



1ST EUROPEAN SORGHUM CONGRESS

WORKSHOP

INNOVATIVE RESEARCH TOWARDS GENETIC PROGRESS

**TACKLING NEW CHALLENGES FOR
EUROPEAN SORGHUM THROUGH
GENETICS
AND NEW BREEDING STRATEGIES**



**OPTIMIZING GENETIC VALUE PREDICTION USING MOLECULAR
MARKERS**



David Pot, PhD, CIRAD FRANCE

BUCHAREST
3-4 NOVEMBER 2016

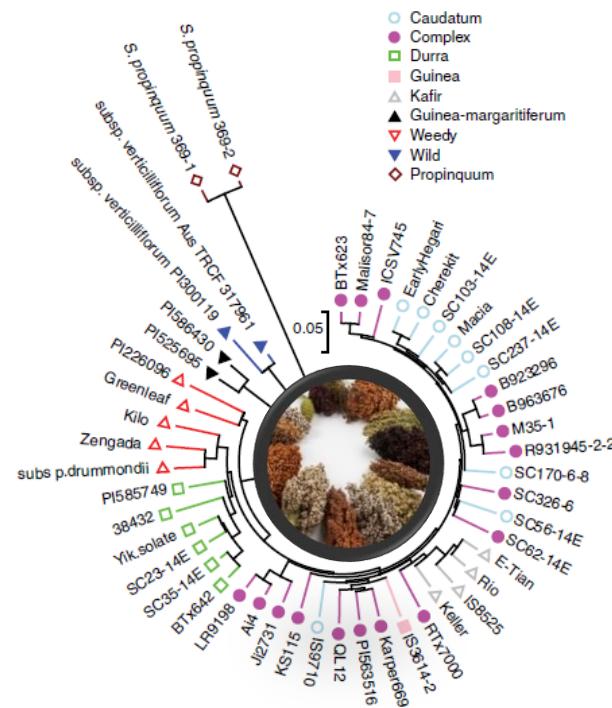
WHOLE GENOME SEQUENCING : AN ACCELERATOR OF SORGHUM GENETICS AND BREEDING

Btx623
sequencing



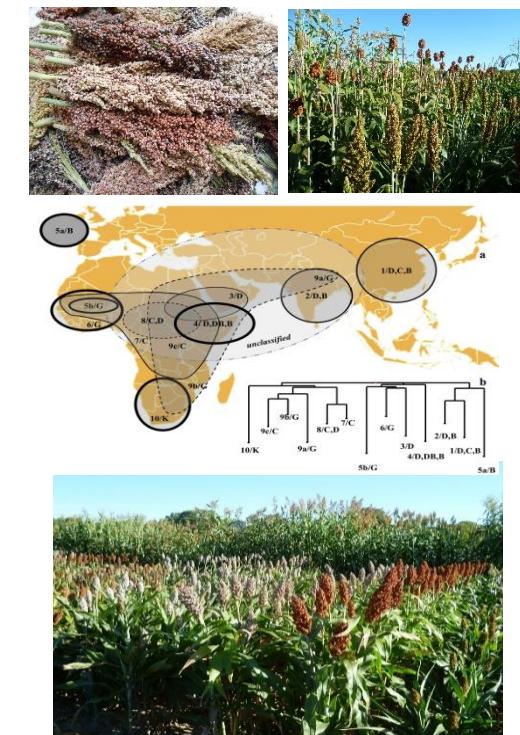
Paterson et al 2009

Enlarging the diversity coverage



Mace et al 2013

To 1000's Genomes :
« Terra » Project

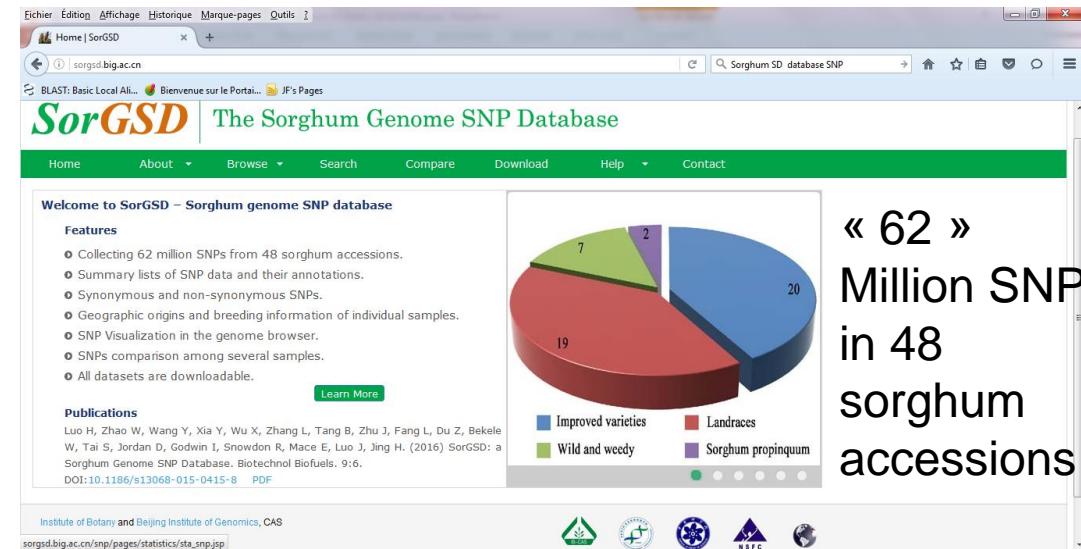


Mockler et al 2016

NUCLEOTIDE DIVERSITY INFORMATION

Btx623 : ATGCATGCATGC
 Keller : ATGCCCTGCCCC
 Tx430 : ATGCATGCATGC
 Ji2731 : ATGCCCTGCATGC
 E-Tian : AAGCATGCATGC
 Rio : AAGCCTGCATGC





Luo et al 2016

- Polymorphism Database and **Next Generation Sequencing (NGS) methodologies**
 - Characterize the diversity (Jeff's talk)
 - Support breeding efforts (**Marker ASSISTED breeding**)

