A Substitution in the Ligand Binding Domain of the Porcine Glucocorticoid Receptor Affects Activity of the Adrenal Gland

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Abstract

Glucocorticoids produced in the adrenal cortex under the control of the hypothalamic-pituitary axis play a vital role in the maintenance of basal and stress-related homeostasis and influence health and well-being. To identify loci affecting regulation of the hypothalamic-pituitary-adrenal (HPA) axis in the pig we performed a genome-wide association study for two parameters of acute and long-term adrenal activity: plasma cortisol level and adrenal weight. We detected a major quantitative trait locus at the position of the glucocorticoid receptor gene (NR3C1) – a key regulator of HPA axis activity. To determine the causal variant(s), we resequenced the coding region of NR3C1 and found three missense single nucleotide polymorphisms (SNPs). SNP c.1829C>T, leading to a p.Ala610Val substitution in the ligand binding domain, showed large (about 0.6× and 1.2× phenotypic standard deviations for cortisol level and adrenal weight, respectively), and highly significant (2.1E-39≤log10(1/p)≤1.7E+0) negative effects on both traits. We were able to replicate the association in three commercial pig populations with different breed origins. We analyzed effects of the p.Ala610Val substitution on glucocorticoid-induced transcriptional activity of porcine glucocorticoid receptor (GR) in vitro and determined that the substitution introduced by SNP c.1829C>T increased sensitivity of GR by about two-fold. Finally, we found that non-coding polymorphisms in linkage disequilibrium with SNP c.1829C>T have only a minor effect on the expression of NR3C1 in tissues related to the HPA axis. Our findings provide compelling evidence that SNP c.1829C>T in porcine NR3C1 is a gain-of-function mutation with a major effect on the activity of the adrenal gland. Pigs carrying this SNP could provide a new animal model to study neurobiological and physiological consequences of genetically based GR hypersensitivity and adrenal hypofunction.

Introduction

Hormones produced by the adrenal gland (corticosteroids and catecholamines) play a vital role in maintaining homeostasis, particularly during stress. Glucocorticoids (in the pig, cortisol) produced by adrenocortical cells of the zona fasciculata under the control of the hypothalamic-pituitary axis facilitate coping with stress and adaptation by influencing various neurobiological, metabolic, and immune processes [1]. Dysfunction of the hypothalamic-pituitary-adrenal (HPA) axis therefore has adverse effects on health status and well-being. In humans, for example, dysregulation of glucocorticoid secretion and signaling has been implicated in the pathogenesis of mood disorders and metabolic syndrome [2,3].

In different species, including the pig, glucocorticoid secretion shows large inter-individual variation that has a considerable genetic component [4,5]. Little is known about the underlying genetic variants; however, their identification and utilization as simple DNA markers in molecular breeding is a promising approach to improving adaptation potential, health, and welfare in pigs and in farm animals in general [6]. Indeed, implementation of traditional breeding programs for these traits, based on phenotypic data, is hampered by their difficult and expensive ascertainment [7].

To identify genes affecting acute and long-term regulation of HPA axis activity in the pig, we previously analyzed polymorphisms in ten functional candidates for potential associations with plasma cortisol level and adrenal weight (herein summarily designated as adrenal activity) in commercial crossbreds. The glucocorticoid receptor gene (nuclear receptor subfamily 3, group C, member 1; NR3C1) was identified as a promising candidate because we found significant association of a non-coding variant (SNP c.*2122G>A) with both traits analyzed [8]. Glucocorticoid receptor (GR) is a ligand-activated transcription factor transducing glucocorticoid signals in almost all organs and tissues, including hypothalamus and pituitary where it mediates feedback inhibition of HPA axis activity by downregulating expression and release of corticotropin-releasing hormone (CRH) and corticotropin...
In humans several rare mutations and common polymorphisms of NR3C1, mainly non-synonymous, have been shown to generate diversity in GR signaling and to contribute to inter-individual variation in glucocorticoid secretion [10,11].

To facilitate a more comprehensive understanding of the genetic architecture of HPA axis activity, here we performed a genome-wide association study (GWAS). This uncovered NR3C1 as a major QTL for plasma cortisol level and adrenal weight.

To identify the causal variant(s) we first concentrated on the analysis of non-synonymous polymorphisms of NR3C1 because they appear to be the main source of inter-individual variation in GR function, and because they are most amenable to functional interpretation. We present here strong genetic and functional evidence that a non-synonymous SNP (c.1829C>T) in NR3C1, leading to the gain-of-function substitution p.Ala610Val in the ligand binding domain of GR, is indeed causally involved.

