

Human crossover interference

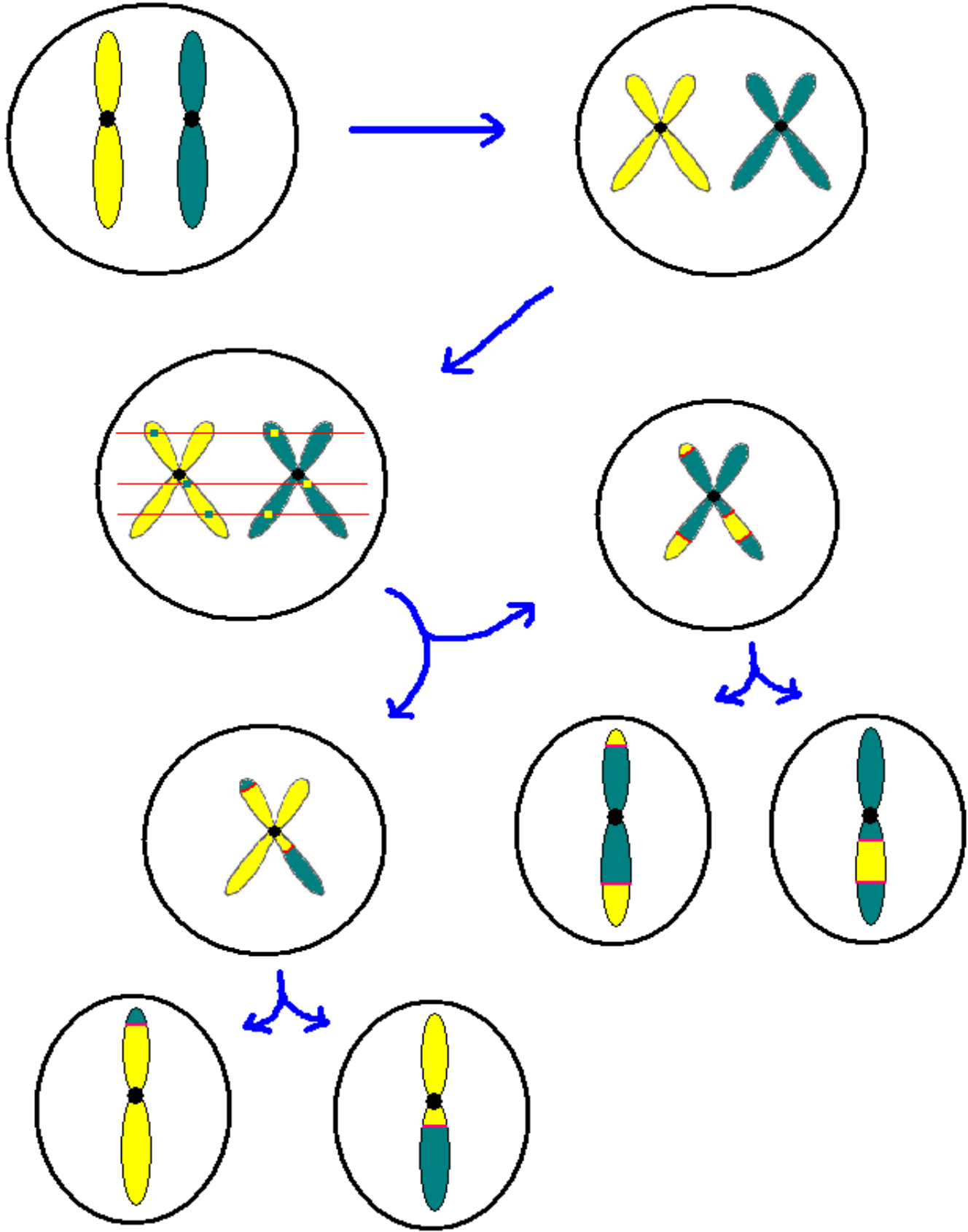
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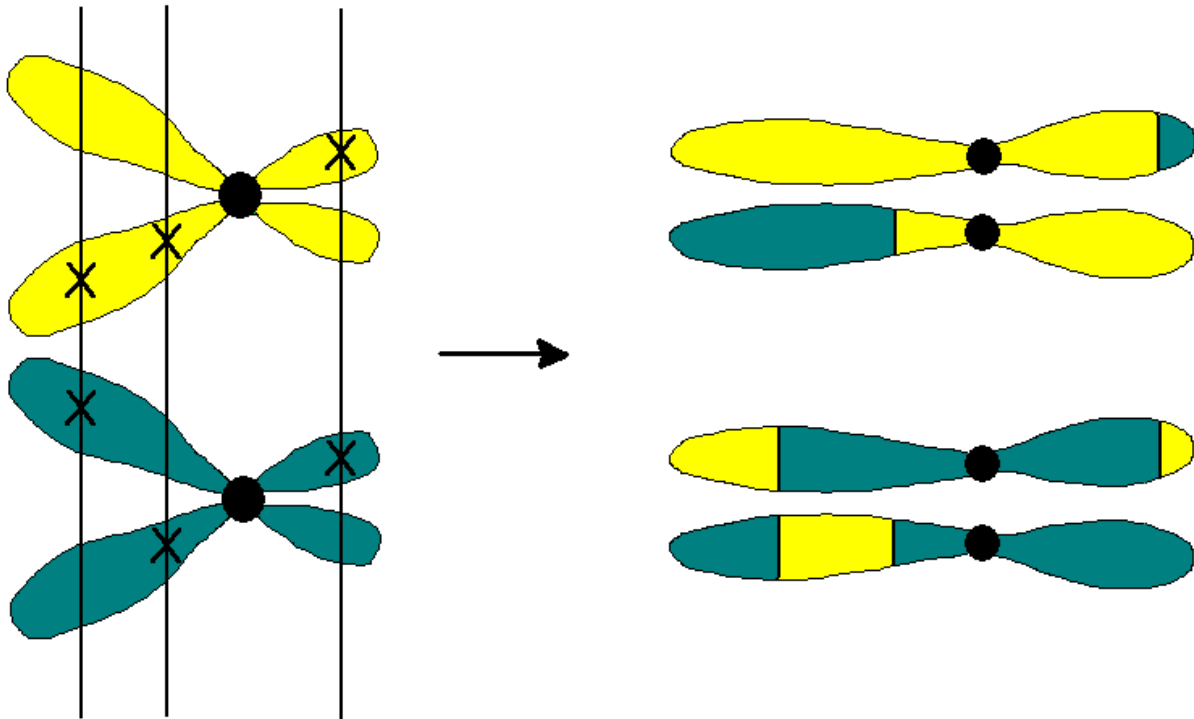
Joint work with James L. Weber,
Marshfield Medical Research Foundation