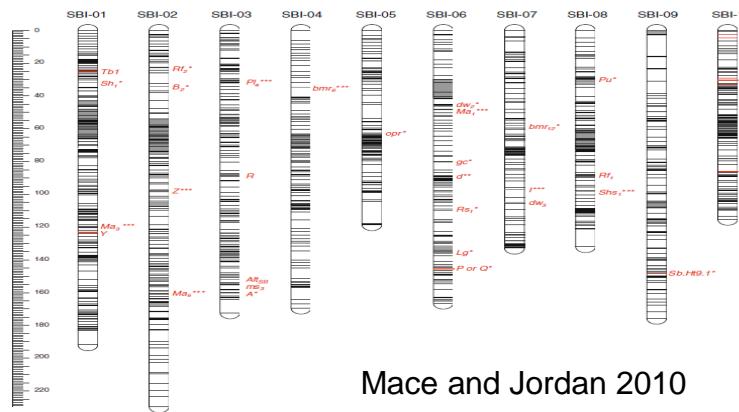


TAKING ADVANTAGE OF GENETIC INFORMATION TO ACCELERATE GENETIC GAINS (PREDICT PHENOTYPES)

- Response to selection: $R_x = I_x * h_x^2 * Sd_x$ (x = grain yield in low Inputs)
With I_x : Selection intensity applied on trait $_x$
 h_x^2 : heritability of trait $_x$,
 Sd_x : Phenotypic standard deviation of trait $_x$
- Indirect selection (select trait $_x$ based on selection on trait $_y$ (y = grain yield in on-station trial)):
 $CR_x = i_y * h_x * h_y * r_{gxy} * Sd_x$
With r_{gxy} : genetic correlation between traits x and y
- Molecular markers allow maximizing h_y . Assuming a high genetic correlation (r_{gxy}) you can expect $CR_x >> R_x$
- Challenge : identify the genomic regions (Markers) controlling the traits of interest (high r_{gxy}) and combine them in new varieties

MAJOR GENES AND QTLS (BIPARENTAL) HAVE ALREADY BEEN IDENTIFIED

Major Genes

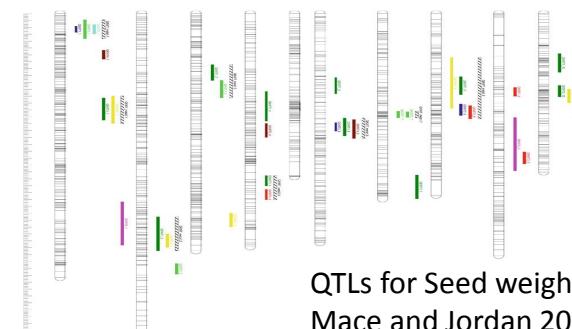


Mace and Jordan 2010

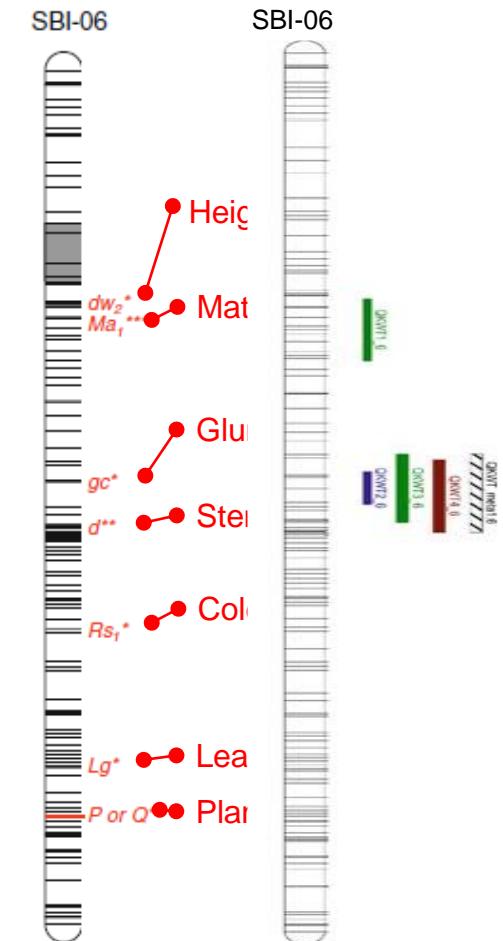
Some genomic regions affecting quantitative traits

QTLS in biparental populations

- Grain yield / size...
- Grain quality
- Biomass yield
- Biomass quality
- Biotic and Abiotic stress