Results

A genome-wide association study identifies a major quantitative trait locus for adrenal activity at the NR3C1 position

The GWAS was performed in a population of commercial German Landrace pigs (n = 564), which were genotyped using the PorcineSNP60 BeadChip (Illumina) and phenotyped with respect to plasma cortisol level and adrenal weight (overview of all genotypic information for 42,359 SNPs that mapped to the 18 autosomes [Scrofa 10.2] was used). A total of 37 SNPs mapping to nine chromosomes were significantly associated with plasma cortisol level at the genome-wide false discovery rate (FDR) q ≤0.05 level (Figure 1A and Table S2). The strongest association signals were found on chromosomes 2 and 7. On chromosome 2 nine significant SNPs, located in an ∼2.6 Mb region between SNPs ALGA0106239 and ALGA0123033, were identified. The two most significantly associated (p = 4.0E-08) SNPs, ALGA0106239 and DRGA0017574, showed complete linkage disequilibrium, and each explained ∼10% of the phenotypic variance (Table S2). On chromosome 7 seven significant SNPs, located in an ∼4.9 Mb region between SNPs H3GA0023063 and MARCO004680, were found. The most significantly associated SNP (p = 3.4E-08), ASGA0036275, explained ∼10% of the phenotypic variance (Table S2).

The GWAS for adrenal weight yielded 62 genome-wide significant SNPs (q≤0.05), located on seven chromosomes (Figure 1B and Table S3). The strongest association signal was found on chromosome 2, where 41 significant SNPs clustered in an ∼9.7 Mb region between SNPs M1GA0024750 and ALGA0123873. However, the majority of the SNPs (39), including all nine SNPs significantly associated with plasma cortisol level, fell in a considerably smaller region (∼3 Mb) between SNPs ALGA0106306 and ALGA0120126. Except on chromosome 2, there was no overlap between the association signals for plasma cortisol level and adrenal weight. SNPs ALGA0106239 and DRGA0017574 again showed the strongest association (p = 3.5E-08), and each explained ∼30% of the phenotypic variance in adrenal weight. Significant SNPs in other genomic regions explained less than 7% of the phenotypic variance (Table S3).

According to the current annotation of the porcine genome (Pore Ensembl Scrofa 10.2) the ∼3 Mb region flanked by SNPs ALGA0106306 and ALGA0120126 contains eight annotated genes, including NR3C1. The two most significant SNPs, ALGA0106239 and DRGA0017574, are located within porcine NR3C1, in introns 6 and 5, respectively.

The GWAS thus confirmed association of NR3C1 with plasma cortisol level and adrenal weight in an additional, purebred, population and suggested that polymorphisms of NR3C1 account for the largest proportion of the variation in adrenal activity.

Resequencing of the region encoding the porcine glucocorticoid receptor alpha isoform reveals a potentially functional missense variant

We resequenced the whole ∼2.38 kb region of NR3C1 encoding the predominant isoform of GR–alpha–in a panel of samples from four different breeds (two each, German Landrace (LR), German Large White (LW), Pietrain (Pi), and Duroc (Du)) and in pooled samples of crossbred animals (Pietrain × German Large White × German Landrace), noted as PiF1 with extreme phenotypes for cortisol secretion and adrenal weight, respectively (three each for high and low cortisol and heavy or light adrenal gland). We identified a total of six SNPs (Figure 2); three synonymous (c.24C>T, c.660G>A, c.1734A>G) and three missense polymorphisms (c.39A>C, c.55G>C, c.1829C>T).

The missense SNP c.39A>C and c.55G>C lead to amino acid substitutions (p.Glu13Asp and p.Val19Leu, respectively) in the N-terminal domain in a region of GR-alpha showing a relatively low level of amino acid sequence conservation (Figure S1) [12]. The third missense SNP, c.1829C>T, leads to an alanine-to-valine substitution (p.Ala610Val) in helix 5 of the ligand binding domain (GR-LBD) (Figure S1). Crystallographic and mutagenesis studies of human and murine GR-LBD reveal that helix 5 is involved in the formation of the ligand binding pocket of GR-alpha [13] and suggest that p.Ala610Val substitution might enhance affinity and, consequently, responsiveness of GR to glucocorticoids by intensifying van der Waals contacts with the ligand [14].

SNP c.1829C>T shows consistent association with cortisol level and adrenal weight across different pig breeds

We sought to assess frequency of the missense SNPs in the most important commercial breeds and in populations included in association analyses, specifically in the LR population initially used for the GWAS and in LW and PiF1, which were also phenotyped with respect to plasma cortisol level and adrenal weight. We therefore genotyped 21 to 35 unrelated LR, LW, Pi, and Du pigs. While SNP c.1829C>T segregated in all four breeds, SNP c.55G>C segregated only in Pi. Moreover, in contrast to LR, LW, and Du, the Pi breed was fixed at SNP c.39A>C. With the exceptions of c.39A>C in Du and c.55G>C in Pi, all three missense SNPs segregated at relatively low frequency, generally about 10% or less. Genotyping of additional 92 LR and 45 LW pigs (in total 127 and 69 individuals, or 15% and 25% of the two populations, respectively) again revealed no carriers of SNP c.55G>C. Therefore, SNP c.55G>C was assumed to be specific for the Pietrain breed. Consequently, SNP c.39A>C and c.1829C>T were genotyped in all three populations (total n = 834 for LR, n = 274 for LW, and n = 537 for PiF1) used for association analyses; SNP c.55G>C was genotyped only in PiF1. In LR and LW the genotype of SNP c.55G>C was assumed to be homozygous GG. Allele distribution of the missense SNPs in the different populations is summarized in Table S4. None of the populations of purebred pigs showed significant deviation from Hardy-Weinberg equilibrium (HWE; p>0.05), indicating that the missense SNPs had no negative impact on viability of their carriers. In PiF1 SNP c.55G>C and c.1829C>T showed
significant deviation from HWE (p<0.05), which is most likely a result of crossbreeding because we frequently observe deviation from HWE in this cross (see [8]).

