



Fine-mapping of a QTL segregating on pig chromosome 2 highlighted epistasis

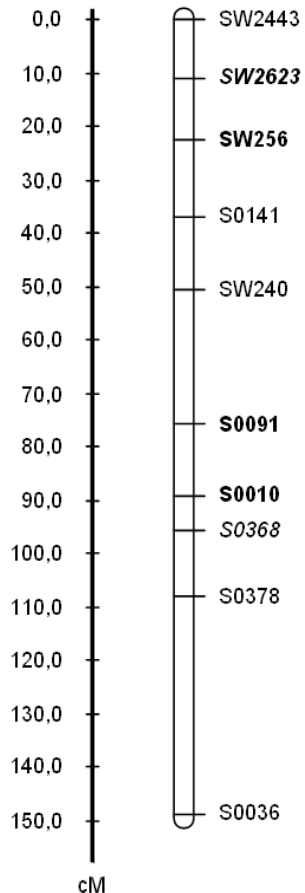
F. TORTEREAU, M.P. SANCHEZ, Y. BILLON, G. BURGAUD, M. BONNET,
J.P. BIDANEL, D. MILAN, H. GILBERT, J. RIQUET

Animal Breeding and Genetics Group, Wageningen University, The Netherlands
INRA Laboratoire de Génétique Cellulaire, Castanet-Tolosan, France
INRA GABI, Jouy-en-Josas, France
INRA GEPA Le Magneraud, Surgères, France



INRA

SSC2 and backfat thickness



IGF2 -intron3-G3072A (Van Laere *et al.*, 2003)

BUT not only explanation (Sanchez *et al.*, 2006)

