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Publications (1)

Vogerd M, Malin J, Tegenthoff M, Lindemuth R, Pongratz D, Tettenborn B, Mortier W, Reichmann H, Burwinkel B, Kubisch C, Kilimann M. Mutation analysis in myophosphorylase deficiency (McArdle's disease). *Annals of Neurology* 1998; 43:326-331.

Projects (0)

No results found.

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