

THE CLINICAL BIOCHEMISTRY OF CHICKS  
WITH AFLATOXICOSIS: SOME EFFECTS  
OF SUPPLEMENTARY CHOLINE,  
FOLATE, THREONINE, LYSINE  
AND LYSINE PLUS ARGinine

CENTRE FOR NEWFOUNDLAND STUDIES

TOTAL OF 10 PAGES ONLY  
MAY BE XEROXED

(Without Author's Permission)

LAURA PARK







THE CLINICAL BIOCHEMISTRY OF CHICKS WITH AFLATOXICOSIS:  
SOME EFFECTS OF SUPPLEMENTARY CHOLINE, FOLATE, THREONINE,  
LYSINE AND LYSINE PLUS ARGinine

A Thesis

by

Laura Park

A Thesis Submitted in partial fulfillment  
of the requirements for the degree of  
Master of Science

Department of Biochemistry  
Memorial University of Newfoundland

Date: December 1983

St. John's

Newfoundland

Permission has been granted to the National Library of Canada to microfilm this thesis and to lend or sell copies of the film.

The author (copyright owner) has reserved other publication rights, and neither the thesis nor extensive extracts from it may be printed or otherwise reproduced without his/her written permission.

L'autorisation a été accordée à la Bibliothèque nationale du Canada de microfilmer cette thèse et de prêter ou de vendre des exemplaires du film.

L'auteur (titulaire du droit d'auteur) se réserve les autres droits de publication; ni la thèse ni de longs extraits de celle-ci ne doivent être imprimés ou autrement reproduits sans son autorisation écrite.

ISBN 0-315-31034-0

FOR MY MOTHER

who might have got an MA instead of ME

(ii)

#### ABSTRACT

Aflatoxin (2.5ug/g diet) was fed to broiler chicks for 24-26 days in five separate feeding trials in which the effects of supplementary choline, folate, threonine, lysine, and lysine plus arginine were examined. Weight gain, feed intake, feed conversion and hepatic lipid responded in a manner typical for aflatoxicosis. Plasma concentrations of LDH, taurine, tyrosine, phenylalanine, arginine, ornithine, citrulline, glutamine, ammonia and perhaps BUN were increased in response to aflatoxin, while plasma levels of threonine, lysine, total protein, albumin/globulin (A/G) ratio, uric acid, cholesterol, calcium, inorganic phosphate, total iron, total iron binding capacity, and percent saturated transferrin, decreased.

Intraperitoneal (IP) administration of choline, but not dietary choline, moderated the influence of aflatoxin on the majority of the biochemical parameters.

Lysine supplementation improved the performance of chicks with aflatoxicosis, while threonine had a negative effect. This may be related to ornithine detoxification of aflatoxin through the opposing effects of these two amino acids on the activity of the enzyme arginase, which catalyzes the conversion of arginine to ornithine and urea.

Plasma lysine concentration varies considerably as a result of genetic differences in lysine metabolism. Data indicates that

(iii)

chicks with high plasma lysine concentration are more resistant to aflatoxicosis than chicks with low plasma concentrations of lysine.

(iy)

#### ACKNOWLEDGEMENTS

I wish to thank my supervisor, Dr. Michael Voigt for his support, encouragement and patient assistance throughout this study. I also thank Doug Hall for the amino acid analysis, and also Alvine Mills and Vernon Whelan for the completion of biochemical analyses on plasma.

I am very grateful to Drs. Ronald Payne, William Davidson, Ian Fraser, and Graham Allaway for their advice and assistance in the electrophoresis, and analysis of lactate dehydrogenase.

Special thanks to Tor Gjesdal for his invaluable assistance in the completion of figures.

I am also grateful to Dr. Norman Haard, Kofi Simpson, Dr. Kasi Shamsuzzaman, Zuzer Ali Shamsuddin, Teik-Mien Tye, Donna Jackman and Peter Reese, for their assistance and moral support without which the work for this thesis would have been much less enjoyable.

Sincere thanks to the Medical Research Grant (MA-7355) and the Memorial University Graduate Fellowship Program for financial assistance.

## TABLE OF CONTENTS

<u>CHAPTER I</u>	page
<u>INTRODUCTION</u> .....	1
Toxin producing moulds .....	2
Feed survey .....	4
History of aflatoxin induced disease .....	6
Structure and metabolism of aflatoxin .....	6
<u>CHAPTER II</u>	
<u>MATERIALS AND METHODS</u> .....	10
Feed collection and storage .....	11
Mycotoxin extraction .....	11
Thin layer chromatography .....	12
Aflatoxin production .....	13
Animal husbandry and necropsy .....	14
Animal performance and biochemical analyses .....	18
Statistical analyses .....	20
<u>CHAPTER III</u>	
<u>EFFECTS OF AFLATOXIN</u> .....	21
Performance .....	22
Hepatic changes .....	26
Clinical parameters .....	29
Plasma free-amino acids .....	33

CHAPTER IV

EFFECT OF CHOLINE AND FOLATE SUPPLEMENTATION.....	38
Introduction.....	39
Effects of choline on performance.....	41
Effect of choline on hepatic lipid, and on selected biochemical constituents in plasma .....	43
Effects of folate .....	47

CHAPTER V

EFFECTS OF LYSINE AND THREONINE SUPPLEMENTATION.....	51
Introduction.....	52
Contrasting effects of threonine and lysine on performance.	56
Lysine/arginine ratio.....	56
Contrasting effects of threonine and lysine on selected amino acids.....	63

CHAPTER VI

EFFECT OF LYSINE/ARGININE SUPPLEMENTATION.....	66
Introduction.....	67
Relationship between plasma lysine/arginine ratio and performance.	69

CHAPTER VII

SUMMARY AND CONCLUSIONS.....	74
------------------------------	----

REFERENCES.....	77
-----------------	----

APPENDIX A: RAW DATA TABLES 1-25.....	86
---------------------------------------	----

## LIST OF TABLES

TABLE		Page
I.	Composition of basal ration.....	15
II.	Content of amino acids in the basal diets for the choline, folate, lysine, threonine and lysine + arginine feeding trials.....	16
III.	Effect of aflatoxin, on LDH activities in various tissues.....	30
IV.	Effect of dietary and IP choline supplements on the performance of chicks with aflatoxicosis.....	42
V.	Effect of dietary and IP choline on hepatic lipid content.....	44
VI.	Effect of dietary and IP choline on the percent change of selected plasma constituents between the chicks receiving aflatoxin and the pair fed controls.....	45
VII.	Effect of dietary and IP choline on the percent change of selected plasma amino acids between the chicks receiving aflatoxin and the pair fed controls.....	46
VIII.	Effect of supplemental folate on the performance of chicks with aflatoxicosis.....	48
IX.	Effect of supplemental folate on hepatic lipid.....	49
X.	Percent change of selected plasma amino acids in chicks receiving a 164% increase in dietary folate... .	50
XI.	Effect of graded levels of lysine or threonine on the final weight of chicks with aflatoxicosis.....	57
XII.	Effect of graded levels of lysine or threonine on feed conversion of chicks with aflatoxicosis .....	58
XIII.	Effect of graded levels of lysine or threonine on feed intake of chicks with aflatoxicosis.....	59
XIV.	Effect of lysine and threonine supplements on plasma levels of selected amino acids in chicks with aflatoxicosis.....	64

(viii)

- XV. Innate differences in the plasma concentrations of lysine, arginine, and lysine/arginine ratio in feeding trials 4 and 5..... 68
- XVI. Effect of graded levels of lysine and arginine on feed intake of chicks receiving aflatoxin:..... 70.
- XVII. Effect of graded levels of lysine and arginine on weight gain of chicks receiving aflatoxin..... 71

(A)