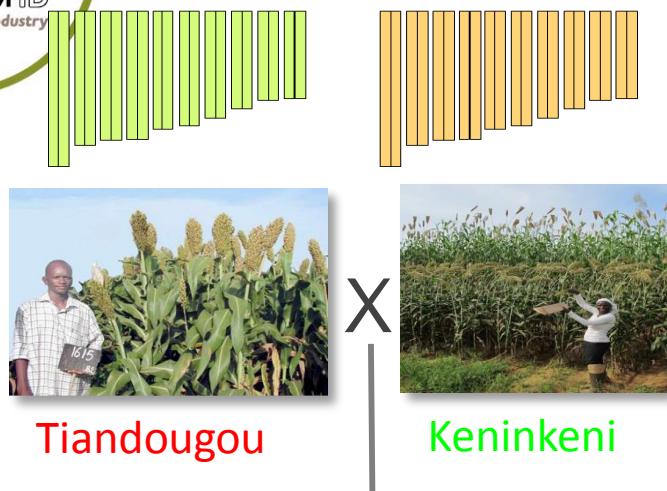


QTLS for Seed weight,
Mace and Jordan 2011

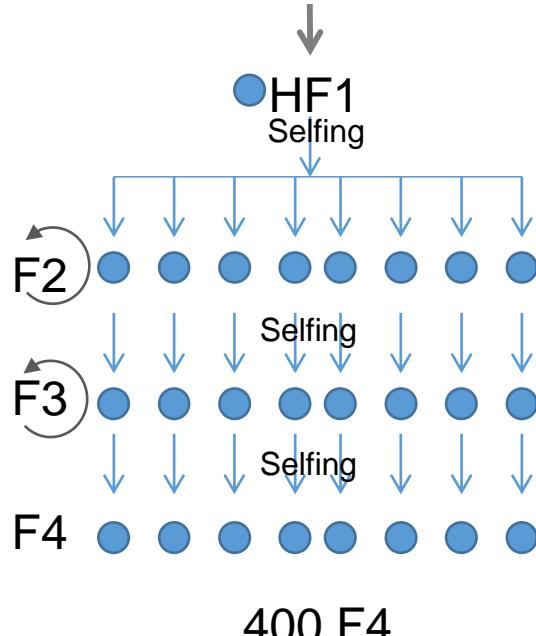




MARKER ASSISTED RECURRENT SELECTION APPLIED TO WITHIN FAMILY BREEDING FOR GRAIN YIELD AND QUALITY IN MALI



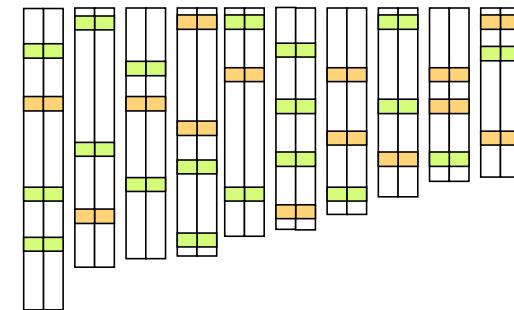
- Solution : **identify** the genomic regions / favorable alleles and combine them



June 2010

- Multi-trait selection
- Combine favorable alleles from the 2 parents

- What we want : combine « 33 » regions



- Not reachable through genealogic / phenotypic selection : million of progenies would be required...



WITHIN FAMILY BREEDING : DEFINING THE MOLECULAR IDEOTYPE



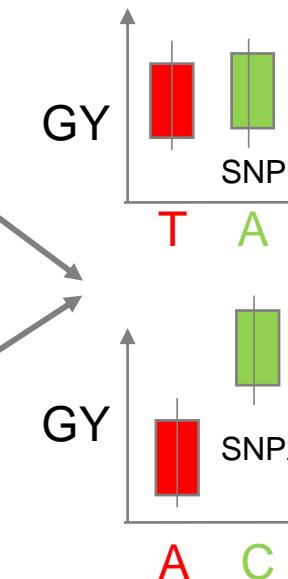
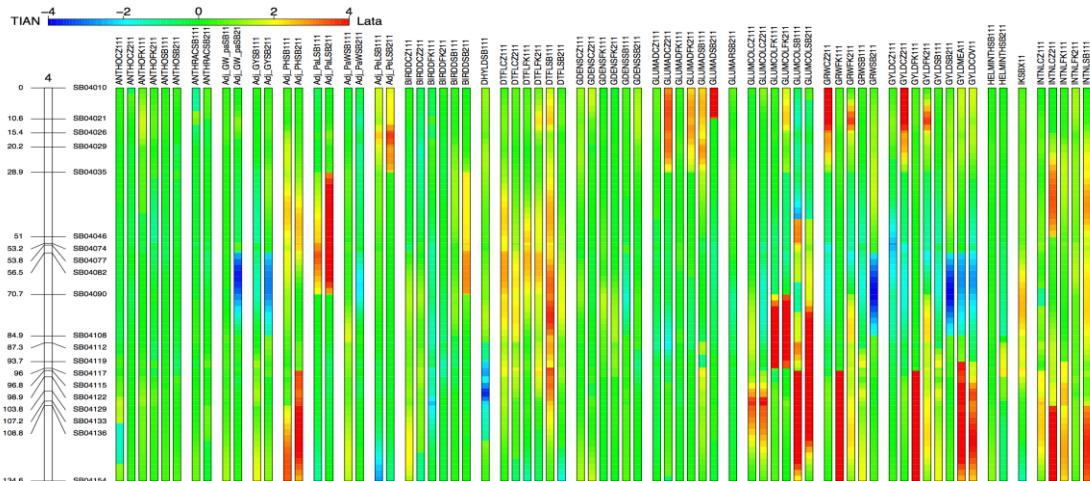
400 F4

MET Phenotyping (1/2011)

