A Genome-Wide Association Study Points out the Causal Implication of SOX9 in the Sex-Reversal Phenotype in XX Pigs

Sarah Rousseau¹, Nathalie Iannuccelli², Marie-José Mercat², Claire Naylies¹, Jean-Claude Thouly³, Bertrand Servin¹, Denis Milan¹, Eric Pailhoux⁴, Juliette Riquet¹*

¹ INRA, UMR444 LGC, Castanet-Tolosan, France, ² IFIP, Pôle génétique, la Motte au Vicomte, Le Rheu, France, ³ INRA, UE0332 Domaine de Galles, ⁴ INRA, UMR1198 BDR, Jouy-en-Josas, France

Abstract

Among farm animals, pigs are known to show XX sex-reversal. In such cases the individuals are genetically female but exhibit a hermaphroditism, or a male phenotype. While the frequency of this congenital disease is quite low (less than 1%), the economic losses are significant for pig breeders. These losses result from sterility, urogenital infections and the carcasses being downgraded because of the risk of boar taint. It has been clearly demonstrated that the SRY gene is not involved in most cases of sex-reversal in pigs, and that autosomal recessive mutations remain to be discovered. A whole-genome scan analysis was performed in the French Large-White population to identify candidate genes: 38 families comprising the two non-affected parents and 1 to 11 sex-reversed full-sib piglets were genotyped with the PorcineSNP60 BeadChip. A Transmission Disequilibrium Test revealed a highly significant candidate region on SSC12 (most significant p-value<4.65.10^-10) containing the SOX9 gene. SOX9, one of the master genes involved in testis differentiation, was sequenced together with one of its main regulatory region Tesco. However, no causal mutations could be identified in either of the two sequenced regions. Further haplotype analyses did not identify a shared homozygous segment between the affected pigs, suggesting either a lack of power due to the SNP properties of the chip, or a second causative locus. Together with information from humans and mice, this study in pigs adds to the field of knowledge, which will lead to characterization of novel molecular mechanisms regulating sexual differentiation and dysregulation in cases of sex reversal.

Citation: Rousseau S, Iannuccelli N, Mercat M-J, Naylies C, Thouly J-C, et al. (2013) A Genome-Wide Association Study Points out the Causal Implication of SOX9 in the Sex-Reversal Phenotype in XX Pigs. PLoS ONE 8(11): e79882. doi:10.1371/journal.pone.0079882

Editor: Stephen Moore, University of Queensland, Australia

Received June 3, 2013; Accepted September 26, 2013; Published November 6, 2013

Copyright: © 2013 Rousseau et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: The project was supported by the French National Research Agency (Agence Nationale de la Recherche, ANR-08-GENM-040 SwAn project) and by BIOPORC. SR PhD is funded by LABOGENA and the Research Ministry (CIFRE grant). The ANR and LABOGENA had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript; BIOPORC has contributed to the construction of the collection used for the design: BIOPORC had no role in the analysis and the preparation of the manuscript.

Competing interests: This study was partly funded by BIOPORC and LABOGENA. There are no patents, products in development or marketed products to declare. This does not alter our adherence to all the PLOS ONE policies on sharing data and materials, as detailed online in the guide for authors.

* E-mail: juliette.riquet@toulouse.inra.fr

Introduction

In mammals, sex differentiation is a genetically and hormonally controlled process. Sex differentiation first relies on the establishment of genetic sex at fertilization by the combination of X or Y chromosome containing gametes to produce XX or XY zygotes. This initial step is then followed by the activation of the appropriate male or female genetic cascades involved in gonad differentiation. In a normal situation, the switch between the male or female pathway is driven by the action of the Y-located testis-determining gene SRY (Sex-determining Region of the Y chromosome) [1]. Once gonad differentiation is engaged, the secretion of testicular anti-Müllerian hormone (AMH) and androgens ensure that sexual characteristics develop correctly into their final form [2]. At any of these steps, flaws in the process can lead to disorders of sexual development. When chromosomal, gonadic and somatic sexes are not consistent, sexual attribution is ambiguous, and both male and female characteristics may be found in a single individual. In mammals, sex reversal has been described in numerous species including humans [3], pigs [4,5], goats [6,7], horses [8], dogs [9,10], mice [11], marsupials [12] and moles [13]. Naturally occurring models of SRY-negative XX sex reversal in XX individuals could provide important clues as to the etiology of some cases of the human disorder and increase our knowledge about sex differentiation in vertebrates. These pathologies can result either from loss-of-function of crucial pro-ovarian genes (for example, FOXL2 in...
and goats present a XX intersex phenotype at a higher frequency than other livestock species. These XX sex-reversed individuals have a normal female karyotype (XX) but some degree of testicular differentiation of their gonads [19]. In goats, SRY-negative XX sex reversal is due to a recessive mutation, referred to as the "PIS" (Polled Intersex Syndrome) mutation, which is associated with the dominant autosomal trait for hornlessness [6,20]. The causative mutation, identified in 2001, is a large 11.7 kb deletion that affects the transcription of PISRT1 (PIS-regulated transcript 1) and FOXL2 (Forkhead box L2). In the homozygous state, this mutation causes the masculinization of all polled genetic females [6]. Despite this successful result in goats, the causal genes in SRY-negative XX sex reversal in pigs remain unidentified.

Pig intersex phenotypes range from true hermaphrodites (48%) to male phenotypes with (50%) or without (2%) ambiguities [4]. As in humans some of these cases result either from the presence of XX and XY cells within the same animal (4%) or from the presence of a small fragment of the Y chromosome (containing the SRY gene) in XX individuals (2%). However, most XX sex-reversed individuals (94%) do not have SRY or any of the other genes normally located on the Y chromosome (SRY-negative XX sex reversal). In these cases, testis induction is therefore due, to alterations of autosomal genes in the absence of SRY [4]. In pigs, the frequency of XX sex reversal varies from 0.08 to 0.75% depending on the herd, and available heritability estimates for this disorder are high, ranging from 0.72 to 0.81 [21]. Previous segregation analyses with pedigrees of limited size are consistent with autosomal recessive transmission of a single gene [4]. A more recent study however, suggests the possible involvement of multiple autosomal regions [22].

