Effects of chronic heat stress and ammonia concentration on blood parameters of laying hens

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ABSTRACT Less evidence is available currently to reveal whether the immune system and productivity of laying hens change under long periods of ammonia exposure in hot climate. The present study was conducted to determine the effects of chronic exposure to high temperature and ammonia concentrations on health, immune response, and reproductive hormones of commercial laying hens. A total of five hundred and seventy six 20-week-old laying hens (Hy-Line Brown) were used in this study. Birds were housed in cages (4 birds per cage) and received 16-wk treatments in 6 artificial environmental chambers. Hens were allocated to 6 treatments: treatment 1 (T1, 20°C, 5 ppm, control group), treatment 2 (T2, 20°C, 20 ppm), treatment 3 (T3, 20°C, 45 ppm), treatment 4 (T4, 35°C, 5 ppm), treatment 5 (T5, 35°C, 20 ppm), and treatment 6 (T6, 35°C, 45 ppm). Blood samples were collected at 22, 26, 30, 34, and 38 wk of age and plasma IgG, IgM, IgA, corticosterone (CORT), total antioxidant capacity (T-AOC), luteinizing hormone (LH), estradiol (E2), and follicular stimulating hormone (FSH) were measured. The results of this study showed that high ambient temperature and excessive ammonia increased the concentration of IgG but decreased the concentration of IgA, T-AOC, LH, FSH, and E2 of hens compared with those of the control birds. From the age of 34 wk, significantly increased concentrations of IgG were observed in hens exposed to moderate and high levels of ammonia. CORT level showed marked differences between the treatments only at the age of 26 wk. In addition, LH and E2 of hens demonstrated significant differences among the treatments in the middle and later stages of the experiment, while FSH levels of the control birds were significantly higher than the others at the age of 38 wk. Excessive ammonia in high temperature was a physiological stress factor that had a negative effect, which inhibited immune function and impacted the reproductive hormones.

Key words: heat stress, ammonia, blood parameter, immunity, laying hen

INTRODUCTION

Aerial ammonia is recognized as one of the most noxious odors in poultry operations, and its negative effect on the environment has been documented by numerous studies (Koerkamp and Bleijenberg, 1998; Koerkamp et al., 1998; Wheeler et al., 2000). Poultry production industry, as the largest contributor to ammonia emissions of all animal husbandry operations, has undergone considerable surveillance from public and regulatory agencies due to their environmental impacts (Hale et al., 2010; Lin et al., 2017). Meanwhile, poultry companies have to be concerned not only for worker health, but also for poultry health because elevated concentrations of atmospheric ammonia have an adverse impact on bird physiology and productivity (Carlile, 1984; Ritz et al., 2006; Almuhanna, 2011; Almuhanna et al., 2011; Kearney et al., 2014; Nemer et al., 2015). Ammonia is known as an irritant alkaline air contaminant, with occupational limits set at 50 ppm (OSHA, 2012) for the 8-h permissible exposure limit or 25 ppm (NIOSH, 2016), and a level of 300 ppm is considered to be immediately dangerous to human life and health (Wheeler et al., 2000).

Exposure of poultry to ammonia has led to significant impacts on health and growth performance. Excessive ammonia has been a physiological stress factor that would...
reduced feed intake and stunt growth (Charles and Payne, 1966; Deaton et al., 1984; Beker et al., 2004; Miles et al., 2004, 2006; Wei et al., 2014), decreased egg production significantly with 7-wk exposure to ammonia at an ammonia concentration of 102 ppm (Charles and Payne, 1966), and adversely affected egg quality (Cotterill and Nordskog, 1954). Moreover, it damaged the respiratory tract and lung atrial wall (Nagaraja et al., 1983; Kristensen and Wathes, 2000), and increased the incidence of diseases and secondary infections such as Newcastle disease, airsacculitis, and the prevalence of Mycoplasma gallisepticum (Anderson et al., 1964; Sato et al., 1973; Oyetunde et al., 1978). Controversial results indicated that excessive ammonia exposure lowered the immunological response of birds, which depended on bird age as well as ammonia level and duration. For broiler chickens, the hemagglutination inhibition antibody titer of Newcastle disease virus was significantly lower when exposed to 26 and 52 ppm ammonia compared to the 0 ppm treatment group for 21 D (Wang et al., 2010). On the contrary, average body weights, air sac scores, lung weights, relative bursa of Fabricius weights, and carcass grades were not significantly different among the birds exposed to ammonia concentration at 0, 25, or 50 ppm for 49 D (Caveny et al., 1981).

