

Supplementary Table 4: High confidence somatic nonsynonymous mutations identified by target gene sequence

Sample	Location	Gene	Variant classification	Reference allele	Variant allele	AA change	Filtered coverage	Variant allele frequency	RefSeq	dbSNP ID	COSMIC ID
PR101	chrX:70360666	MED12	Stop_Gained	C	T	p.Q2076*	52	0.077	NM_005120.2		
PR101	chr5:112162851	APC	Non_Synonymous_Coding	G	A	p.M485I	5924	0.469	NM_001127510.2		
PR102	chr3:178922301	PIK3CA	Non_Synonymous_Coding	G	A	p.R357Q	3977	0.137	NM_006218.2		COSM276752
PR103	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	2846	0.223	NM_003482.3		
PR103	chr12:49432245	KMT2D	Non_Synonymous_Coding	T	C	p.N2965S	6929	0.489	NM_003482.3		
PR103	chr14:95916416	SYNE3	Non_Synonymous_Coding	C	T	p.R434H	2238	0.492	NM_152592.3	rs183604014	
PR104	chr7:91731945	AKAP9	Non_Synonymous_Coding	G	A	p.R3712Q	4052	0.456	NM_005751.4	rs186148498	
PR104	chr17:37673806	CDK12	Non_Synonymous_Coding	C	T	p.S987F	8795	0.479	NM_016507.2	rs201661022	
PR105	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	5408	0.224	NM_003482.3		
PR105	chr16:72828867	ZFXH3	Non_Synonymous_Coding	T	C	p.M2572V	4624	0.51	NM_006885.3	rs147672861	
PR105	chrX:70360579	MED12	Non_Synonymous_Coding	A	G	p.I2047V	548	0.969	NM_005120.2		
PR106	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	312	0.417	NM_005269.2	rs2229300	
PR106	chr16:72984669	ZFXH3	Non_Synonymous_Coding	G	A	p.S972L	942	0.499	NM_006885.3	rs77124117	
PR106	chr19:5131419	KDM4B	Non_Synonymous_Coding	G	A	p.A550T	1370	0.503	NM_015015.2		
PR107	chr12:49421002	KMT2D	Non_Synonymous_Coding	G	A	p.P4916L	110	0.073	NM_003482.3		
PR107	chr15:23019819	NIPA2	Non_Synonymous_Coding	C	G	p.W59C	34	0.118	NM_030922.6		
PR107	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	4141	0.205	NM_003482.3		
PR107	chr3:38888862	SCN11A	Non_Synonymous_Coding	A	G	p.C1567R	4641	0.467	NM_014139.2	rs201595463	
PR108	chr7:91715706	AKAP9	Stop_Gained	T	A	p.C3063*	420	0.043	NM_005751.4		COSM28887
PR108	chr10:89653841	PTEN	Non_Synonymous_Coding	A	G	p.R47G	91	0.077	NM_000314.4		
PR108	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	4216	0.228	NM_003482.3		
PR108	chr7:91731945	AKAP9	Non_Synonymous_Coding	G	A	p.R3712Q	2572	0.478	NM_005751.4	rs186148498	
PR108	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	365	0.499	NM_005269.2	rs2229300	
PR108	chr7:91646406	AKAP9	Non_Synonymous_Coding	G	A	p.R1276Q	2288	0.505	NM_005751.4	rs146797353	
PR108	chr8:71037059	NCOA2	Non_Synonymous_Coding	T	A	p.T1320S	1423	0.515	NM_006540.2		
PR108	chr16:72831357	ZFXH3	Coding_Deletion	CTTGTTG	C	p.QQ1740-	2233	0.563	NM_006885.3		COSM1731926
PR109	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	5095	0.242	NM_003482.3		
PR109	chr16:72829385	ZFXH3	Non_Synonymous_Coding	T	C	p.N2399S	5581	0.467	NM_006885.3		
PR110	chr12:49421003	KMT2D	Non_Synonymous_Coding	G	A	p.P4916S	148	0.061	NM_003482.3		
PR110	chr7:91682269	AKAP9	Non_Synonymous_Coding	T	A	p.S1856R	113	0.071	NM_005751.4		
PR110	chr16:72828408	ZFXH3	Non_Synonymous_Coding	C	A	p.A2725S	5104	0.073	NM_006885.3		
PR110	chr17:37619359	CDK12	Non_Synonymous_Coding	T	A	p.S345R	32	0.094	NM_016507.2		
PR110	chr17:47696426	SPOP	Non_Synonymous_Coding	A	C	p.F133V	3470	0.223	NM_001007230.1		COSM219965
PR110	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	3688	0.368	NM_003482.3		
PR110	chr7:91712512	AKAP9	Non_Synonymous_Coding	A	G	p.Q2730R	4012	0.44	NM_005751.4	rs80191629	
PR110	chr5:112174665	APC	Non_Synonymous_Coding	T	C	p.V1125A	2958	0.494	NM_001127510.2	rs377278397	
PR110	chr10:89711909	PTEN	Non_Synonymous_Coding	A	G	p.Y176C	1004	0.536	NM_000314.4		
PR110	chr2:138000044	THSD7B	Non_Synonymous_Coding	G	A	p.R692Q	3741	0.985	NM_001080427.1	rs76693568	COSM148945
PR111	chr17:7578427	TP53	Non_Synonymous_Coding	T	G	p.H168P	4091	0.058	NM_000546.5		COSM44808
PR111	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	3900	0.255	NM_003482.3		
PR111	chr14:95916404	SYNE3	Non_Synonymous_Coding	G	A	p.A438V	3236	0.5	NM_152592.3	rs376830751	COSM198717
PR111	chr12:49444996	KMT2D	Frame_shift	A	AG	p.-823?	93	0.677	NM_003482.3		

