Stress and Chicken Broiler Meat Quality.

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MARY: Effects of ante-mortem stress, chemically generated by subcutaneous injections of epinephrine, on post-mortem broiler muscle WARY: Effects of ante-mortem stress, chemically generated by subcutaneous injections of opinopulation, and subsequent meat quality were studied. Also the effects of long and short term natural stress on meat quality characteristics have ⁿ evaluated.

Blood lactate and glucose levels have been measured as well as tissue lactate, glycogen and glucose at time of slaughter and 24 hours post $h_{e_{III}}$. The color (e.g. L* and a* values) of the raw meat has been estimated.

 $h_{jections}^{jections}$ of epinephrine resulted in higher terminal muscle pH in birds killed 4 to 16 hours post injection. Also these birds showed ificantly darker breast meat (lower L* values). ^{wy darker} breast meat (lower L* values). ^{Inthe long} and short term stress experiment indicated only some minor effects on meat quality characteristics.

NIRODUCTION: Variation in poultry meat quality is partly affected by processes like stunning, scalding and picking, duration of the Way of cutting up of the birds, etc. An important part of poultry meat quality however is already determined at the moment of stunning killing. The negative effects of ante mortem stress on poultry meat quality are neither well documented, nor understood or consistent.

Simpson and Goodwin (1975), Lee et al. (1976) and Froning et al. (1978) found some negative effects on meat quality caused by heat Also $A_{1,A_{1,S_{0}}}$ pre slaughter free struggle has shown to affect meat quality negatively (Kahn en Nakamura (1970), Ma *et al.* (1971), Grey *et al.* η_{4} , Froning et al. (1978) and Ngoka et al. (1982).

Froning et al. (1978) and Ngoka et al. (1982). Rener et al., (1978), Ngoka et al., (1982) and Ngoka and Froning, (1982) showed an effect of stress caused by heat, free struggle and $W_{0,q}$ and et al., (1978), Ngoka *et al.*, (1978), Ngoka *et al.*, (1978), W_{0,q} when the occurrence of darker turkey meat color.

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kahn and Nakamura (1970) concluded that minimizing postmortem glycolysis by epinephrine injection 5 hours prior to slaughter resulted higher up the total total provided that minimizing postmortem glycolysis by epinepinnie injection of total provided in increased with the most tender meat. Woods and Richards (1975) found that epinephrine injections resulted in increased in the total provided to the second provided to sele toughness.

 h_{e}^{work} attempting to relate antemortem stress to poultry meat quality is often contradictory and incomplete. Much of the contradiction $b_{e} d_{ue}$ h_{be} h_{be} h

The purpose of this study was to compare the physiological effects of epinephrine injections to subsequent meat quality attributes. Purpose of this study was to compare the physiological effects of epinephrine injections to subsequence a maximum stress ^{Na} may be ^{al} may be obtained.

A second part of the stress study was to introduce natural stress and to compare the physiological and meat quality data with the results h_{θ} chere. he chemical induced stress study. In this study only results of the experiments to estimate chemical induced stress will be presented.

MATERIALS AND METHODS: Broilers chickens used in these experiments were reared at Spelderholt Centre. The trials were executed WERIALS AND METHODS: Broilers chickens used in these experiments were reared at Spendemon Centre. All birds of a trial to the experimental set up given in table 1. Epinephrine was injected subcutaneously on the inner side of the thigh. All birds of a trial the injected at the same time.

TRIAL	NUMBER of birds	AGE (days)	EPINEPHRINE INJECTION (mg/kg live weight)	STUNNING / KILLING TIME POST INJECTION (hours)
1	105	57	0.33	0, 0.5, 1, 2, 4, 8, 12, 16, 24, 30 and 48
2	105	64	4.0	0, 3, 6, 9 and 12
3	105	39	0.83	0, 0.5, 1, 2, 4, 8, 12, 16, 24, 30 and 48
4	105	46	1.5	0, 0.5, 1, 2, 4, 8, 12, 16, 24, 30 and 48

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Table 1. Experimental set up of the experiments.

At the moment of slaughtering 8 birds were electrically stunned and killed. Blood samples were taken during bleeding of the birds for any samples were taken during bleeding of the birds and the birds and the same samples were taken during bleeding of the birds and the birds and the birds are taken during bleeding of estimation of glucose, lactate, triiodothyronin (T-3) and thyroxin (T-4). Immediately after killing from 4 out of 8 birds breast meat was same to measure initial pH, initial tissue lactate and initial tissue actate actate actate and initial tissue actate acta to measure initial pH, initial tissue lactate and initial tissue glycogen. One breast half of the other birds was used to measure ultimate tissue lactate and ultimate tissue glycogen

The other breast half from all 8 birds was used to measure Warner-Bratzler shear force and color.

Ph was measured according to the method described by Jeacocke (1977).

L* and a* color values were measured with the "Minolta Chroma meter II" Reflectance Colorimeter with a 8 mm measuring head

Shear was measured using the Warner-Bratzler shear and an Overload Dynamics testing machine. The sampling and measuring method has be

Glucose, glycogen and lactate were measured according to modified methods, based upon the enzymatic Boehringer Mannheim methods

T-3/T-4 were measured with kits supplied by Diagnostic Products Corporation, Los Angeles, Ca, Usa

RESULTS AND DISCUSSION: The results of the meat quality determinations and muscle- and blood analyses are given in the figure 12. 1 to 12.

