

Oral administration of D-Aspartate decrease body temperature in broiler chicks

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Orally Administered D-Aspartate Depresses Rectal Temperature and Alters Plasma Triacylglycerol and Glucose Concentrations in Broiler Chicks

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1 L-Aspartate (L-Asp), D-aspartate (D-Asp) or their chemical conjugates plays important physiological roles in regulating food intake, plasma metabolites and thermoregulation in animals. However, there are very few studies available in layers and no reports have been found in broilers. Broilers are very important commercial birds for meat production, so effects of L- or D-Asp in broilers would provide new physiological insight of this strain. Therefore, the purpose of this study was to determine the effect of oral administration of L- or D-Asp on feed intake, rectal temperature and some plasma metabolites in broiler chicks. Broiler chicks (5 days old) were orally administered with different doses (0, 3.75, 7.5 and 15 mmol/kg body weight) of L- or D-Asp. At 120 min after administration of L- or D-Asp, the blood was immediately collected through the jugular vein. The rectal temperature of chicks was measured at 30, 60 and 120 min after administration using a digital thermometer with an accuracy of $\pm 0.1^{\circ}\text{C}$, by inserting the thermistor probe in the rectum to a depth of 2 cm. A repeated-measures two-way ANOVA was applied for the analysis of feed intake and rectal temperature. Plasma metabolites were statistically analyzed by one-way ANOVA and regression equations. The study showed that oral administration of both L- and D-Asp did not alter feed intake. However, D-Asp, but not L-Asp, dose-dependently decreased the rectal temperature in chicks. It was also found that D-Asp increased plasma glucose and decreased triacylglycerol concentrations. The changes in plasma metabolites further indicate that D-Asp treatment modulates the energy metabolism in broiler chicks. In conclusion, D-Asp may be a beneficial nutrient not only for layers but also for broilers, since orally administered D-Asp lowered rectal temperature without reducing feed intake.

Key words: broiler chicks, D-aspartate, L-Aspartate, plasma metabolites, rectal temperature

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Introduction

High ambient temperature (HT) induced heat-stress is a serious concern over the globe. The Intergovernmental Panel on Climate Change (IPCC) (2015) reported that the global surface temperature is increasing in this century. Therefore, probably summer time is becoming more unbearable in many tropical and subtropical countries. It is well known that summer heat stress is causing a great economic loss in commercial poultry sector. Donkoh (1989) reported

that continuous exposure of broilers to HT markedly affected their performance and physiological functions. In addition, HT induces quick increment of deep body temperature even in chicks (Chowdhury *et al.*, 2012) and causes oxidative stress (Chowdhury *et al.*, 2014). To cool down the temperature in the poultry house using fossil energy leads further increase in the environmental temperature. Therefore, it is very important to find out other strategies to control the body temperature in the chicken. For instance, suitable nutrients or drugs to reduce high body temperature when exposed to HT should be clarified. Nutrients, especially amino acids, widely applied as anti-stress agents with regard to psychological and physiological stress (Yamane *et al.*, 2009; Hamast *et al.*, 2010; Erwan *et al.*, 2012, 2014b). Amino acid in particular essential amino acids supplementation has been performed in order to overcome heat-stress problems in

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birds (Mendes *et al.*, 1997; Brake *et al.*, 1998; Willemsen *et al.*, 2011; Dai *et al.*, 2012). However, not only essential amino acids, but also non-essential amino acids may be useful to mitigate heat-stress problems. Recently, Chowdhury *et al.* (2015) showed that a non-essential amino acid, L-citrulline, reduced body temperature in young chicks and suggested its possibility to use as a hypothermic agent under heat-stress in chickens.

