Expression and regulation of glucocorticoid-induced leucine zipper in the developing anterior pituitary gland

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Abstract

The expression profile of glucocorticoid-induced leucine zipper (GILZ) in the anterior pituitary during the second half of embryonic development in the chick is consistent with in vivo regulation by circulating corticosteroids. However, nothing else has been reported about the presence of GILZ in the neuroendocrine system. We sought to characterize expression and regulation of GILZ in the chicken embryonic pituitary gland and determine the effect of GILZ overexpression on anterior pituitary hormone levels. Pituitary GILZ mRNA levels increased during embryogenesis to a maximum on the day of hatch, and decreased through the first week after hatch. GILZ expression was rapidly upregulated by corticosterone in embryonic pituitary cells. To determine whether GILZ regulates hormone gene expression in the developing anterior pituitary, we overexpressed GILZ in embryonic pituitary cells and measured mRNA for the major pituitary hormones. Exogenous GILZ increased prolactin mRNA above basal levels, but not as high as that in corticosterone-treated cells, indicating that GILZ may play a small role in lactotroph differentiation. The largest effect we observed was a twofold increase in FSH β subunit in cells transfected with GILZ but not treated with corticosterone, suggesting that GILZ may positively regulate gonadotroph development in a manner not involving glucocorticoids. In conclusion, this is the first report to characterize avian GILZ and examine its regulation in the developing neuroendocrine system. We have shown that GILZ is upregulated by glucocorticoids in the embryonic pituitary gland and may regulate expression of several pituitary hormones.

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Introduction

In a study profiling gene expression during cellular differentiation of the chicken embryonic pituitary gland, our laboratory identified glucocorticoid-induced leucine zipper (GILZ) as an anterior pituitary transcript that increased between embryonic day (e) 12 and e17 (Ellestad et al. 2006). This expression pattern is consistent with in vivo regulation of GILZ in the anterior pituitary by glucocorticoids, as circulating corticosteroids increase between e12 and e17 in the developing chicken embryo (Scott et al. 1981, Jenkins et al. 2007, Porter et al. 2007). Corticosterone (CORT), the predominant glucocorticoid in rodents and birds, increases 100-fold during chicken embryonic development, from subnanomolar concentrations on e10 (0.3 nM) to e17 (40 nM), when the levels are highest (Jenkins et al. 2007, Porter et al. 2007). Corticosteroids are thought to play a role in the initiation of hormone expression during pituitary cell ontogeny in both mammals and birds. In both rats and chickens, CORT stimulates GH and prolactin (PRL) expression during somatotroph and lactotroph differentiation respectively (Hemming et al. 1984, Nogami et al. 1995, Morpurgo et al. 1997, Dean & Porter 1999, Dean et al. 1999, Bossis & Porter 2000, Fu & Porter 2004). Glucocorticoids also repress pro-opiomelanocortin (POMC) expression during negative feedback on adrenocorticotropic hormone (ACTH) release from pituitary corticotrophs (Drouin et al. 1989), although when this feedback is established during development is not clear. In the pituitary, GILZ may be mediating effects of glucocorticoids as well as exhibiting independent effects in this tissue.

GILZ was first identified as a glucocorticoid-induced transcript in mouse thymocytes, and its expression was initially thought to be restricted to lymphoid tissues (D’Adamio et al. 1997). Subsequently, GILZ has been shown to be widely expressed and has been detected in numerous tissues and cell types (Cannarile et al. 2001, Berrebi et al. 2003, Kolbus et al. 2003, Shi et al. 2003, Zhao et al. 2006). GILZ is thought to be a transcription factor and has been shown experimentally to bind DNA (Shi et al. 2003) as well as interact with other leucine zipper transcription factors such as c-Fos and c-Jun (Mittelstadt &
GILZ has been implicated in apoptosis (Ayroldi et al. 2001, 2002, 2007). GILZ has been involved in immunosuppression (Berrebi et al. 2003) and cell proliferation (Ayroldi et al. 2007), transport (Soundararajan et al. 2005), parturition (Zhao et al. 2006), and cell fate choice (Shi et al. 2003, Levine et al. 2007, Zhang et al. 2007).