Until recently, the identification of genes causing inherited disorders was performed using genome-wide pedigree-based linkage analyses, or by focusing on candidate genes using population-based association studies. With the recent development of commercial high throughput SNP genotyping BeadChips, these two approaches have merged into a single approach referred to as genome-wide association studies (GWAS). GWAS methods have been shown to be more powerful than linkage based methods [23]. The power of GWAS has been demonstrated by the identification of several genes underlying a range of diseases in humans, dogs and cattle which had been difficult to study previously [24,25].

In this contribution we report the results of a GWAS study of XX intersex in French Large White pigs. We genotyped nuclear families identified by one or several sex reversed offspring using the Illumina PorcineSNP60 BeadChip tool to identify chromosome regions and markers associated with sex reversal. A unique association signal was identified and suggests a probable role for the SOX9 gene in sex reversal in pigs. The SOX9 region has been described as a gene desert, presenting extremely conserved elements among mammals. It is now clear that SOX9 is expressed by means of a complex regulatory region thought to cover at least 1 Mb. Among these sequences, Sekido and Lovell-Badge first located a 3.2-kb testis-specific enhancer of Sox9 (TES) that was [26] further refined to 1.4 kb and is referred to as the Testis Enhancer Sequence core element (TESCO). Even if SOX9 was retained as the main candidate gene, TESCO was also candidate for the causative mutation in pigs.

Materials and Methods

Animals and Phenotypes

DNA samples were collected from experimental families in the 1990s, and more recently from nuclear families during the national selection program of the French Large-White breed. The collection of experimental XX sex-reversed cases was established over a 6 year period starting in 1992 in an experimental herd (Domaine de Galles, Avord, France) of the Institut National de la Recherche Agronomique (INRA). This collection has been described previously [4] and some cases were used for in-depth descriptions of XX sex-reversal, including fetal development [5]. Briefly, all cases (n=30) were tested for the presence of the Y chromosome, especially SRY. Then, the parents of true XX sex-reversed cases (94%) were retained in the experimental herd and mated to determine the mode of inheritance of the intersex condition in the pig. The reproductive systems of all intersex cases were examined post-mortem in adults (sporadic cases) or at 5 weeks postnatally (familial cases). As a result, 12 nuclear families comprising 1 to 9 affected piglets and their two non-affected parents were selected.

In addition, samples from 26 half/full-sib families were collected between 2006 and 2010 from the Large-White population of the French breeding companies belonging to the BIOPORC association. In addition to blood sampling, the cases were described by the farmer, the technician or the veterinary surgeon using standard nomenclature. The phenotype was defined using only external anatomical criteria: enlarged vulva, presence of a penile clitoris, number of testicles (0, 1 or 2) located in a scrotum-like structure, and presence of a midventral penile sheath. Animals presenting one or more of these criteria were considered as affected and included in the sample. Samples were also collected from their non-affected parents. These two phenotypic categories, affected and non-affected, were used for association analysis. The animals in the experimental families produced between 1992 and 1998 were described by Pailhoux & al. in the frame of another study [4]. All procedures were approved by the animal welfare commission of the Institute at the time of this study. In our study, only the DNA collected from these animals was used. The second set of samples was collected by the BIOPORC association. The collection of blood samples and external phenotyping of the animals were performed in compliance with the guidelines of the French Ministry of Agriculture and Fisheries.

Sample genotyping

Genomic DNA was extracted from blood samples using standard protocols. For each sample, the presence/absence of the SRY gene was determined by PCR amplification with

PLOS ONE | www.plosone.org 2 November 2013 | Volume 8 | Issue 11 | e79882
suggestive association were determined. We applied a than 95% (4944 SNPs), and markers with a minimum allele quality control and genome-wide association analyses population frequency of allele B (PFB). Both LRR and BAF surrounding the Markov Model (HMM) including the pedigree information when the PFB file was calculated based on the BAF of each and 1.91 \times 10^{-6} \) (significant, \( \log(1/p) = 5.97 \)) and 1.91 \times 10^{-5} \) (suggestive, \( \log(1/p) = 4.72 \)).

Identification of CNV's

PennCNV software [29] was used to detect pig CNVs surrounding the SOX9 gene on SSC12. Analysis was performed with all the SNP of the chip (62,163) but only the results obtained on SSC12 were considered. PennCNV software integrates, in a joint-calling algorithm, a Hidden Markov Model (HMM) including the pedigree information when available, the total signal intensity (Log R ratio - LRR) and allelic intensity ratio (B allele frequency - BAF) at each SNP marker, the distance between neighboring SNPs and the population frequency of allele B (PFB). Both LRR and BAF were exported from GenomeStudio (Illumina San Diego, USA) and the PFB file was calculated based on the BAF of each marker. CNV calling was performed using the default parameters of the HMM model with 0.01 as UF factor. The marker positions on SSC12 were derived from the swine genome sequence assembly (10.2) (http://www.ensembl.org/ Sus_scrofa/Info/Index).

TESCO and SOX9 sequencing

Five affected pigs, carrying different mutated haplotypes (including the 3 most frequent) were selected for sequencing. These five unrelated individuals were selected from five different nuclear families of the design. In addition, five individuals were selected as controls in families from another project in which no cases of intersexuality were reported. Pairs of primers, covering the entire SOX9 gene and TESCO regulatory region, were chosen to amplify and sequence these 2 candidate regions (Table S1). The SOX9 pig sequence is available and annotated in Ensembl (Sscrofa10.2:12:9028879:9033246:1), whereas the TESCO pig sequence was obtained by aligning TESCO human sequence against the pig draft genome (Sscrofa10.2:12:9045199:9047423:1). To sequence PCR products, an aliquot (1–12 µl) was purified in a single step (45 min at 37°C followed by 30 min at 80°C) using 0.5U of Shrimp Alkaline Phosphatase (Promega) and 0.8 U of Exonuclease I (New England Biolabs). Sequencing was performed on a 3730 ABI capillary DNA sequencer using a Big Dye terminator V3.1 cycle sequencing kit (Applied Biosystems, Inc.). Sequences were analyzed using the CodonCode Aligner software v4.0.4 (http://www.codoncode.com/aligner/), and any polymorphisms were detected by comparison with the reference sequence. Multiple alignments of the core region of TESCO was performed with multalin software [30]. In total, five distinct sequences (alleles) were obtained for SOX9 (Genbank Accession numbers KF422597, KF422598, KF422599, KF422600, KF422601), and 3 for the TESCO region (Genbank Accession numbers KF422602, KF422603, KF422604).

