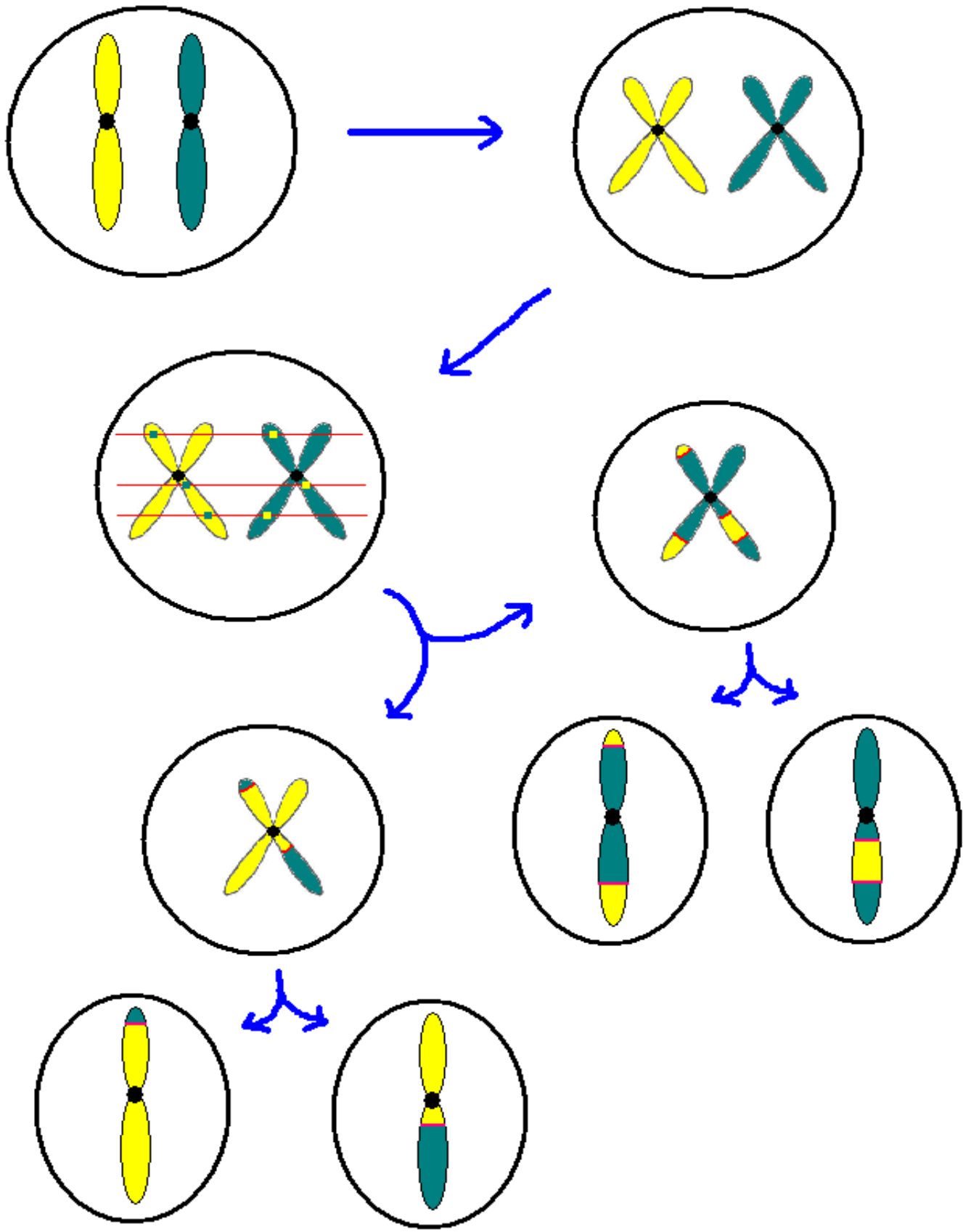


Human meiotic interference

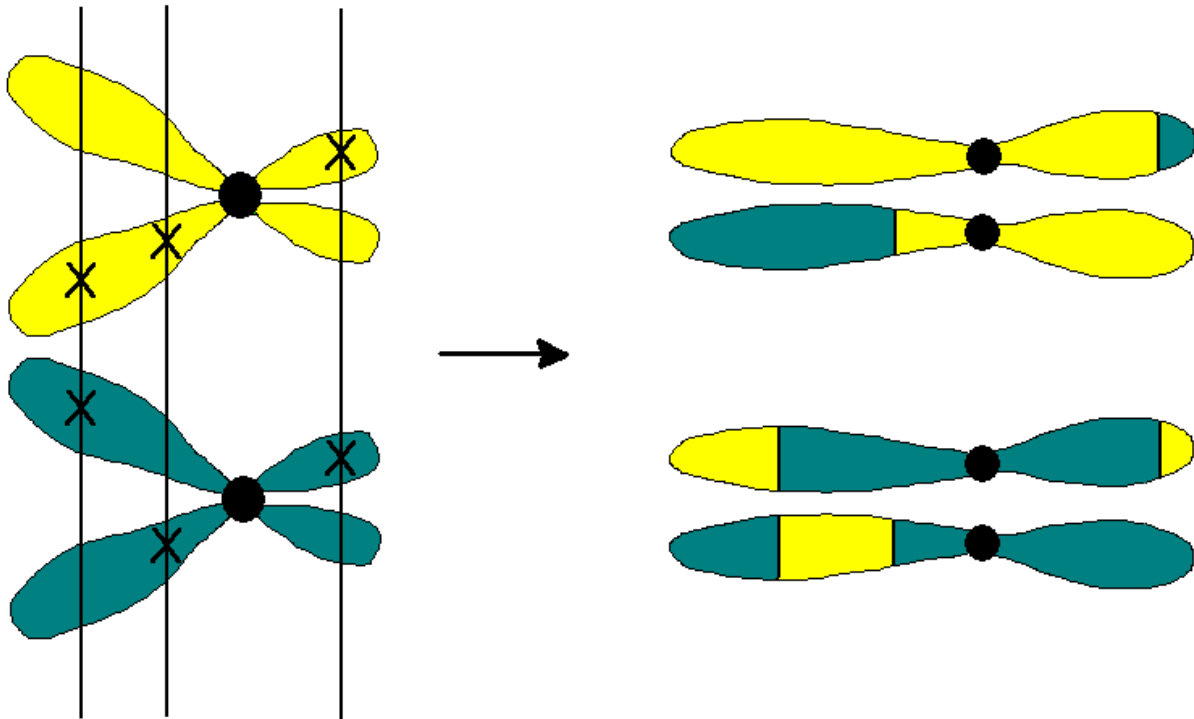
Karl W Broman
James L Weber



Meiosis



Interference



- Strand choice
→ Chromatid interference
- Spacing
→ Chiasma (crossover) interference

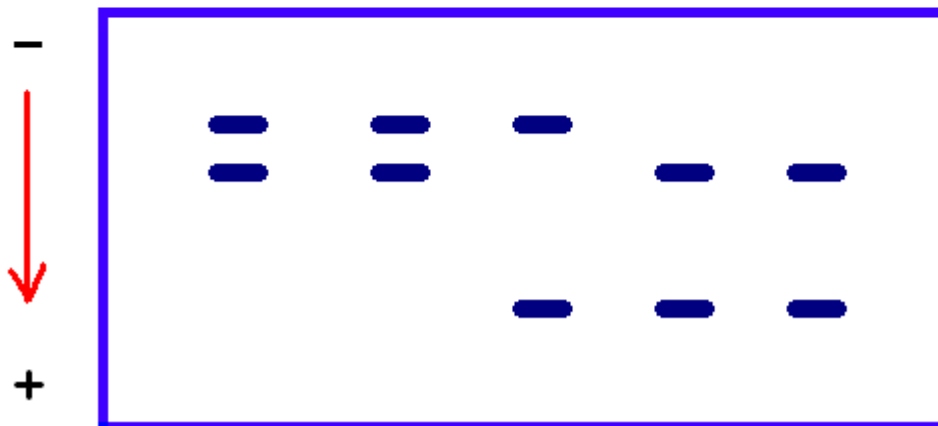
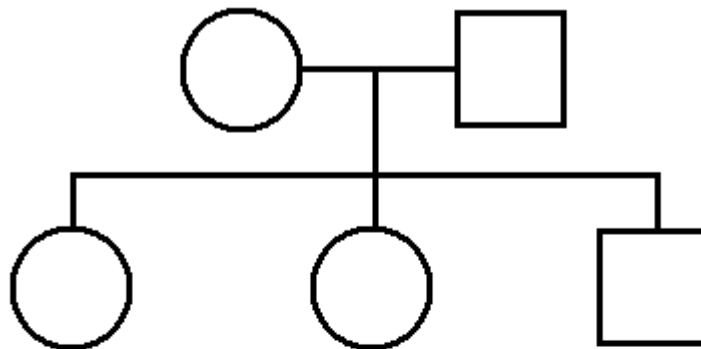
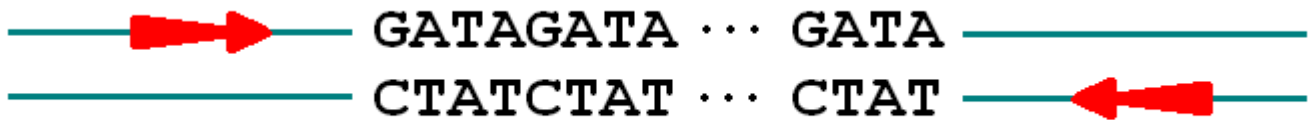
Why study interference?

- Estimate the probability of a double crossover in a small interval
- Obtain a model of meiosis for simulation and analysis
- Compare human meiosis to that of other organisms

Goals

- Demonstrate the presence of interference in human meiosis
- Obtain an empirical map function
- Find a good model

Genetic markers: STRPs or microsatellites



Model organisms

- Lots of meioses
- A few linked markers
- Look at frequency of rare multiple recombination events