Amino acids can exist in two-mirror-image form (L- or D-form) termed enantiomers. Aspartate (Asp), which is a non-essential amino acid, plays important roles as a constituent of protein synthesis, a precursor of specific neurotransmitters and physiological modulators (Spinelli *et al.*, 2006; Schell *et al.*, 1997; Errico *et al.*, 2009). L- and D-Asp have been shown to occur in various brain regions of chickens (Neidle and Dunlop, 1990) and pigeons (Kera *et al.*, 1996). As far as the physiological roles in correlation with D-Asp in mammals and birds are concerned, important biological functions of this amino acid are now postulated (Topo *et al.*, 2010). Though some evidence to date suggests that D-Asp plays important physiological roles in adult organisms, little is known about the function of D-Asp during an early development when its level is physiologically high (Homma, 2007; Errico *et al.*, 2008).

It has been reported that the injection of D-Asp released the antidiuretic hormone arginine vasopressin (AVP) in rats (Koyuncuoğlu *et al.*, 1984). Studies in rats have further shown that D-Asp not only lowered body temperature after oral administration (Koyuncuoğlu *et al.*, 1982a; Koyuncuoğlu and Berkman, 1982), but also decreased plasma triacylglycerol (TG) (Koyuncuoğlu *et al.*, 1982b). We have previously shown that intracerebroventricular (i.c.v.) injection of L- or D-Asp clearly attenuated stress responses in layer chicks (Yamane *et al.*, 2009; Erwan *et al.*, 2012, 2014b) and oral administration of D-Asp decreased feed intake, body temperature and altered some plasma metabolites in layer chicks (Erwan *et al.*, 2013a, 2014a). Hence, L- or D-Asp may act as a modulator of important physiological functions in chicks. However, some physiological functions are different between layer and broiler chicks. Tachibana *et al.* (2001) reported that agouti-related peptide, a potent stimulator of feed intake (Rossi *et al.*, 1998), stimulated feed intake only in layer type chicks but not broiler chicks. Therefore, it is important to know whether D-Asp can regulate body temperature and influence feed intake in broilers. The purpose of this study was to evaluate the effect of L- or D-Asp on feed intake, body temperature and plasma metabolites in broiler chicks.

Materials and Methods

Animals and Drugs

One-day-old broiler chicks (Cobb) (*Gallus gallus domesticus*) were purchased from a local hatchery (Charoen Pokphand Jaya Farm Ltd, Pekanbaru, Indonesia) and housed in a wooden cage (50×35×33 cm) in a group (25 birds) at a constant temperature of 30±1°C with continuous lighting. Feed (metabolizable energy: >3,050 kcal/kg, protein: >

23.5%: commercial starter diet, 311-VIVO, Pokphand Tbk, Medan, Indonesia) and water were provided *ad libitum*. One day before the experiment (4 days old), thirty two chicks (Means±S.E.M. were 92.1±2.0 g in Experiment 1 and 92.8±1.7 g in Experiment 2) were reared individually in cages (20×25×25 cm) and assigned for treatment and control groups on the basis of their body weight in order to produce uniform groups in both Experiments 1 and 2. This study was performed in accordance with the guidelines for animal experiments carried out at the Faculty of Agriculture and Animal Science, State Islamic University of Sultan Syarif Kasim Riau, Pekanbaru, Indonesia.

Both of L- and D-Asp were purchased from Wako Pure Chemical Industries (Osaka, Japan).

Administration of L- or D-Asp

Following an acclimatization period with individual rearing for 24 hr, chicks were randomly selected and divided into four groups, each group consisting of 8 chicks. The birds were provided with *ad libitum* access to the diet during the whole experimental period. On the day of the experiment, each chick (5 days old) was orally administered either a solution of L-Asp (Experiment 1) or D-Asp (Experiment 2) for the treatment groups, or distilled water (DW) for the control group, via an elastic plastic needle on a small syringe. Based on our recent report on L- or D-Asp in layer chicks (Erwan *et al.*, 2013a, 2014a), oral administration of L- or D-Asp in Experiments 1 and 2, has been designed to use 3.75, 7.5 and 15.0 mmol/kg body weight as the low, medium and high doses, respectively.

