Supporting Information for '**Damaging variants in *FOXI3* cause microtia and craniofacial microsomia'**

This appendix contains Supplemental Figures S1-2.

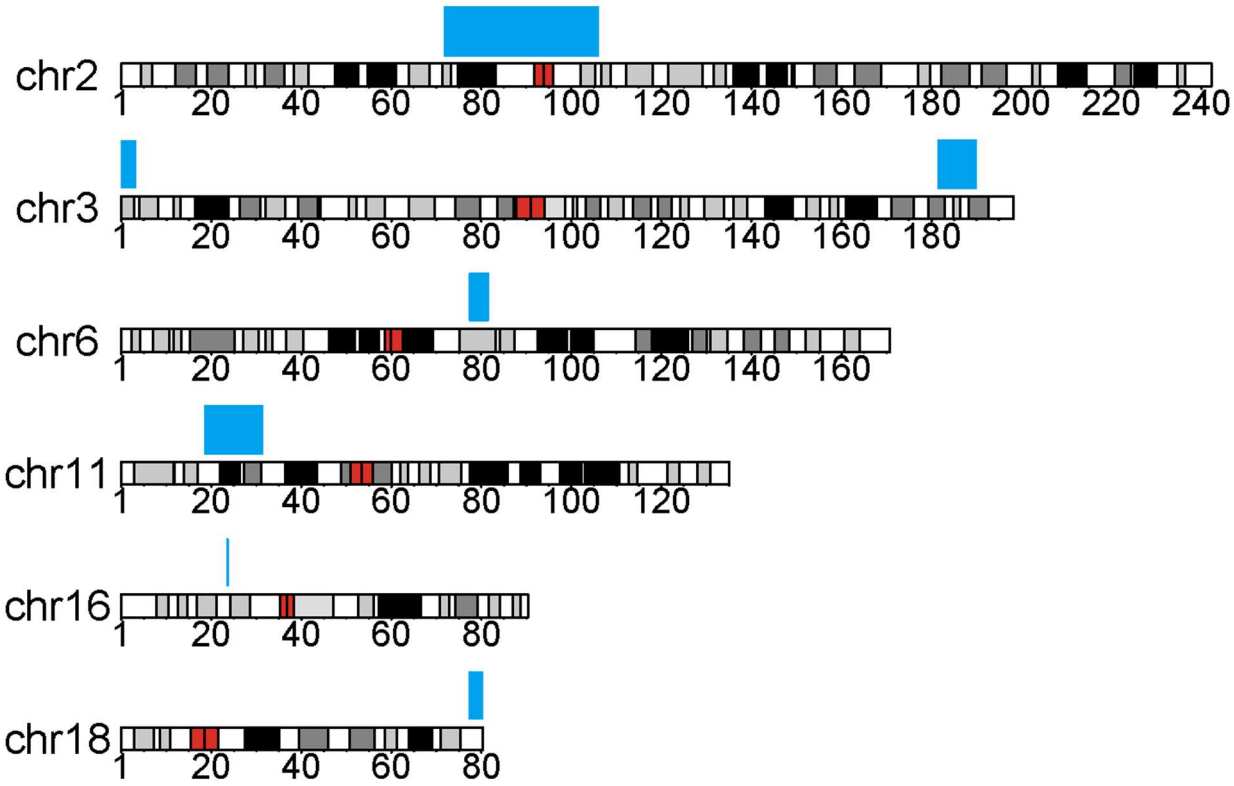

Figure S1. Mapping shared haplotypes in Kindred 1 using rare variants. Locations of shared heterozygous haplotypes in 4 individuals from Kindred 1 (II-5, IV-1, IV-3, IV-6) with WGS are shown in blue. Coordinates are in hg38.

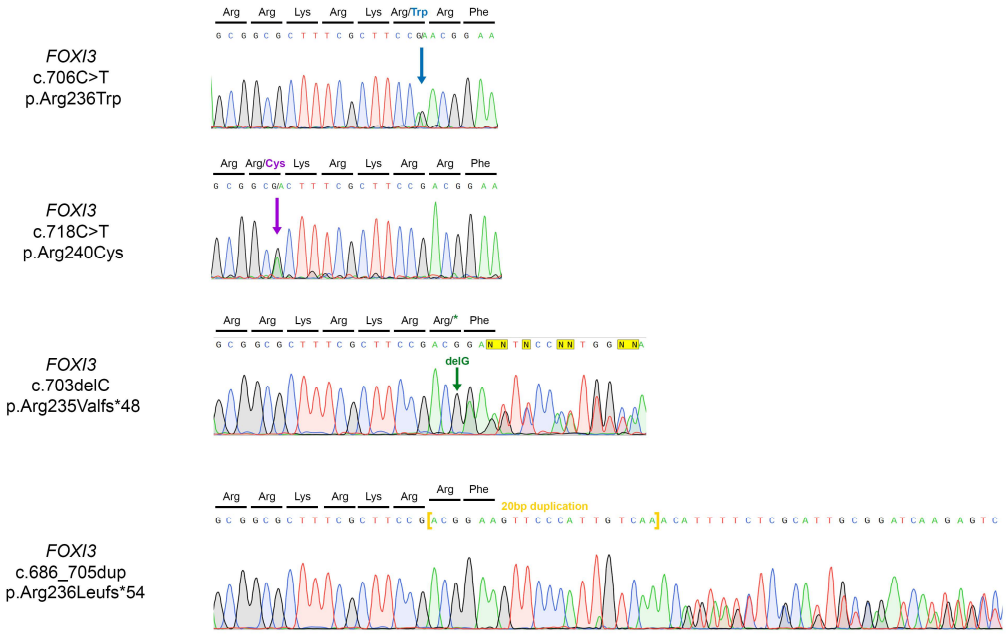

Figure S2. Confirmation of *FOXI3* variants by Sanger Sequencing.