

SUPPLEMENTARY TABLES

Supplementary Table 1. Number of mutations and differentially expressed genes in each patient

Patient N° (this study)	N° of passed mutations	N° of moderate mutations	N° of high mutations	N° of DEGs (FC > 2; adj p -value < 0.01)
1	86	34	9	1982
2	105	39	5	3170
3	71	29	7	2808
4	68	21	5	2652
5	49	20	6	4029
6	53	18	6	1034
7	87	31	8	2889
8	77	31	6	1439
9	83	27	3	3528
10	55	21	1	2159
11	83	31	11	1215
12	114	43	16	3438
13	89	27	6	1797
14	63	21	2	3563
15	68	24	2	2453

Patient samples that complied with the histological selection criteria for homogeneity and tumor cellularity were renumbered from 1 to 15. Mutations were classified as defined by Ensembl. Duplicate RNA-seq of tumor and patient-matched normal tissue was used to define differentially expressed genes (DEGs) by counting the reads with Genomic Alignments (1.17.3) and then using DESeq2 (1.17.3) with R (3.5.0). The results have been obtained following the general guidelines given in the DESeq2 tool. We used a Log2FC of -1/+1 (+2/-2) and a cutoff for the adjusted p-value (Q-value) < 0.01. All main figures have been generated using the high, moderate and modifier mutations specified in Supplementary Data File 1.

Supplementary Table 2. Novel mutations previously not reported for prostate cancer patients

Patient 1		Total number of somatic mutations: 86	
Gene	Type of Variant	New Mutation	Function
<i>OR10H1</i>	Missense	T57M	olfactory receptor
<i>CDHR5</i>	Splicing mut	Splice donor exon 6	member of the cadherin superfamily; enriched in colorectal cancer, liver cancer, pancreatic cancer, renal cancer, stomach cancer
<i>HSD17B6</i>	Missense	R52Q	17 Beta-hydroxysteroid dehydrogenase type 6; involved in intra-tumoral synthesis of DHT
<i>TRMT10A</i>	Missense	R275Q	tRNA (Guanine-1)-methyltransferase
<i>BANP</i>	Missense	R246H	forms complex with p53 & negatively regulates p53 transcription, tumor suppressor and cell cycle regulator
<i>DGAT2</i>	Missense	L91R	Diacylglycerol O-Acyltransferase 2
<i>CASD1</i>	Missense	I755S	O-acetyltransferase
<i>TTC28</i>	Frameshift	L1164LLTC.....	During mitosis, may be involved in the condensation of spindle midzone microtubules, leading to the formation of midbody
<i>ITGB7</i>	Frameshift & splice region	PCERHR644PGL CR.....	Integrin Subunit Beta 7 (ECM signaling)
<i>FEZ1</i>	Stop gain	E65*	leucine-zipper protein, its expression is altered in multiple human tumors; candidate tumor suppressor
<i>TINAGL1</i>	Missense	A424V	Inhibits TNBC progression and metastasis; prognostic marker in renal cancer (favourable) and thyroid cancer (favourable)
<i>ZFR2</i>	Missense	Q632H	Zinc-finger RNA binding protein
<i>SLC35B2</i>	Missense	S218F	Predicted membrane protein Transporter; Prognostic marker in liver (unfavorable), prostate (favorable) & breast cancer (unfavorable)
<i>PPP2R5D</i>	Missense	L183Q,	protein phosphatase 2 regulatory subunit B Delta isoform
<i>WDFY1</i>	Missense	L149P	phosphatidylinositol 3-phosphate binding protein (signaling)