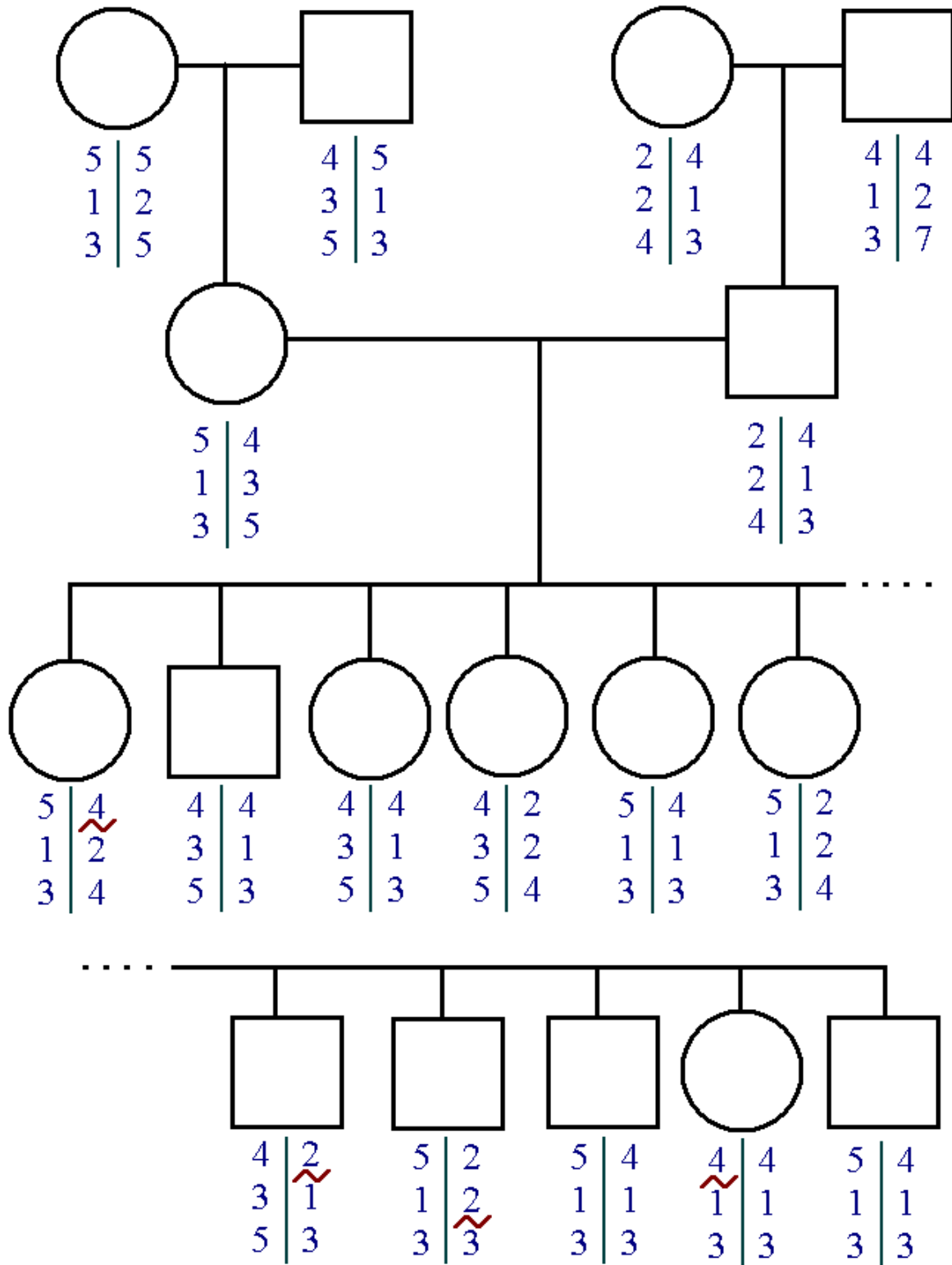
Drosophila data (Morgan et al 1935)

Event	Count	Event	Count
0000	10,431	1001	46
1000	771	0101	53
0100	1,579	0011	25
0010	1,221	1110	1
0001	1,994	1101	1
1100	4	1011	1
1010	7	0111	1
0110	4	1111	1

Human data

- www.marshmed.org/genetics
- 8 CEPH families
 - three generations
 - 11 to 15 progeny
 - 92 meioses
- ~8,000 STRP markers
 - 90 ± 7 % typed
- Average spacing
 - female: 0.6 ± 1.2 cM
 - male: 0.4 ± 1.0 cM
 - sex-ave: 0.5 ± 0.9 cM
- Data cleaning
 - Removed 764/964,425 (~0.08%) genotypes resulting in tight double recombinants