Contd...

Supplementary Table 4: Contd...

Sample	Location	Gene	Variant classification	Reference allele	Variant allele	AA change	Filtered coverage	Variant allele frequency	RefSeq	dbSNP ID	COSMIC ID
PR111	chrX:66937337	AR	Non_Synonymous_Coding	G	A	p.V731M	2059	0.997	NM_000044.3	rs137852571	COSM1468985
PR112	chr17:47696425	SPOP	Non_Synonymous_Coding	A	C	p.F133C	2480	0.189	NM_001007230.1		COSM242645
PR112	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	137	0.263	NM_005269.2	rs2229300	
PR112	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCCTCAGGTGGTGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	3359	0.286	NM_003482.3		
PR112	chr16:72832153	ZFXH3	Non_Synonymous_Coding	C	T	p.M1476I	1266	0.509	NM_006885.3	rs200133878	
PR113	chr17:37687207	CDK12	Non_Synonymous_Coding	G	C	p.V1371L	272	0.04	NM_016507.2		
PR113	chr12:57864588	GLI1	Non_Synonymous_Coding	C	T	p.P689S	178	0.067	NM_005269.2		
PR113	chr16:72828408	ZFXH3	Non_Synonymous_Coding	C	A	p.A2725S	5110	0.068	NM_006885.3		
PR113	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCCTCAGGTGGTGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	10 048	0.241	NM_003482.3		
PR113	chr12:49445021	KMT2D	Frame_shift	C		p.-815?	173	0.422	NM_003482.3		
PR114	chr2:137988662	THSD7B	Non_Synonymous_Coding	G	T	p.R560M	124	0.04	NM_001080427.1		
PR114	chr3:73433708	PDZRN3	Non_Synonymous_Coding	G	A	p.P670L	261	0.05	NM_015009.1		
PR114	chr7:91739463	AKAP9	Non_Synonymous_Coding	T	C	p.M3905T	4221	0.589	NM_005751.4	rs77447750	
PR115	chr14:95932477	SYNE3	Non_Synonymous_Coding	T	C	p.I140V	7018	0.496	NM_152592.3		
PR115	chr12:49424475	KMT2D	Non_Synonymous_Coding	C	T	p.C4583Y	3132	0.522	NM_003482.3		
PR201	chr19:5144388	KDM4B	Non_Synonymous_Coding	T	C	p.S956P	9471	0.031	NM_015015.2		
PR201	chr17:37619118	CDK12	Non_Synonymous_Coding	G	T	p.S265I	75	0.04	NM_016507.2		
PR201	chr3:100467047	TFG	Non_Synonymous_Coding	G	T	p.G292V	197	0.041	NM_006070.5		
PR201	chr2:138400097	THSD7B	Non_Synonymous_Coding	A	C	p.K1251Q	95	0.042	NM_001080427.1		
PR201	chr12:56428856	IKZF4	Non_Synonymous_Coding	A	G	p.K500R	88	0.045	NM_022465.3		
PR201	chr7:91714912	AKAP9	Non_Synonymous_Coding	C	T	p.P2979L	2084	0.047	NM_005751.4		
PR201	chr16:72821079	ZFXH3	Non_Synonymous_Coding	T	G	p.D3699A	1342	0.047	NM_006885.3		
PR201	chr5:112173687	APC	Non_Synonymous_Coding	A	C	p.V799S	105	0.048	NM_001127510.2		
PR201	chr2:138163363	THSD7B	Non_Synonymous_Coding	G	T	p.R863L	348	0.052	NM_001080427.1		
PR201	chr17:37687207	CDK12	Non_Synonymous_Coding	G	T	p.V1371L	111	0.054	NM_016507.2		
PR201	chr19:50549280	ZNF473	Non_Synonymous_Coding	G	A	p.G527D	3327	0.057	NM_001006656.1		
PR201	chr3:100467073	TFG	Non_Synonymous_Coding	G	C	p.A301P	198	0.061	NM_006070.5		
PR201	chr5:112111327	APC	Non_Synonymous_Coding	T	A	p.S142T	63	0.063	NM_001127510.2		
PR201	chr12:49421002	KMT2D	Non_Synonymous_Coding	G	A	p.P4916L	126	0.063	NM_003482.3		
PR201	chr12:49425616	KMT2D	Non_Synonymous_Coding	G	C	p.A4291G	157	0.064	NM_003482.3		
PR201	chr19:5071040	KDM4B	Non_Synonymous_Coding	C	T	p.H216Y	240	0.071	NM_015015.2		
PR201	chrX:66766249	AR	Non_Synonymous_Coding	G	A	p.G421R	84	0.071	NM_000044.3		
PR201	chr4:87217	ZNF595	Non_Synonymous_Coding	C	A	p.T607N	626	0.073	NM_182524.2		
PR201	chr12:56428449	IKZF4	Non_Synonymous_Coding	C	G	p.S364R	112	0.08	NM_022465.3		
PR201	chr3:73437120	PDZRN3	Non_Synonymous_Coding	T	C	p.Q506R	285	0.084	NM_015009.1		
PR201	chr16:72828408	ZFXH3	Non_Synonymous_Coding	C	A	p.A2725S	2338	0.147	NM_006885.3		
PR201	chr4:86822	ZNF595	Coding_Deletion	GGAGAAACCCTACAATGTGAAGATGT GGCAAAGCTTTCATATGGTCGCNAGC CTGAATGAACATAAGAATATTTACTGTGA	C	p.EKPYKCEECGKAF IWSASLINEKNIH TG476-	212	0.302	NM_182524.2		
PR201	chr17:37673806	CDK12	Non_Synonymous_Coding	C	T	p.S987F	10 151	0.506	NM_016507.2	rs201661022	
PR201	chr4:87313	ZNF595	Frame_shift	G	GA	p.-639?	184	0.783	NM_182524.2	rs60154095	
PR201	chr7:91714911	AKAP9	Non_Synonymous_Coding	C	T	p.P2979S	2076	0.947	NM_005751.4	rs1063242	
PR201	chrX:66765516	AR	Non_Synonymous_Coding	C	A	p.S176R	1550	0.995	NM_000044.3		