<http://kbroman.homepage.com>

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

Model organisms

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

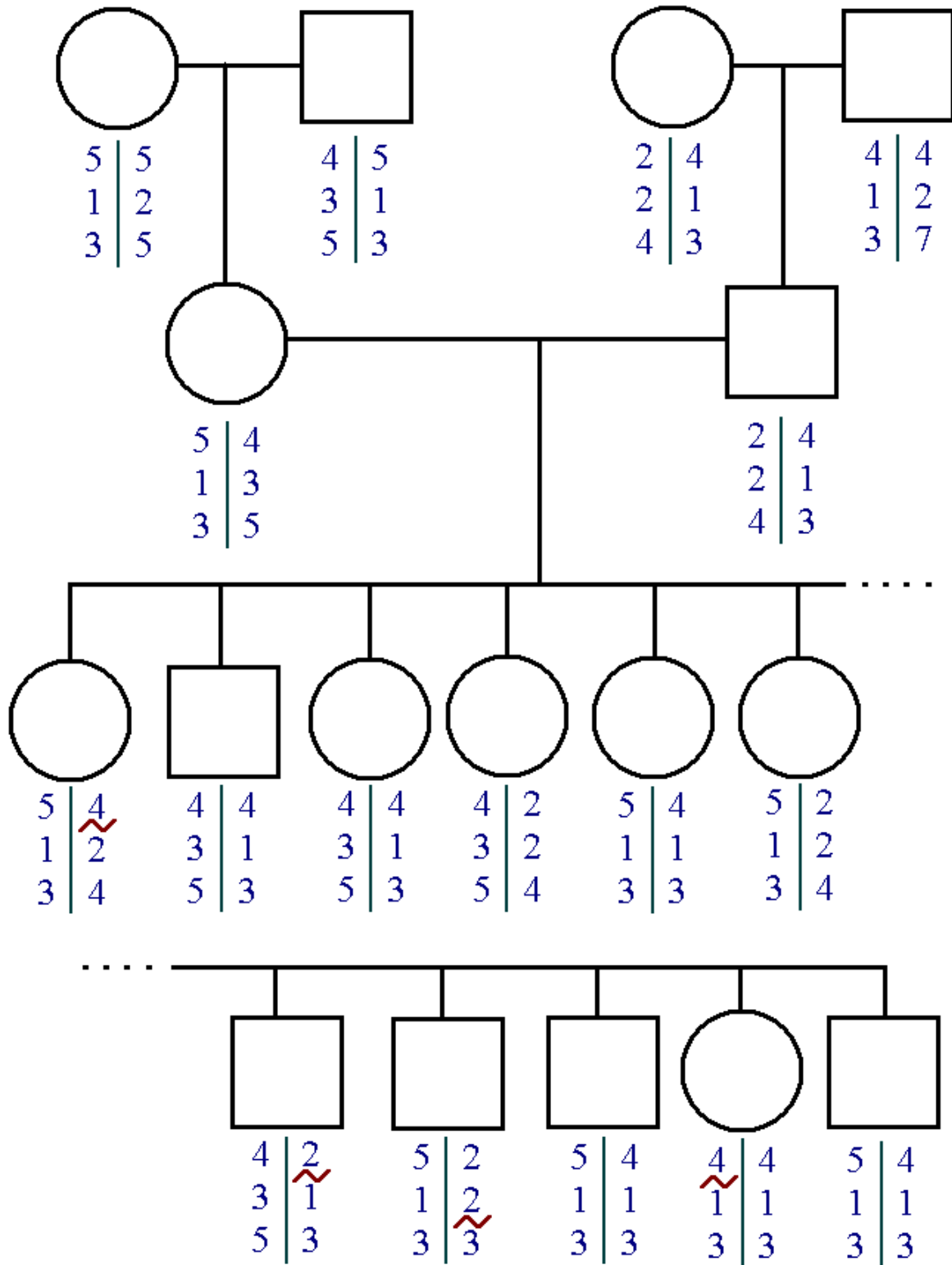
Drosophila data (Morgan et al 1935)

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

- research.marshfieldclinic.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

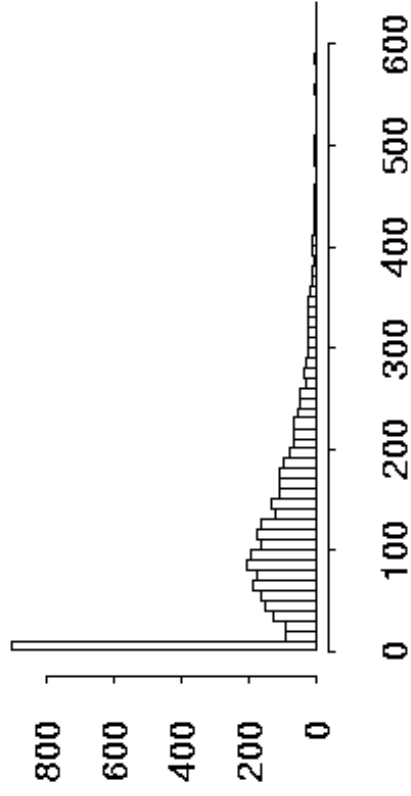


CRI-MAP *chrompic* output

CEPH individual 1331-11
maternal chromosome 10

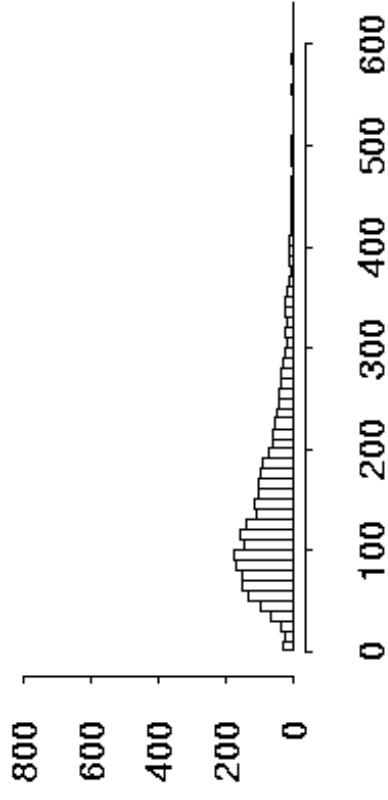
```
11111111--- 11-1111-11- --11--1111i
1-11---11- 111111--i- 11-1111-11
--111111-11 1111-111111 11--111111
111-1111i-i 11111111111 10000-0-00
0--o00-000 0000-00000 0000--0000
0000--0000 0000-00000 000-0--0--
--0-11-11- -1111ii1i-1 ---1-i-1-i
1111-i--11 11111-11i1 -11i-11111
-1-----i111 1i11111-111 -11i1-111-
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--1i111i
1i1i-11111 i-0---0000 00000-000o
o0-00o
```


