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Goat genome assembly, Availability of an international 50K SNP chip and RH panel: an update of the **International Goat Genome** Consortium projects

Gwenola Tosser-Klopp on behalf of IGGC









Outline

- IGGC presentation / history
- Goat Genome Assembly
- RH panel (discussed on monday)
- International goat SNP chip
- Next projects: Brian Sayre





IGGC

- International Goat Genome Consortium
- Created in March, 2010
- www.goatgenome.org
- Coordination: Wenguang Zhang & Gwenola Tosser-Klopp
- 3 ongoing projects
- Open meeting on Monday afternoon (3pm:5pm, Towne room) to discuss further projects





Capra hircus genome assembly



Yunnan black goat



Cashmere goat

Yunnan black goat (XX) lead to high quality reference genome of the domestic goat generated by combining Illumina new-generation short reads sequencing and the optical mapping technology of large DNA molecules which was used to generate the super-scaffolds.

Cashmere goat transcriptomes of primary and secondary fiber-growing follicles were generated







284,683 primary super-scaffolds

ψ.	Contig ₀		Scaffold.		Primary Super-scaffold	
ą.	bp	Number₽	bp	Number.	bp	Number
N90.	4,410.	141,869	440,999	1,348	582,523	976₽
N80.	7,994	100,335	846,998	922.	1,175,001	664.
N70.	11,323	73,948	1,253,003	664₽	1,739,998	481.
N60.	14,862	54,526	1,694,371	482.	2,447,724	352₽
N50 ₀	18,720	39,408	2,212,139	344.	3,057,189	254₽
Total₽	2,522,851,955	542,1 <mark>45</mark>	2,662,658,003	285,383	2,662,728,047	284,683

^{*}Total number: the number of contig/scaffold sequences with length > 100bp

The assembled base pairs (SOAPdenovo software) total 2.66 Gb, which is about 92% of the estimated goat genome size (~ 2.9 Gb)







349 super-scaffolds, using optical mapping

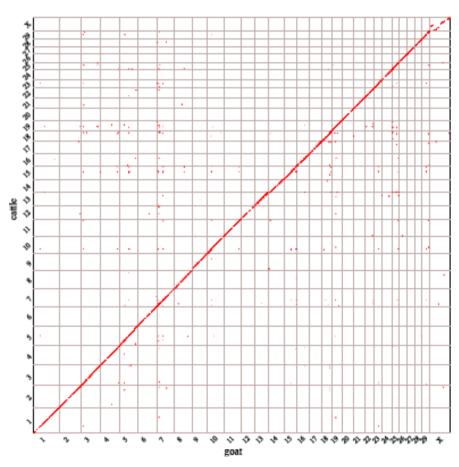
Super scaffold		Size (bp)	Number
N10		37,201,881	6
N20		31,8,804	13
N20		26,309,660	22
N40		22,009,453	32
N50		18,182,911	45
N60		14,726,152	60
N70		10,332,770	81
N80		6,331,148	112
N90		2,857,184	171
Total Number			349
Total Size	2,525,731,503		







30 pseudo-chromosomes



Based on the high colinearity between bovine and goats (Cribiu, E. P. et al. Cytogenet Cell Genet 2001), we used bovine genome to assemble the 315 super-scaffolds together with 422 extra scaffolds which were not included in super-scaffolds into 30 pseudo-chromosomes for the goat.







SNP chip project











Data

- INRA, France:
 - RRL of 6 goats (454) and whole genome sequencing (HiSeq) of 13 Alpine, Saanen and Creole
- –Malaysian Agricultural Research and Development Institute,Malaysia & DNA Landmarks :
 - Whole genome sequencing of 64 Boer, Savanna and Kacang meat and indigenous goats
- -University of Utrecht, Netherlands
 - RRL of 17 Saanen dairy goats. 120 millions of 32 bp paired-ends sequences
- -Italy, Spain, USA: ESTs + genes

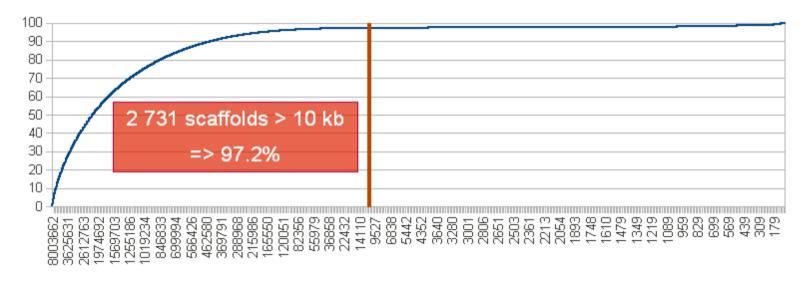




Goat scaffolds were used to map the « SNP » sequences

BGI scaffolds:

- 285 375 scaffolds/contigs => total length = 2, 662 Gpb
- length from 100 pb to 19 Mpb



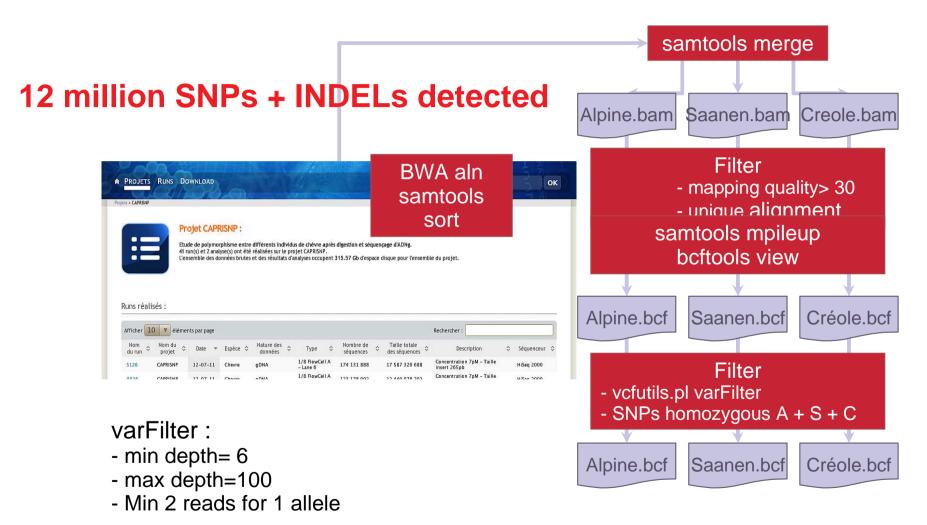
Bejing Genome Institute / Kumming Institute of Zoology / Inner Mongolia Agricultural University, China (W. Zhang, W. Wang)







Pipeline – INRA + NL data









Malaysian data & ESTs

Malaysian data

EST

BWA alignment (bwasw)

BWA alignment (bwasw)

SNP position extraction

SIM4 alignment validation

File to implement the database

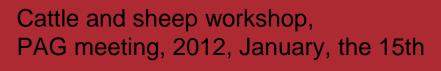
SNP position extraction

File to implement the

database

3.5 million SNPs

7000 SNPs

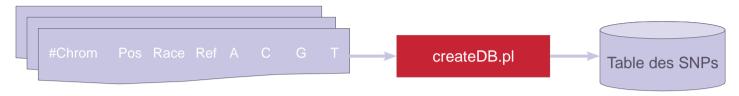


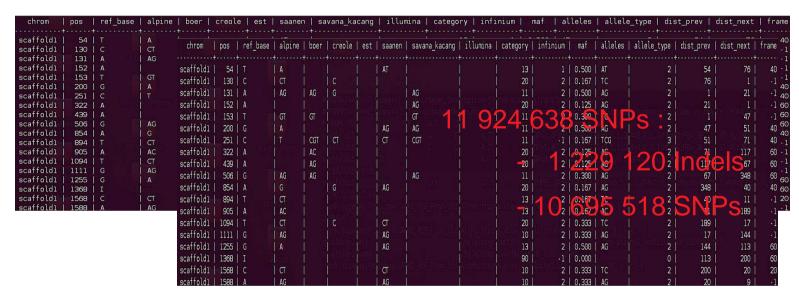






Data base filling





Infinium

I : A/T et $C/G \Rightarrow 2$ probes

II: other => 1 probe

Category

1: EST

2: Heteroz. in 5 breeds

3 & 4 : Heteroz. in 4 breeds

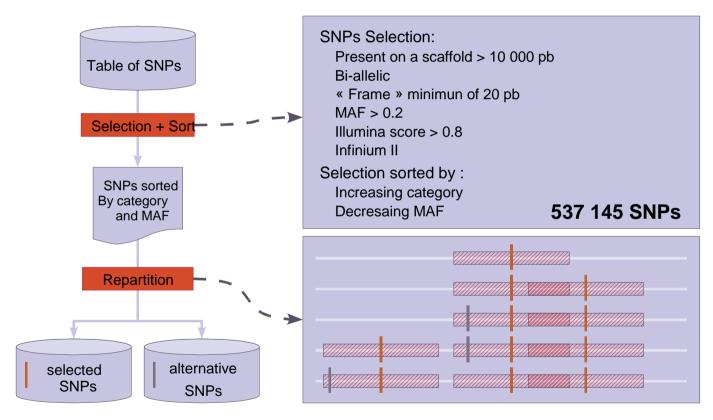
5 & 6: Heteroz. in 3 breeds...







