Analysis of pedigree data and whole-genome sequences in 12 cattle breeds reveals extremely low within-breed Y-chromosome diversity

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Summary

In this article, we analyzed pedigree information on males from 12 bovine breeds born in France between 2015 and 2019. We report an overall small number of paternal lineages with, for example, a minimal number of ancestors accounting for 95% of the Y-chromosome pool of their breed ranging from only 2 to 15 individuals. Then, we mined whole-genome sequence data from 811 sires (2 ≤ n ≤ 510 per breed) and built a median-joining network using 1411 SNPs. Most branches were breed-specific and in agreement with the geographic and genetic relatedness of these populations. The within-breed haplotype diversity was lower than expected based on genealogical information, which supports the existence of major male founder effects predating pedigree recording. In addition, we observed de novo mutation events among the descendants of the same ancestors, which are of interest to define paternal sub-lineages. Our results pave the way to future studies on the estimation of the effects of Y-chromosome haplotypes on male reproductive performances and on the conservation of Y-chromosome diversity.

Keywords: cattle, haplogroup, haplotype, paternal lineage, whole-genome sequence, Y1, Y2, Y-chromosome

In mammals, the Y chromosome is essential for male sex determination and fertility. In spite of its potential interest for breeding more fertile males, only a few research studies have focused on the genetic diversity and gene composition of the Y chromosome in cattle (Yue et al. 2015; Hughes et al. 2020). This lack of interest for the Y chromosome is due to the fact that breeding schemes have long focused on production traits (milk or meat) only, and massively used a few bulls with high genetic breeding value through artificial insemination (AI), without other consideration. In addition, the presence of a large number of highly repetitive and palindromic sequences, have posed a technological challenge in terms of sequencing and assembly of this chromosome’s sequence. Nevertheless, the absence of chromosomal recombination outside the pseudo-autosomal region preserves the original haplotypes, and the transmission of the Y chromosome can therefore be followed by studying paternal lines in pedigrees. In this study, we investigated Y-chromosome diversity in 12 beef and dairy breeds bred in France (Table 1) using both pedigree and molecular marker information.

As a preliminary analysis, we mined the genealogies of males born between 2015 and 2019 using the Lineage program, one of the latest updates of the PEDIG package (Boichard 2002). We considered only the individuals with both parents known and, for each breed, we calculated the contributions to the present Y-chromosome pool of the most ancient paternal ancestors recorded within each pedigree.

Regardless of the breed, the number of male lineages was very small and the main ancestor (defined as the individual with the highest contribution to the Y pool of its breed among the most ancient paternal ancestors) accounted for 23.2–58% of these (MainA, Table 1). However, we observed significant differences between dairy versus dual-purpose and beef breeds for all the metrics calculated. Dairy breeds showed a higher use of AI and, as a consequence, a smaller number of ancestors contributing to more than 1% of the paternal lineages (NbA > 1), smaller minimal numbers of ancestors accounting for 95% of the paternal lineages (MinNbA95), and higher contributions from the main ancestors (MainA; Table 1, Table S1). This phenomenon is particularly significant in our dataset for Holstein cattle in which 98.3% of the Y-chromosomes of the investigated population derive from only two ancestors: Wis Burke Ideal (HOLUSAM000001013415 born in 1947 with a contribution of 47.5%) and Rosafe Pearl Hannibal.
Then, we verified all individual genotypes that generated (Sargolzaei 2015) to create a preliminary median-joining network. We used Fimpute3 available for at least 90% of the variants (with heterozygous genotypes considered as missing). We used Fimpute3 without pedigree information to impute missing genotypes (Sargolzaei et al. 2014), and Popart v1.7 (Leigh & Bryant 2015) to create a preliminary median-joining network. Then, we verified all individual genotypes that generated web-like structures in the network and manually edited putative errors of variant calling or imputation according to the principle of parsimony (number of corrected genotypes n = 56/1 144 321). The final median-joining network was based on information from 811 sires (2 ≤ n ≤ 453 per breed; Tables S2 & S3) genotyped for 1411 SNPs, which to our knowledge represents the largest dataset analyzed to date in cattle in terms of individuals combined with number of markers. In total, 1266 SNPs (89.7%) supported the existence of the two main North-East (Y1) and South-West (Y2) European haplogroups that were initially described by Edwards et al. (2011) based on sequence information from three Y-chromosome genes and two microsatellite markers (Fig. 1). Among these 1266 SNPs, 66 were included in the panel of 68 markers used by Chen et al. (2018) to differentiate the Y1 and Y2 haplogroups. Within these haplogroups, most of the branches were breed-specific and reflected the geographic and genetic relatedness of these populations as previously evaluated using autosomal markers (Gautier et al. 2010; Park et al. 2015). For example, some Blonde d’Aquitaine bulls shared identical Y-haplotypes with Limousin and Salers bulls, and the same occurred between Montbéliarde and Simmental or between Abondance and Tarentaise. The within-breed haplotype diversity was lower than expected based on pedigree information with independent influential ancestors carrying the same Y-chromosome haplotype. For example, in Normande, we did not observe any variation between the Y-chromosomes of 30 bulls descending from five ancestors. These results support the existence of major male founder effects dating back to the creation of the breeds, and thus several decades before the advent of modern reproduction and selection techniques. The availability of 453 male whole genomes for the sole Holstein breed enabled us to