Contd...

Supplementary Table 4: Contd...

Sample	Location	Gene	Variant classification	Reference allele	Variant allele	AA change	Filtered coverage	Variant allele frequency	RefSeq	dbSNP ID	COSMIC ID
PR202	chr19:5150397	KDM4B	Non_Synonymous_Coding	C	T	p.T1017M	161	0.043	NM_015015.2		
PR202	chr12:49444986	KMT2D	Non_Synonymous_Coding	T	C	p.Q827R	27	0.074	NM_003482.3		
PR202	chr5:112175043	APC	Non_Synonymous_Coding	T	C	p.V1251A	107	0.075	NM_001127510.2		
PR202	chrX:70339696	MED12	Non_Synonymous_Coding	G	T	p.G122V	53	0.075	NM_005120.2		
PR202	chr16:72821887	ZFX3	Frame_shift	C	CG	p.-3429?	1432	0.214	NM_006885.3		
PR202	chr7:91646406	AKAP9	Non_Synonymous_Coding	G	A	p.R1276Q	1608	0.631	NM_005751.4	rs146797353	
PR202	chr4:87313	ZNF595	Frame_shift	G	GA	p.-639?	142	0.965	NM_182524.2	rs60154095	
PR203	chr16:72991626	ZFX3	Non_Synonymous_Coding	C	T	p.E807K	216	0.042	NM_006885.3		
PR203	chr15:23006222	NIPA2	Stop_Lost	T	A	p.*361L	836	0.045	NM_030922.6		
PR203	chr12:49445310	KMT2D	Non_Synonymous_Coding	G	A	p.P719L	59	0.085	NM_003482.3	rs185660524	
PR203	chr5:112173644	APC	Non_Synonymous_Coding	C	G	p.H785D	1781	0.127	NM_001127510.2		
PR203	chr17:47696425	SPOP	Non_Synonymous_Coding	A	G	p.F133S	1070	0.133	NM_001007230.1		COSM247573
PR203	chr9:21974694	CDKN2A	Non_Synonymous_Coding	C	G	p.G45R	2296	0.501	NM_001195132.1		
PR203	chr16:72831357	ZFX3	Coding_Deletion	C TTGTTG	C	p.QQ1740-	1886	0.512	NM_006885.3		COSM1731926
PR203	chr7:91690697	AKAP9	Non_Synonymous_Coding	G	A	p.A1909T	2874	0.558	NM_005751.4	rs200844952	
PR203	chr4:87313	ZNF595	Frame_shift	G	GA	p.-639?	110	0.936	NM_182524.2	rs60154095	
PR203	chr7:91714911	AKAP9	Non_Synonymous_Coding	C	T	p.P2979S	1553	0.958	NM_005751.4	rs1063242	
PR205	chr19:50549280	ZNF473	Non_Synonymous_Coding	G	A	p.G527D	14 424	0.041	NM_001006656.1		
PR205	chr12:49427950	KMT2D	Non_Synonymous_Coding	C	T	p.R3547H	6749	0.045	NM_003482.3		
PR205	chr17:37687204	CDK12	Non_Synonymous_Coding	C	A	p.L1370M	437	0.046	NM_016507.2		
