

# Analysis of candidate genes for prolificacy in two commercial sow populations



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## Background

So far, at least 54 quantitative trait loci (QTL) and twelve candidate genes with association for reproductive traits in sows have been identified (Buske et al. 2006; Figure 1). In our approach, the genes encoding **Properdin (BF)**, **Glutathione peroxidase 5 (GPX5)**, **Fucosyltransferase 1 (FUT1)** and **Estrogen receptor 2 (ESR2)** were chosen as candidate genes to search for genetic variation influencing prolificacy in two different commercial sow populations. These genes have physiological relevance and are located in chromosomal regions, which have shown genetic effects on prolificacy in crossbreed populations.

## Material and Methods

**Traits:** Total number born (TNB) piglets; Number born alive (NBA) piglets.

**Animal populations:** *Population 1:* 129 F<sub>1</sub>-hybrid, paternal half-sib sows (German Landrace sows x Duroc boar) with at least four litters.

*Population 2:* 61 F<sub>2</sub>-sows with extreme high (TNB 15.7±2.2, NBA 14.1±2.1) and 62 F<sub>2</sub>-sows with extreme low litter size (TNB 10.4±2.4, NBA 9.8±2.3) of a cross between (Large White x Landrace) sows x Leicoma boars were selected (second to fourth litter) from a population comprising 447 F<sub>2</sub>-sows. All sows were mated with Piétrain boars to produce fattening piglets.

**Genotyping:** Gene tests were adopted from Jiang and Gibson (1998) for BF, Bertani et al. (1999) for GPX5, Klukowska et al. (1999) for FUT1, and Muñoz et al. (2004) for ESR2.

**Statistical analyses:** In *population 1*, the effect of genotype on TNB and NBA piglets was studied using analysis of variance (ANOVA).

In *population 2*, the Chi-square test with the extension of the Fisher's Exact test was used to test differences in the distribution of genotypes between the two extreme performance groups. Additionally, ANOVA including the Duncan test was used to investigate whether a special genotype was associated with TNB or NBA piglets (Buske et al. 2005).

## Results

*Population 1:* Significant association of the ESR2 gene variants was found for NBA piglets of the F<sub>1</sub>-sows (Table 1). Sows with the AB genotype had on average 0.68 more NBA piglets than sows with the AA genotype (p=0.034). No BB genotype was detectable, because the sire was AA homozygous. For the other genes, no association with litter size parameters was observed, possibly due to extremely unbalanced genotype frequencies.

**Table 1.** *Population 1:* Association analysis between genotypes and litter size parameters (TNB, NBA) for the ESR2 gene in 129 F<sub>1</sub> half-sib sows (German Landrace x Duroc)

Genotype	No. of sows	No. of litters	LS Means TNB (≥ 2. litter)	SE	LS Means NBA (≥ 2. litter)	SE
AA	42	162	12.02	0.32	10.96 <sup>a</sup>	0.30
AB	87	329	12.60	0.25	11.64 <sup>b</sup>	0.23

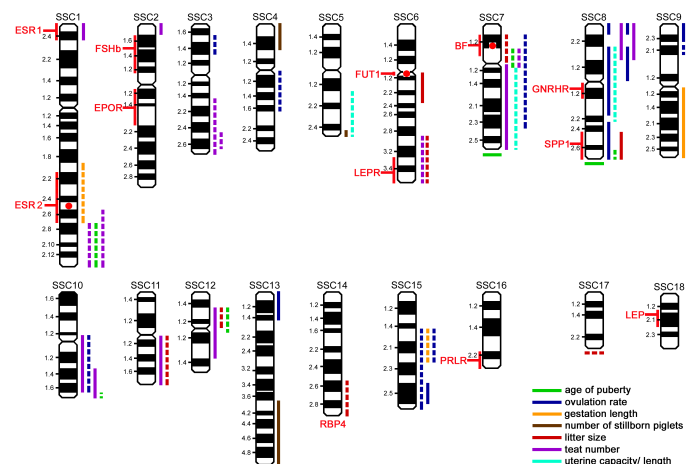
a,b LS Means in the same column with different superscripts significantly differ at p< 0.05. SE = Standard error; TNB = Total number born piglets; NBA = Number born alive piglets.

\* For each sow, second up to fourth or fifth (when available) litter was analyzed. First litter was ignored.

## Conclusions

Concerning the ESR2 gene, an association with NBA piglets was found in *population 1*. The FUT1 gene was associated with litter size in *population 2*, in which sows of extreme performance groups were genotyped. However, it must be stated that the observed associations may also represent effects of linked genes. The results clearly show that population specific knowledge of gene effects may be helpful for more efficient selection strategies. To evaluate gene effects of candidate genes on litter size parameters more reliably, an improvement would be to perform an additional investigation with planned matings to obtain all genotypes in balanced genotype frequencies in the sow population. Furthermore, a population-wide candidate gene approach with a larger data set would be desirable to avoid overestimation of effects, which can occur by chance in association analyses with comparatively low animal numbers.

**Figure 1.** QTL and candidate genes with association to prolificacy in pigs



Red circles within the chromosome = investigated genes for this study; bold solid lines = level of significance p < 0.05; dashed lines = level of significance p > 0.05; cytogenetic positions of the lines at the end of the chromosomes and for RBP4 were not evaluable.

BF = properdin; ESR1 = estrogen receptor 1; ESR2 = estrogen receptor 2; EPOR = erythropoietin receptor; FSHb = follicle stimulating hormone beta; FUT1 = fucosyltransferase 1; GNRHR = gonadotropin releasing hormone receptor; LEPR = leptin; LEPR = leptin receptor; PRLR = prolactin receptor; RBP4 = retinol-binding protein 4; SPP1 (OPN) = secreted phosphoprotein 1

Reference: Buske et al. (2006); modified

*Population 2:* The analysis of genotype distribution pattern among F<sub>2</sub> hybrid sows with extremely high and low litter size has identified significant differences for the FUT1 gene. The number of sows with the AB genotype was significantly increased in the high performance group in comparison to the low performance group (Table 2). For the BF gene, the A allele was the unfavorable one, leading to significantly less offspring (ANOVA; Duncan test).

**Table 2.** *Population 2:* Genotype distribution of candidate genes in groups of F<sub>2</sub> hybrid sows (Large White x Landrace) x Leicoma with extremely high and low litter size

Genotype	BF Performance		GPX5 Performance		FUT1* Performance		ESR2 Performance	
	Low	High	Low	High	Low**	High**	Low	High
AA	3	0	4	3	2	1	8	8
AB	13	7	24	17	11	24	54	53
BB	46	54	34	41	48	34	0	0

\* Three sows could not be genotyped for the FUT1 gene.

\*\* Significantly different at p< 0.05, Chi-square-test.

As estrogen receptors are important for prolificacy and the investigated polymorphism in the ESR2 gene is located in a coding region leading to the amino acid exchange Met 949 Val, ESR2 can be considered as a potential genetic marker to enhance NBA piglets of sows in future analyses.

## References

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**Acknowledgement:** We thank the H. Wilhelm Schaumann Stiftung (Hamburg, Germany) for the grant for B. Buske