Expression of thyroid hormone regulator genes in the yolk sac membrane of the developing chicken embryo

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Abstract. Thyroid hormones (THs) are essential for the correct development of nearly every structure in the body from the very early stages of development, yet the embryonic thyroid gland is not functional at these stages. To clarify the roles of the egg yolk as a source of THs, the TH content in the yolk and the expression of TH regulator genes in the yolk sac membrane were evaluated throughout the 21-day incubation period of chicken embryos. The yolk TH content (22.3 ng triiodothyronine and 654.7 ng thyroxine per total yolk on day 4 of incubation) decreased almost linearly along with development. Real-time PCR revealed gene expression of transthyretin, a principal TH distributor in the chicken, and of a TH-inactivating iodothyronine deiodinase (DIO3), until the second week of incubation when the embryonic pituitary-thyroid axis is generally thought to start functioning. The TH-activating deiodinase (DIO2) and transmembrane transporter of thyroxine (SLCO1C1) genes were expressed in the last week of incubation, which coincided with a marked increase of circulating thyroxine and a reduction in the yolk sac weight. DIO1, which can remove iodine from inactive THs, was expressed throughout the incubation period. It is assumed that the chicken yolk sac inactivates THs contained abundantly in the yolk and supplies the hormones to the developing embryo in appropriate concentrations until the second week of incubation, while THs may be activated in the yolk sac membrane in the last week of incubation. Additionally, the yolk sac could serve as a source of iodine for the embryo.

Key words: Chicken, Iodothyronine deiodinase, Real-time PCR, Thyroid hormones, Yolk sac

The avian yolk sac, or more precisely the yolk sac membrane, is not only a “bag” enveloping the yolk containing all the nutrients necessary for normal embryo development, but also a metabolic organ that possesses a number of enzymes converting these nutrients to facilitate embryonic use [1–3]. The two main structural components of the yolk sac membrane are the epithelial cell layer and the capillary layer. The former is a monolayer of endodermal cells in direct contact with the yolk, whereas the latter is of mesodermal origin and consists of vascular endothelial cells and smooth muscle cells [4]. During incubation of the chicken egg, the yolk is absorbed by the epithelial cells and transported to the embryo via vascular cells [1].

Unlike in the case of mammals where the mother can supply nutrients to the fetus through the placenta, avian embryos have to nurture themselves with whatever is confined in their own eggshells. The yolk contains lipophilic hormones, such as steroids and thyroid hormones (THs), deposited by the mother when the egg was formed in the ovary [5]. The THs, 3,5,3′-triiodothyronine (T₃, potent TH) and 3,5,3′,5′-tetraiodothyronine (T₄ or thyroxine, less active TH), are synthesized only in the thyroid gland. As in mammals, the THs are essential for the correct development of nearly every structure in the body in birds from the very early stages of development, yet the embryonic thyroid gland is not functional at these stages [6, 7]. It is therefore generally accepted that the yolk functions as a source of THs for the avian embryo by supplying them through the circulation before the embryo’s own thyroid gland starts to function [5].

In all vertebrate species studied so far, more than 99% of THs in the circulation are transported by binding to distributor proteins [8]. In the case of adult birds, THs are distributed mainly by binding to transthyretin (TTR, previously called pre-albumin) and albumin (ALB) [9]. By applying the comprehensive serial analysis of gene expression approach to study the total yolk sac membrane in the last week of a 21-day incubation of chicken embryos, the expression of TTR and ALB was shown [3], indicating a possibility of distribution of yolk THs to the embryo. Moreover, since THs do not readily cross the lipid bilayer membrane, the majority of THs are transported through the cell membrane via transmembrane transporters belonging to different transporter families, including members of the monocarboxylate transporter (MCT) and organic anion transporter (OATP) families [10]. Furthermore, the balance between activation and inactivation of THs within the cell is determined—largely independent of their circulating concentrations—by three types of
iodothyronine deiodinases (DIO1, DIO2, and DIO3, also known as D1, D2, and D3) located in non-thyroidal tissues [11, 12]. Activation of THs occurs through the removal of iodine (deiodination) at the 5′-position of T4 by DIO1 or DIO2, producing T3. Inactivation of THs occurs through the 5-deiodination of T3 and T2 by DIO1 or DIO3, producing reverse T3 (rT3) and diiodothyronine (T2), respectively [11, 12]. Gene expression of DIOs in yolk sac epithelial cells from embryonic day (E) 2 to E4 of chicken was revealed by microarray analysis [4], indicating a possibility of these enzymes regulating the balance of T3 and T4 distribution to the embryo or partially acting in the yolk sac membrane regardless of their respective concentrations in the yolk.

