Family pedigree and electropherogram of heterozygous IVS11-2A>C(c.1957-2A>C) mutation in the GLI2 gene. Full-black filled box indicates index case with Culler-Jones syndrome phenotype, shaded boxes indicate father and brother who are also heterozygous for the identical mutation with incomplete phenotype, empty boxes indicate mother and sister with wild type.

Ectopic Posterior Pituitary, Polydactyly, Midfacial Hypoplasia and Multiple Pituitary Hormone Deficiency due to a Novel Heterozygous IVS11-2A>C(c.1957-2A>C) Mutation in the GLI2 Gene
Demiral M et al.
Page: 319-328
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