CEPH pedigree



CRI-MAP chrompic output

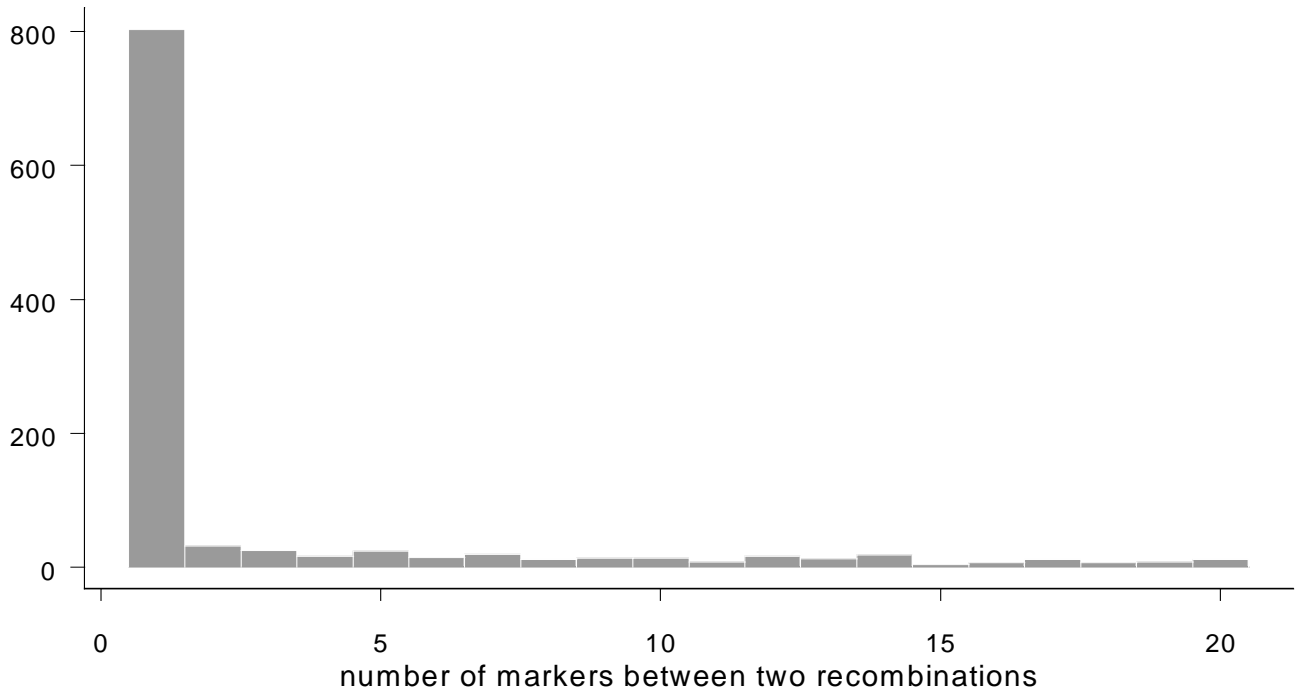
CEPH individual 1331-11

maternal chromosome 10

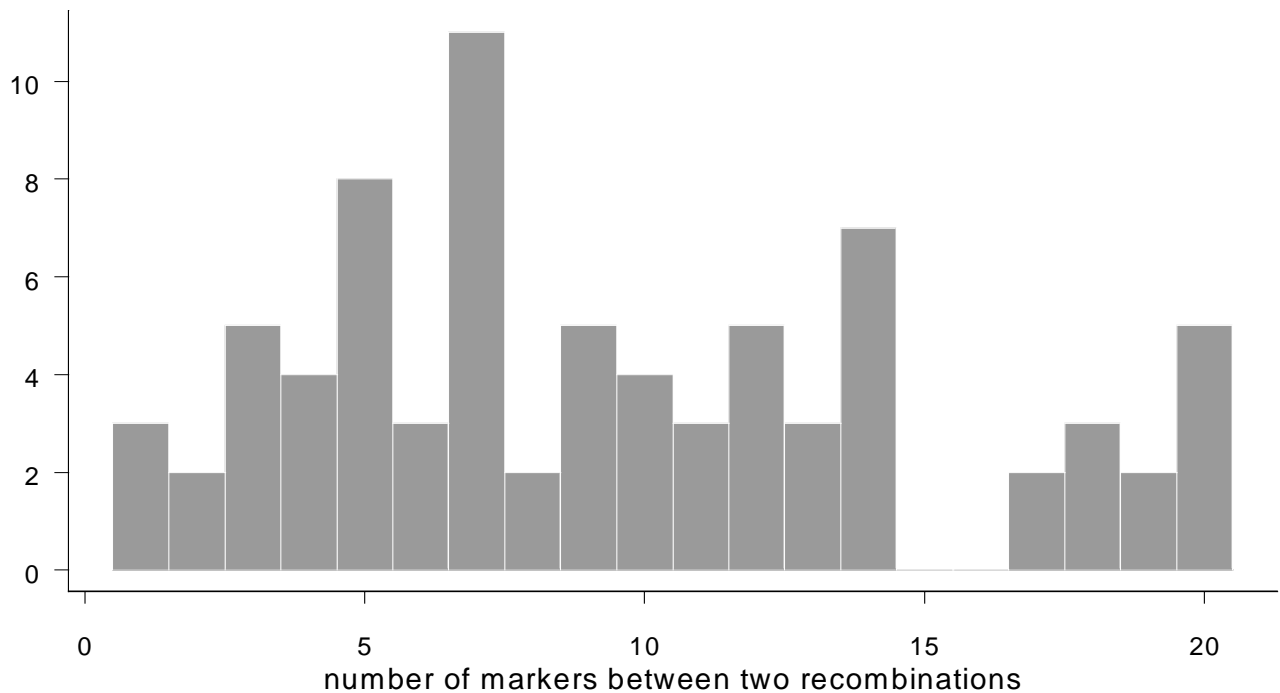
```
11111111--- 11-1111-11- --11--1111i
1-11---11- 111111--i- 11-1111-11
--111111-11 1111-11111 11--111111
111-1111i-i 1111111111 10000-0-00
0--o00-000 0000-00000 0000--0000
0000--0000 0000-00000 o00-0--0--
--0-11-11- -1111ii1i-1 ---1-i-1-i
1111-i--11 11111-11i1 -11i-11111
-1-----i111 1i11111-111 -11i1-111-
11-1111111i 111-i111i- 11111111-i-
11111111-1i 1i-111i11- 1i--1-11-1
111-1i-1-1 1-1----1-1 1i-1ii1i11
1i--1--1i- 11i11--111 11--1i1111i
1i1i-11111 i-0---0000 00000-000o
o0-00o
```

