

New methods for construction of genetic linkage maps in hexaploid sweetpotato

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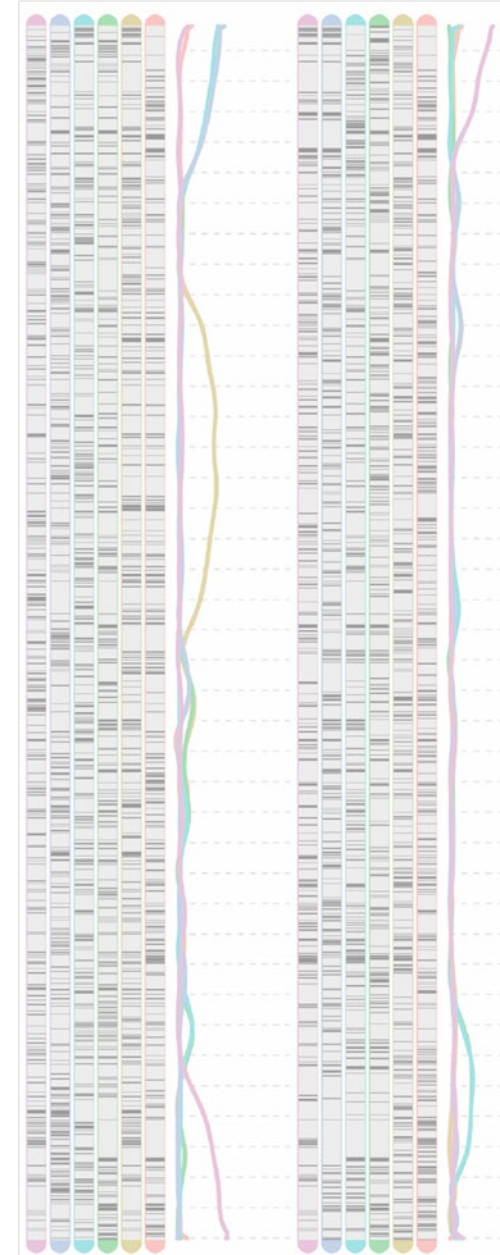
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BILL & MELINDA
GATES *foundation*

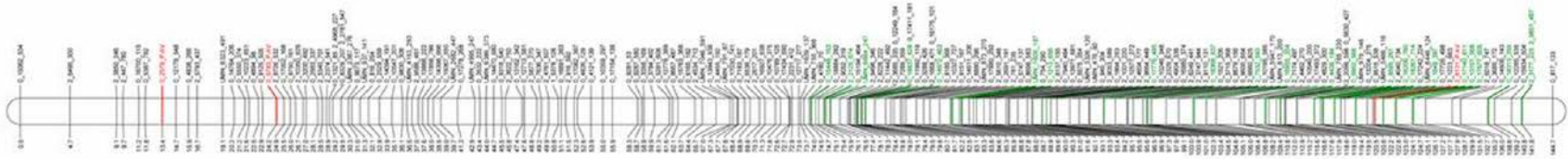
Overview

- Why do we need genetic maps?
- The general concept of genetic mapping in diploids.
 - Accessing genetic variation.
 - Genetic distance.
- Polyploid species: why are they different?
- Genotyping polyploid species.
- Genetic mapping in polyploid species.
 - The phasing problem: How to find the origin of haplotypes?
 - Recombination fraction estimation.
- Building a sweetpotato map.

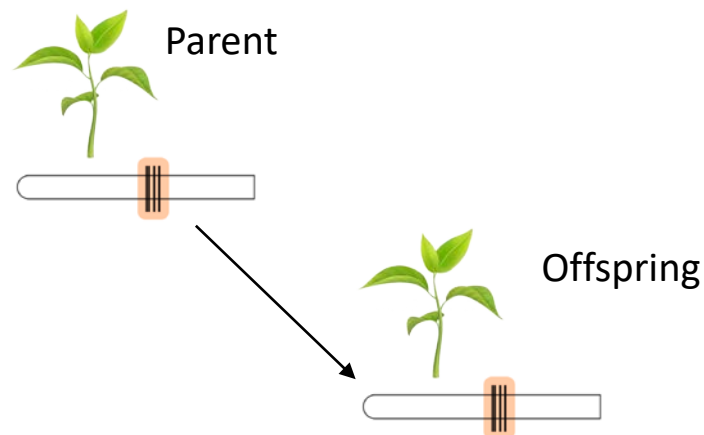


Why do we need a genetic map?

- Genetic maps are linear arrangements of markers representing chromosomes of a species

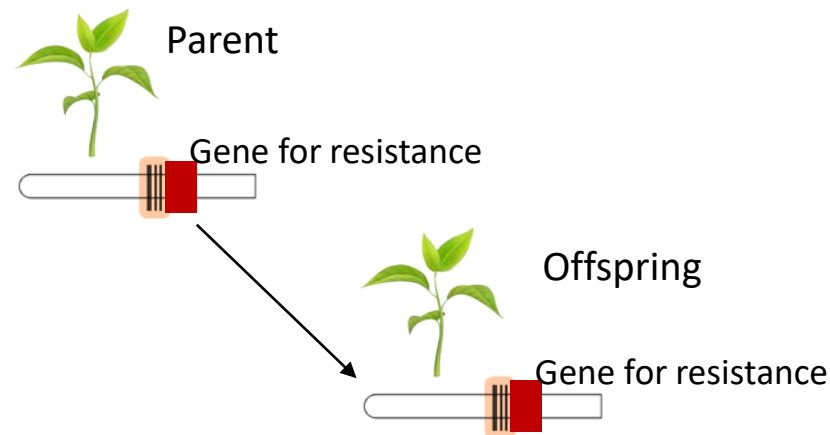


- Inheritance history** of the mapping population: close markers are inherited together



Why do we need a genetic map?

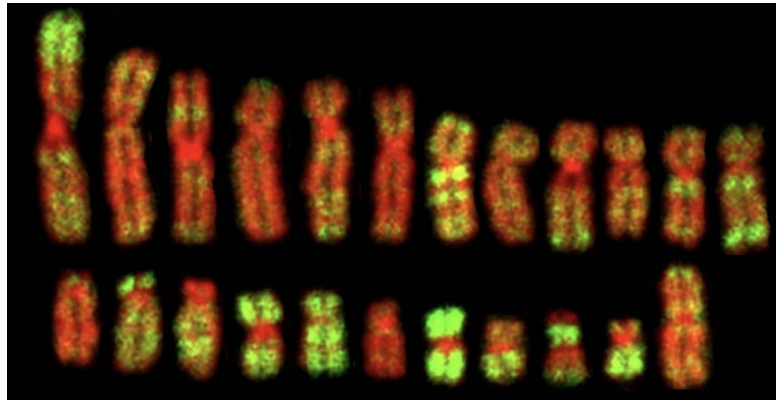
- We use markers as proxy to access important traits such as **virus and weevil resistance**, **beta-carotene** and **yield**.



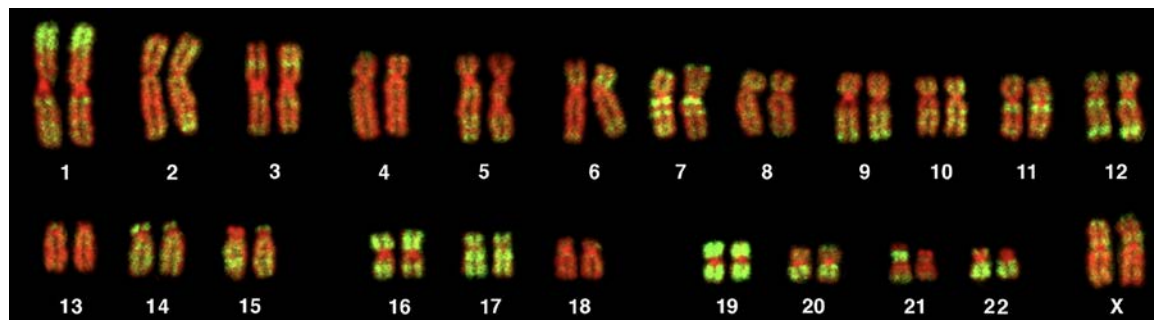
- Since we have **thousands of markers** spread across the genome, we try to find associations between them and **traits measured in the field**. The final goal, is to select **better genotypes based on the markers**. This can **speed up** the breeding process.