PR205	chr2:138400100	THSD7B	Non_Synonymous_Coding	G	C	p.E1252Q	312	0.048	NM_001080427.1		
PR205	chr5:112176407	APC	Non_Synonymous_Coding	T	A	p.S1706T	548	0.049	NM_001127510.2		
PR205	chr12:49424422	KMT2D	Non_Synonymous_Coding	G	T	p.P4601T	201	0.05	NM_003482.3		
PR205	chr3:178942538	PIK3CA	Non_Synonymous_Coding	A	G	p.I782S	791	0.053	NM_006218.2		
PR205	chr12:56428447	IKZF4	Non_Synonymous_Coding	A	T	p.S364C	431	0.053	NM_022465.3		
PR205	chr12:56428445	IKZF4	Non_Synonymous_Coding	A	T	p.H363L	404	0.054	NM_022465.3		
PR205	chr19:5119672	KDM4B	Non_Synonymous_Coding	G	A	p.R375Q	403	0.062	NM_015015.2		
PR205	chr16:72991377	ZFX3	Non_Synonymous_Coding	C	T	p.G890R	496	0.067	NM_006885.3		
PR205	chr12:49444990	KMT2D	Non_Synonymous_Coding	G	T	p.P826T	292	0.068	NM_003482.3		
PR205	chr2:138400097	THSD7B	Non_Synonymous_Coding	A	C	p.K1251Q	318	0.097	NM_001080427.1		
PR205	chr16:72828408	ZFX3	Non_Synonymous_Coding	C	A	p.A2725S	8996	0.105	NM_006885.3		
PR205	chr16:72828405	ZFX3	Frame_shift	GA	G	p.-2725	8808	0.107	NM_006885.3		
PR205	chr12:49445334	KMT2D	Non_Synonymous_Coding	G	A	p.P711L	44	0.136	NM_003482.3		
PR205	chr17:7577147	TP53	Non_Synonymous_Coding	A	T	p.L264Q	98	0.184	NM_000546.5		
PR205	chr7:91682269	AKAP9	Non_Synonymous_Coding	T	A	p.S1866R	83	0.193	NM_005751.4		
PR205	chr17:37627784	CDK12	Non_Synonymous_Coding	C	G	p.P567A	92	0.196	NM_016507.2		
PR205	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGACACAGGGGTGAC	T	p.ESPLSPPEE446E	7007	0.463	NM_003482.3		
PR206	chr8:71078935	NCOA2	Non_Synonymous_Coding	T	C	p.N199S	4974	0.035	NM_006540.2		
PR206	chr3:178928073	PIK3CA	Non_Synonymous_Coding	G	A	p.G451R	4034	0.037	NM_006218.2		
PR206	chr14:95932477	SYNE3	Non_Synonymous_Coding	T	C	p.I140V	8892	0.041	NM_152592.3		
PR206	chr3:178916833	PIK3CA	Non_Synonymous_Coding	A	G	p.T74A	7255	0.106	NM_006218.2		
PR206	chr17:37649079	CDK12	Coding_Deletion	CATT	C	p.I729-	13 486	0.22	NM_016507.2		

Contd...

