

Mitotic recombination

Genetics 371B Lecture 19

2 Nov. 1999

Rare relative to meiotic recombination

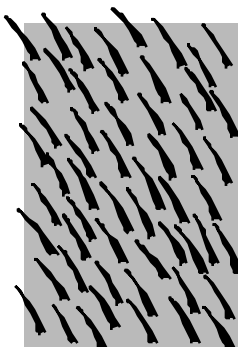
Discovery: Curt Stern, 1936

Linked genes **singed bristles** and **yellow body**

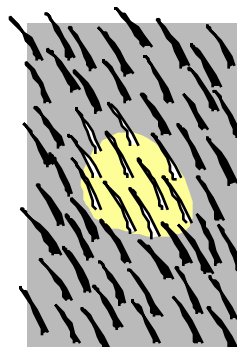
$\frac{+}{sn} \frac{y}{+}$ double heterozygote in trans configuration

Exercise: Design an experiment to confirm the trans configuration

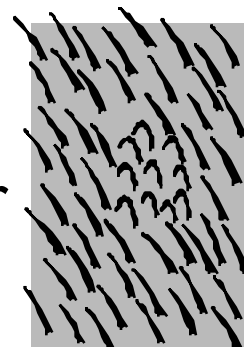
Normal



Occasionally:

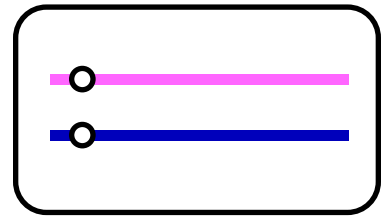
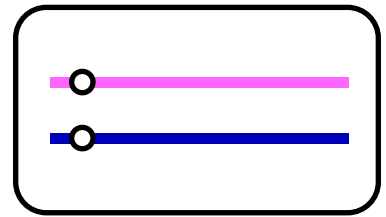
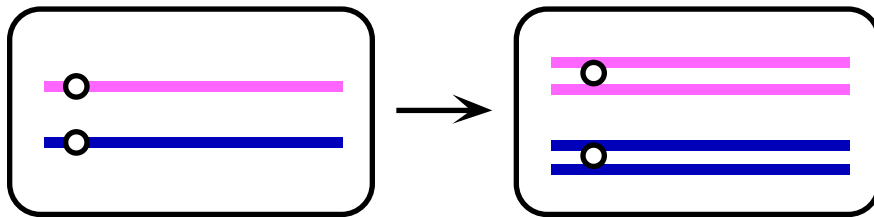