Using the genotypic information collected in LR, LW, and PIF1 we performed population-based single-marker association analysis. Whereas SNP c.39A>C and c.55G>C showed no or only inconsistent associations, SNP c.1829C>T was consistently and, in general, highly significantly associated with decreased cortisol level and adrenal weight in all three populations (Tables 1,2). The estimated allele substitution effect was—with an average of about -0.6 phenotypic standard deviations for cortisol level and about -1.2 phenotypic standard deviations for adrenal weight, respectively—very pronounced. Heterozygous carriers generally showed intermediate values, indicating that SNP c.1829C>T is associated with an additive effect. The estimated phenotypic variance explained by SNP c.1829C>T was ~10% for cortisol level and ~25% for adrenal weight, in a similar range as found for the QTL detected by the GWAS.

The genotype-based analysis was complemented by haplotype-based association analysis. To examine whether the missense polymorphisms account for the previously found association of SNP c.*2122G>A with plasma cortisol level and adrenal weight, we also included genotype information of this SNP in the haplotype analysis (for more information see [9]). Based on these data, a total of five common haplotypes (frequency >1%) were inferred (Table S3). Each of the missense variants was located on a separate haplotype. We compared the four common haplotypes carrying allele A at the SNP c.*2122G>A, including the three haplotypes carrying the missense polymorphism, with the wild-type haplotype carrying the G allele, by means of the haplotype trend regression test. Estimated haplotype substitution effects are presented in Table 3. In accordance with the genotype-based analysis, haplotype A-G-T-A, carrying the derived T allele of SNP c.1829C>T, stood out as the only one showing consistent effects, significantly decreasing cortisol level and adrenal weight with similar magnitudes in all three populations. This finding suggests that SNP c.1829C>T is most likely underlying the previously found association of SNP c.*2122G>A with cortisol and adrenal weight [8].

Finally, we performed linkage disequilibrium (LD) and haplotype block analysis of the 3 Mb region between SNPs ALGA0106239 and DRGA0017574 on chromosome 2 using data from the GWAS on the 35 significant and 36 intervening SNPs, and SNP data from genotyping NR3C1 in the LR population. The results are presented in Figure S2. The three segregating NR3C1 SNPs, c.39A>G, c.1829C>T, and c.*2122G>A, formed a 102 kb haplotype block spanning the NR3C1 gene, together with three GWAS SNPs, including the two most significant SNPs, ALGA0106239 and DRGA0017574. Among all analyzed SNPs, ALGA0106239 and DRGA0017574 showed the highest LD (r² = 0.781) with SNP c.1829C>T.

Together, these results provide strong genetic evidence that SNP c.1829C>T is responsible for the association of NR3C1 with plasma cortisol level and adrenal weight indicated in the present study by the GWAS and previously by the analysis of SNP c.*2122G>A [8].

The p.Ala610Val substitution increases sensitivity of porcine GR to dexamethasone in vitro

As mentioned above, mutagenesis studies of GR indicated that the p.Ala610Val substitution might influence responsiveness of GR.
to glucocorticoids, which, in view of the key role of GR in feedback regulation of HPA axis activity, might explain phenotypic effects associated with SNP c.1829C>T. To examine the impact of the p.Ala610Val substitution on transcriptional activity of porcine GR, we compared the ability of the two allelic receptor variants to drive luciferase expression from the mouse mammary tumor virus promoter (MMTV) in COS-7 cells after stimulation with various dexamethasone concentrations (Figure 3). The transactivation assay revealed that the p.Ala610Val substitution enhances transcriptional activity of porcine GR in a dose-dependent manner ($p = 0.007$). The two GR variants have similar baseline transcriptional activity (Figure 3A). At 0.1 and 1 nM dexamethasone, respectively, the p.610Val variant induced 2-fold and 1.25-fold higher luciferase expression compared to the wild-type p.610Ala variant; however, at 0.1 nM dexamethasone the difference reached significance ($p = 0.012$). At higher concentrations of dexamethasone, transcriptional activity reached a plateau that was similar for both GR variants. Analysis of the dose-response curves (Figure 3B) revealed that the half-maximal effective concentration (EC50) of dexamethasone was 2-fold lower for the p.610Val variant compared to the p.610Ala variant (0.48 nM and 0.94 nM, respectively; $p = 0.026$). These results demonstrate that the p.610Val variant of porcine GR has increased sensitivity to glucocorticoids compared to the p.610Ala variant and, thus, that SNP c.1829C>T is a gain-of-function mutation.