LIST OF FIGURES

FIGURE	Page
1. Structures of aflatoxin B1, B2, G1, and G2.....	8
2. Metabolism of aflatoxin B1.....	9
3. Effect of aflatoxin on the performance of chicks....	23
4. Daily weight gain of chicks with aflatoxicosis.....	24
5. Daily feed consumption of chicks with aflatoxicosis.....	25
6. Effect of aflatoxicosis on hepatic weight, percent moisture, and percent lipid.....	27
7. Effect of aflatoxicosis on the plasma levels of selected hepatic enzymes.....	28
8. Comparison of LDH isozyme patterns from various tissues.....	31
9. Effect of aflatoxicosis on the clinical biochemistry of chicks.....	32
10. Effect of aflatoxicosis on plasma concentrations of selected amino acids.....	35
11. Role of ornithine in the formation of physiological nucleophiles.....	54
12. Effect of graded levels of threonine on plasma lysine/arginine ratio of chicks with aflatoxicosis.....	61
13. Effect of graded levels of lysine on plasma lysine/arginine ratio of chicks with aflatoxicosis.....	62
14. Relationship between plasma lysine concentration and aflatoxin toxicity.....	73

(x)

ABBREVIATIONS

AAA.....	aromatic amino acid
A/G.....	albumin/globulin ratio
Ala.....	alanine
Alb.....	albumin
ALT.....	alanine aminotransferase
AMN.....	ammonia
AP.....	alkaline phosphatase
App.....	appendix
Asn.....	asparagine
Asp.....	aspartic acid
AST.....	aspartate aminotransferase
Arg.....	arginine
BCAA.....	branched chain amino acid
BUN.....	blood urea nitrogen
C.....	celsius
Ca.....	calcium
Chol.....	cholesterol
Cit.....	citrulline
Cth.....	cystathionine
Cys.....	cysteine
dwb.....	dry weight basis
Eth.....	ethanolamine
Fig.....	figure

g.....	gram
GAL.....	galactosyltransferase
Gln.....	glutamine
GPr.....	globulin protein
Glu.....	glutamic acid
Gluc.....	glucose
Gly.....	glycine
Hcy.....	homocysteine
His.....	histidine
HOL.....	hydroxylysine
HOP.....	hydroxyproline
HPLC.....	high pressure liquid chromatography
Ile.....	isoleucine
IP.....	intraperitoneal
LDH.....	lactate dehydrogenase
Leu.....	leucine
Lys.....	lysine
Met.....	methionine
NSP.....	ninhydrin positive substance
NRC.....	national research council
Orn.....	ornithine
P.....	inorganic phosphate
PCV.....	packed cell volume
Phe.....	phenylalanine
Pro.....	proline

(xii)

SAL.....	sialyltransferase
Ser.....	serine
SFe.....	percent saturated transferin
Tau.....	taurine
TB.....	total bilirubin
TFe.....	total iron
TIBC.....	total iron binding capacity
Thr.....	threonine
TLC.....	thin layer chromatography
TPr.....	total protein
Tri.....	triglyceride
Trp.....	tryptophan
Tyr.....	tyrosine
U.....	unit
UA.....	uric acid
Val.....	valine

CHAPTER I  
INTRODUCTION

In this chapter, general aspects of toxin producing moulds are reviewed. An historical account of aflatoxicosis is included, with a discussion of the chemical structure of aflatoxin and current knowledge of aflatoxin metabolism. The preliminary work which led to the final choice of this thesis topic is also presented. Additional reviews which appear before chapters III, VI, and V contain more detailed information of particular relevance to the subject matter of the chapter in question.

#### Toxin producing moulds

Mycotoxins are poisonous substances produced in a substrate as the result of the growth and metabolism of moulds. Production of these toxic metabolites is dependent on the presence of a toxin producing mould in combination with the appropriate temperature, oxygen concentration, moisture level, and substrate. Optimal conditions for mould growth are not necessarily optimal conditions for toxin production (1). It is estimated that 30-40% of all moulds are capable of producing mycotoxins under certain conditions. Crops may be infested by moulds prior to harvest, usually after a plant's natural defense mechanisms have been weakened by insect infestation or harsh weather conditions. Fungal infections also occur post-harvest, and are a result of inadequate dehydration or improper storage and handling. Control of moisture and temperature are particularly critical in this respect (1). A given toxin may be produced by a single or many different mould species and the different mycotoxins vary considerably in chemical form and properties. Although the toxin producing moulds

are not restricted to any single group of moulds, the genera in which they occur most frequently are Aspergillus, Penicillium, and Fusarium. A group of closely related metabolites produced by strains of Aspergillus flavus and Aspergillus parasiticus are known collectively as aflatoxin. These fungi are ubiquitous, and thus the potential for the outgrowth of aflatoxin producing fungi in foodstuffs and animal feeds is widespread. Corn and peanuts are particularly susceptible to aflatoxin contamination and both are major ingredients of animal feeds. Aflatoxin is not only the most prevalent, but also the most toxic and carcinogenic mycotoxin known. Aflatoxin contamination of grains is normally associated with warm climates although the lower limiting temperature for growth of A. flavus is 12C (1). Hence, Newfoundland is not generally thought of as a region in which aflatoxin contamination should be significant. However since all of the grain utilized in Newfoundland is imported, the environmental conditions encountered during transportation and storage, as well as the quality of the original product may be more relevant to the levels of aflatoxin or other mycotoxins in Newfoundland feeds. A snowballing effect has been demonstrated in which there is an accumulative increase in the content of mycotoxins with each step of their journey from the field to the feed trough (grain elevator, truck, shipping car, warehouse, barn loft etc.) (2). This suggests that the greater the geographical separation between the sites of production and consumption of a given product, the greater the expectation of mycotoxin contamination. Even if a product contains

negligible concentrations of a toxin at the site of production, the levels may have increased to biologically important concentrations by the time the feed is consumed in Newfoundland. In addition there is an absence of routine testing of feeds for aflatoxin or other mycotoxins in Newfoundland, and the symptoms of aflatoxicosis are non-specific and thus not easily diagnosed. Therefore, Newfoundland appears to be a more likely target for aflatoxicosis than might be predicted at first glance.

#### Feed survey

Before choosing a mycotoxin to study intensively through feeding trials, a preliminary survey of mycotoxins occurring in animal feeds in Newfoundland was undertaken. One hundred and eleven samples of feed including rations for poultry, cattle, and swine from across the province were analyzed for the presence of fourteen different mycotoxins (zearalenone, sterigmatocystin, roridin A, T-2 toxin, penicillic acid, patulin, diacetoxyscirpenol, verrucarin A, ochratoxin A, citrinin, and the four aflatoxins B-1, B-2, G-1, and G-2). Thirty-seven milk samples were also analyzed for M-1, a metabolite of aflatoxin B-1 that occurs in the milk of animals fed aflatoxin contaminated feed. Tentative positives for aflatoxin B-1 (often in combination with B-2 and G-1) were obtained from forty percent of the feed samples tested in the initial screening process, but none of these samples were consistently positive throughout all of the subsequent confirmatory tests. Only four of these samples were associated with adverse performance (decreased egg production, feed

refusal) which could be attributable to aflatoxicosis. Of the thirty-seven milk samples tested one positive M-1 was obtained. Tentative positives for patulin, verrucarin A, and penicilllic acid were obtained in 6%, 8%, and 17% of the feed samples respectively, while less than 5% of the samples showed tentative positives for T-2, zearalenone, sterigmatocytin, ochratoxin and citrinin. Roridin A and diacetoxyscirpenol were negative in all samples. Positive identification of these toxins by comparison of sample extracts with mycotoxin standards subjected to thin layer chromatography was difficult because of the large number of standards to be run with each sample. In addition, the large number of different components in the feed extracts (due to the broad specificity of the extraction procedure) influenced the colour and Rf of different components of the extract on the chromatograms, making identification by comparison with individual standards unreliable. Towards the end of this survey it became apparent that the TLC methods of screening for multiple mycotoxins are inadequate to permit the confirmation of specific mycotoxins. For large scale surveys, TLC methods have been rapidly replaced by computerized HPLC systems which can store the HPLC profile of mycotoxin standards for future reference. Using this system, sample extracts are simply injected into the HPLC unit, and all mycotoxins present are identified and quantitated automatically. Because the object of the project sponsored by MRC was to determine the influence of diet on avian mycotoxicosis, an expensive change in the methodology of the survey could not be justified, and the research was redirected.

toward feeding studies utilizing aflatoxin and poultry.

