

2018 - 1

Lysosomal storage disorder gene variants in multiple system atrophy

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Brain

Nr. 7(141) / 2018 / ISSN 0006-8950 / ISSN e 1460-2156

Disponibil online 20 July, 2021. Descarcări-0. Vizualizări-292

2017 - 1

Genotype-phenotype correlations and expansion of the molecular spectrum of AP4M1-related hereditary spastic paraplegia

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Orphanet Journal of Rare Diseases

Nr. 1(12) / 2017 / ISSN 1750-1172

Disponibil online 11 December, 2017. Descarcări-1. Vizualizări-731