Patient 2		Total number of somatic mutations: 105	
Gene	Type of Variant	New Mutation	Function
<i>NBR1</i>	Missense	V96I	Autophagy cargo receptor
<i>RHPN2</i>	Missense	V73M	Rhopilin-Like Rho-GTPase Binding Protein
<i>SH3PXD2A</i>	Missense	T847P	Adapter protein adhesion
<i>MUC2</i>	Missense	T2305A	Member of mucin family
<i>CHN2</i>	Missense	R47W	Role in cell proliferation and migration
<i>CCDC169</i>	Missense	R197T	Coiled-Coil Domain-Containing Protein 169
<i>SPTLC2</i>	Frameshift Ins	P467SSS...,	Serine palmitoyltransferase
<i>TRIM49C</i>	Missense	K363E	Uncharacterized
<i>BRF1</i>	Missense	K282Q	RNA Polymerase III Transcription Initiation Factor Subunit
<i>HIST1H3J</i>	Missense	H114Y	AR signaling
<i>ZBTB42</i>	Missense	F82C	Transcriptional repressor
<i>LRRCC1</i>	Missense	E525K	Leucine Rich Repeat & Coiled-Coil Centrosomal Protein 1
<i>ZNF44</i>	Missense	E348A	Uncharacterized
<i>PPP1R26</i>	Missense	C186F	Phosphatase

Patient 3		Total number of somatic mutations: 71	
Gene	Type of Variant	New Mutation	Function
<i>GSG1L2</i>	Missense	V72L	Germ Cell-Specific Gene 1-Like Protein 2-Like, Uncharacterized
<i>ASB15</i>	Missense	V358I	Participate in the ubiquitin-proteasome system for the degradation of proteins in the cell cycle and signal transduction pathways
<i>SH3PXD2A</i>	Missense	T847P	Role in cell migration

<i>TOPAZ1</i>	Missense	R85H	Required for progression to the post-meiotic stages of spermatocyte development
<i>ZNF280A</i>	Missense	R457Q	Uncharacterized
<i>OTOG</i>	Missense	R2294C	Component of the acellular membranes of the inner ear
<i>LTK</i>	Missense	R221H	Tyrosine kinase
<i>LRRTM4</i>	Missense	Q87K	Development and maintenance of the vertebrate nervous system
<i>JAK2</i>	Missense	P121L	Tyrosine kinase
<i>RGPD2</i>	Missense	L12V	RNA transport
<i>KRT5</i>	in-frame del	GGGLGGGLA517G	Member of keratin gene family
<i>TEAD4</i>	Missense	I242L	Hippo signaling, EMT
<i>TTC34</i>	Missense	G538S	Uncharacterized
<i>CCDC71L</i>	Missense	G28V	Uncharacterized
<i>DUOXA1</i>	Missense	F130L	Involved in hydrogen peroxide production necessary for thyroid hormonogenesis
<i>VWCE</i>	Missense	A58V	Regulatory element in the beta-catenin signaling pathway

Patient 4		Total number of somatic mutations: 68	
Gene	Type of Variant	New Mutation	Function
<i>DERL3</i>	Missense	V106I	Degrades misfolded glycoproteins in the ER
<i>PDGFB</i>	Missense	R217W	catalytic receptors with intracellular tyrosine kinase activity. Regulate embryonic development, angiogenesis, cell proliferation and differentiation.
<i>TEKT4</i>	Missense & Stop gain	L366V	Contributes to sperm motility
<i>TET3</i>	Missense	L1035M	Methylcytosine Dioxygenase
<i>PRH2</i>	Missense	I42L	Inhibitor of calcium phosphates
<i>TAS1R3</i>	Missense	G314D	G-protein coupled receptor
<i>KRTAP3-2</i>	Frameshift del	A16APP...	Keratin associated protein
<i>DRG2</i>	Missense	A135T	GTP binding protein
<i>NPFF</i>	Missense	R21H	Morphine modulating peptide