#### History of aflatoxin induced disease

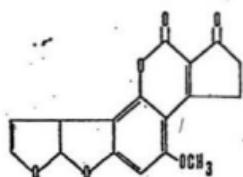
Aflatoxicosis received world wide attention in 1960 when A. flavus contaminated peanut meal was implicated as the cause of the mysterious 'turkey X disease' which was responsible for the deaths of at least 6,000 chicks and 12,000 ducklings in addition to 100,000 turkeys in the United Kingdom. The disease was characterized by sudden loss of appetite, subcutaneous haemorrhages and a high and rapid mortality in young birds. At postmortem the livers were pale, fatty, and showed extensive biliary proliferation. Partridge and pheasant poult together with cattle, pigs and sheep were also affected. Almost simultaneously, many trout bred in commercial hatcheries in the United States developed hepatomas and this outbreak was eventually attributed to the same fungal toxin (3). Since 1960, numerous studies have been conducted to determine the chemical nature of these A. flavus toxins and the clinical problems encountered as a result of their consumption. Four major chemically related aflatoxins designated B1, G1, B2, and G2 commonly occur together in contaminated feeds and feedstuffs. The proportions in which these metabolites are produced may vary considerably depending on environmental conditions, particularly temperature. Aflatoxin produced in this laboratory consisted of 88% B1, but the proportion of G1 is reported to be increased substantially at lower temperatures (1).

#### Structure and metabolism of aflatoxin

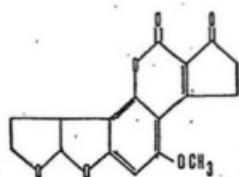
The aflatoxins fluoresce strongly in ultraviolet light (ca

365nm); B1 and B2 produce a blue fluorescence whereas G1 and G2 produce a green fluorescence. The structures of these compounds are shown in Fig.1. Because aflatoxin B1 is the most prevalent as well as the most potent of the aflatoxins, the metabolites of aflatoxin B1 have been studied most intensively, and will be reviewed below. B2, G1, and G2 undergo most of the same reactions to produce the corresponding B2, G1, or G2 derivative (3). In general the detoxification of xenobiotics by the liver results in the formation of more polar products which may or may not be subsequently conjugated with amino acids, glucuronic acids, sulfate or bile acids as an aid to their excretion. The aflatoxin molecule lends itself to biodegradation in at least 6 ways (3), as shown in Fig. 2. Reactions involving the intact molecule have been confirmed by the isolation of the metabolites which are cited as examples.

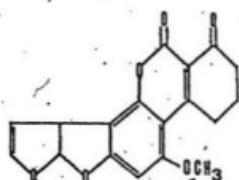
The predominant biotransformations vary between animals of different species, a fact which partially explains the wide variation in species susceptibility to aflatoxicosis. A number of dietary factors have been found to influence the metabolism, and thus the toxicity of a given dose of aflatoxin. In the following chapters, the influence of specific dietary factors that are relevant to the corresponding nutritional study will be discussed in more detail. The feeding trials examining the effects of supplemental choline, folate, threonine, lysine, and lysine plus arginine on chicks with aflatoxicosis and are referred to as feeding trials 1, 2, 3, 4, and 5, respectively.



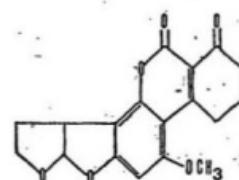
AFLATOXIN B1



AFLATOXIN B2



AFLATOXIN G1

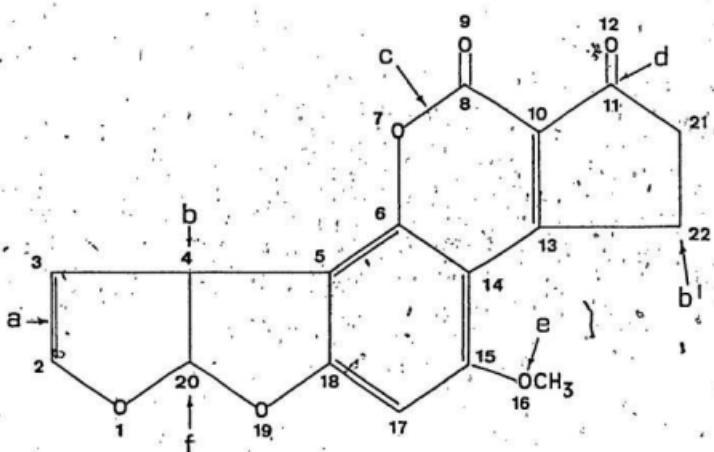


AFLATOXIN G2

Fig. 1. Structures of Aflatoxin B1, B2, G1, and G2.

Fig. 2. Metabolism of Aflatoxin Bl.

- a) Reductive or hydrolytic attack on the 2,3, double bond. Examples include the 2,3-dihydroxy, the 2-hydroxy (also known as B2a or hemiacetal) and 2,3-epoxide derivatives. The 2,3-epoxide is assumed to be an important reactive intermediate and has never been isolated although there is good evidence of its formation.
- b, b') Hydroxylation at one or more points in the molecule. Both aflatoxin- M1 and Q1 are examples of hydroxylated derivatives (4-hydroxy and 22-hydroxy respectively).
- c) Hydrolytic fission of the coumarin lactone.
- d) Cyclopentenone reduction. Aflatoxin Ro, also referred to as aflatoxicol, results from reduction of the 11,12 double bond.
- e) O-demethylation of the methoxy-coumarin structure. Aflatoxin P1 refers to the phenolic compound derived in this manner.
- f) Opening of the bisfuranoid structure.



CHAPTER II  
MATERIALS AND METHODS

#### Feed collection and storage

Farmers and feed manufacturers in Newfoundland were requested to submit samples of feedstuffs for analysis of mycotoxins from June 1979 to August 1980. Initial contact was made by telephone, from a list provided by Dr. A. Smith (provincial veterinarian). Packages containing an explanatory letter, sample submission forms, plastic bags for sample submission and return address labels with guaranteed postage were mailed to all individuals who responded favourable to the initial telephone communication. The respondents included three feed manufacturers and 144 farmers involved with swine, cattle, sheep, horses, and poultry. The regular submission of unbiased samples was requested in addition to bias samples. Bias samples were defined as feeds associated with feed refusal or undesirable animal performance and/or feed having a mouldy or otherwise unusual appearance or odour. A total of 111 feed samples were submitted over the entire 14 month period, and 35% of these were bias samples. Milk samples were obtained from five separate commercial dairies and included samples of raw, whole, 1%, 2%, skim, and chocolate milk. Milk was obtained over a period of 3 months starting in November 1979. All feed and milk samples were stored in sealed bags or bottles at 30C.

#### Mycotoxin extraction

Feed samples were extracted by the procedure for the

extraction of mycotoxins described by Patterson and Roberts (4). This procedure produced two separate extracts from each feed sample, one containing basic mycotoxins (ochratoxin, and citrinin) and a second containing acidic mycotoxins (aflatoxin, zearalenone, roridin A, T-2 toxin, penicilllic acid, patulin, verrucarin A, sterigmatocystin, and diacetoxyscirpenol).

Milk samples were extracted by the aflatoxin M<sub>1</sub> procedure of Stubblefield (5).

Thin layer chromatography

Aflatoxin M<sub>1</sub> extracts of the milk samples were evaporated to dryness under nitrogen and redissolved in 100ul chloroform. Twenty ul aliquots were developed on one dimensional precoated silica gel TLC plates (Macherey-Nagel, Germany; SIL G-25HR) in isopropanol/acetone/chloroform (2:10:88) and visualized under longwave UV light. Positive samples were confirmed using two dimensional TLC with the multiple solvent systems recommended for aflatoxin by Patterson and Roberts (4).