Measurement of Feed Intake and Rectal Temperature

Feed intake (at 30, 60 and 120 min) was determined by measuring the reduction in the amount of feed consumed from a pre-weighed feeder. The rectal temperature of chicks was also measured at 30-, 60- and 120-min of the experimental time with a digital thermometer with an accuracy of ±0.1°C (Thermalert TH-5, Physitemp Instruments Inc., USA) by inserting the thermistor probe into the rectum through cloaca to a depth of 2 cm from anus. It took about 5 sec to measure rectal temperature by inserting the probe.

Blood Collection and Analysis of Plasma Metabolites

At 120 min after administration of L- or D-Asp, all birds in each group (n=6-8 per group) were sacrificed and their bloods were collected for analysis of plasma metabolites. Blood was collected through jugular vein into heparinized eppendorf tubes and centrifuged at 5,000×g for 15 min. Plasma was stored at -20°C until assay. The plasma metabolites (TG, glucose, total cholesterol (TCHO) and total protein (TP)) were measured with Microlab 300 (Vital Scientia, Netherland) as per the manufacturer's instructions. All the samples were assayed together and in a random sequence for each metabolite.

Statistical Analysis

For the rectal temperature, a repeated-measures two-way analysis of variance (ANOVA) was applied. Plasma metabolites were statistically analyzed by one-way ANOVA and regression equations. The Tukey test was done as a post hoc test. Significant differences were denoted as $P<0.05$.

Values were presented as means \pm S.E.M. Statistical analysis was carried out using the commercially available package StatView (Version 5, SAS Institute, Cary, USA, 1998). All data in each group were first subjected to a Thompson rejection test to eliminate outliers ($P < 0.05$), and the remaining data were used for the analysis among groups.

Results

Experiment 1: Effects of L-Asp on Feed Intake, Rectal Temperature and Plasma Metabolites

Oral administration of several doses of L-Asp did not significantly alter feed intake at 2 h in chicks (Means (g) \pm S.E.M. were 47.8 ± 1.8 in the control group, 47.5 ± 2.8 in the low L-Asp group, 47.9 ± 1.3 in the medium L-Asp group, and 50.3 ± 1.5 in the high L-Asp group, respectively). As shown in Fig. 1, rectal temperatures of chicks were not changed significantly by the oral administration of L-Asp. Table 1 shows the effect of oral administration of several doses of L-Asp on the concentration of plasma metabolites. There was no significant effect of L-Asp on the concentration of plasma metabolites.

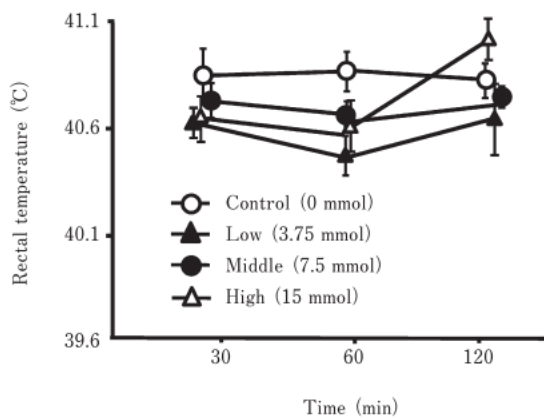


Fig. 1. Effect of orally L-Asp (3.75, 7.5 and 15.0 mmol/kg) administration on rectal temperatures in chicks during 120 min of the experimental period (Experiment 1). The number of chicks used in each group ranged between 5–7. Values are means \pm S.E.M.