SNP selection



28,500 pb minimum between 2 SNPs







60 000 selected SNPs

1: EST

2: Heteroz. in 5 breeds

3 et 4 : Heteroz. in 4 breed

5 et 6: Heteroz. in 3 breed

10: Heteroz. S & A

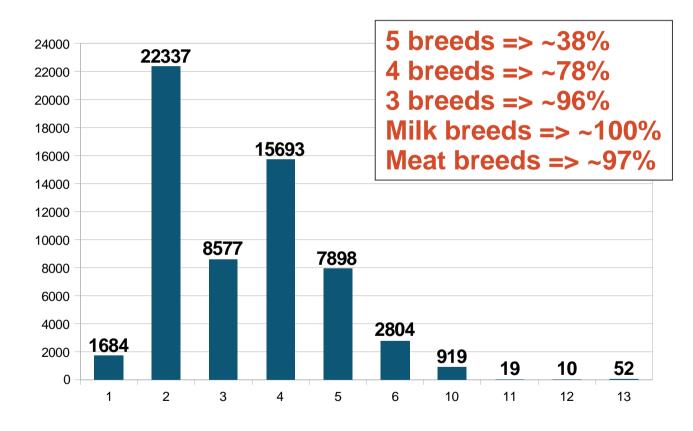
11 : Heteroz. (A or S

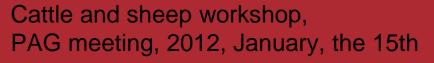
&(C or B or KS)

