Point mutation of the mitochondrial tRNA<sub>Leu</sub> gene (A 3243 G) in maternally inherited hypertrophic cardiomyopathy, diabetes mellitus, renal failure, and sensorineural deafness

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Abstract

The A 3243 G mutation of the mitochondrial tRNA<sub>Leu</sub> gene was found to segregate with maternally inherited diabetes mellitus, sensorineural deafness, hypertrophic cardiomyopathy, or renal failure in a large pedigree of 35 affected members in four generations. Presenting symptoms almost consistently involved deafness and recurrent attacks of migraine-like headaches, but the clinical course of the disease varied within and across generations. The A 3243 G mutation has been previously reported in association with the mitochondrial encephalomyopathy, lactic acidosis, and stroke-like episode syndrome (MELAS) and with diabetes mellitus and deafness. To our knowledge, however, hypertrophic cardiomyopathy is not a common feature in people with the A 3243 G mutation and renal failure has not been hitherto reported in association with this mutation. The present observation gives additional support to the variable clinical expression of mtDNA mutations in humans.

In the last few years, several mitochondrial DNA (mtDNA) mutations have been described in association with maternally inherited human diseases. Among them, the heteroplasmic A 3243 G mutation in the tRNA<sub>Leu</sub> gene has been described in mitochondrial encephalomyopathy, lactic acidosis, and stroke-like episodes (MELAS), in non-MELAS associated phenotypes including progressive external ophthalmoplegia (PEO), and in maternally inherited diabetes mellitus and hearing loss. Here, we describe hypertrophic cardiomyopathy, renal failure, and endocrine dysfunction as uncommon manifestations of the A 3243 G mutation. This observation gives additional support to the variable clinical expression of mtDNA mutations in human.

III-13 was referred for a personal and family history of hypertrophic cardiomyopathy, diabetes mellitus, and deafness (fig 1). Hearing loss occurred in the third decade, diabetes mellitus at the age of 36 years (requiring insulin administration four years later), and severe concentric hypertrophic cardiomyopathy at 44 years of age. He also had frequent migraine-like headaches and general weakness but no renal or ocular involvement was noted.

III-13 belongs to a large family of 40 affected subjects with maternally inherited multigenerational involvement over four generations (fig 1). Affected subjects were of both sexes and all had recurrent headaches and hearing loss. Several patients (14/40) had adult onset insulin dependent diabetes mellitus (IDDM) and renal failure, originally ascribed to IDDM. Renal failure was, however, present in one patient without diabetes mellitus (II-4). Focal sclerosis of glomeruli was observed in a needle biopsy of the kidney (III-12). Several people had severe cardiac involvement, including fatal hypertrophic cardiomyopathy (3/40, III-11, III-12, III-24), heart block (2/40, II-1, II-3), or sudden death (2/40, II-4, II-5). Neurological involvement included general seizures (2/40, II-5, III-5), brain infarctions, mild mental retardation, ataxia, and cerebellar atrophy (1/40, III-26) or calcification of the basal ganglia (1/40, II-5). Other endocrine manifestations included thyroid goitre (6/40) without thyroid insufficiency or hyperthyroidism (II-1, II-4, II-6, III-7, III-16, III-18), hypoparathyroidism (1/40, II-5), and hypothalamic hypocorticism (1/40, III-37). All affected females had hyperproctisis. Liver cirrhosis without evidence of alcoholism was present in II-8 and III-19. Mental retardation, cerebellar ataxia (1/40) and muscle weakness (2/40) were seldom observed. All children of generation IV (except IV-27, IV-28, IV-29, and IV-42) were asymptomatic at the time of examination, but IV-27, IV-28, and IV-29 had headaches and hearing loss. IV-42 suffered from recurrent “pallor attacks” without abnormal movements or mental retardation from 18 months of age. Interestingly,
Figure 1  Segregation of the A 3243 G mutation in a four generation pedigree.

while I-3 had hearing loss only, 8/9 affected subjects in generation II had deafness plus one (or more) additional organ involvement (diabetes mellitus, renal failure, cardiomyopathy) and 5/9 died before the age of 65 years. In generation III, hearing loss occurred before 30 years of age while it occurred after the age of 40 years in II-4, II-6 and II-9.