	GY	PH
P1	: 5	130
P2	: 6	170
F4.1	: 2	175
F4.2	: 5	140
F4.3	: 3	120
F4.n	: 8	180

P1	: ATGCATGC
P2	: AAGCCTGC
F4.1	: ATGCATGC
F4.2	: ATGCCTGC
F4.3	: AAGCATGC
F4.n	: AAGCCTGC

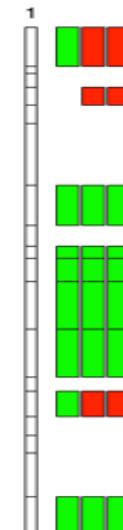
SNP1 SNP2



No effect

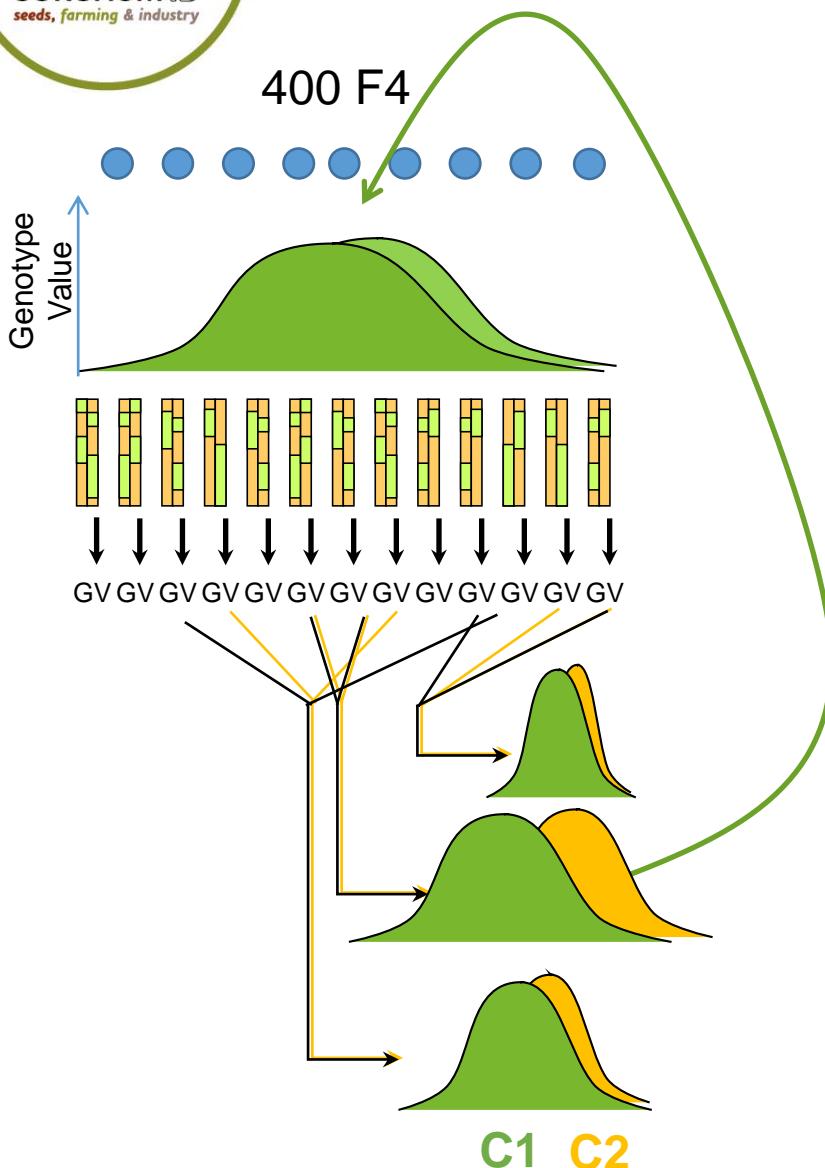
Effect

- Molecular target definition

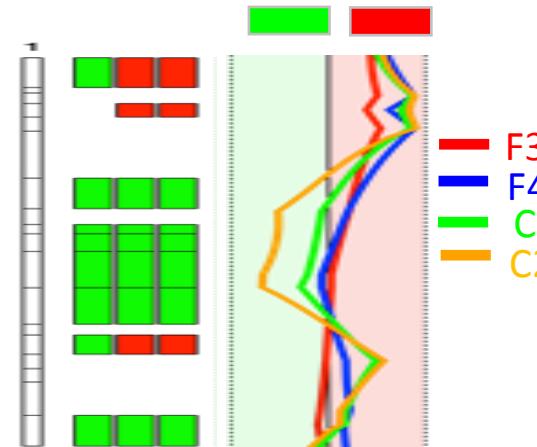




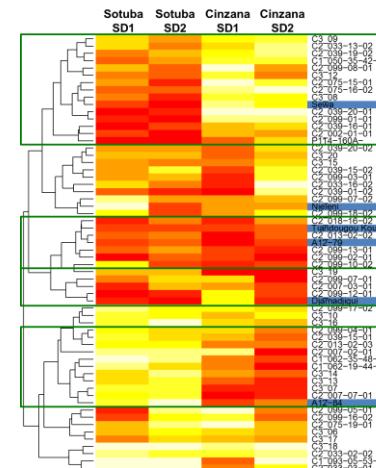
WITHIN FAMILY BREEDING : CONVERGING TOWARDS THE MOLECULAR IDEOTYPE



- Evolution towards molecular ideotype (6/2011 – 1/2013)



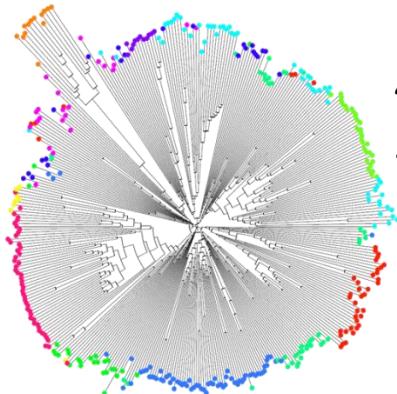
- Field evaluation (On-Station and on-Farm)



- Comparison with local varieties
- Within family MARS works (faster, more accurate)
- But information is not easily transferred to broad base populations



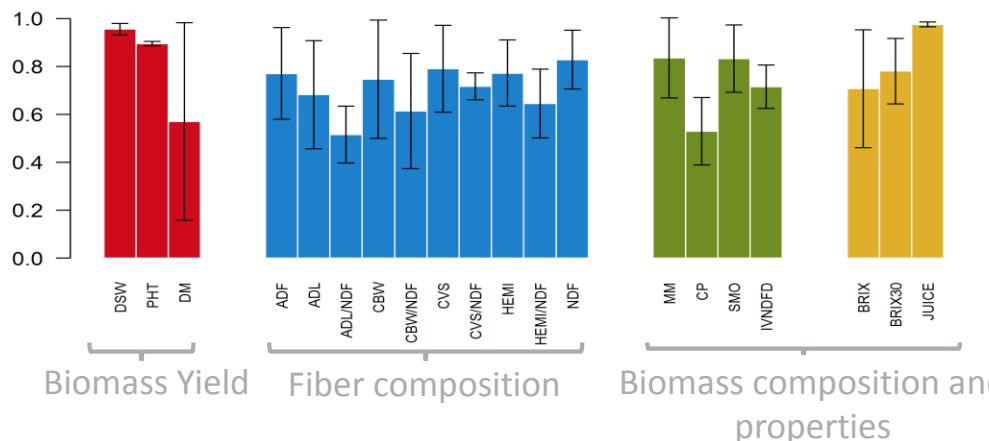
Multi-Environments / Panels



413 accessions
 5 trials (100 - 362 genotypes)
 • 2 Montpellier
 • 2 Mali
 • 1 Mali Off Season