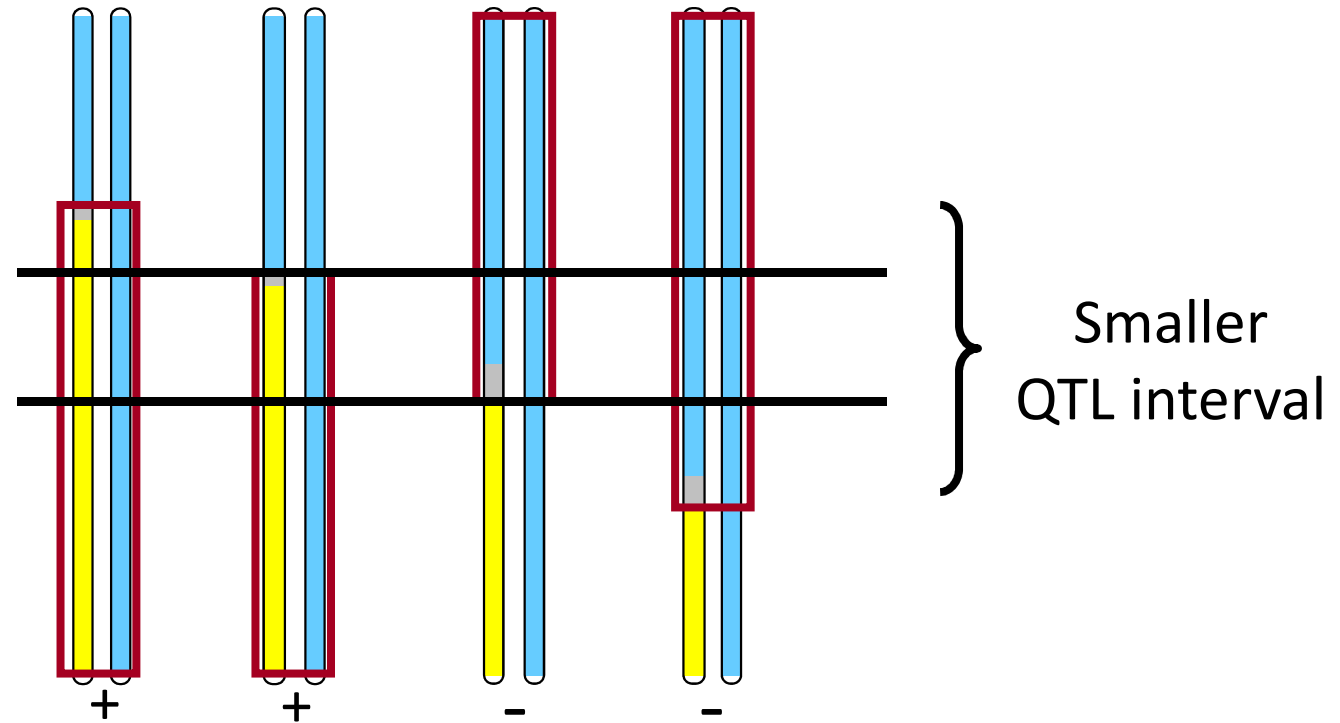
Reports of other QTLs

(Lee *et al.*, 2003 ; Geldermann *et al.*, 2010 ; Tortereau *et al.*, 2011)

