

Hugo_Symbol	Entrez_Gen	Chromosome	Start_position	End_position	Variant_Classification	Reference_Seq	Tumor_Seq	Tumor_Seq	Genome_Build	Annotation	Transcript_ID	Transcript_ID	Transcript_ID	cDNA_Change	Codon_Change	Protein_Change	Other_Transcripts	Refseq_mRNA_Id
CTD-2192116.20	0	19	12691164	12691164	3'UTR	C	C	T	g.chr19:12691164	ENST000000000000	-	0	625				ZNF490_ENST0000000311437.6_3'UTR	
AKR1A1	10327	1	46032770	46032770	3'UTR	C	C	T	g.chr1:46032770	ENST000000000000	+	0	528				AKR1A1_ENST0000000351829.4_Intron1 AKR1A1_EN	
AKAP3	10566	12	4724854	4724854	3'UTR	G	G	A	g.chr12:4724854	ENST000000000000	-	0	3137				AKAP3_ENST0000002288.NM_001278309.1	
FCER1A	2205	1	159277781	159277781	3'UTR	C	C	A	g.chr1:159277781	ENST000000000000	+	0	932				FCER1A_ENST000000368.NM_002001.3	
ABL1	25	9	133761129	133761129	3'UTR	G	G	A	g.chr9:133761129	ENST000000000000	+	0	3833				NM_005157.4	
OR2L13	284521	1	248262672	248262672	5'UTR	G	G	C	g.chr1:248262672	ENST000000000000	+	0	332				OR2L13_ENST000000358.NM_175911.2	
Unknown	0	5	62072618	62072618	IgR	G	G	A									IPO11 (148209 upstream): None (None downstream)	
KCNIP3	30818	2	95976299	95976299	Intron	T	T	G	g.chr2:95976299	ENST000000000000	+	2	316				KCNIP3_ENST000000360.NM_013434.4	
CA8	767	8	61121269	61121269	Intron	T	T	C	g.chr8:61121269	ENST000000000000	-	8	1173				NM_004056.4	
SUMF2	25870	7	56140772	56140772	Missense	J_G	G	A	g.chr7:56140772	ENST000000000000	+	3	395	c.364G>A	c.(364-366) p.E122K		SUMF2_ENST000000437.NM_001042469.1 NM_C	
NR3C2	0	4	149357871	149357871	Missense	J_C	C	T	g.chr4:149357871	ENST000000000000	-	2	504	c.142G>A	c.(142-144) p.V48I		NR3C2_ENST000000511528.1_Missense_Mutation	
FRG1B	0	20	29628245	29628245	Missense	J_G	G	A	g.chr20:29628245	ENST000000000000	+	6	627	c.247G>A	c.(247-249) p.A83T		FRG1B_ENST000000358464.4_Missense_Mutation	
RTP4	64108	3	187086270	187086270	Missense	J_A	A	G	g.chr3:187086270	ENST000000000000	+	1	151	c.41A>G	c.(40-42) p.E14G		NM_022147.2	
CSMD1	64478	8	4495039	4495039	Missense	J_T	T	C	g.chr8:4495039	ENST000000000000	-	2	682	c.127A>G	c.(127-129) p.I43V		CSMD1_ENST000000400186.3_Missense_Mutation	
CTNNA2	1496	2	80136918	80136918	Missense	J_A	A	C	g.chr2:80136918	ENST000000000000	+	11	1775	c.1051A>C	c.(1051-105) p.N351H		CTNNA2_ENST000000496558.1_Missense_Mutatio	
CUL9	23113	6	43164595	43164595	Missense	J_G	G	A	g.chr6:43164595	ENST000000000000	+	11	2882	c.2798G>A	c.(2797-27) p.R933H		CUL9_ENST00000037264.NM_015089.2	
DYRK1B	9149	19	40318202	40318202	Missense	J_C	C	T	g.chr19:40318202	ENST000000000000	-	7	1370	c.902G>A	c.(901-903) p.C301Y		DYRK1B_ENST000000348817.3_Missense_Mutatio	
OTOF	9381	2	26697530	26697530	Missense	J_A	A	C	g.chr2:26697530	ENST000000000000	-	26	3265	c.3139T>G	c.(3139-314) p.T1047V		OTOF_ENST00000033858.NM_194248.2	