Left tail of the distribution of # markers between recombinations

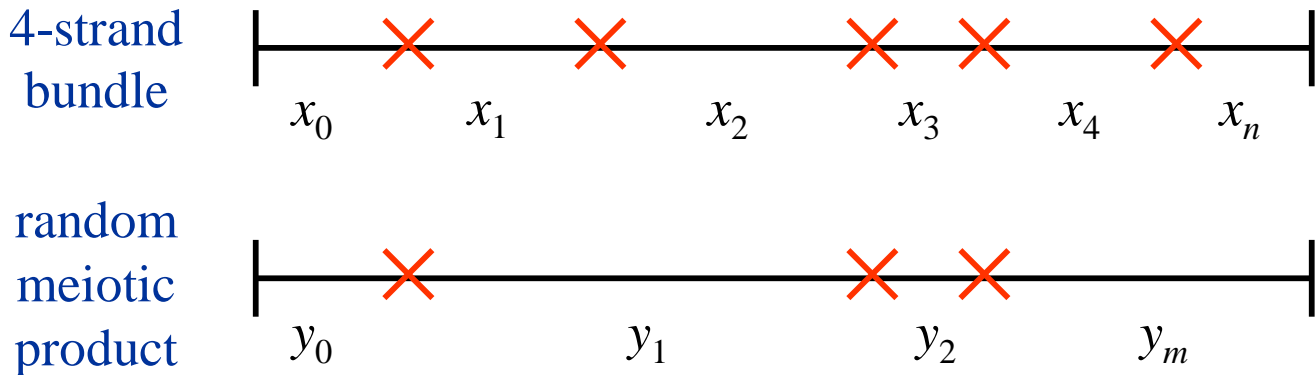
Raw Data



Clean Data



Models



- Count-location model

$$n \sim (p_0, p_1, p_2, \dots)$$

locations | $n \sim$ iid uniform

- Gamma model

x_i 's \sim stationary gamma renewal process
(shape = v , rate = $2v$)

y_i 's \sim mixtures of gammas

Genetic distance

distance (cM) = average # crossovers
in 100 meiotic products

per Morgan $\left\{ \begin{array}{l} 2 \text{ chiasmata on 4-strand bundle} \\ 1 \text{ crossover on meiotic product} \end{array} \right.$

