



Reference transcripts based on build GRCh37 (hg19) interrogated by Lysosome, Peroxisome, GSD Panels.

Lysosomal Storage Disease Panel	
Gene	GenBank Accession Number
<i>ACP2</i>	NM_001610.2
<i>AGA</i>	NM_000027.3
<i>ARSA</i>	NM_000487.5
<i>ARSB</i>	NM_000046.3
<i>ARSH</i>	NM_001011719.1
<i>ASAH1</i>	NM_004315.5
<i>ATP13A2</i>	NM_022089.3
<i>CHIT1</i>	NM_003465.2
<i>CLN3</i>	NM_000086.2
<i>CLN5</i>	NM_006493.2
<i>CLN6</i>	NM_017882.2
<i>CLN8</i>	NM_018941.3
<i>CTNS</i>	NM_001031681.2
<i>CTSA</i>	NM_000308.3
<i>CTSD</i>	NM_001909.4
<i>CTSF</i>	NM_003793.3
<i>CTSK</i>	NM_000396.3
<i>DNAJC5</i>	NM_025219.2
<i>FUCA1</i>	NM_000147.4
<i>GAA</i>	NM_000152.4
<i>GALC</i>	NM_000153.3
<i>GALNS</i>	NM_000512.4
<i>GBA</i>	NM_000157.3
<i>GFAP</i>	NM_002055.4
<i>GLA</i>	NM_000169.2
<i>GLB1</i>	NM_000404.3
<i>GM2A</i>	NM_000405.4
<i>GNPTAB</i>	NM_024312.4
<i>GNPTG</i>	NM_032520.4
<i>GNS</i>	NM_002076.3
<i>GRN</i>	NM_002087.3
<i>GUSB</i>	NM_000181.3
<i>LAMP2</i>	NM_002294.2
<i>HEXA</i>	NM_000520.5
<i>HEXB</i>	NM_000521.3
<i>HGSNAT</i>	NM_152419.2
<i>HYAL1</i>	NM_153281.1
<i>IDS</i>	NM_000202.7

Lysosomal Storage Disease Panel	
Gene	GenBank Accession Number
<i>IDUA</i>	NM_000203.4
<i>KCTD7</i>	NM_153033.4
<i>LIPA</i>	NM_000235.3
<i>MAN2B1</i>	NM_000528.3
<i>MANBA</i>	NM_005908.3
<i>MCOLN1</i>	NM_020533.2
<i>MFSD8</i>	NM_152778.2
<i>NAGA</i>	NM_000262.2
<i>NAGLU</i>	NM_000263.3
<i>NEU1</i>	NM_000434.3
<i>NPC1</i>	NM_000271.4
<i>NPC2</i>	NM_006432.3
<i>PANK2</i>	NM_153638.2
<i>PSAP</i>	NM_002778.3
<i>PPT1</i>	NM_000310.3
<i>SGSH</i>	NM_000199.4
<i>SLC17A5</i>	NM_012434.4
<i>SMPD1</i>	NM_000543.4
<i>SUMF1</i>	NM_182760.3
<i>TPP1</i>	NM_000391.3

Neuronal Ceroid Lipofuscinosis (Batten Disease) Panel	
Gene	GenBank Accession Number
<i>ATP13A2</i>	NM_022089.3
<i>CLN3</i>	NM_000086.2
<i>CLN5</i>	NM_006493.2
<i>CLN6</i>	NM_017882.2
<i>CLN8</i>	NM_018941.3
<i>CTSD</i>	NM_001909.4
<i>CTSF</i>	NM_003793.3
<i>CTSK</i>	NM_000396.3
<i>DNAJC5</i>	NM_025219.2
<i>GRN</i>	NM_002087.3
<i>KCTD7</i>	NM_153033.4
<i>MFSD8</i>	NM_152778.2
<i>PANK2</i>	NM_153638.2
<i>PPT1</i>	NM_000310.3
<i>TPP1</i>	NM_000391.3

Targeted Genes for Lysosome, Peroxisome, GSD Panels (continued)

Peroxisomal Disorder Panel	
Gene	GenBank Accession Number
<i>ABCD1</i>	NM_000033.3
<i>ABCD2</i>	NM_005164.3
<i>ABCD3</i>	NM_002858.3
<i>ACOX1</i>	NM_004035.6
<i>ACOX3</i>	NM_003501.2
<i>AGPS</i>	NM_003659.3
<i>AMACR</i>	NM_014324.5
<i>CAT</i>	NM_001752.3
<i>DNM1L</i>	NM_012062.4
<i>ECH1</i>	NM_001398.2
<i>GNPAT</i>	NM_014236.3
<i>HSD17B4</i>	NM_001199291.1
<i>PEX1</i>	NM_000466.2
<i>PEX2</i>	NM_000318.2
<i>PEX3</i>	NM_003630.2
<i>PEX5</i>	NM_001131023.1
<i>PEX6</i>	NM_000287.3
<i>PEX7</i>	NM_000288.3
<i>PEX10</i>	NM_153818.1
<i>PEX11B</i>	NM_003846.2
<i>PEX12</i>	NM_000286.2
<i>PEX13</i>	NM_002618.3
<i>PEX14</i>	NM_004565.2
<i>PEX16</i>	NM_057174.2
<i>PEX19</i>	NM_002857.3
<i>PEX26</i>	NM_017929.5
<i>PHYH</i>	NM_006214.3
<i>SCP2</i>	NM_002979.4
<i>SUGCT</i>	NM_001193311.1
<i>TRIM37</i>	NM_015294.4

Glycogen Storage Disease Panel	
Gene	GenBank Accession Number
<i>AGL</i>	NM_000642.2
<i>ALDOA</i>	NM_184043.2
<i>ENO3</i>	NM_001976.4
<i>EPM2A</i>	NM_005670.3
<i>G6PC</i>	NM_000151.3
<i>GAA</i>	NM_000152.4
<i>GBE1</i>	NM_000158.3
<i>GYG1</i>	NM_004130.3
<i>GYS1</i>	NM_002103.4
<i>GYS2</i>	NM_021957.3
<i>LAMP2</i>	NM_002294.2
<i>LDHA</i>	NM_005566.3
<i>NHLRC1</i>	NM_198586.2
<i>PFKM</i>	NM_000289.5
<i>PGAM2</i>	NM_000290.3
<i>PGK1</i>	NM_000291.3
<i>PGM1</i>	NM_002633.2
<i>PHKA1</i>	NM_002637.3
<i>PHKA2</i>	NM_000292.2
<i>PHKB</i>	NM_000293.2
<i>PHKG2</i>	NM_000294.2
<i>PRKAG2</i>	NM_016203.3
<i>PYGL</i>	NM_002863.4
<i>PYGM</i>	NM_005609.3
<i>SLC2A2</i>	NM_000340.1
<i>SLC37A4</i>	ENST00000545985.1