Searching for functional coding and non-coding polymorphisms of NR3C1 associated with SNP c.1829C>T

To further prove causality of SNP c.1829C>T, we investigated whether other polymorphisms in LD might affect function of NR3C1 and, thus, potentially influence adrenal activity. We resequenced the region of NR3C1 encoding the GR-alpha isoform in three additional individuals (one each PiF1, LR, and LW)

### Table 1. Association of the three identified missense SNPs in porcine NR3C1 with plasma cortisol level (ng/ml) in three different commercial populations.

| SNP     | Population1 (n) | p-value | LSM ± SE2 (n) | a3 (% var4) |
|---------|----------------|---------|---------------|-------------|
| c.39A>C |                |         |               |             |
|         | AA             | 0.096   | 74.2 ± 1.8 (765) | 73.4 ± 7.0 (21) |
|         | AC             | 0.048   | 93.7 ± 1.8 (444) | 81.5 ± 6.0 (28) |
|         | CC             | 0.347   | 81.4 ± 3.6 (218) | 89.8 ± 9.0 (15) |
| c.55G>C |                |         |               |             |
|         | GG             | 0.339   | 91.8 ± 2.1 (291) | 94.8 ± 2.6 (181) |
|         | GC             | 0.018   | 85.3 ± 3.7 (186) | 70.3 ± 6.0 (44) |
|         | CC             | 0.018   | 85.3 ± 3.7 (186) | 70.3 ± 6.0 (44) |
| c.1829C>T |             |         |               |             |
|         | CC             | 4.2E-14 | 78.1 ± 2.9 (121) | 31.9 ± 13.6 (4) |
|         | CT             | 1.0E-06 | 95.5 ± 4.7 (45)  | 55.3 ± 11.4 (7) |
|         | TT             | 0.02    | 2.6 ± 0.0 (3)    | 5.3 ± 0.0 (3)   |

1LR-German Landrace, PiF1-(Pietrain x German Large White x German Landrace), LW-German Large White.
2Least-squares means (LSM) with different superscripts a,b,c differ significantly at $p<0.01$.
3Allele substitution effect in fractions of phenotypic standard deviation.
4Phenotypic variance in percent explained by the SNP.

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### Table 2. Association of the three identified missense SNPs in porcine NR3C1 with weight of the left adrenal gland (g) in three different commercial populations.

| SNP     | Population1 (n) | p-value | LSM ± SE2 (n) | a3 (% var4) |
|---------|----------------|---------|---------------|-------------|
| c.39A>C |                |         |               |             |
|         | AA             | 0.394   | 2.3 ± 0.03 (658) | 2.23 ± 0.11 (15) |
|         | AC             | 0.889   | 2.3 ± 0.03 (377) | 2.29 ± 0.08 (18) |
|         | CC             | 0.002   | 2.3 ± 0.04 (193) | 2.6 ± 0.10 (15) |
| c.55G>C |                |         |               |             |
|         | GG             | 0.075   | 2.28 ± 0.03 (250) | 2.35 ± 0.03 (145) |
|         | GC             | 0.018   | 85.3 ± 3.7 (186) | 70.3 ± 6.0 (44) |
|         | CC             | 0.018   | 85.3 ± 3.7 (186) | 70.3 ± 6.0 (44) |
| c.1829C>T |             |         |               |             |
|         | CC             | 2.1E-39 | 2.4 ± 0.02 (563) | 1.76 ± 0.19 (3) |
|         | CT             | 4.3E-17 | 2.35 ± 0.02 (358) | 1.52 ± 0.12 (6) |
|         | TT             | 1.6E-12 | 2.43 ± 0.04 (163) | 1.96 ± 0.23 (2) |

1LR-German Landrace, PiF1-(Pietrain x German Large White x German Landrace), LW-German Large White.
2Least-squares means (LSM) with different superscripts a,b,c differ significantly at $p<0.01$.
3Allele substitution effect in fractions of phenotypic standard deviation.
4Phenotypic variance in percent explained by the SNP.

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homozygous for the A-G-T-A haplotype; apart from SNP c.1829C>T, no other coding polymorphisms (missense or synonymous) were found.

To detect possible cis-acting effects of non-coding polymorphisms in LD with SNP c.1829C>T on expression of NR3C1 in tissues related to the HPA axis, we analyzed its allelic expression imbalance (AEI) in hippocampus, hypothalamus, pituitary, and whole adrenal gland by pyrosequencing using ten samples per tissue (five samples for each diplotype A-G-T-A/A-G-C-G and A-G-T-A/A-G-C-A). In all four examined tissues, expression of the T allele was significantly higher compared to the C allele (Figure 4). However, the magnitude of the expression imbalance was very low, ranging between 1.05 [log2(AEI) = 0.07] and 1.10-fold [log2(AEI) = 0.13]. Further, allelic ratios in genomic DNA fluctuated tightly around one [i.e., log2(T/C)<0.0], indicating that the porcine NR3C1 locus is not affected by copy-number variation (Figure 4).