Fibres of decreased size with few internal nuclei and a number of intermyofibrillar and subsarcolemmic basophilic deposits were observed in the muscle biopsy of patient III-13. PAS staining showed abundant glycogen matrix and red oil O staining fatty infiltrations. The modified Gomori stain showed numerous ragged red fibres and histoenzymological studies showed normal ATPase activity, non-homogeneous NADH-tetrazolium reductase activity (predominantly in subsarcolemmic regions), and, in ragged red fibres, increased SDH but decreased COX activities.

Screening for the A 3243 G mutation was carried out using oligonucleotide primers nt 3214–3238 and nt 3779–3803 (amplification conditions were: 95°C for 30 seconds, 55°C for 30 seconds, 72°C for 30 seconds). Amplification products (590 bp) were either digested with ApaI or submitted to direct sequencing on an automatic DNA sequencer (Applied Biosystems 373A) using Taq Dye Deoxy TM Terminator cycle sequencing kit (Applied Biosystems). Heteroplasmy for the A 3243 G mutation was shown by ApaI digestion and direct sequencing of amplified mtDNA from patient III-13 skeletal muscle and leucocytes. A heteroplasmic A 3243 G mutation was also present in circulating leucocytes of patients II-6, II-7, II-9, III-16, III-18, III-23, III-26, III-36, III-37, III-38, IV-42, IV-43, and IV-44 (fig 2). The relative amount of normal
and mutant mtDNA was quantitated by Southern blot and densitometry analysis of Apal restriction fragments in lymphocytes (fig 2). No correlation between the amount of mutated molecules and the clinical profile of the patients could be observed, but an unaffected obligate carrier (III-25) did not apparently bear the A 3243 G mutation in his circulating leucocytes.

In this hitherto unreported four generation pedigree (40 patients) the association of symptoms (sensorineural hearing loss, migraine-like headaches, diabetes mellitus, hypertrophic cardiomyopathy, renal failure, and thyroid goitre) along with the maternal inheritance of the disease were suggestive of a mitochondrial disorder. Histopathological and molecular investigations in III-3 allowed us to ascribe this maternally inherited disease to a point mutation of the mitochondrial tRNA^{Leu} gene (A 3243 G).

The A 3243 G mutation of the mitochondrial tRNA^{Leu} gene has been previously reported in patients with MELAS, PEO, A dystonia, and occasionally with hypertrophic cardiomyopathy. Hitherto, however, renal failure, thyroid goitre, or hyperparathyroidism have not been reported in association with the A 3243 G mutation. The present observation emphasises the variable clinical expression of the A 3243 G mutation, a variability which could be ascribed to random segregation of mutated mitochondria in the different tissues, to a particular mtDNA haplotype, to the nuclear genetic background, or even to epigenetic phenomena. It is worth noting, however, that a variety of clinical symptoms have been reported in association with other tRNA^{Leu} mutations, including myoclonic epilepsy and ragged red fibres (C 3256 T), hypoparathyroidism and diabetes mellitus (A 3252 G), and myopathy with cardiomyopathy (A 3260 G).

All affected subjects tested bore the A 3243 G mutation in their circulating leucocytes. Again, the failure to detect the tRNA^{Leu} mutation in leucocyte DNA of III-25 can be ascribed to random drift. Yet, why the disease apparently worsened with generations is unknown. Although this phenomenon may be because of ascertainment bias, it could also be related to either random drift or replicative advantage of mutant mtDNA molecules, as shown in the MELAS syndrome. It appears therefore that the clinical expression of the A 3243 G mutation is particularly broad. This mutation should be considered in patients presenting an unexplained association of symptoms with an early onset and a progressive course, when hearing loss or diabetes mellitus is present.