Results

Informativeness of the SNP panel

Thirty-eight half/full-sib families comprising 89 sex-reversed piglets were genotyped with the Illumina PorcineSNP60 BeadChip. After quality controls, 173 individuals and 47,155 autosomal SNP were retained for transmission disequilibrium (TDT) analysis. The number of SNPs per chromosome ranged from 1,051 on SSC18 to 5,216 on SSC1. MAFs of polymorphic markers were uniformly distributed between 0.01 and 0.5 (Figure S1). The power of the TDT was directly proportionate to the number of heterozygous parents for markers associated with causative variants, and the level of linkage disequilibrium (LD) between causative variants and SNP markers. The proportion of heterozygous parents for each of the 47,155 SNPs is represented in Figure S2. At least half of the parents of the design were not informative for more than 85% of the SNPs. The LD estimation was computed as reported by Dupuis & al [31] to assess the level of genome coverage provided by the 47,155 remaining SNPs. Each SNP was considered as a pseudocausative polymorphism in order to identify the SNP (and its distance) with the highest \( r^2 \) value (\( r^2_{\text{max}} \)) among the remaining markers on the chip located within a 2 Mb surrounding window. Calculations were performed using only the parents of the design. The cumulative frequency distribution of \( r^2_{\text{max}} \) is reported in the Figure S3(A). In our pedigree, \( r^2_{\text{max}} \) values were lower than 0.5 for 28% of the SNPs and only 25% of the SNPs were in total linkage disequilibrium (\( r^2_{\text{max}} =1 \)) with another SNP on the chip. The average distance for SNP with a \( r^2_{\text{max}} \) of 0.5 was 300 kb (Figure S3(B)). The number of remaining markers and the informativeness of the design indicate that the coverage of the SNP panel may not be optimal for all chromosomal regions.
Transmission Disequilibrium Test

Due to the two periods of birth of the founders, the population structure was quantified using genome-wide pairwise identity-by-state distances estimated by PLINK. All animals were grouped in a same cluster; these results indicate that despite the differences in the collection periods the two populations have not diverged. The SNPs showing significant genome-wide ($p < 1.06 \times 10^{-6}$) and chromosome-wide ($1.06 \times 10^{-6} < p < 1.33 \times 10^{-5}$) association with the intersexuality (IS) phenotype as determined by TDT are presented in Table 1 and the Manhattan plot in Figure 1. All these SNPs were located on SSC12 between position 6,459,472 and 9,597,291 bp, the most significant marker being located at position 8,881,355 bp (M1GA0024789) (Table 1). Because all these SNPs were localized within the same short interval, it seemed likely that they were linked to a single causal locus. No other significant or suggestive regions were detected by the genome scan. No signals were identified close to PISRT1 and FOXL2 on chromosome 13, in spite of very informative markers in that region (ALGA0119502: 39 informative meioses ($p=0.53$) and ASGA0089628: 43 informative meioses ($p=1$)).

Detection of CNV

To investigate whether a chromosomal rearrangement (deletion or duplication) surrounding the SOX9 gene could be causal, genotyping data obtained with the 60K SNP chip were analyzed with PennCNV. The analyses of SNP arrays provide normalized total signal intensity and allelic intensity ratios which represent overall copy numbers and allelic contrasts. To detect CNV, we jointly analyzed the 60KSNP genotypes of the 89 affected piglets. Successive windows of 3 markers, corresponding to 150kb windows on average (given the density of the chip) were analyzed. Among all the affected individuals of our design, no structural variations were identified in the 3.14 Mb segment encompassing SOX9.

Haplotypic characterization of SOX9 surrounding interval

To further refine the map position of the causative mutation, 35 SNPs of the chip located between positions 8,566,755 and 9,604,096 bp (interval containing “the significant markers” bounded by an additional marker on each side) were selected.

### Table 1. Significant SNP association hits in the Transmission Disequilibrium Test for sex-reversal.

| SNP name       | Chr | Position (bp) | $\chi^2$ | p-value          | log(1/p) |
|----------------|-----|---------------|----------|-----------------|----------|
| ASGA0052652    | 12  | 6,459,472     | 21.6     | 3.36E-06        | 5.47     |
| ASGA0052953    | 12  | 8,586,694     | 28.47    | 9.51E-08        | 7.02     |
| ASGA0098350    | 12  | 8,826,005     | 24       | 9.63E-07        | 6.02     |
| ALGA0112008    | 12  | 8,835,743     | 23.4     | 1.32E-06        | 5.88     |
| ALGA0108362    | 12  | 8,836,960     | 23.4     | 1.32E-06        | 5.88     |
| MARC0011974    | 12  | 8,851,013     | 22.7     | 1.89E-06        | 5.72     |
| M1GA0024789    | 12  | 8,881,355     | 38.82    | 4.65E-10        | 9.33     |
| MARC0009109    | 12  | 8,894,982     | 34.68    | 3.89E-09        | 8.41     |
| ALGA0109673    | 12  | 8,897,127     | 23.82    | 1.06E-06        | 5.98     |
| ALGA0064738    | 12  | 9,053,320     | 34.77    | 3.71E-09        | 8.43     |
| ASGA0053002    | 12  | 9,128,147     | 33.91    | 5.78E-09        | 8.24     |
| HSGA0033370    | 12  | 9,257,006     | 33.91    | 5.78E-09        | 8.24     |
| ASGA0097657    | 12  | 9,469,336     | 33.24    | 8.14E-09        | 8.09     |
| MARC0041079    | 12  | 9,597,291     | 22.35    | 2.27E-06        | 5.64     |

