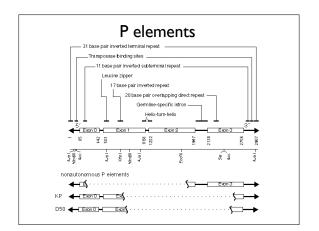
## P elements

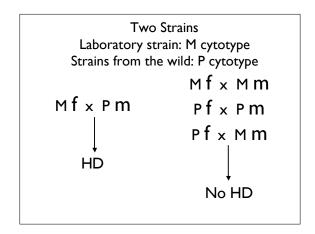
Problem Set 4 for this and the last lecture.

Quiz coming up next week: covers material up to today--7 lectures (includes lecture Chromosomes 4 from Sept. 12)



## Hybrid Dysgenesis

- I. Heat sensitive sterility
- 2. Increased rates of mutagenesis-unstable alleles
- 3. Increased rates of chromosomal rearrangements
- 4. All effects in the germline, not in somatic tissues



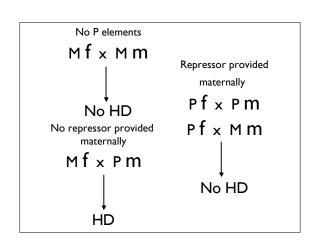
M strains have no P elements.

P strains have many P elements.

P elements produce an inhibitor of transposition, which is inherited maternally in germline.

Therefore, inhibition of transposition a maternal effect.

All somatic cells produce an inhibitor.



Rubin and colleagues showed that HD caused by a transposable element.

Rationale: white gene had been cloned, so induced HD white mutations and looked for insertions in white gene.

Mf x Pm

HD m x Basc/Basc f

contains M X chromosome no P elements an inversion, Bar and a w allele.

Screened for white-eyed flies w\*/Basc, where w\* is new HD allele.

### Isolated seven new white alleles

- I. All seven white mutants spontaneously reverted to red.
- 2. All seven contained insertions that the wild-type *white* gene lacked.
- 3. All of the insertions were related (Ac like).
- 4. Many elements present in P strains, but none in the M strains
- 5. The largest element in the genome was 2.9 kbp.

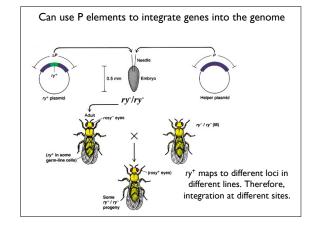
# Does the 2.9 kb element contain transposase activity? Inject plasmid containing 2.9 kb P element snw/Y

Singed bristle assay:  $sn^w$  is a weak allele of singed.  $sn^w$  has two P elements in the singed promoter. When one is excised, get a more extreme phenotype  $(sn^e)$ , When the other is excised, get wild-type  $(sn^+)$ .

Injected  $sn^w/Y \ m \times sn^-/sn^- f$ 

 $sn^w/sn^-$  females Also get wild type  $(sn^+/sn^-)$  and more extreme phenotype  $(sn^e/sn^-)$ 

2.9 kb element has transposase activity.



# Uses of P elements

- I. New alleles of gene
- 2. Insert genes into genome e.g., test whether cloned correct gene by mutant rescue.
- 3. Genetic mapping
- 4. Reverse genetics
- 5. Enhancer trapping
- 6. Missexpression of genes.

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# The bipartite GAL4-UAS system



- GAL4 is a transcriptional activator from yeast that recognizes a DNA sequence called the UAS (upstream activating sequence)
- We can use this to control expression of YFG in a tissue specific manner by using enhancer elements specific for the tissue we are interested in