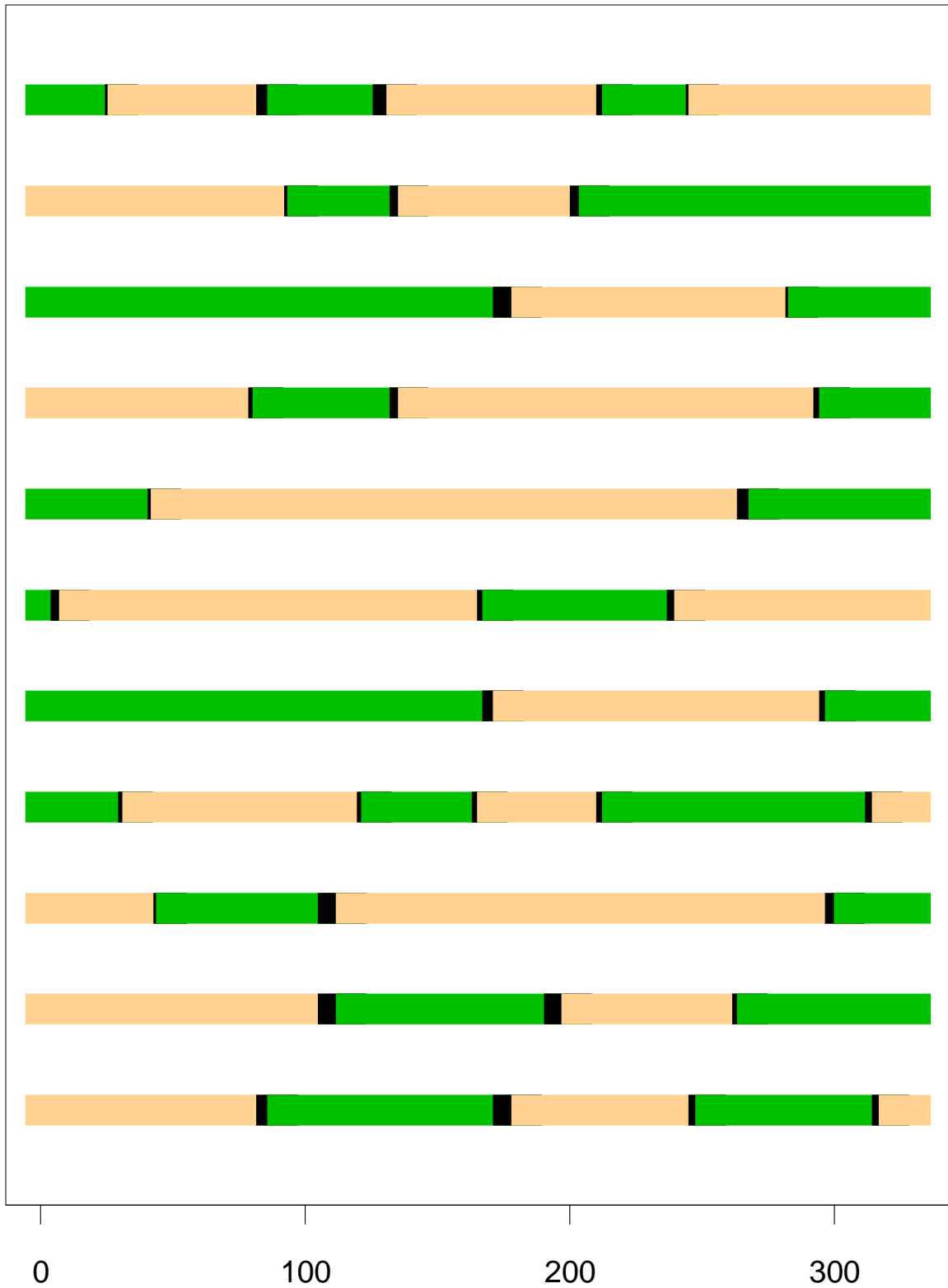
Map function

recombination fraction as a function of genetic distance

Haldane $r(d) = \frac{1}{2} [1 - \exp(-2d)]$

Kosambi $r(d) = \frac{1}{2} \tanh(2d)$

Another view of the data



Model fitting

- Count-location model

$m_i = \# \text{ crossovers}$

$n_i = \text{underlying } \# \text{ chiasmata}$

$n_i \sim (p_0, p_1, p_2, \dots)$

$m_i | n_i \sim \text{binomial}(n_i, 1/2)$

MLEs via a version of the EM algorithm

Model fitting

- Gamma model

$$x_1, x_2, \dots \sim f(v, 2v)$$

$$x_0 \sim g = 2[1 - F(v, 2v)]$$

x_i 's independent

$$y_1, y_2, \dots \sim \sum (1/2)^k f(kv, 2v)$$

$$y_0 \sim 1/2 g + \sum (1/2)^{(k+1)} g * f(kv, 2v)$$

y_i 's independent

- MLE of v using y_i 's
- g calculated numerically
- Convolutions calculated numerically
- Maximization performed using a quasi-Newton method

Distributions of # XOs / chr

Maternal chromosome 1

	0	1	2	3	4	5	> 5	X²
Obs.	2	7	12	24	23	14	10	
Pois.	3	9	17	20	17	12	14	9.2
C-L	2	7	14	22	23	16	9	0.8
Gamma	1	5	14	23	23	16	10	1.2

Maternal chromosome 4

	0	1	2	3	4	5	> 5	X²
Obs.	1	16	36	15	15	9	0	
Pois.	7	18	23	20	13	7	4	14.4
C-L	4	16	26	25	15	6	1	12.8
Gamma	4	15	26	24	15	6	1	7.1

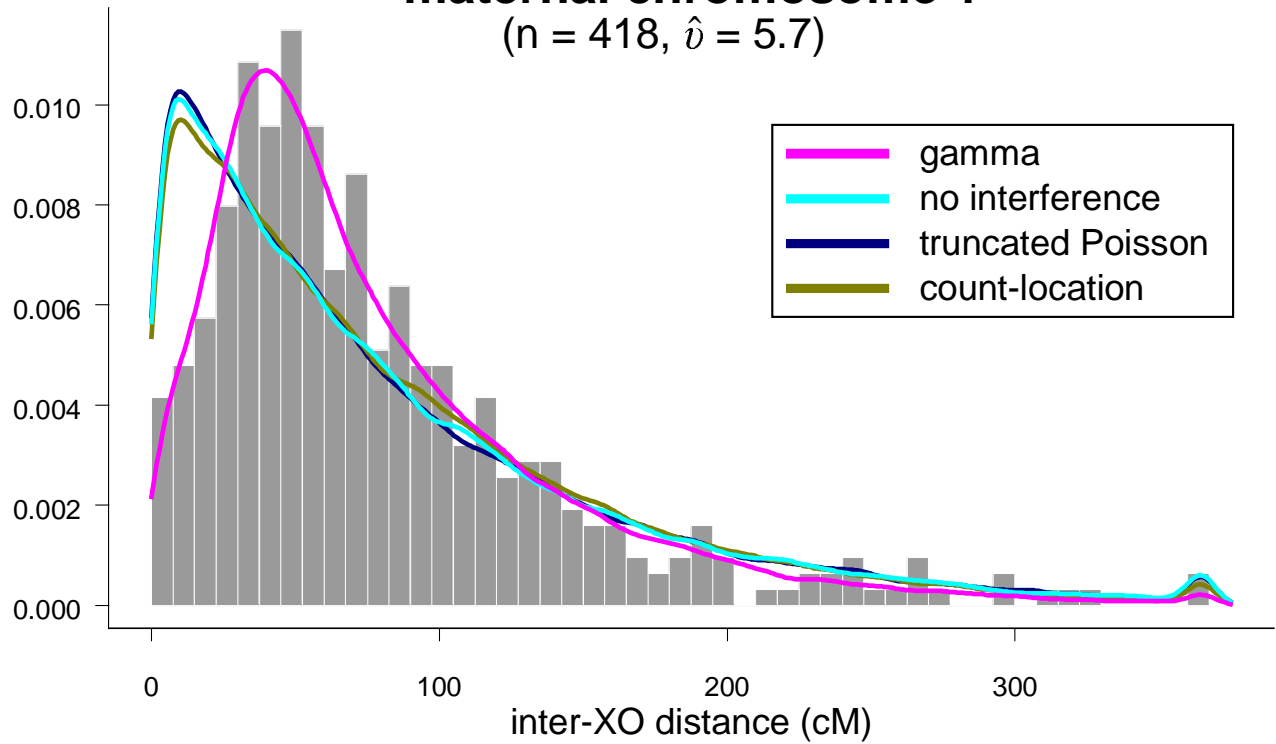
Evidence for interference:

maternal 3, 9, 12, 14, 15, 17

paternal 1, 4, 5, 9, 14

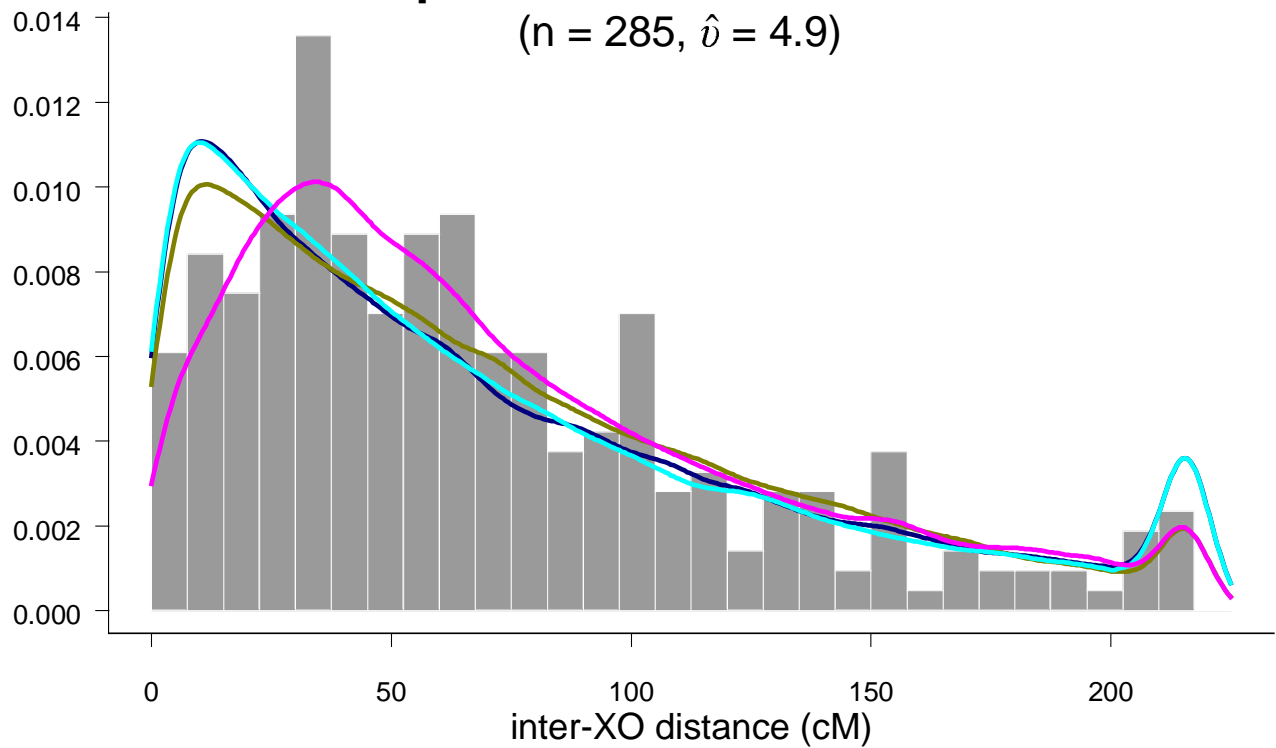
maternal chromosome 1

($n = 418, \hat{\nu} = 5.7$)