Chromosome-wide significant SNP ($1.06 \times 10^{-6} < p < 1.33 \times 10^{-5}$) are in italic, genome-wide significant SNP ($p < 1.06 \times 10^{-6}$) in normal font, and the SNP corresponding to the lowest p-value is highlighted in bold. Positions were defined on the Sscrofa10.2 draft sequence.

doi: 10.1371/journal.pone.0079882.g001

| SNP name       | Chr | Position (bp) | $\chi^2$ | p-value          | log(1/p) |
|----------------|-----|---------------|----------|-----------------|----------|
| ASGA0052652    | 12  | 6,459,472     | 21.6     | 3.36E-06        | 5.47     |
| ASGA0052953    | 12  | 8,586,694     | 28.47    | 9.51E-08        | 7.02     |
| ASGA0098350    | 12  | 8,826,005     | 24       | 9.63E-07        | 6.02     |
| ALGA0112008    | 12  | 8,835,743     | 23.4     | 1.32E-06        | 5.88     |
| ALGA0108362    | 12  | 8,836,960     | 23.4     | 1.32E-06        | 5.88     |
| MARC0011974    | 12  | 8,851,013     | 22.7     | 1.89E-06        | 5.72     |
| M1GA0024789    | 12  | 8,881,355     | 38.82    | 4.65E-10        | 9.33     |
| MARC0009109    | 12  | 8,894,982     | 34.68    | 3.89E-09        | 8.41     |
| ALGA0109673    | 12  | 8,897,127     | 23.82    | 1.06E-06        | 5.98     |
| ALGA0064738    | 12  | 9,053,320     | 34.77    | 3.71E-09        | 8.43     |
| ASGA0053002    | 12  | 9,128,147     | 33.91    | 5.78E-09        | 8.24     |
| HSGA0033370    | 12  | 9,257,006     | 33.91    | 5.78E-09        | 8.24     |
| ASGA0097657    | 12  | 9,469,336     | 33.24    | 8.14E-09        | 8.09     |
| MARC0041079    | 12  | 9,597,291     | 22.35    | 2.27E-06        | 5.64     |

Chromosome-wide significant SNP ($1.06 \times 10^{-6} < p < 1.33 \times 10^{-5}$) are in italic, genome-wide significant SNP ($p < 1.06 \times 10^{-6}$) in normal font, and the SNP corresponding to the lowest p-value is highlighted in bold. Positions were defined on the Sscrofa10.2 draft sequence.

doi: 10.1371/journal.pone.0079882.g001
to characterize the haplotypes segregating among affected offspring. The phases of each descendant were reconstructed using familial allelic transmission from parents to offspring. Comparing the haplotype-based genotypes, no single homozygous chromosome segment could be identified among the affected piglets of the design. In total 43 different

Figure 2. Log(1/p) values of the SNP localized in the 6Mb-12Mb region on SSC12. The non-significant markers are presented in grey, suggestive ones in medium grey, the significant SNP in black. The suggestive (dotted line) and significant (solid line) thresholds are indicated in red. The two vertical lines delimit the minimum interval defined by the first and least significant SNP in the region. At the bottom, genes and transcripts annotated in Ensembl (Sscrofa10.2) are reported.
doi: 10.1371/journal.pone.0079882.g002
haplotypes segregate within the 38 families. However, among the affected offspring, the frequency of haplotype “8” (50.8%) was much higher than the frequencies of other haplotypes (Figure 3). In addition the frequencies of two other haplotypes, haplotypes 2 (6.7%) and 17 (6.3%) were slightly higher than the others.

No IBS segment was found in common in the 43 haplotypes, but most of them shared segments of variable size (2 to 30 adjacent markers) identical to the most frequent haplotype. These results suggest that a mutation may be shared by several haplotypes, including the three more frequent ones.

Candidate region sequencing

The minimum interval mapped by genetic analysis covered 882,642 kb and the causative mutation could be locate anywhere within it. Before undertaking an exhaustive search for all polymorphisms in this region, we chose to target the regions annotated in the interval: the SOX9 gene and its regulatory region TESCO. We could not exclude that a substitution of an amino-acid residue might increase the lifespan or the DNA affinity of the protein, or block the nuclear exportation signal and lead to abnormal nuclear retention of SOX9 in the XX gonads [32]. To test whether the causal mutation might be located in the SOX9 gene or in its regulatory TESCO region, these elements were sequenced in five individuals. Four were homozygous or heterozygous carriers for haplotypes “8”, “2” and “17”, and the fifth was heterozygous for two rare haplotypes. The aim was (1) to search for candidate mutations that might affect the functional structure of the gene or (2) to identify a small segment shared between the “carrier” haplotypes. Even if a candidate mutation was not identified in the sequences of SOX9 and TESCO, the presence of a single haplotype shared among all affected individuals would suggest that the mutation is close to these two elements.

Additionally, five healthy animals were also sequenced. The genomic sequence of SOX9, including the 5'UTR, the 3 exons,
The 2 introns and the 3 'UTR regions, was obtained using 7 pairs of primers and the 1.4kb core region of TESCO was obtained with 4 pairs of primers. In the genomic sequence of SOX9, 14 different polymorphisms were identified (10 substitutions and 4 indels from 1 to 18 nucleotides). All these polymorphisms were located outside of the gene exons and outside of the intron / exon junctions. One Indel and 14 substitutions were characterized in the 1.4 kb TESCO region. Consequently, no functional candidate mutation was highlighted (Figure S4). The distribution of the polymorphisms among the sequenced individuals (Figure 4) showed five different alleles for SOX9 and 3 for TESCO. No shared allele was present between the different "carrier" chromosomes, and identical alleles were identified on "carrier" and "control" chromosomes. Among the control animals, two were homozygous for the same alleles as those characterized in case pigs (Figure S4).