| Purpose | Breed          | Number of males | AI | NbA > 1 | MinNbA95 | MainA  |
|---------|----------------|-----------------|----|---------|----------|--------|
| Dairy   | Holstein       | 8 259 344       | 93%| 3       | 2        | 50.8%  |
|         | Montbéliarde   | 1 866 789       | 91%| 6       | 6        | 36.2%  |
|         | Normande       | 986 395         | 97%| 4       | 4        | 53.2%  |
|         | Abondance      | 111 424         | 80%| 5       | 5        | 44.3%  |
|         | Brown Swiss    | 75 638          | 95%| 5       | 5        | 34.2%  |
|         | Tarentaise     | 39 927          | 83%| 5       | 4        | 58.0%  |
| Dual    | Simmental      | 80 741          | 94%| 12      | 11       | 38.5%  |
|         | Charolaise     | 1 893 381       | 32%| 9       | 11       | 23.2%  |
|         | Limousine      | 1 359 913       | 16%| 12      | 15       | 39.5%  |
|         | Blonde d’Aq.   | 805 828         | 30%| 11      | 15       | 36.1%  |
|         | Salers         | 199 433         | 13%| 9       | 9        | 46.1%  |
|         | Rouge des Prés1| 98 254          | 25%| 10      | 9        | 28.6%  |

P-value dairy vs beef²

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identify 41 carriers of a third Y-chromosome haplotype in Holstein, which originates from the bull HOLUSAM000000827071 (born before 1950) that is the paternal great-grand sire of the influential AI bull Ivanhoe Star (HOLUSAM000001441440; Fig. 1). Nowadays, this rare haplotype account for only 1.2% of the Y paternal lines (See Suppl. Table S1). Moreover, our results show that only five of the 1411 SNPs differentiated the three basal Y-chromosome haplotypes found in the Holstein breed and we report six additional mutation events that occurred among the descendants of the two main paternal lines from Rosafé Pearl Hannibal, (HOLUSAM000001322381, n = 4) and Wis Burke Ideal (HOLUSAM000001013415, n = 2; Fig. 1). Adding these variants to custom SNP chips (Boichard et al. 2018) would be very useful to distinguish sub-lineages and estimate the effects on male-related traits of other types of mutations that may have accumulated over time, and require long-read sequencing technologies or quantitative PCR to be detected. Previously published analyses demonstrated that: (i) the ampliconic region of the Y chromosome can experience substantial loss or gain of gene copies within a few generations (Yue et al. 2015); (ii)}
some of these copy number variants are associated with scrotal circumference and male fertility in Holsteins (Yue et al. 2013, 2014); and (iii) consequently that the estimation of the impact of the Y-chromosome on traits based on paternal lineage is inaccurate (Yue et al. 2015).

In addition, we observed very rare carriers of the Holstein Y1 haplotype within the Simmental (1/70) and Montbéliarde (1/45) breeds, which result from planned introgressions during the 1970s and 1980s to improve milk production. Based on pedigree information, there is a single Holstein paternal lineage still segregating at a low rate (0.3%, seventh most important lineage) in the population of Montbéliarde males born between 2015 and 2019. The founder of this lineage is the Red Holstein bull Janes Royal (HOLUSAM000001599157, born in 1970), a remote descendant of the ancestor bull HOLUSAM000000798226 (born around 1950), the lineage of which has disappeared from the present Y-chromosome pool of French Holstein.

Given the large genetic distance between the Y1 and Y2 haplogroups, it is likely that they have accumulated many mutations in addition to the SNPs we are studying, with some of them potentially affecting male reproduction. The presence of Y1 and Y2 haplotypes on a similar genomic background in Montbéliarde represent a unique opportunity to study the impact of the Y variations on fertility traits in the future, when the number of Y1-haplotype carriers among Montbéliarde AI bulls will be sufficient. Estimating the phenotypic consequences of the variants associated with the different Y haplogroups and haplotypes found in West-European cattle breeds is a prerequisite before starting conservation introgression programs to improve Y-chromosome variability and/or male reproductive performances. Finally, we also identified six additional examples of Y1-haplotype carriers within breeds that are normally fixed for the Y2 haplotype. After verification, three of them were attributed to errors in pedigree data, breed affiliation, or in the correspondence between DNA samples and animal ID (Data S1). These results call for a systematic verification of the identity (and breed affiliation) of the individuals whole-genome sequenced for example by studying the concordance between genotypes derived from SNP array and from WGS, prior to any analysis. Such errors may have important consequences on imputation and subsequent GWAS or on filtering of breed-specific variants when searching for mutations responsible for recessive genetic defects (Michot et al. 2016).

In conclusion, using pedigree information and whole-genome sequence data, we report an extremely low within-breed Y-chromosome diversity in 12 dairy and beef cattle breeds. This situation due to ancient bottlenecks was particularly exacerbated in dairy breeds due to the increased use of modern reproduction techniques. The large set of informative SNPs identified in this study are of particular interest for future studies dedicated to the estimation of the effects of the various Y-chromosome haplotypes on male reproductive performances and to the conservation of Y-chromosome diversity.

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Conflict of interest

The authors report no conflict of interest.

Data availability statement

The whole-genome sequence data of 4110 taurine cattle individuals analysed in this study are part of the 1000 Bull Genomes Run 8.0. Part of these are already publicly available in the BioProject database (https://www.ncbi.nlm.nih.gov/bioproject/) at accession numbers PRJNA238491, PRJEB9343, PRJNA176557, PRJEB18113, PRNA343262, PRJNA324822, PRJNA324270, PRJNA277147, PRJEB5462 (Table S4). The complete dataset with all the Y-chromosome variants of the 811 males used in this study is available from the corresponding author on reasonable request.

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Supporting information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Table S1. List of ancestors accounting for at least 1% of the paternal lineage of their breed.

Table S2. Details on Y-chromosome SNPs considered in this study.

Table S3. Genotype information for the distinct haplotypes identified in this study.

Table S4. Haplogroups, sub-haplogroups and accession numbers of the 811 bulls.

Data S1. Investigations on Y1-haplotype carriers within breeds that are normally fixed for the Y2-haplotype.