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A number sign (#) is used with this entry because of evidence that autosomal recessive deafness-1A (DFNB1A) is caused by homozygous or compound heterozygous mutation in the GJB2 gene (), which encodes the gap junction protein connexin-26 (CX26), on chromosome 13q12. Autosomal dominant deafness-3A (DFNA3A; 601544) is an allelic disorder. See also DFNB1B (), which is caused by mutation in the GJB6 ...

DEAFNESS, AUTOSOMAL RECESSIVE 1A; DFNB1A - OMIM

Password requirements: 6 to 30 characters long; ASCII characters only (characters found on a standard US keyboard); must contain at least 4 different symbols;

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Nakamura et al. (1995) reported 2 families with autosomal recessive hereditary spastic paraplegia, mental impairment, and thin corpus callosum. In the first family, 3 affected brothers had onset in the second decade of gait disturbance resulting in wheelchair use by age 21 years.

OMIM Entry - # 604360 - SPASTIC PARAPLEGIA 11, AUTOSOMAL ...

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