Patient 5		Total number of somatic mutations: 49	
Gene	Type of Variant	New Mutation	Function
<i>SCCPDH</i>	Missense & splice region	Y102C	Oxidoreductase
<i>WRN</i>	Missense	V1155L	Helicase
<i>ZBED5</i>	Missense & Stop gain	Q329*	Not well characterized
<i>HDAC1</i>	Missense	R77S	Histone deacetylase
<i>TPSB2</i>	Missense	R22P	Serine protease
<i>ZNF777</i>	Missense	P158A	Not well characterized
<i>IL15</i>	Start lost	M1L	Cytokine activity
<i>FEZ2</i>	Missense	K278E	Involved in axonal outgrowth
<i>FKBP9</i>	Missense	H620Q	Calcium ion binding
<i>IFITM2</i>	frameshift del	R86RRW	Cytokine signaling
<i>PIM3</i>	in-frame del	AT226	Ser/thr kinase

Patient 6		Total number of somatic mutations: 53	
Gene	Type of Variant	New Mutation	Function
<i>ZP1</i>	Missense	Y271S	Structural component of zona pellucida
<i>FASTKD3</i>	Missense	R216H	Kinase
<i>MAST3</i>	Missense	M786V	Kinase

<i>TGFB3</i>	Missense & splice region	E119K	TGF beta/Smad signaling network
<i>DLL1</i>	Stop gain	C44*	Notch ligand
<i>ZBED8</i>	Missense	A523S	Uncharacterized

Patient 7		Total number of somatic mutations: 87	
Gene	Type of Variant	New Mutation	Function
<i>COLCA2</i>	Missense & splice region	T34N	Colorectal Cancer Associated 2
<i>FASTKD3</i>	Missense	R216H	Kinase
<i>ATG101</i>	Missense	T133K	Autophagy
<i>HTATIP2</i>	Missense	R49S	Oxidoreductase required for tumor suppression
<i>ALDH2</i>	Missense	R338Q	Oxidoreductase
<i>TTC29</i>	Frameshift	N499K*	Uncharacterized
<i>ARHGAP45</i>	Missense	L371Q	GTPase activator
<i>EMC9</i>	Missense	F30Y	ER membrane protein
<i>CCDC168</i>	Missense	D5448G	Uncharacterized
<i>HLA-DRB1</i>	Missense	D31G	Antigen presentation
<i>SLC5A9</i>	Missense	A386V	Glucose transport
<i>LRRC18</i>	Missense	A217T	Sperm maturation
<i>SHROOM2</i>	Missense	A1406V	Endothelial cell morphology

Patient 8		Total number of somatic mutations: 77	
Gene	Type of Variant	New Mutation	Function
<i>ZNF354C</i>	Missense	V176I	Transcriptional repressor
<i>ZMYM6</i>	Missense	S300C	Regulates cell morphology
<i>RABGGTB</i>	Frameshift & stop gain	R65RMN....	RAB GTPase Binding
<i>ALDH2</i>	Missense	R338Q	Oxidoreductase
<i>SCRN1</i>	Missense	R50W	Peptidase
<i>RASSF8</i>	Missense	R208H	Tumor suppressor
<i>ITPRIPL2</i>	Missense	P171A	Intracellular calcium signaling
<i>ZBTB14</i>	Missense	H355Q	Not well characterized
<i>CYB5A</i>	Missense	F63V	Cytochrome C oxidase
<i>EFNA1</i>	Missense	E55A	Receptor tyrosine kinase
<i>SLC11A1</i>	Missense	A505S	Fe and Mn transporter
<i>OR5M9</i>	Missense	A298P	Signaling by GPCR

Patient 9		Total number of somatic mutations: 83	
Gene	Type of Variant	New Mutation	Function
<i>METTL4</i>	Missense	Y79F	Probable methyltransferase
<i>ROGDI</i>	Missense	V244M	Uncharacterized
<i>ZIC2</i>	Missense & in-frame del	S422C	Transcription repressor/activator
<i>MAPK7</i>	Missense	R400H	Role in proliferation
<i>AMIGO3</i>	Missense	R293W	May regulate cell-cell interaction
<i>GABPB2</i>	Missense	R117Q	May function as TF for purine rich repeats
<i>OLFM3</i>	Missense	N251S	Uncharacterized
<i>OR5AP2</i>	Missense	I220T	Olfactory receptor
<i>DBR1</i>	in-frame del	DD541D	RNA lariat debranching enzyme that hydrolyzes 2'-5' prime branched phosphodiester bonds