Raw Data

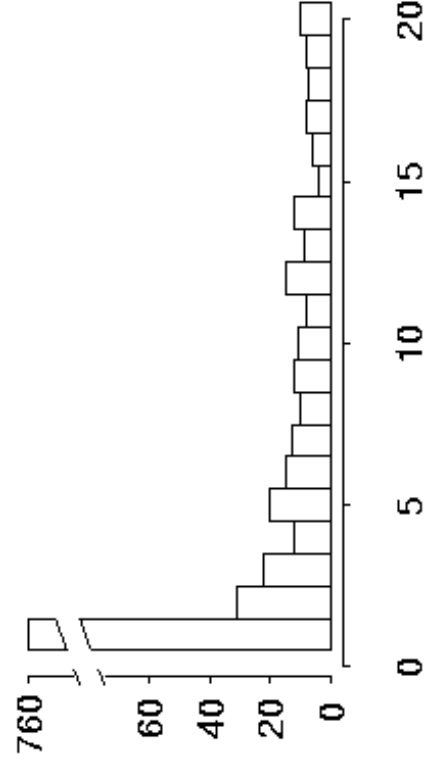


No. markers between recombinations

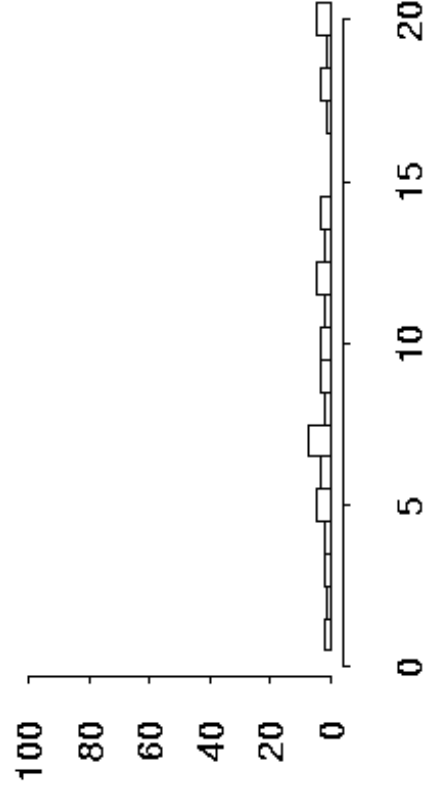
Clean Data



No. markers between recombinations

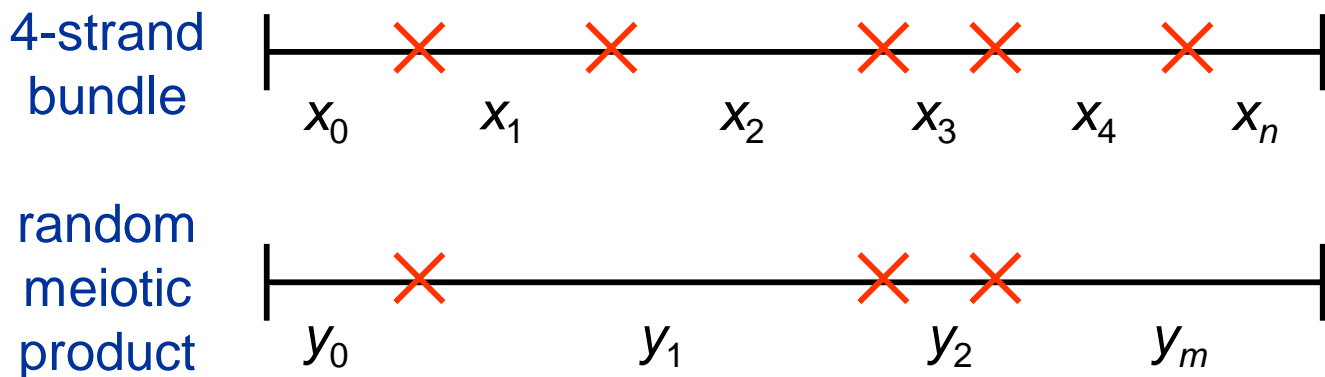


No. markers between recombinations



No. markers between recombinations

Models



- Count-location model

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

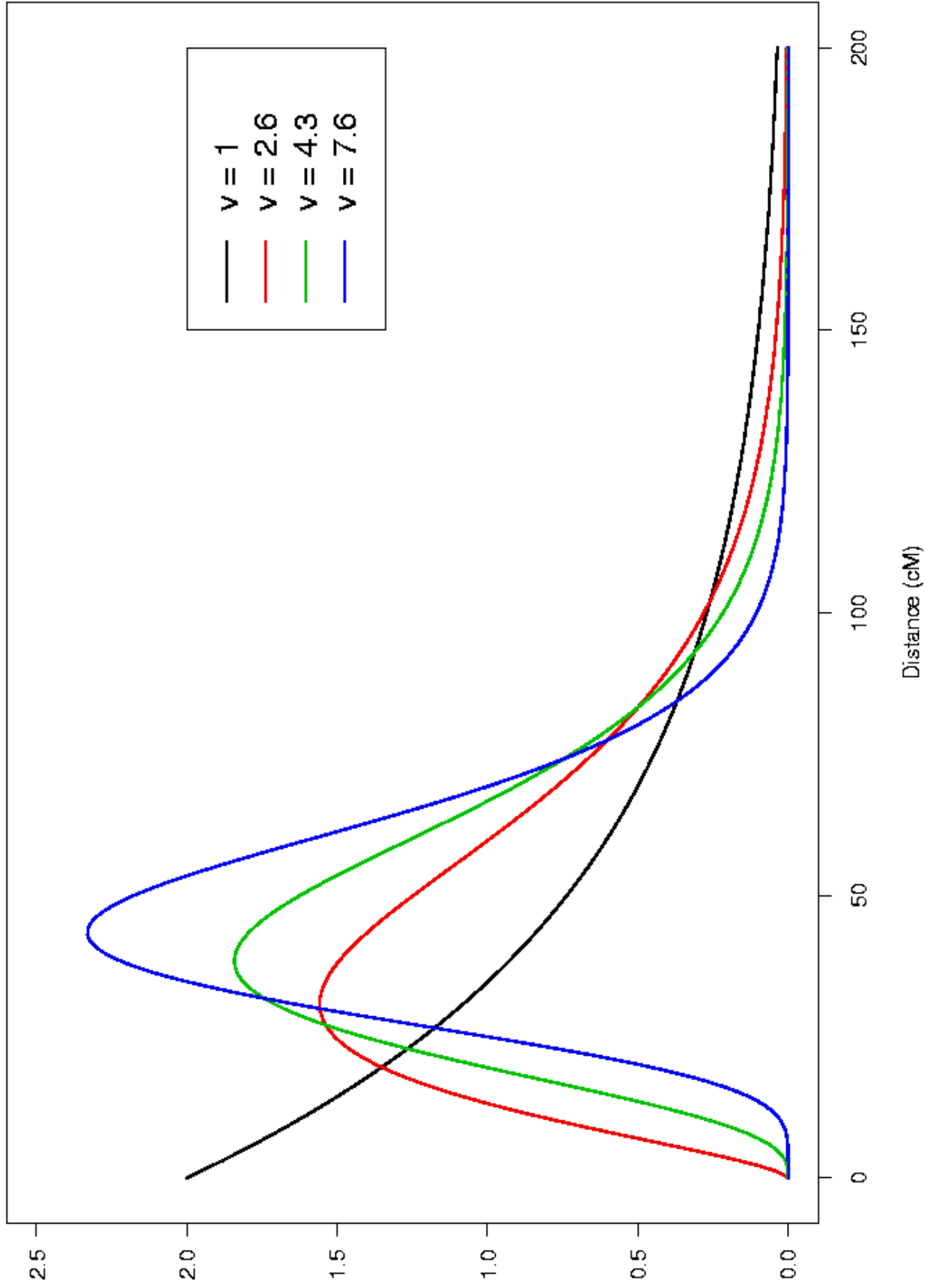
locations | $n \sim$ iid uniform

- Gamma model

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

y_i 's \sim mixtures of gammas