However, the fluctuation of these TH regulator molecules during the whole period of incubation of chicken embryos has never been characterized collectively. In the present study, we have analyzed the expression levels of the DIO1, DIO2, DIO3, TTR, and ALB genes in the same animal throughout the whole period of incubation to gain a broad-range view of the roles of the yolk sac membrane in processing and supplying THs to the developing embryo. In addition, the expression of SLC16A2 (MCT10 gene), SLC16A10 (MCT8 gene), SLCO1C1 (OATP1C1 gene), which are assumed to be responsible for transport of yolk THs in and out of the yolk sac membrane, was determined. The present results demonstrated the expression of DIO3 and TTR in the earlier period, DIO2 and SLCO1C1 in the later period, and DIO1 and ALB in both periods of incubation, and the chronological atypical expression of SLC16A2 and SLC16A10.

Materials and Methods

Animals and sampling

Fertilized eggs from layer strain (Hy-Line) chicken were purchased from Japan Layer, Gifu, Japan. The eggs were incubated at 37.8°C and 70% relative humidity. The day of onset of incubation was labeled embryonic day 0 (EO). All the embryos (chicks) were wipping with their beaks outside the eggshell at the time of E21 sampling. The yolk sac membrane was sampled daily from E4 embryos to 3-day-old chicks (C3) (n = 7–9). The membrane was rinsed thoroughly with sterile phosphate-buffered saline (PBS) with several changes to remove remaining yolk, put on a piece of filter paper to absorb excess PBS, weighed on an electronic balance, and then immediately frozen in liquid nitrogen and stored at −85°C until RNA extraction.

The yolk contents were collected every other day (n = 5), put into plastic tubes, weighed on an electronic balance, and stored at −30°C for measuring the yolk TH concentrations. Blood was collected from the extraembryonic blood vessel on E10–E16, from the embryonic heart on E17–E20, and by quick decapitation on E21–C3 (n = 7–9). E10 was the earliest day when we successfully collected blood that was not contaminated with other body fluids. Serum was prepared and stored at −60°C until measurement of the T4 concentration. All animal manipulations were approved by the Animal Research and Welfare Committee at Gifu University.

Quantification of mRNA expression

Total RNA was extracted from frozen yolk sac membranes using TRIzol reagent (Invitrogen, Carlsbad, CA, USA). RNase-free DNase I (gDNA Remover, TOYOBO, Osaka, Japan) was used to eliminate genomic DNA according to the manufacturer’s instructions. Total RNA (1 μg) was then reverse transcribed into first-strand cDNA using ReverTra Ace (TOYOBO) and a random primer (TOYOBO).

The mRNA levels were determined by real-time PCR, performed in an MX3000P Real-Time PCR system (Agilent Technologies, Santa Clara, CA, USA) with the two-step standard cycling program for the Brilliant III Ultra-Fast SYBR Green QPCR Master Mix (Agilent). The volumes of reagents used for a single reaction were 10 μl of QPCR Master Mix, 0.8 μl of each 10 μM forward and reverse primer, 7.4 μl of sterile water, and 1 μl of template cDNA. The cycling program consisted of 3 min of initial denaturation at 95°C, followed by 40 cycles of 5 sec of denaturation at 95°C and 20 sec of annealing and extension at 60°C. The amplification program was followed by a dissociation curve analysis to detect any nonspecific amplification. Each experimental and standard sample was assayed in duplicates. Relative expression values were calculated according to the standard curve method, using the software installed in the MX3000P System, with a serial dilution of pooled cDNAs as the standard. No-sample controls were included in the reactions to confirm the specificity of the reverse transcription and PCR amplification.

Using real-time PCR, we measured the expression of two housekeeping genes (GAPDH and RNA18S or 18S ribosomal RNA) in the yolk sac membrane throughout development and found RNA18S to be the most suitable since it did not show significant changes throughout development, with a difference of 12.4 ± 7.5% (mean ± standard deviation) between the mean expression values of each day. The primers used for real-time PCR are listed in Table 1. The primers for DIO2, DIO3, TTR, SLC16A2, SLCO1C1, and RNA18S are previously reported sequences for Gallus gallus. The primers for DIO1, ALB, and SLC16A10 were designed from G. gallus genomic sequences in the GenBank database using the online primer design software Primer3 [13]. The PCR-amplified products from these primers were sequenced by using the same primers; the determined sequences were 98–100% matched with the corresponding G. gallus sequences in the GenBank database.

