

**This table includes regions of low-quality coverage for the Holoprosencephaly panel and therefore variant calling may be limited.**

<b>Gene</b>	<b>Genomic Region (Hg19)</b>	<b>Gene Region</b>	<b>Canonical Transcript</b>
<i>PTCH1</i>	chr9:98270589-98270643	Exon 1	NM_001083603.5
<i>ZIC2</i>	chr13:100634319-100634672	Exon 1	NM_007129.5

If you are a clinician and have a strong clinical suspicion of a disorder associated with a condition in a gene listed in this table, please contact the laboratory and additional orthogonal sequencing may be considered.