To our knowledge, no studies investigating the presence or regulation of GILZ in the neuroendocrine system, a well-established target of glucocorticoid action, have been conducted. Further, GILZ has not been characterized in an avian species, and it is unknown whether avian GILZ is regulated in a manner similar to mammalian GILZ. The chicken has several advantages over traditional mammalian models for studying pituitary gland development, including the ability to precisely control the beginning of embryonic development and therefore perform experiments with hundreds of pituitary glands at a given age. Pituitary development is highly conserved among vertebrates; therefore, understanding regulation and potential action of GILZ in the developing pituitary gland of the chick embryo will provide information about pituitary development that is applicable to mammalian species as well. The objectives of this study were to determine the sequence and tissue distribution of chicken GILZ mRNA, to determine GILZ ontogeny and regulation by glucocorticoids in the embryonic pituitary gland, and to determine whether GILZ plays a role in regulating expression of the anterior pituitary hormones.

Materials and methods

Animals and pituitary collection

Broiler strain chicken embryos used for these experiments were purchased from Allen’s Hatchery (Seaford, DE, USA). Eggs were placed in a 37-5 °C humidified incubator on embryonic day (e) 0 and removed on the appropriate day of the 21-day incubation period for each experiment. For cell culture experiments, pituitary glands were isolated with a dissecting microscope from embryos on e11 and dissociated into individual cells using trypsin digestion and mechanical agitation (Porter et al. 1995). Anterior pituitaries from ~60 embryos were isolated and pooled for each of the four replicate trials of a given experiment (n = 4). For the GILZ ontogeny experiment, whole anterior pituitary glands were collected from e10, e12, e14, e16, e18, e20, post-hatch day (d) 1, d3, d5, and d7 birds using a dissecting microscope. To yield adequate mRNA from each developmental age, pituitaries from 3 e10, 2 e12, 2 e14, and 2 e16 embryos were pooled for each replicate sample; one pituitary per replicate was used for the older ages. Four replicate samples were collected for each age (n = 4). Pituitaries were immediately frozen in liquid N2 and stored at −80 °C until RNA extraction. All procedures were approved by the Institutional Animal Care and Use Committee at the University of Maryland.

Sequencing of chicken GILZ cDNA and comparison with other species

The putative chicken GILZ clone was part of a neuroendocrine cDNA library used to construct a custom cDNA microarray and was initially identified through random sequencing of the library in combination with BLAST comparisons (Cogburn et al. 2004, Carre et al. 2006, Ellestad et al. 2006). The plasmid containing chicken GILZ (pgp1n.pk005.a10, GenBank accession numbers BI39162) was purified according to a standard protocol (Sambrook et al. 1989), and the clone was sequenced in its entirety in both directions. In order to fully sequence the insert, three sequencing reactions were performed on each of the sense and antisense strands. Oligonucleotide primers (Sigma Genosys) used for sequencing were Sp6microarray+, Gilzsense1, and Gilzsense2 for the forward reactions, and T7microarray+, Gilzantisense1, and Gilzantisense2 for the reverse reactions (Table 1). Sequencing reactions were performed by the University of Maryland’s Center for Biosystems Research DNA Sequencing Facility with AmpliTaq-FS DNA polymerase and Big Dye terminators with dITP in an Applied Biosystems DNA Sequencer (Model 3100; Foster City, CA, USA). Overlapping sequences were assembled into the full-length chicken GILZ cDNA using Vector NTI 9.0 (Invitrogen), and the same software was used to make comparisons between chicken GILZ nucleotide and predicted amino acid sequences with those for human (accession numbers NM_004089 and NP_004080), mouse (accession numbers NM_010286 and NP_034416), dog (accession numbers XM_549177 and XP_549177), zebrafish (accession numbers XM_703382 and XP_708474), and Xenopus (accession numbers BC043841 and AAH43841) obtained from GenBank. Human, mouse, and Xenopus sequences are clones mRNAs, and dog and zebrafish sequences are predicted transcripts based on computational analysis. Biology Workbench (http://workbench.sdsc.edu/) was used to create the alignment (CLUSTALW tool), color-coding (BOXSHADE tool), and unrooted phylogenetic tree diagrams shown in Fig. 1.

Cell culture

Hormones and other chemicals were obtained from Sigma–Aldrich, and cell culture reagents were obtained from Invitrogen. Cells were maintained in a 37-5 °C, 5%
CO₂ atmosphere. Dispersed pituitary cells were cultured in serum-free medium (DMEM/F12) supplemented with 0.1% BSA, 5 μg/ml human insulin, 5 μg/ml human transferrin, 100 U/ml penicillin G, and 100 μg/ml streptomycin sulfate.

For the GILZ mRNA induction experiments, e11 pituitary cells (1 × 10⁶/well) cultured in 12-well plates (Corning Incorporated, Corning, NY, USA) for 24 h were treated for the final 1.5, 3, 6, or 12 h with CORT (1 nM) in the absence and presence of cycloheximide (CHX; 10 μg/ml). In wells receiving CHX plus CORT, cells were pretreated for 1.5 h with CHX prior to addition of CORT. Untreated cells and cells receiving CHX alone for 1.5 h served as controls. In all experiments, cells were harvested by trypsinization and stored at −80 °C until total RNA was extracted.

