

modERN Call

White Lab

02-11-2016

modERN fly ChIP-seq datasets

17 new since last report, 48/80 for FY.

194 lines complete data sets

- ab, **Abd-B**, achi, acj6^l, Antp-mimic, Atf-2, Atf3, Bab2, br, brk, Bteb2, btn, **cad**, Camta-mimic, cato, CG10462, CG10565, CG10631, CG11398, CG11762, CG11902, CG12236, CG12744, CG13624, CG14965, CG1602, CG1620, CG1647, CG16863, CG1792, CG18476, CG2120, CG32264, CG31627, CG3163, CG33213, CG3838, CG4402(aka CG34406), CG4282, CG4820, CG4854, CG5204, CG6792(aka Plzf), CG7786, CG8089, CG8319, CG8944, CG9305, CG9609, CG9876, CHES-1-like, chif*, chn, **cic**, cnc, corto, crebA-mimic, crc-mimic, crg-1, **crp**, cyc, da, **dac**, Dad, **Dfd**, Dif, Dip3, disco, dl*, dm, dpn-mimic, dsf, dsx, **EcR**, Eip75B-MiMIC, Eip78C, Eip93F, emc, ems, en, ERR, esn, E(spl)m3, Ets21C, Ets65A(ets3), Ets97D, E(var)3-9, **eve**, **exd**, ey, eyg, foxo, ftz-f1, fru-mimic, fu2, GATAd, gcm2, gzfz, grh, **grn**, gro, **h**, her, HLH54F, HLHm7, HmgD, Hmx-MiMIC, Hnf4, Hr38, Hr39, **Hr4**^l, Hr46, Hr51, **Hr78**, Hr83, Hr96, hsf, insv, jim, **jing**, Jra*, jumu-mimic, kn, lbe, lilli, **lola***, Lpt, luna-mimic, lz, Mad, maf-s, mam-mimic, Max, med, Mes4, Met, Mio, Mnt, mod(mdg4), myb, **N**, NC2alpha, NC2beta, Neu2, NK7.1, **nmo**, OdsH^l, ovo, p53-mimic, pb, pdm3, **pdp1**(&mimic), pho, Pif1A, Pif1B, pnt-MiMIC, psq^l, pum-mimic, Rel, repo, sage, sens, shn, side-mimic, sima, slou, slp2, Smox, Sox102F, Sox14, Sox15, Sry-delta, **Stat92E**, Su(H), Su(var)3-7, sv, svp, tai, **tio**, **tll**, toe, topi, trh, trl-mimic, tup, tx, **usp**, vfl, vri, Vsx2, woc, Zfh2, ZnT49B.

- **14 from modENCODE**

- 180 from modERN

* multiple isoforms run

! Multiple time-points collected

XX: verifying data with rerun.

Current fly ChIP-seq

MiMIC being expanded for ChIP-seq ([ADD1](#), CG16779, CG9727, *dsx*[†], *foxo*, *Snoo*)

87 tagged lines being expanded for ChIP-seq, 48 new lines from last report.

Target Stages:

Embryo: [Az2](#), *bcd*, *Beaf-32*, CG10543, [CG10654](#), CG11617, CG11723, CG12104, [CG12155](#)^{!!}, [CG12942](#), [CG13123](#), CG16815, [CG13775](#), [CG15073](#), CG15479(aka Mabi), CG15601, CG16629, [CG17568](#), CG1832, CG18011, [CG18764](#), [CG30403](#), [CG3065](#), [CG31388](#), [CG4617](#), [CG4424](#), [CG5245](#), CG6683, CG7556, CG8388, CG9727, CG9817, [CG9883](#)^{!!}, *Chrac-16*, *Creb17-A*, *dmrt93B*, *dpn*, *E(bx)*, [E\(spl\)m-beta](#), [E\(spl\)m-gamma](#), *fd96Cb*, [Fer1](#), *fkh*, *HmgZ*, *ind*, [kay](#)^{!!}, *Kr*^{!!}, *meics*, *oc*, *odd*, *org-1*, *pita*, [Six4](#), *Sp1*, *su(Hw)*, *su(var)2-10-RH*, *TFAM*, [tin](#), *unpg*, *Usf*, *Vsx1*, [Xbp1](#), [YL-1](#).