Figure 1 and 2 show the initial and terminal muscle pH of the epinephrine treated birds. The values found initially do not show an effect of the treated birds. The values found initially do not show an effect of the treated birds. significant differences. Terminal muscle pH significantly shows an effect of the treatment of the broilers with epinephrine. Birds slaughter 4 to 12 hours after epinephrine administration show a increase in terminal muscle and terminal muscle pH significantly shows an effect of the treatment of the broilers with epinephrine. Birds slaughter this is 4 to 12 hours after epinephrine administration show a increase in terminal muscle pH. The bird recovers after about 16 hours. This is agreement with the data given in figure 10 (initial tissue glycogen) and 12 (initial tissue glyco agreement with the data given in figure 10 (initial tissue glycogen) and 12 (terminal tissue lactate). Epinephrine administration causes and the terminal lactate concentration will be low. Under the second decrease in tissue glycogen and the terminal lactate concentration will be low. Under these circumstances the terminal pH will be relatively high information and terminal pH will be relatively high with terminal pH will be relatively high with terminal p Terminal tissue glycogen was near 0 mg/kg for all the treated groups indicating that all the available glycogen has been transferred. A high initial glycogen content obviously results in a high lactate content in the muscle and thus in the solution of the solution of the solution.

Meat quality characteristics are also affected by epinephrine administration. Especially the L* color value shows a decrease when brilling (figure a) are slaughtered 4 to 12 hours after epinephrine administration. Under these circumstances the meat is darker. The a* color value $(figure^{4})$ not significantly affected by the epinephrine administration. The Warren Device the meat is darker. The a* color value $(figure^{4})$ not significantly affected by the epinephrine administration. Under these circumstances the meat is darker. The a* color value (figure in value) to epinephrine administration in the period from 4 to 12 hours post morter. The effected is the show a decrease in value of the show a decrease in value (figure in value) and the show a decrease in value (figure in val

The blood values for glucose, lactate, T-3 and T-4 are shown in figure 6 to 9. Blood glucose (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the blood lactate values (figure 6) shows a significant increase in the bl birds killed between 0.5 and 8 hours after the injection. The blood lactate values (figure 7) are not affected. Only in trial 2 a very high after the injection of the blood lactate values (figure 7) are not affected. Only in trial 2 a very high after the injection of the blood lactate values (figure 7) are not affected. for blood lactate and also for initial tissue lactate (figure 11) has been monitored. In this trial the mortality during the period direct difference of the p Due to the increased metabolic rate in the animal caused by the administration of epinephrine the T-3 concentration in the blood we ted (figure 8). T-4 values (figure 9) were not. Further studies are required to evolve the terms of the terms of the blood we have the terms of terms of the terms of the terms of terms of the terms of terms o

affected (figure 8). T-4 values (figure 9) were not. Further studies are required to explain these effects.

The results of these study have been the starting point for further study on the effects of natural stress on meat quality characteristics¹⁰⁹ metabolites in muscle and blood from stressed broiler chickens. Results of this study will be published in the near future (Fletcher, et al, 1991)

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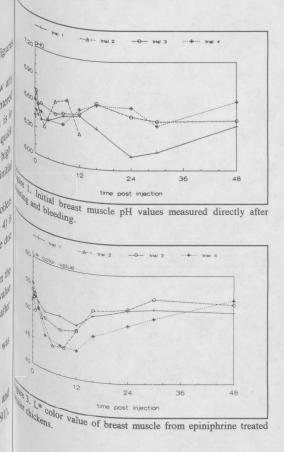
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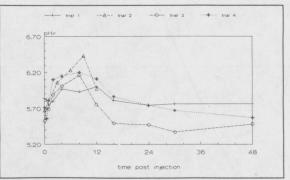


Figure 2. Terminal breast muscle pH.

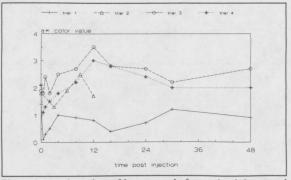


Figure 4. a* color values of breast muscle from epinephrine treated broiler chickens.



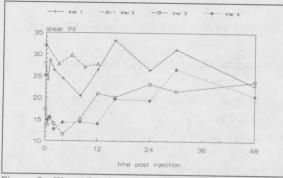


Figure 5. Warner-Bratzler shear forces of breast meat from epinephrine treated broiler chickens.

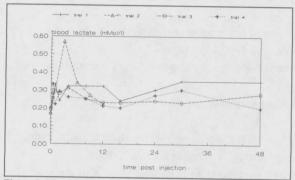


Figure 7. Blood lactate content in samples obtained during bleeding.

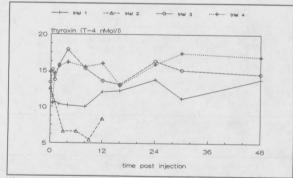


Figure 9. T-4 contents of blood samples obtained during bleeding

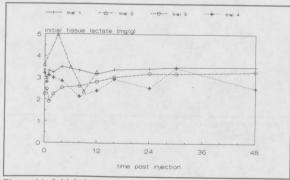


Figure 11. Initial tissue lactate in epinephrine treated broiler breast muscle.

