# Molecular epidemiology of Coxiella burnetii reveals the circulation of three main genotype clusters in French ruminants farms

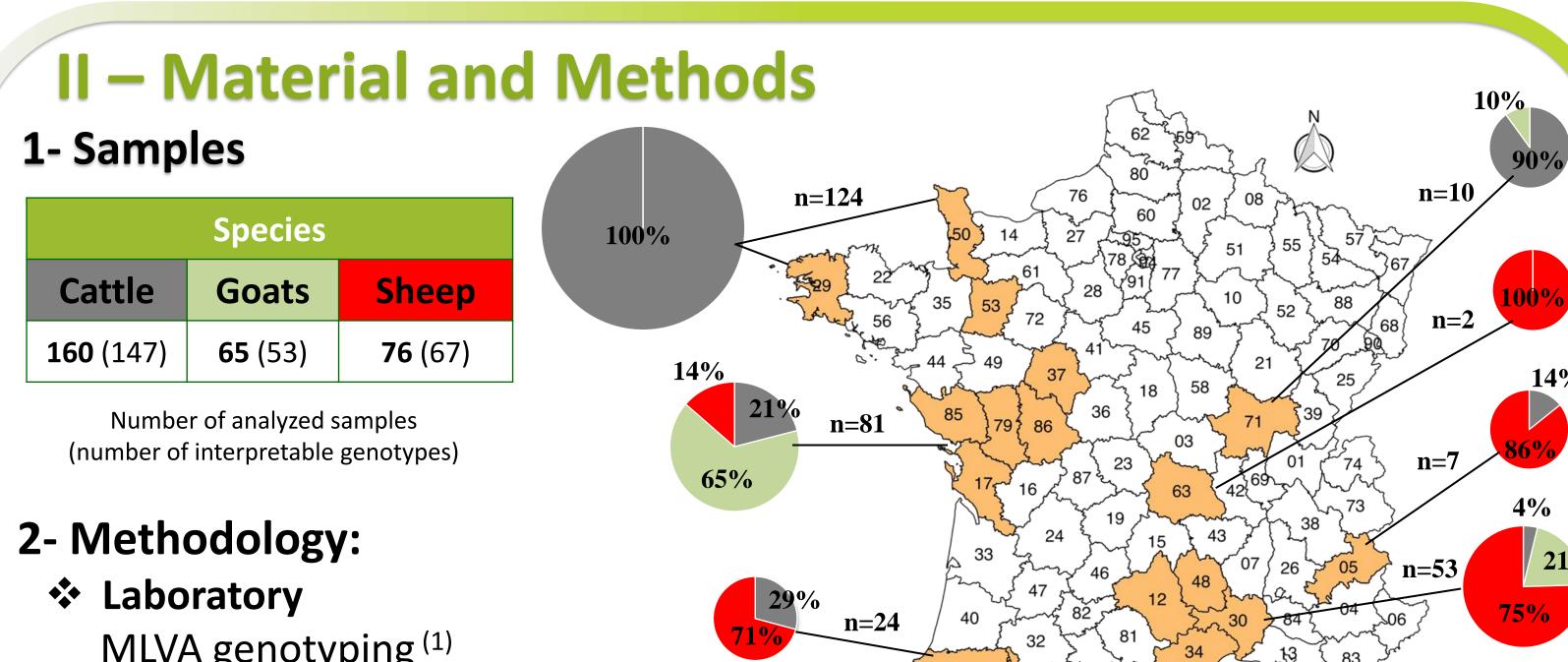
Joulié A.<sup>1,2,3</sup>, Sidi-Boumedine K.<sup>3</sup>, Bailly X.<sup>1</sup>, Gasqui P.<sup>1</sup>, Barry S.<sup>1</sup>, Jaffrelo L.<sup>4</sup>, Poncet C.<sup>4</sup>, Abrial D.<sup>1</sup>, Leblond A.<sup>2</sup>, Rousset E.<sup>3</sup>, Jourdain E.<sup>1</sup>

- <sup>1</sup> EPIA, INRA, 63122 Saint-Genès Champanelle, France;
- <sup>2</sup> Université de Lyon, VetAgro Sup, Marcy l'Etoile, France;
- <sup>3</sup> Anses (French Agency for Food, Environmental and Occupational Health and Safety), Laboratory of Sophia-Antipolis, Animal Q Fever Unit, Sophia-Antipolis, France;
- <sup>4</sup> GDEC, INRA, 63039 Clermont-Ferrand, France

## Context and objectives

Q fever is a worldwide zoonosis caused by the bacterium, Coxiella burnetii. In domestic ruminants, Q fever main clinical manifestations are abortions and stillbirths. Human outbreaks are generally associated with infected ruminants and especially small ruminants. A molecular characterization of circulating strains within French ruminants herds is essential to trace the origins of Q fever outbreaks and investigate potential links between genotypes and virulence traits. This study aimed to:

- Identify potential relationships between *C. burnetii* genotypes and domestic ruminant host species;
- Assess the distribution of *C. burnetii* genotypes within major ruminant breeding areas with reference to other European countries.