All feed extracts were initially screened on one dimensional silica gel TLC plates. An aliquot containing the various reference standards was also applied to each plate. Extracts were dried under nitrogen and redissolved in 100ul chloroform. Twenty ul of the extract containing the basic mycotoxins were developed in toluene/ethyl acetate/90% formic acid (60:30:10) and visualized under

longwave UV light. Twenty  $\mu$ l of the extract containing acidic mycotoxins were developed in chloroform/acetone (90:10) and followed by the visualization treatments recommended by Patterson and Roberts (4) for the general screen. Two dimensional TLC, combined with chromatogenic derivative formation were used to confirm the presence of specific mycotoxins in a sample extract, using the solvent system and visualization sequence recommended by Patterson and Roberts (4) for the specific toxin.

#### Aflatoxin production

Aflatoxin used in this study was produced by Aspergillus parasiticus NRRL 2999 on sterile polished rice by the method of Shotwell et al. (6) as modified by West, et al. (7). However, to prevent adhesion of the rice, the sterile rice and water were combined (2:1, respectively) at 100C and the mixture was allowed to steam under reduced heat until the water was absorbed (ca 20 min.). After cooling, the rice was aseptically transferred into 250 ml Erleameyer flasks, inoculated and incubated under the conditions described by West et al. The mouldy rice was heated at 100C for 3 min. to inactivate the fungus and then dried and ground to a fine powder using a ballmill. The powder was analyzed for total aflatoxin content by the spectrophotometric method of Nabney and Nesbitt (8) as modified by Wiseman et al. (9). The rice powder contained aflatoxin in the following ratio: B1:G1:B2:G2 (88:9:2:1). Rice powder was added to the

basal ration to obtain 2.5ug aflatoxin /g of feed. Rice powder added to the feed never exceeded 0.2% of the total diet.

Animal husbandry and necropsy

Day-old chicks (cockerels, Hubbard/Hubbard) were reared in wire-floored brooder batteries (31cm X 50cm X 23cm) with constant illumination at a light intensity of 56 footcandles. Water was available from separate containers for each cage of chicks from the day of hatch until termination of the experiment. The lysine plus arginine feeding trial terminated after 26<sup>a</sup> days while the other feeding trials concluded after 24 days. The basal ration consisted of a commercial broiler starter ration free of all medications. The formulation and amino acid composition of the rations are listed in Tables I and II, respectively. Four separate 3x3 factorial experiments were completed in which the influence of added supplements of choline, folate, threonine or lysine on aflatoxicosis in chicks was examined. The particular levels were chosen to provide supplementation substantially above the NRC requirement for broiler chicks and that provided by common premixes used by industry (Hoffman-LaRoche) and universities (UGA), yet with care to avoid levels which could produce toxicity. The levels of threonine, lysine and arginine supplements were also chosen with reference to information provided in the reports of Austic (79,80), which discuss the effects of supplementation of these amino acids to broiler chicks at a variety of levels and

<sup>a</sup> Termination was delayed in this feeding trial because freeze clamping equipment required for liver preparation by a co-worker for a separate project was not available on day 24.

Table I. Composition of the basal ration.

Ingredient	Composition <sup>a</sup>
Ground yellow corn	57.46
Soybean oil meal (dehulled)	30.84
Poultry by-product meal	4.9
Fat (vegetable)	2.98
Ground limestone	1.15
Dicalcium phosphate	1.74
Salt	0.04
Methionine (DL)	0.15
Trace mineral mix <sup>b</sup>	0.05
Vitamin premix <sup>c</sup>	0.25
Protein	23.0
Metabolizable energy <sup>d</sup>	3.120 MJ/Kg
Calcium <sup>d</sup>	1.09
Phosphorus (available) <sup>d</sup>	0.51

<sup>a</sup> Percentage unless indicated otherwise.

<sup>b</sup> Supersweet Feeds Mineral Premix (International Multifoods Corp., Minneapolis, Minnesota) provided the following minerals (mg/kg of diet): manganese 55, zinc 80, iron 80, copper 11, iodine 0.38.

<sup>c</sup> Vitamin premix provided the following amounts of vitamins per kg of diet: vitamin A, 4,400 IU; vitamin D<sub>3</sub>, 880 ICU (International chicken Units, 0.025 ug of cholecalciferol = 1 ICU); vitamin E, 11 IU; riboflavin, 4.4 mg; calcium pantothenate, 9.6 mg; nicotinic acid, 44 mg; choline chloride 220 mg; vitamin B<sub>12</sub>, 46.6 ug; pyridoxine hydrochloride, 2.2mg; menadione sodium bisulfite 3.49 mg; folic acid, 0.55 mg; D-biotin, 0.11 mg; thiamine mononitrate, 2.2 mg; ethoxyquin, 125 mg. Choline or folate were deleted from the premix for the corresponding studies. Vitamins were supplied by Hoffman-La Roche Inc. (Chemical Division, Nutley, New Jersey).

<sup>d</sup> Calculated values (National Research Council, 1977, Nutrient Requirements of Poultry, 7th edition, National Academy of Science, Washington D.C.).

Table II. Content of amino acids in the basal diets for the choline, folate, lysine, threonine and lysine + arginine feeding trials.

Amino acid	Feeding Trials				Lysine moles/g. of diet * NRC (%)	Threonine moles/g. of diet NRC(%)	Lysine + Arginine moles/g. of diet NRC (%)
	Choline/Folate moles/g. of diet * NRC (%)	lysine moles/g. of diet	NRC (%)	Lysine + Arginine moles/g. of diet NRC (%)			
Lysine	1366±8	NR	1324±7	NR	1393±8	NR	125
Alanine	265±4	NR	264±5	NR	265±8	NR	268
Ammonia	78.5±6	95	78.0±5	9	78.1±2	93	75.5
Aspartic Acid	18.6±6	NR	16.1±4	NR	17.0±1	NR	169
Glycine	1658.8	14.0 <sup>d</sup>	1451.1	14.0 <sup>d</sup>	1556.6	15.0 <sup>d</sup>	1394
Glutamic Acid	2759.4	NR	2714.2	NR	2745.7	NR	268
Half cystine	22.2±6	77.0 <sup>e</sup>	21.2±7	54.0 <sup>e</sup>	14.7±6	52.0 <sup>e</sup>	1264
Histidine	36.8±7	16.3	36.3±5	16.1	37.5±5	16.5	156
Hydroxyproline	10.9±2	NR	21.2±3	NR	28.6±2	NR	7.42
Isoleucine	60.9±9	99	62.5±1	102	5.8±7.4	96	57.1
Leucine	1365.8	134	1374.2	133	1385.6	134	133
Leucine	94.3±6	103	93.4±5	102	94.5±7	102	72.5
Neotyrosine	26.1±3.0	76 <sup>f</sup>	23.5±3.5	63 <sup>f</sup>	26.4±0.93	62.0 <sup>e</sup>	28.3
Phenylalanine	60.1±4.2	13.8	61.0±5.1	14.0	59.3±3.2	13.6	61.2
Folime	12.1±6.1	NR	12.1±5.3	NR	12.1±2.8	NR	12.0
Serine	1035.8	14.0 <sup>f</sup>	1025.7	14.0 <sup>d</sup>	1035.9	15.0 <sup>d</sup>	1394
Threonine	74.6±3.4	1.18	67.2±4.7	1.07	80.7±2.1	1.02	102
Tyrosine	27.5±2.1	11.0 <sup>d</sup>	27.3±2.0	11.0 <sup>d</sup>	27.7±2.1	11.0 <sup>d</sup>	69
Valine	39.5±1.2	11.3	81.4±3.4	11.6	77.6±3.1	11.0	75.3

Means ± standard deviations (for the choline and folate experiments the values are from nine analyses; for the lysine experiment from five analyses; for the threonine experiment from duplicate analyses). Percentage of the NRC dietary requirement (1977).

NRC = no NRC requirement.

<sup>a</sup>Analytical values are low, when compared to the calculated contents from feed composition.

<sup>b</sup>Calculated contents (x NRC) methionine + cysteine = 97.2; methionine + cytosine = 97.2.

combinations.

