

## Lissencephaly Panel

### Sequence Analysis and Exon-Level Deletion/Duplication Testing of 26 Genes

**Panel Gene List:** ACTB\*, ACTG1, ARX\*, ATP6V0A2, B3GALNT2, B4GAT1\*\*, CIT, DCX, FKRP\*\*, FKTN, GMPPB, ISPD, KATNB1, LAMB1, LARGE1, NDE1, PAFAH1B1, POMGNT1\*, POMGNT2, POMT1, POMT2, RELN, TMEM5, TUBA1A\*, VLDLR, WDR62

\*Only large deletion/duplications may be detected for the ACTB, ARX, POMGNT1 and TUBA1A genes

\*\*No deletion/duplication analysis for the B4GAT1 and FKRP genes

#### Clinical Features:

Lissencephaly is characterized by a thickened cortex and the absence of folds or gyri (agyria) or the presence of abnormally wide gyri (pachygyria). Lissencephalies can present in different forms and with varying severity, ranging from complete agyria to mixed agyria and pachygyria, or simplified gyri with subcortical band heterotopia (double cortex). Pathogenic variants in distinct genes lead to overlapping forms of lissencephaly, including classic smooth lissencephaly, cobblestone lissencephaly, lissencephaly with agenesis of the corpus callosum, and lissencephaly with cerebellar hypoplasia. Subcortical band heterotopia is a mild form of lissencephaly characterized by normal gyri but the presence of an abnormal and often symmetric band of gray matter under the cortex. Lissencephalies uniformly cause developmental delay, epilepsy, and intellectual disability.<sup>1</sup> The spectrum of disorders associated with lissencephaly includes a group of congenital muscular dystrophies called the alpha-dystroglycanopathies including Walker-Warburg syndrome which is characterized by cobblestone lissencephaly, cerebellar abnormalities, eye defects, muscle weakness.<sup>2,3</sup>

#### Genetics:

The various forms of lissencephaly demonstrate clinical and genetic heterogeneity.<sup>1,2</sup> Classic lissencephaly occurs as an autosomal dominant, autosomal recessive, or X-linked trait. PAFAH1B1 (LIS1) is among the most prominent genes associated with lissencephaly. Loss of PAFAH1B1 alone causes epilepsy and developmental delay, but with absence of severe dysmorphism. Deletion of this gene along with adjacent ones in the terminal end of the short arm of chromosome 17 causes Miller-Dieker syndrome, characterized by epilepsy, dysmorphic features, severe developmental delay and, occasionally, heart defects and omphalocele. Somatic mutations in PAFAH1B1 and DCX have been described.<sup>4,5</sup> Cobblestone lissencephaly and lissencephaly with cerebellar hypoplasia segregate as autosomal recessive disorders and lissencephaly with agenesis of the corpus callosum is an X-linked disorder. Lissencephaly can sometimes occur together with heterotopias or polymicrogyria, depending on the affected gene and the type of pathogenic variant.

## Test Methods:

Using genomic DNA from the submitted specimen, the complete coding regions and splice site junctions of the genes on this panel are enriched using a proprietary targeted capture system developed by GeneDx for next-generation sequencing with CNV calling (NGS-CNV). The enriched targets are simultaneously sequenced with paired-end reads on an Illumina platform. Bi-directional sequence reads are assembled and aligned to reference sequences based on NCBI RefSeq transcripts and human genome build GRCh37/UCSC hg19. After gene specific filtering, data are analyzed to identify sequence variants and most deletions and duplications involving coding exons. Alternative sequencing or copy number detection methods are used to analyze regions with inadequate sequence or copy number data. Reportable variants include pathogenic variants, likely pathogenic variants and variants of uncertain significance. Likely benign and benign variants, if present, are not routinely reported but are available upon request.

## Test Sensitivity:

The clinical sensitivity of sequencing and deletion/duplication analysis of the genes included in this panel depends in part on the patient's clinical phenotype. Specific information about the diagnostic yield for each gene in selected populations is summarized in the following table(s). The technical sensitivity of sequencing is estimated to be > 99% at detecting single nucleotide events. It will not reliably detect deletions greater than 20 base pairs, insertions or rearrangements greater than 10 base pairs, or low-level mosaicism. The copy number assessment methods used with this test cannot reliably detect copy number variants of less than 500 base pairs or mosaicism and cannot identify balanced chromosome aberrations. Assessment of exon-level copy number events is dependent on the inherent sequence properties of the targeted regions, including shared homology and exon size. Gene specific exclusions for exon-level deletion/duplication testing for this panel are: B4GAT1 and FKRP genes, no copy number testing, ACTB, ARX, POMGNT1 and TUBA1A genes, only whole gene deletions or duplications may be detected.