For GILZ overexpression experiments, dispersed e11 anterior pituitary cells (3 × 10⁶/well) were co-transfected with 2 μg golgi-targeted green fluorescent protein (GFP) expression vector (Pecot & Malhotra 2004) and either 2 μg empty expression vector (pCMV-Sport6.1; Invitrogen) or 2 μg GILZ-expression vector (pgp1n.pk005.a10) for 6 h using Lipofectamine 2000 (Invitrogen) and Opti-MEM I medium. Cells were transfected in suspension for 2 h and then plated into 6-well culture plates for the remaining 4 h. Cells were harvested by trypsinization and stored at −80 °C until total RNA was extracted.

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Flow cytometry

Successfully transfected cells in GILZ overexpression studies were identified by flow cytometric detection of GFP (fluorescence detection 530/30 nm). GFP-positive cells were sorted using a high performance flow cytometer and cell sorter (MoFlo; DakoCytomation, Carpinteria, CA, USA) equipped with an argon ion laser tuned to 488 nm (Innova 300, Coherent, Palo Alto, CA, USA). To enhance recovery of sorted cells, the enrichment sort mode was used. Sorting of the GFP-positive population (19.3% of cells; $n=4$ replicate experiments) resulted in collection of $\sim 70,000$ GFP-positive cells per group. Additionally, GFP-negative cells were collected, and both were pelleted at 1000 g for 10 min and snap-frozen in liquid nitrogen. Cells were stored at $-80^\circ$C until total RNA was extracted.

Northern blotting

Total RNA samples isolated from indicated tissues using TRIZOL (Invitrogen) were quantified by measuring the absorbance at 260 nm with u.v. spectrophotometer. RNA (10 μg) was separated by formaldehyde agarose gel (1%) electrophoresis, transferred to a Zeta-Probe Blotting Membrane (Bio-Rad), and cross-linked to the membrane with a u.v. Stratalinker (Stratagene, La Jolla, CA, USA). The membrane was probed with a $^{32}$P-labeled full-length chicken GILZ cDNA probe generated by PCR using the pgg1n.pk005.a10 plasmid with Sp6microarray+/T7microarray+ primers (Sigma-Genosys, Table 1) and the following conditions: initial denaturation of 94°C for 3 min; 35 cycles of 94°C for 45 s, 52°C for 45 s, and 72°C for 3 min; and a final extension of 72°C for 7 min. The labeled PCR product was purified with a spin column-30 (Sigma–Aldrich) and hybridized to the membrane overnight at 65°C using PerfectHyb Plus 1X Hybridization Buffer (Sigma–Aldrich). The membrane was washed twice with 2× SSC and 0-1% SDS at room temperature for 10 min and thrice with 0-2× SSC and 0-1% SDS at 65°C for 10 min. Bands were visualized using a phosphorimager (Molecular Dynamics PhosphorImager; Amersham Biosciences). The membrane was then stripped with 0-02× SSC and 0-1% SDS at 90°C and re-hybridized with a $^{32}$P-labeled chicken β-actin (ACTB) probe described previously (Fu & Porter 2004).

Reverse transcription-PCR (RT-PCR) and 3′-rapid amplification of cDNA ends (RACE)

Total RNA (2 μg) was reversed transcribed (20 μl reactions) using SuperScript III Reverse Transcriptase (Invitrogen) and either a GILZ-specific reverse primer (GILZantisense1; Table 1; Sigma Genosys) for standard RT-PCR or the Anchored-dT-one-V primer (Table 1; Sigma Genosys) for 3′-RACE. As a negative control for genomic DNA contamination, a reaction was conducted with each sample in which all components were added except reverse transcriptase (RT$^-$). For standard RT-PCR, the RT reactions were diluted tenfold prior to PCR analysis using GILZsense1 and GILZantisense2 primers (Table 1; Sigma Genosys). PCR cycling parameters were as follows: 94°C for 3 min; 35 cycles of 94°C for 45 s, 59-5°C for 45 s, and 72°C for 3 min; and a final extension at 72°C for 7 min. To perform 3′-RACE, two consecutive PCRs were conducted. In the first, undiluted cDNA was amplified using GILZsense1 and Anchored-dT-one-V primers (Sigma-Genosys). This reaction was then purified with a spin column-100 (Sigma–Aldrich) and diluted 1:1000 for subsequent amplification using GILZwd2 and Anchored-dT-one-V primers. PCRs were conducted as for RT-PCR, except the annealing temperature was 48°C. Products were visualized using agarose gel (1%) electrophoresis and ethidium bromide staining and sequenced to confirm they were GILZ.