Confirmation, fine-mapping and description of its mode of segregation

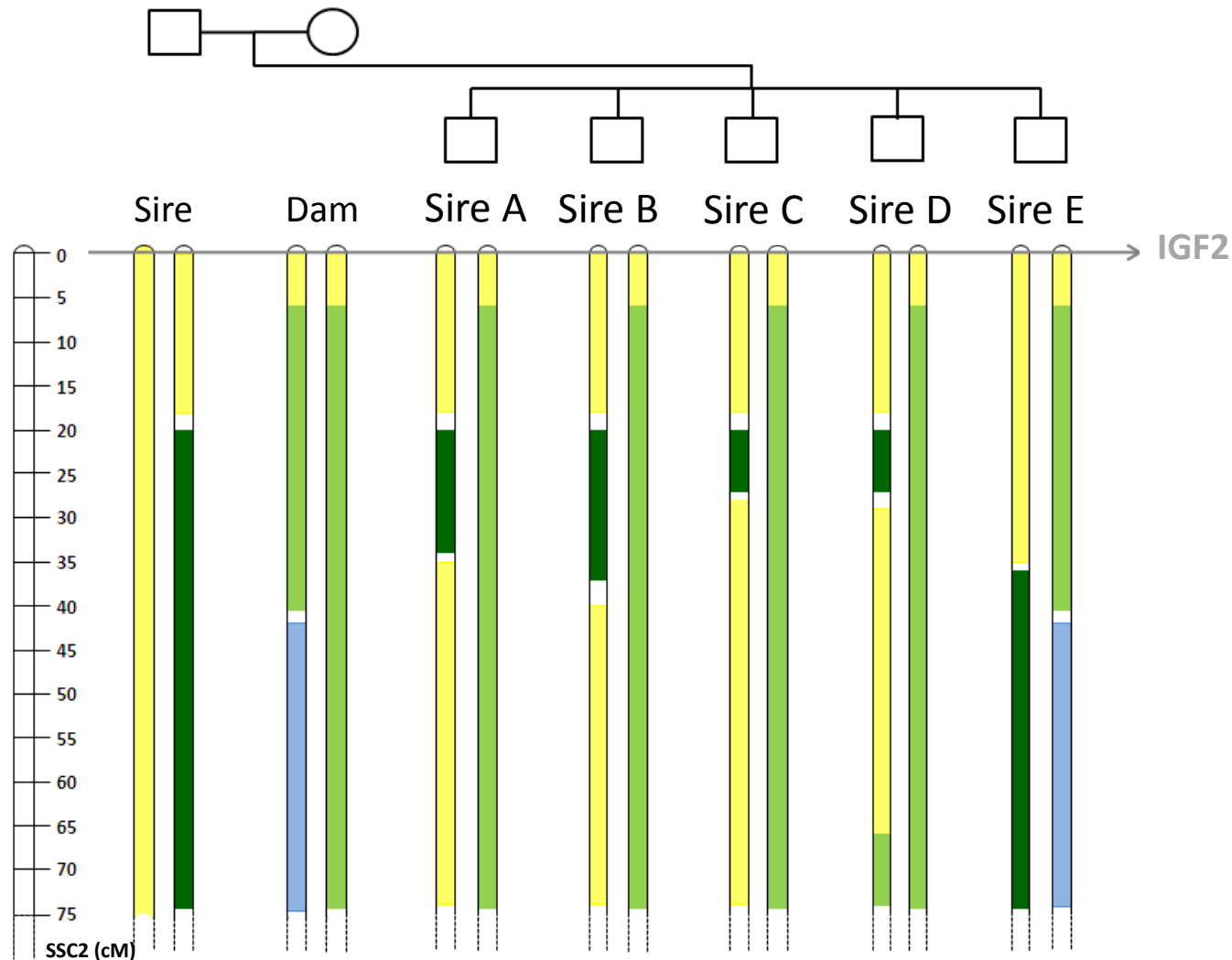
Marker-assisted backcross design

Creation of animals carrying different recombination points in the candidate region



QTL segregation

Tested sires



Only G/G animals,
to avoid the
segregation of the
mutation

By construction :

- A unique **IBD MS** haplotype
- Various **LW** haplotypes

Progeny-testing results

Progeny testing with LW females,
about 100 offspring per sire

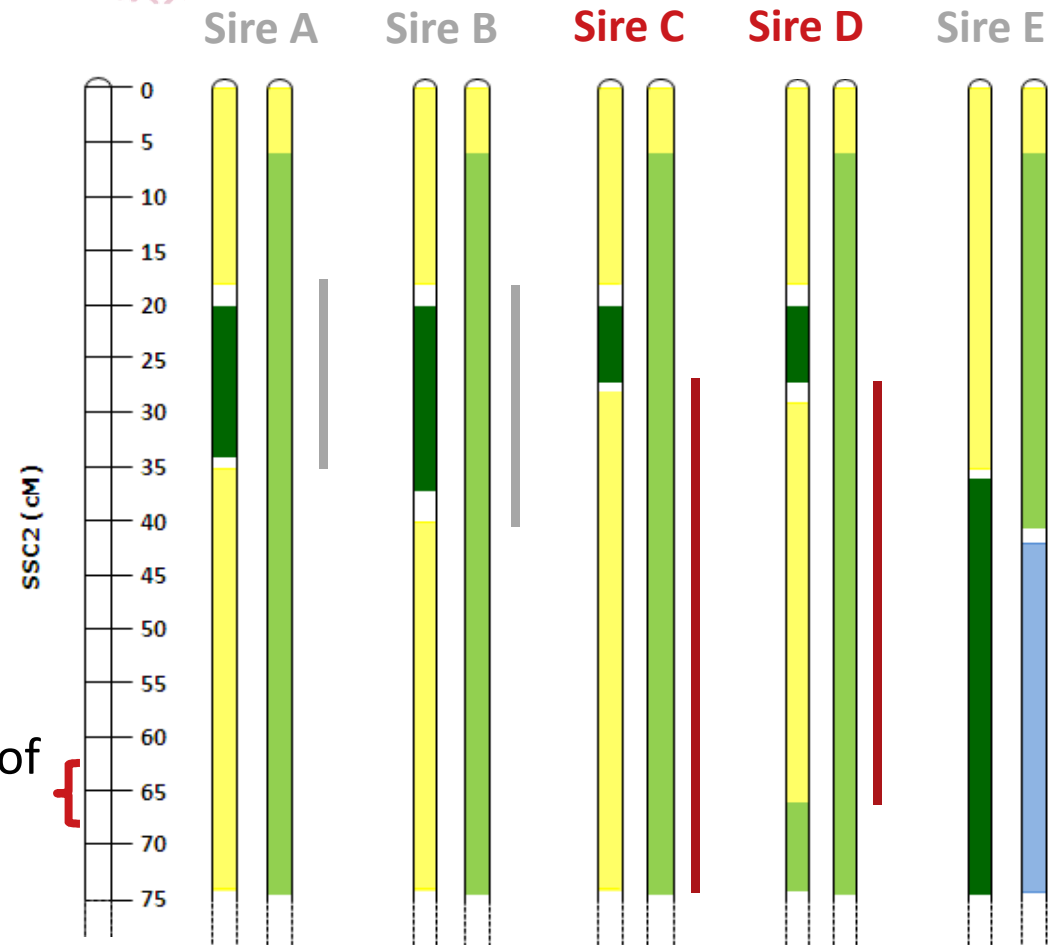
Results with US BFT measurements :