The general concept of genetic mapping: *diploid organisms*

- Basic chromosome number: the number of different chromosomes that make up a single complete set. In humans, this number is 23.

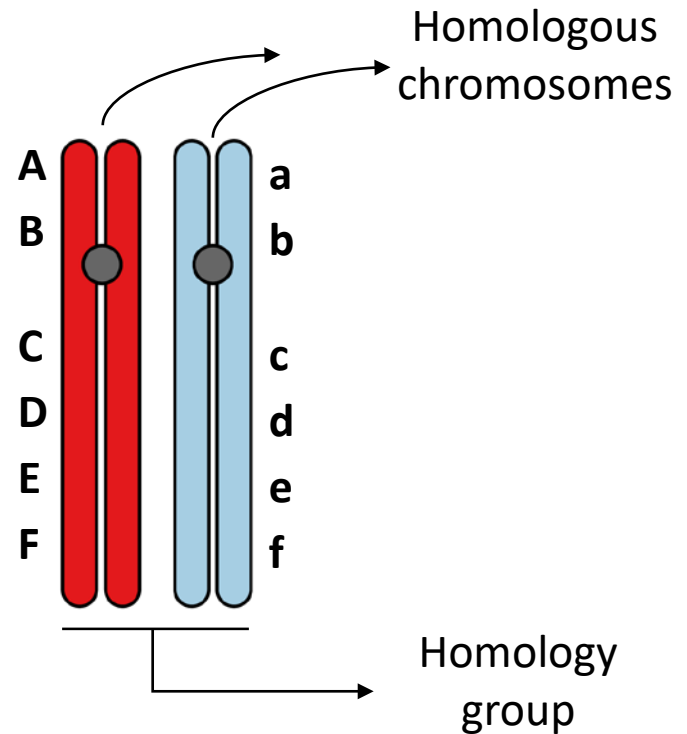
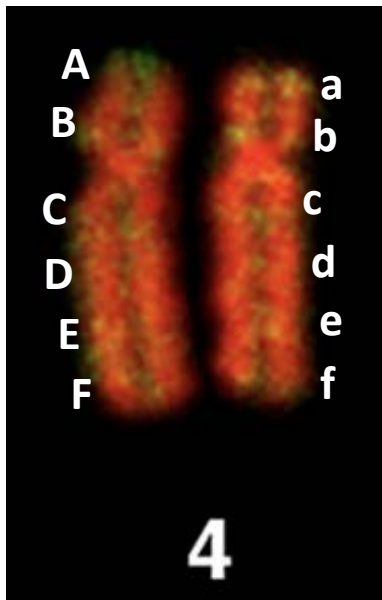


- Ploidy level: Number of basic chromosome sets. In humans, the ploidy level is two. Thus we are diploids.

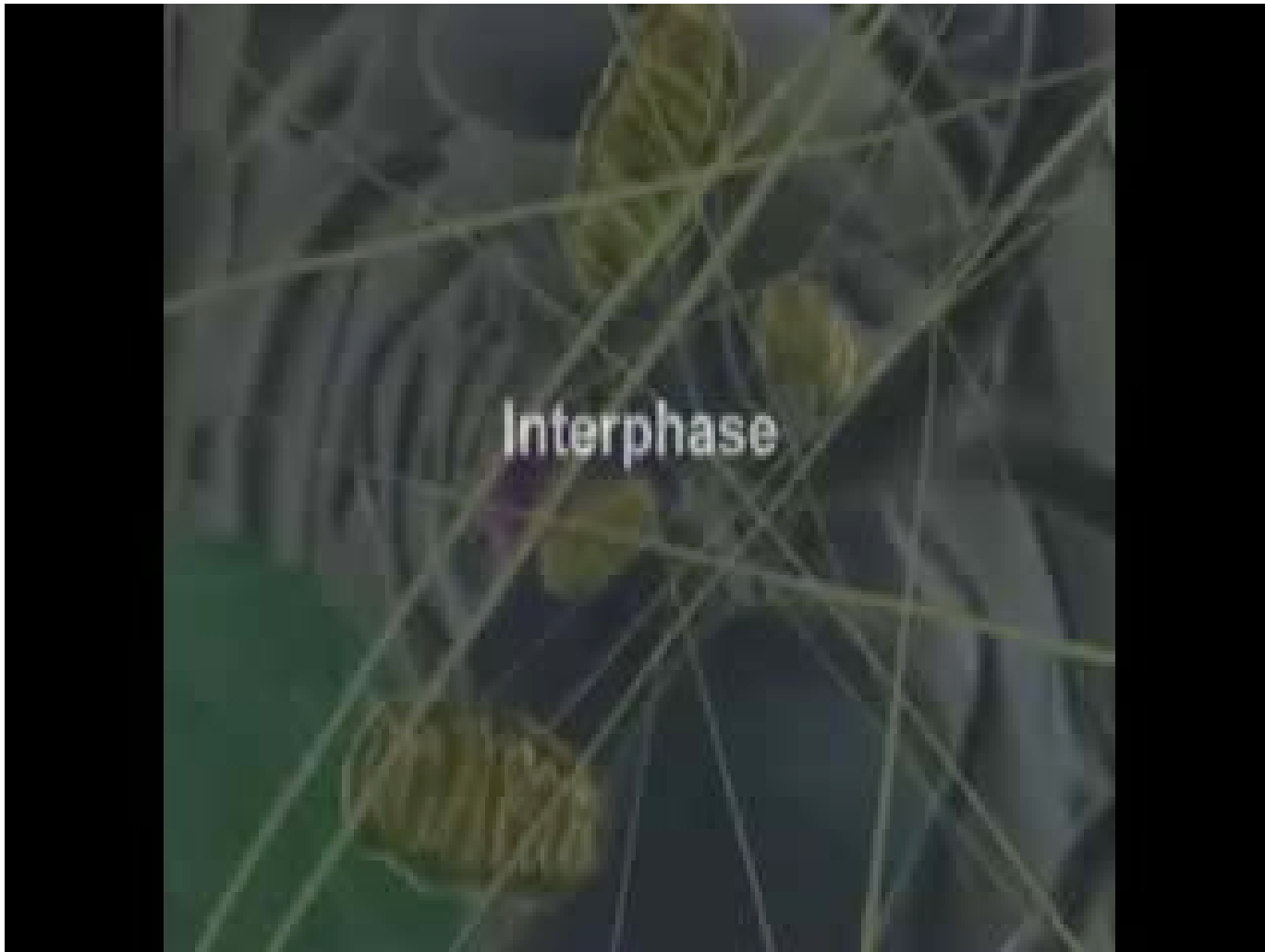


Accessing genetic variation.

- Genotyping: measurement of variations (alleles) in **homologous chromosomes** within a locus
- Molecular markers: access the allelic variation in each one of the homologous chromosomes

