14365-Review

BY FRANCESCA MARINI

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Genetic test in MEN1 syndrome: An evolving story

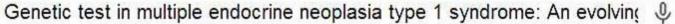
Francesca Marini, Francesca Giusti, Maria Luisa Brandi

11 Abstract

Multiple endocrine neoplasia type 1 (MEN1) is an autosomal dominant inherited tumour syndrome expressing various endocrine and non-endocrine lesions and tumours. Since the identification of the causative gene, the oncosuppressor gene MEN1, in 1997, genetic testing has revealed an important approach for the early and differential diagnosis of the disease. The finding of a MEN1 mutation in a patient has important clinical implications for relatives since it allows very early disease diagnosis and identification of carriers, even

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