Heritabilities



GBS : 190ksnp

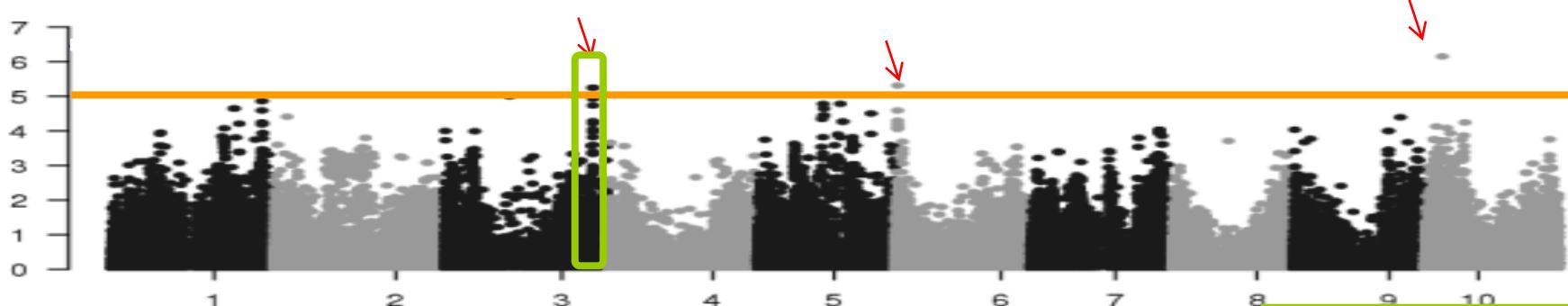
Btx623	:	ATGCATGCATGC	
Keller	:	ATGCC <color>CT</color> TGC	C
Tx430	:	ATGCATGC	C
Ji2731	:	ATGCC <color>CT</color> TGCATGC	
E-Tian	:	AAGCATGCATGC	
Rio	:	AAGCC <color>CT</color> TGCATGC	

SNP Indels

GWAS : GENE IDENTIFICATION, BUT NEEDS FOR MULTI-ENVIRONMENTS ASSESSMENTS

GWAS for Cell Wall lignin content

$-\log_{10}(p)$



Chrom	Pos(pb)	LOD	Gènes candidats		
			Début	Fin	Fonction
3	67 247 746	5.25	67 242 057	67 244 587	Phenylpropanoids /lignin
6	1 273 618	5.31	1 208 904	1 216 444	TF/TF MYB
10	7 794 293	6.15	7 751 084	7 755 352	TF/TF MADS

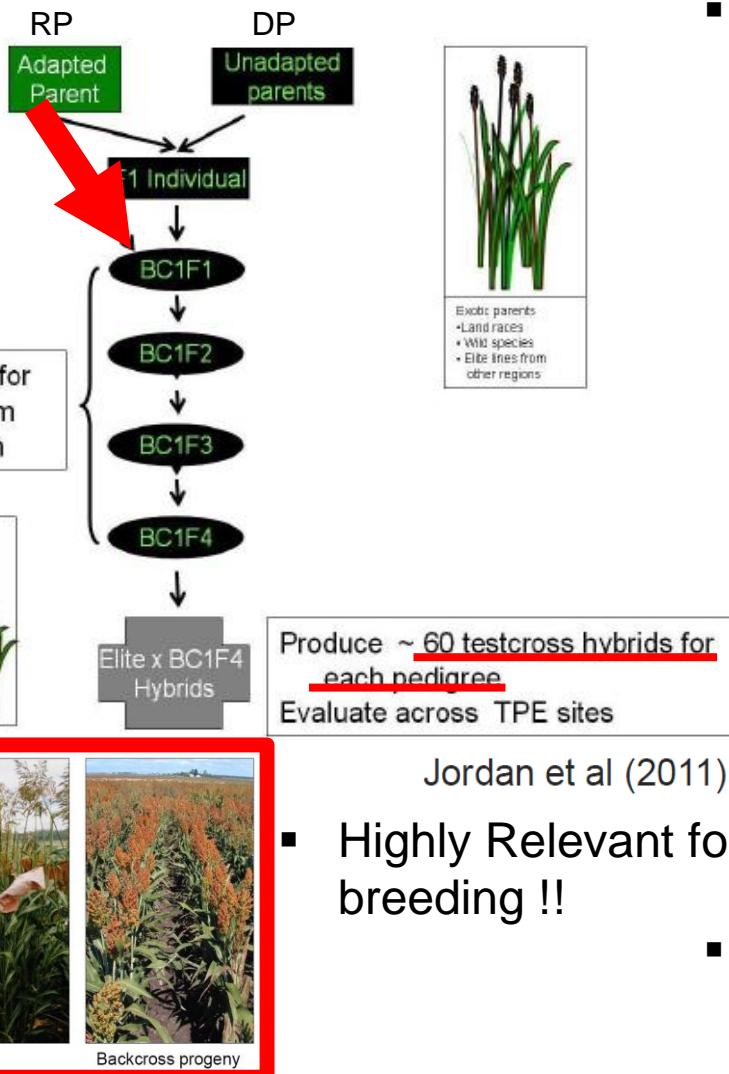
LACCASE Is Necessary and Nonredundant with PEROXIDASE for Lignin Polymerization during Vascular Development in *Arabidopsis*^{CH}

Qiao Zhao,^a Jin Nakashima,^a Fang Chen,^a Yanyan Yin,^b Chunxiang Fu,^c Jian Zeng-Yu Wang,^c and Richard A. Dixon^{a,2}

