MUTATION (ITALIAN EDITION)

Miles Fischel

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J Peripher Nerv Syst. Sep;17(3) doi: /j x. A novel autosomal dominant GDAP1 mutation in an Italian CMT2 family.

J Cardiovasc Med (Hagerstown). Aug;14(8) doi: /JCM. 0bea R25C mutation in the NKX gene in Italian patients affected.

Search for mutations in a cystic fibrosis patient, compound heterozygous for –1G?A and another uncharacterized molecular defect, revealed the presence.

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The presence of a specifically mutated XPD allele, irrespective of its homozygous, hemizygous, or heterozygous condition, was always associated with the same degree of cellular UV hypersensitivity. These observations are consistent with those obtained by fluorescence complementation assay. Int J Cancer 72 : — ClinicallyknownasETC,Pfizerdidnotmovetrialsforward,probablybecaus Such drugs could benefit people in chronic painwho often struggle to find relief with existing treatments. Close Figure Viewer. Surprisingly,however,theseverityoftheclinicalsymptomsdidnotcorrel 2.