Finally, to determine whether SNP c.1829C>T is in LD with a splicing polymorphism affecting structure of the GR-alpha isoform, we designed overlapping amplicons covering the cDNA sequence of NR3C1 from a putative porcine homolog to human exon 1C (which shows widespread expression in humans [15]) and through exons 2–8 to exon 9-alpha (Figure 5). Using cDNA from adrenal gland and pituitary of three individuals each homozygous for the carrier haplotype A-G-T-A and the wild-type haplotype A-G-C-G, the overlapping fragments were amplified, electrophoretically separated (Figure 5), and sequenced. We found no haplotype-specific differences in the splicing pattern, ruling out the existence of splicing polymorphisms in LD with SNP c.1829C>T.

### Table 3. Haplotype effects of porcine NR3C1 on plasma cortisol level (ng/ml) and adrenal weight (g) in three different commercial populations.

| Haplotype | Population | Cortisol level | p-value | Adrenal weight | p-value |
|-----------|------------|----------------|---------|----------------|---------|
| A-G-C-A   | LR         | 0.424          | 0.07    | 3.0E-4         |         |
|           | PiF1       | 0.01±0.26      | 0.998   | 0.03           | 0.003   |
|           | LW         | 6.12±5.50      | 0.269   | 0.03           | 0.583   |
| C-G-C-A   | LR         | 0.94±6.73      | 0.942   | 0.04           | 0.639   |
|           | PiF1       | 0.95±6.29      | 0.134   | 0.06           | 0.421   |
|           | LW         | 5.50±8.74      | 0.531   | 0.21           | 0.021   |
| A-C-C-A   | PiF1       | 0.54±3.01      | 0.859   | 0.02           | 0.456   |
| A-G-T-A   | LR         | 21.53±2.70     | 8.4E-15 | 0.46           | 5.2E-35 |
|           | PiF1       | 19.87±4.01     | 7.6E-7  | 0.43           | 6.4E-18 |
|           | LW         | 14.43±5.24     | 0.007   | 0.44           | 1.4E-11 |

1 Haplotype c.39-c.55-c.1829-c.*2122.
2 LR-German Landrace (n = 673), PiF1-(Pietrain × (German Large White × German Landrace) (n = 391), LW-German Large White (n = 208).
3 Estimated substitution effects ± standard errors of indicated haplotypes compared to haplotype A-G-C-G.

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Figure 3. The p.Ala610Val substitution increases sensitivity of porcine GR to glucocorticoids in vitro. Transcriptional activity of the two GR-alpha variants was investigated in COS-7 cells cotransfected with the pGL4.36 reporter construct expressing firefly luciferase under the control of the glucocorticoid-inducible mouse mammary tumor virus promoter after induction with dexamethasone at the indicated concentrations. (A) Fold induction of normalized luciferase expression in dexamethasone- vs. vehicle-treated cells. Significant differences (p<0.05) are indicated by an asterisk; (B) Estimated dose-response curve and EC50. Data are means ± SEM of four separate experiments performed in triplicate.

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Collectively, our data showed no considerable effects of non-coding polymorphisms in LD with SNP c.1829C>T on the function of porcine NR3C1, which leaves SNP c.1829C>T as the most likely causal variant.

Discussion

Genetic studies of the inter-individual variation in HPA axis activity are comparatively rare; thus, knowledge of its molecular genetic background is limited in most species. Here, we contribute to the understanding of the genetics of this axis in the pig by identifying a polymorphism in NR3C1 with a major effect on adrenal activity and by detecting several other genomic regions/QTL affecting plasma cortisol level and adrenal weight, most of which are novel.

Identification of a robustly replicating SNP-trait association is a crucial first step in identifying phenotypically causal genetic variants [16]. We show that a missense SNP c.1829C>T in NR3C1 and the carrier haplotype are consistently and strongly associated with variation in plasma cortisol levels and adrenal weight in three populations with different breed origins. Importantly, not only direction, but also magnitudes, of the effects of SNP c.1829C>T were highly reproducible, implying either direct involvement of this SNP, or, at least, very tight linkage disequilibrium between the SNP and the causal variant.