<i>GAK</i>	Missense & splice region	D1016Y	Cyclin G associated kinase
<i>GOPC</i>	Missense	A203V	Protein trafficking

Patient 10		Total number of somatic mutations: 55	
Gene	Type of Variant	New Mutation	Function
<i>ACSM4</i>	Missense	R325W	Fatty acid beta-oxidation
<i>RAB29</i>	Missense	P131A	Rab GTPase
<i>PPP1R10</i>	Missense & in-frame del	H896Q	Phosphatase
<i>TBL2</i>	Missense	H400Y	Beta transducin
<i>EHD3</i>	Missense	F18L	Membrane re-organization
<i>PMEL</i>	Missense	T419K	Melanosome biogenesis
<i>DNAH12</i>	Missense	A1501T	Dynein

Patient 11		Total number of somatic mutations: 83	
Gene	Type of Variant	New Mutation	Function
<i>TMEM132C</i>	Missense	V51G	Protein phosphatase
<i>SRM</i>	Missense	V112M	Spermidine biosynthesis
<i>WHRN</i>	Missense	S175T	Actin cytoskeletal assembly
<i>ADAMTS4</i>	Missense	R447H	Metalloproteinase
<i>SIGLEC12</i>	Missense	P198L	Protein carbohydrate interaction
<i>ANXA1</i>	Frameshift del	aa315*	Tumor suppressor. Inhibits phospholipase A2; anti-inflammatory
<i>STEAP1</i>	Frameshift del	aa244*	Cell surface antigen expressed at cell-cell junctions
<i>OR4E2</i>	Frameshift del	aa147*	Olfactory receptor
<i>ZFP30</i>	Missense	E159G	Not well characterized
<i>ISM1</i>	Missense	D181N	Uncharacterized
<i>SLC6A18</i>	Missense	G33E	Sodium transporter
<i>RNH1</i>	Missense	D436N	Ribonuclease inhibitor
<i>UBE2R2</i>	Missense	D232N	Beta-catenin degradation
<i>RHOBTB1</i>	Missense	V580I	Rho GTPase
<i>MMP15</i>	Missense	P575Q	Matrix Metalloproteinase
<i>SLC6A13</i>	Splice donor & intron	Splice site	GABA transporter

Patient 12		Total number of somatic mutations: 114	
Gene	Type of Variant	New Mutation	Function
<i>KCMF1</i>	Missense	Y82C	E3 Ub Ligase function
<i>SH3PXD2A</i>	Missense	T847P	Adapter protein podosome formation
<i>HOXB7</i>	Missense	T173M	Homeobox transcription factor
<i>CCDC168</i>	Stop gain	S4604*	Uncharacterized
<i>WNK2</i>	Missense	R874W	Mitogen-Activated Protein Kinase Kinase Kinase
<i>KRTAP19-8</i>	Stop gain	R50*	Keratin associated protein
<i>FAM198A</i>	Missense	P42Q	Uncharacterized
<i>KIR3DL2</i>	Missense, frame-shift ins & del	N239S	Killer cell immunoglobulin-like receptor
<i>SEC11A</i>	Missense	M156I	Peptidase
<i>ANKRD31</i>	Stop gain	L276*	Uncharacterized
<i>COPB1</i>	Missense	K450Q	Vesicle transport
<i>TEAD4</i>	Missense	I242L	Transcription factor, hippo pathway, EMT
<i>SKOR2</i>	Missense	F947S	Transcriptional repressor
<i>DENND5B</i>	frameshift	AKRTG461VWLW...,	Guanine nucleotide exchange factor

