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sample of ageing dementia with hearing loss. Dementia with hearing loss (DwHL) is a common and frequent sequela to Alzheimer's disease (AD) and Lewy body disease (LBD). If a link with AD, LBD, or with the neurological manifestations of mitochondrial complex 1 deficiency (C1D) and deficiency of the ATP synthase (ATP) remains to be proven, it is not known if mitochondrial genome mutations are important in the pathogenesis of this disorder. In a population sample with chronic idiopathic bilateral sensorineural hearing loss, we sequenced the mitochondrial genome of a panel of 64 clinically defined individuals with clinical signs of AD, LBD, and C1D. The most frequent mutation, 16168A > G in the NADH dehydrogenase (ND)6 gene, was found in 8.7% of patients, i.e. in 4.2% of those who fulfilled clinical and neuropathological criteria for C1D. The other mutations found in the ND6 gene, 16180T > G 50b96ab0b6

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