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Publications (3)

Poncet A, Smirnov V, Dollfus H, Defoort-Dhellemmes S, Jobic F, Schorderet D, Devos A, Le Moing A, Todorova M, Bocquet B, Pelletier V, Drumare I, Meunier I, Vaclavik V, Grunewald O, Dhaenens C. Contribution of Whole-Genome Sequencing and Transcript Analysis to Decipher Retinal Diseases Associated with MFSD8 Variants. *Int J Mol Sci* 2022; 23:4294.

Habibi I, El Matri L, El Matri K, Turki A, Helfenstein M, Vaclavik V, Wyrsh S, Todorova M, Falfoul Y, Schorderet D. Correction: Clinical and Genetic Findings of Autosomal Recessive Bestrophinopathy (ARB). *Genes (Basel)*. 2020; 11:503.

Habibi I, El Matri L, El Matri K, Turki A, Helfenstein M, Vaclavik V, Wyrsh S, Todorova M, Falfoul Y, Schorderet D. Clinical and Genetic Findings of Autosomal Recessive Bestrophinopathy (ARB). *Genes* 2019; 10:1-17.

Projects (0)

No results found.

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