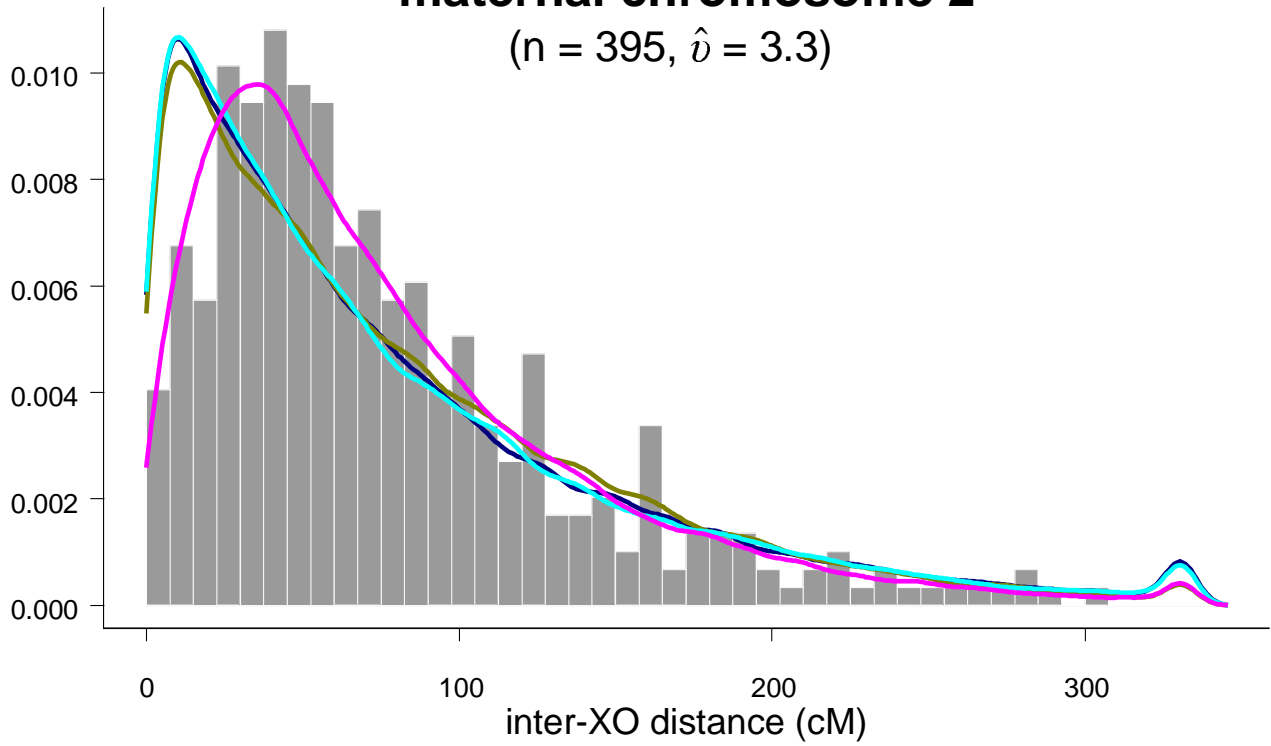
paternal chromosome 1

($n = 285, \hat{\nu} = 4.9$)



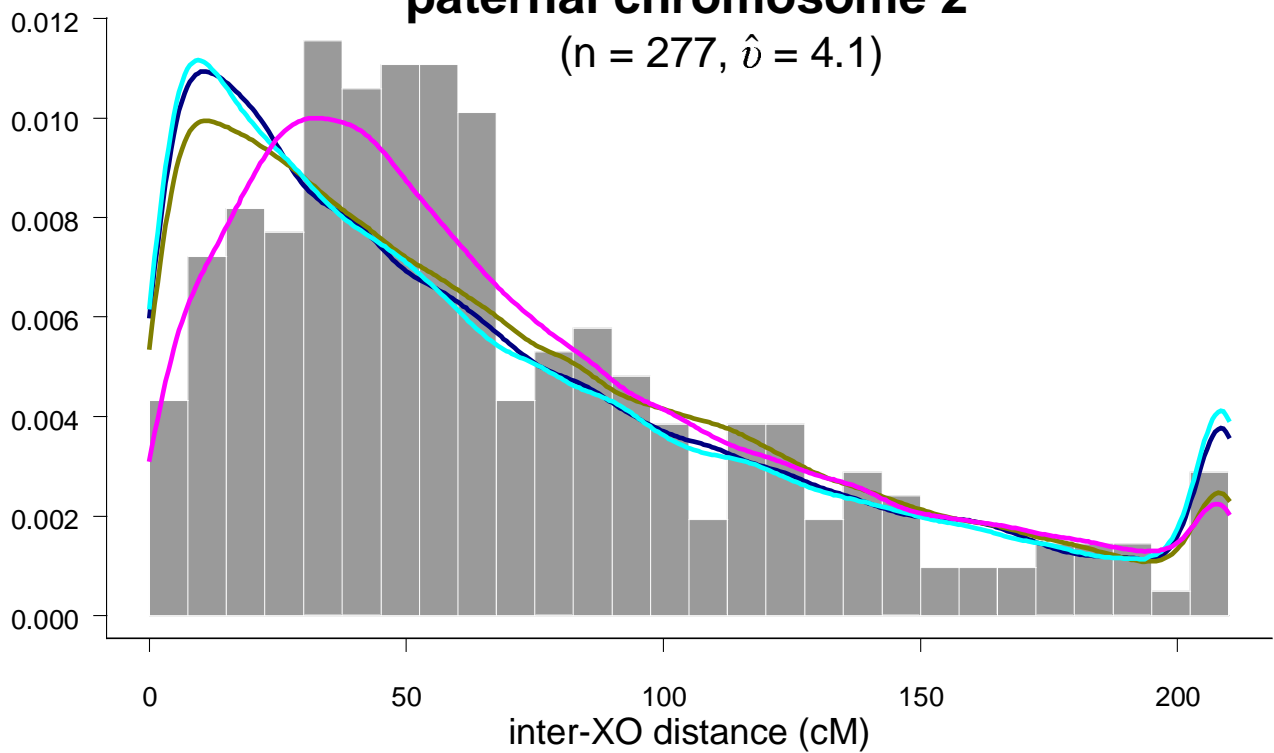
maternal chromosome 2

($n = 395, \hat{v} = 3.3$)



paternal chromosome 2

($n = 277, \hat{v} = 4.1$)



Discussion

- **Approximations**
 - Correct marker order
 - Correct genetic distances
 - All crossovers observed
 - Interval censoring unimportant
 - No individual variation in recombination
 - Interference constant across chromosome
- **Conclusions**
 - Gamma model fits well
 - Count-location model fits poorly
 - Gamma parameter, $\hat{v} \approx 3-5$
(stronger than Kosambi, $v \approx 2.6$)
- **Further work**
 - Interference across the centromere
 - Variation between chromosomes