- 2 segregating sires

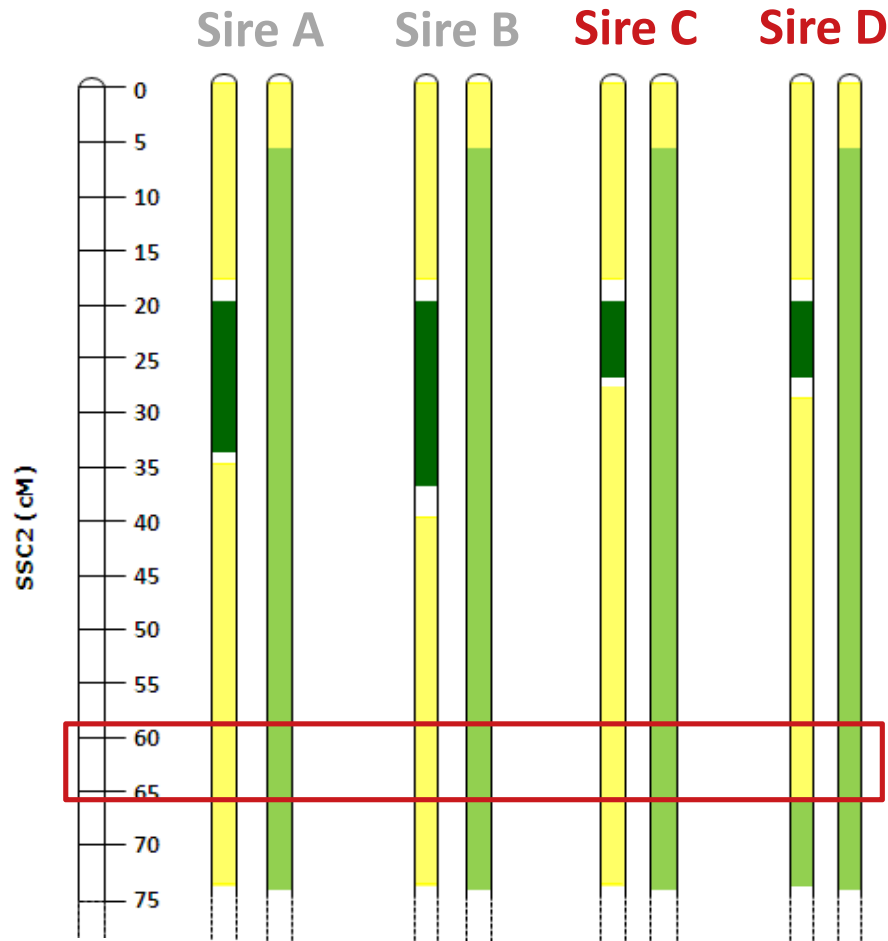
-3 non-segregating sires

No overlapping region ...

Most likely position of
the detected QTLs



Full-sibs with identical contrasts



At the most likely position:
same contrasts between
two IBD haplotypes



Interaction with another
locus in the genome

Whole genome scan

Genotyping of 578 microsatellites covering all the autosomes and SSCX

Selection of candidate microsatellites :

	Interaction candidate region			Genotype		
	Marker	Chromosome	Position	All1	All2	
Sire A	S0155	1	95	4	2	Excluded
Sire B	S0155	1	95	4	2	
Sire C	S0155	1	95	4	1	
Sire D	S0155	1	95	4	2	
Sire A	SW2551	1	96	1	1	No informative
Sire B	SW2551	1	96	1	1	
Sire C	SW2551	1	96	1	1	
Sire D	SW2551	1	96	1	1	
Sire A	SW2512	1	144	9	10	Identical genotypes
Sire B	SW2512	1	144	9	10	
Sire C	SW2512	1	144	7	10	Identical genotypes
Sire D	SW2512	1	144	7	10	