BNC1	646	15	83933158	83933158	Missense	J_C	C	T	g.chr15:83933158	ENST000000000000	-	4	930	c.845G>A	c.(844-846) p.G282E		BNC1_ENST00000056970.NM_001717.3	
SEZ6L2	26470	16	29883824	29883824	Missense	J_G	G	A	g.chr16:29883824	ENST000000000000	-	15	3039	c.2512C>T	c.(2512-251) p.R838W		SEZ6L2_ENST0000005374.NM_001114099.2 NM_2	
CTSS	1520	1	150724429	150724429	Missense	J_A	A	G	g.chr1:150724429	ENST000000000000	-	5	715	c.455T>C	c.(454-456) p.L152P		CTSS_ENST000000480766.NM_001199739.1 NM_C	
CCDC181	57821	1	169391215	169391215	Missense	J_G	G	C	g.chr1:169391215	ENST000000000000	-	4	961	c.454C>G	c.(454-456) p.L152V		CCDC181_ENST000000367806.3_Missense_Mutati	
NEB	4703	2	152500345	152500345	Missense	J_T	T	G	g.chr2:152500345	ENST000000000000	-	57	8145	c.7943A>C	c.(7942-794) p.D2648A		NEB_ENST000000604864.NM_001164507.1	
PHLPP2	23035	16	71712830	71712830	Missense	J_T	T	C	g.chr16:71712830	ENST000000000000	-	7	1829	c.1096A>G	c.(1096-105) p.N366D		PHLPP2_ENST000000567016.1_Missense_Mutatio	
GABRG2	0	5	161580192	161580192	Missense	J_T	T	G	g.chr5:161580192	ENST000000000000	+	10	1706	c.1246T>G	c.(1246-124) p.C416G		GABRG2_ENST00000036.NM_000816.3 NM_1985	
ATXN7L2	127002	1	110033878	110033878	Missense	J_G	G	A	g.chr1:110033878	ENST000000000000	+	10	1708	c.1693G>A	c.(1693-165) p.V565M		ATXN7L2_ENST000000405.NM_153340.4	
OR4P4	81300	11	55406231	55406231	Missense	J_T	T	C	g.chr11:55406231	ENST000000000000	+	1	398	c.398T>C	c.(397-399) p.I133T		NM_001004124.1	
FRG1B	0	20	29628243	29628243	Missense	J_T	T	C	g.chr20:29628243	ENST000000000000	+	6	625	c.245T>C	c.(244-246) p.L82S		FRG1B_ENST000000358464.4_Missense_Mutation	
CARM1	10498	19	11022906	11022906	Missense	J_C	C	T	g.chr19:11022906	ENST000000000000	+	5	795	c.605C>T	c.(604-606) p.A202V		CARM1_ENST000000344.NM_199141.1	
MSLNL	401827	16	824911	824911	Missense	J_C	C	T	g.chr16:824911	ENST000000000000	-	7	1660	c.1661G>A	c.(1660-166) p.R554Q		MSLNL_ENST000000442466.1_Missense_Mutation	
DACH1	1820	13	72147042	72147042	Missense	J_G	G	A	g.chr13:72147042	ENST000000000000	-	4	1657	c.1235C>T	c.(1234-12) p.T412I		DACH1_ENST0000003131.NM_080759.4	
CDK5	1020	7	150754212	150754212	Missense	J_C	C	G	g.chr7:150754212	ENST000000000000	-	2	754	c.73G>C	c.(73-75) p.Gap.E25Q		CDK5_ENST00000029751.NM_004935.3	
MAZ	4150	16	29819095	29819095	Missense	J_A	A	C	g.chr16:29819095	ENST000000000000	+	2	1095	c.989A>C	c.(988-990) p.D330A		MAZ_ENST00000056340.NM_001042539.1	
NOP14	8602	4	2954016	2954016	Missense	J_G	G	A	g.chr4:2954016	ENST000000000000	-	6	921	c.856C>T	c.(856-858) p.L286F		NOP14_ENST000000502735.1_Missense_Mutatio	