Quantitative real-time RT-PCR (qRT-PCR)

Total RNA was extracted from caudal and cephalic pituitary lobes from late embryonic chicks or cultured cells with the RNeasy Mini Kit (Qiagen) and quantified using the Quant iT RiboGreen RNA Quantitation Reagent (Invitrogen). RNA (20 ng) isolated from flow cytometric-sorted GFP-positive cells was amplified with the procedure (Phillips & Eberwine 1996) and modifications (Porter & Ellestad 2005, Ellestad et al. 2006) described elsewhere. Two-step qRT-PCR was used to quantify mRNA levels for ACTB, GH, PRL, POMC, α-glycoprotein subunit (αGSU), TSH β subunit (TSHβ), LH β subunit (LHβ), FSHβ, and GILZ, as appropriate. RT reactions (20 μl) were carried out using SuperScript III (Invitrogen) with random primers (Invitrogen) for amplified RNA (200 ng), or the Anchored-dT-one-V primer (Sigma Genosys) with T-RNA (500 ng) for all others. As a negative control for genomic DNA contamination, a pool of all the RNA from a given experiment was made and the reaction conducted as the others except reverse transcriptase was not added. To control for potential GILZ plasmid carryover through the RNA extraction, a pool of RNA from cells transfected with the GILZ-expression vector was also made for the GILZ overexpression studies, and the reaction conducted as the others except reverse transcriptase was not added. All RT reactions were diluted to 100 μl prior to PCR analysis. Primers (Sigma Genosys) used in the PCRs were designed with Primer Express Software (Applied Biosystems) and are listed in Table 1. PCRs (20 μl) contained 2 μl diluted cDNA, 400 nM each primer, PCR buffer, 0-12 U/μl Taq Polymerase, 200 nM dNTPs, 40 nM fluorescein (Invitrogen), and SYBR Green I Nucleic Acid Gel Stain.
Developmental regulation of pituitary GILZ  ·  L E ELLESTAD and others

We identified GILZ as an anterior pituitary transcript that is upregulated during the second half of embryogenesis in the chick using a neuroendocrine system cDNA microarray (Ellestad et al 2006). As an avian homolog for GILZ has never been reported, we sequenced the cDNA insert of the GILZ clone used to print the microarray (GenBank accession number BI391625) in its entirety in both directions. This sequence was then compared with available GILZ sequences from other species (Table 2 and Fig. 1). Chicken GILZ mRNA (GenBank accession number DQ917420) was determined to be 1118 bp long and encode a predicted protein consisting of 139 amino acids. Along the entire length of chicken GILZ cDNA, there is 55–60% identity with human, mouse, dog, zebrafish, and Xenopus GILZ; however, comparisons among the coding regions indicate a much higher identity of 75–80%. Similarly, the protein homology between chicken GILZ and other vertebrates is high (75–80%), especially when positive amino acid substitutions are considered (80–85%). Importantly, chicken GILZ contains the leucine zipper motif characterized by four conserved leucine residues and one conserved asparagine (Fig. 1A). Phylogenetically, chicken GILZ lies between mammalian and non-mammalian vertebrate species (Fig. 1B). A notable difference is that chicken GILZ contains a substantial truncation of ~500 bp in the 3′-untranslated region compared with that in other vertebrates. We verified through northern blotting that the size of the major chicken GILZ transcript in all tissues examined is 1.1 kb (Fig. 2A). The apparent discrepancy in transcript size in various tissues was due to differences in migration of different RNA samples through the gel and does not represent alternative splicing in the different tissues, as the band detected for ACTB was also shifted in the same direction in those samples. Most notably, the transcripts for GILZ and ACTB detected in the heart, liver, and brain appear larger than those in the other tissues, but are not. The difference in migration of RNA extracted from these tissues may be due to higher levels of impurities in the samples, possibly glycogen, resulting from higher levels of this compound in the starting material. In order to further confirm the presence of this truncation and the length of the transcript, 3′-RACE was performed with two sequential GILZ sense primers and an oligo-dT primer (Table 1). As is evident from the picture, the first reaction yielded a smear surrounding the predicted size (789 bp) and the second reaction yielded a clean band of appropriate size (407 bp), which was sequenced and confirmed to be GILZ. This demonstrated that chicken GILZ is indeed shorter in the 3′-untranslated region than mammalian GILZ (Fig. 2D). Further, there is a polyadenylation signal (AAUAAA) just upstream of the 3′-end of our sequence. These results indicate that the major form of chicken GILZ in the tissues examined contains this 3′-truncation, and may imply

Data analysis

The qRT-PCR data, expressed as fold induction relative to basal or relative to pituitary lobe or age with the highest levels for a given gene, were log2-transformed prior to statistical analysis. All data were analyzed by ANOVA using the MIXED models procedure of SAS (SAS Institute, Cary, NC, USA), and differences between groups were determined using the test of least significant difference (PDIFF; SAS).