W3L: CG7839, [HLH106](#), [net](#), [rgr](#).

WPP: *bsh*, *caup*^{!!}, CG18619, CG32006, CG30431, CG33017, CG4318, [CG5245](#), [CG7368](#), CG9139, [E\(z\)](#), [fd85E](#), *Nfl*, [salr](#).

AM: CG33017, [CG3919](#), CG7045, *Clk*.

AF: CG30403, [tj](#).

Repeat: *Abd-B*(for *sue*), *cic*, [crp](#), [dac](#), *grn*, *insv*, *jing*, *nmo-small*, *tio*

[†]: Collect at different time-point if fail

^{!!}: failed once

[07](#): Chromatin extracted

[05](#): IPed

[30](#): Librariied

17: @ HGAC

00: Processing

[04](#): awaiting reps or PCR

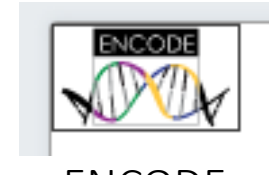
DCC

- Most current datasets for dm3, ce10 and ce11, have been uploaded
 - > 6k files.
- Currently reprocessing data with dm6
- Look into a threshold for RSC/NSC values
- Finish uploading Ab comparisons for validation

DCC

- Items from Last weeks call:

- Icons added to pages.
 - Do we want one?



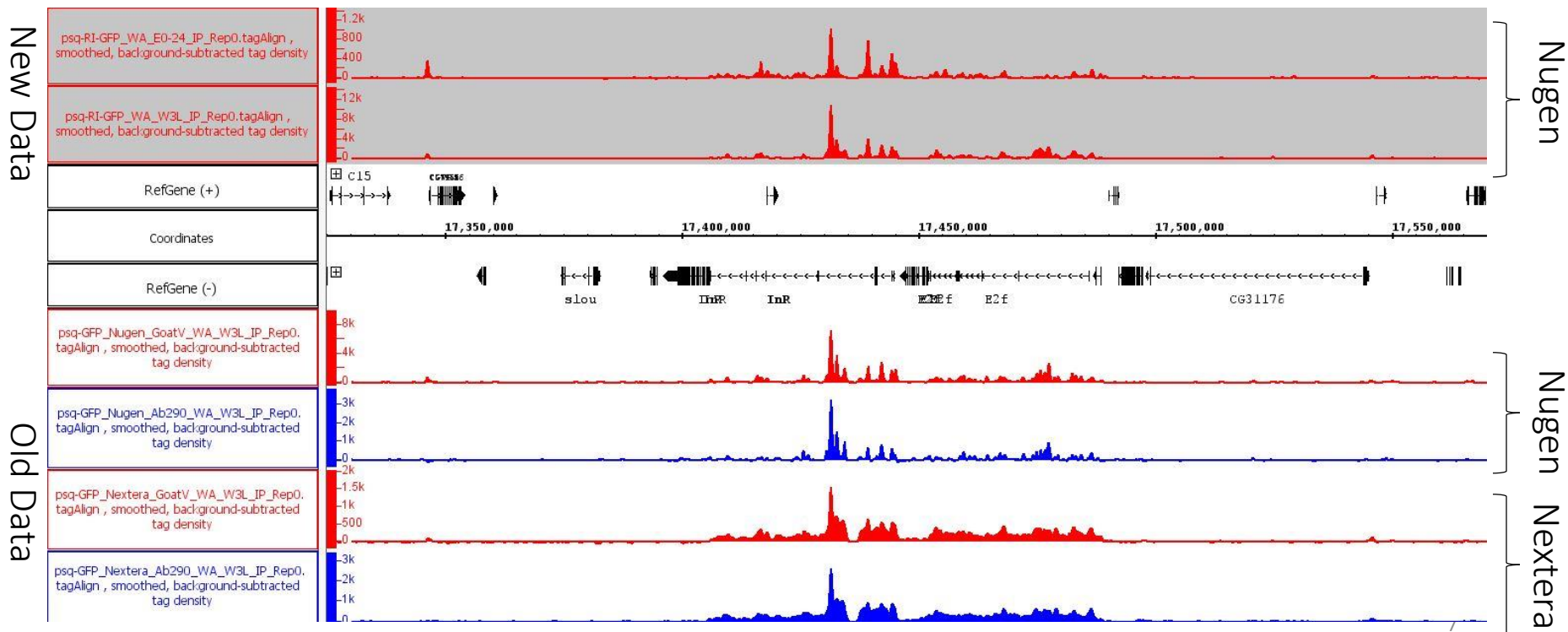
- Anyone attending TAGC?
 - Esther willing to present poster describing available data
- ENCODE epitope tagging standards:
 - DCC wants official answer if we'll comply with ENCODE line validation.
 - Would require IP/WB of each line.

