Transcription factor modulation upon prostate

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cancer stimulus with KB698 Treatment

QC

Downstream CHIP-seq data was processed through the Genialis platform

Collection title : Kronos_AR_&_FOXA1

Alignment: BWA ALN + batch mode on (Aquas parameters)

Build : hg19

Quality threshold : 5
Seed length : 32

(Default remaining settings)

Peak calling: MACS2 batch + batch mode on (Aquas parameters)

Promoter regions Bed file : HG19_TSS_ALL_-1000_+1000.bed

Use TagAlign files : Yes Quality filtering threshold : 30

Background : Pooled Input

(Default remaining settings)

QC

Alignment Quality

	FriP	Peak count	# Reads in promotors	Fraction reads in Promoters
Treat_FOXA1	0.17	50332	671070	0.044
Control_FOXA1	0.046	23062	725856	0.029
Treat_AR	0.14	46018	1925153	0.085
Control_AR	0.086	37057	1484441	0.06
	Total reads	Total mapped reads	% Mapped	
Treat_FOXA1	39225299	34444571	87.81%	
Control_FOXA1	45886429	43290108	94.34%	
Treat_AR	48498882	45126663	93.05%	
Control AR	44073133	40871557	92.74%	

Meme-CHIP was used to extract the motifs from the top 1000 peaks in each sample respectively

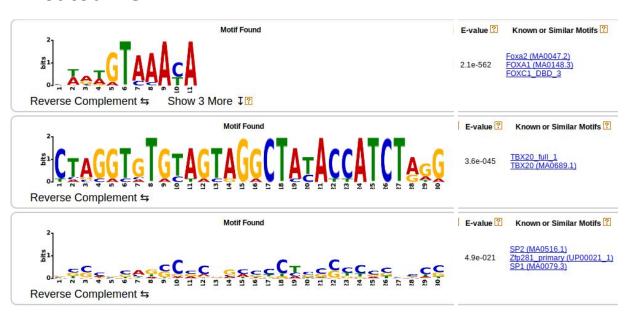
Narrow peak files were extracted and loaded on to Rstudio to sort and isolate the top 1000 peaks

The files were then converted into fasta files to use as the meme-chip input

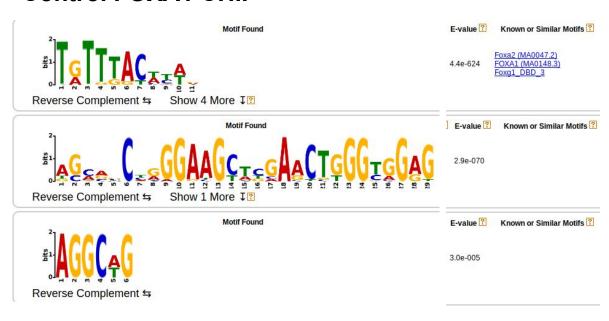
```
$ bedtools getfasta [OPTIONS] -fi <input FASTA> -bed <BED/GFF/VCF>
```

