



## Christina Gerth-Kahlert

### Contact

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## Publications (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

## Projects (0)

No results found.

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