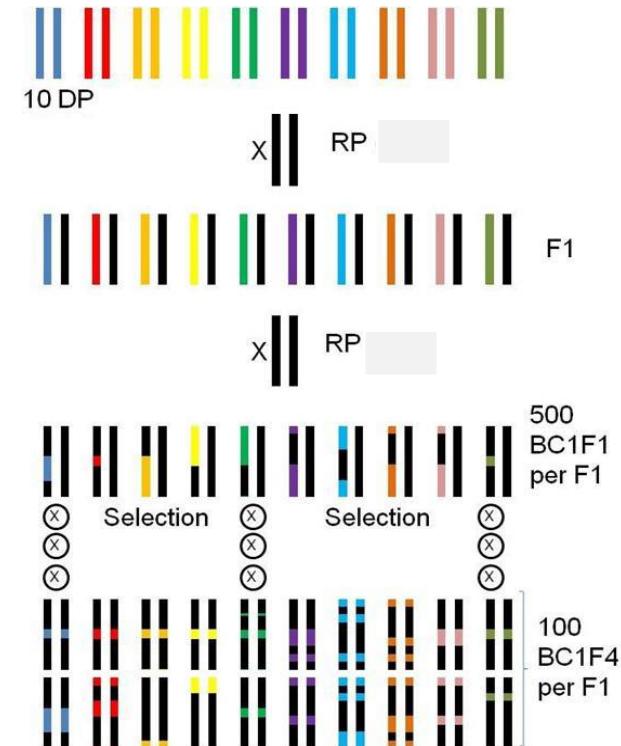


- Refined genomic regions / genes identification
- GXE + Transfer to breeding programmes (exotic lines...) + Power issues (allelic frequencies)

MERGING GENETIC AND BREEDING OBJECTIVES FOR BROAD BASED PANELS : COMBINING BIPARENTAL AND GWAS MAPPING



- And high resolution genetic mapping

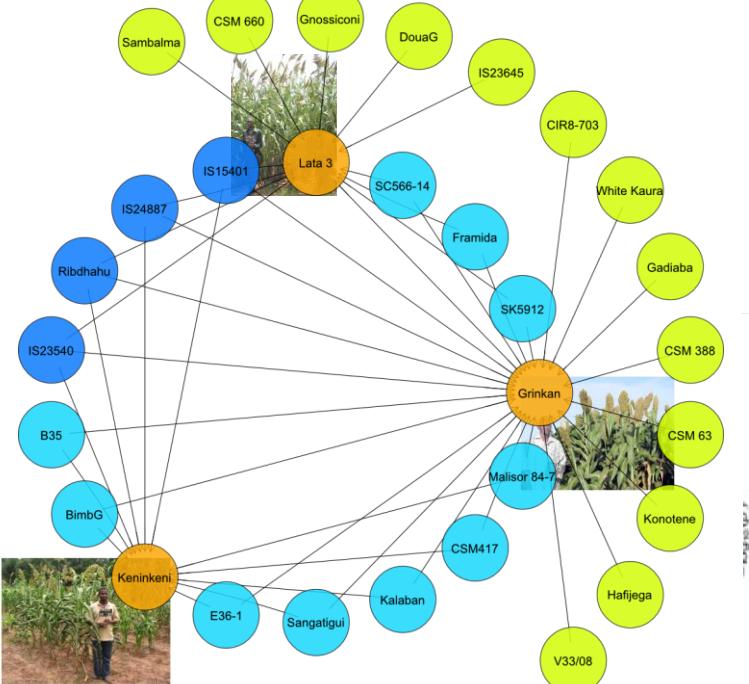


Jordan et al (2011).

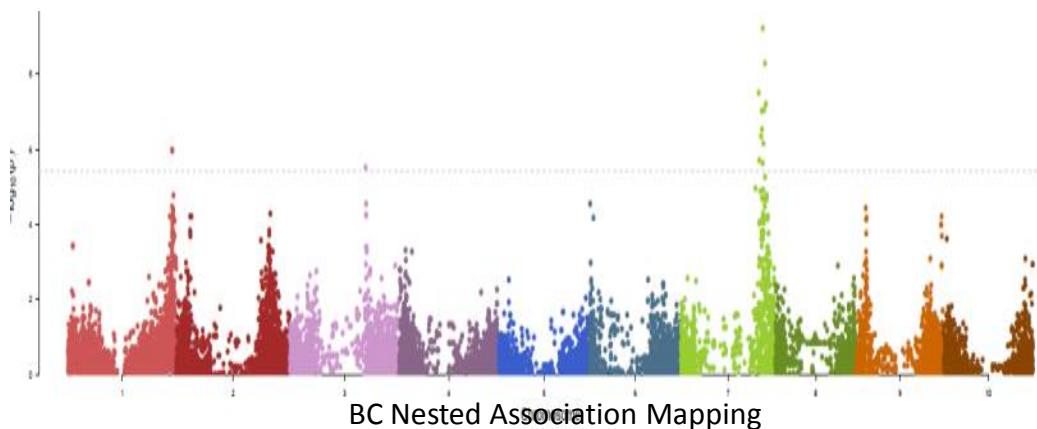
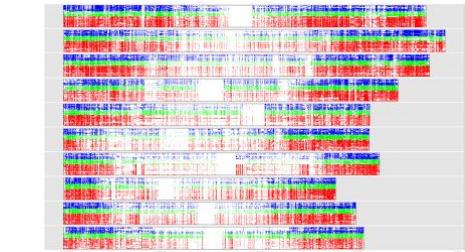
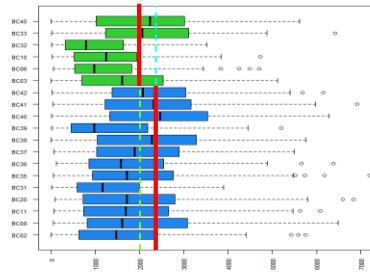
- Highly Relevant for breeding !!
- See also Morris et al 2015 : NAM for Btx623 and Tx430



