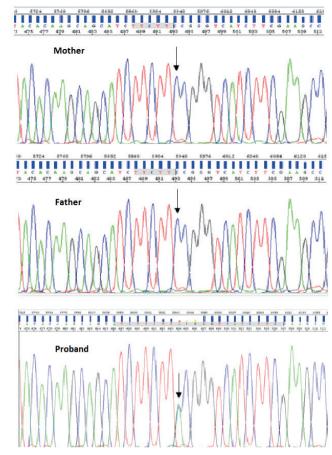
Supplementary material to article by L. Guerra et al. "Keratoderma-Deafness-Mucocutaneous Syndrome Associated with Phe142Leu in the GJB2 Gene"



**Fig. S1. Mutation identification.** Electropherograms of *GJB2* exon 2 sequence encompassing the c.426C>A variant, present at the heterozygote state in the proband and absent in her parents (*arrows*).