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Publikationen (2)

Delas F, Koller S, Feil S, Dacheva I, Gerth-Kahlert C, Berger W. Novel Mutation in Conserved Ultraviolet-Protective Tryptophan (p.Trp131Arg) Is Linked to Autosomal Dominant Congenital Cataract. Int J Mol Sci 2023; 24

Fierz F, Landau K, Kottke R, Wichmann W, Sturm V, Weber K, Gerth-Kahlert C. The "Eyelet Sign" as an MRI Clue for Inflammatory Brown Syndrome. J Neuroophthalmol. 2021

Projekte (0)

Keine Resultate gefunden.

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