Results

Chicken GILZ is highly homologous to GILZ in other vertebrate species

We identified GILZ as an anterior pituitary transcript that is upregulated during the second half of

(Invitrogen) diluted 1:10 000 and were carried out in the MyiQ Single-Color Real-Time PCR Detection System (Bio-Rad). A two-step PCR cycle was used: initial denaturation at 95 °C for 5 min followed by 40 cycles of 95 °C for 15 s and 60 °C for 60 s. Dissociation curve analysis and gel electrophoresis were conducted to ensure that a single PCR product of appropriate size was amplified in each reaction and absent from the no RT controls. Except for the developmental ontogeny experiments, in which data were analyzed as previously described (Ellestad et al 2006), the amount of mRNA for each target gene was normalized to the previously described (Ellestad et al 2006). The amount of mRNA for each target gene was normalized to the previously described (Ellestad et al 2006). The amount of mRNA for each target gene was normalized to the previously described (Ellestad et al 2006). The amount of mRNA for each target gene was normalized to the previously described (Ellestad et al 2006). The amount of mRNA for each target gene was normalized to the previously described (Ellestad et al 2006).

The Ct value is the threshold cycle when the amount of amplified product reaches a fixed threshold for fluorescence due to binding of SYBR green to the double-stranded PCR product. Data were then transformed using the following equation mRNA level = 2^(-ΔCt).

For the caudal and cephalic pituitary lobe expression studies, the transformed value for each sample was divided by the mean of the transformed value for the lobe with the highest expression level for a particular gene. Data for each gene are expressed relative to the lobe with the highest expression level. When GILZ was measured during pituitary development, the transformed value for each value was divided by the mean of the transformed value for d1, such that data are expressed relative to d1, when GILZ expression was found to be highest. For cell culture experiments, the transformed value for each sample was divided by the mean of the transformed value for the basal wells in each experiment such that data are presented as a fold induction in mRNA when compared with basal levels (receiving no treatment and transfected with empty vector, as appropriate).

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that there are differences in the regulation of GILZ expression at the post-transcriptional level between mammals and birds.

**Chicken GILZ is ubiquitously expressed**

Both northern blot and RT-PCR analyses were used to determine the tissue distribution of GILZ in the chicken. In both methods, GILZ was detected in all tissues examined (Fig. 2A and B), namely pituitary, hypothalamus, brain, heart, lung, adrenal, spleen, thymus, bursa of Fabricius, liver, and kidney. Given that GILZ is present in the pituitary, we sought to further define localization within this tissue because the avian anterior pituitary consists of two anatomically distinct caudal and cephalic lobes. In the chicken, four of the major pituitary cell types are unevenly distributed among these two lobes. GH-producing somatotrophs reside primarily in the caudal lobe, while PRL-producing lactotrophs, TSH-producing thyrotrophs, and ACTH-producing corticotrophs are mainly localized within the cephalic lobe (Kansaku et al. 1995, Lopez et al. 1995, Gerets et al. 2000, Nakamura et al. 2004, Muchow et al. 2005). Gonadotrophs, which produce LH and FSH, are initially present in the caudal lobe but spread throughout the entire gland as the pituitary develops (Maseki et al. 2004). We measured mRNA levels of GH, PRL, TSHβ, and GILZ in caudal
and cephalic portions of the pituitary glands from chicks just prior to hatch. It should be noted that the higher variation in GILZ mRNA relative to that for the pituitary hormones measured is not due to normalization and appears to reflect true biological variability in the samples collected. Regardless, results from qRT-PCR analysis indicate that GILZ mRNA is equivalently expressed in both lobes in a manner similar to LH and FSH, and confirm the caudal distribution of GH mRNA and cephalic localization of PRL and TSHβ mRNA (P<0.05; n=3; Fig. 2C).