Previous studies have reported that the ammonia level inside poultry houses was affected by multiple factors, such as housing system, ventilation, manure management, bird density (Tasistro et al., 2007; David et al., 2015; Zhao et al., 2015), and components of the aerial environment including temperature, humidity, dust, and pathogens. All these may interact with ammonia (Koon et al., 1963; Dennis and Gee, 1973; Feddes et al., 1985; Kristensen and Wathes, 2000). Wathes et al. (1997) reported that the mean ammonia concentration of cage houses was as high as 24.2 ppm in summer. Miles et al. (2011) found that the maximum ammonia level was up to 7 times greater at 40.6°C vs. 18.3°C. A study carried out in floor systems revealed that most houses exceeded concentrations of 25 ppm ammonia, and in some areas went up to 80 ppm (David et al., 2015). All these studies indicated that high temperature was an important factor resulting in excessive ammonia. Additionally, both acute and chronic heat stress (≥35°C) have been documented to have an adverse influence on egg production and feed intake of laying hens (Deaton et al., 1981; Mahmoud et al., 1996; Mashaly et al., 2004). In terms of physiological health, previous studies have shown that heat stress impaired the activity of antioxidant enzymes and the redox system of the bird (Altan et al., 2003; Lin et al., 2008). Heat stress also resulted in the disruption of hormones responsible for ovulation and a decrease in responsiveness of granulosa cells to luteinizing hormone (LH) (Donoghue et al., 1989; Novero et al., 1991).

Various parameters have been used to evaluate the health and immune response in birds. Immunoglobulins G, M, and A are the 3 major classical Ig secreted by immune B-cells for the host against non-self antigens (Mestecky et al., 1999; Suzuki et al., 2007; Gomes et al., 2014). Total antioxidant capacity (T-AOC) was considered as an important indicator of animal health (Aslan et al., 2005; Gao et al., 2013; Zeng et al., 2014).

Laying hens live longer than broilers in poultry houses and may develop different disorders when suffering from extended periods of ammonia exposure (Beker et al., 2004). Although there are a multitude of researches on the adverse effects of ammonia on the health and performance of poultry, currently less evidence is available to reveal whether the immune system and productivity of laying hens change under long periods of ammonia exposure in hot climate. Therefore, this study was carried out to evaluate the effects of heat stress and ammonia on blood parameters and the interaction between them.

MATERIALS AND METHODS

Experimental Chambers

Six artificial environmental chambers (each 24.8 m², 4.5 × 5.5 m) in the College of Animal Science and Technology (Hebei Agricultural University, China) were used for the experiment. Cages (50 × 40 × 40 cm, L × W × H) were divided into 3 groups (8 cages per group) and were set evenly in the chamber for hen rearing (Figure 1). The chambers were computer programmed to keep the environmental conditions as required. Sensors installed in these chambers were used to monitor temperature, humidity, ammonia, and carbon dioxide concentration every 10 s, which were then displayed on the computer screen for observation.

The supply system of ammonia gas has been studied in detail earlier (Kristensen et al., 2000) in order to maintain the desired ammonia concentrations. Compressed anhydrous ammonia was stored in a cylinder and its flow rate was adjusted by a flow meter. This design enabled each chamber to be filled with gaseous ammonia independently of the others.

Light emitting diode lamps were fixed on the ceiling of the chamber and could be adjusted to provide the required light intensity. Temperature was adjusted...
automatically by an air conditioner, and air inlet and outlet were trepanned for mechanical ventilation. Feeder and drinker were provided in each chamber and video cameras were fitted.

### Bird Management and Experimental Design

A total of 576 Hy-Line Brown hens acquired from a commercial farm were used for this study. The study proposal was approved by the Laboratory Animal Ethical Committee of China Agricultural University. Birds were acclimatized for 2 wk in the chambers at the age of 20 wk before treatment. There were a total of 6 treatments of 3 ammonia concentration exposures (≤5, 20, and 45 ppm) under 2 temperatures (20 ± 2°C, 35 ± 2°C) as shown in Table 1. T1 was the control group. Birds were randomly assigned to 6 groups and were transferred to the cages (4 birds per cage). Each treatment had 3 replications with 32 birds per replicate.