<i>MUC2</i>	Missense	S1943R	Member of the mucin family
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Patient 13		Total number of somatic mutations: 89	
Gene	Type of Variant	New Mutation	Function
<i>CYP24A1</i>	Missense	Y145H	Cytochrome p 450
<i>MRPL46</i>	Missense	R7W	Mitochondrial ribosomal protein
<i>RIBC2</i>	Missense	R112C	Uncharacterized
<i>INF2</i>	Missense	P1163H	(de)polymerization of actin filaments
<i>OR10V1</i>	Missense	R15C	Olfactory receptor
<i>NKAPL</i>	Missense	M266I	Transcriptional repressor of Notch signaling
<i>CAMK2G</i>	Missense	I45T	Calmodulin dependent protein kinase
<i>TANC1</i>	Missense	D1438N	Not well characterized

Patient 14		Total number of somatic mutations: 63	
Gene	Type of Variant	New Mutation	Function
<i>UBE2J1</i>	Missense & splice region	Y227H	Ubiquitin conjugating enzyme
<i>TET2</i>	Stop gain	S774*	Methylcytosine Dioxygenase
<i>SLC39A11</i>	Missense	S153R	Metal ion transporter
<i>SPATA21</i>	Missense	R227C	Differentiation of haploid spermatids
<i>PRR36</i>	Missense	P832H	Uncharacterized
<i>GOLGA8K</i>	Missense	L311V	Uncharacterized
<i>TAAR1</i>	Missense	K188N	G-Protein coupled receptor
<i>OR2W1</i>	Missense	I256L	Olfactory receptor
<i>HMCN2</i>	Missense	G3327S	Calcium ion binding

Patient 15		Total number of somatic mutations: 68	
Gene	Type of Variant	New Mutation	Function
<i>ATP1A4</i>	Missense	V287A	Cation transport ATPase
<i>SCLT1</i>	Missense	T468M	Sodium channel regulator
<i>MCEMP1</i>	Missense	T159M	Not well characterized
<i>MARCH11</i>	Missense	R311C	Membrane-bound E3 ubiquitin ligases
<i>CASP5</i>	Missense	R24H	Aspartate protease (caspase)
<i>ZRANB3</i>	Missense	M162L	DNA annealing helicase and endonuclease
<i>C5orf30</i>	Missense	K108E	Putative role in protein trafficking <i>via</i> interaction with UNC119 and UNC119B cargo adapters
<i>P2RX7</i>	Missense	G27V	Ligand gated ion channel, receptor for ATP
<i>LRRC45</i>	Stop gain	E564*	Centrosome linker required for cohesion
<i>CHRNA7</i>	Missense	C450Y	Muscle-type acetylcholine receptor

Red, mutations considered to have important cell biological effects.

Supplementary Table 3. Annotated differentially expressed (>2-fold) lncRNAs and miRs in each patient.