Control Experiments

- Precipitated by increase in overlapped peaks in new datasets
- Possible causes:
 - Library kit effect
 - GoatV effect
 - Comparison of TF antibody vs GFP antibody
 - Background peaks

Library kit and GoatV controls

- Not caused by Library or goatV
 - Ab290 (blue tracks) show similar traces to goatV (red tracks)

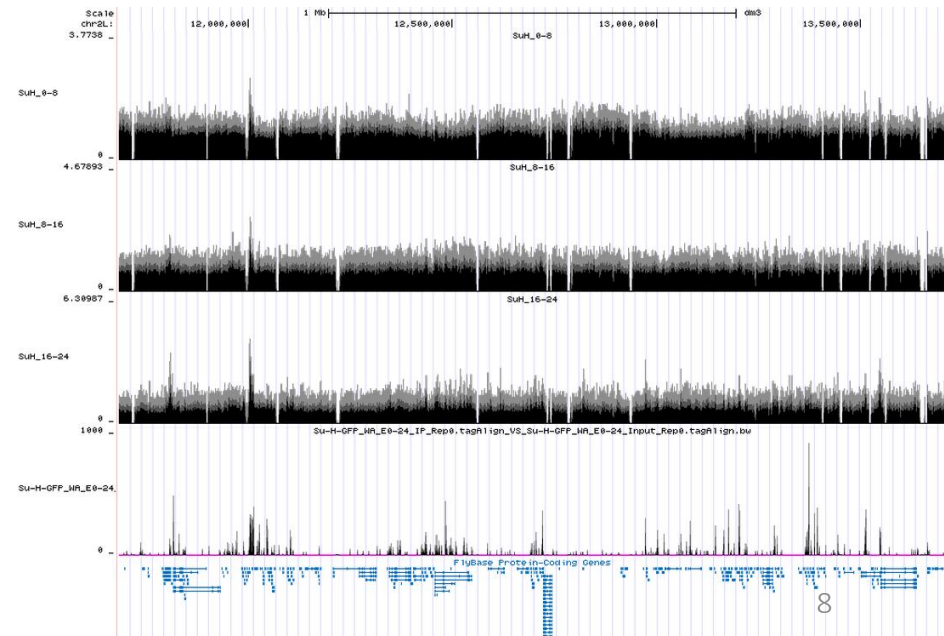
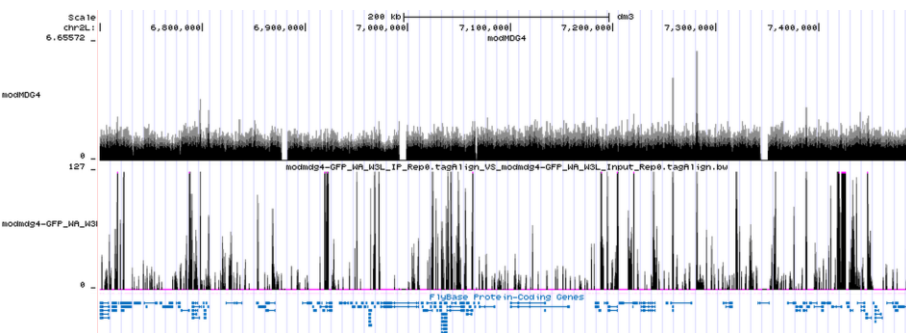


TF antibody vs GFP antibody

Difficult to tell with the small number of datasets with same factor and time point