Different genotypes

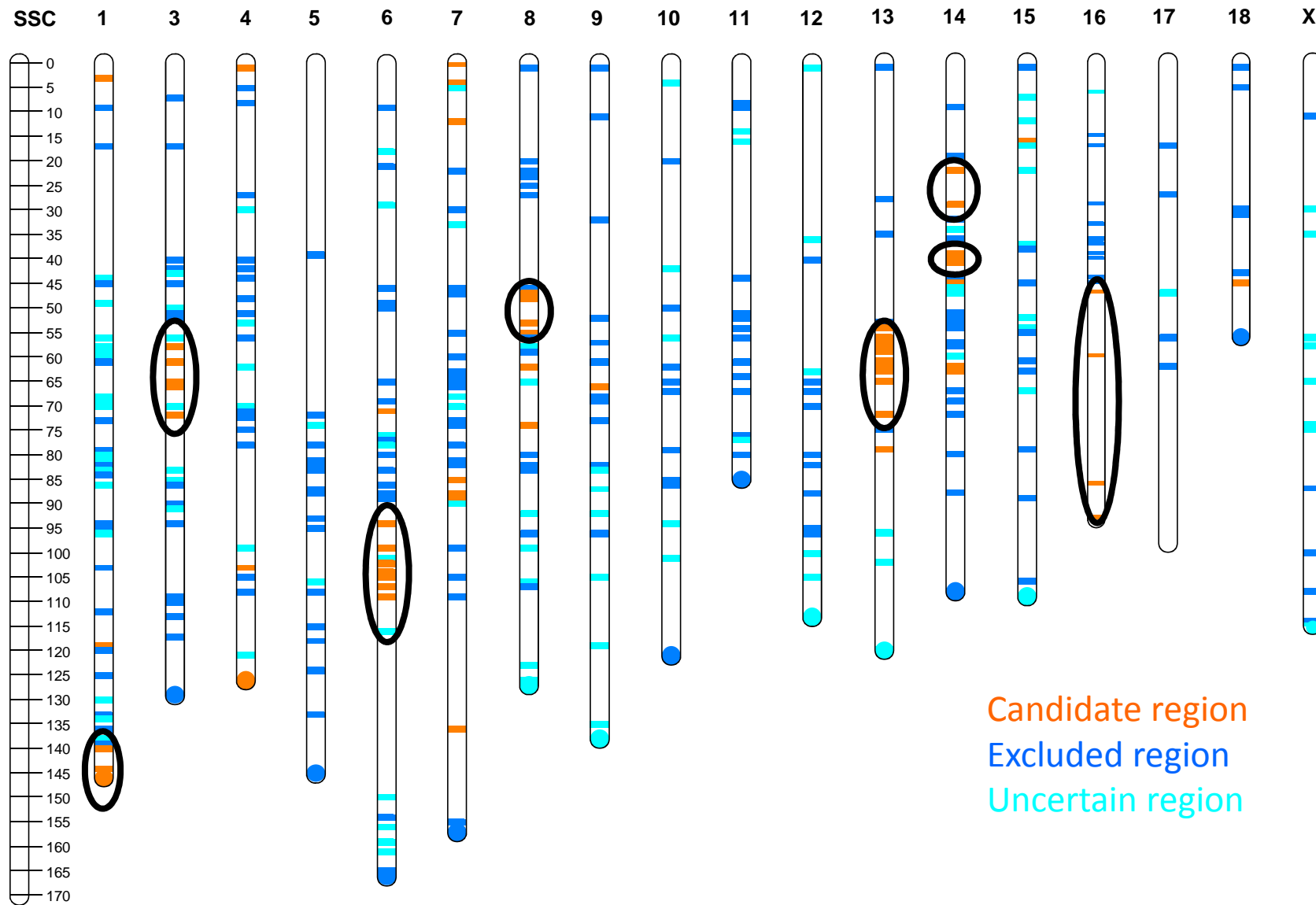
Whole genome scan

Genotyping of 578 microsatellites covering all the autosomes and SSCX

Selection of candidate microsatellites :

	Interaction candidate region			Genotype		
	Marker	Chromosome	Position	All1	All2	
Sire A	S0155	1	95	4	2	Excluded
Sire B	S0155	1	95	4	2	
Sire C	S0155	1	95	4	1	
Sire D	S0155	1	95	4	2	
Sire A	SW2551	1	96	1	1	No informative
Sire B	SW2551	1	96	1	1	
Sire C	SW2551	1	96	1	1	
Sire D	SW2551	1	96	1	1	
Sire A	SW2512	1	144	9	10	Identical genotypes
Sire B	SW2512	1	144	9	10	
Sire C	SW2512	1	144	7	10	Different genotypes
Sire D	SW2512	1	144	7	10	