Patient	Differentially expressed lncRNAs	Differentially expressed miRs
1	HOTTIP, SCHLAP1, NEAT1, PCA3, DANCER, MEG3, ARLNC1	MIR205HG, MIR4300HG, MIR4458HG, MIR646HG, MIR9-3HG, MIR99AHG
2	SCHLAP1, NEAT1, PCAT1, PCA3, DANCER, MEG3	MIR1-1HG-AS1, MIR205HG, MIR222HG, MIR23A, MIR27A, MIR5687, MIR5704
3	PCAT1, PCA3, DANCER, ARLNC1	MIR181A1HG, MIR205HG, MIR22HG, MIR27A, MIR3142HG, MIR4458HG, MIR5687, MIR5704, MIR9-3HG
4	HOTTIP, HOTAIRM1, NKILA, SCHLAP1, PCAT1, PCA3, ARLNC1	MIR1-1HG-AS1, MIR2052HG, MIR222HG, MIR3142HG, MIR3189, MIR34AHG, MIR3648-2, MIR3687-2, MIR4300HG, MIR4435-2HG, MIR4500HG, MIR4796, MIR5704, MIR578, MIR646HG, MIRLET7BHG
5	HOTTIP, NKILA	MIR100HG, MIR222HG, MIR3189, MIR4453HG, MIR4458HG, MIR5704
6	HOTTIP, NEAT1, ARLNC1	MIR205HG, MIR3142HG, MIR4429, MIR4435-2HG, MIR561
7	HOTTIP, NKILA, SCHLAP1, NEAT1, PCAT1, PCA3, DANCER, MEG3	MIR1-1HG-AS1, MIR205HG, MIR222HG, MIR22HG, MIR23A, MIR27A, MIR3142HG, MIR5687, MIR5704
8	HOTTIP, SCHLAP1, PCAT1, PCA3, DANCER, MEG3	MIR1-1HG-AS1, MIR205HG, MIR222HG
9	HOTTIP, NEAT1, PCAT1, PCA3, DANCER, ARLNC1	MIR100HG, MIR1-1HG-AS1, MIR155HG, MIR193BHG, MIR2052HG, MIR205HG, MIR222HG, MIR3648-2, MIR5704, MIR5706, MIR99AHG, MIRLET7BHG
10	HOTTIP, SCHLAP1, NEAT1, PCAT1, PCA3, MEG3, ARLNC1	MIR1-1HG-AS1, MIR222HG, MIR22HG, MIR34AHG, MIR5687, MIR5704
11	SCHLAP1, PCAT, ARLNC1	MIR1251, MIR137HG, MIR22HG, MIR23A, MIR27A, MIR3648-2, MIR578, MIR646HG
12	HOTTIP, HOTAIRM1, SCHLAP1, NEAT1, PCAT1, PCA3, DANCER, MEG3, ARLNC1	MIR100HG, MIR1-1HG-AS1, MIR17HG, MIR200CHG, MIR205HG, MIR222HG, MIR25, MIR3142HG, MIR4429, MIR4697HG, MIR503HG, MIR5706, MIR635, MIR646HG, MIR99AHG
13	NKILA, SCHLAP1, PCAT1, MEG3	MIR1-1HG-AS1, MIR205HG, MIR22HG, MIR34AHG, MIR4697HG
14	SCHLAP1, PCAT1, PCA3, MEG3, ARLNC1	MIR100HG, MIR1-1HG, MIR1-1HG-AS1, MIR1255A, MIR193BHG, MIR205HG, MIR222HG, MIR27B, MIR4697HG, MIR5687, MIR99AHG
15	NKILA, SCHLAP1, PCAT1, PCA3, MEG3, ARLNC1	MIR205HG, MIR3189, MIR34AHG, MIR4664, MIR4697HG, MIR5687, MIR646HG, MIR99AHG

Functionally relevant annotated miRs and lncRNAs were taken from[1-9]. Red, upregulated in tumors; black, downregulated in tumors relative to the patient-matched normal prostate tissue. All patient-specific deregulated miRs and lncRNAs can be found in Supplementary Data File 2.