### Discussion

This study focused on the identification of the gene(s) underlying XX sex reversal in the pig. A TDT was conducted using the Illumina PorcineSNP60 array to detect risk loci. In this family-based study, we found genome-wide association with intersexuality for 9 SNPs on SSC12. All these markers are localized in a same small interval that also contains the strong candidate gene, SOX9. Some of the experimental families included in the present analysis had been analyzed in 2001 for the expression level of SOX9 in gonads at 4 developmental stages (50 and 70 dpc; 5 weeks after birth and adult) [5]. This included gonads of XX males with genital ambiguities (mainly

| Case | TESCO | SOX9 |
|------|-------|------|
| 8    | 361   |       |
| 8    | 627   |       |
| 51   | 971   |       |
| 966  | 637   |       |
| 1028 | 936   |       |
| 1228 | 1288  |       |
| 1268 | 1268  |       |
| 1268 | 1268  |       |
| 1334 |       |       |
| 134  | 217   |       |
|       | 5'UTR |       |
|       | 247   |       |
|       |       | 3'UTR |
|       |       |       |

**Figure 4. Polymorphisms identified for the 5 cases and 5 controls individuals used for sequencing.** For each case, the number in red corresponds to the haplotype number defined with the 35 SNP of the chip surrounding SOX9. For each individual the phases defined in TESCO and SOX9 with the detected polymorphisms is reported. Positions were defined from position 9033579 (=1) on the pig draft sequence for SOX9 and 9047123 (=1) for TESCO. Five different alleles were detected for SOX9 (Genbank Accession numbers: KF422597, KF422598, KF422599, KF422600, KF422601), and 3 for the TESCO (KF422602, KF422603, KF422604).

doi: 10.1371/journal.pone.0079882.g004
testicular tissue) and of true hermaphrodites for which testis-like gonads and ovarian-like gonads had been studied separately. Results had shown that the level of SOX9 increases in testis-like gonads of XX males and of true hermaphrodites, whereas it remains at a normal female level in the ovarian-like tissues of the latter. However, whereas these results alone were not proof for genetic causality, a SOX9 genetic determinism has now stronger support with the association signal obtained here. No additional suggestive regions were identified in accordance with a monogenic autosomal recessive mode of inheritance as previously proposed [4].

In the mouse, the transcription factor SOX9 has been shown to be the direct target of the protein encoded by SRY [26]. It has several relevant functions during development and is essential for testis differentiation. In the mouse, Sox9 is initially expressed in the bi-potential gonad of both sexes at 10.5 day post conception (dpc). It then becomes highly up-regulated in XY gonads and down-regulated in XX gonads at 11.5 dpc [33]. In males, it plays an essential role during Sertoli cell differentiation from supporting cell precursors, which is the preliminary step in testis differentiation [34]. In females, the down-regulation of SOX9 is maintained in the granulosa cells throughout fetal development. In pigs, SOX9 is expressed in a similar pattern to that described in mice [35]. Porcine SOX9 expression during embryonic and fetal development begins at 21 dpc, in both sexes and then increases in the testis from 28dpc on. As in mice, expression in the female is intense as early as 21 dpc, and then decreases from 28 dpc, the key stage of gonadal switch in pigs.

Both loss- and gain-of-function studies in the mouse support the implication of the Sox9 gene in gonadal differentiation. A conditional deletion of Sox9 in the gonads results in XY male- to-female sex reversal [36,37] whereas misexpression of a Sox9 transgene in XX mice induces female-to-male sex reversal [17,38]. Similar phenotypes have been found in humans. Mutations within and outside of SOX9 leading to haploinsufficiency are known to cause a bone disorder, campomelic dysplasia, in both sexes and gonadal dysgenesis in 75% of XY individuals presenting as females [39,40]. Even if female-to-male sex reversal in humans is rare, Huang & al. reported in 1999 [16] evidence supporting that SOX9 duplication, through a de novo mosaic 46,XX,dup(17) (q23.1;q24.3), can cause XX sex reversal. In 2011, Cox & al. described a familial 46, XX Developmental Testicular Disorder due to an approximately 178-kb duplication located 600 kb upstream of SOX9 [41] and Vetro & al., a 96 kb triplication 500 kb upstream of SOX9 [42]. Several copy number variants (CNV) and translocations within a large gene desert region upstream of SOX9 have been characterized as responsible for some of these developmental disorders affecting the skeleton and the genitalia [43,44]. It is now clear that Sox9 is expressed in various tissues by means of a complex regulatory region thought to cover at least 1 Mb [45]. Based on the large amount of information obtained in other species, we looked for candidate CNVs within the interval detected by TDT that might disrupt regulatory regions. We thus mined our 60K genotyping data for CNVs, but were not able to identify any structural variations. No CNVs have been reported in this interval in previously published studies in different pig breeds [46,47]. Although CNV detection is feasible with this technology, it is impaired by low marker density, irregular distribution of SNPs along chromosomes and the lack of non-polymorphic probes specifically designed for CNV identification. Hence, only the largest CNVs are expected to be detected using the Porcine 60K SNP Chip. We cannot therefore exclude the possibility that CNVs of smaller sizes might exist in the candidate interval. Results obtained in humans with a dense SNP chip reported much higher numbers of CNVs with smaller average sizes than with a less dense SNP chip. Based on our analysis, we cannot exclude that the causal mutation is a small deletion, but it is unlikely that a major structural variation is responsible for the sex-reversed cases studied.

To further refine the localization of the mutation, haplotype analysis was performed on the entire interval. Forty-three different haplotypes were identified among the affected individuals. Although the hypothesis tested by the TDT analysis was that intersexuality is determined by a single autosomal recessive locus, no shared segment was identified among the different “carrier” haplotypes. However one of the haplotypes (haplotype “8”) was shown to be more common than any other, and more than half of the 43 haplotypes had a shared segment of 2 to 30 adjacent markers, with haplotype “8”. Although these similarities are consistent with the strong association found in this study, some of the haplotypes identified in sex-reversed individuals were very different from the major haplotype “8”.