BC-NAM : GRAIN YIELD AND QUALITY FOR WESTERN AFRICA



47 populations, 4717 BC₁F₄ families



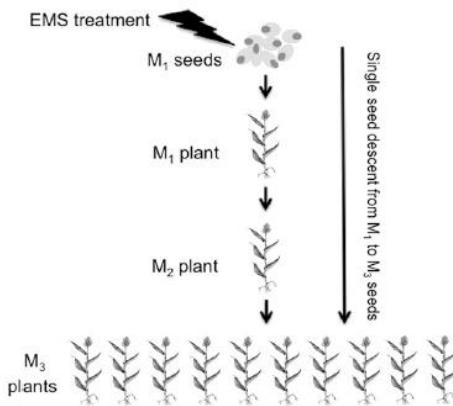
Direct variety development through PPB

Nom	Pedigree	Dicila	Koutiala	Mande	MOYEN	Rang
Fadda	12A/Lata Hybrid Check	94	242	99	138	1
Lagri	Lata//Grin-8-39-1-1	87	193	92	120	3
Essi	Lata//SC566-6-44-1-1	102	172	89	122	2
Lango	Lata/Ngol-3-6-1-1	89	150	77	104	11
Tieble	CSM335 Local check	72	148	75	100	14
Djala	Lata//DouaG-1-2-1-1	96	172	71	110	7
Lani	Lata//Gnos-7-13-1-1	81	194	67	107	8
Samboni	Lata//Samb-5-1-1-1	82	196	67	106	9
Dili	Lata/Ridb-3-9-1-1	94	183	57	102	13



CREATING NEW VARIABILITY THROUGH CHEMICAL MUTATIONS

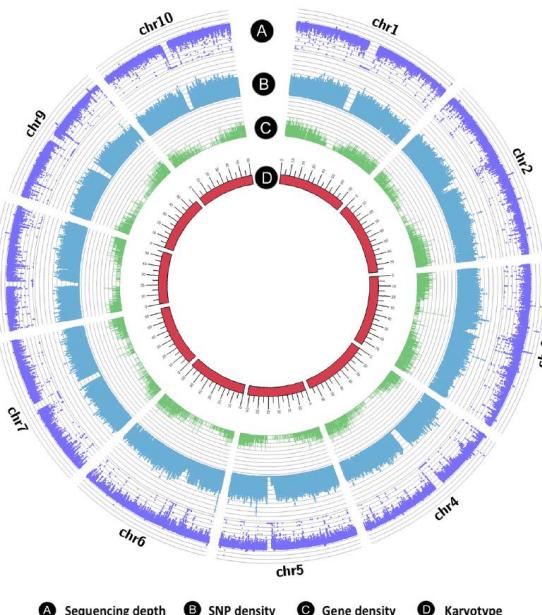
EMS Mutant libraries development



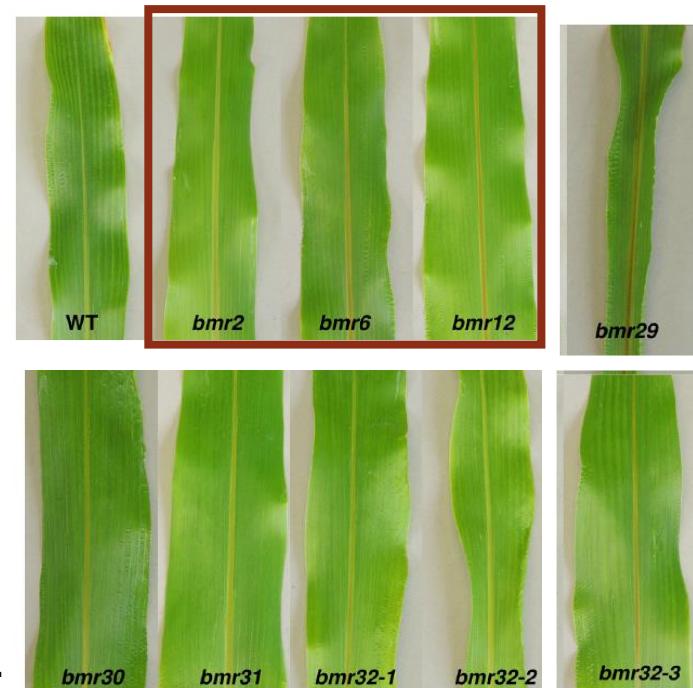
Jiao et al 2016 : 6400 M4 mutants



256 mutants lines sequenced.
A ressource for gene validation



Identification of mutants relevant for breeding

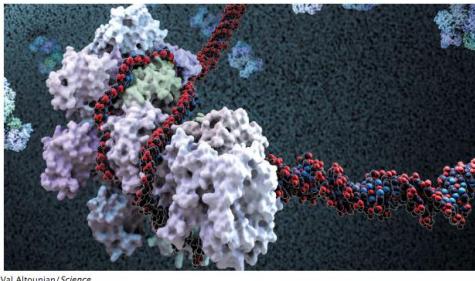


Sattler et al 2014

CREATING NEW VARIABILITY THROUGH GENOME EDITING

SPECIAL COLLECTION

The CRISPR Revolution



Val Altounian/Science

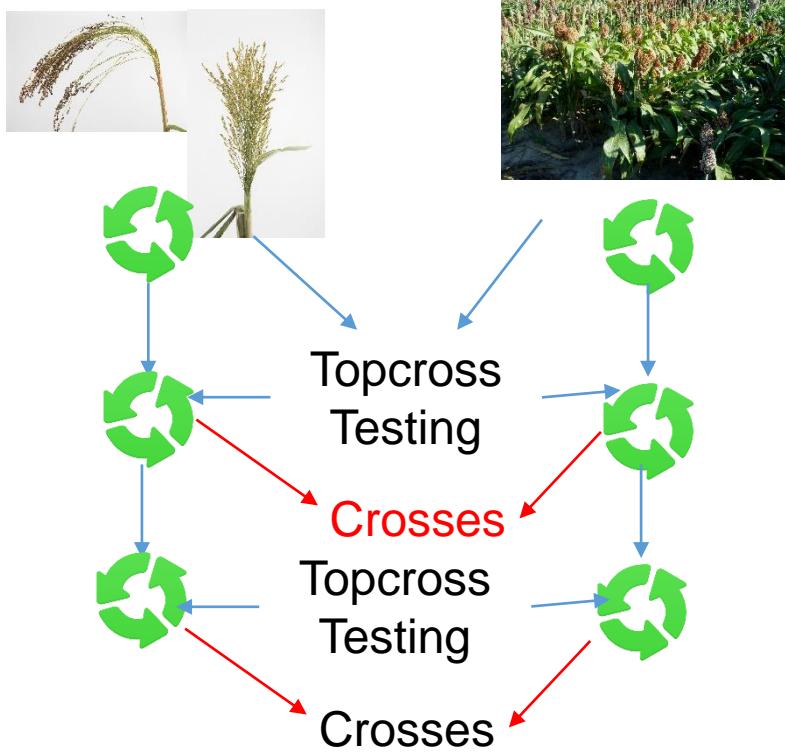
Demonstration of CRISPR/Cas9/sgRNA-mediated targeted gene modification in *Arabidopsis*, tobacco, sorghum and rice