All birds were provided with a corn-soybean basal diet; feed and water were provided ad libitum. Ambient relative humidity was maintained at 40 to 60% and carbon dioxide level was kept below 1,000 ppm during the experimental period.

The duration of light exposure during the growing period from 20 wk of age was 13.5 h (4:00–17:30) per day for hens in the first week and then stepped up gradually per week until it reached 16 h (4:00–20:00) at 31 wk of age. From then on, permanent illumination of 16 h was employed from 32 to 38 wk of age. Besides, light intensity of 30 lx was equalized at bird head level following the Hy-Line Brown Layers Guide Manual (Hy-Line International; http://www.hyline.com).

### Sample Preparation and Collection

For each sampling, 4 birds were randomly selected from each replicate and marked with tags for the following operation. Wing venipuncture was performed for blood collection after fasting for 12 h starting at week 22, 26, 30, 34, and 38. Plasma was prepared after centrifugation and stored at −20°C.

### Measured Contents and Methods

The concentrations of plasma Ig (IgG, IgA, and IgM), corticosterone (CORT), T-AOC, and reproductive hormones (LH, estradiol [E2], and follicular stimulating hormone [FSH]) were detected using commercially available ELISA kits (Jianglai Biological Technology Co. Ltd., Shanghai, China).

### Statistical Analysis

All statistical analyses were performed using linear mixed models parameterized with SPSS (IBM SPSS Statistics 25.0; IBM Corporation, Armonk, NY, USA). The data were analyzed with fixed effects of temperature, ammonia concentration, week of age, and the random effect of replicate. The model equation was as follows:

\[
Y_{ijku} = \mu + T_i + AC_j + WOA_k + R_{u} + T \times AC_{ij} + T 
\times WOA_{ik} + AC \times WOA_{jk} + \epsilon_{ijk},
\]

where \(AC\) = ammonia concentration; \(Y_{ijku}\) = parameters investigated; \(\mu\) = model constant; \(T_i\) = effect of temperature \((i = 1–2)\); \(AC_j\) = effect of ammonia concentration \((j = 1–3)\); \(WOA_k\) = effect of age \((k = 1–5)\); \(R_u\) = effect of replicate \((u = 1–3)\); \(T \times AC_{ij}\) = effect of interaction between temperature and ammonia concentration; \(T \times WOA_{ik}\) = effect of interaction between temperature and week of age; \(AC \times WOA_{jk}\) = effect of interaction between ammonia concentration and week of age; and \(\epsilon_{ijk}\) = the residual error term.

Effects in the statistical model were tested simultaneously and they were removed from the original model when they were not significant. When the effect was statistically different \((P < 0.05)\), further analysis was needed. Duncan’s multiple range (least significant difference) test was applied for post-hoc group comparisons.

### RESULTS

Statistical analysis showed that for all the blood parameters, the effect of replicate was not significant and it was excluded from the original model.

### Plasma Ig Concentrations

As shown in Table 2, temperature, ammonia concentration, and bird age had significant effects on IgG and IgA \((P < 0.05)\). However, IgM was not affected by temperature, ammonia concentration, and bird age \((P > 0.05)\). The concentration of IgG was increased with increasing temperature. However, the concentration of IgA was decreased under 35°C treatment. Compared with the treatment of low ammonia concentration \((≤5 \text{ ppm})\), the level of IgG was significantly increased under ammonia concentrations of 20 and 45 ppm \((P < 0.05)\), while the concentrations of IgA were significantly decreased \((P < 0.05)\). Furthermore, IgG increased steadily with age \((P < 0.05)\).

Less change was observed in the plasma concentration of IgG and no significant difference was found among the 6 treatments from 22 to 30 wk of age (Figure 2). A significant increase of IgG occurred in all treatments at 34 wk of age \((P < 0.05)\), and this kept increasing until 38 wk of age. Exposure to high temperature and ammonia resulted in a significant elevation in IgG \((P < 0.05)\). At 34 wk of age, plasma IgG concentrations of hens in T6, T5, and T3 were significantly higher than those of T1 \((P < 0.05)\), and the differences increased at 38 wk.
of age ($P < 0.05$). IgM and IgA levels of birds varied from 22 to 38 wk of age. Compared to other treatments, the IgA level in T6 was significantly lower at week 38 ($P, 0.05$).