Dist'n of distance between chiasmata



Model fitting

- Count-location model

$m_i = \#$ crossovers

$n_i =$ underlying # 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

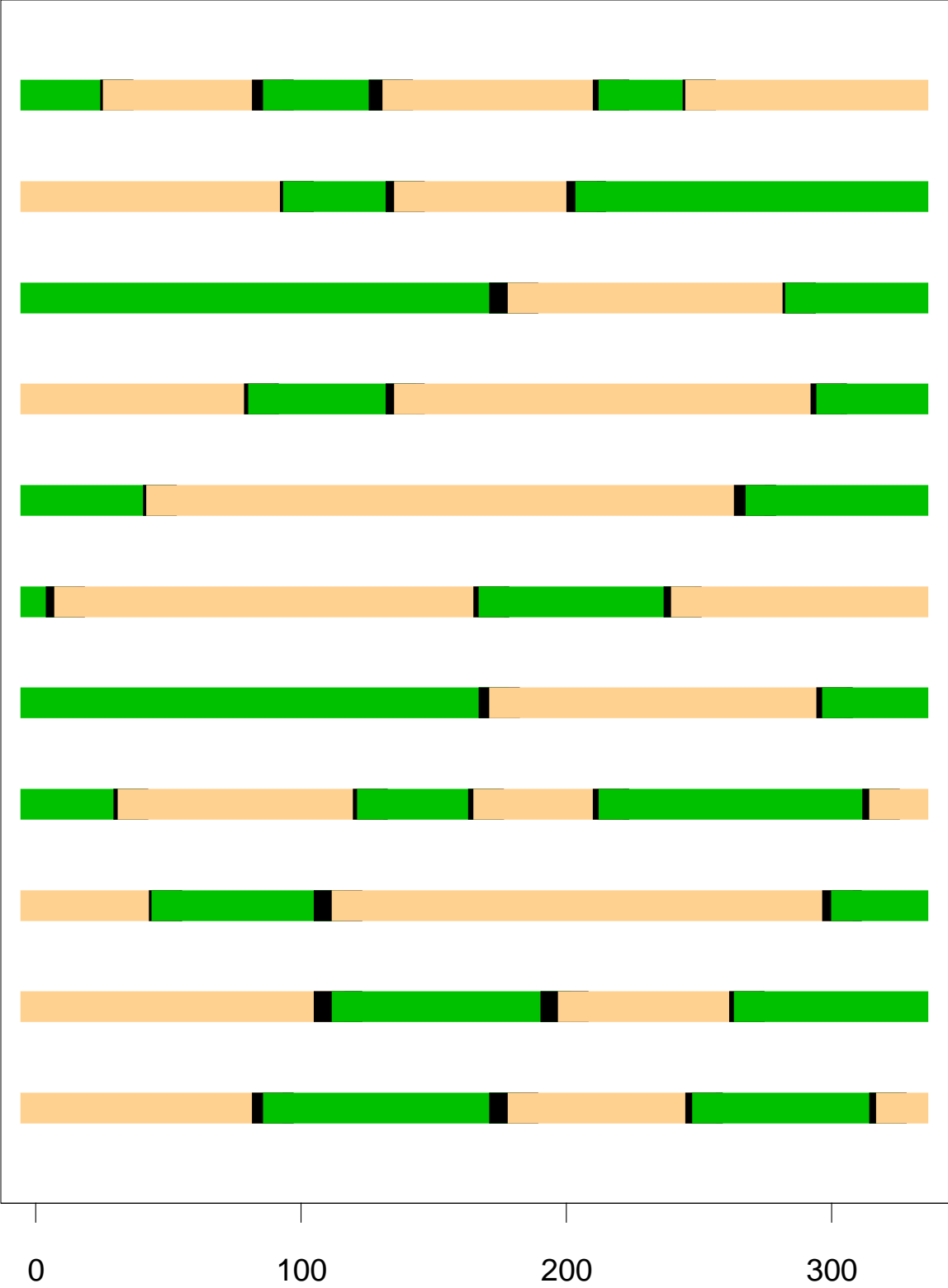
$$y_1, y_2, \dots \sim \sum (1/2)^k f(kv, 2v)$$

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

y_i 's independent

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

Another view of the data



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

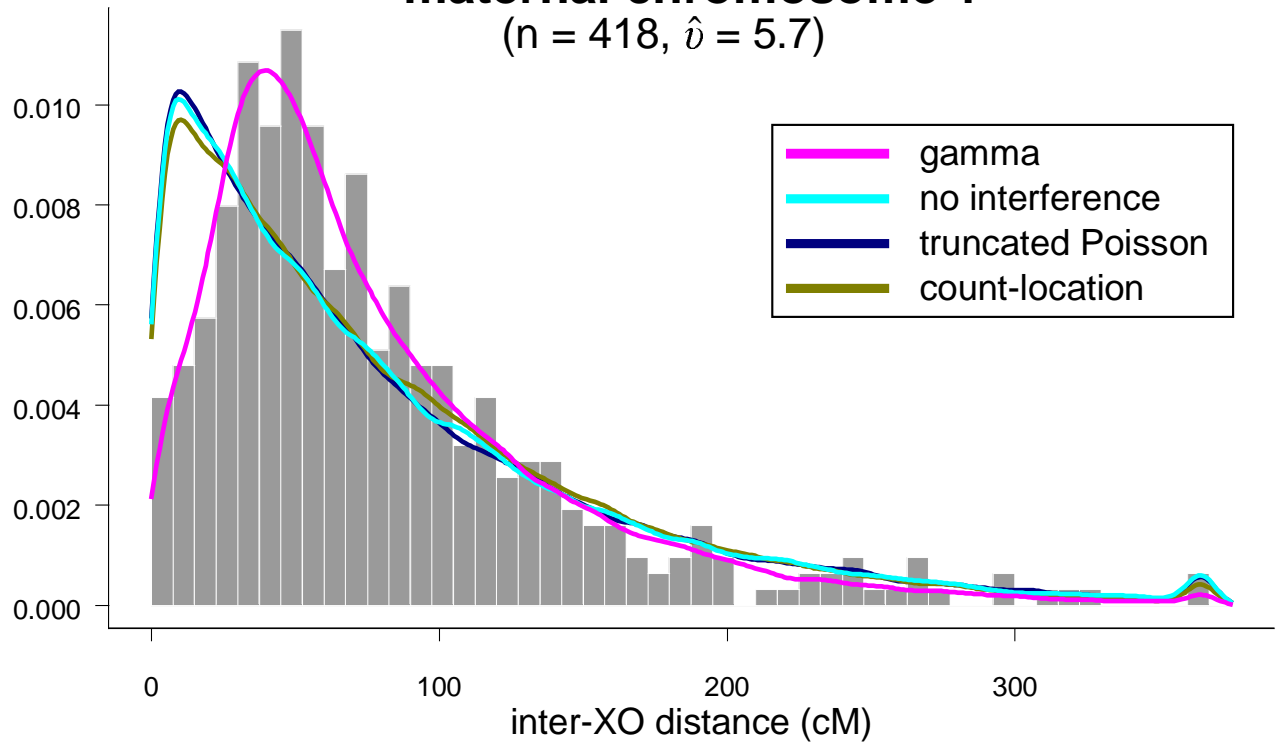
Evidence for interference:

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

paternal 1, 4, 5, 9, 14

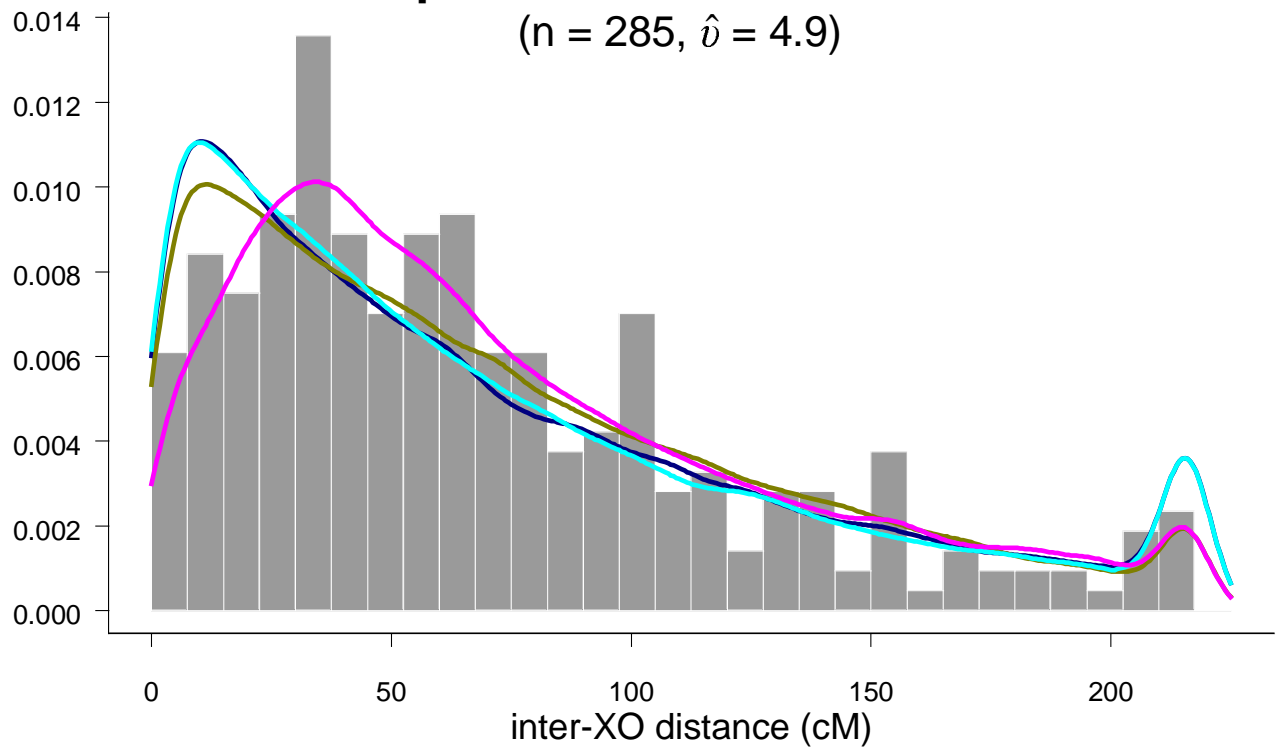