**GILZ is developmentally regulated in the pituitary gland**

The mature anterior pituitary gland consists of five major cell types that emerge during development in a temporally specific manner. Initiation of hormone transcription in these cells occurs during the second half of embryogenesis and continues during early neonatal life in both mammals and birds (Japon et al. 1994, Sasaki et al. 2003, Ellestad et al. 2006). Pituitary αGSU, POMC, and TSHβ mRNAs steadily increase during the second half of embryogenesis, expression of GH, PRL, and FSHβ mRNA increases most substantially during the last third of embryogenesis, and LHβ mRNA is not abundant until after hatch (Ellestad et al. 2006, Ellestad and Porter, unpublished). We characterized pituitary GILZ mRNA during this time in order to determine whether GILZ is regulated in a manner consistent with a potential role in pituitary development. GILZ mRNA was measured during the latter half of chicken embryonic development and the first week post-hatch, from e10 through d7. Expression increased between e12 and e14 and continued to dramatically increase during the final third of embryonic development to a maximum immediately post-hatch, and then decreased through d7 (P<0.05; n=4; Fig. 2E). The expression profile measured in this experiment indicates that GILZ may play a developmental role in this tissue, and it is most consistent with a putative role in the development of gonadotrophs, somatotrophs, and lactotrophs based on timing of their appearance.

**GILZ is upregulated by glucocorticoids in chicken embryonic pituitary cells**

The expression profile of GILZ in the developing chicken pituitary gland is consistent with upregulation by glucocorticoids, because circulating CORT increases in a similar manner to GILZ during embryogenesis (Jenkins et al. 2007, Porter et al. 2007). To determine whether glucocorticoids can induce GILZ in embryonic pituitary cells from birds at an age when circulating CORT is comparably low, we analyzed GILZ mRNA in e11 pituitary cells treated with CORT (1 nM) by qRT-PCR. This dose of CORT was chosen because it reflects circulating levels of CORT on e14, the embryonic age at which GILZ expression first begins to increase in the chicken (Jenkins et al. 2007, Porter et al. 2007). CORT treatment induced GILZ mRNA after 3 and 6 h, and there was a further increase after 12 h (P<0.05; n=4; Fig. 3A). In an attempt to evaluate whether GILZ is a direct transcriptional target of CORT, we cultured e11 anterior pituitary cells for 1.5, 3, and 6 h with CORT in the presence of CHX, an inhibitor of protein synthesis (Fig. 3B). Similar to the results shown in Fig. 3A, GILZ was upregulated almost threefold after 1.5 h CORT treatment, and this upregulation was maintained through 6 h (P<0.05; n=4). However, because treatment with CHX alone for 1.5 h stimulated GILZ mRNA in these cells to a level similar to that with CORT alone for 1.5 h (P<0.05; n=4; Fig. 3C, compare white and black bars at 0 h and black bar at 0 h with white bar at 1.5 h), it...
GILZ can upregulate POMC mRNA in embryonic anterior pituitary cells. These results also demonstrate that the CMV promoter was able to substantially drive expression of GILZ in these cells (P<0.05). Overexpression of GILZ had no effect on GH mRNA in the absence or presence of CORT, but cells transfected with GILZ in the absence of CORT had intermediate levels of PRL mRNA when compared with basal cells transfected with empty vector and cells treated with CORT (P<0.05). Similarly, cells transfected with GILZ in the absence of CORT had significantly, although modestly, higher POMC mRNA than basal cells transfected with empty vector (P<0.05), but this effect was not apparent in the presence of CORT. This indicates that CORT may be inducing factors that prevent the slight upregulation of POMC by GILZ. Together, these results imply that GILZ may play a minor role in the induction of PRL by glucocorticoids during pituitary development and that GILZ can upregulate POMC mRNA in embryonic pituitary cells in the absence of glucocorticoids.
We also measured mRNA levels for αGSU, TSHβ, LHβ, and FSHβ, which have not been shown to be regulated by corticosteroids during pituitary development, to determine whether GILZ might regulate glycoprotein hormone expression (Fig. 4). Neither GILZ overexpression nor CORT affected levels of αGSU or TSHβ in e11 cells. Expression of GILZ increased FSHβ and LHβ mRNA under basal and CORT-treated conditions respectively (P<0.05), although CORT itself had no effect. These results indicate that GILZ may positively regulate expression of pituitary gonadotropins, and they suggest that GILZ may have effects on gene expression that are independent from its regulation by glucocorticoids.