We checked that the mRNAs quantified in the present study were virtually expressed in the yolk sac epithelial cells by scraping the cells off from the yolk sac membrane and analyzing the expression of these genes. The yolk sac membrane from E15–E20 eggs, thoroughly rinsed in sterile PBS as described above, was soaked in 0.05% trypsin-0.02% EDTA solution (Wako Pure Chemical, Osaka, Japan) at 4°C while scraping the epithelial cells gently with forceps under a dissecting microscope. Because some of the epithelial cells were still tightly adhered to the remaining membrane, we were unable to separate the vascular layer from the epithelial cell layer. The scraped cells were collected, rinsed three times in PBS by centrifugation (150 × g, 10 min, 4°C), and processed for RNA extraction and reverse transcription as described above. AmpliTaq Gold 360 DNA polymerase (Applied Biosystems, Foster City, CA, USA) was used for the PCR amplification according to the manufacturer’s instructions. The cycling program consisted of 5 min of initial denaturation at 95°C, followed by 35 cycles of 30 sec of denaturation at 95°C, 30 sec of annealing at 58°C, and 30 sec of extension at 72°C. The PCR products were electrophoresed on a 10% polyacrylamide gel, stained with ethidium bromide, and photographed.
Measurement of yolk THs by enzyme immunoassay

THs were extracted from the frozen yolk using the methanol/chloroform extraction procedure of Wilson and McNabb [7]. In brief, whole yolks (n = 5 for each day) were diced with scissors and homogenized in a plastic syringe with an 18-gauge needle by drawing and expelling the needle rod vigorously. The homogenized yolk (2 g) was transferred to a 50 ml polypropylene centrifuge tube, and 8 ml of methanol containing 1 mM propylthiouracil (Sigma-Aldrich, St. Louis, MO, USA) was added. The tube was shaken on a mixer (150 rpm, 10 min) and then centrifuged (1,700 × g, 10 min, 4°C), and the supernatant was decanted into another 50 ml tube. The precipitate was resuspended in 4 ml of methanol, shaken for 10 min, and centrifuged, and the supernatant was decanted into a third 50 ml tube. The two separate supernatants each received 20 ml of chloroform and 2 ml of 2 M ammonium hydroxide and were shaken and centrifuged in the same way. The upper phase of the two tubes was collected and combined in another 50 ml tube. The lower phase remaining in each tube was extracted again with ammonium hydroxide, and the upper phase from each was added to the 50 ml tube. The precipitate was resuspended in 4 ml of methanol, shaken for 10 min, and centrifuged, and the supernatant was decanted into a third 50 ml tube. The two separate supernatants each received 20 ml of chloroform and 2 ml of 2 M ammonium hydroxide and were shaken and centrifuged in the same way. The upper phase of the two tubes was collected and combined in another 50 ml tube. The lower phase remaining in each tube was extracted again with ammonium hydroxide, and the upper phase from each was added to the 50 ml tube. The sample was dried under a filtered air stream for 15°C. Once dried, the sample was resuspended in 4 ml of ammonium hydroxide, vortexed well, and centrifuged, and the supernatant was decanted into a polypropylene tube. The whole extraction procedure was then repeated for this supernatant with 4 ml of chloroform, and the upper phase was collected, dried, and resuspended in 300 μl of 75% ethanol. The reconstituted extracts were stored at −20°C until assayed for TH concentrations. The serum T_4 concentration was measured directly without extraction or dilution.

T_3 and T_4 were measured using enzyme immunoassay kits (Immunospec, Canoga Park, CA, USA). According to the manufacturer’s document, the cross-reactivity with related compounds other than T_3 and T_4 is less than 1.5%. The linearity of the yolk extract and the serum sample to the standard curve was confirmed by measuring serially diluted samples with the assay buffer provided in the immunoassay kit. The intra- and inter-assay variations were 6.2–12.4% and 3.8–12.9%, respectively.

To assess the recoveries of the above-described TH extraction procedure, fixed amounts of T_3 (2.0 ng) and T_4 (15.0 ng) were added to the thawed yolk before its homogenization. The yolk was processed as described above and the extracts were measured by enzyme immunoassay. The recoveries were 92.4 ± 6.6% for T_3 and 71.5 ± 5.9% for T_4 (mean ± standard error of five separate trials), which were numerically better than the recoveries reported in the original paper (61% for T_3 and 63% for T_4) [7].