Candidate regions



Detection of interactions

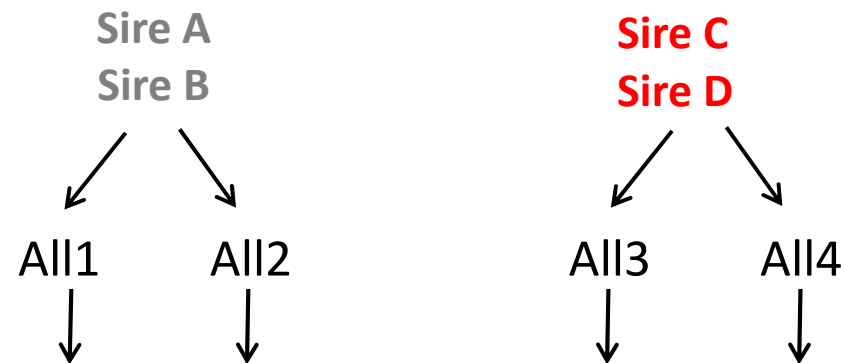
Selection of 1 or 2 microsatellite per candidate region

Genotyping of all the offspring of the 4 sires

Interaction analysis :

group of sires

alleles at the candidate
interacting region



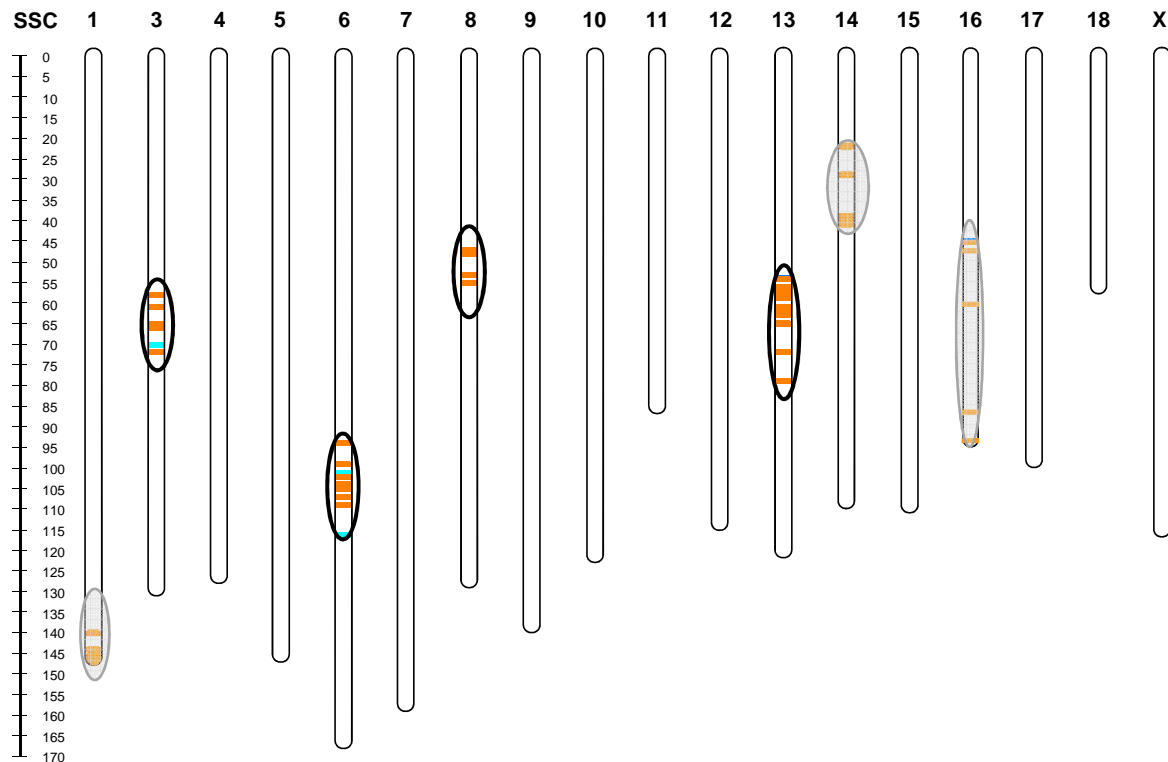
Detection of a QTL on SSC2 ?