In each experiment, one-third of the chicks received a diet containing 2.5ug aflatoxin/g of feed. The remaining chickens provided two sets of controls. One group of controls was pair-fed to the feed intake of the birds that received aflatoxin, while the remaining chicks were allowed to consume feed ad libitum. Each of the three primary treatments was subdivided into three groups in which the concentration or mode of administration of the supplemented vitamin or amino acid was varied. In the experiment with choline, the chicks received either the basal ration without the addition of choline, which supplied 106% of the NRC requirement of choline, or else equivalent dosages of choline via diet or intraperitoneal (IP) injection to achieve 175% of the NRC requirement of choline. In the folate experiment, the basal ration provided 244% of the NRC requirement of folate and was supplemented to produce two additional diets containing folate at concentrations of 344 and 644% of the NRC requirement. Similarly, in the threonine experiment the basal diet provided 128% of the NRC requirement, and was supplemented to produce two additional diets containing 155 and 179%. In the lysine experiment, the basal ration containing 102% of the NRC requirement was supplemented to 122 and 146%. A fifth, 4x3 factorial experiment was completed in which the influence of concurrent dietary administrations of lysine and arginine were studied. Each of the three

primary groups (aflatoxin, pair-fed, and ad libitum controls) were subdivided into four groups in which the concentrations and proportions of lysine and arginine were varied. The basal ration provided 102% and 94% of the NRC requirements of lysine and arginine, respectively. This ration was supplemented to produce three additional diets, containing lysine and arginine in the following percentages of the NRC requirement: 102% lysine and 122% arginine, 122% lysine and 122% arginine, and 146% lysine combined with 122% arginine. In all five experiments, each experimental treatment was given to four replicate pens of chicks from the day after hatch. The pens initially contained 6 chicks each and were reduced to 5 chicks each on the seventh day.<sup>a</sup> The experimental treatments were assigned in a completely randomized design. At the termination of the feeding trials, a 5 ml aliquot of blood was obtained from each chick by cardiac puncture using a syringe containing 100 U of sodium heparin. After bleeding, the chicks were killed by cervical dislocation. Livers were excised. All samples were stored at -30C until analyzed with the exception of the livers from feeding trial 5, which were stored at -60C after freeze clamping in liquid nitrogen.

#### Animal performance and biochemical analyses

Feed intake and mortality were recorded daily, while weight gain was recorded daily for the first two experiments, but at three day intervals for the remaining experiments. The following biochemical

<sup>a</sup>Six chicks were initially provided/cage so that chicks lost due to congenital abnormalities or early mortality could be replaced.

parameters were measured at the termination of the experiment: calcium (10), inorganic phosphate (11), total iron, total iron binding capacity (TIBC) and transferrin iron (12), glucose (13), triglyceride-glycerol (14), glycerol (bacterial lipase modification of procedure 15), cholesterol (16), protein (17), albumin (18), blood urea nitrogen (BUN, 19), uric acid (20), bilirubin (21), creatine (22), alkaline phosphatase (AP, 23), lactate dehydrogenase (LDH, 24), aspartate aminotransferase (AST, 25), alanine aminotransferase (ALT, 26), sialytransferase and galactosyltransferase (SAL, GAL, 27) and hepatic lipid (28). Hepatic moisture was determined by drying for 24h at 110C. LDH isozyme patterns in various tissues, plasma and packed blood cells were analyzed by polyacrylamide gel electrophoresis and stained for enzyme activity (29). The plasma concentrations of free-amino acids and ninhydrin positive substances (NPS) were determined using the supernatants from mixtures of 0.8 ml of plasma and 0.8 ml of 2.5% sulfosalicylic acid in lithium citrate buffer (0.15N pH2.2) using a Beckman 121-MB amino acid analyzer and employing a physiological fluid program. The amino acid profiles of feed were also determined using the Beckman 121-MB from 6N hydrochloric acid digestions (in vacuo, 110C, 24h). The digestions for methionine and cysteine employed a hydrogen bromide/performic acid oxidation prior to the hydrochloric acid digestions (30), while the analyses for tyrosine contained 0.1% thioglycolic acid and 0.05% phenol in the hydrochloric

acid digestions (31). Sigma Kit No. 565 (St. Louis, Missouri) was used for the iron profiles. Cholesterol, triglyceride-glycerol and ALT evaluations were assayed with Abbott Laboratory kits (South Pasadena, California). Glycerol profiles were determined with Boehringer Mannheim Kit No. 148270 (Mannheim, West Germany). Except for SAL and GAL, the remaining blood analyses were performed using the Technicon SAM 12/60 autoanalyzer.

Statistical analyses

All performance and biochemical results within a given feeding trial were statistically evaluated by implementing SAS software packages (Statistical Analysis System Institutter Inc., Cary, North Carolina) within an IBM 370/158 or AMDAHL computer system. The statistical procedures included evaluation for the occurrence of interaction in the major effects as well as analysis of variance, Duncan's multiple range test and the Waller-Duncan K-ratio t-test. Statements of significance are based on  $P \leq 0.05$ .

Data presented in chapter III are averages of the values (mean and standard deviation) obtained from the chicks receiving the basal ration (no supplementation) in each of the individual feeding trials.

CHAPTER III  
EFFECTS OF AFLATOXIN

Performance

Before one can study the biochemical effects of dietary supplements on chicks with aflatoxicosis, one must first describe the biochemical changes produced by aflatoxin. This is also necessary in order to demonstrate that a typical aflatoxin lesion has been produced in a given experimental trial. Anorexia, impaired performance and accumulation of hepatic lipid are widely recognized as typical responses to ingested aflatoxin in poultry and most other species of animals (32). The data presented in this chapter, unless otherwise specified, are the combined values obtained from the aflatoxin and control groups receiving the basal ration in the five separate feeding trials. The actual numerical values obtained in the individual experiments from which these figures were derived, are shown in App. 1-25. Fig. 3 shows the effect of aflatoxin on performance of chicks after 24 days of experimental treatment. Weight gain was significantly reduced in the pair fed controls, and further reduced by aflatoxin. This indicates the presence of both an anorexic and a toxic component of this response. The toxic component (decreased weight gain over and above the decrease attributable to the anorexic effects of aflatoxin) was significantly different in only two (folate, and lysine) out of the five feeding trials, and will be discussed in detail in chapter VI. A decreased rate of weight gain (Fig. 4) was apparent by the 6-8th day in the birds receiving aflatoxin, while a decreased feed intake was observed by the 10th day (Fig. 5).

Fig. 3. Effect of aflatoxin on the performance of chicks. Values are expressed as a percentage of the ad libitum control and are averages of the data from the aflatoxin and control groups which received the basal ration (no supplementation) in the five separate feeding trials.

Weight gain, feed consumption, and feed conversion (feed consumption/weight gain) are impaired by aflatoxin. Weight gain is significantly reduced compared to the ad libitum controls, and further reduced compared to the pair-fed controls. This reduction was significant in some feeding trials but not in others and is discussed in detail in chapter VI.

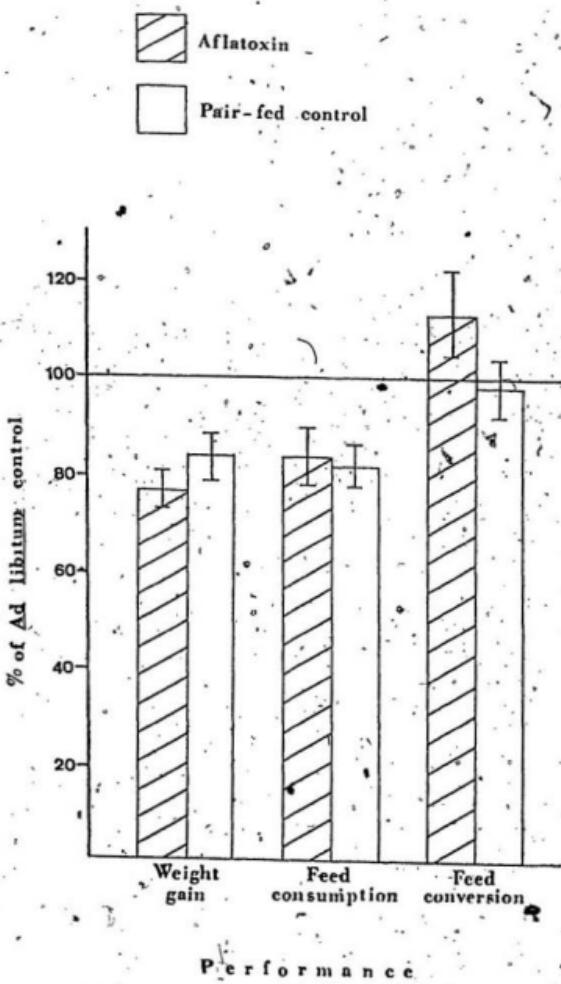


Fig. 4. Daily weight gain of chicks with aflatoxicosis. Values are averages of the data from the aflatoxin and control groups which received the basal ration (no supplementation) in the five separate feeding trials.