Supplementary Table 4: Contd...

Sample	Location	Gene	Variant classification	Reference allele	Variant allele	AA change	Filtered coverage	Variant allele frequency	RefSeq	dbSNP ID	COSMIC ID
PR206	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGGACAGGGGTGAC	T	p.ESPLSPPPEE446E	5541	0.243	NM_003482.3		
PR206	chr7:107823354	NRCAM	Non_Synonymous_Coding	C	T	p.G772D	5502	0.486	NM_001037132.2		
PR207	chr3:38888625	SCN11A	Non_Synonymous_Coding	G	T	p.L1646I	6475	0.127	NM_014139.2		
PR208	chr12:49445341	KMT2D	Non_Synonymous_Coding	G	A	p.P709S	100	0.04	NM_003482.3		
PR208	chr12:57865549	GLI1	Non_Synonymous_Coding	C	T	p.P1009L	81	0.074	NM_005269.2		
PR208	chr16:72828408	ZFXH3	Non_Synonymous_Coding	C	A	p.A2725S	8350	0.082	NM_006885.3		
PR208	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGGACAGGGGTGAC	T	p.ESPLSPPPEE446E	7658	0.303	NM_003482.3		
PR208	chr3:73433697	PZRN3	Non_Synonymous_Coding	C	G	p.G674R	390	0.454	NM_015009.1		
PR210	chr16:72828408	ZFXH3	Non_Synonymous_Coding	C	A	p.A2725S	9425	0.06	NM_006885.3		
PR210	chr19:5144410	KDM4B	Non_Synonymous_Coding	C	G	p.P963R	105	0.105	NM_015015.2		
PR210	chr16:72821887	ZFXH3	Frame_shift	C	CG	p.-3429?	3674	0.201	NM_006885.3		
PR210	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGGACAGGGGTGAC	T	p.ESPLSPPPEE446E	7636	0.323	NM_003482.3		
PR210	chr14:95942049	SYNE3	Non_Synonymous_Coding	C	T	p.R37H	7754	0.401	NM_152592.3		
PR210	chr14:95932477	SYNE3	Non_Synonymous_Coding	T	C	p.I140V	10 415	0.499	NM_152592.3		
PR210	chr7:91646406	AKAP9	Non_Synonymous_Coding	G	A	p.R1276Q	2452	0.529	NM_005751.4	rs146797353	
PR210	chr8:71078935	NCOA2	Non_Synonymous_Coding	T	C	p.N199S	4847	0.56	NM_006540.2		
PR210	chr16:72991697	ZFXH3	Codon_Change_Plus_Coding	GCGCGCCGCCGCGAGCCA	G	p.VAAAAA777A	390	0.985	NM_006885.3		
PR212	chr3:178917586	PIK3CA	Non_Synonymous_Coding	G	C	p.R154T	30	0.1	NM_006218.2		
PR212	chrX:70348499	MED12	Non_Synonymous_Coding	A	T	p.I1136F	799	0.149	NM_005120.2		
PR212	chr19:5144410	KDM4B	Non_Synonymous_Coding	C	A	p.P963H	83	0.205	NM_015015.2		
PR212	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGGACAGGGGTGAC	T	p.ESPLSPPPEE446E	6707	0.211	NM_003482.3		
PR212	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	168	0.357	NM_005269.2	rs2229300	
PR212	chr12:49420078	KMT2D	Non_Synonymous_Coding	C	T	p.R5224H	3045	0.5	NM_003482.3	rs3782356	
PR213	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGGACAGGGGTGAC	T	p.ESPLSPPPEE446E	5577	0.203	NM_003482.3		
PR213	chrX:70349256	MED12	Non_Synonymous_Coding	T	G	p.V1223G	3524	0.442	NM_005120.2		
PR213	chr12:49420078	KMT2D	Non_Synonymous_Coding	C	T	p.R5224H	4335	0.493	NM_003482.3	rs3782356	
PR214	chr3:178917586	PIK3CA	Non_Synonymous_Coding	G	T	p.R154M	86	0.058	NM_006218.2		
PR214	chr2:138000044	THSD7B	Non_Synonymous_Coding	G	A	p.R692Q	6899	0.492	NM_001080427.1	rs76693568	COSM148945
PR215	chr16:72828408	ZFXH3	Non_Synonymous_Coding	C	A	p.A2725S	12 278	0.059	NM_006885.3		
PR215	chr7:91699374	AKAP9	Non_Synonymous_Coding	A	C	p.K2121Q	55	0.091	NM_005751.4		
PR215	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	45	0.178	NM_005269.2	rs2229300	
PR215	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGGACAGGGGTGAC	T	p.ESPLSPPPEE446E	8010	0.342	NM_003482.3		
PR215	chr3:38962696	SCN11A	Non_Synonymous_Coding	T	G	p.N255H	1634	0.523	NM_014139.2	rs371647806	
PR302	chr8:23540243	NKX3-1	Non_Synonymous_Coding	G	A	p.P54S	96	0.052	NM_006167.3		
PR302	chr16:72828408	ZFXH3	Non_Synonymous_Coding	C	A	p.A2725S	7963	0.061	NM_006885.3		
PR302	chr7:91682269	AKAP9	Non_Synonymous_Coding	T	A	p.S1866R	60	0.117	NM_005751.4		
PR302	chr8:71036150	NCOA2	Non_Synonymous_Coding	G	A	p.T1421M	5550	0.451	NM_006540.2		
PR302	chr3:38941459	SCN11A	Non_Synonymous_Coding	G	A	p.L650F	7862	0.483	NM_014139.2		COSM136332
PR304	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	115	0.461	NM_005269.2	rs2229300	
PR305	chrX:70360666	MED12	Stop_Gained	C	T	p.Q2076*	50	0.06	NM_005120.2		
PR305	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	309	0.469	NM_005269.2	rs2229300	
PR306	chr12:49444990	KMT2D	Non_Synonymous_Coding	G	C	p.P826A	106	0.057	NM_003482.3		