Results

Among the 7 tested regions:

3 have been excluded

4 remained candidate interacting regions



Analysis of a F2 design

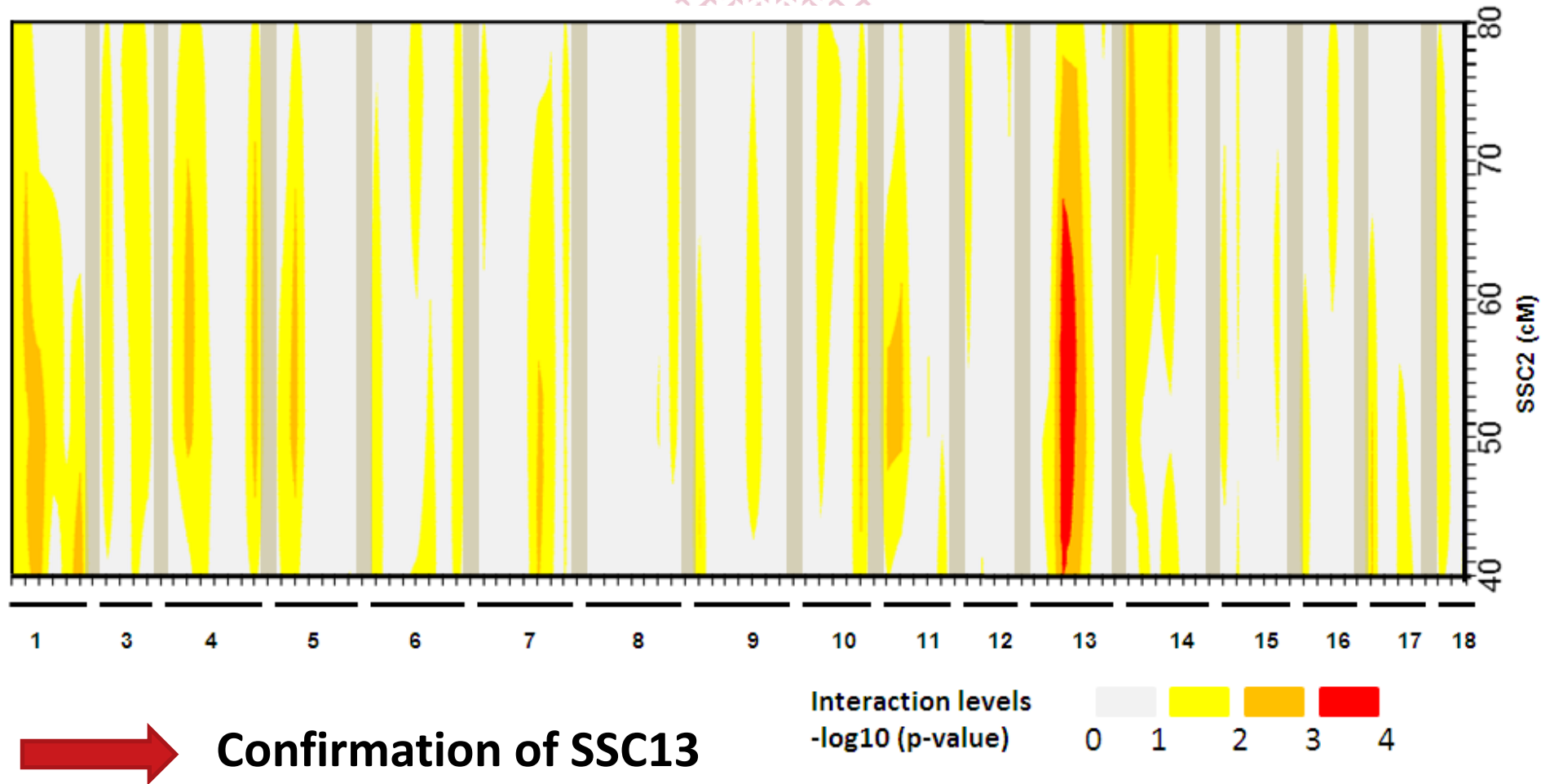
- PORQTL F2 design (Bidanel et al., 2001),
same breeds
ancestor of the BC pedigree