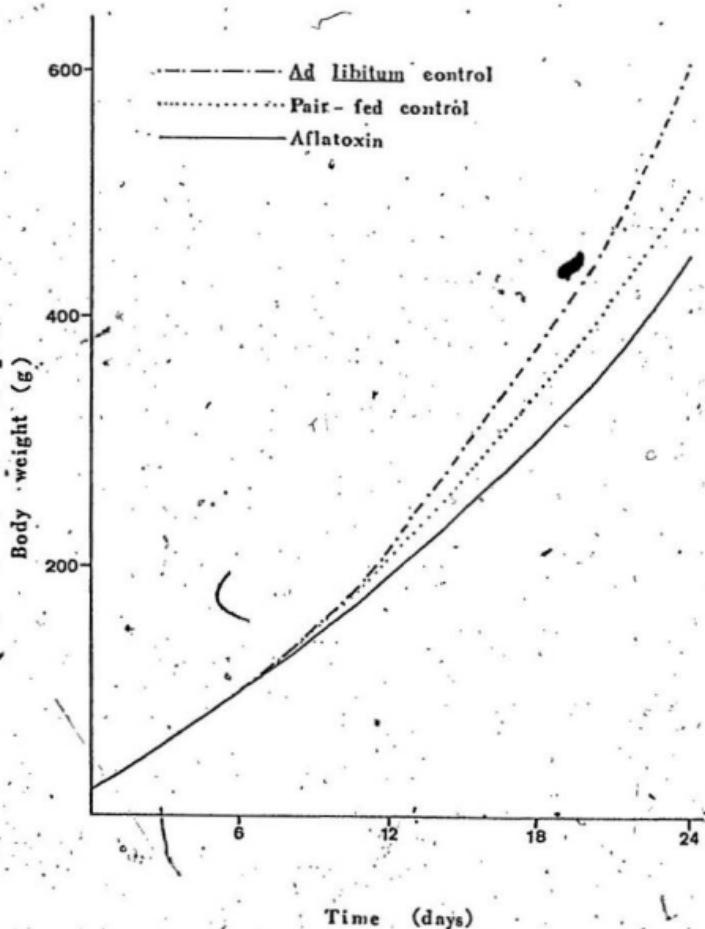
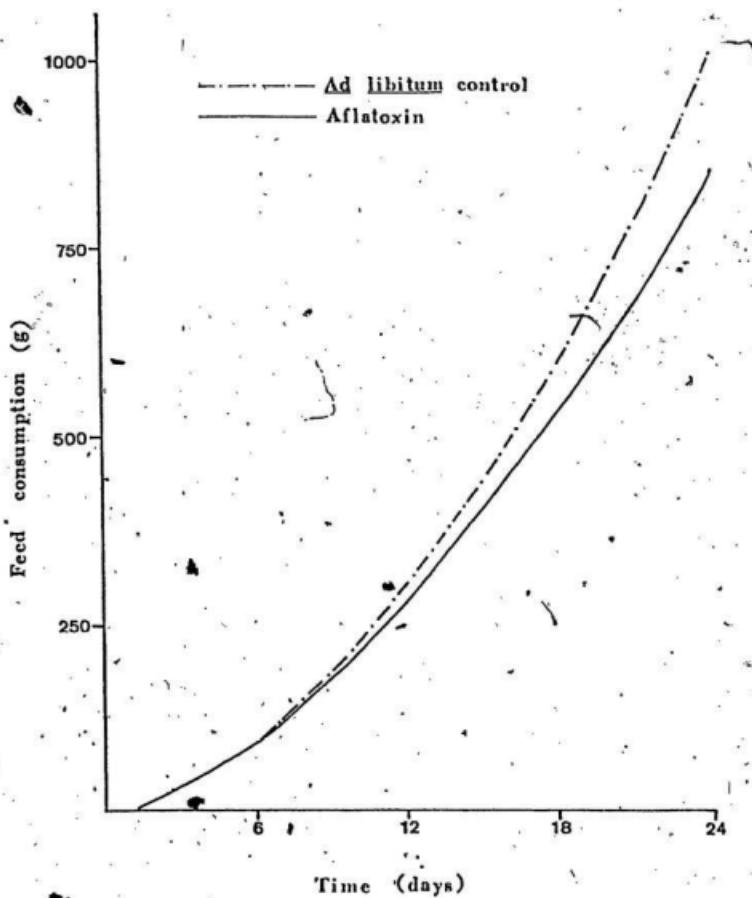


Fig. 5. Daily feed consumption of chicks with aflatoxicosis. Values are averages of the data from the aflatoxin and control group which received the basal ration (no supplementation) in the five separate feeding trials.



Hepatic changes

Hepatic weight, moisture, and lipid content are shown in Fig. 6. The % lipid/g liver can be seen to have been increased by an average of 80%, with an increase in total weight but without an increase in % moisture. Because the liver is considered to be the primary target tissue of aflatoxin, the plasma concentration of a series of hepatic enzymes were measured in plasma as an indication of hepatic damage (feeding trials 1-4 only). Fig. 7 shows that of the enzymes measured, LDH alone was found to increase significantly. An elevation in the activity of LDH of 80-100% was consistent throughout all four experiments. The distribution of aflatoxin-derived radioactivity in the rat, mink, rhesus monkey and swine shows that liver, and to a lesser degree kidney and heart, are the most susceptible tissues for macromolecular binding (33), while in poultry, particularly in broiler chicks, the metabolites are more widely dispersed in all body tissues (34,35). This suggests that the hepatotoxicity of aflatoxin may be less in poultry than in certain other species. Garlich et al. have reported an increase in AP in laying hens with 20ug/g of dietary aflatoxin (36). In ducklings, Brown and Abrams found a slight increase in plasma levels AP, AST, and ALT after receiving 0.5ug/g of dietary aflatoxin for 4 weeks. The elevations progressed to large increases in AST and ALT at 8 weeks, while LDH was markedly elevated at both time intervals (37). Aflatoxin has previously been shown to cause increases in plasma bilirubin, ALT, AST, and cholesterol in goats, cattle, rabbits (38), which are all indications of liver damage. In this

Fig. 6. Effect of aflatoxicosis on hepatic weight, percent moisture, and percent lipid. Values are expressed as a percentage of the ad libitum control and are averages of the data from the aflatoxin and control groups which received the basal ration (no supplementation) in the five separate feeding trials, with the exception of hepatic weight which represents data from feeding trials 1-4 only. Livers from feeding trial 5 were freeze clamped, and total liver weight was not measured.

Hepatic weight and percent lipid are increased by aflatoxicosis.