or

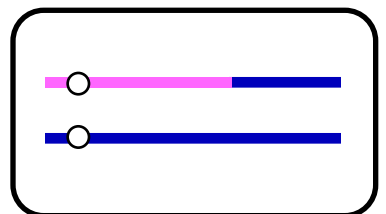
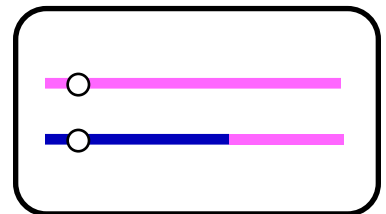
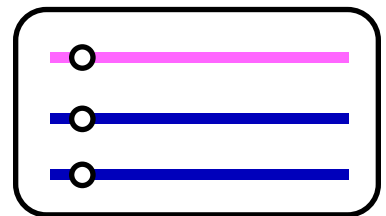
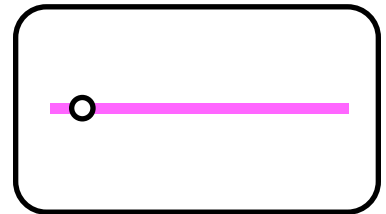
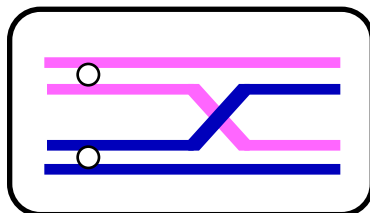
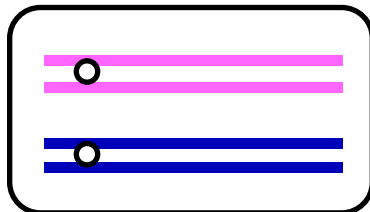
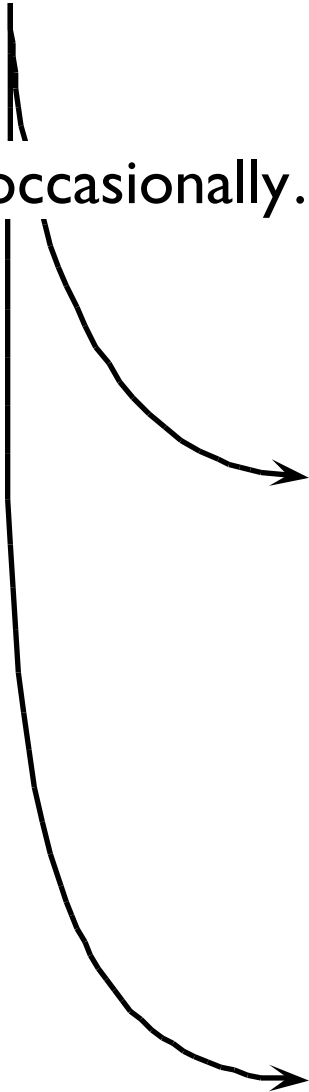


Stern's explanation

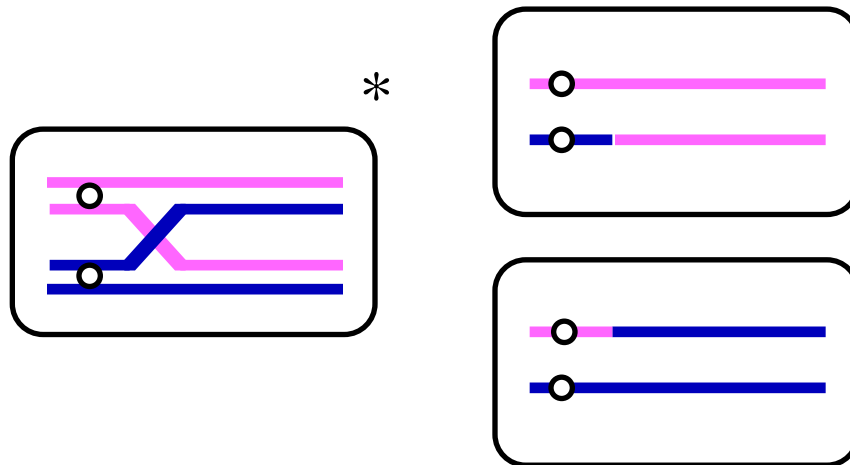
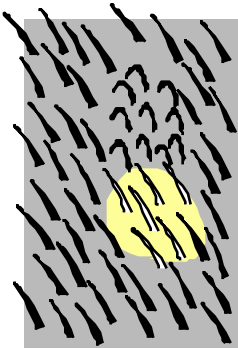
◆ Normal mitosis



◆ but occasionally...



Rare twin spots:



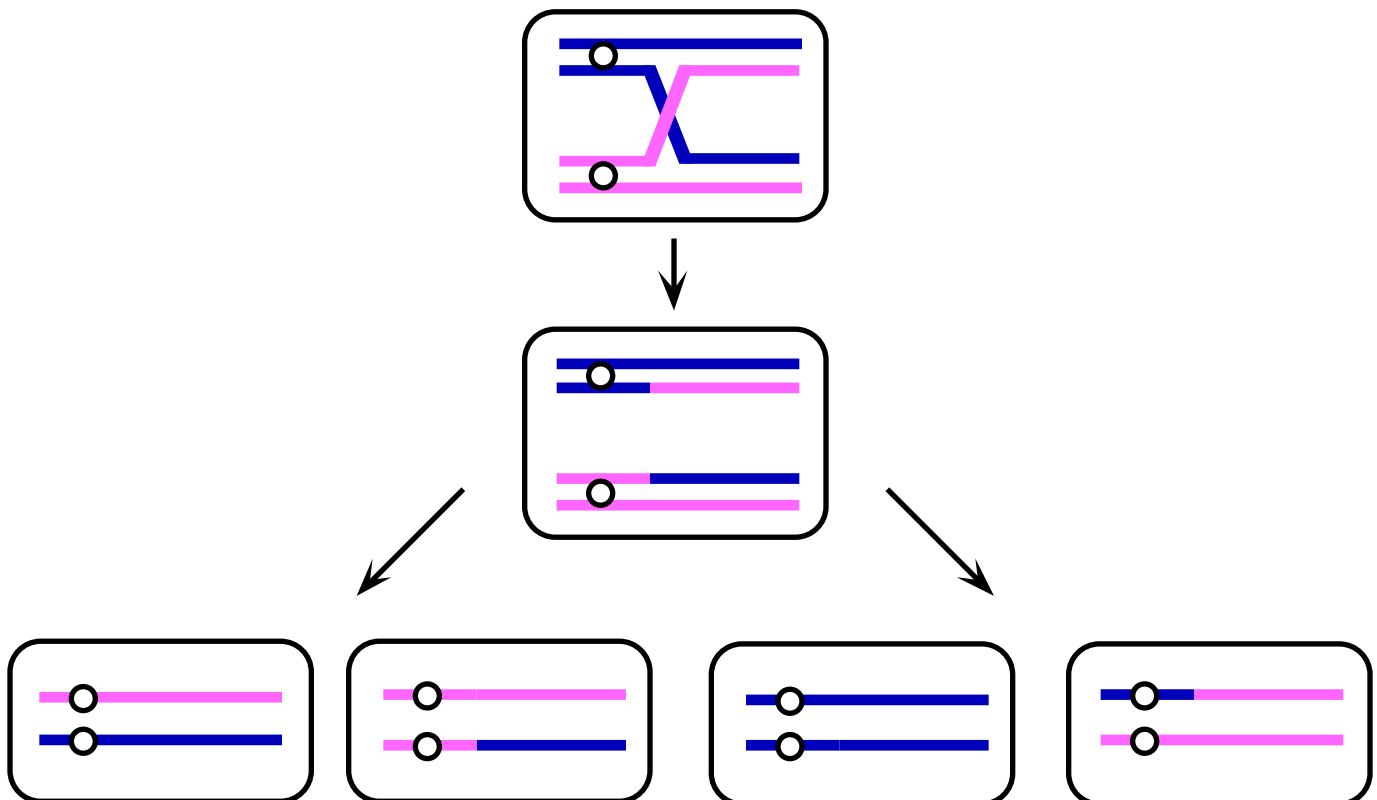
Exercise: This cell is shown to be undergoing mitotic recombination after completion of S phase (how can we tell from the diagram)? How can you tell from the products of the division that the recombination did indeed occur post-S phase?

Significance for human health?

Suppose we're talking about a recessive disease allele...

“Loss of heterozygosity”

e.g., Retinoblastoma, Wilms tumor



Sporadic cases—

Inherited form—

Explanation?

“2-hit kinetics”

$Rb^+/Rb^+ \longrightarrow Rb^+/rb \longrightarrow rb/rb$

“1-hit kinetics”

$Rb^+/rb \longrightarrow rb/rb$

Applications

- ◆ Mapping – refrequency of "spots" proportional to map distance

- ◆ Mapping *centromeres* – can you get twin spots?



Caution: These are *mitotic recombination* frequencies!

- ◆ Studying development, recessive lethal alleles

- ◆ Assay for genotoxic agents – “SMART”