PCDHGA2	0	5	140736478	140736478	Missense	J_G	G	A	g.chr5:140736478	ENST000000000000	+	1	1711	c.1711G>A	c.(1711-17) p.G571S		PCDHGA2_ENST00000003.NM_018917.2	
SLC25A11	8402	17	4843151	4843151	Missense	J_A	A	C	g.chr17:4843151	ENST000000000000	-	1	395	c.55T>G	c.(55-57) p.Tc.p.S19A		SLC25A11_ENST0000005003.NM_001165417.1 NM_C	
SPHKAP	80309	2	228882624	228882624	Missense	J_C	C	A	g.chr2:228882624	ENST000000000000	-	7	2992	c.2946G>T	c.(2944-294) p.K982N		SPHKAP_ENST000000344.NM_001142644.1	
ATRNL1	26033	10	117221456	117221456	Missense	J_C	C	A	g.chr10:117221456	ENST000000000000	+	22	3454	c.3328C>A	c.(3328-332) p.L110I		ATRNL1_ENST000000423.NM_207303.2	
DTPY19L4	286148	8	95777417	95777417	Missense	J_G	G	C	g.chr8:95777417	ENST000000000000	+	9	976	c.877G>C	c.(877-879) p.E293Q		NM_181787.2	
DTNB	1838	2	25861940	25861940	Nonsense	J_G	G	A	g.chr2:25861940	ENST000000000000	-	3	340	c.91C>T	c.(91-93) p.Cg.p.R31*		DTNB_ENST00000040718.NM_001256303.1 NM_C	
ARHGAP26	23092	5	142283200	142283200	Nonsense	J_C	C	A	g.chr5:142283200	ENST000000000000	-	8	1153	c.798C>A	c.(796-798) p.Y266*		ARHGAP26_ENST000000.NM_001135608.1	
RP11-707M1.1	0	11	49831701	49831701	RNA	G	G	T	g.chr11:49831701	ENST000000000000	+	0	1926					
HPCAL1	3241	2	10560207	10560207	Silent	C	C	T	g.chr2:10560207	ENST000000000000	+	4	850	c.324C>T	c.(322-324) p.Y108Y		HPCAL1_ENST000000307.NM_134421.2	
MAGEA12	4111	X	151900285	151900285	Silent	G	G	A	g.chrX:151900285	ENST000000000000	-	3	869	c.516C>T	c.(514-516) p.I172I		MAGEA12_ENST00000003.NM_001166386.1	
ARHGAP15	55843	2	144525663	144525663	Silent	C	C	A	g.chr2:144525663	ENST000000000000	+	14	1517	c.1350C>A	c.(1348-135) p.I450I		CTD-2252P21.1_ENST000000.NM_018460.3	
LY9	4063	1	160784539	160784539	Silent	C	C	T	g.chr1:160784539	ENST000000000000	+	4	1090	c.1060C>T	c.(1060-106) p.L354L		LY9_ENST000000368035.NM_001261456.1 NM_C	
DPP9	91039	19	4689663	4689663	Silent	G	G	A	g.chr19:4689663	ENST000000000000	-	15	1945	c.1668C>T	c.(1666-166) p.L556L		DPP9_ENST00000059467.NM_139159.4	
RAP1GAP	5909	1	21930202	21930202	Silent	C	C	T	g.chr1:21930202	ENST000000000000	-	16	1640	c.1632G>A	c.(1630-162) p.A544A		RAP1GAP_ENST000000374761.2_Intron1 RAP1GAP	
ENOX2	10495	X	129837113	129837113	Silent	G	G	A	g.chrX:129837113	ENST000000000000	-	5	582	c.165C>T	c.(163-165) p.A55A		ENOX2_ENST0000004922.NM_182314.1	
IGSF3	3321	1	117122123	117122123	Silent	T	T	A	g.chr1:117122123	ENST000000000000	-	10	3990	c.3225A>T	c.(3223-322) p.T1075T		IGSF3_ENST00000036948.NM_001007237.1	
FAM160B2	64760	8	21959775	21959775	Silent	G	G	A	g.chr8:21959775	ENST000000000000	+	15	1987	c.1941G>A	c.(1939-194) p.P647P		NM_022749.5	
BAZZA	11176	12	56993811	56993811	Silent	C	C	T	g.chr12:56993811	ENST000000000000	-	26	5071	c.4872G>A	c.(4870-487) p.Q1624Q		BAZZA_ENST000000549884.1_Silent_p.Q1654Q B	