Statistical analysis

Data were expressed as the mean ± standard error. Comparisons among groups were performed with one-way analysis of variance followed by the Tukey-Kramer multiple comparison test, using the JMP statistical package (ver. 10, SAS Institute Japan, Tokyo). A probability value of P < 0.05 was used throughout as the criterion to accept a significant difference of means.
Results

First, we have confirmed by standard PCR that all the genes determined in the present study were expressed in the yolk sac epithelial cells, as shown in Fig. 1. Expression of FOXA2 (forkhead box protein A2) [14] and CDH1 (E-cadherin) [4, 15], markers of endoderm-derived cells, was found in the epithelial cells scraped off from the yolk sac membrane, whereas PDGFRA (platelet-derived growth factor receptor alpha) [15], a marker of mesoderm-derived cells, was not amplified. In the rest of the yolk sac membrane (containing epithelial cell remnants and vascular layer; see Materials and Methods), however, PDGFRA was expressed as well as FOXA2 and CDH1 (data not shown).

Then, we examined the concentration of circulating THs as well as their content in the yolk. As shown in Fig. 2, the serum T4 concentration was very low (0.4–0.9 ng/ml) from E10 to E14, but then increased gradually from E15 (1.8 ng/ml), peaked significantly on E20 (36.6 ng/ml), and then dropped on E21 to 9.7 ng/ml, almost the same level as before the peak. The yolk weight (wet weight without yolk sac membrane) was 19.0 g on E4 and then gradually decreased along with development. As the embryos started hatching, the yolk weight decreased markedly from 6.19 g on E20 to 3.86 g on C1 and reached 0.66 g on C2. The total yolk TH content also showed significant decreases along with development; the yolk T3 content decreased from 22.3 ng (on E4) to 2.2 ng (on C2), and the yolk T4 content decreased from 654.7 ng (on E4) to 50.0 ng (on C2). The yolk TH concentrations did not change substantially during the incubation period (not shown in Fig. 2); 1.0 ng (T3) and 30.4 ng (T4) per gram yolk on E4, and 1.1 ± 0.4 ng (T3) and 27.9 ± 4.2 ng (T4) per gram yolk in the mean ± standard deviation of all data from E4 to C2. The weight of the yolk sac membrane (wet weight without yolk) was 0.67 g on E4 and increased gradually, especially on and after E11 (1.79 g), and then peaked on E17 (4.67 g). The yolk sac membrane shrunk thereafter, reaching 0.68 g on C3, which was almost the same weight as on E4.

Real-time PCR was performed to examine whether DIOs are expressed in the yolk sac membrane. As shown in Fig. 3, all three DIOs were expressed. DIO1 was found to be highly expressed as early as E4 and increased about 2-fold to reach its maximum levels on E5–E6. It then decreased gradually and was only slightly detectable by the time of hatching. The cause of the transient decrease on E13 could not be determined. DIO3 expression was very low on E4 but increased suddenly (~100,000-fold) to its maximum levels on E5–E6. It then decreased gradually and became very low again from E13 (~1/3,000 of E6 level) onwards. In contrast to these two DIOs, DIO2 expression was very low between E4 and E13, increased from E14, and peaked at around the time of hatching (5.5-fold increase from E14 to E20).

Next, we examined the expression level of TH distributors in the yolk sac membrane. As shown in Fig. 4, TTR was highly expressed as early as E4 and then increased about 2-fold, showing a peak on E5 and another peak on E10 with slightly lower levels in between, the lowest being on E8 at almost the same level as on E4. It is noteworthy that TTR expression became very low from E13 onwards (1/80 of E10 level on E15), which resembled the expression profile of DIO3. In contrast, ALB expression showed peaks in both the earlier (on E10) and later (on E15; ~1.7-fold higher than the E10 level) periods of development, and was very low from E19 onwards (1/60 of E15 level on E20). We again could not determine the reason for the transient decrease seen on E13.

Lastly, we measured the expression level of genes responsible for the transport of yolk THs. As shown in Fig. 5, TH transporter gene SLCO1C1 showed a mostly convex expression profile from E16 to E21. On the other hand, SLC16A2 and SLC16A10 both showed an atypical expression profile; SLC16A2 was predominantly expressed on E4–E6, E13–E15, and C1 but had very low expression in between

Fig. 1. Expression of thyroid hormone regulator genes in epithelial cells scraped off from the yolk sac membrane on embryonic day (E)18. CDH1 and FOXA2 were used as markers of endoderm-derived cells; PDGFRA (410 bp, no band was detected) was used as a marker for mesoderm-derived cells. M: molecular weight marker. Electrophoretogram in a 10% polyacrylamide gel.
These periods. *SLC16A10* was predominantly expressed on E4–E9, E13–E14, and E19–C1. Taken together, the results indicate that the expression of the three membrane transporters seems to be independent during embryo development.

