Supplementary Material: First Case Report of Primary Carnitine Deficiency Manifested as Intellectual Disability and Autism Spectrum Disorder

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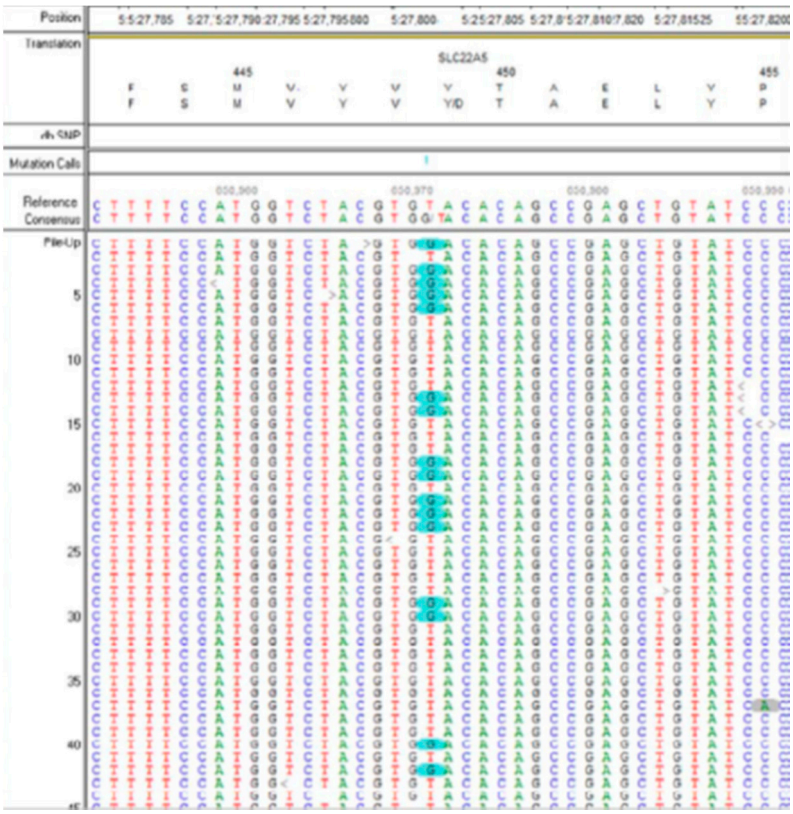

Figure S1. Identification of c.1345T>G (p.Y449D, p.Tyr449Asp) mutation in the *SLC22A5* gene.