**Plasma CORT and T-AOC Levels**

Table 3 shows that CORT level was affected by temperature and week of age ($P < 0.05$) significantly and a 2-way interaction was found between them. All factors significantly affected the levels of T-AOC ($P < 0.05$). Levels of CORT and T-AOC were significantly decreased with increased temperature ($P < 0.05$). Furthermore, the concentrations of T-AOC were significantly decreased with increased ammonia concentrations ($P < 0.05$). T-AOC increased with age and significant differences were found ($P < 0.05$). Levels of CORT peaked (73.06 ng/mL) at 26 wk of age ($P < 0.05$), compared to that at 22 wk of age (63.37 ng/mL), and decreased (62.51 ng/mL) until 38 wk of age.

Blood CORT and T-AOC concentrations of the laying hens exposed to different ammonia concentrations and temperatures are shown in Figure 3. An increase in CORT was observed after 4 wk of treatment at the age of 26 wk. Then, CORT levels decreased at 30 wk of age and this was maintained until the end of the experiment at 38 wk. Significant differences among the treatments were found only at 26 wk of age ($P < 0.05$). Birds exposed to high ammonia concentrations had significantly higher CORT levels than those in T1 ($P < 0.05$). Less change was observed in plasma T-AOC at 22 and 26 wk of age. Then, plasma T-AOC of T1 increased and was significantly higher compared to the other groups at 30 wk of age. At the age of 34 wk, the T-AOC levels of birds in all treatments increased significantly and at 38 wk they reached their maximum. From 30 wk of age, the plasma T-AOC level of birds in T1 was significantly higher ($P < 0.05$) compared to the high temperature and ammonia treatments and the difference increased significantly with age.

![Figure 2](image_url)

**Figure 2.** (A) Plasma IgG, (B) IgM, and (C) IgA concentrations of laying hens exposed to different temperatures and ammonia concentrations at 22, 26, 30, 34, and 38 wk of age. Data represent mean ± SE. $abc$ Values marked with different letters are significantly different ($P < 0.05$).
Plasma LH, E2, and FSH

As presented in Table 4, temperature, ammonia concentration, and age significantly affected the levels of LH and E2 ($P < 0.05$). Temperature and ammonia concentration had a significant influence on FSH. Moreover, LH, E2, and FSH levels were significantly decreased with higher temperature ($P < 0.05$). High ammonia concentrations of 45 ppm significantly decreased the levels of these 3 hormones compared to those at low ammonia concentration ($≤5$ ppm) ($P < 0.05$).

The change of plasma LH, FSH, and E2 concentrations in different treatments with age of birds is shown in Figure 4: A (LH), B (FSH), and C (E2). Significant differences of LH levels between T1 and T6 were observed at 30, 34, and 38 wk of age ($P < 0.05$). Hens treated with high temperature and ammonia concentration presented decreased LH levels. Plasma FSH of birds in T2 to T6 remained constant from 22 to 38 wk of age. In contrast, the FSH levels of birds in the control group increased with age. As a result, birds in the control group had a significantly higher level of FSH than birds subjected to high temperature and ammonia treatments at the age of 38 wk ($P < 0.05$). E2 levels showed a similar trend to LH. At each sampling week, T1 had the highest level of E2 in all treatments while T6 had the lowest level of E2.

DISCUSSION

The effect of temperature and ammonia on immune parameters and reproductive-related hormones of laying hens was investigated in this study. The results showed that the blood parameters of laying hens were affected by high temperature, ammonia concentration, and age, and there was an interaction between them.

Heat stress and ammonia exposure caused a significant increase in plasma levels of IgG at the age of 34 and 38 wk of laying hens. Previous studies have indicated that stressors, such as heat and ammonia, are able to elevate IgG levels (Nasrin et al., 2013; Honda et al., 2015; Wu et al., 2017), consistent with the results of this study. On the other hand, increased levels of IgM were observed at the age of 22 wk and it decreased from 26 wk in high temperature and ammonia treatments in this study. Honda et al. (2015) reported that birds exposed to 19-D heat stress of $38^\circ C$ had higher plasma IgM levels. Meanwhile, Wu et al. (2017) indicated that plasma IgM level reduced after 45 wk of 30-ppm ammonia pollution. The early increase of IgM at the beginning of egg laying may have been caused by high temperature and late reduction was due to ammonia exposure. As the experiment went on,