To summarize, expression of the TH regulator genes could be divided into two periods, approximately before and after E10–E13. *DIO3* and *TTR* belong to the earlier period, whereas *DIO2* and *SLCO1C1* belong to the later period. *DIO1, ALB, SLC16A2,* and *SLC16A10* were expressed in both these periods.

**Discussion**

The present study has given us two perceptions to gain clues to unraveling the functions of the avian yolk sac membrane in processing...
and supplying yolk THs to the developing embryo. First, our results have confirmed previous observations that maternal THs in the yolk are available to the embryo [5]. Second, our study has shown that yolk THs could be metabolized before their distribution to the embryo or action in the yolk sac membrane.

Our results indicated that the yolk weight decreased significantly...
and almost linearly during the incubation period, as previously reported for chicken [16]. The total yolk TH content decreased in comparable tendency, and thus the yolk is likely to supply its components, including THs, to the developing embryo. THs in the circulation of adult chicken are mostly bound to TTR and ALB [9], and TTR synthesized in the visceral yolk sac of the rat is reported to be secreted toward the fetal circulation [17]. Therefore, the observed expression of TTR and ALB in the chicken yolk sac membrane in the present study is supportive evidence that the yolk supplies THs to the embryo through the circulation.

It is generally accepted that the thyroid gland of the chicken embryo starts secreting THs somewhere between E10 and E12, under the control of the thyroid-stimulating hormone from the pituitary gland [6, 18]. The present study showed that the expression of TTR, whose protein has high affinity for THs, was very low in the yolk sac membrane from E13 onwards. The role of the yolk sac as a source of circulating THs may thus weaken from around this time. Despite this, as clearly shown in Fig. 2, yolk THs were not yet depleted at this stage but went on to decrease further. A possibility is that ALB, whose gene expression increases between E14 and E18, contributes to TH distribution to the embryo in turn. Hilfer and Searls [19] reported that chicken embryos decapitated on E3 before the onset of pituitary function showed 100-fold less T₄ in the thyroid gland on E20, but only 2-fold less T₄ in the blood, compared with the levels in the control embryo, and concluded that most of the circulating T₄ was derived from the yolk. Their results seem to indicate that the yolk contains a sufficient amount of THs that can be supplied to the embryo even after the onset of thyroid function. The role, transporting route, and fate of yolk THs after the onset of thyroid function are thus intriguing and worthy of clarification.
Hitherto, experimental data in mammals have shown that maternal hyperthyroidism and hypothyroidism severely affect fetal brain development [20, 21]. An appropriate amount of THs is therefore essential for the correct development of the embryo. The yolk TH concentrations revealed in the present study (1.0 ng T₃ and 30.4 ng T₄ per gram yolk on E4) are relatively high, being more or less...
comparable with the peak of circulating THs (T3; ~2.6 ng/ml [22]; T4; 36.6 ng/ml; Fig. 2) at the time of hatching of the chicks, when active thermoregulation is necessary. It is assumed that the TH-inactivating DIO3, whose gene was expressed in almost the same period as TTR and ALB (E4–E12), could regulate the amount of TTR- and ALB-borne T3, thereby preventing unfavorable premature TH signaling in the earlier embryo. Such role of DIO3 is well documented in the liver of embryonic chicken in the last half of incubation, where high expression of the enzyme in the liver keeps the embryonic plasma T3 level low, in spite of the gradually increasing plasma T4, until the last days of embryonic development. Plasma T3 markedly increases to its peak level at the time of hatching, when hepatic DIO3 activity decreases by 98% from E17 to E21 [22, 23].

The expression of DIO2, the TH-activating deiodinase gene, was high between E14 and E21. This period corresponded to the increase of circulating T4, beginning at around E14 and reaching its peak on E20. The expression of SLC16C1, the gene of transmembrane transporter OATP1C1 with high preference for T3 [10], was also high at around this period. It may be responsible for transporting plasma T3 into the yol sac epithelial cells. Taking these results together, we speculate that the increased T4 in the circulation is taken into T3, thereby preventing unfavorable premature TH signaling.

Yolk sac carbohydrate levels and gene expression of key gluconeogenic and glycogenic enzymes during chick embryonic development. Poult Sci 2012; 91: 444–453. [Medline] [CrossRef]

The present results warrant further detailed immunocytochemical and cell biological studies of the yol sac membrane for elucidation of the subcellular mechanisms that regulate the processing and supply of THs from the yolk into the circulation.

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