- Interaction analyses :

SSC2 QTL region

~ 100 microsatellites covering all the autosomes

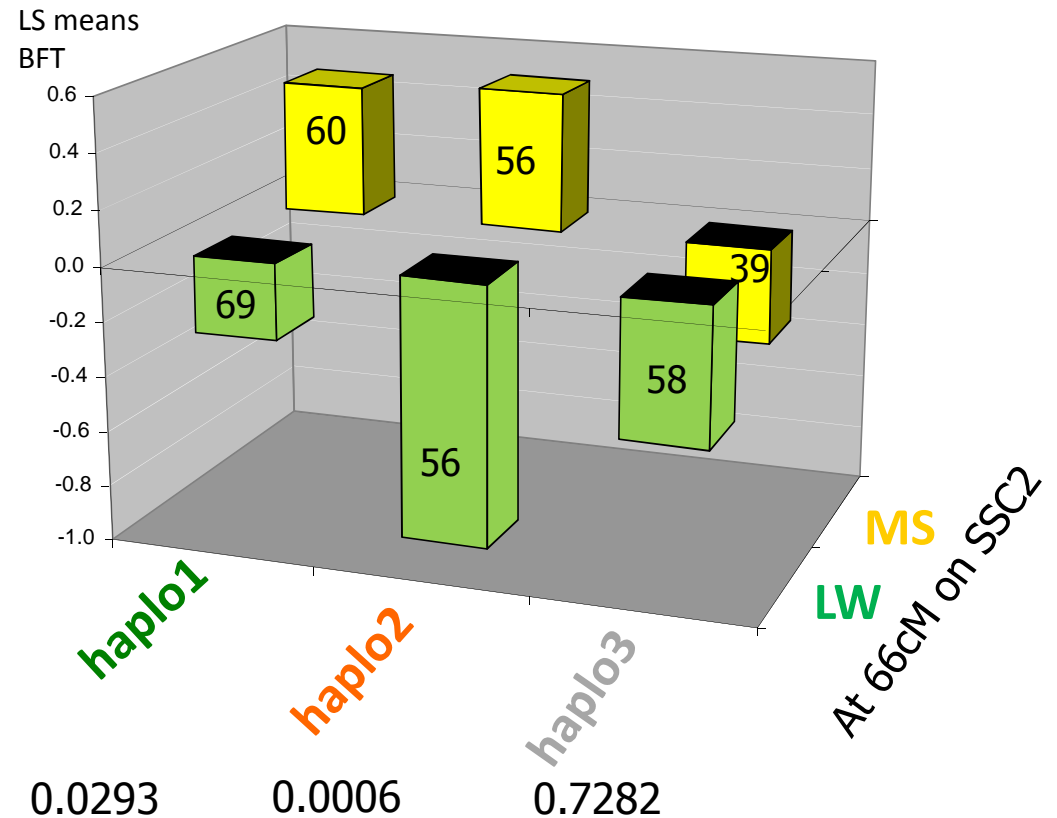
Confirmation in a F2 design



Interaction SSC2 – SSC13

Analysis of the SSC13 region in the BC pedigree:

Sire A haplo1 / haplo3
Sire B haplo1 / haplo3
Sire C haplo1 / haplo2
Sire D haplo2 / haplo2



Conclusion

- QTL underlying fatness traits on SSC2: 37 – 67 cM
- Interactions with at least SSC13
- Interactions with SSC3, SSC6, SSC8 to be confirmed / invalidated



Thank you for your attention

Conclusion

- QTL underlying fatness traits on SSC2: 37 – 67 cM
- Interactions with at least SSC13
- Interactions with SSC3, SSC6, SSC8 to be confirmed / invalidated