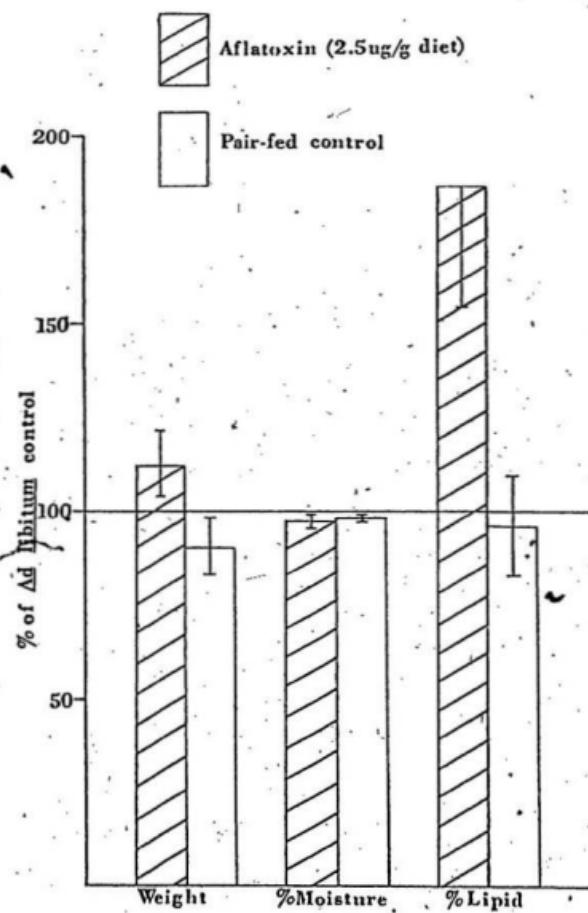
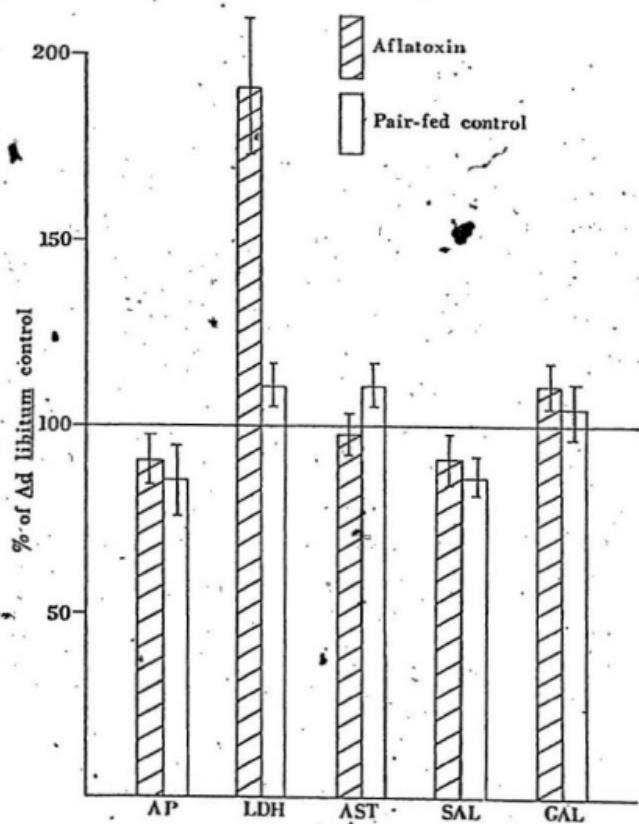


Fig. 7. Effect of aflatoxicosis on plasma levels of selected hepatic enzymes. Values are expressed as a percentage of the ad libitum control and are averages of the data from the aflatoxin and control chicks which received the basal ration (no supplementation) from three or more separate feeding trials as detailed below; Alkaline phosphatase, lactate dehydrogenase, and aspartate aminotransferase are averages from feeding trials 1-4. Sialyltransferase and Galactosyltransferase are averages of data from feeding trials 1-3 only. The analyses were not completed on plasma from the corresponding feeding trials omitted from the calculations, due to the high cost of the assays relative to the amount of new information one would expect the assays to provide.

Only LDH was increased significantly by aflatoxicosis.



series of experiments with broiler chicks, the failure of aflatoxin to induce plasma increases in the other hepatic enzymes, suggests that the plasma LDH probably originates from some tissue other than the liver. LDH activities (Table III) and zymograms (Fig. 8) from various tissues, packed blood cells, and plasma suggest that the elevated LDH activity in plasma originated from hemocytes and not from hepatocytes. This implication is consistent with the unchanged plasma levels of the other hepatic enzymes, as well as a previous report that the fragility of red blood cells increases with aflatoxicosis (39,40). The numbers of the various types of leucocytes, which contain about 50 fold higher level of LDH than erythrocytes, or thrombocytes (41) are markedly altered by aflatoxicosis and thus may contribute to the rise in plasma LDH.

#### Clinical parameters

To establish a clinical picture of the aflatoxin lesion, a series of standard clinical analyses were measured in plasma as shown in Fig. 9. The analyses selected were those which can be obtained rapidly by a single administration of a plasma sample to a standard clinical autoanalyzer (available in the clinical laboratory of any hospital). It was hoped that a distinct pattern or "biochemical fingerprint" could be determined for aflatoxicosis which could then be used to distinguish aflatoxicosis from other syndromes in commercial poultry operations. This fingerprint would also provide a means of monitoring the effects the dietary supplements on the clinical status of the chicks. BUN, and total bilirubin concentrations in plasma were found to increase in response to aflatoxin, with no change in plasma glucose

Table III. Effect of aflatoxin on LDH activities in various tissues.

	Aflatoxin	Pair-fed	<u>Ad libitum</u>
Tissue	Protein (mg/g or ml)		
Liver	80	68	78
RBC	161	154	166
Plasma	12	25	24
	LDH activity (rate/g or ml)		
Liver	129	456	437
RBC	19	25	29
Plasma	0.64	0.46	0.41
	LDH specific activity (rate/mg protein)		
Liver	1.60	6.76	5.64
RBC	0.12	0.17	0.17
Plasma	0.055	0.019	0.017

LDH activities were calculated from linear kinetic data using several incremental aliquots of each sample. Samples were pooled from forty or more individual animals.

Rate = absorbance at 360 nm / minute, or umoles pyruvate reduced / minute.

Fig. 8. Comparison of LDH isozyme patterns from various tissues.

Anaerobic tissues (skeletal muscle, liver) predominantly synthesize the polypeptide chain A which is more efficient under anaerobic conditions, while aerobic tissues (heart, RBC) predominantly synthesize chain B which is more efficient under aerobic conditions. Each isozyme (1-5) is a tetramer, formed from a combination of the two different monomers in all possible combinations, with LDH-1 formed from four B monomers, and LDH-5 from four A monomer's.

Fig. 8. shows that liver contains predominantly isozymes 3, 4, and 5, while RBC contains predominantly 2, and 3. Likewise, plasma contains predominantly isozymes 2 and 3. No increase in isozymes 3, 4, and 5 is apparent in the plasma of the birds receiving aflatoxin as would be expected if the increased plasma LDH was of hepatic origin.

Two bands were obtained from the plasma of the birds receiving aflatoxin due to leakage of the sample from the well in which it was applied into an adjacent well at the origin.