Functional testing of candidate causal variants provides valuable clues to differentiate between phenotypically important and co-segregating, neutral genetic variant(s) [17]. Our experimental data demonstrated that SNP c.1829C>T is a functional mutation, increasing sensitivity of the porcine GR to glucocorticoids in vitro by about two-fold. Although the overall transactivation capacity of GR is not affected by the p.Ala610Val substitution, due to its increased sensitivity, the p.610Val variant becomes activated at lower glucocorticoid levels. This earlier activation could be expected to enhance feedback inhibition of the HPA axis activity and, consequently, lead to reduced secretion of ACTH, which is a major regulator of adrenal function and growth [18]. Inhibition of HPA axis activity is most likely persistent, as indicated by the reduction of adrenal weight associated with SNP c.1829C>T. A negative relationship between GR sensitivity and HPA axis activity was recently described by Zhang et al. [19] who observed markedly decreased basal and post-stress ACTH and corticosterone levels, reduced adrenal weight, and adrenocortical thinning in a knock-in mouse line carrying an artificial gain-of-function substitution, p.Met610Leu, in GR. Notably, the substitution engineered by Zhang et al. [20] is located in the GR-LBD next to the position corresponding with the p.Ala610Val substitution in the porcine GR-LBD (i.e., p.Ala611Val in murine GR). This substitution induces similar dose-dependent increases in transactivation activity of GR in vitro [20]. Hence, the p.Met610Leu knock-
in mouse model provides substantial support for a causal role of SNP c.1829C>T.

However, in contrast to a knock-in mouse model, in an outbred species like the pig variation in a complex trait usually cannot be unambiguously attributed to a particular genetic variant, at least not before contribution of other variants in LD is considered. The only other detected missense polymorphisms of \(NR3C1\), SNPs c.39A>C and c.55G>C, were located on different haplotypes and showed no or only inconsistent effects on adrenal activity. Consequently, assuming that other variants of \(NR3C1\) in LD with SNP c.1829C>T affect adrenal activity, these necessarily have to be functional non-coding polymorphisms. For their detection we followed an experimental approach and analyzed association of SNP c.1829C>T with allele expression imbalance and splicing of \(NR3C1\) in tissues related to HPA axis. We found no evidence for splicing polymorphisms and only modest allele expression imbalance of \(NR3C1\), which argues against any significant contribution of polymorphisms in LD with SNP c.1829C>T to the variation in adrenal activity, especially in view of evidence from genome-wide association studies that non-coding variants tend to have weak phenotypic effects [17]. However, we cannot formally exclude existence of other contributing genetic variants acting in other temporally-spatial or functional contexts.

Taken together, our findings provide several mutually reinforcing lines of evidence supporting the conclusion that SNP c.1829C>T in porcine \(NR3C1\) is a gain-of-function mutation with a major effect on activity of the adrenal gland. This SNP is the first genetic variant in the pig for which a causal relationship to adrenal activity has been established and adds to a growing, but still limited, list of genetic variants known to be directly responsible for variation in complex traits reviewed in [21].

So far only two natural variants of GR, p.Asn363Ser and p.Asp401His (commonly designated N363S and D401H, respectively), that potentially increase its sensitivity to glucocorticoids for variation in complex traits reviewed in [21]. However, we cannot formally exclude existence of other contributing genetic variants acting in other temporally-spatial or functional contexts.

In the mouse, apart from the p.Met610Leu knock-in model, gene-targeting studies have been largely based on glucocorticoid p.Ala610Val variant.

Materials and Methods

Animals and phenotypes

Ethics statement. Animal care and tissue collection processes followed the guidelines of the German Law of Animal Protection, and the experimental protocol was approved by the Animal Care Committee of the Leibniz Institute for Farm Animal Biology (FBN, Dummerstorf, Germany).

Three commercial pig populations were used for sample and phenotype collection: a population of purebred German Landrace pigs (\(n = 834\)) and a population of German Large White (\(n = 274\)) pigs, both consisting only of barrows; and a Pietrain \(\times\) (German Large White \(\times\) German Landrace) cross consisting of barrows (\(n = 291\)) and females (\(n = 246\)). The average age at sampling was \(~170\) days.

Sample collection was performed in the experimental slaughter facility of the FBN between 0930 and 1130 h. Tissues used for RNA preparation were quickly removed, dissected, frozen in liquid nitrogen and stored at \(-80^\circ\)C. For dissection of brain areas of interest (hippocampus, hypothalamus, and amygdala) the stereotaxic atlas of the pig brain served as a reference [26]. For tissue sampling and weighing, left adrenal gland was taken.

A 50 ml sample of trunk blood was collected from each pig during the exsanguination in a plastic tube containing 1 ml of 0.5 M EDTA. After plasma preparation samples were stored at \(-80^\circ\)C until use. Plasma cortisol levels (total) were determined in duplicate using commercially available enzyme-linked immunoassay (DRG, Marburg, Germany) according to manufacturer’s protocol. The intra- and inter-assay coefficients of variation were lower than 7.0% and 9.8%, respectively. An overview of analyzed phenotypes is given in Table S1. Before association analysis outliers with cortisol levels \(\geq 200\) ng/ml were removed. Remaining data were approximately normally distributed.

Samples of Pi and Du animals used for assessment of allele frequency and of Hardy-Weinberg equilibrium were also collected in the experimental slaughter facility of the FBN.