- MLVA genotyping (1)
  - $\rightarrow$  DNA chips  $\rightarrow$  2 markers > 500 pb

➤ DNA sequencer → 15 markers < 500pb

- **Bioinformatics and statistics** 
  - ➤ Parsimony network → Proportion of host species per sub-cluster

> Ascending hierarchical classification -> Genotype clustering

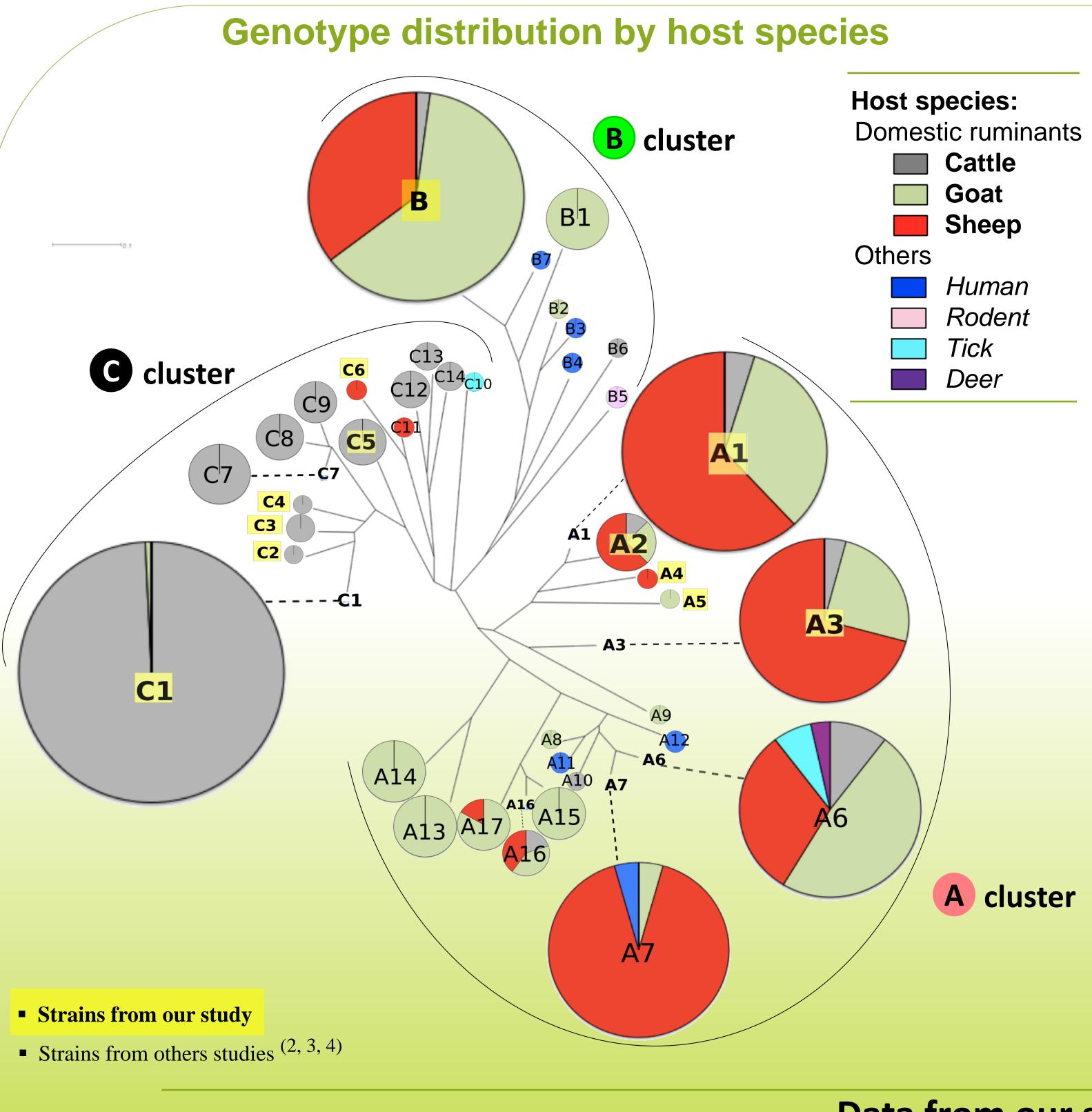
- >MANOVA Genotypic diversity associated with ruminant species and/or breeding areas

