

Figure S4

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```
require(dplyr)
require(GenomicRanges)
require(TFCookbook)

load("ZFP3.motif.RData")
```

HEK293 cells data

```
ChIP.H293.peaks <- read.table("ENCFF107KSN.bed") %>%
  GenomicRanges::makeGRangesFromDataFrame(seqnames.field = "V1",
                                           start.field    = "V2",
                                           end.field      = "V3") %>% unique()

ZFP3.sites <- TFCookbook::matchPEM(PEM = ZFP3.Core.PEM,
                                   subject = ChIP.H293.peaks,
                                   genome   = "hg38",
                                   out      = "positions",
                                   E.cutoff = -5)

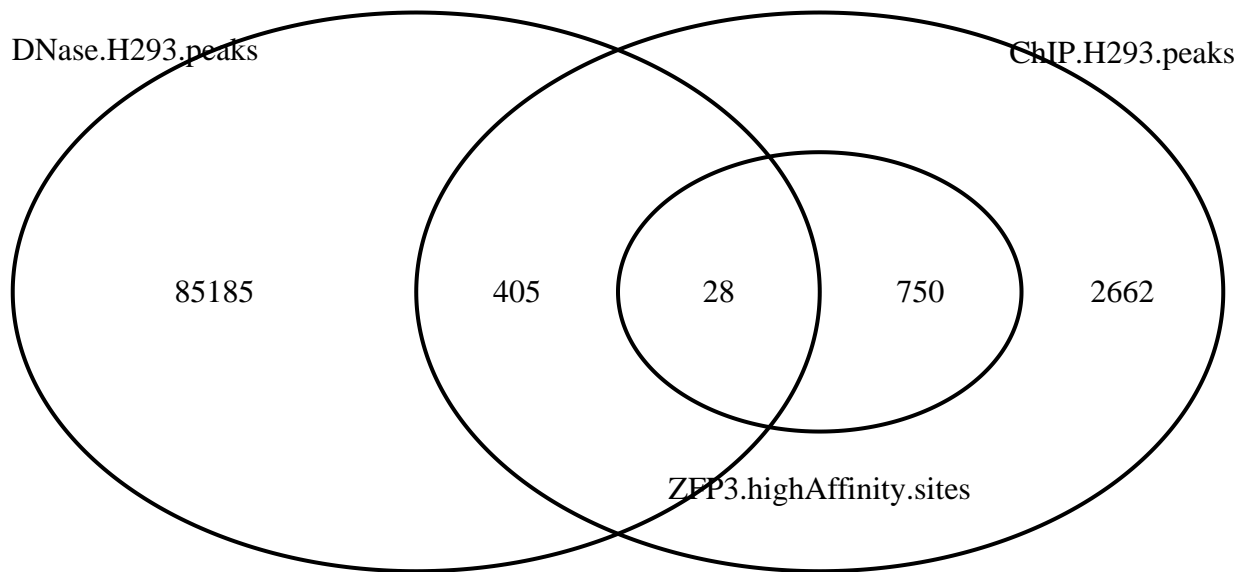
ZFP3.sites$predicted.Core.Energy <- ZFP3.sites$predicted.Energy
ZFP3.sites$predicted.Energy      <- NULL
ZFP3.sites$predicted.Upstream.Energy <- TFCookbook::predictEnergy(ZFP3.sites$Sequence, ZFP3.Upstream.PEM)
ZFP3.sites$predicted.Full.Energy  <- ZFP3.sites$predicted.Upstream.Energy + ZFP3.sites$predicted.Core.Energy

ZFP3.highAffinity.sites <- subset(ZFP3.sites, predicted.Upstream.Energy < (-4))

DNase.H293.peaks <- read.table("DNase-seq signals/ENCFF2850XK.H293T.bed") %>%
  GenomicRanges::makeGRangesFromDataFrame(seqnames.field = "V1",
                                           start.field    = "V2",
                                           end.field      = "V3") %>% unique()

ChIPpeakAnno::findOverlapsOfPeaks(ZFP3.highAffinity.sites,
```

```
ChIP.H293.peaks,
DNase.H293.peaks,
minoverlap = 0.85) %>%
ChIPpeakAnno::makeVennDiagram()
```



```
## $p.value
##      ZFP3.highAffinity.sites ChIP.H293.peaks DNase.H293.peaks pval
## [1,]                0                1                1      1
## [2,]                1                0                1      1
## [3,]                1                1                0      0
##
## $vennCounts
##      ZFP3.highAffinity.sites ChIP.H293.peaks DNase.H293.peaks Counts
## [1,]                0                0                0      0
## [2,]                0                0                1 85185
## [3,]                0                1                0 2662
## [4,]                0                1                1  405
## [5,]                1                0                0      0
## [6,]                1                0                1      0
## [7,]                1                1                0  750
## [8,]                1                1                1   28
##      count.ZFP3.highAffinity.sites count.ChIP.H293.peaks count.DNase.H293.peaks
## [1,]                0                0                0
## [2,]                0                0                85185
## [3,]                0                2662                0
```

```
## [4,] 0 406 409
## [5,] 0 0 0
## [6,] 0 0 0
## [7,] 814 757 0
## [8,] 28 29 30
## attr(,"class")
## [1] "VennCounts"
```

SK-N-SH cells data