Experiment 2: Effects of D-Asp on Feed Intake, Rectal Temperature and Plasma Metabolites

Oral administration of several doses of D-Asp did not significantly alter feed intake in chicks (Means (g) \pm S.E.M. were 46.6 ± 2.1 in the control group, 46.7 ± 2.8 in the low L-Asp group, 49.0 ± 2.6 in the medium L-Asp group, and 47.9 ± 2.0 in the high L-Asp group, respectively). However, D-Asp significantly declined rectal temperatures in broiler chicks ($F(3, 20) = 5.81$, $P < 0.005$) as shown in Fig. 2. D-Asp further showed a significant effect of time ($F(3, 2) = 25.09$, $P < 0.0001$) and an interaction between the dose and time ($F(11, 40) = 4.70$, $P < 0.005$) on declining body temperature, implying that the effect of D-Asp was more prominent with the progress of time and the low dose of the treatment caused more reduction in body temperature. Table 2 shows the changes in plasma metabolites due to the oral administration of D-Asp. A significant ($P < 0.005$) negative correlation was detected between the administered doses of D-Asp and the plasma concentration of TG ($143.4 - 4.1 X$, $R^2 = 0.295$). Plasma TG concentration significantly decreased by 15 mmol/kg body weight D-Asp compared with the control

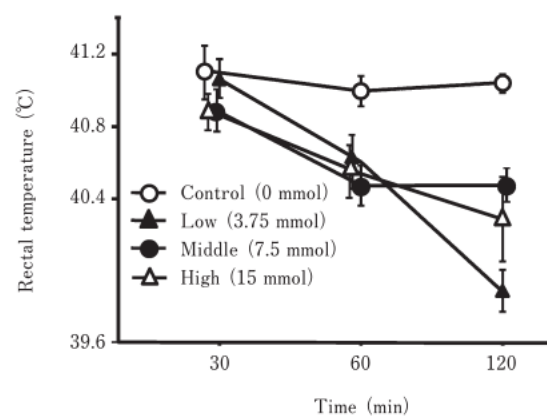


Fig. 2. Effect of orally D-Asp (3.75, 7.5 and 15.0 mmol/kg) administration on rectal temperatures in chicks during 120 min of the experimental period (Experiment 2). The number of chicks used in each group ranged between 5–7. Values are means \pm S.E.M.

Table 1. Effects of oral administration of several doses of L-Asp on plasma metabolites in chicks (Experiment 1)

Parameters	L-Asp (mmol/kg)				R2
	0	3.75	7.5	15	
Glucose (mg/100 ml)	139 ± 11	127 ± 5	127 ± 4	125 ± 3	0.058
Total cholesterol (mg/100 ml)	192 ± 8	192 ± 9	181 ± 11	194 ± 13	0.0001
Total protein (g/100 ml)	1.57 ± 0.1	1.54 ± 0.1	1.26 ± 0.1	1.54 ± 0.2	0.022
Triacylglycerol (mg/100 ml)	150 ± 11	157 ± 23	154 ± 14	140 ± 14	0.04

Values are means \pm SEM. The number of samples used for analysis was 6–8.

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Table 2. Effects of oral administration of several doses of D-Asp on plasma metabolites in chicks (Experiment 2)

Parameters	D-Asp (mmol/kg)				R2
	0	3.75	7.5	15	
Glucose (mg/100 ml)	124±4 ^a	131±2 ^{ab}	129±3 ^{ab}	140±3 ^b	0.348
Total cholesterol (mg/100 ml)	204±5	210±7	232±22	200±12	0.0001
Total protein (g/100 ml)	1.67±0.1	1.62±0.1	1.63±0.1	1.74±0.1	0.023
Triglycerol (mg/100 ml)	147±15 ^a	107±15 ^{ab}	136±9 ^a	75±8 ^b	0.295

Means with different superscripts in the same row were significantly different at $P<0.05$.

Values are means±SEM. The number of samples used for analysis was 6–8.

($F(3, 25)=6.51$, $P<0.005$). On the other hand, a significant positive correlation was observed between the administered doses of D-Asp and the plasma concentration of glucose (124.6 ± 1.0 X, $R^2=0.348$). Plasma glucose concentration was significantly increased by 15 mmol/kg body weight D-Asp compared with the control ($F(3, 23)=5.00$, $P<0.01$). The values of TP and TCHO showed no significant changes in any of the treated groups.