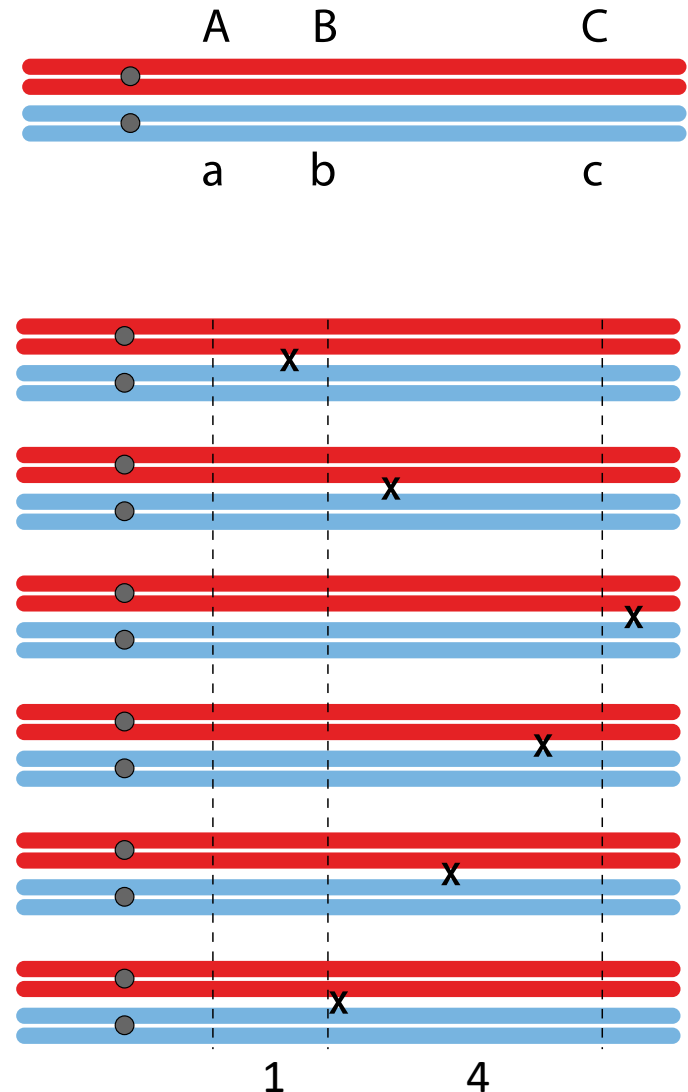


The general concept of genetic mapping: *the crossing over process*

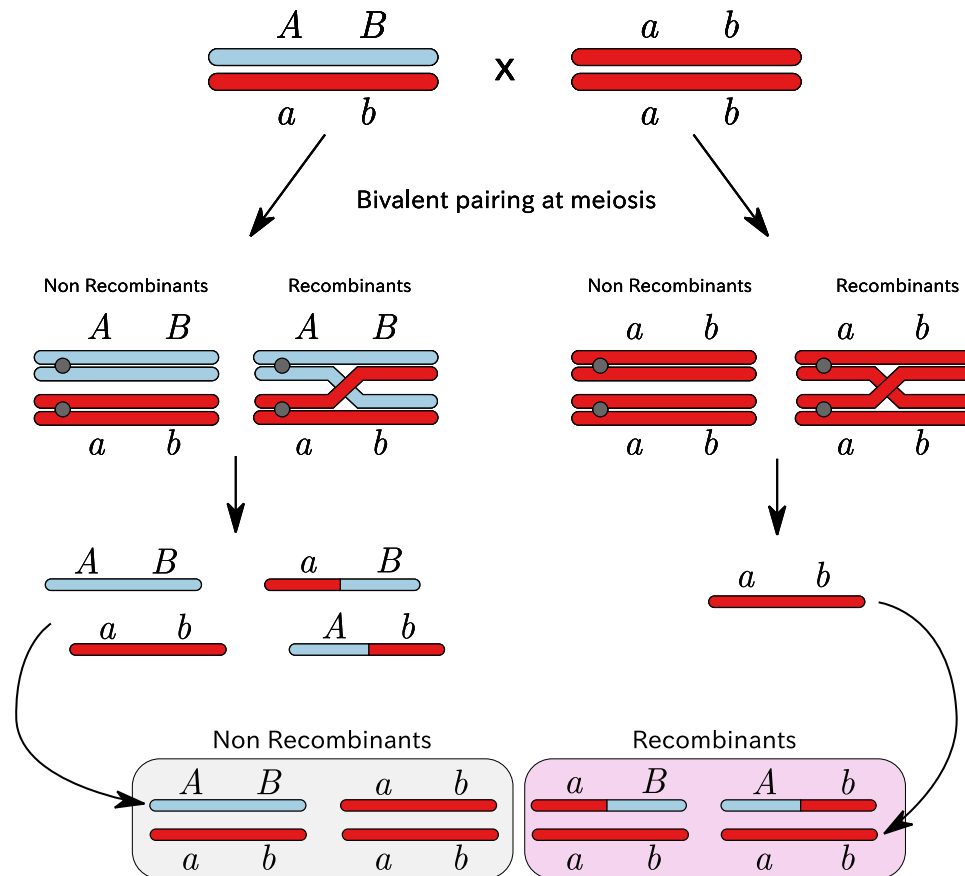


The general concept of genetic mapping

- Suppose we have a chromosome with **three markers**, two of them are **close** to each other (A and B) and the third (C) is **further away**.
- The process described in the video will happen several times and the **gametes** will be stored in pollen and ovule cells
- The number of crossing overs detected between two points will be **proportional to their physical distance**.



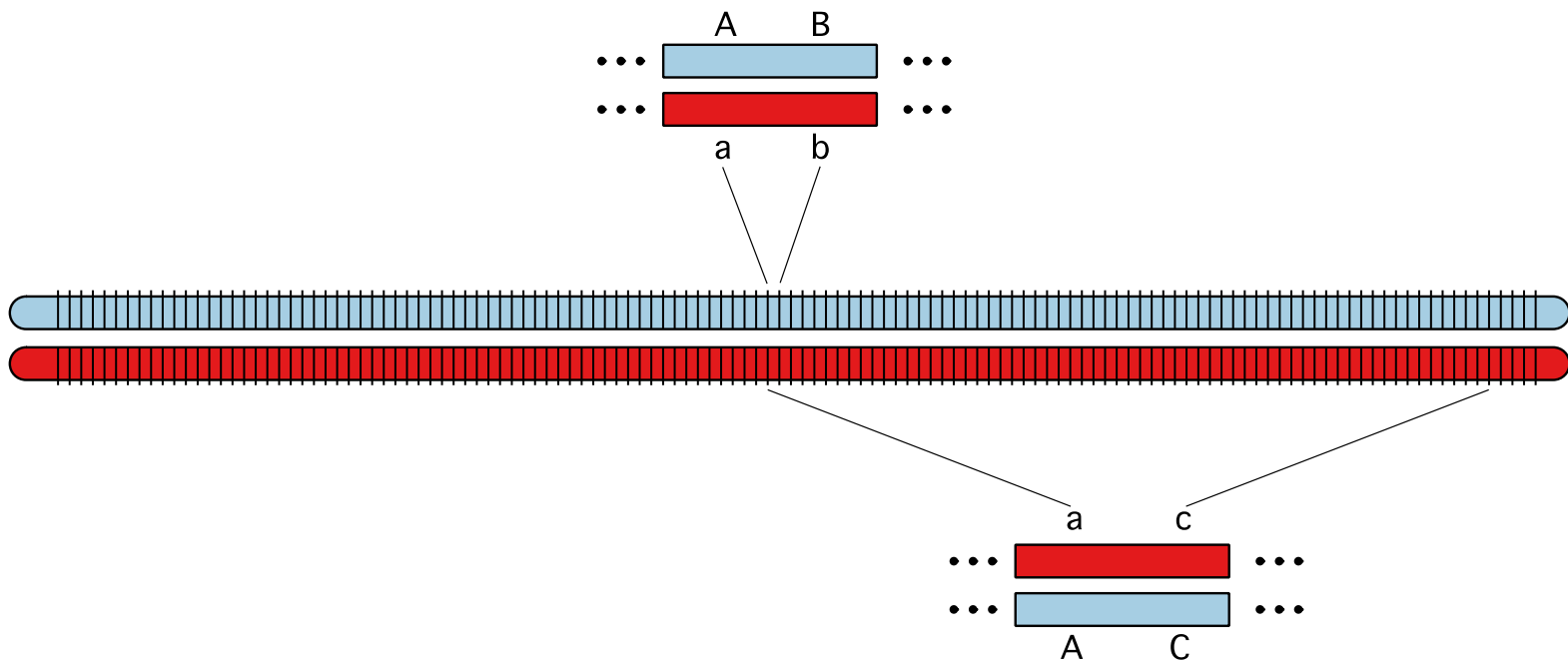
The general concept of genetic mapping



$$\hat{r} = \frac{\# \text{recombinants}}{\# \text{total}}$$

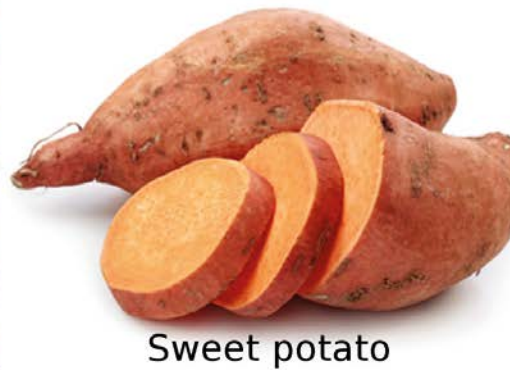
The general concept of genetic mapping

- The two markers presented are part of a bigger picture. However, the reasoning of count crossing-overs works for any pair of markers



Polyploid species

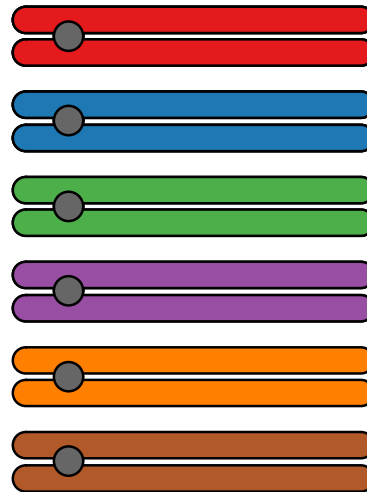
Examples of polyploid crops



Polyploid species: why are they different?