12: Heteroz. C et (B or KS

13: Heteroz. A or S

20 : other 90 : INDEL



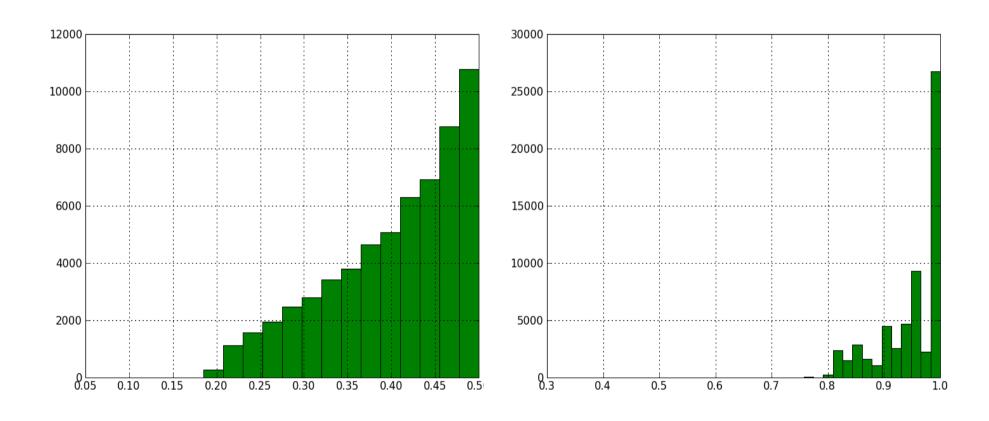








60 000 SNPs - MAF - Illumina Score



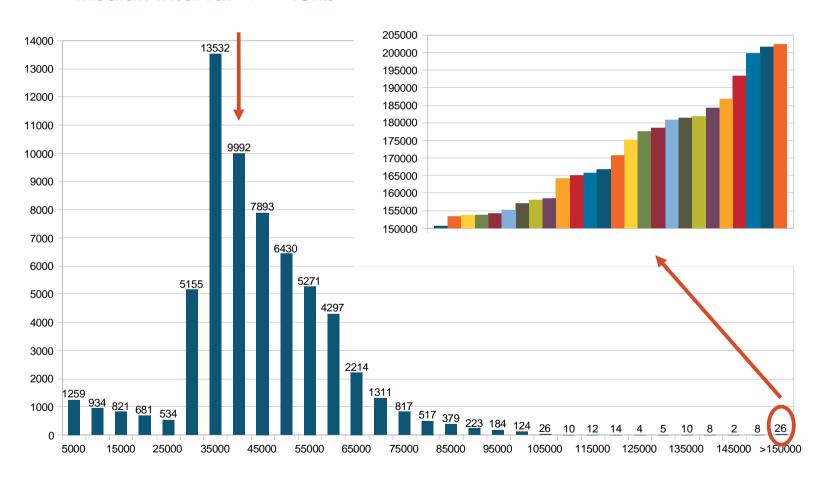






60 000 SNPs - Spacing

median interval => ~ 40kb



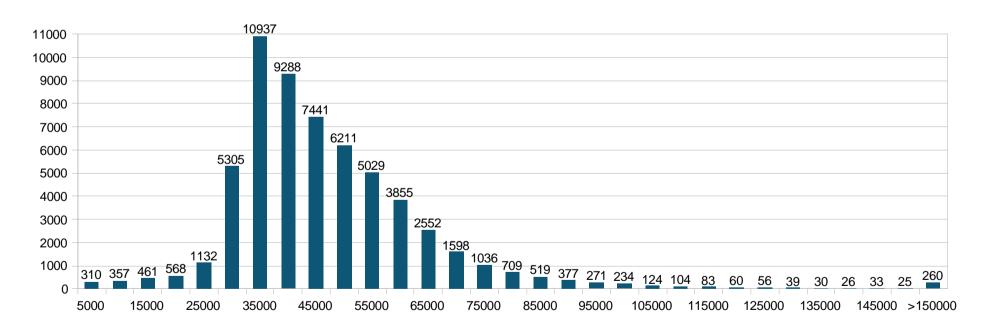






60 000 SNPs – Spacing on cattle genome (UMD3)

59 001 SNPs localised on a bovine chromosome









Chip manufacturing and cluster files

- Illumina iSelect design
- 288 animals were used for cluster file generation and quality control
- Includes the animals used for SNP discovery
- Breeds: Alpine, Saanen, Creole, Katjang,
 Savanna, Boer, Skopelos, Angora, Jinlan





SNP chip characteristics

- 53,348 synthesized loci
- 52,295 successful loci
- 8,000 ordered samples in September 2011
- Cluster files (.egt) available:
 Gwenola.Tosser@toulouse.inra.fr
- Annotation and publication of the loci coming soon





A chip useful for many breeds

Breed	Samples	SNPs MAF>0.05
Alpine	53	51339
Angora	26	47195
Boer	30	48494
Creole	38	50216
Jinlan	13	45648
Katjang	13	33873
Saanen	57	51689
Savanna	20	46629
Skopelos	27	50908
Yunling	1	17335







Upcoming projects of IGGC

- Hapmap project
- Resequencing
- Integration of RH and genome data
- •
- Open meeting on monday





Acknowledgements

SNP discovery:

- Henri Heuven
- Saadiah Jamli, Tun-Ping Yu
- Carole Moreno, Philippe Mulsant, Isabelle Palhière, Rachel Rupp, Gwenola Tosser-Klopp
- Marcel Amills, Patrice Martin, Eric Pailhoux, Brian Sayre, Alessio Valentini, (ESTs)
- Julien Sarry, Aurélie Tircazes
- UNCEIA, Capgenes and Apis-gene (French breeding organizations)

Genome sequence:

Jun Wang, Wen Wang, Wenguang Zhang

Bioinformatics:

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- Ibouniyamine Nabihoudine
- Curt Van Tassell for testing his spacing software on the data

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John McEwan









African Goat Production Value Chain Development Project

USDA-ARS and ILRI Sponsored Workshop Nairobi, Kenya, November 2011







African Goat Production Workshop

- A workshop was held in Nairobi, Kenya in November 2011 sponsored by the USDA-ARS and ILRI.
- The aims of this workshop were:
 - Bring together research experts for improvement of goat production in Africa
 - 2. Determine the potential of applying genome-based tools to value chain development projects in goat production
 - 3. Determine the current needs for characterization of goat populations and utilization of genome-based tools





General Project Development Concept

- Use emerging technologies to characterize and improve the adapted germplasm
- Development of a refined, high quality genome sequence and genome-based tools, if needed
- Development of genetic signatures for goat populations
- Determine the needs of the local producers for development of the goat production value chain
- Based on producer needs, develop improved germplasms using a genetic signature based approach







Outcomes

- Appears feasible to use high level genome-based tools for improved selection and sustainability of adapted germplasm
- Advantageous to have multiple independent genome sequence assemblies for the goat to improve the error checking and quality of the reference genome sequence
- Samples from African goat populations will be characterized with the current SNP panel to get an initial characterization of the populations
- Meet in October 2012, with support groups to develop local value chain assessments and determine methods for genetic signature utilization.





We thank you for your attention