maternal chromosome 1

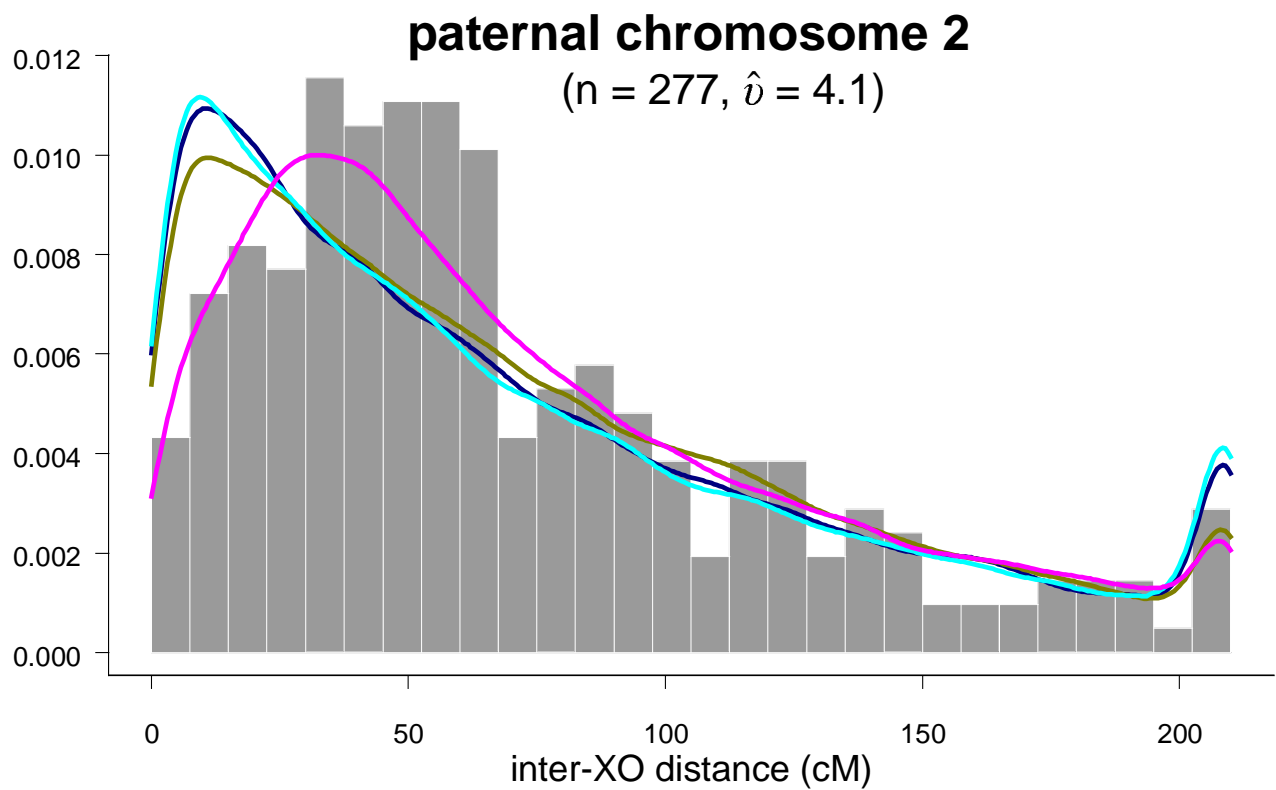
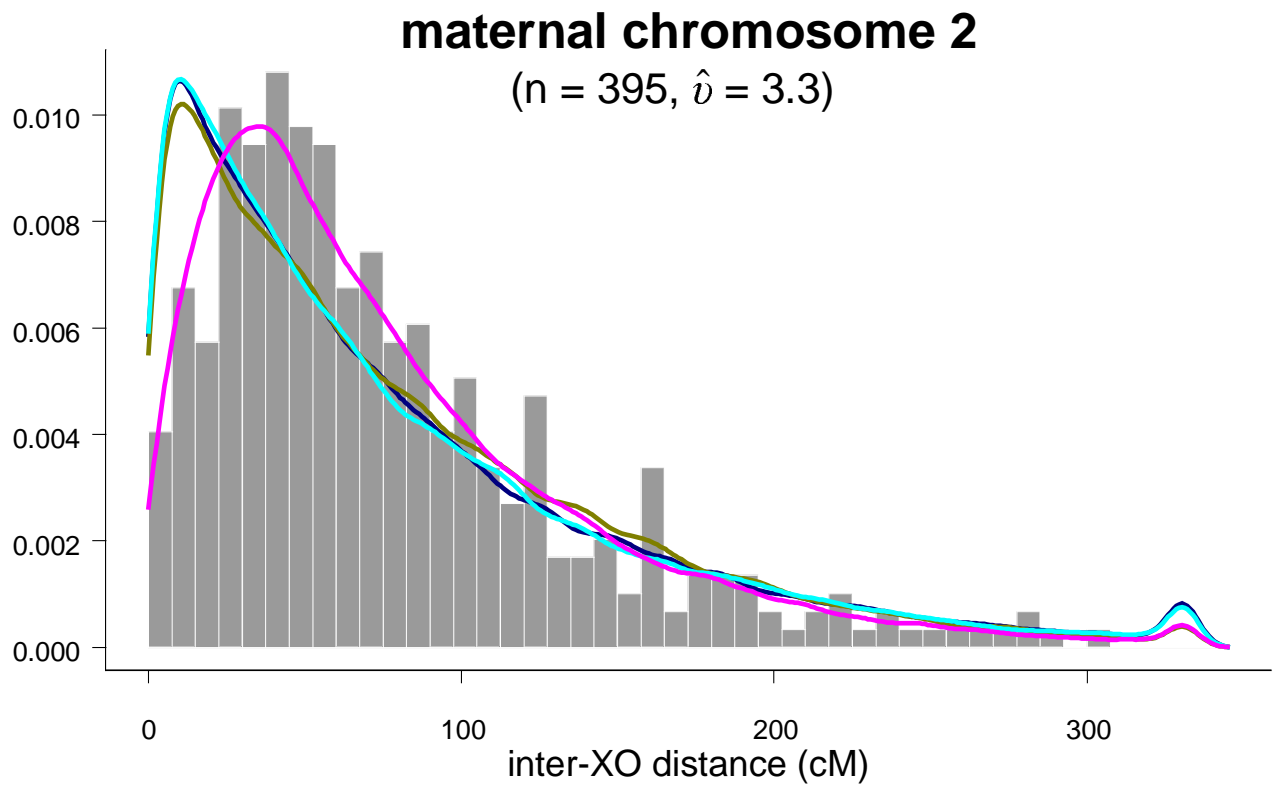
($n = 418, \hat{\nu} = 5.7$)