- Polyploids are organisms with multiple sets of chromosomes.

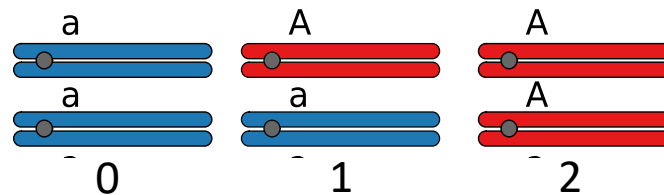
Hexaploid



- Two problems:
 - How to access the genetic variation?
 - How to count crossing overs?

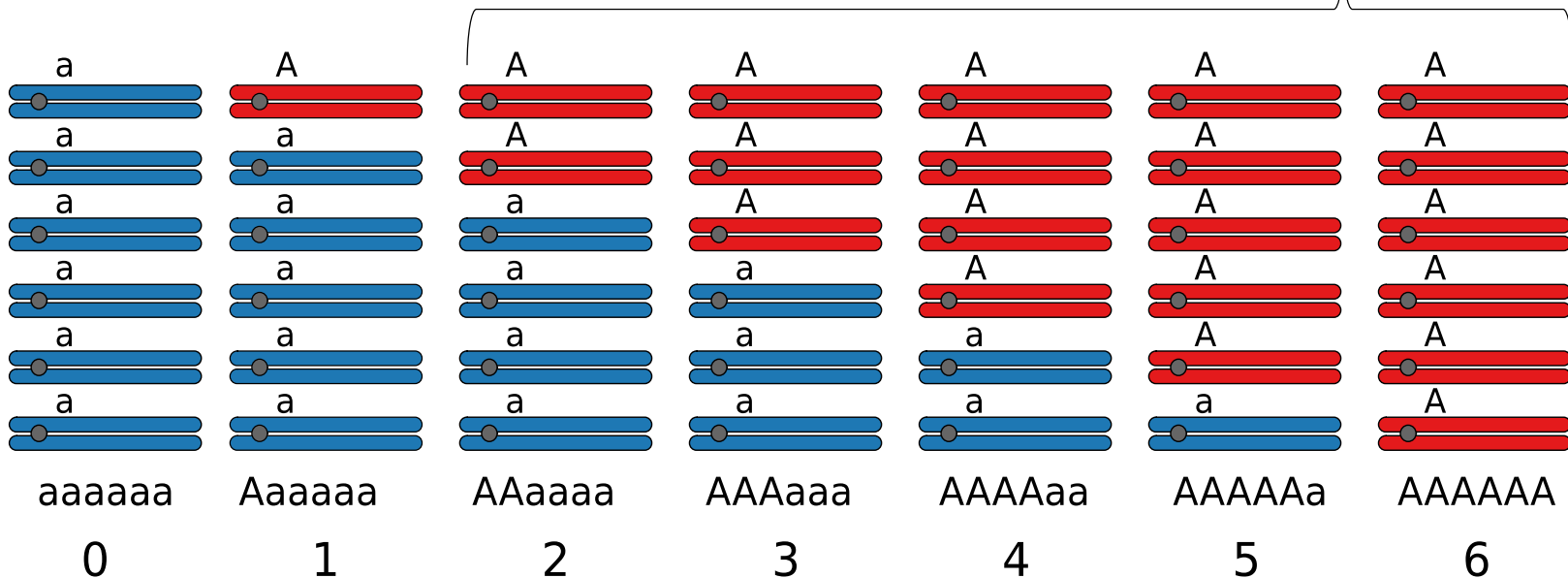
Genotyping polyploid species

- In diploids, we usually refer to the genotypes using the quantity of a reference alleles, in this case, A



- In polyploids, this number is called **dosage**

Multidose markers

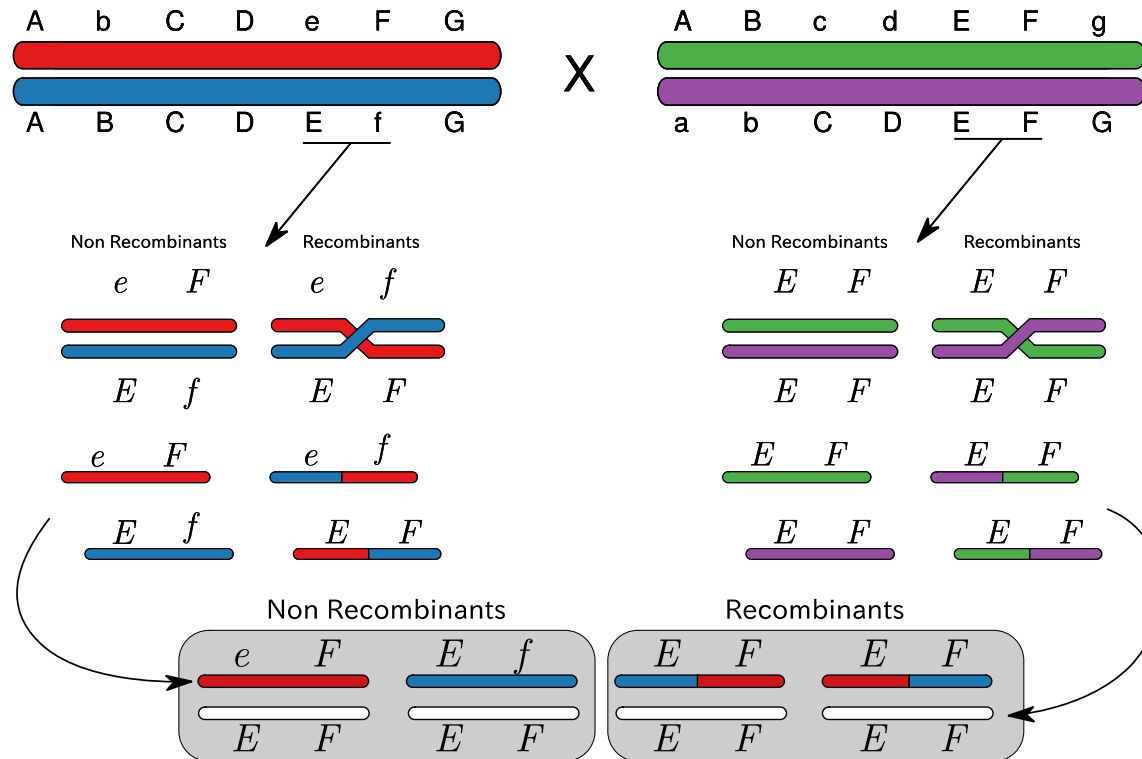


Genotyping polyploid species

- Example of input file for *MAPPoly* software

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<div> <div> <div>Paste</div> <div>Cut</div> <div>Copy</div> <div>Format</div> </div> <div> <div>Calibri (Body)</div> <div>12</div> <div>A</div> <div>A</div> </div> <div> <div>B</div> <div>I</div> <div>U</div> </div> <div> <div>Wrap Text</div> <div>Merge & Center</div> </div> <div> <div>General</div> <div>\$</div> <div>%</div> <div></div> </div> <div> <div>Conditional Formatting</div> <div>Format as Table</div> <div>Cell Styles</div> <div>Insert</div> </div> </div>											
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	A	B	C	D	E	F	G	H	I	J	K
1		Ind_1	Ind_2	Ind_3	Ind_4	Ind_5	Ind_6	Ind_7	Ind_8	Ind_9	Ind_10
2	M_1	1	2	1	2	2	1	1	2	2	1
3	M_2	0	1	0	1	1	1	0	0	1	1
4	M_3	2	2	1	2	2	2	1	0	1	2
5	M_4	1	0	1	0	1	1	2	1	0	1
6	M_5	0	0	0	0	0	1	1	1	0	1
7	M_6	0	0	0	0	0	1	1	1	0	1
8	M_7	2	1	1	1	1	3	0	1	2	1
9	M_8	2	4	2	4	3	2	3	3	4	3
10	M_9	1	0	1	0	0	1	0	1	1	0
11	M_10	3	2	2	3	3	3	2	1	3	1
12	M_11	1	0	1	0	0	1	0	1	1	0
13	M_12	2	1	1	1	1	3	0	1	2	1
14	M_13	2	2	1	1	3	3	2	2	1	3
15	M_14	2	1	2	1	1	1	1	1	1	1
16	M_15	0	0	1	0	0	0	0	1	0	0