Table 3. Plasma CORT and T-AOC levels of laying hens at 22, 26, 30, 34, and 38 WOA exposed to different ammonia concentrations under $20^\circ C$ and $35^\circ C$.

| Item$^1$ | CORT (ng/mL) | T-AOC (U/mL) |
|---|---|---|
| T ($^\circ C$) | | |
| 20 | 68.09 ± 1.03$^a$ | 9.11 ± 0.32$^a$ |
| 35 | 66.56 ± 0.86$^b$ | 8.15 ± 0.27$^b$ |
| AC (ppm) | | |
| ≤5 | 65.90 ± 0.94 | 9.94 ± 0.47$^a$ |
| 20 | 65.14 ± 1.05 | 8.53 ± 0.33$^b$ |
| 45 | 68.66 ± 1.45 | 7.41 ± 0.22 |
| WOA | | |
| 22 | 63.37 ± 1.30$^b$ | 5.87 ± 0.12$^a$ |
| 26 | 73.06 ± 1.50$^a$ | 6.04 ± 0.13$^a$ |
| 30 | 69.84 ± 1.31$^a$ | 6.58 ± 0.19$^a$ |
| 34 | 64.05 ± 1.59$^b$ | 11.17 ± 0.32$^b$ |
| 38 | 62.51 ± 1.34$^b$ | 13.67 ± 0.49$^a$ |

$^1$T, temperature; AC, ammonia concentration; WOA, week of age.

$^2$NS, no significance.

$^a$–cMeans within main effects without a common letter differ ($P < 0.05$).

Abbreviations: CORT, corticosterone; T-AOC, total antioxidant capacity.

$^1$T, temperature; AC, ammonia concentration; WOA, week of age.

$^2$NS, no significance.

Figure 3. (A) Plasma corticosterone (CORT) and (B) total antioxidant capacity (T-AOC) levels of laying hens exposed to different temperatures and ammonia concentrations at 22, 26, 30, 34, and 38 wk of age. Data represent mean ± SE. a–eValues marked with different letters are significantly different ($P < 0.05$).
ammonia concentration had a greater effect on IgM than that of high temperature at peak egg production. The decrease of IgA caused by ammonia and heat stress may be explained by the result reported by Nasrin et al. (2013), who showed that heat stress and Newcastle disease virus resulted in increased IgA concentrations in the trachea, cecal tonsils, and Harderian gland. In other words, the concentration of IgA in stressed birds increased in lymphoid organs, where immune system activation occurs, but not in the peripheral blood. Therefore, the different changes of Ig (IgG, IgM, and IgA) may be determined by the type of stress and its intensity and duration. In addition, different types of stress had different effects on different periods of egg laying. Overall, the present study showed that heat and ammonia stress in the early laying period of laying hens impacts the immune system and activity for a long period of time.

Corticosterone is a common hormone that is released in many stress situations by the hypothalamus–pituitary–adrenal axis. It is a glucocorticoid that has been reported to dysregulate immune responses and is potentially harmful to the health of animals if it remains at a high level for a long period of time (Post et al., 2003). Shini et al. (2009) demonstrated that oral CORT treatment affected the physiology of hens, reduced performance, and may model the effects of production stressors. Negative correlation between plasma CORT levels and immune response under heat stress has been reported in birds (Elenkov et al., 2000; Quinteiro-Filho et al., 2010; Calef et al., 2014). In our study, high temperature and ammonia exposure significantly increased the plasma level of CORT after 4-wk treatment, which indicated that both were stressors during the growth of laying hens. However, there were no statistically significant changes of measured CORT level in laying hens from 30 wk of age, which suggested that hens may have the capability to adapt to moderate and high levels of chronic ammonia stimulation and heat stress after a certain period of time. Therefore,

Table 4. Plasma LH, FSH, and E2 levels of laying hens at 22, 26, 30, 34, and 38 WOA exposed to different ammonia concentrations under 20°C and 35°C.