Contd...

Supplementary Table 4: Contd...

Sample	Location	Gene	Variant classification	Reference allele	Variant allele	AA change	Filtered coverage	Variant allele frequency	RefSeq	dbSNP ID	COSMIC ID
PR306	chr2:137988661	THSD7B	Non_Synonymous_Coding	A	C, G	p.R560G	32	0.094	NM_001080427.1		
PR306	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	117	0.496	NM_005269.2	rs2229300	
PR307	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	377	0.44	NM_005269.2	rs2229300	
PR307	chr7:91668025	AKAP9	Non_Synonymous_Coding	G	C	p.C1544S	2296	0.472	NM_005751.4		
PR307	chr7:91709065	AKAP9	Non_Synonymous_Coding	A	G	p.I2540V	1978	0.502	NM_005751.4		
PR307	chr16:72823230	ZFXH3	Non_Synonymous_Coding	G	C	p.Q1421E	7856	0.516	NM_006885.3		
PR307	chr16:72829373	ZFXH3	Non_Synonymous_Coding	G	A	p.S2403F	4584	0.517	NM_006885.3		
PR308	chr12:49425853	KMT2D	Non_Synonymous_Coding	C	T	p.R4212Q	3047	0.493	NM_003482.3		
PR309	chr2:138417295	THSD7B	Non_Synonymous_Coding	A	C	p.N1450T	2559	0.032	NM_001080427.1		
PR309	chr16:72821079	ZFXH3	Non_Synonymous_Coding	T	G	p.D3699A	3002	0.033	NM_006885.3		
PR309	chr19:50549280	ZNF473	Non_Synonymous_Coding	G	A	p.G527D	10 005	0.034	NM_001006656.1		
PR309	chr12:49431013	KMT2D	Non_Synonymous_Coding	C	G	p.V3376L	594	0.042	NM_003482.3		
PR309	chr3:178916786	PIK3CA	Non_Synonymous_Coding	T	C	p.L58P	1431	0.043	NM_006218.2		
PR309	chr3:38904766	SCN11A	Non_Synonymous_Coding	T	C	p.T1326A	2150	0.045	NM_014139.2		
PR309	chr5:112173291	APC	Non_Synonymous_Coding	A	G	p.Q667R	736	0.045	NM_001127510.2		
PR309	chr3:73450149	PDZRN3	Non_Synonymous_Coding	G	A	p.A393V	1574	0.047	NM_015009.1		COSM1425294
PR309	chr12:49436006	KMT2D	Non_Synonymous_Coding	T	C	p.E1992G	64	0.047	NM_003482.3		
PR309	chr2:138163363	THSD7B	Non_Synonymous_Coding	G	A	p.R863Q	915	0.048	NM_001080427.1		
PR309	chr16:72991476	ZFXH3	Non_Synonymous_Coding	C	T	p.A857T	1570	0.049	NM_006885.3	rs146607776	COSM1379726
PR309	chr5:112176066	APC	Non_Synonymous_Coding	A	G	p.K1592R	1204	0.052	NM_001127510.2		