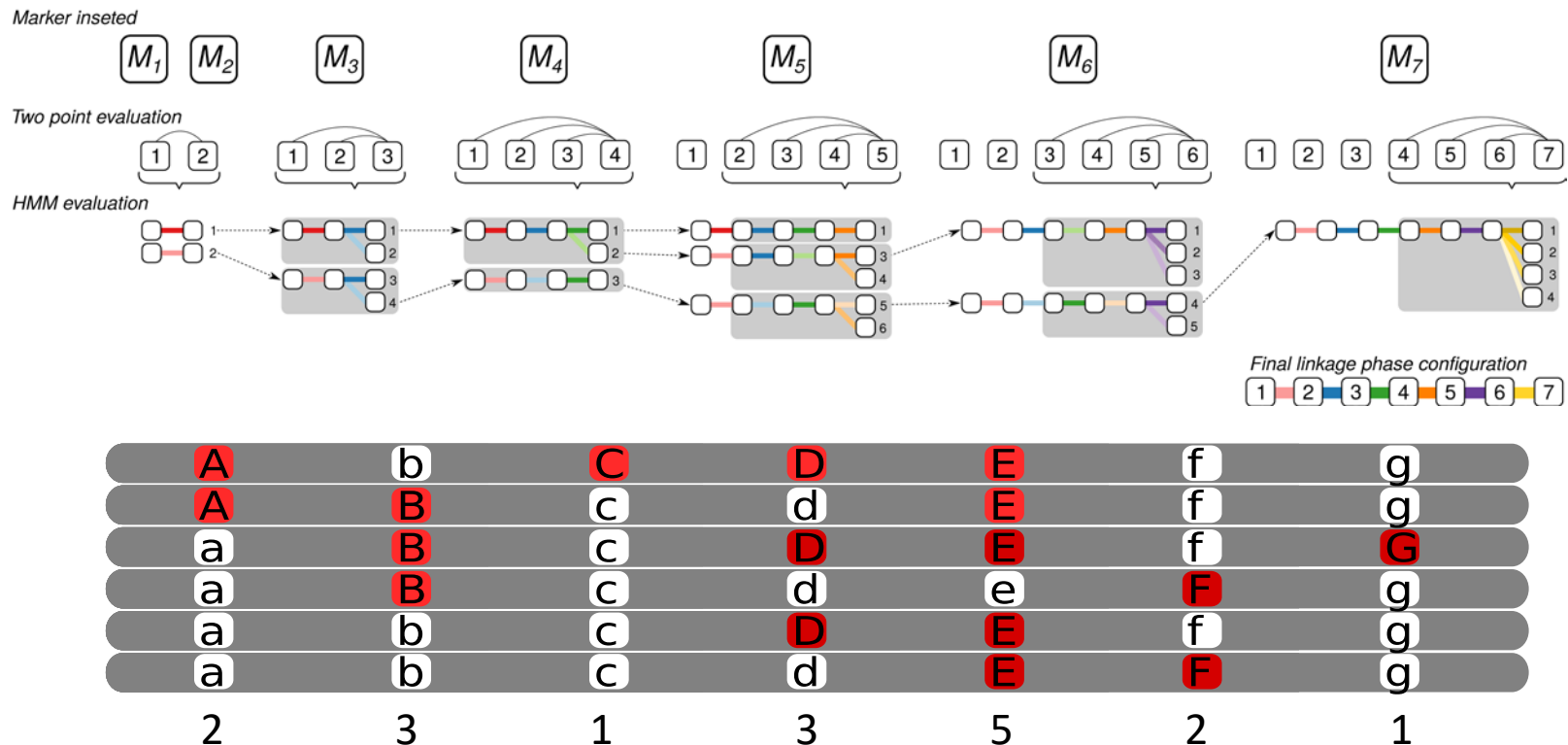
The importance of haplotype phasing in polyploids



- In this case, we **know how the alleles are arranged** in parental chromosomes. Thus, we can distinguish recombinant and non recombinant gametes.
- If this information is not available, we can **infer** that gametes e-F and E-f are non recombinants, since they will be more frequent than E-F.

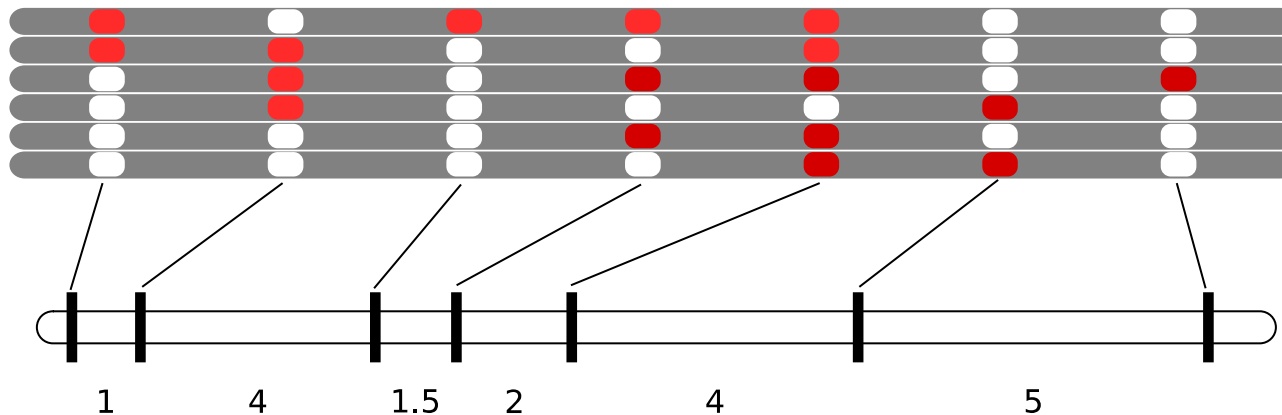
Haplotype phasing and assembly of homology groups

- In polyploids, we do not know how alleles are arranged in parental chromosomes
- We know the dosage, but we need to infer the **linkage phase configuration** of the alleles



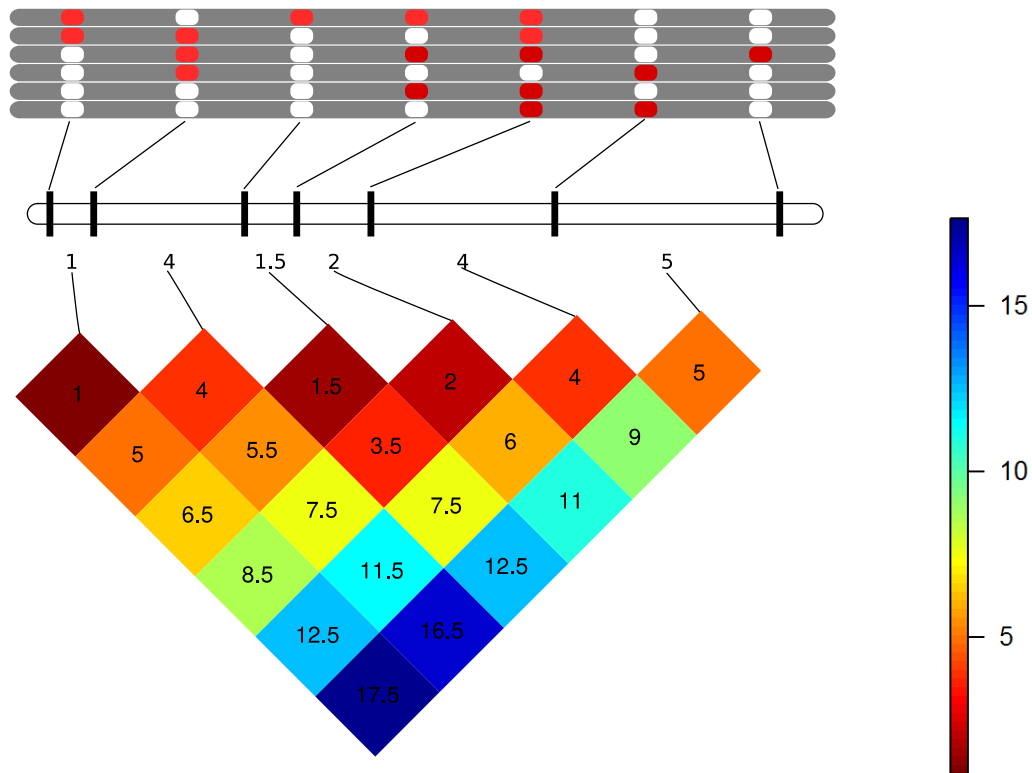
Recombination fraction estimation

- Once we have the linkage phase configuration, we can distinguish between recombinant and parental gametes, and **count the number of crossing over between markers**.
- The procedure is not as straightforward as the one used in diploids, but the idea is the same