Discussion

We confirmed that oral administration of L- or D-Asp did not affect feed intake in broiler chicks. Our current findings on L-Asp concerning feed intake in broiler chicks is in accordance with other reports where it was shown that either peripheral or central administration of L-Asp did not influence feed intake in layer chicks (Maruyama *et al.*, 1972; Bungo *et al.*, 2002; Erwan *et al.*, 2013a). However, the effect of D-Asp on feed intake differed with our recent study (Erwan *et al.*, 2013a), since feed intake was significantly decreased when layer chicks were orally administrated with D-Asp. The reason for these discrepancies on feed intake behavior due to D-Asp is unknown. However, feed intake and growth rate is 1.5 to 2-fold greater in broilers than in layers at 2 or 5 days old (Masic *et al.*, 1974; Mahagna and Nir, 1996). It is well known that broiler chicks eat more food than layer-type chicks and then grow faster (NRC, 1994). Furuse *et al.* (2007) suggested that layers have higher levels of anorexigenic neuropeptides compared to broilers. Shibana *et al.* (2001) revealed that agouti-related protein (AGRP) stimulated feed intake in layer-type chicks under an ad libitum feeding condition but not broiler chicks and suggested that the orexigenic effect of AGRP is different between the two breeds possibly due to the higher expression of its antagonist α -melanocyte stimulating hormone in layers. Therefore, neuropeptide regulation of feed intake is different between broiler and layer type chicks. Although D-Asp dependent neuropeptide regulation for food intake is still unknown, we could hypothesize that D-Asp may influence the food intake regulatory circuitry in layers, but not broilers, due to having different food intake regulatory system in broilers and layers. Therefore, strain specific feeding regulation and variation in metabolic rate may cause the different responses for L- and D-Asp. Furthermore, it has been well

documented that L- or D-Asp has a function as neurotransmitters to influence the central nervous system to regulate feed intake in rats. L-Asp is not only concentrated in nerve endings (Gundersen *et al.*, 1991), but is also found localized and accumulated in common synaptic vesicles (Gundersen *et al.*, 1998; Fleck *et al.*, 2001). Thus, it could be further speculated that the administered dose of D-Asp might not have influenced physiological actions on neurocircuits in controlling feeding behavior in broiler chicks.

We also demonstrated here that L-Asp did not influence body temperature in broiler chicks. The result was accordance with the previous findings that oral administration of L-Asp did not decrease rectal temperature in rats (Koyuncuoğlu *et al.*, 1982a; Koyuncuoğlu and Berkman, 1982) and in layer chicks (Erwan *et al.*, 2014a). Interestingly, we have found that when L-Asp was chemically conjugated with lauric acid, a medium-chain fatty acid, then it enabled decreasing body temperature in layer chicks (Erwan *et al.*, 2013b). Thus, it could be suggested that the chemical structure of L-Asp is not designed to influence body temperature but esterified L-Asp may have that potential.

Previous research findings revealed that the functions of different enantiomers are increasing (Gong *et al.*, 2005; Yamane *et al.*, 2009; Yoshida *et al.*, 2015). It seems to be evident from the following reports that there were different responses due to the difference in enantiomers of L- or D-Asp. Koyuncuoğlu and Berkman (1982) revealed that in rats, the concomitant oral administration of L-Asp seemed to antagonize the effect of D-Asp. The injection of D-Asp but not L-Asp released arginine vasopressin (AVP) in rats (Koyuncuoğlu *et al.*, 1984). We have revealed the differences of L-Asp and D-Asp on stress response, feed intake and regulation of body temperature (Erwan *et al.*, 2012, 2013a, 2013b, 2014). Further it was found that the mechanism of stress attenuating function of L- and D-Asp possibly occurs through different receptors (Erwan *et al.*, 2012, 2014b). Koyuncuoğlu *et al.* (1982b) demonstrated that rectal temperature significantly decreased when rats received D-Asp or D- plus L-Asp in a 1:1 ratio. Since the decreased body temperature by D-amino acids was antagonized by naloxone, an opioid antagonist (Koyuncuoğlu *et al.*, 1982a), the decrease in rectal temperature might be caused by opioid