RNA extraction and cDNA synthesis

Isolation of total RNA was performed using TRI reagent (Sigma, Taufkirchen, Germany). After DNase I treatment (Roche, Mannheim, Germany), RNA was cleaned using the NucleoSpin RNA II Kit (Macherey-Nagel, Düren, Germany). Quantity and purity of RNA were determined using the NanoDrop ND-1000 spectrophotometer (NanoDrop, Peqlab, Germany), and integrity was checked on 1% denaturing agarose gels.

First-strand cDNA was synthesized using SuperScript III MMLV reverse transcriptase (Invitrogen, Karlsruhe, Germany) in a reaction containing 1.5 µg RNA, a mixture of 500 ng random hexamers (Promega, Mannheim, Germany) and 500 ng of oligo (dT)11VN primer, according to manufacturer’s protocol.

Detection of polymorphisms

Genotyping using the PorcineSNP60 BeadChip (Illumina Inc., San Diego, CA, USA) was performed in accordance with manufacturers protocol for the SNP Infinium HD assay (http://www.illumin.com). In brief, 200 ng of DNA were used for genome-wide amplification and subsequent fragmentation. DNA was hybridized to the 62,163 locus-specific 50mers covalently linked to the beads distributed on the surface of the microarray. Single-base extension of oligos on the BeadChip, using captured DNA as a template, was performed, incorporating detectable
labels on the BeadChip. Signals of each wavelength were determined using an Illumina iScan that converted the images to intensity data. Intensity data for each SNP were normalized and assigned a cluster position and a resulting genotype with the GenomeStudio software (Illumina Inc.); a quality score was generated for each genotype. Samples with call rates <95% were removed. Markers were excluded if they had low minor-allele frequency (MAF) < 5%. The average call rate for all samples was 99.8% ± 0.2.

The sequence of porcine NR3C1 encoding GR-alpha was retrieved from NCBI from the sequence of BAC clone CH242-105G5 (Accession: CU928713). For resequencing, five overlapping fragments of ~600 bp covering the target region were designed and amplified in a standard PCR mix containing genomic DNA or cDNA as template, 0.2 and amplified in a standard PCR mix containing genomic DNA or cDNA as template, 0.2 and amplified in a standard PCR mix containing genomic DNA or cDNA as template, 0.2 and amplified in a standard PCR mix containing genomic DNA or cDNA as template, 0.2 μM of each primer (Tables S6 and S7), 50 μM each of dNTP, and 0.5 U SupraTherm Tag Polymerase in 1x supplied PCR buffer containing 1.5 mM MgCl2 (Genecraft, Ludwigshafen, Germany). The temperature profile consisted of 40 cycles of denaturation at 95°C for 15’s, annealing at appropriate Ta for 60 s, and extension at 72°C for 60 s. PCR products were sequenced using Big Dye Terminator Cycle sequencing kit V3.1 (Applied Biosystems, Darmstadt, Germany) and analyzed on ABI 3130 automated sequencer. Sequence polymorphisms were detected using the Multiple SeqDoc online tool [27] (http://research.imb.uq.edu.au/seqdoc/multi.html).

For genotyping SNP c.39A>C and c.55G>C, PCR-RFLP assays were established. Polymorphic regions were amplified in a standard PCR-reaction as described above using primers GRf1 and GRr1, and 10 μl of amplification products were digested overnight using 2.5 U of MboII and 10 U BglII restriction enzymes, respectively, according to manufacturer’s recommendations (Fermentas, St. Leon-Rot, Germany). Resulting RFLPs were analyzed on 2% ethidium bromide-stained agarose gels.

SNP c.1829C>T was genotyped using pyrosequencing. The polymorphic region was amplified using primers GRf7 (biotinylated) and GRr7 and as described above, except that DreamTag Polymerase (Fermentas) was used. The sequencing reaction was performed using primer GRseq1 and the Pyro Gold Reagent Kit (Biotage, Uppsala, Sweden) in the PSQ 96MA Pyrosequencing instrument (Biotage, Uppsala, Sweden) according to manufacturer’s instructions.

Genotyping of SNP c.*2122G>A was described previously [8]. Information on primers and amplicons designed to detect DNA polymorphisms are summarized in Tables S6 and S7.

Expression constructs and transactivation assay
To construct GR expression plasmids, a 2.38 kb GR-alpha fragment was amplified from cDNA of two individuals homozygous for the c.1829C>T SNP using primers GR_C-Flag_fw and GR_wt_rev (Table S6) and the proofreading PrimeSTAR HS DNA Polymerase (MoBiTec, Gottingen, Germany). The Ncol restriction sites, nested in the amplification primers, were used for insertion of the GR-alpha fragment into the pCMV-Tag 1 plasmid (Agilent Technologies, Waldbronn, Germany). The resulting expression plasmids, pCMV-GRA610 and pCMV-GRV610, were checked by sequencing for orientation and absence of mutations other than SNP c.1829C>T.