The second targeted analysis that we performed within the candidate interval was the re-sequencing of the SOX9 gene and TESCO region on different “carrier” and “control” chromosomes. The different elements regulating site- and stage-specific transcription of the SOX9 gene and all the core regulation elements are still not characterized in detail; however in 2008, Sekido and Lovell-Badge [26] identified a 3.2 kb element responsible for the testis-specific enhancer (TES) of Sox9, located 13 kb upstream of the mouse transcription start of the gene. A 1.4 kb core region within TES (TESCO) has been found to be highly conserved across mouse, rat, dog and human genomes, and a 180-bp sequence within TESCO is conserved even in amphibians [48]. No functional candidate mutations were detected within these sequences. The most informative results of this comprehensive analysis of the SOX9 gene and TESCO variability were (1) the absence of a shared segment between the different “carrier” chromosomes, and to the contrary (2) the presence of identical alleles among “carrier” and “control” chromosomes. Altogether, these results suggest that the causative mutation is not in the sequenced regions, meaning that it is not "in" or "near" the SOX9 gene or TESCO.

Up to now haplotype analysis and sequencing have not revealed a single haplotype associated with the causal mutation. Some affected individuals carry rare haplotypes very different from haplotype “8”. Different hypotheses can be proposed to explain these results: (1) First, these results could be explained if some individuals were incorrectly phenotyped as could be the case when managing collections from numerous breeding farms. Although an information sheet

SOX9 is Implicated in XX Sex-Reversal in Pigs.
describing the disease was drawn up to ensure that phenotyping was as consistent as possible, we cannot exclude that a few of the "affected animals" were in fact, unaffected. Pig intersex phenotypes range from true hermaphrodites to male phenotypes with or without ambiguities [4]. Most of the sex-reversed animals used in this study were included based on an external morphological description only, and for the TDT all the phenotypic variants were pooled in a single category of affected animals. The wide range of possible phenotypes indicates that the "expressivity" of the trait is variable, and a small number of animals might be poorly phenotyped and misclassified (2). Secondly the absence of a common segment shared by different "carrier" haplotypes could also be due to the informativeness of the markers used for the analysis. The data obtained so far indicate that none of the SNPs on the chip is in total linkage disequilibrium with the causal mutation. SNP markers on the Illumina PorcineSNP60 BeadChip were selected in order to achieve pan-genetic analyzes, regardless of the breed and whatever the nature of the traits of interest [49]. The LW breed was one of the seven economically important pig breeds used to validate the SNPs selected for the chip, and the highest number of polymorphic loci was obtained for the LW samples. All the families of the XX sex reversal design were from LW breed, which is the best situation (from the breed point of view) to take fully advantage of the informativeness of the markers on the chip. An additional quality criterion used to select the SNPs used on the chip was the MAF value obtained in a validation population panel. The average MAF for the SNPs on the chip is 0.274, and 90% of the SNPs have a MAF >0.15 (criteria of the three first waves of selection). The frequency of XX sex reversed animals in pig populations is less than 1%, and consequently the expected frequency of the mutated allele must be low. It was therefore unlikely that the SNPs on the chip would show strong linkage disequilibrium with the causal mutation. If the chip is of good quality for the analysis of purebred LW animals a priori, its use for mapping loci influencing rare trait variations is not necessarily optimal. This sub-optimal situation could explain why no common homozygous haplotype for all the XX sex reversed animals was identified (3). Thirdly, it is necessary to clarify whether allelic (or a genetic) heterogeneity is possible within the analyzed pedigree. The signal obtained on chromosome 12 in the region of SOX9 targets one mutation affecting gene regulation and certainly explains most cases of XX sex-reversed animals in this design. However, if a second independent gene mutation, also inducing a deregulation of SOX9, segregates in the pedigree, some animals could have two mutated copies but be double-heterozygous "affected/+" or "-affected/+" with two different haplotypes. In the SOX9 region, these individuals would be heterozygous for several polymorphisms. Finally, we also cannot exclude that another gene in another chromosomal region might induce an intersex phenotype that is therefore independent of SOX9. This hypothesis could not however be confirmed in the present design, as most individuals shared part of haplotype "8". Further analyses should be conducted by excluding these pigs, leading to a highly reduced number of affected pigs in the residual test population.

So far, the genetic determination of only a small percentage of sex-reversed pigs is known. The presence and expression of the SRY gene (due to chromosomal rearrangement or trisomy) does not however, explain the vast majority of the cases. Here we report the identification of a strong association with genetic markers near the SOX9 gene. To date, this result has not been reported in pigs and the identification of the causative mutation will add to the knowledge and understanding of sexual differentiation in mammals. The data published here support the continued exploration of the regions that regulate SOX9 and further mapping of the interval to find the causal mutation. At first, a special effort should be made to increase the power of the design by adding (1) additional individuals and families and (2) SNP markers. Having a larger number of cases would allow help to refine the phenotypes into different sub-categories and understand the variability of the phenotype expression by identifying other potential regulatory genes. In parallel the full re-sequencing of cases and controls in the candidate interval will provide more informative SNP markers presenting a higher LD with the causal mutation. However, the identification of the candidate "8" haplotype means that its frequency can now be estimated in different populations and therefore the number of potentially sex-reversed unexplained cases predicted. From a functional point of view, this haplotype can be used as of now to produce controlled crosses for a detailed exploration of the underlying physiological mechanisms.

Supporting Information

Figure S1. Frequency distributions of minor allele frequencies (MAF).
(TIF)

Figure S2. Cumulative frequency of SNP as a function of the heterozygous parents frequency.
(TIF)

Figure S3. Estimation of the linkage disequilibrium among the founders of the design.
(TIF)

Figure S4. Localization of the detected polymorphisms in TESCO and SOX9.
(TIF)

Table S1. List of primer pairs used for the amplifications of the SOX9 and TESCO regions.
(TIF)

Acknowledgements

We gratefully acknowledge BIOPORC gathering the French pig breeding companies (ADN, NUCLEUS, Gene+ and Pen Ar Lan) for their contribution in the constitution of the collection. We also thank LABOGENA for its technical assistance with the microarray genotyping. We thank J. Demars, H. Gilbert and P. multisant for helpful discussion and revision of the manuscript.
The authors also thank Byron C Jones for its precious help for the English revision of the manuscript.