system. Previous reports demonstrated that the injection of D-Asp increased AVP in rats (Koyuncuoğlu *et al.*, 1984). The homologous nonapeptide AVP in birds is arginine vasotocin (AVT) (Acher *et al.*, 1970). Similarly, to the function in mammal, AVT, a neurohypophyseal hormone in non-mammalian vertebrates, is involved in water balance in birds (Stallone and Braun, 1986). However, plasma levels of AVT increased in hypertonic solution and this response was further enhanced by naloxone injection as the doses increased (Saito *et al.*, 1999). Furthermore, i.c.v. injection of AVT increased body temperature (Tachibana *et al.*, 2004). These facts suggest that decreased body temperature induced by orally administered D-Asp could not be explained by increased AVT through the opioid system. Further investigations are needed to determine the relationships between D-Asp and possible factors to regulate body temperature. This finding raised the possibility that D-Asp may be a useful hypothermic agent at HT not only for layer chicks but also for broiler chicks. Importantly, D-Asp did not reduce feed intake but caused to reduce rectal temperature in broiler chicks, which clearly indicate that the body temperature reduction by D-Asp may be due to its specific action on central or peripheral thermoregulatory mechanisms in broiler chicks. We assume that D-Asp dependent declining in rectal temperature may not be connected with metabolic function related regulation in body temperature since D-Asp reduced body temperature both in layer and broiler chicks. As the brain is the center of thermoregulation (Yahav, 2015), D-Asp may have some function to control the thermal center in the brain. D-Asp may have some direct influence on this thermal center or indirect influence through peripheral sensory nervous system.

In Experiment 1, it was found that L-Asp did not influence any plasma metabolites studied. This finding confirmed our previous report in layer chicks (Erwan *et al.*, 2014a). In addition, Tada *et al.* (2008) demonstrated that a subchronic oral administration of L-Asp with a dietary concentration of up to 5.0% had no influence on serum glucose in rats. Moreover, Delaney *et al.* (2008) and Karaman *et al.* (2011) also revealed that oral administration of L-Asp or N-acetyl-L-aspartic acid, an N-acetylated derivative of L-Asp, had no effect on TG, glucose, or TP. Thus, the present data corroborate those reported by others indicate that L-Asp or its derivative may not change the plasma metabolites in chicks especially in short term administration.

Consistent with previous research (Erwan *et al.*, 2014a), plasma TG was found to be significantly decreased with the oral administration of D-Asp in the current study. Studies with rat (Koyuncuoğlu *et al.*, 1982b) and dogs (Shida *et al.*, 1977) clearly showed the effect of decreasing plasma TG resulted hypothermia. It was further confirmed in this study that oral administration of D-Asp increased glucose (Table 2) as observed in layer chicks (Erwan *et al.*, 2014a). A positive correlation between the dose of D-Asp and glucose suggest that D-Asp might have increased plasma glucose. These findings are consistent with earlier report that revealed that hypothermia resulted in increasing glucose (Shida *et al.*,

1977). In addition, Marley and Stephenson (1975) revealed that hypothermia was associated with increased blood glucose concentration in chicks. Therefore, the present results indicate that D-Asp dependent hypothermia may cause to increase energy demand in broiler chicks and thereby increase plasma glucose and decrease plasma TG.

In conclusion, oral administration of D-Asp, but not L-Asp, could cause hypothermia in broiler chicks without affecting feed intake and altered some plasma metabolites – namely, TG, and glucose. These results suggest the possibility of D-Asp as a potential hypothermic agent in chickens to mitigate heat-stress problems.

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