to note that neither of the patients was obese and each had clinically ascertained lipodystrophy. Anecdotally, we have heard of additional cases of sleep apnea in extended kindreds. A more systematic assessment of the presence of obstructive sleep apnea in people with familial partial lipodystrophy or other lipodystrophies...
would increase confidence in the possible mechanistic association of these conditions.

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