Discussion

This is the first report to characterize GILZ in an avian species, and the first to investigate expression, regulation, and a potential function for GILZ in neuroendocrine development. We sequenced a chicken GILZ cDNA clone, and it contained an insert that was only 1118 bp long, ~650 bp shorter than GILZ in other vertebrates. The insert contained the entire predicted open reading frame for GILZ, and the majority of the truncation (500 bp) lies within the 3′-untranslated region and the remainder within the 5′-untranslated region. The estimated transcript size for chicken GILZ was confirmed by northern blotting in all tissues and the 3′-end of the transcript was verified using 3′-RACE, indicating that this is the predominant form of GILZ in chickens. This truncation may imply an alternative regulation of GILZ mRNA at the post-transcriptional level in birds, as elements within the 3′-untranslated region are thought to modulate mRNA stability (Hughes 2006). Despite this truncation, the coding region of GILZ mRNA and the predicted protein are highly homologous with those from other species, including the conserved leucine zipper region.
facilitating protein–protein interactions (homo- or heterodimerization) that regulate DNA-binding characteristics of this family of transcription factors.

In the chicken, GILZ mRNA was detected in every tissue we examined, including the pituitary. In mammals, GILZ has been shown to be expressed in a variety of tissues and cell types, including heart, brain, kidney, liver, lung, pancreas, spleen, muscle, uterus, mesenchymal cells, pre-adipocytes, and white and red blood cells (Cannarile et al. 2001, Berrebi et al. 2003, Kolbus et al. 2003, Shi et al. 2003, Zhao et al. 2006). However, this is the first report indicating expression of GILZ in the neuroendocrine system of any species. Within the pituitary, GILZ was detected in both caudal and cephalic lobes, indicating that GILZ expression is not restricted to any particular cell type or is present in cell types that are distributed throughout the avian gland, such as gonadotrophs (Maseki et al. 2004).

In the developing chicken embryo, circulating glucocorticoids increase from subnanomolar concentrations on e12 to 1–5 nM on e14, and increase further to 40–50 nM on e17 before decreasing slightly on e20 (Scott et al. 1981, Jenkins et al. 2007, Porter et al. 2007). After hatch, circulating glucocorticoids remain in the 10–30 nM range during the first week of neonatal life, with serum levels peaking on d1, and declining through d4 before rising again on d5 (Latour et al. 1995). The developmental profile of pituitary GILZ mRNA during the second half of embryonic and early post-hatch development measured in the current study reflect these levels of serum CORT and are consistent with an in vivo regulation of GILZ by circulating glucocorticoids. We found that GILZ expression increased slightly between e12 and e14, then dramatically increased through d1, after which mRNA levels decreased through the remainder of the experiment. These data are consistent with our previous report, where GILZ mRNA was shown to increase between e12 and e17 by both microarray and qRT-PCR analysis (Ellestad et al. 2006). This potential upregulation of GILZ by CORT in the chicken pituitary was confirmed by in vitro experiments, in which cultured e11 pituitary cells treated with CORT at a level similar to that found in the e14 embryo had significantly higher levels of GILZ mRNA than those left untreated.

In cultured chick embryonic pituitary cells, GILZ mRNA was upregulated by CORT within 1–5 h. The magnitude of induction observed in these cells, between two- and fourfold, is consistent with induction of the mammalian GILZ transcript by the synthetic glucocorticoid dexamethasone in non-lymphoid tissues (Kolbus et al. 2003, Chen et al. 2006). In lymphoid cells, however, glucocorticoids appear to induce GILZ gene expression more strongly (D’Adamo et al. 1997, Cannarile et al. 2001, Asselin-Labat et al. 2005). One mode of action of glucocorticoids is to bind the intracellular glucocorticoid receptor (GR), which then acts as a ligand-activated transcription factor that binds to glucocorticoid response elements (GREs) in the regulatory regions of direct target genes. In the developing chick anterior pituitary, GR mRNA and protein are present by e8 and both increase in expression during embryonic development (Bossis et al. 2004, Ellestad et al. 2006, Kwok et al. 2007). GR protein is found in almost all chicken anterior pituitary cells (Bossis et al. 2004) indicating that GILZ may be a direct transcriptional target of GR in these cells. In mammalian cells, GILZ mRNA is increased in as little as 15 min after exposure to dexamethasone and is induced even in the presence of a protein synthesis inhibitor (Asselin-Labat et al. 2005, Chen et al. 2006) further supporting that GILZ is likely a direct glucocorticoid transcriptional target. This is consistent with the presence of multiple GREs in the human GILZ promoter (Asselin-Labat et al. 2005). However, because it was induced by CHX in chicken embryonic pituitary cells, it is impossible to determine from the current experiments if GILZ is indeed a direct transcriptional target in the developing pituitary gland. Further, the GILZ gene falls within an unassembled region of the chicken genome, so in silico promoter analysis for GREs in this species has not yet been possible. Nonetheless, our present findings indicate that GILZ is rapidly upregulated by CORT in the developing pituitary gland.