**Author Contributions**

Conceived and designed the experiments: DM EP JR. Performed the experiments: SR NI. Analyzed the data: SR BS.

**References**

1. Sinclair AH, Berta P, Palmer MS, Hawkins JR, Griffiths BL et al. (1990) A gene from the human sex-determining region encodes a protein with homology to a conserved DNA-binding motif. Nature 346: 240–244. doi: 10.1038/346240a0. PubMed: 1695712.

2. Eggers S, Sinclair A (2012) Mammalian sex determination—insights from humans and mice. Chromosome Res 20: 215–238. PubMed: 22290220.

3. Hughes IA, Hous K, Ahmed SF, Lee PA (2006) Consensus statement on management of intersex disorders. J Pediatr Urol 2: 148–162. doi: 10.1016/jjpui.2006.03.004. PubMed: 18947601.

4. Pailloux E, Pelliniemi L, Barbosa A, Parma P, Kuopio T et al. (1997) Relevance of intersexuality to breeding and reproductive biotechnology programs; XX sex reversal in pigs. Theriogenology 47: 93–102. doi: 10.1016/S0093-691X(96)00343-3.

5. Pailloux E, Parma P, Sundström J, Vigier B, Servel N et al. (2001) Time course of female-to-male sex reversal in 38,XX fetal and postnatal pigs. Dev Dyn 222: 325–340. doi:10.1002/dvdy.1194. PubMed: 11747069.

6. Pailloux E, Vigier B, Chaffaux S, Servel N, Taourit S et al. (2001) A 11.7-kb deletion triggers intersexuality and polledness in goats. Nat Genet 29: 453–458. doi:10.1038/ng769. PubMed: 11726932.

7. Pailloux E, Vigier B, Vaiman D, Schibler L, Vaiman A et al. (2001) Contribution of domestic animals to the identification of new genes involved in sex determination. J Exp Zool 290: 700–708. doi: 10.1002/jez.1120. PubMed: 11748616.

8. Leal TL, McGee RB (2012) Disorders of sexual development in the domestic horse. Equus caballus. Sex Dev 6: 61–71. doi: 10.1159/000334048. PubMed: 22095202.

9. Meyers-Wallen VN, Schlafer D, Barr I, Lovell-Badge R, Keyzner A (1999) Sry-negative XX sex reversal in purebred dogs. Mol Reprod Dev 53: 266–273. doi:10.1002/(SICI)1098-2795(199907)53:3. PubMed: 10369387.

10. Meyers-Wallen VN (2012) Gonadal and sex differentiation abnormalities of dogs and cats. Sex Dev 6: 46–60. doi:10.1159/000332740. PubMed: 22005957.

11. Nagamine CM, Morohashi K, Carlisle C, Chang DK (1999) Sex reversal caused by Mus musculus domesticus Y chromosomes linked to variant expression of the testis-determining gene Sry. Dev Biol 216: 182–194. doi: 10.1006/dbio.1999.9436. PubMed: 10588871.

12. Whitworth DJ, Shaw G, Renfree MB (1996) Gonadal sex reversal of the developing marsupial ovary in vivo and in vitro. Development 122: 4057–4063. PubMed: 9012525.

13. Jiménez R, Burgos M, Sánchez A, Sinclair AH, Alarcón FJ et al. (1993) Fertile females of the mole Talpa occidentalis are phenotypic intersexes with ovotestes. Development 118: 1303–1307. PubMed: 8269856.

14. Parma P, Radi O, Vidal V, Chaboissier MC, Dellamora E et al. (2006) R-spondin1 is essential in sex determination, skin differentiation and malignancy. Nat Genet 38: 1304–1309. doi:10.1038/ng1907. PubMed: 17041800.

15. Berta P, Hawkins JR, Sinclair AH, Taylor A, Griffiths BL et al. (1999) Genetic evidence equating SRY and the testis-determining factor. Nature 348: 448–450. doi:10.1038/348448a0. PubMed: 22471409.

16. Huang B, Wang S, Ning Y, Lamb AN, Bartley J (1999) Autosomal XX sex reversal caused by duplication of SOX9. Am J Med Genet 87: 349–353. doi:10.1002/(SICI)1098-6562(19991203)87:4. PubMed: 10588843.

17. Bishop CE, Whitworth DJ, Qin Y, Agoulnin AI, Agoulnin IU et al. (2000) A transgenic insertion upstream of sox9 is associated with dominant XX sex reversal in the mouse. Nat Genet 26: 490–494. doi:10.1038/s6258. PubMed: 11101582.

18. Koopman P, Gubbay J, Vivian N, Goodfellow P, Lovell-Badge R (1991) Male development of chromosomally female mice transgenic for Sry. Nature 351: 117–121. doi:10.1038/351117a0. PubMed: 2030730.

19. Hunter RH (1996) Aetiology of intersexuality in female (XX) pigs, with novel molecular interpretations. Mol Reprod Dev 45: 392–402. doi: 10.1002/(SICI)1098-2795(199611)45:3. PubMed: 8916051.

20. Pannetier M, Elzaïat M, Thépôt D, Pailloux E (2012) Telling the story of XX sex reversal in the goat: highlighting the sex-crossroad in domestic mammals. Sex Dev 6: 33–45. doi: 10.1002/(SICI)1098-2795(199611)45:3. PubMed: 8916051.

21. Larzul C, Delaunay I, Schwob S, Mercat MJ (2008) Paramètres génétiques des principales anomalies congénitales porcines. 40èmes Journées de la Recherche Porcine, Paris FRANCE 40: 141–142.

22. Pailloux E, Mandon-Pepin B, Coulot C (2001) Mammalian gonadal differentiation: the pig model. Reprod Suppl 58: 65–80. PubMed: 11980203.

23. Risch N, Merikangas K (1996) The future of genetic studies of complex human diseases. Science 273: 1516–1517. doi:10.1126/science.273.5281.1516. PubMed: 8601636.

24. Charlier C, Coppicieters W, Rollin F, Desmecht D, Agerholm JS et al. (2008) Highly effective SNP-based association mapping and management of recessive defects in livestock. Nat Genet 40: 449–454. doi:10.1038/ng.96. PubMed: 18344986.