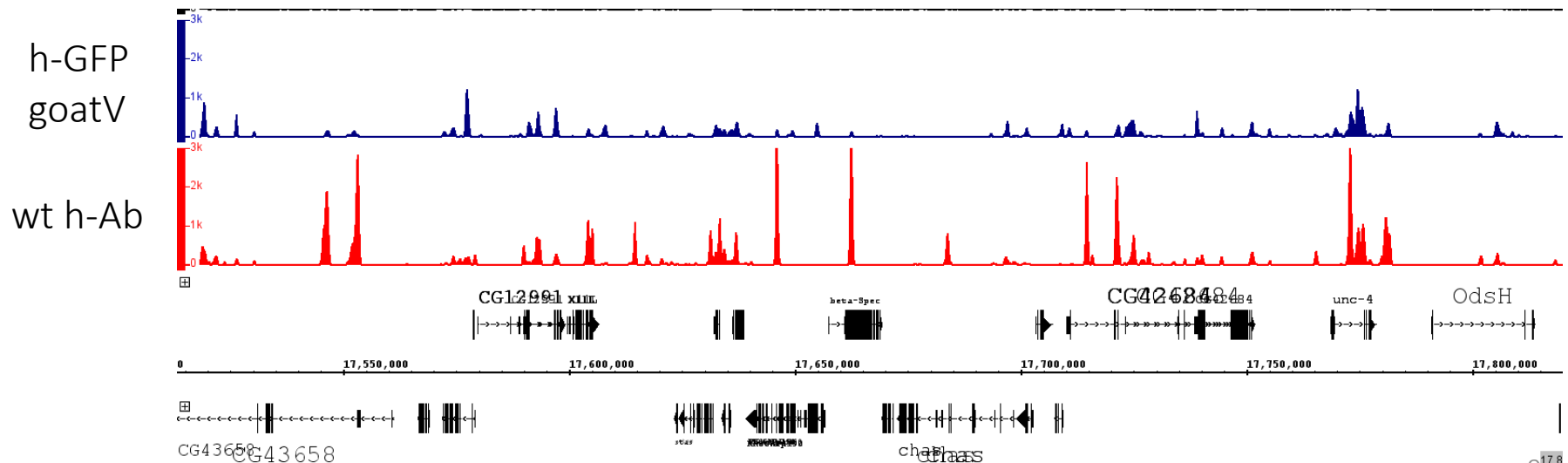
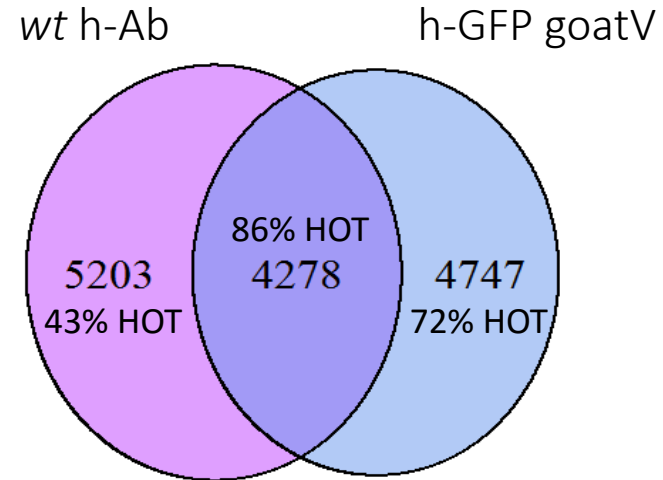
- mod(MDG4)
 - GFP: 5580
 - Iso1: 235
 - 207 in common.

Su(H)
GFP: 3762
Iso1: 347
114 in common.



