

Input Variant	Chromosomal Variant	ClinVar Status
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NM_000118.3:c.14C>T	NC_000009.11:g.130616621G>A	
NM_000104.3:c.142C>G	NC_000002.11:g.38302390G>C	
NM_022124.5:c.1053C>T	NC_000010.10:g.73377069C>T	
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NM_007194.3:c.319+43dup	NC_000022.10:g.29130348dup	
NM_000138.4:c.7819+8A>C	NC_000015.9:g.48712876T>G	
NM_000070.2:c.606T>C	NC_000015.9:g.42680058T>C	
NM_000152.3:c.2780C>T	NC_000017.10:g.78092585C>T	
NM_001242896.1:c.1095T>C	NC_000022.10:g.32200161T>C	
NM_000251.2:c.1077-10T>C	NC_000002.11:g.47656871T>C	
NM_003494.3:c.3065G>A	NC_000002.11:g.71797762G>A	
NM_000492.3:c.1584G>A	NC_000007.13:g.117199709G>A	
NM_001242896.1:c.3358A>G	NC_000022.10:g.32266603A>G	
NM_177438.2:c.4206+11del	NC_000014.8:g.95566106del	
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NM_025243.3:c.435C>T	NC_000002.11:g.228563996G>A	
NM_025243.3:c.549T>C	NC_000002.11:g.228563882A>G	
NM_004004.5:c.-22-12C>T	NC_000013.10:g.20763754G>A	
NM_000138.4:c.3442C>G	NC_000015.9:g.48779530G>C	
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NM_206933.2:c.11736G>A	NC_000001.10:g.215901702C>T	
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NM_000051.3:c.378T>A	NC_000011.9:g.108106443T>A	
NM_206933.2:c.11907A>T	NC_000001.10:g.215901531T>A	
NM_000059.3:c.10234A>G	NC_000013.10:g.32972884A>G	
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NM_000525.3:c.868G>A	NC_000011.9:g.17408771C>T	
NM_000527.4:c.1166C>T	NC_000019.9:g.11222295C>T	
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NM_000546.5:c.423C>G	NC_000017.10:g.7578507G>C	
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NM_000546.5:c.464C>A	NC_000017.10:g.7578466G>T	
NM_000546.5:c.658T>C	NC_000017.10:g.7578191A>G	

NM_000546.5:c.770T>A	NC_000017.10:g.7577511A>T	
NM_000551.3:c.488T>G	NC_000003.11:g.10191495T>G	
NM_001018005.1:c.62G>T	NC_000015.9:g.63335090G>T	
NM_001018005.1:c.715G>A	NC_000015.9:g.63354787G>A	
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NM_007294.3:c.4096+3A>G	NC_000017.10:g.41243449T>C	
NM_007294.3:c.5098A>G	NC_000017.10:g.41215945T>C	
NM_174936.3:c.1069C>T	NC_000001.10:g.55523076C>T	
NM_206933.2:c.12145G>A	NC_000001.10:g.215853640C>T	
NM_206933.2:c.4823A>G	NC_000001.10:g.216262417T>C	
NM_206933.2:c.9433C>T	NC_000001.10:g.215990476G>A	
NM_206933.2:c.9885T>G	NC_000001.10:g.215972322A>C	
NM_000203.3:c.1225G>C	NC_000004.11:g.996555G>C	
NM_000203.3:c.235G>A	NC_000004.11:g.981673G>A	
NM_000203.3:c.1582C>G	NC_000004.11:g.997190C>G	
NM_000203.3:c.296C>T	NC_000004.11:g.981734C>T	
NM_000203.3:c.709C>T	NC_000004.11:g.995586C>T	
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NM_000203.3:c.1293C>T	NC_000004.11:g.996623C>T	
NM_000203.3:c.898G>A	NC_000004.11:g.995875G>A	
NM_000352.3:c.3120-19C>T	NC_000011.9:g.17428496G>A	
NM_000352.3:c.3612C>T	NC_000011.9:g.17424246G>A	
NM_000352.3:c.323G>A	NC_000011.9:g.17491737C>T	
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NM_000352.3:c.1572G>A	NC_000011.9:g.17464325C>T	
NM_000352.3:c.3728C>T	NC_000011.9:g.17419911G>A	
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NM_000352.3:c.3542C>A	NC_000011.9:g.17426074G>T	
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NM_000352.3:c.3000C>T	NC_000011.9:g.17428597G>A	
NM_000352.3:c.102G>A	NC_000011.9:g.17498222C>T	
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NM_000465.2:c.609A>C	NC_000002.11:g.215645989T>G	
NM_000465.2:c.2212A>G	NC_000002.11:g.215593522T>C	
NM_000465.2:c.1933T>C	NC_000002.11:g.215595203A>G	
NM_000465.2:c.1738G>A	NC_000002.11:g.215610518C>T	
NM_000465.2:c.620A>G	NC_000002.11:g.215645978T>C	
NM_000465.2:c.2191C>G	NC_000002.11:g.215593543G>C	

NM_000558.3:c.-66-?_*111+?del	NC_000016.9:g.226650_227521del	
NM_000558.3:c.-66-?_*111+?dup	NC_000016.9:g.226650_227521dup	
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NM_004364.3:c.756G>T	NC_000019.9:g.33792565C>A	
NM_004364.3:c.693C>G	NC_000019.9:g.33792628G>C	
NM_004364.3:c.1041G>A	NC_000019.9:g.33792280C>T	
NM_004364.3:c.612G>C	NC_000019.9:g.33792709C>G	
NM_004364.3:c.667G>A	NC_000019.9:g.33792654C>T	
NM_004364.3:c.308_313dup	NC_000019.9:g.33793008_33793013dup	
NM_004364.3:c.388G>A	NC_000019.9:g.33792933C>T	
NM_006494.2:c.1230T>C	NC_000019.9:g.42753034A>G	
NM_006494.2:c.615C>G	NC_000019.9:g.42753649G>C	
NM_006494.2:c.981C>T	NC_000019.9:g.42753283G>A	
NM_006494.2:c.1104T>C	NC_000019.9:g.42753160A>G	
NM_006494.2:c.1192G>A	NC_000019.9:g.42753072C>T	

