Supplementary material for Mendez et al., "Oculocutaneous albinism type 1B associated with a functionally significant tyrosinase gene polymorphism detected with Whole Exome Sequencing", *Ophthalmic Genetics*, 2021.

Table S1. OCA gene panel

AP3B1, AP3D1, BLOC1S3, BLOC1S6, C10ORF11, DTNBP1, FRMD7, GPR143,

HPS1, HPS3, HPS4, HPS5, HPS6, LYST, MC1R, MLPH, MYO5A, OCA2, OCA5,

RAB27A, SLC24A5, SLC45A2, TYR, TYRP1