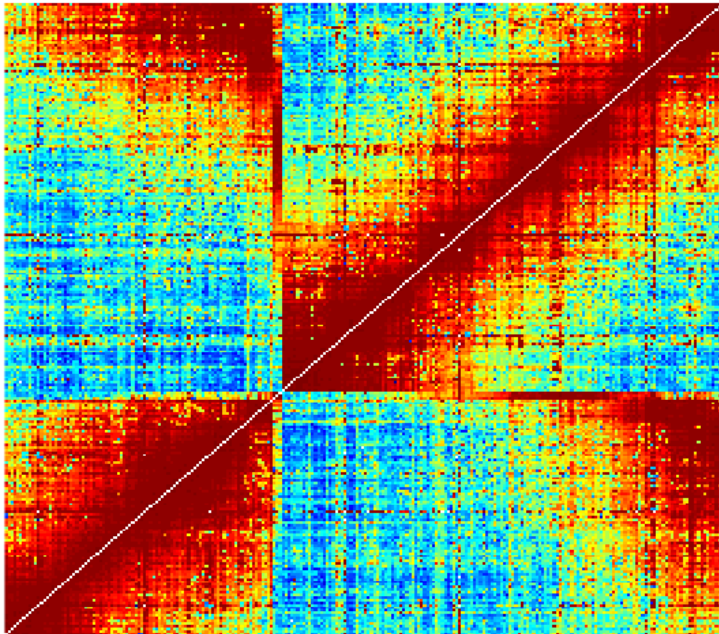
Recombination fraction matrix

- Important way to diagnose problems in genetic maps



Recombination fraction matrix

- Important diagnostic tool way to find problems in genetic maps



Map with a inversion

Map without inversion

Biparental Population - BT

- Beauregard x Tanzania
- 315 individuals, 26164 high quality SNPs



Beauregard

X

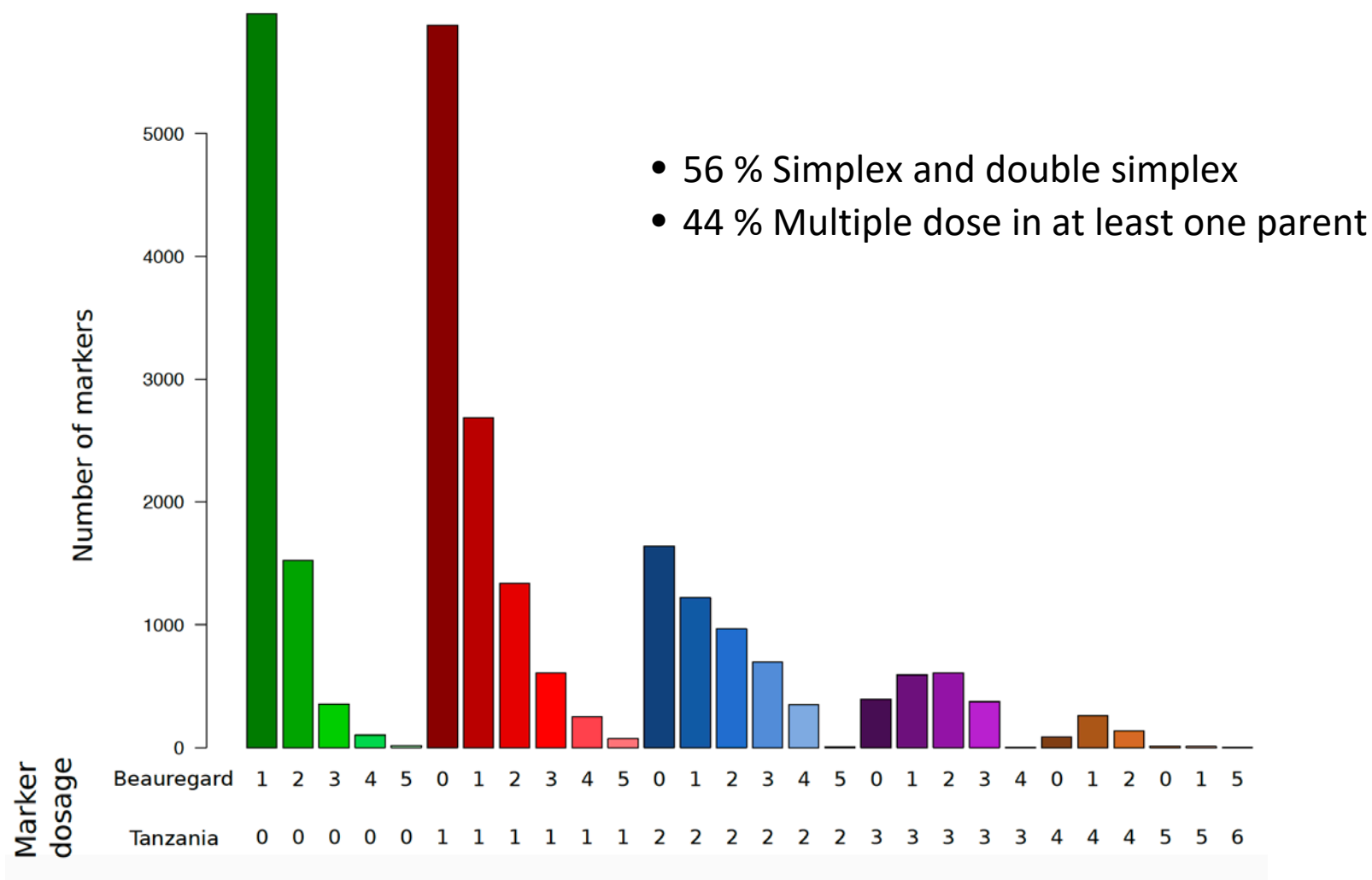


Tanzania

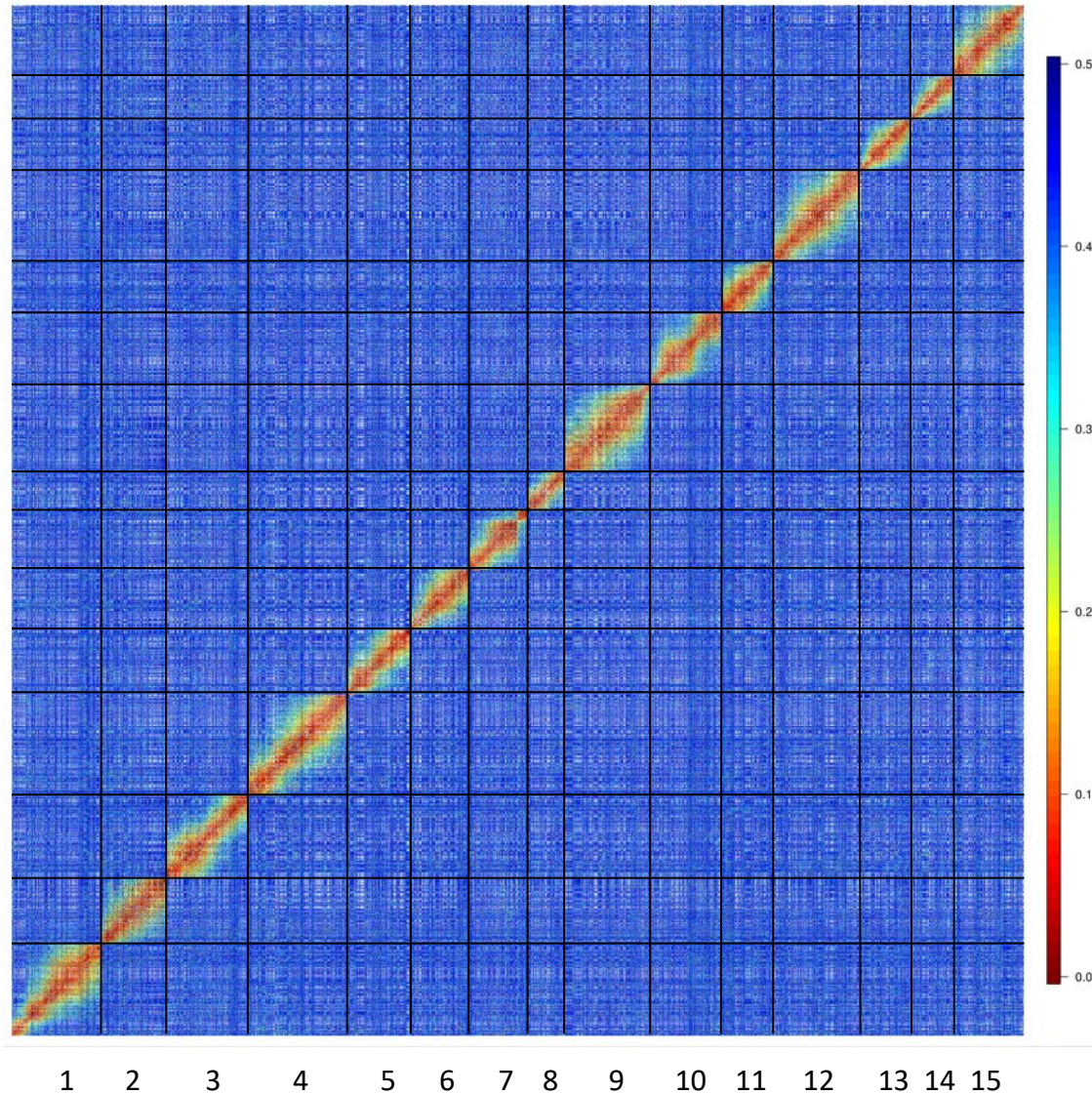


Results - Genotyping Calling - BT population

- Distribution of SNP dosage in both parents for all 26,164 SNPs.