Wenzhi Jiang¹, Huanbin Zhou², Honghao Bi², Michael Fromm³, Bing Yang² and Donald P. Weeks^{1,*} 2013

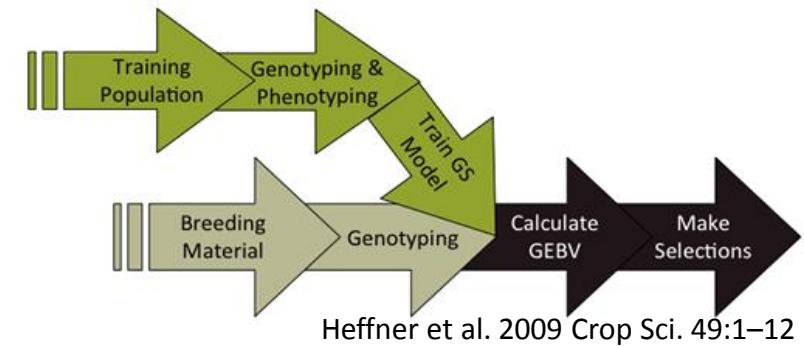
- CRISPR Cas9 can be used for :
 - Introduction of large deletions / Rearrangements
 - Introduction of single point mutations
- Interest for Gene function validation ! => acceleration of gene discovery and development of breeding tools
- New Breeding Technology (Creation of targeted variability)

FUTURE GENOMIC ASSISTED BREEDING STRATEGIES

- Heterotic groups identified
 - Towards dedicated Reciprocal Recurrent Selection schemes



- Marker assisted breeding strategies
 - Classical QTL / GWAS detection and allele follow up and stacking
 - Genomic selection : 1) Calibration of genotypic value based on markers and ii) prediction (no interest in « Genes »)



- Take advantage of the New Breeding Technologies (NBT)
 - Functional validation
 - Variability development



SORGHUM GENETICS AND BREEDING IN EU : PERSONAL THOUGHTS

- Sorghum breeding is efficient : an asset to provide producers with genetic materials adapted to their needs and the ones of the end-users (See Jeanson's Talk)
- European sorghum breeding has specificities (target environments). It requires specific research efforts
- Works performed in the US / Africa / India / Australia... are highly valuable and have to be considered
- No need to COORDINATE. But huge benefits to COLLABORATE / AGREGATE results and EXCHANGE EXPERIENCES : **we need a place for that !!**



PEOPLE AND FUNDINGS

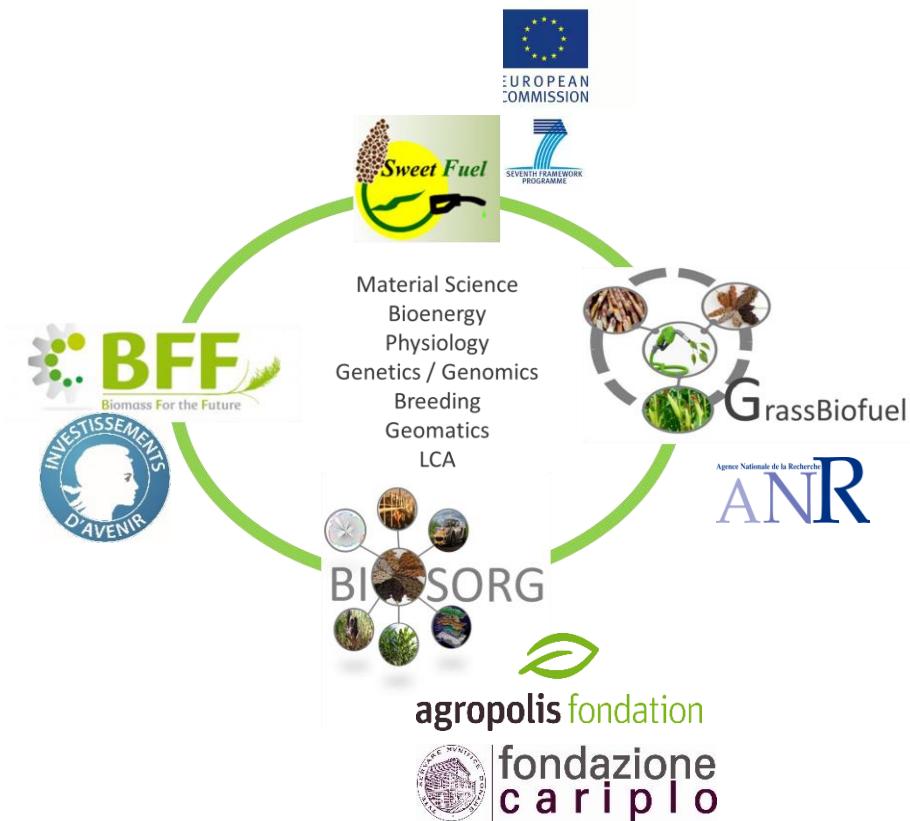
- Gilles Trouche



- Jean-François Rami



Sorghum Biomass Projects



QUESTIONS ?