PR309	chr13:48934201	RB1	Non_Synonymous_Coding	T	C	p.M219T	538	0.052	NM_000321.2		
PR309	chr13:48934204	RB1	Non_Synonymous_Coding	T	C	p.L220P	534	0.052	NM_000321.2		
PR309	chr17:37687207	CDK12	Non_Synonymous_Coding	G	C	p.V1371L	323	0.053	NM_016507.2		
PR309	chrX:70346837	MED12	Non_Synonymous_Coding	G	A	p.V902I	668	0.054	NM_005120.2		COSM1124669
PR309	chr2:138425344	THSD7B	Non_Synonymous_Coding	A	G	p.M1522V	1208	0.058	NM_001080427.1		
PR309	chr16:72828408	ZFXH3	Non_Synonymous_Coding	C	A	p.A2725S	9128	0.058	NM_006885.3		
PR309	chr5:112174845	APC	Non_Synonymous_Coding	C	T	p.T1185I	461	0.059	NM_001127510.2		
PR309	chr5:112179113	APC	Non_Synonymous_Coding	G	A	p.A2608T	494	0.059	NM_001127510.2		
PR309	chr5:112177895	APC	Non_Synonymous_Coding	A	G	p.K2202E	275	0.062	NM_001127510.2		
PR309	chr7:91631131	AKAP9	Non_Synonymous_Coding	G	A	p.E634K	699	0.062	NM_005751.4	rs368006874	
PR309	chr3:178943760	PIK3CA	Non_Synonymous_Coding	A	T	p.Q809H	302	0.063	NM_006218.2		
PR309	chr12:56428449	IKZF4	Non_Synonymous_Coding	C	G	p.S364R	560	0.066	NM_022465.3		
PR309	chrX:70347260	MED12	Non_Synonymous_Coding	T	C	p.L975P	2086	0.07	NM_005120.2		
PR309	chr7:91609645	AKAP9	Non_Synonymous_Coding	G	C	p.E117Q	113	0.08	NM_005751.4		
PR309	chr8:23540200	NKX3-1	Non_Synonymous_Coding	C	T	p.R68H	191	0.089	NM_006167.3		
PR309	chr16:72828173	ZFXH3	Non_Synonymous_Coding	G	A	p.S2803F	357	0.098	NM_006885.3		
PR309	chr17:7577147	TP53	Non_Synonymous_Coding	A	T	p.L264Q	61	0.098	NM_000546.5		
PR309	chr12:57865549	GLI1	Non_Synonymous_Coding	C	T	p.P1009L	29	0.172	NM_005269.2		
PR309	chr7:91682269	AKAP9	Non_Synonymous_Coding	T	G	p.S1866R	47	0.255	NM_005751.4		
PR309	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGGGACAGGGGTGAC	T	p.ESPLSPPEE446E	8861	0.35	NM_003482.3		
PR309	chr16:72831357	ZFXH3	Coding_Deletion	CTTGTG	C	p.QQ1740-	2881	0.431	NM_006885.3		COSM1731926
PR309	chr16:72832153	ZFXH3	Non_Synonymous_Coding	C	T	p.M1476I	1414	0.592	NM_006885.3	rs200133878	

Contd...

Supplementary Table 4: Contd...