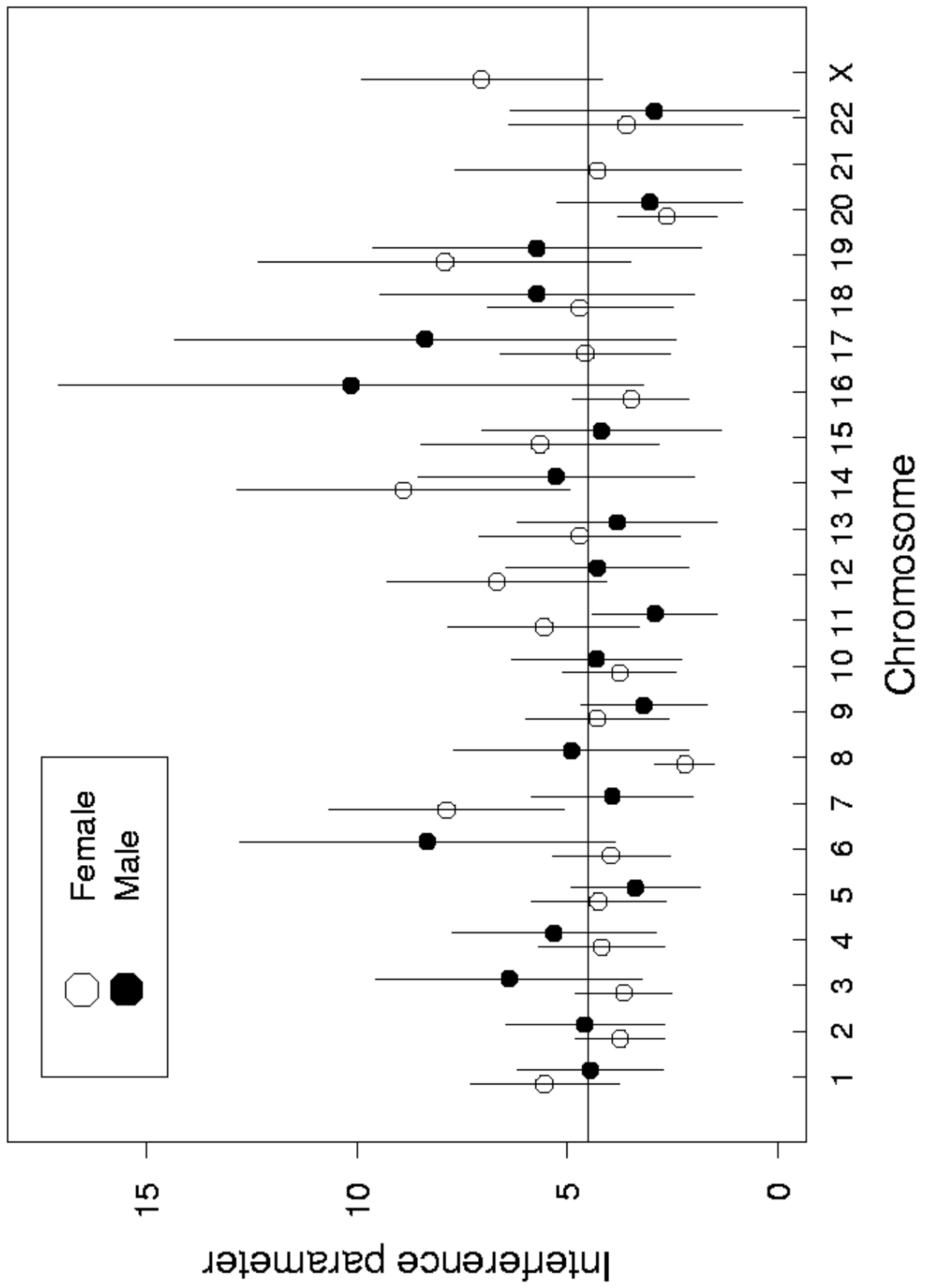


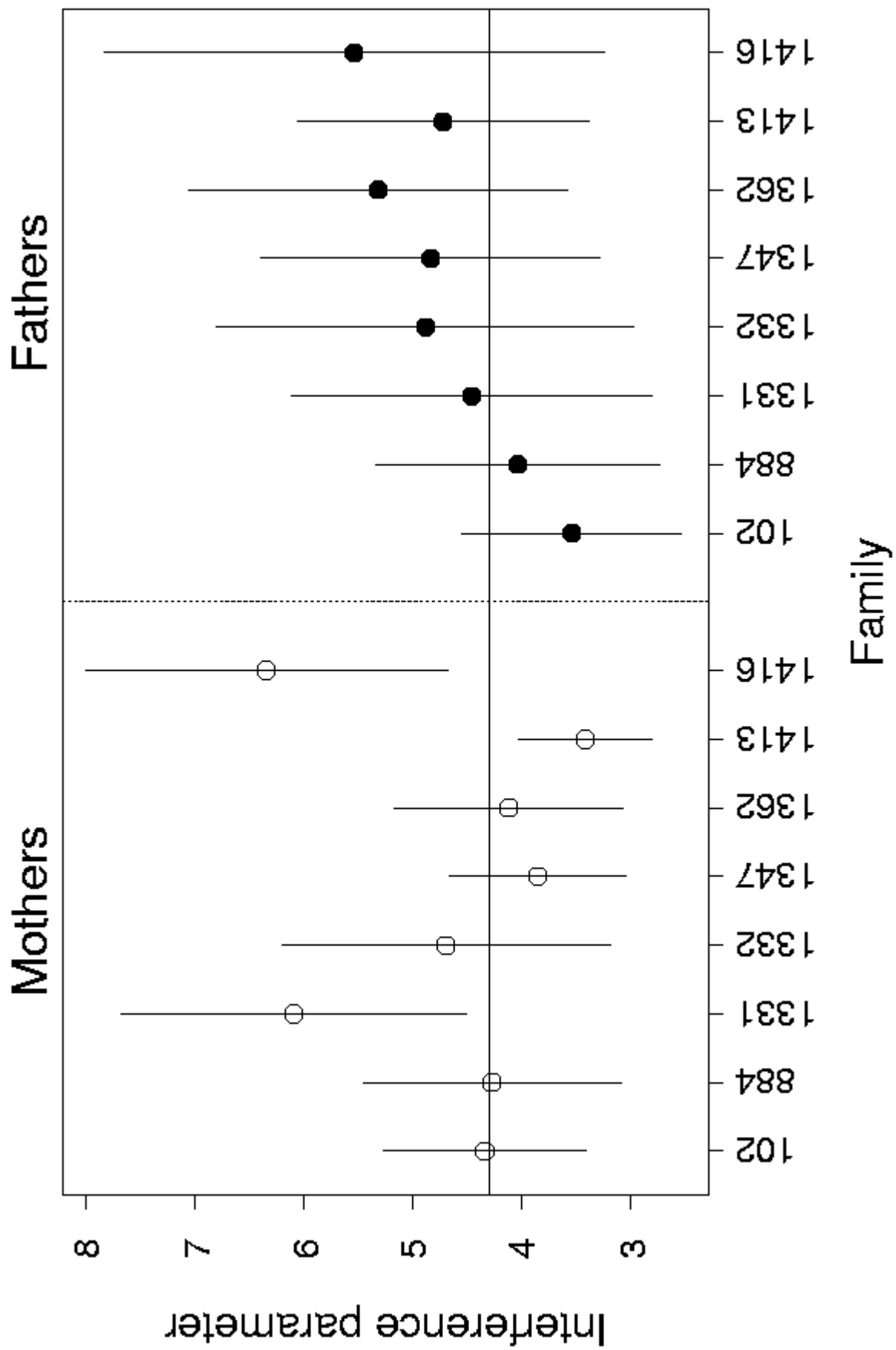
paternal chromosome 1

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

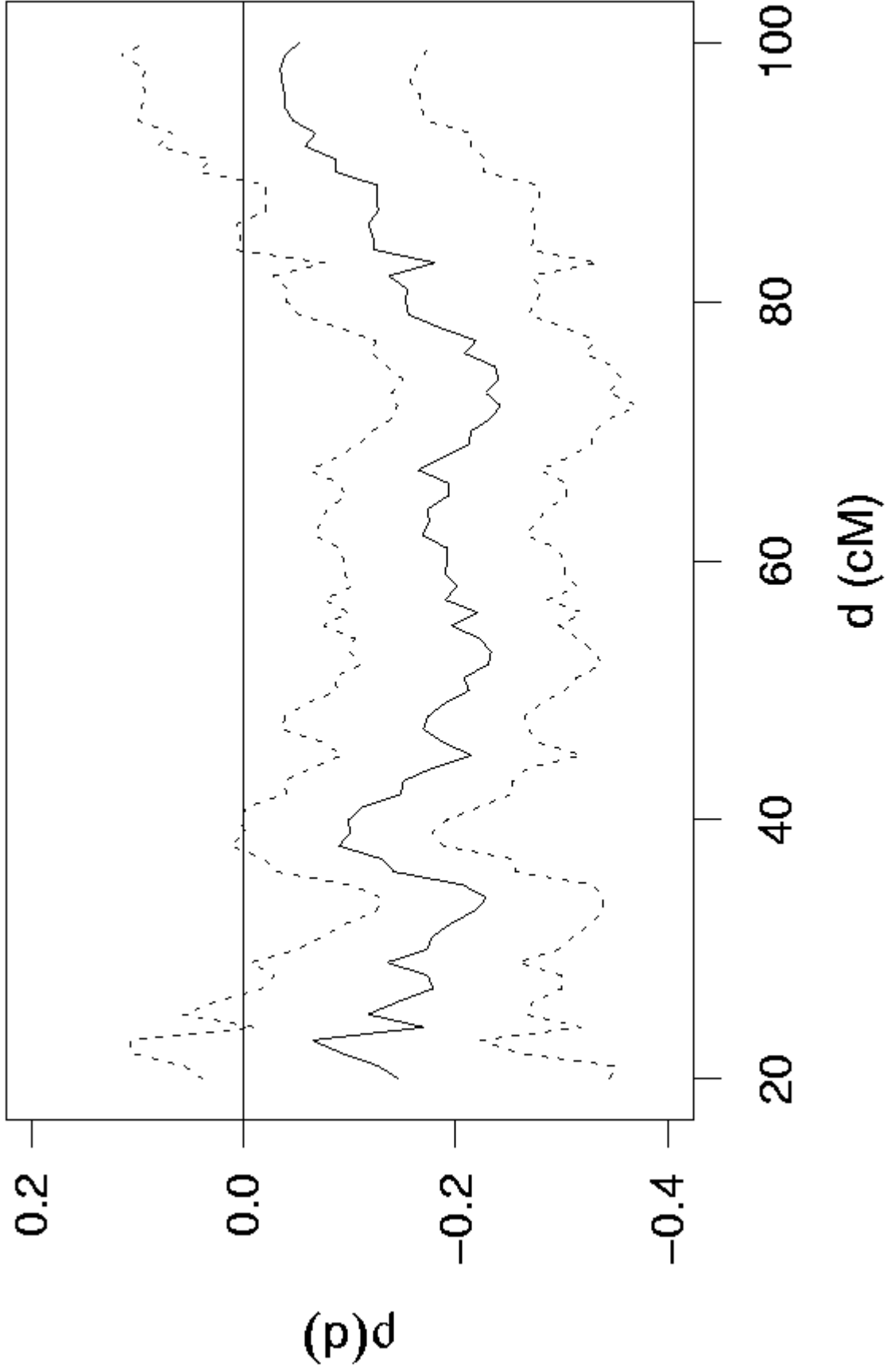


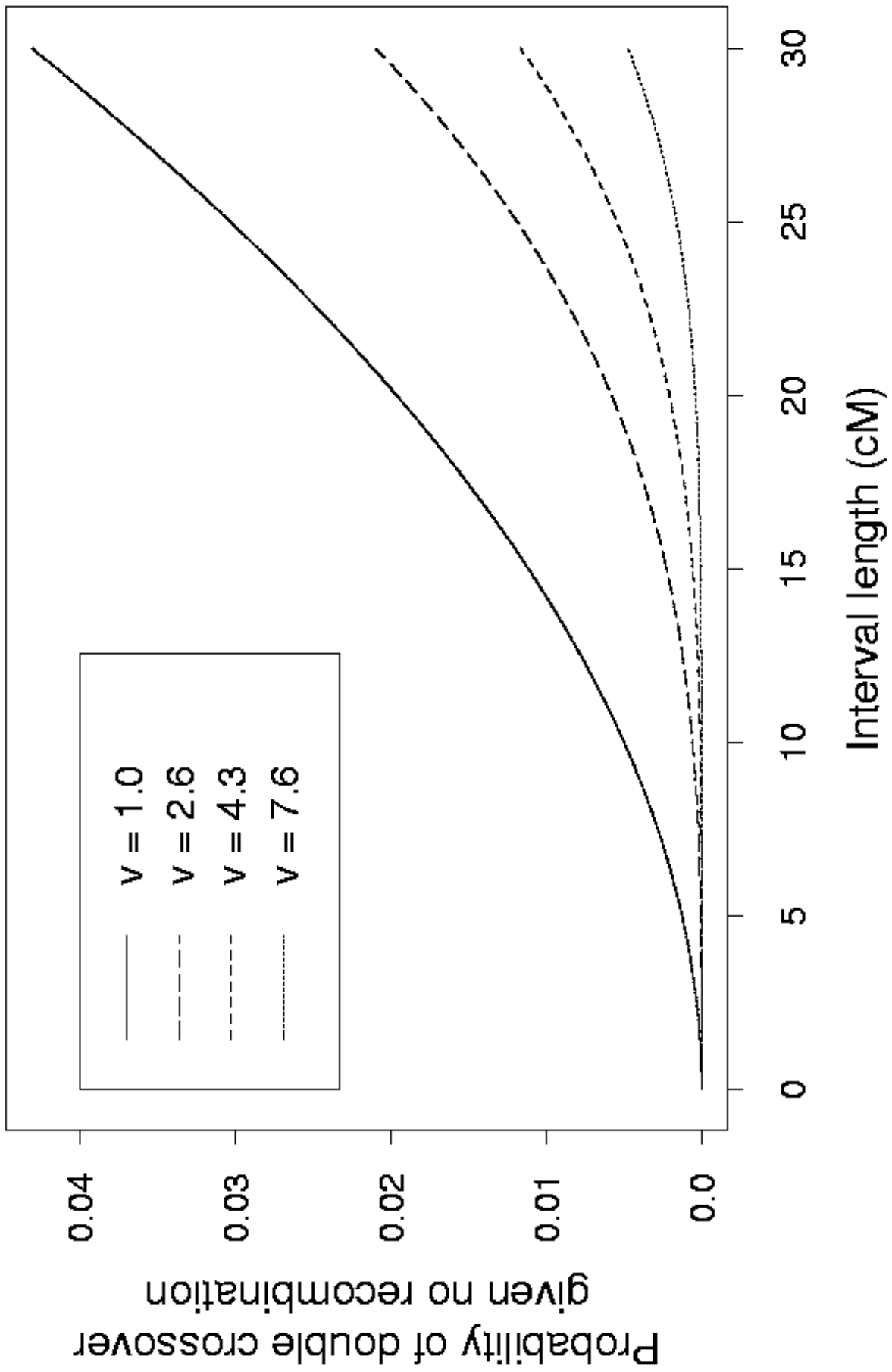






$$\rho(d) = \text{corr}(x_p, x_q | \max\{x_p, x_q\} \leq d, \min\{L_p, L_q\} \geq d)$$





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, $\nu \approx 4.3$
(stronger than Kosambi, $\nu \approx 2.6$)
 - Little variation between chr or sexes
 - Possible variation between mothers
 - Interference across the centromere