TRPV5	56302	7	142626649	142626649	Silent	G	G	A	g.chr7:142626649	ENST000000000000	-	4	709	c.361C>T	c.(361-363) p.L121L		TRPV5_ENST0000004426.NM_019841.4	
WSCD1	23302	17	5993659	5993659	Silent	C	C	T	g.chr17:5993659	ENST000000000000	+	4	951	c.561C>T	c.(559-561) p.G187G		WSCD1_ENST000000317744.5_Silent_p.G187G W	
ENG	2022	9	130587621	130587621	Silent	C	C	T	g.chr9:130587621	ENST000000000000	-	6	985	c.705G>A	c.(703-705) p.T235T		ENG_ENST000000373203.4_Silent_p.T235T	
HADHB	3032	2	26502095	26502095	Silent	G	G	A	g.chr2:26502095	ENST000000000000	+	9	827	c.723G>A	c.(721-723) p.Q241Q		HADHB_ENST00000	

Refseq_prot_id	SwissProt_acc_id	SwissProt_entry_id	Description	UniProt_AApos
∑ST00000372070.3_intron	P14550	AK1A1_HUMAN	aldo-keto reductase family 1, member A1 (aldehyde reductase)	
NP_001265238.1	O75969	AKAP3_HUMAN	A kinase (PRKA) anchor protein 3	
NP_001992.1	P12319	FCERA_HUMAN	Fc fragment of IgE, high affinity I, receptor for; alpha polypeptide	
NP_005148.2	P00519	ABL1_HUMAN	c-abl oncogene 1, non-receptor tyrosine kinase	
NP_787107.1	Q8N349	OR2LD_HUMAN	olfactory receptor, family 2, subfamily L, member 13	
NP_038462.1	Q9Y2W7	CSEN_HUMAN	Kv channel interacting protein 3, calсенил	
NP_004047.3	P35219	CAH8_HUMAN	carbonic anhydrase VIII	
NP_001035934.2 NP_056226Q8NBJ7		SUMF2_HUMAN	sulfatase modifying factor 2	103
∫_p.V481 NR3C2_ENST000005.P08235		MCR_HUMAN	nuclear receptor subfamily 3, group C, member 2	48
_p.A83T FRG1B_ENST00000439954.2_Missense_Mutation_p.A88T				
NP_071430.2	Q96DX8	RTP4_HUMAN	receptor (chemosensory) transporter protein 4	14
n_p.I43V CSMD1_ENST000001Q96P27		CSMD1_HUMAN	CUB and Sushi multiple domains 1	43
∩n_p.N351H CTNNA2_ENST00CP26232		CTNA2_HUMAN	catenin (cadherin-associated protein), alpha 2	351
NP_055904.1	Q8IWT3	CUL9_HUMAN	cullin 9	933
m_p.C301Y DYRK1B_ENST000Q9Y463		DYRK1B_HUMAN	dual-specificity tyrosine-(Y)-phosphorylation regulated kinase	301
NP_919224.1	Q9HC10	OTOF_HUMAN	otofelin	1047
NP_001708.3	Q01954	BNC1_HUMAN	basonuclein 1	282
NP_001107571.1 NP_963866Q6UXD5		SE6L2_HUMAN	seizure related 6 homolog (mouse)-like 2	838
NP_001186668.1 NP_00407CP25774		CATS_HUMAN	cathepsin 5	152
ion_p.L152V CCDC181_ENST00000491570.1_5'UTR CCDC181_ENST00000367805			coiled-coil domain containing 181	
NP_001157979.1	P20929	NEBU_HUMAN	nebulin	2648
n_p.N401D PHLPP2_ENST000Q6ZVD8		PHLP2_HUMAN	PH domain and leucine rich repeat protein phosphatase 2	366
NP_000807.2 NP_944494.1	P18507	GBRG2_HUMAN	gamma-aminobutyric acid (GABA) A receptor, gamma 2	408
NP_699171.3	Q576C5	AT7L2_HUMAN	ataxin 7-like 2	565
NP_001004124.1	Q8NGL7	OR4P4_HUMAN	olfactory receptor, family 4, subfamily P, member 4	133