During development, glucocorticoids play a role in functional maturation of several tissues, including lung (Cole et al. 2004), small intestine (Thomson & Keelan 1986), and pituitary (Nogami et al. 1995, Dean & Porter 1999, Fu & Porter 2004). In mammals and birds, glucocorticoids stimulate GH and PRL expression during somatotroph and lactotroph differentiation respectively, in the course of anterior pituitary development (Hemming et al. 1984, Nogami et al. 1995, Morpurgo et al. 1997, Dean & Porter 1999, Dean et al. 1999, Bossis & Porter 2003, Fu & Porter 2004). The timing of this stimulation and, in the case of GH, requirement for ongoing protein synthesis, implicates an indirect mechanism requiring induction of an intermediary protein (Nogami et al. 1997, Porter et al. 2001, Bossis & Porter 2003). Glucocorticoids are also known to repress POMC expression during negative feedback on ACTH release from corticotrophs (Drouin et al. 1989), although the time during hypothalamic–pituitary–adrenal axis ontogeny when this feedback is first established is unknown. Pituitary GILZ shows developmental regulation consistent with the increase in circulating glucocorticoids that occurs during the second half of embryonic development, and GILZ is upregulated by CORT in embryonic pituitary cells. These observations led us to hypothesize that GILZ may play a role in mediating levels of the pituitary hormones, expression of which is initiated by corticosteroids during
development. In addition, GILZ is present in the embryonic pituitary gland at the earliest age we investigated, e10, even when circulating CORT levels are low, and GILZ may play a role in initiating expression of hormones that are not induced by glucocorticoids.

GILZ overexpression in embryonic pituitary cells did not affect GH mRNA, either in the presence or absence of CORT, indicating it is not involved in somatotroph development. On the other hand, GILZ overexpression did modestly increase PRL to 165% and POMC to 150% of mRNA levels in the absence of CORT. In the case of PRL, GILZ overexpression increased PRL mRNA to a level intermediate between basal cells and those receiving CORT. This indicates that, while GILZ alone is capable of positively regulating PRL expression in embryonic pituitary cells, it likely plays a minor role in the induction of PRL by CORT and the regulation of other factors by glucocorticoids is also required. The increase of POMC expression in the presence of exogenous GILZ, although slight, may contribute to the lack of a negative effect of CORT on POMC mRNA in these experiments. At this embryonic age, GILZ that is induced by CORT may positively regulate POMC mRNA in a manner that prevents its downregulation by glucocorticoids. Overexpression of GILZ also upregulated levels of LHβ and FSHβ. The twofold increase in FSHβ mRNA in the absence of CORT was the greatest effect we observed in these experiments, although this effect was diminished in the presence of CORT. While cortisol does repress the reproductive axis in mammals (Breen & Karsch 2004, Gore et al. 2006), these genes have not previously been shown to be developmentally regulated by glucocorticoids. Consistent with this, CORT did not affect levels of gonadotropin β-subunit mRNA in chicken embryonic pituitary cells. This indicates that any positive regulation of GILZ may have on gonadotropin β-subunit mRNA levels during embryogenesis is independent of their regulation by glucocorticoids. The distribution of GILZ in the avian pituitary, the ontogenic profile of FSHβ during embryonic development (Ellestad and Porter, unpublished), and the increase in FSHβ mRNA in the presence of exogenous GILZ are all consistent with a role for this gene in pituitary gonadotroph development. It should be noted that a role for GILZ in regulation of the pituitary hormones at later embryonic stages, or even post-hatch, cannot be ruled out by the current experiments.

In conclusion, this is the first report in which an avian GILZ homolog has been characterized and the first demonstration that GILZ is present and regulated by glucocorticoids in the anterior pituitary gland of any species. Due to the spatial and temporal conservation of pituitary gland development among vertebrates, the results presented should be relevant in mammalian systems. We have elucidated GILZ mRNA expression profiles during both embryonic and neonatal development. The expression profile is consistent with a potential role in initiation of hormone transcription during embryogenesis, particularly for those hormones that are known to be positively regulated by corticosteroids during development, namely GH and PRL. GILZ does not appear to play a role in somatotroph development, but may contribute to lactotroph development by playing a minor role in initiation of PRL expression in combination with other factors regulated by glucocorticoids. Finally, GILZ may also contribute to gonadotroph development by positively regulating gonadotropin β-subunit mRNA, primarily that of FSHβ, during embryogenesis.

Declaration of interest

The authors declare that there is no conflict of interest that would prejudice the impartiality of this scientific work.

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