Gene	Protein	Inheritance	Disease Associations
<i>ACTB</i> *	Actin, Beta	AD	80% of Baraitser-Winter syndrome <sup>6</sup>
<i>ACTG1</i>	Actin, Gamma-1	AD	20% of Baraitser-Winter syndrome <sup>6</sup>
<i>ARX</i> *	Aristaless-related homeobox protein	XL	70-95% of XLAG7,8, 3-10% in XLID <sup>9,10</sup>
<i>ATP6V0A2</i>	Lysosomal H(+)-ATPase	AR	21-24% of autosomal recessive cutis laxa type II <sup>11,12</sup>

	V0 subunit A2		
<i>B3GALNT2</i>	Beta-1,3-N-Acetylgalactosaminyltransferase 2	AR	Rare in alpha-dystroglycanopathies <sup>13</sup> Rare in congenital muscular dystrophy <sup>14</sup>
<i>B4GAT1</i> ( <i>B3GNT1</i> )**	Beta-1,4-Glucuronyltransferase 1	AR	Rare in alpha-dystroglycanopathies <sup>15,16</sup>
<i>CIT</i>	Citron rho-interacting serine/threonine kinase	AR	Rare in MCPH <sup>17,18,19</sup>
<i>DCX</i>	Doublecortin	XL	Up to 100% XL lissencephaly 10% of classic lissencephaly 85% females and ~30% males with SBH <sup>1,8,20</sup>
<i>FKRP</i> **	Fukutin-related protein	AR	~2% of cobblestone lissencephaly <sup>3,21</sup> 9% of alpha-dystroglycanopathies <sup>22</sup> 6% of limb-girdle MD <sup>23</sup> 2.5% of congenital muscular dystrophy <sup>14</sup>
<i>FKTN</i>	Fukutin	AR	~7% of alpha-dystroglycanopathies <sup>24</sup> Rare in congenital muscular dystrophy <sup>14</sup> 1.5% of limb-girdle MD <sup>25</sup> Does not include the Japanese founder mutation in the 3' UTR
<i>GMPPB</i>	GDP-Mannose Pyrophosphorylase B	AR	Rare in alpha-dystroglycanopathies <sup>27,28</sup> Rare in congenital muscular dystrophy <sup>14</sup>
<i>ISPD</i>	Isoprenoid synthase domain-containing protein	AR	~6% of cobblestone lissencephaly <sup>24</sup> ~30% of Walker-Warburg syndrome and ~11% of alpha-dystroglycanopathies <sup>29,30</sup> Rare in LGMD <sup>31</sup>
<i>KATNB1</i>	Katanin regulatory subunit B1	AR	<1% of MCD <sup>32,33</sup>
<i>LAMB1</i>	Laminin, Beta-1	AR	Rare in cobblestone lissencephaly <sup>34</sup>
<i>LARGE1</i>	Like-glycosyltransferase	AR	2-5% of cobblestone lissencephaly <sup>3,21</sup> ~1% of alpha-dystroglycanopathies <sup>22,24</sup>
<i>NDE1</i>	nudE neurodevelopment protein 1	AR	Rare <sup>35,36,37</sup>

<i>PAFAH1B1</i> ( <i>LIS1</i> )	Platelet-Activating Factor Acetylhydrolase 1b, Regulatory Subunit 1	AD	~40-65% of classic lissencephaly <sup>1,38</sup>
<i>POMGNT1</i> *	Protein O-Mannose Beta 1-2-N-Acetylglucosaminyltransferase	AR	11-18% of cobblestone lissencephaly <sup>3,21</sup> 8-10% of alpha-dystroglycanopathies <sup>22,24</sup>
<i>POMGNT2</i>	Protein O-Mannose Beta-1,4-N-Acetylglucosaminyltransferase 2	AR	Rare in alpha-dystroglycanopathies <sup>39</sup>
<i>POMT1</i>	Protein O-Mannosyltransferase 1	AR	27-34% of cobblestone lissencephaly <sup>3,21</sup> 9-21% of alpha-dystroglycanopathies <sup>22,24</sup>
<i>POMT2</i>	Protein O-Mannosyltransferase 2	AR	8-11% of cobblestone lissencephaly <sup>3,21</sup> 9-11% of alpha-dystroglycanopathies <sup>22,24</sup>
<i>RELN</i>	Reelin	AR	Rare <sup>40,41</sup>
<i>TMEM5</i>	Transmembrane protein 5	AR	~6% of cobblestone lissencephaly <sup>21</sup> Rare in alpha-dystroglycanopathies <sup>42</sup>
<i>TUBA1A</i> *	Tubulin, Alpha-1A	AD	1% of classic lissencephaly 30% of lissencephaly with cerebellar hypoplasia <sup>43,44</sup>
<i>VLDLR</i>	Very low density lipoprotein receptor	AR	Rare cerebellar hypoplasia with simplified gyri <sup>45,46</sup>
<i>WDR62</i>	WD repeat-containing protein 62	AR	Unknown <sup>47</sup>

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