| Item  | LH (pg/mL) | FSH (mIU/mL) | E2 (pg/mL) |
|-------|------------|--------------|------------|
|       | T (°C)     |              |            |
| 20    | 53.89 ± 0.69a | 8.34 ± 0.16a | 94.64 ± 1.49a |
| 35    | 45.48 ± 0.80b | 7.68 ± 0.14b | 81.95 ± 1.30b |
| AC (ppm) |          |              |            |
| ≤5    | 51.99 ± 1.02a | 8.55 ± 0.20a | 99.32 ± 0.78a |
| 20    | 49.88 ± 0.90b | 7.87 ± 0.17b | 86.64 ± 1.50b |
| 45    | 47.19 ± 1.02b | 7.61 ± 0.17b | 78.92 ± 1.60c |
| WOA |           |              |            |
| 22    | 49.37 ± 0.95b | 8.70 ± 0.23 | 84.09 ± 1.74c |
| 26    | 55.49 ± 0.96a | 8.15 ± 0.22 | 85.08 ± 1.89b,c |
| 30    | 49.02 ± 1.37b,c | 7.63 ± 0.22 | 91.38 ± 2.29b,c |
| 34    | 48.38 ± 1.37b,c | 8.17 ± 0.23 | 93.53 ± 2.64a,b |
| 38    | 45.58 ± 1.40c | 8.31 ± 0.27 | 87.39 ± 2.76a,b,c |

P-value

|        | T     | AC    | WOA   |
|--------|-------|-------|-------|
| T × AC | <0.05 | NS    | <0.05 |
| T × WOA| <0.05 | NS    | <0.05 |
| AC × WOA| <0.05 | NS    | NS    |

\(^{a–c}\)Means within main effects without a common letter differ (P < 0.05).

Abbreviations: E2, estradiol; FSH, follicular stimulating hormone; LH, luteinizing hormone.

1T, temperature; AC, ammonia concentration; WOA, week of age.

2NS, no significance.

Figure 4. (A) Plasma luteinizing hormone (LH), (B) follicular stimulating hormone (FSH), and (C) estradiol (E2) levels of laying hens exposed to different temperatures and ammonia concentrations at 22, 26, 30, 34, and 38 wk of age. Data represent mean ± SE. \(^{a–d}\)Values marked with different letters are significantly different (P < 0.05).
CORT was not considered to be a stress indicator during prolonged exposure of high temperatures and ammonia from the age of 30 wk until the end of the experiment. The SEMs of CORT were relatively large, which may indicate that CORT varied widely among individuals. Oxidative stress was part of the stress response of birds to heat exposure and other stressors (Dröge et al., 2002; Lin et al., 2006). Devi et al. (2000) and Thomas et al. (2000) reported that increased activities of antioxidant enzymes were considered to be a protective response against oxidative stress. In the present study, the T-AOC of layer hens in the control group was significantly higher than that of other treatments at 30 wk of age. It indicated that stress suppressed the T-AOC release of birds under chronic treatment. This result is in agreement with a research in ducks which mainly released birds under chronic treatment. This result is not conducive to the health of laying hens. Only in the treatment would increase the stress of laying hens, which may have been caused by depressing ovarian functions due to the decreased secretion of LH and FSH (Younghen et al., 1991; You et al., 1995; Rozenboim et al., 2007). Another possible mechanism for the reduction of ovarian function might be that heat stress can cause a reduction in blood flow to the ovary of cattle (Wolfenson et al., 1981, 1997), which may be another explanation for the result of our study. All these results of the 3 reproductive hormones may have an influence on ovarian function, and then may result in a decrease in reproductive performance or reproductive failure.

This study indicated that heat stress and high concentration of ammonia were negative factors for the immune system and increased the incidence of diseases. Based on the changes of IgG, ammonia concentration and temperature should be more strictly controlled during peak egg production. In addition, according to the indication of Ig, a normal temperature at the beginning of the laying process should be maintained, while ammonia concentration should be focused on during peak egg production. The difference of CORT concentration in 6 treatments indicated that high temperature and ammonia concentration would increase the stress of laying hens, which is not conducive to the health of laying hens. Only in the middle and late stages of this experiment, the blood T-AOC concentrations of laying hens showed significant differences, which indicated that under heat stress, the hen’s antioxidant capacity decreased significantly with ammonia concentration increasing after long periods of treatment. The reproductive hormone data presented in this study suggested that potential reproductive failure associated with high environmental temperature and ammonia concentration might be caused directly by depressing ovarian functions or decreasing the related hormones secreted by hypophysis.

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