COS-7 cells were seeded in 96-well plates at 2.5 × 10^4 cells/well in DMEM supplemented with 10% FBS. The next day, cells were co-transfected with 200 ng pCMV-GRA610 or pCMV-GRV610, 100 ng pGL4.36, and 2 ng pRL-SV40 using Lipofectamine 2000. The pGL4.36 reporter (Promega) expresses firefly luciferase under the control of the glucocorticoid-inducible MMTV promoter. The constitutive expression of Renilla luciferase by the pRL-SV40 plasmid (Promega) was used to normalize for transfection efficiency.

Twenty-four hours after transfection, cells were treated with dexamethasone (Sigma) in different concentrations ranging from 0 to 100 nM, and, 24 h later, firefly and Renilla luciferase activities were measured under the use of the Dual-Glo Luciferase Assay System (Promega) in a DTX 880 Multimode Detector (Beckman Coulter, Krefeld, Germany). Four separate experiments were performed in triplicate. The effect of the p.Ala610Val substitution on the normalized expression of the firefly luciferase in the transactivation assay was tested by ANOVA using the MIXED procedure of the SAS V9.2 software package (SAS Institute, Cary, USA). The model included fixed effects of GR variant, dexamethasone concentration, and their interaction. Estimated least-squares means (LSM) were compared using t-test, and p-values were adjusted by simulation. Dose-response curve and EC50 were estimated using GraphPad Prism 5 (GraphPad Software Inc, San Diego, CA).

Analysis of NR3C1 transcription
To analyze allelic expression imbalance, the ratio of alleles of SNP c.1829C>T in cDNA and corresponding genomic DNA (gDNA) samples were measured using pyrosequencing. Pyrosequencing was performed as described above in two separate experiments, in duplicate. Ratios obtained from cDNA were normalized using ratios obtained from gDNA within each experiment. After log2 transformation, allelic expression imbalance was tested by comparing ratios obtained from cDNA with ratios obtained from gDNA using a two tailed t-test.

For the analysis of splicing of the GR-alpha isoform, six overlapping amplicons covering the whole open reading frame (ORF) were designed. The sequence corresponding to human exon 1C was retrieved from the sequence of BAC clone CH242-105G5. Reverse-transcription PCR was performed using standard conditions described in the previous paragraph, and products were visualized on 2% agarose gels. Identity of the amplicons was confirmed by direct sequencing. Information on primers and amplicons used for the analysis of the integrity of GR-alpha ORF are summarized in Tables S6 and S7.

Population genetic and association analysis
Genome-wide and single-marker association analyses were performed using mixed linear models implemented in the JMP Genomics 5 and SAS V9.2 software packages (SAS Institute), respectively. The model included fixed effect of the SNP genotype and, to account for relatedness, the random effect of sire. For adrenal weight the model included body weight as a covariate. For both traits gender was included as a fixed effect for the analysis of the F1 population. LSM for NR3C1 genotypes were compared by t-test, and p-values were adjusted by Tukey-Kramer correction. Allele substitution effects were estimated by linear regression on the number of copies of the derived allele. FDR (q-value, [28]) was computed using JMP Genomics 5.

Haplotype inference was performed using the expectation-maximization algorithm implemented in the HAPLOTYPE procedure of SAS V9.2. Haplotype trend regression was performed by including one variable for each haplotype indicating the number of copies of the haplotype in question in the models described above.

Hardy-Weinberg equilibrium was analyzed using the ALLELE procedure of SAS V9.2. Linkage disequilibrium between SNPs and
haplotype blocks were determined using Haplovew V4.2 [29]. Haplotype blocks were defined using the four gametes rule criteria.

**Supporting Information**

**Figure S1 Alignment of the glucocorticoid receptor amino acid sequence among different vertebrates.** Two regions of the glucocorticoid receptor amino acid sequence are shown where DNA polymorphisms in the porcine NR3C1 introduce amino acid substitutions. The position of the three variable residues found in pigs is indicated by arrows, and the homologous residues of other species by gray bars. Evolutionarily conserved residues are indicated by dots. Residues of helix 5 of the ligand binding domain are underlined.

**Figure S2 Linkage disequilibrium analysis of the QTL region on chromosome 2.** The values in the boxes show linkage disequilibrium between SNPs (\(r^2\)) and the box color reflects conserved residues. Residues of helix 5 of the glucocorticoid receptor amino acid sequence among different vertebrates.

**Table S1 Descriptive statistics of the analyzed traits.**

**Table S2 SNPs with genome-wide significant evidence for association with plasma cortisol level.**

**Table S3 SNPs with genome-wide significant evidence for association with adrenal weight.**

**Table S4 Frequencies of the three identified missense SNPs in porcine NR3C1 in commercial breeds.**

**Table S5 Frequencies of inferred haplotypes of porcine NR3C1 in three different commercial populations.**

**Table S6 Primer information.**

**Table S7 Information on amplicons used for the analysis of splicing pattern, resequencing, and genotyping of NR3C1.**

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**Author Contributions**

Conceived and designed the experiments: EM. Performed the experiments: HR SP SF. Contributed reagents/materials/analysis tools: EM HR SP SF. Wrote the paper: EM.

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