25. Lequarré AS, Andersson L, Andrè C, Fredholm M, Hitte C et al. (2011) LUPA: a European initiative taking advantage of the canine genome architecture for unravelling complex disorders in both human and dogs. Vet J 189: 155–159. doi:10.1016/j.tvjl.2011.06.013. PubMed: 2175267526.

26. Sekido R, Lovell-Badge R (2008) Sex determination involves synergistic action of SRY and SF1 on a specific SOX9 enhancer. Nature 453: 930–934. doi:10.1038/nature06944. PubMed: 18454134.

27. Purcell S, Neale B, Todd-Brown K, Thomas L, Ferreira MA et al. (2007) PLINK: a tool set for whole-genome association and population-based linkage analyses. Am J Hum Genet 81: 559–575. doi:10.1016/j.na.2007.05.010. PubMed: 17701901.

28. Lande R, Kruglyak L (1995) Genetic dissection of complex traits: guidelines for interpreting and reporting linkage results. Nat Genet 11: 241–247. doi:10.1038/31195-241. PubMed: 7581446.

29. Wang K, Li M, Hadley D, Liu R, Gissendracher J et al. (2007) PennCNV: An integrated hidden Markov model designed for high-resolution copy number variation detection in whole-genome SNP genotyping data. Genome Res, 17: 1665–1674. PubMed: 17921354.

30. Corpet F (1986) Multiplicity sequence alignment and hierarchical clustering. Nucleic Acids Res 16: 10881–10889. doi:10.1093/nar/16.22.10881. PubMed: 2849754.

31. Dupuis MC, Zhang Z, Druet T, Denux JO, Charlier C et al. (2011) Results of a haplotype-based GWAS for recurrent laryngeal neuropathy in the horse. Mamm Genome 22: 613–620. doi:10.1007/s00335-011-9337-3. PubMed: 216947232.

32. Gasca S, Canizares J, De Santa Barbara P, Mejean C, Poulat F et al. (2002) A nuclear export signal within the high mobility group domain of Sox9 is implicated in XX Sex-Reversal in Pigs.
determination in the mouse. Development 131: 1891–1901. doi: 10.1242/dev.01087. PubMed: 1505661537.

37. Lavery R, Lardenois A, Ranc-Jiannotamedi F, Pauper E, Gregoire EP et al. (2011) XY Sox9 embryonic loss-of-function mouse mutants show complete sex reversal and produce partially fertile XY oocytes. Dev Biol 354: 111–122. doi:10.1016/j.ydbio.2011.03.029. PubMed: 21466799.

38. Vidal VP, Chaboissier MC, De Rooij DG, Schedl A (2001) Sox9 induces tests development in XX transgenic mice. Nat Genet 28: 216–217. doi:10.1038/90046. PubMed: 11431689.

39. Foster JW, Dominguez-Steglich MA, Guioli S, Kwok C, Weller PA et al. (1994) Campomelic dysplasia and autosomal sex reversal caused by mutations in an SRY-related gene. Nature 368: 525–530. PubMed: 7990924.

40. Wagner T, Wirth J, Meyer J, Zabel B, Held M et al. (1994) Autosomal sex reversal and campomelic dysplasia are caused by mutations in and around the SRY-related gene SOX9. Cell 79: 1111–1120. doi: 10.1016/0092-8674(94)90041-8. PubMed: 8001137.

41. Cox JJ, Willatt L, Homfray T, Woods CG (2011) A SOX9 duplication and familial 46,XX developmental testicular disorder. N Engl J Med 364: 91–93. doi:10.1056/NEJMc1010311. PubMed: 21208124.

42. Vetro A, Ciccone R, Giorda R, Patricelli MG, Della Mina E et al. (2011) males SRY negative: a confirmed cause of infertility XX. J Med Genet 48: 710–712 doi:10.1136/jmedgenet-2011-100036. PubMed: 21653197.

43. White S, Ohnesorg T, Notini A, Roeszler K, Hewitt J et al. (2011) Copy number variation in patients with disorders of sex development due to 46,XY gonadal dysgenesis. PLOS ONE 6: e17793. doi:10.1371/journal.pone.0017793. PubMed: 21408189.

44. Jakubiczka S, Schröder C, Ullmann R, Volleth M, Ledig S et al. (2010) Translocation and deletion around SOX9 in a patient with acampomelic campomelic dysplasia and sex reversal. Sex Dev 4: 143–149. doi: 10.1159/000302403. PubMed: 20453475.

45. Bagheri-Fam S, Barrionuevo F, Dohrmann U, Günther T, Schüle R et al. (2006) Long-range upstream and downstream enhancers control distinct subsets of the complex spatiotemporal Sox9 expression pattern. Dev Biol 291: 382–397. doi:10.1016/j.ydbio.2005.11.013. PubMed: 16458883.

46. Wang J, Jiang J, Fu W, Jiang L, Ding X et al. (2012) A genome-wide detection of copy number variations using SNP genotyping arrays in swine. BMC Genomics 13: 273. doi:10.1186/1471-2164-13-273. PubMed: 22726314.

47. Ramayo-Cal das Y, Castelló A, Pena RN, Alves E, Mercadé A et al. (2010) Copy number variation in the porcine genome inferred from a 60 k SNP BeadChip. BMC Genomics 11: 593. doi: 10.1186/1471-2164-11-593. PubMed: 20969757.

48. Bagheri-Fam S, Sreenivasan R, Bernard P, Knowler KC, Sekido R et al. (2012) Sox9 gene regulation and the loss of the XY/XX sex-determining mechanism in the mole vole Ellobius lutescens. Chromosome Res 20: 191–199. doi:10.1007/s10577-011-9269-5. PubMed: 22215485.

49. Ramos AM, Crooijmans RP, Affara NA, Amaral AJ, Archibald AL et al. (2009) Design of a high density SNP genotyping assay in the pig using SNPs identified and characterized by next generation sequencing technology. PLOS ONE 4: e6524. doi:10.1371/journal.pone.0006524. PubMed: 19654876.