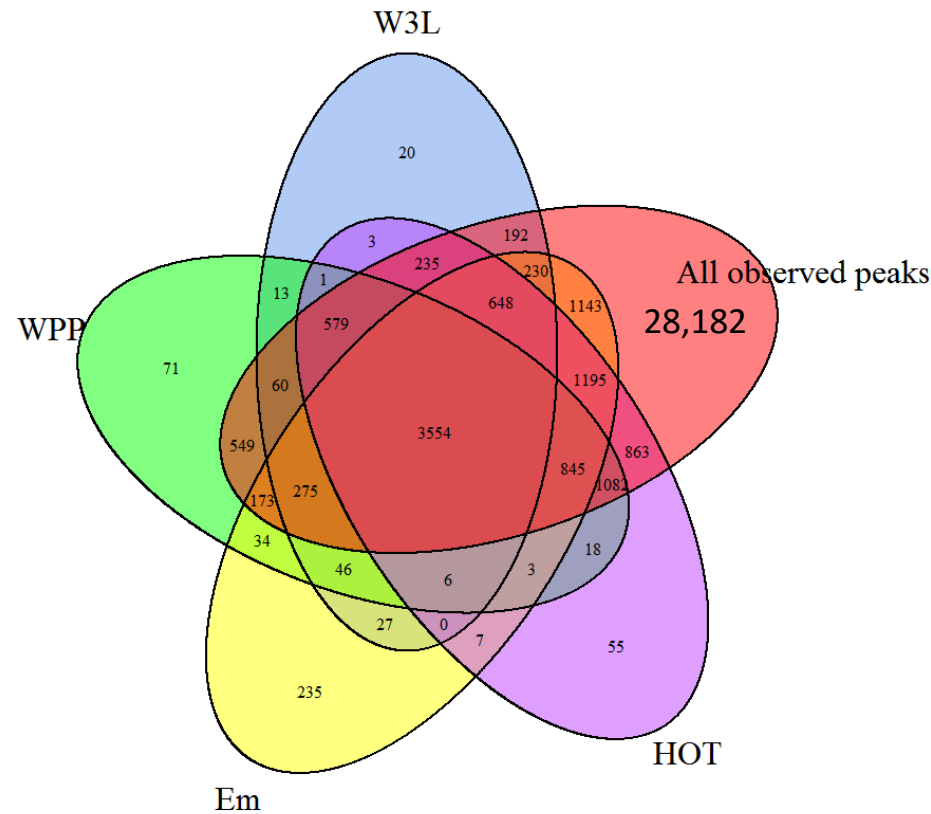
TF antibody vs GFP antibody

- Using current protocol, repeated experiment with TF Ab used for modENCODE
 - ~45% overlap.
 - Only 548 of overlapped peaks are not HOT regions, tend to be smaller peaks



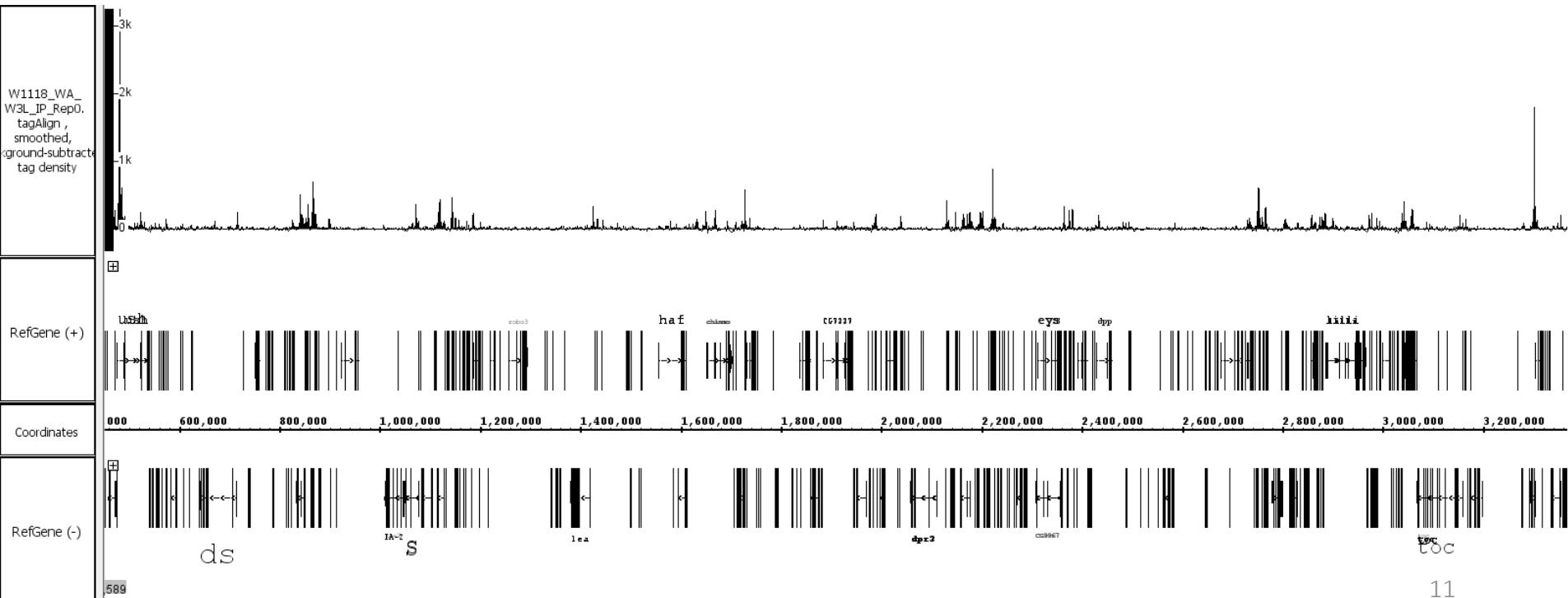
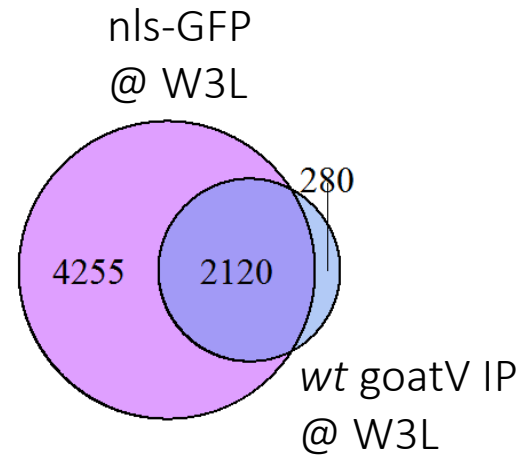
Background Peaks