Supplementary Table 4. Sequencing statistics for RNA-seq and EXOME-seq

Normal	RNA-SEQ		EXOME-SEQ			Tumor	RNA-SEQ		EXOME-SEQ		
	Reads	Align %	Reads	Align %	Duplicate reads %		Reads	Align %	Reads	Align %	Duplicate reads %
1N_rep1	36603682	92.26	42475200	96.07	9.12	1T_rep1	49485994	91.34	55706200	96.12	9.12
1N_rep2	38501720	92.64				1T_rep2	52252504	91.78			
2N_rep1	35578814	90.50	56298000	96.51	11.09	2T_rep1	34340289	86.04	44136000	96.09	10.30
2N_rep2	63894243	91.60				2T_rep2	53856558	91.50			
3N_rep1	33037647	88.49	45167400	95.96	11.62	3T_rep1	39206644	83.38	45636400	95.90	9.85
3N_rep2	38061308	87.49				3T_rep2	44685255	87.41			
4N_rep1	39635442	87.12	45418800	96.31	8.21	4T_rep1	47273179	84.77	49255300	96.06	8.41
4N_rep2	54890556	92.23				4T_rep2	53502497	91.40			
5N_rep1	40283379	91.73	49694100	96.43	7.78	5T_rep1	64602422	94.25	43400000	96.26	13.96
5N_rep2	46638779	90.20				5T_rep2	47515766	91.17			
6N_rep1	46031895	86.04	34157000	96.06	9.98	6T_rep1	33455857	87.00	45420600	96.29	8.19
6N_rep2	55861482	91.92				6T_rep2	76033143	92.16			
7N_rep1	31882647	87.96	43500400	96.04	11.28	7T_rep1	47721526	86.91	47714500	96.02	11.74
7N_rep2	42675237	87.76				7T_rep2	38438256	93.03			
8N_rep1	35333114	83.44	54276700	96.33	9.09	8T_rep1	74598149	88.50	48035100	96.09	14.18
8N_rep2	80370444	94.85				8T_rep2	44121100	91.54			
9N_rep1	80833193	88.13	46282300	95.93	10.72	9T_rep1	39677083	90.05	48773100	96.04	7.44
9N_rep2	41562421	94.28				9T_rep2	38703772	93.42			
10N_rep1	53431330	85.30	53645600	96.29	8.11	10T_rep1	42928847	86.65	55294500	96.26	8.08
10N_rep2	51151259	91.93				10T_rep2	54227448	91.26			
11N_rep1	42821733	83.82	51965800	96.32	9.47	11T_rep1	48657860	82.43	57373400	96.42	11.73
11N_rep2	56544524	92.71				11T_rep2	53899463	92.03			
12N_rep1	37996964	91.08	55279200	96.12	16.20	12T_rep1	44233768	91.96	52977500	96.37	8.51
12N_rep2	40528066	91.61				12T_rep2	47182837	92.44			
13N_rep1	48075036	87.48	49494300	95.97	5.36	13T_rep1	51068154	87.50	60609000	96.16	11.58
13N_rep2	44318278	93.21				13T_rep2	48949742	91.89			
14N_rep1	36167783	86.15	46894500	96.11	9.67	14T_rep1	45615157	85.63	56130300	96.08	14.89
14N_rep2	45548318	90.53				14T_rep2	35440307	89.91			
15N_rep1	46655594	88.21	53319000	96.06	12.62	15T_rep1	43724856	85.14	49329900	96.03	11.78
15N_rep2	43936010	90.63				15T_rep2	66277516	91.99			

Strand specific paired-end RNA sequencing (Illumina Hiseq 2500, 125/150bp) of Ribo-depleted total RNA from biological duplicates (adjacent tissues sections). Quality of sequencing was assessed using FastQC tool. Exome sequencing was performed using Exome capture kit (Agilent) and subsequently sequenced on Hiseq2500 for 125 bp paired-end.

Supplementary Table 5. Number of target genes co-deregulated with the indicated cognate transcription factor in each prostate cancer sample.

P No	Total DEGs	ERG	FOXA1	HOXB13	MAZ	MYB	MYC	NKX2-1	PAX5
1	1982	462						191	
2	3170	756					1038	340	
3	2808	596					952		
4	2652	516				591	797		234
5	4029	910		291		947			
6	1034	174							69
7	2889	586			556		892	324	
8	1439	292					444		97
9	3528	762				759	1083	388	
10	2159	379		151			606		
11	1215	200				242			
12	3438	781			696		1105	377	232
13	1797	367							
14	3563	755	2337	245		736	1130		
15	2453	452	1602		480		759		

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