

SUPPLEMENTARY TABLE S1. LIST OF GENES RELATED TO THYROID DISORDERS

<i>Gene</i>	<i>Condition</i>
<i>ALB</i>	Familial dysalbuminemic hyperthyroxinemia (FDH)
<i>ALMS1</i>	Alstrom syndrome
<i>CDCA8</i>	Thyroid dysgenesis
<i>DIO1</i>	High T4
<i>DIO2</i>	High T4
<i>DIO3</i>	Low rT3
<i>DUOX1</i>	Thyroid dysmorphogenesis
<i>DUOX2</i>	Thyroid dysmorphogenesis
<i>DUOXA2</i>	Thyroid dysmorphogenesis
<i>FGF8</i>	Combined pituitary hormone deficiency
<i>FOXE1 (TTF2)</i>	Thyroid dysgenesis
<i>GLIS3</i>	Thyroid dysgenesis
<i>HHEX</i>	Thyroid dysgenesis
<i>IGSF1</i>	Central hypothyroidism
<i>IRS4</i>	Central hypothyroidism
<i>IYD (DEHAL1)</i>	Thyroid dysmorphogenesis
<i>JAG1</i>	Thyroid dysgenesis
<i>KDM6A</i>	Kabuki Syndrome
<i>KMT2D</i>	Kabuki Syndrome
<i>NCOR2</i>	(Corepressor of TH receptors)
<i>NKX2-1 (TTF1)</i>	Thyroid dysgenesis
<i>NKX2-3</i>	Thyroid dysgenesis
<i>NKX2-5</i>	Thyroid dysgenesis
<i>NKX2-6</i>	Thyroid dysgenesis
<i>NTN1</i>	Thyroid dysgenesis
<i>PAX8</i>	Thyroid dysgenesis
<i>POU1F1</i>	Combined pituitary hormone deficiency
<i>PROP1</i>	Combined pituitary hormone deficiency
<i>PTRH2</i>	Infantile-onset multisystem neurologic, endocrine, and pancreatic disease
<i>RYR2</i>	Hyperemesis
<i>SALL1</i>	Townes–Brocks syndrome
<i>SECISBP2</i>	SBP2 deficiency
<i>SERPINA7(TBG)</i>	TBG deficiency
<i>SLC16A2(MCT8)</i>	MCT8 deficiency
<i>SLC26A4(PDS)</i>	Thyroid dysmorphogenesis
<i>SLC5A5(NIS)</i>	Thyroid dysmorphogenesis
<i>SLC30A10</i>	SLC30A10 deficiency
<i>SLCO1C1 (OATP1C1)</i>	OATP1C1 (T4 transporter) deficiency
<i>STAMBP</i>	Microcephaly-capillary malformation syndrome
<i>TBLIX</i>	Hypersensitivity to thyroid hormone
<i>TBX1</i>	Thyroid dysgenesis
<i>TG</i>	Thyroid dysmorphogenesis
<i>THRA</i>	Resistance to TH
<i>THRB</i>	Resistance to TH
<i>TPO</i>	Thyroid dysmorphogenesis
<i>TRH</i>	Central hypothyroidism
<i>TRHR</i>	Resistance to TRH
<i>TRIP11</i>	(Interactor of TH receptor)
<i>TRIP12</i>	(Interactor of TH receptor)
<i>TSHB</i>	Central hypothyroidism
<i>TSHR</i>	Thyroid dysgenesis/resistance to TSH
<i>TTR</i>	Dystransthyretinemic hyperthyroxinemia
<i>UBR1</i>	Johanson–Blizzard syndrome