- Over-expression of nls-GFP yielded ChIP-seq peaks
 - Most were initially called as HOTspots
- Unclear if this was due to overexpression of GFP



Background Peaks

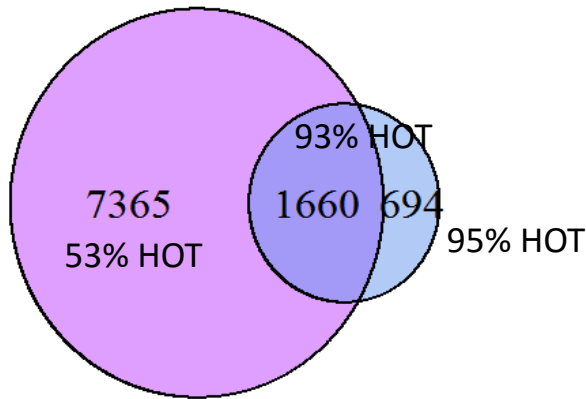
- IP using goatV on *wt Drosophila*
 - 2400 peaks called.
 - Fewer than nls-GFP
 - 97% peaks are HOTspots



Background Peaks

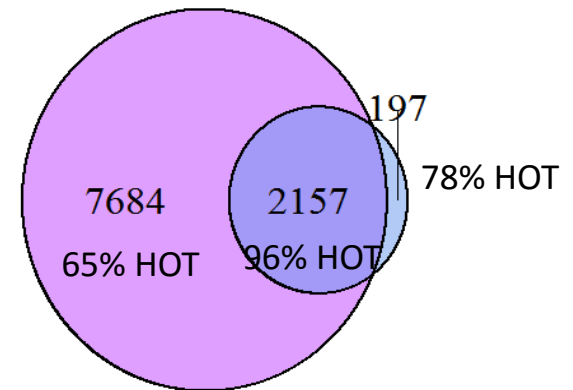
- GFP background peaks seen in non-GFP datasets
 - not really GFP background?

wt h-Ab vs *wt goatV*



wt h-Ab 60% HOT

h-GFP goatV vs *wt goatV*



h-GFP goatV 74% HOT

Future work

- Repeat nls-GFP and *wt-goatV* tests again
 - Will the same peaks be observed?
- Are the *wt-goatV* peaks a subset of the nls-GFP peaks?
- Are these regions the most intense when seen in TF datasets?
- Can “hyper-ChIP-able” regions be subtracted from data prior to peak calling?
- Could check background with another Ab, eg aFLAG
- Test a few more TF-Ab vs GoatV.
 - Both comparing to old data and repeating with TF Ab.