Files were then uploaded to Meme-Chip with default parameters

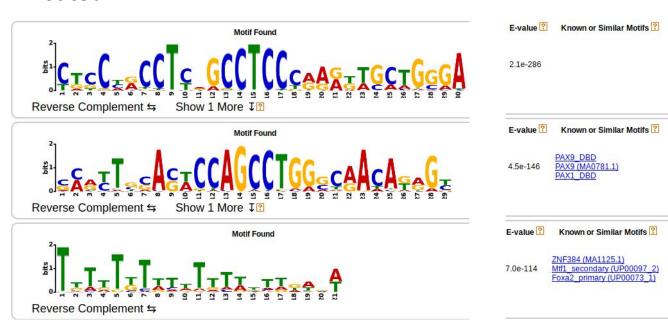
Treated FOXA1 CHIP



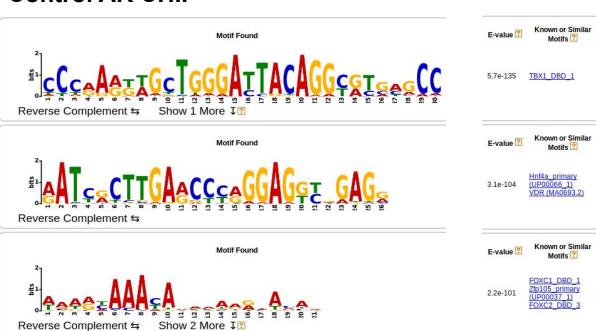
Control FOXA1 CHIP



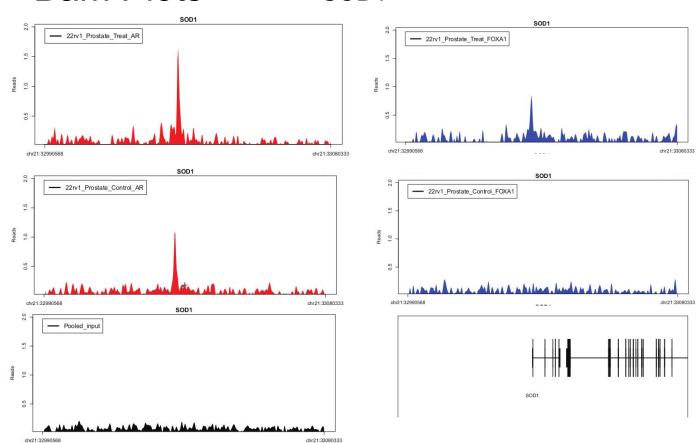
Treated AR CHIP



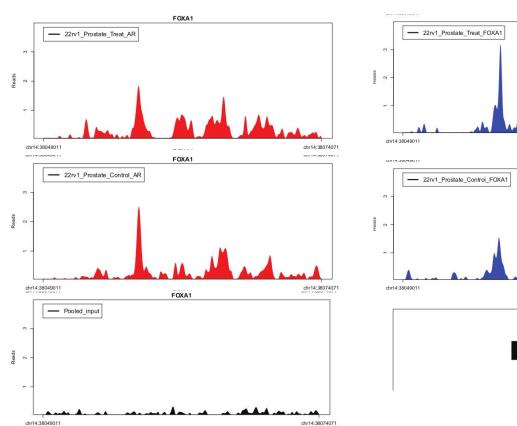
Control AR CHIP

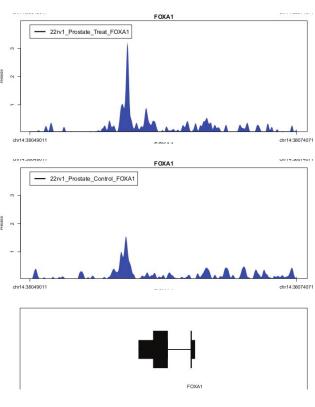


SOD1

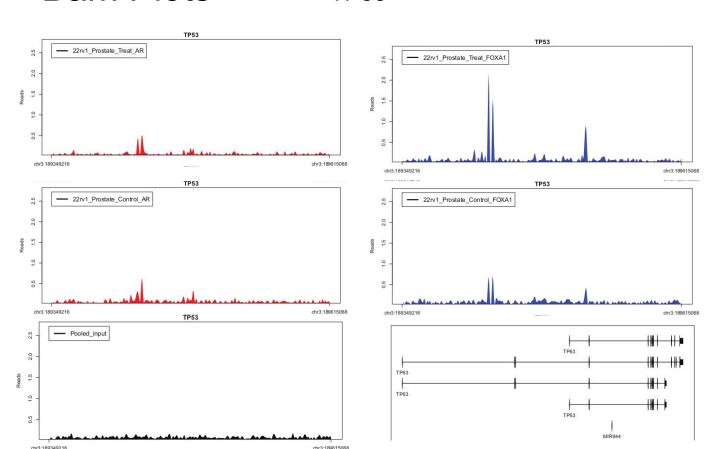


FOXA1

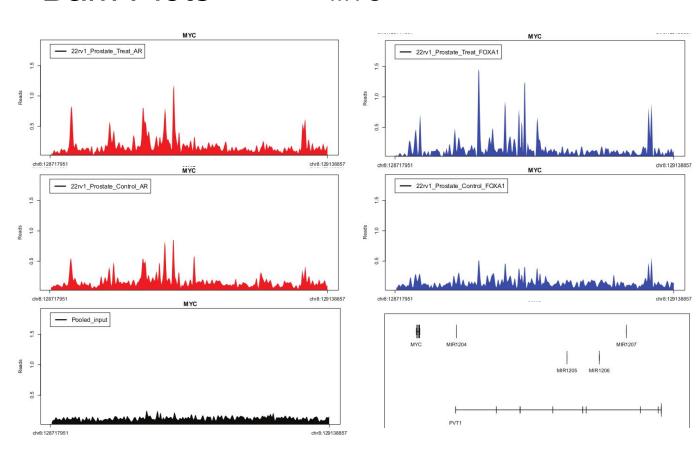




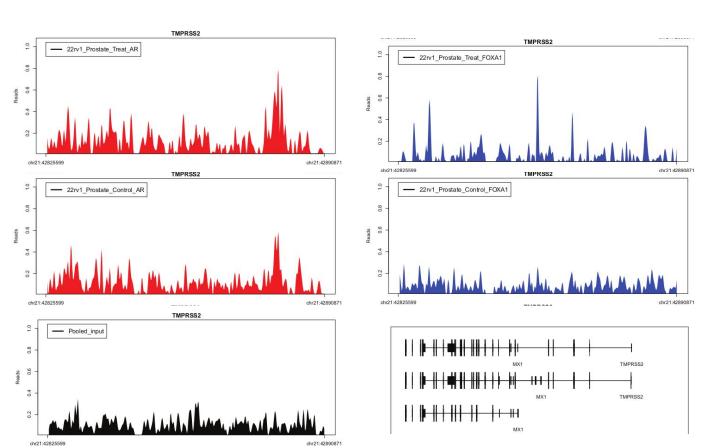
TP53



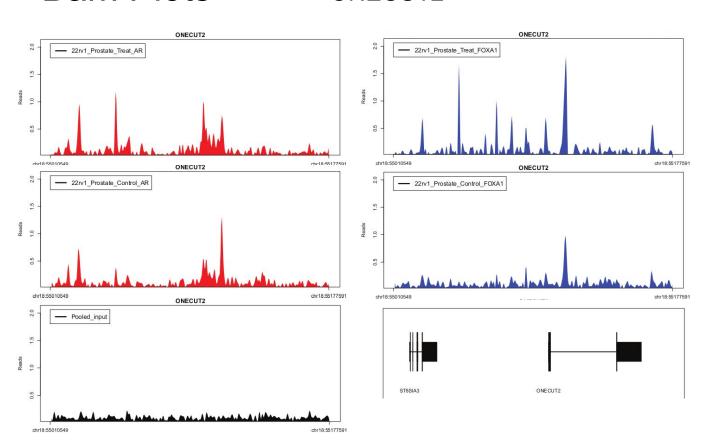
MYC



TMPRSS2



ONECUT2



UCSC Genome Browser (link)

http://torta.bcmt.bcm.edu/cgi-bin/hgTracks?db=hg19&lastVirtModeType=default&lastVirtModeExtraState=&virtModeType=default&virtMode=0&nonVirtPosition=&position=chr21%3A32990568%2D33080333&hgsid=644_VmV8XA3tCsxG8S34EeGjTa8Dv731