Ordering with MDS – 15 linkage groups

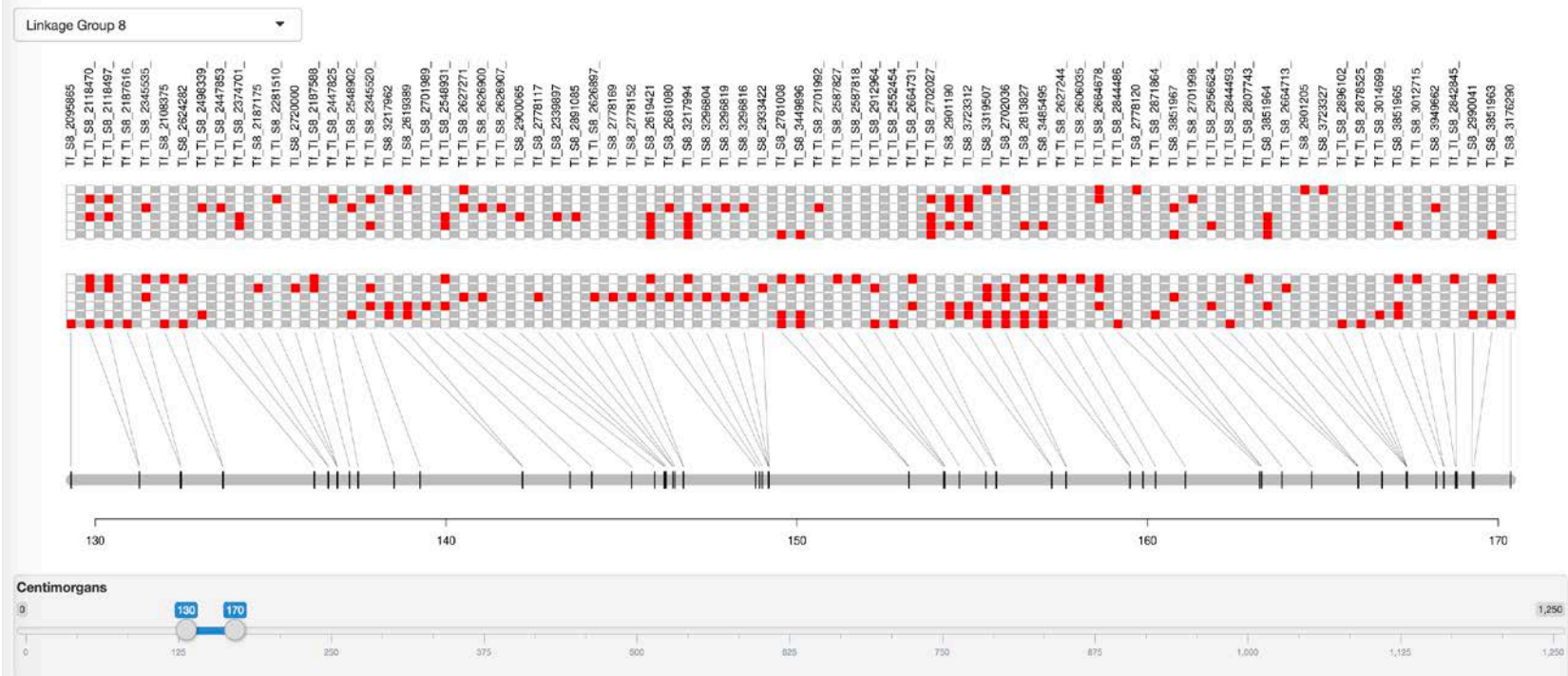


Numbers indicate the associated chromosomes in *I. trifida* and *I. triloba* reference genomes

Genetic mapping – Linkage group 8 – 873 SNPs

- Differently from several polyploid maps, we have an unified map which considers both parents at the same time.
- The map is constructed for the population, not for the parents. Also, it enables to detect QTLs considering both parents in the same model

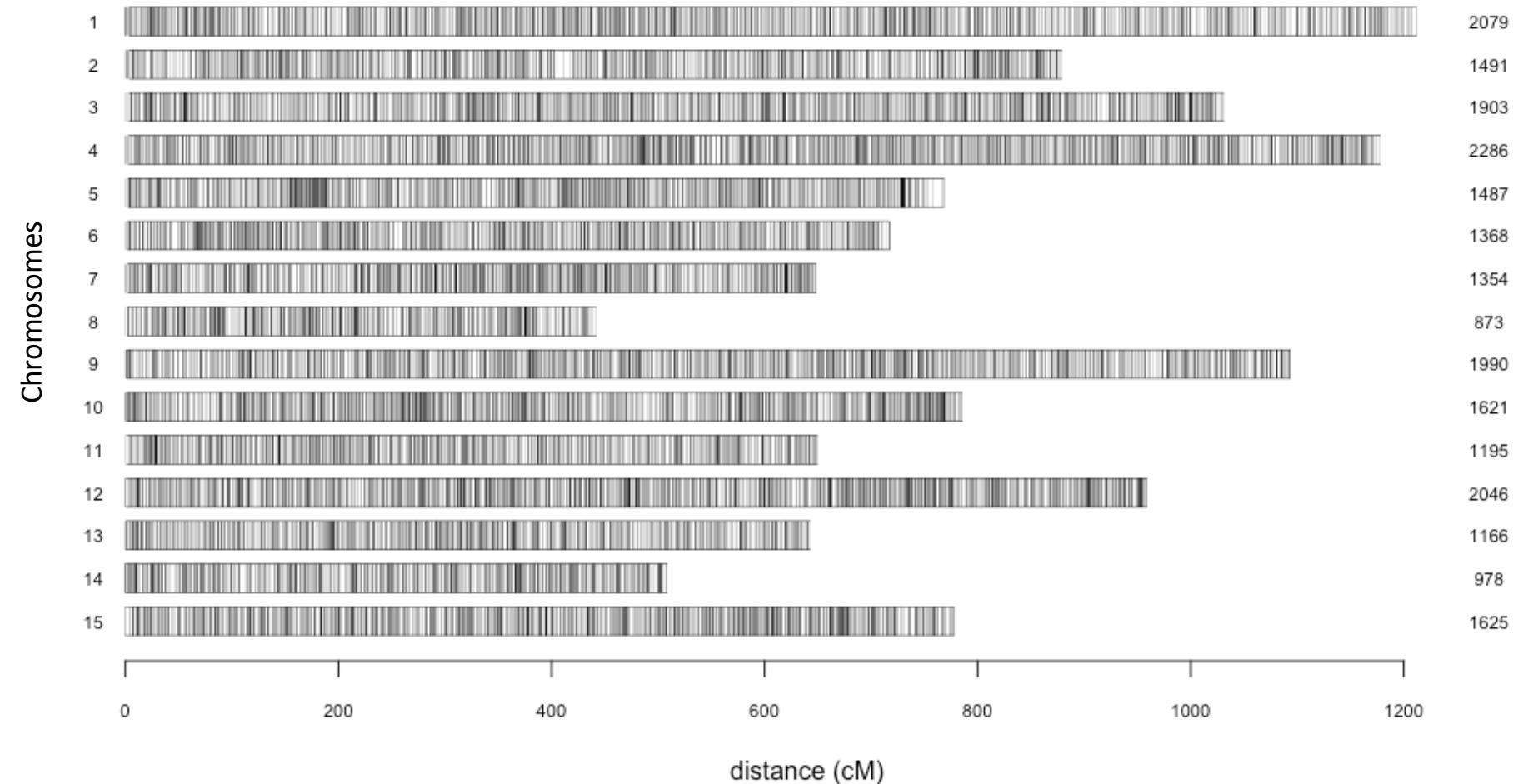
Sweetpotato genetic map - Beauregard x Tanzania (BT)