Sample	Location	Gene	Variant classification	Reference allele	Variant allele	AA change	Filtered coverage	Variant allele frequency	RefSeq	dbSNP ID	COSMIC ID
PR310	chr2:137988662	THSD7B	Non_Synonymous_Coding	G	C	p.R560T	49	0.102	NM_001080427.1		
PR311	chr16:72993791	ZFXH3	Non_Synonymous_Coding	G	A	p.P105L	1658	0.154	NM_006885.3	rs199521581	COSM130307
PR312	chr16:72821887	ZFXH3	Frame_shift	C	CG	p.-3429?	3536	0.154	NM_006885.3		
PR313	chr2:137988662	THSD7B	Non_Synonymous_Coding	G	A	p.R560K	39	0.077	NM_001080427.1		
PR313	chr7:91712775	AKAP9	Non_Synonymous_Coding	A	C	p.T2818P	62	0.113	NM_005751.4		
PR313	chr12:49432474	KMT2D	Non_Synonymous_Coding	C	T	p.G2889R	31	0.129	NM_003482.3		
PR313	chr3:73657760	PDZRN3	Non_Synonymous_Coding	G	A	p.R267W	1517	0.477	NM_015009.1	rs373980887	
PR315	chr19:5144410	KDM4B	Non_Synonymous_Coding	C	T	p.P963L	21	0.238	NM_015015.2		
PR315	chr7:91739463	AKAP9	Non_Synonymous_Coding	T	C	p.M3905T	2857	0.476	NM_005751.4	rs77447750	
PR315	chr19:5134026	KDM4B	Non_Synonymous_Coding	C	T	p.T680M	3383	0.514	NM_015015.2	rs149752315	
PR315	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	498	0.552	NM_005269.2	rs2229300	
PR401	chr12:49444992	KMT2D	Non_Synonymous_Coding	G	C	p.S825C	155	0.052	NM_003482.3		
PR401	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	63	0.349	NM_005269.2	rs2229300	
PR401	chr8:71071746	NCOA2	Non_Synonymous_Coding	A	G	p.L373P	24	0.375	NM_006540.2		
PR401	chr12:49444996	KMT2D	Frame_shift	A	AG	p.-823?	127	0.701	NM_003482.3		
PR403	chr16:72821887	ZFXH3	Frame_shift	C	CG	p.-3429?	3159	0.172	NM_006885.3		
PR403	chr12:49445021	KMT2D	Frame_shift	C	CG, CGG	p.-815?	85	0.376	NM_003482.3		
PR403	chr12:57865558	GLI1	Non_Synonymous_Coding	G	T	p.G1012V	517	0.472	NM_005269.2	rs2229300	
PR404	chr16:72991476	ZFXH3	Non_Synonymous_Coding	C	T	p.A857T	2706	0.04	NM_006885.3	rs146607776	COSM1379726
PR404	chr16:72991494	ZFXH3	Non_Synonymous_Coding	C	T	p.G851S	2731	0.044	NM_006885.3		
PR404	chr16:72991473	ZFXH3	Non_Synonymous_Coding	C	T	p.E858K	2757	0.051	NM_006885.3		
PR404	chr12:49436073	KMT2D	Non_Synonymous_Coding	C	T	p.D1970N	138	0.094	NM_003482.3		
PR404	chr16:72991700	ZFXH3	Non_Synonymous_Coding	G	A	p.A782V	54	0.148	NM_006885.3	rs62639304	
PR404	chr16:72991706	ZFXH3	Non_Synonymous_Coding	G	A	p.A780V	53	0.151	NM_006885.3		
PR404	chr12:49446101	KMT2D	Codon_Change_Plus_Coding	TTCTCAGGTGGTGGGACACAGGGGTGAC	T	p.ESPLSPPEE446E	6052	0.282	NM_003482.3		
PR414	chr12:49416542	KMT2D	Non_Synonymous_Coding	C	T	p.R5390Q	5172	0.485	NM_003482.3		
PR414	chr19:50549441	ZNF473	Non_Synonymous_Coding	G	A	p.E581K	10 668	0.487	NM_001006656.1	rs150885598	
PRY01	chr12:49445021	KMT2D	Frame_shift	C	CG, CGG	p.-815?	191	0.424	NM_003482.3		
PRY02	chr19:50549280	ZNF473	Non_Synonymous_Coding	G	A	p.G527D	2445	0.034	NM_001006656.1		
PRY02	chr16:72821887	ZFXH3	Frame_shift	C	CG	p.-3429?	1603	0.224	NM_006885.3		
PRY02	chr4:87313	ZNF595	Frame_shift	G	GA	p.-639?	62	0.968	NM_182524.2	rs60154095	
PRY04	chr16:72827613	ZFXH3	Non_Synonymous_Coding	C	T	p.V2990I	509	0.063	NM_006885.3		
PRY04	chr16:72821887	ZFXH3	Frame_shift	C	CG	p.-3429?	1684	0.249	NM_006885.3		
PRY04	chr16:72821887	ZFXH3	Frame_shift	C	CG	p.-3429?	1684	0.249	NM_006885.3		
PRY04	chr4:87313	ZNF595	Frame_shift	G	GA	p.-639?	66	0.879	NM_182524.2	rs60154095	

PTEN: phosphatase and tensin homolog; AR: androgen receptor