_p.L82S FRG1B_ENST00000439954.2_Missense_Mutation_p.L87S				
NP_954592.1	Q86X55	CARM1_HUMAN	coactivator-associated arginine methyltransferase 1	202
∫_p.R203Q	Q96KJ4	MSLNL_HUMAN	mesothelin-like	203
NP_542937.2	Q9UI36	DACH1_HUMAN	dachshund homolog 1 (Drosophila)	462
NP_004926.1	Q00535	CDK5_HUMAN	cyclin-dependent kinase 5	25
NP_001036004.1	P56270	MAZ_HUMAN	MYC-associated zinc finger protein (purine-binding transcript)	330
∫_p.L286F NOP14-AS1_ENST0P78316		NOP14_HUMAN	NOP14 nucleolar protein	286
NP_061740.1				
NP_001158889.1 NP_003556Q02978		M2OM_HUMAN	solute carrier family 25 (mitochondrial carrier; oxoglutarate t	19
NP_001136116.1	Q2M3C7	SPKAP_HUMAN	SPHK1 interactor, AKAP domain containing	982
NP_997186.1	Q5VV63	ATRNL1_HUMAN	attractin-like 1	1110
NP_861452.2	Q7Z388	D19L4_HUMAN	dpy-19-like 4 (C. elegans)	293
NP_001243232.1 NP_068707Q60941		DTNB_HUMAN	dystrobrevin, beta	31
NP_001129080.1	Q9UNA1	RHG26_HUMAN	Rho GTPase activating protein 26	266
NP_602293.1	P37235	HPCL1_HUMAN	hippocalcin-like 1	108
NP_001159858.1	P43365	MAGAC_HUMAN	melanoma antigen family A, 12	172
NP_060930.3	Q53QZ3	RHG15_HUMAN	Rho GTPase activating protein 15	450
NP_001248385.1 NP_002336Q9HBG7		LY9_HUMAN	lymphocyte antigen 9	354
NP_631898.3	Q86TI2	DPP9_HUMAN	dipeptidyl-peptidase 9	527
'_ENST00000374765.4_intron P47736		RPGP1_HUMAN	RAP1 GTPase activating protein	460
NP_872114.1	Q16206	ENOX2_HUMAN	ecto-NOX disulfide-thiol exchanger 2	55
NP_001007238.1	O75054	IGSF3_HUMAN	immunoglobulin superfamily, member 3	1075
NP_073586.5	Q86V87	F16B2_HUMAN	family with sequence similarity 160, member B2	647
AZ2A_ENST00000379441.3_Si Q9UIF9		BAZ2A_HUMAN	bromodomain adjacent to zinc finger domain, 2A	1656
NP_062815.2	Q9NQA5	TRPV5_HUMAN	transient receptor potential cation channel, subfamily V, member 5	121
'SCD1_ENST00000573634.1_Si Q658N2		WSCD1_HUMAN	WSC domain containing 1	187
	P17813	EGLN_HUMAN	endoglin	235
NP_000174.1	P55084	ECHB_HUMAN	hydroxyacyl-CoA dehydrogenase/3-ketoacyl-CoA thiolase/en	241
NP_001193600.1 NP_078833Q7Z5B4		RIC3_HUMAN	RIC3 acetylcholine receptor chaperone	95
NP_003667.1	O15121	DEGS1_HUMAN	delta(4)-desaturase, sphingolipid 1	75
NP_005152.1	P35414	APJ_HUMAN	apelin receptor	72
NP_878919.2	Q658P3	STEAP3_HUMAN	STEAP family member 3, metalloredutase	71
NP_001186794.1 NP_078988Q96L93		KI16B_HUMAN	kinesin family member 16B	1142
∫8_ENST00000405249.1_Silen O00222		GRM8_HUMAN	glutamate receptor, metabotropic 8	103
NP_115991.1	Q969M2	CXA10_HUMAN	gap junction protein, alpha 10, 62kDa	456
NP_003707.2	Q9ULU8	CAPS1_HUMAN	Ca++-dependent secretion activator	707
NP_001121185.1	Q8WXF7	ATLA1_HUMAN	atlastin GTPase 1	307
NP_000259.1 NP_057502.2 P35240		MERL_HUMAN	neurofibromin 2 (merlin)	38