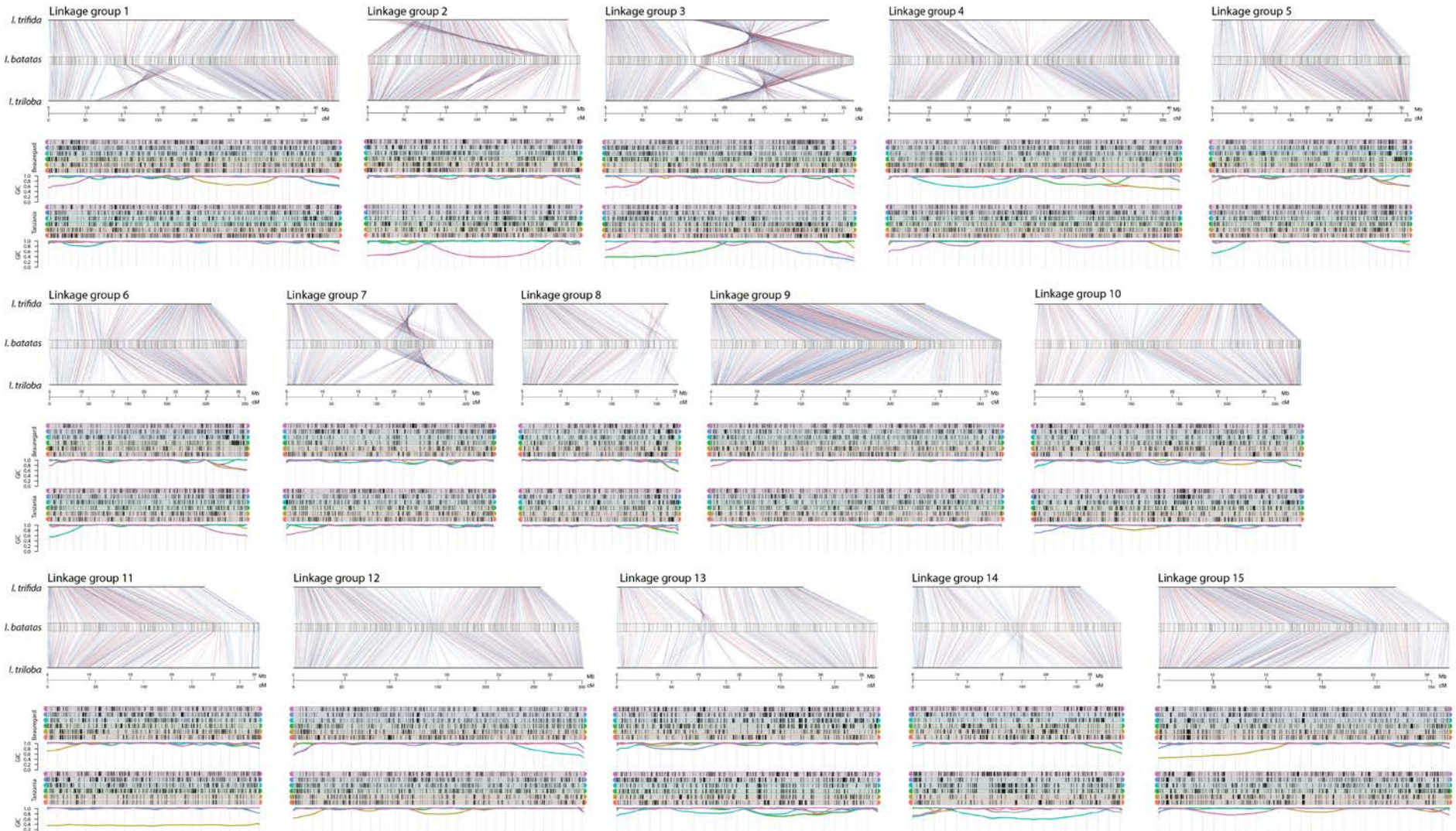
Interactive version: https://gt4sp-genetic-map.shinyapps.io/polymap_shiny/

Genetic mapping – 15 chromosomes – 23,462 SNPs

chromosome

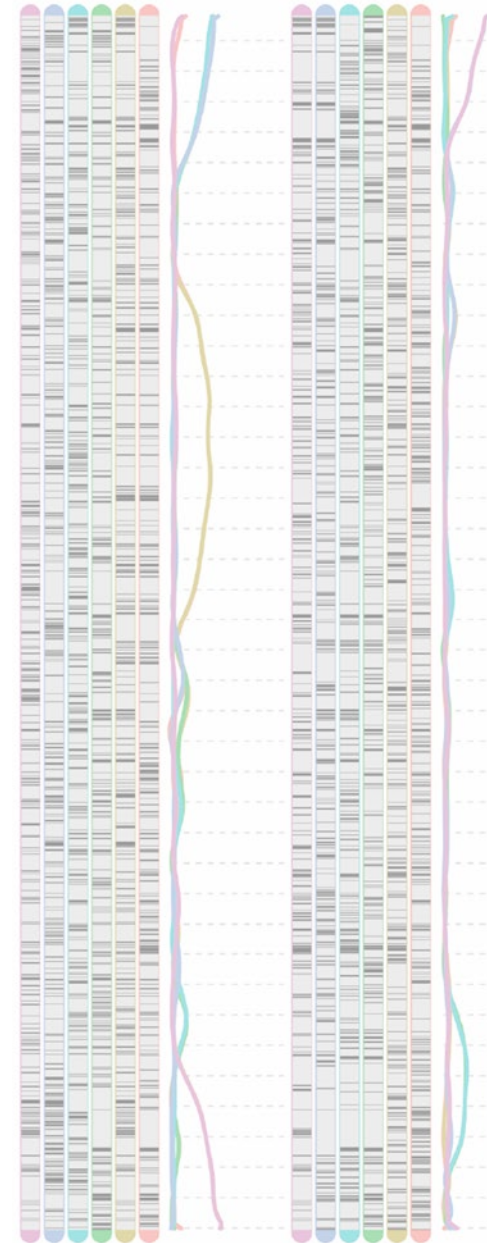


Sweetpotato map - 22,793 SNPs



Summary

- Use **adequate models** and software to construct maps in polyploids.
- **Dosage-based markers** (single dose + multiple dose) are essential in polyploid mapping
- Maps should be constructed **integrating both parents**
- Only maps **based in homology groups** will provide the adequate framework to QTL mapping



Demonstration in  Studio[®]