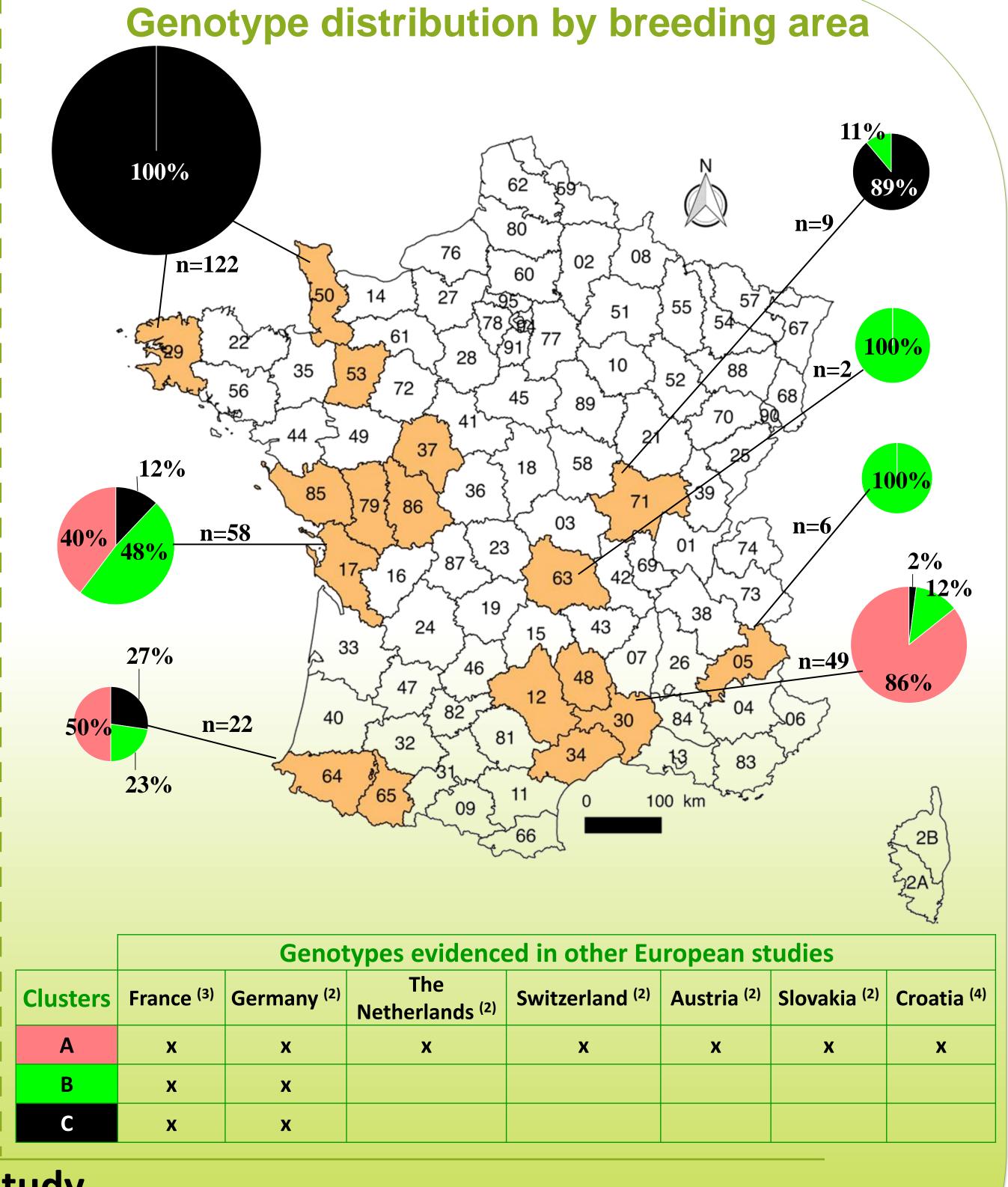
Relative number of animal sampled for each

host species by breeding areas

Unscaled pie charts

### Results





### Data from our study

> 3 main clusters : A B C

▶12 sub-clusters identified

65% sheep belong to A cluster (n=76)

✓ 63% goat belong to B cluster (n=48)

✓ 98% cattle belong to C cluster (n=144)

Genotypic diversity was significantly associated with both

host species and breeding areas (Manova, p<0.01)

### IV – Perspectives

Our species-associated patterns are consistent with those observed in other European studies (2,3,4). Further epidemiological studies are needed to complete our findings. Phenotypic approaches will allow establishing links between virulence of genotyped strains and ruminant species.



Aurélien Joulié INRA Auvergne-Rhône-Alpes Site de Theix F-63122 Saint-Genès Champanelle FRANCE Tél: 00 33 (0)4 73 62 47 87 Mail: aurelien.joulie@clermont.inra.f

http://www.ara.inra.fr/

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### **References:**

>(1) Arricau-Bouvery et al., 2006. Molecular characterization of Coxiella burnetii isolates by infrequent restriction site-PCR and MLVA typing. BMC microbiology 6, 38. >(2) Frangoulidis et al., 2014. Molecular analysis of Coxiella burnetii in Germany reveals evolution of unique clonal clusters. International Journal of Medical Microbiology, 304(7), 868-876. >(3) Prigent et al., 2015. Validation study for using lab-on-chip technology for Coxiella burnetii multi-locus-VNTR-analysis (MLVA) typing: application for studying genotypic diversity of strains from domestic ruminants in France. Microbes and Infection, 17(11), 782-788.

VNTR analysis. Veterinary microbiology, 173(3), 340-347.