```
ChIP.SKN.peaks <- read.table("ENCFF049VST.bed") %>%
  GenomicRanges::makeGRangesFromDataFrame(seqnames.field = "V1",
                                           start.field   = "V2",
                                           end.field     = "V3") %>% unique()

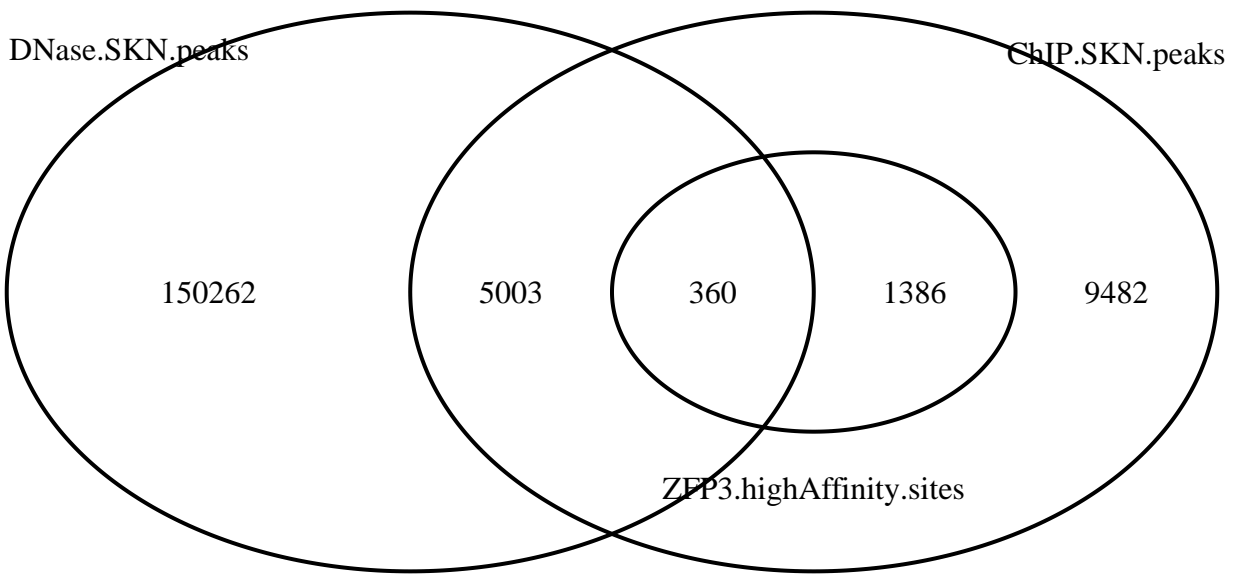
ZFP3.sites <- TFCookbook::matchPEM(PEM = ZFP3.Core.PEM,
                                   subject = ChIP.SKN.peaks,
                                   genome  = "hg38",
                                   out     = "positions",
                                   E.cutoff= -5)

ZFP3.sites$predicted.Core.Energy <- ZFP3.sites$predicted.Energy
ZFP3.sites$predicted.Energy      <- NULL
ZFP3.sites$predicted.Upstream.Energy <- TFCookbook::predictEnergy(ZFP3.sites$Sequence, ZFP3.Upstream.PEM)
ZFP3.sites$predicted.Full.Energy  <- ZFP3.sites$predicted.Upstream.Energy + ZFP3.sites$predicted.Core.Energy

ZFP3.highAffinity.sites <- subset(ZFP3.sites, predicted.Upstream.Energy < (-4))

DNase.SKN.peaks <- read.table("DNase-seq signals/ENCFF7520ZB.SKN.bed") %>%
  GenomicRanges::makeGRangesFromDataFrame(seqnames.field = "V1",
                                           start.field   = "V2",
                                           end.field     = "V3") %>% unique()

ChIPpeakAnno::findOverlapsOfPeaks(ZFP3.highAffinity.sites,
                                   ChIP.SKN.peaks,
                                   DNase.SKN.peaks,
                                   minoverlap = 0.85) %>%
ChIPpeakAnno::makeVennDiagram()
```



```
## $p.value
##      ZFP3.highAffinity.sites ChIP.SKN.peaks DNase.SKN.peaks pval
## [1,]                0                1                1      1
## [2,]                1                0                1      1
## [3,]                1                1                0      0
##
## $vennCounts
##      ZFP3.highAffinity.sites ChIP.SKN.peaks DNase.SKN.peaks Counts
## [1,]                0                0                0      0
## [2,]                0                0                1 150262
## [3,]                0                1                0   9482
## [4,]                0                1                1   5003
## [5,]                1                0                0      0
## [6,]                1                0                1      0
## [7,]                1                1                0   1386
## [8,]                1                1                1    360
##      count.ZFP3.highAffinity.sites count.ChIP.SKN.peaks count.DNase.SKN.peaks
## [1,]                0                0                0
## [2,]                0                0                150262
## [3,]                0                9482                0
## [4,]                0                5119                5088
## [5,]                0                0                0
## [6,]                0                0                0
## [7,]               1479                1430                0
## [8,]                389                408                372
## attr(,"class")
```

```
## [1] "VennCounts"
```