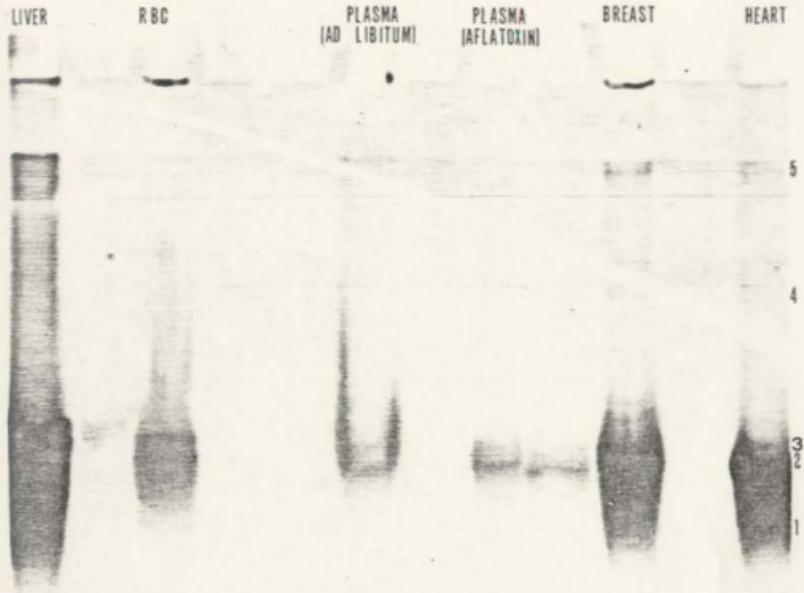
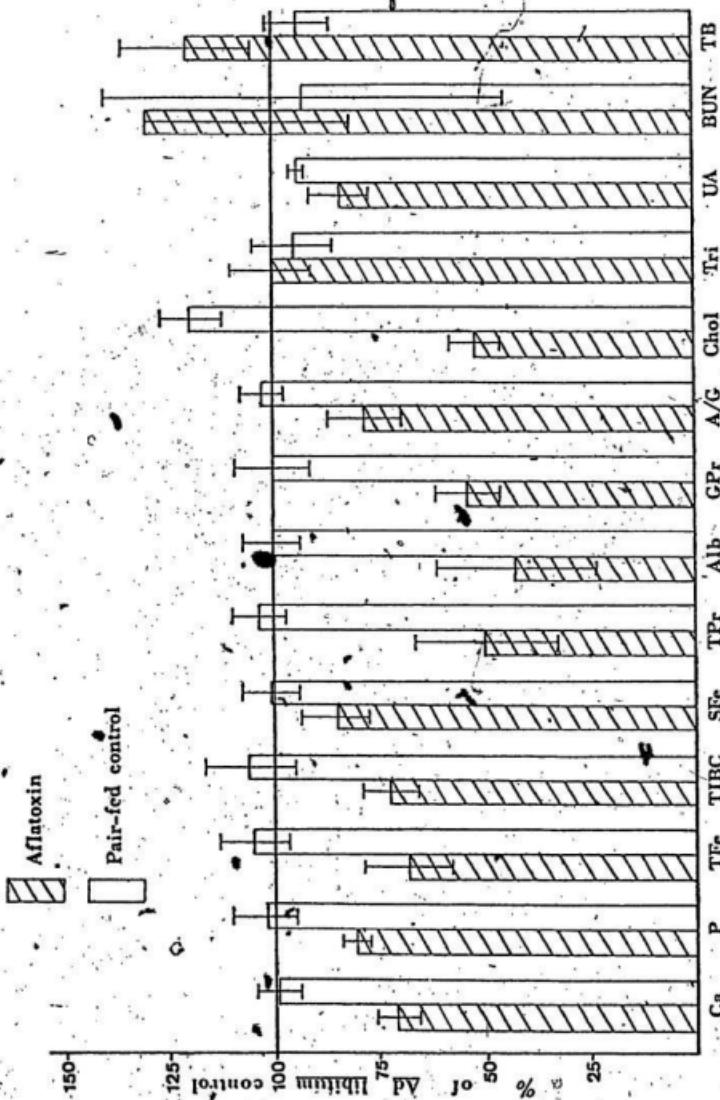


Fig. 9. Effect of aflatoxicosis on the clinical biochemistry of chicks. Values are expressed as a percentage of the ad libitum control and are averages of data from chicks which received the basal ration (no supplementation) in feeding trials 1-4. The screen for clinical parameters was not completed on plasma from feeding trial 5.



and triglyceride concentration, while the uric acid, total protein, albumin, globulin, albumin/globulin ratio, cholesterol, calcium, inorganic phosphate, total iron, total iron binding capacity, and percent saturated transferin concentrations decreased. BIL and total bilirubin were not present in sufficient concentrations to permit reliable quantitation, although there appeared to be an elevation in these compounds. The findings for calcium, total protein, and glucose are consistent with the responses reported in the literature (37,39,42) while the decrease in total iron and TIBC supports the observation by Lanza *et al.* that aflatoxin reduces iron absorption (43). The suppression in albumin, globulin, and albumin/globulin ratio has also been reported to occur in ducklings and laying hens (37). Other hepatotoxins usually induce an increase in the globulin fraction, and therefore the electrophoretic pattern of plasma proteins obtained with aflatoxin may be of diagnostic significance.

The suppression in plasma concentration of calcium, phosphorus, total protein, albumin, total iron, and cholesterol, but with elevated LDH, is a profile that is typical of malabsorption syndromes (44). Osborne and Hamilton have shown that aflatoxicosis lowers the concentration of bile secreted (45) and decreases the formation of pancreatic lipase (46), thereby impairing the digestion and absorption of lipids, and lipid soluble vitamins. This relationship between aflatoxin and malabsorption will be discussed in more detail in chapter IV.

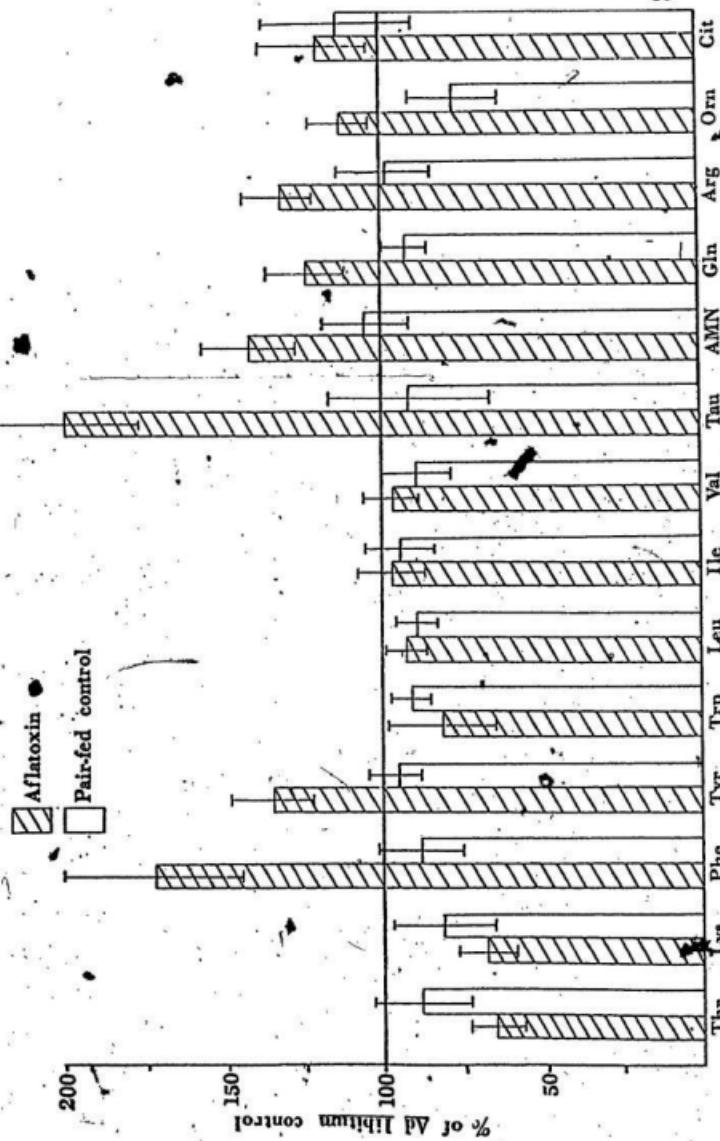
#### Plasma free-amino acids

Plasma amino acid response to aflatoxin was also monitored. An

earlier study reported that aflatoxin (1.25 or 2.5ug/g. of diet), reduced the concentrations of all amino acids in plasma of broiler chicks (Cobb/Cobb). However, the protocol for the previous amino acid determinations involved hydrochloric acid hydrolysis of supernatants obtained from mixtures of sulfosalicylic acid and plasma (47). The acid hydrolysis was necessary since a non-physiological amino acid program was employed. As a result, the values which were obtained included free-amino acids, conjugates, peptides, and proteins not precipitated by sulfosalicylic acid. The amino acid profiles reported in this study were determined using a physiological program, and represent free-amino acid contents. Certain amino acids or amino acid ratios responded to aflatoxin in a remarkably consistent fashion throughout the five experiments, and are shown in Fig. 10. Data for all other amino acids and ninhydrin positive substances showed a non specific or variable response to aflatoxin and are shown in App. 21-25.

Elevated phenylalanine and tyrosine, without an increase in tryptophan, but with a reduction in branched chain amino acids (BCAA) in plasma has been reported by several authors as a result of impaired liver function. This has been rationalized by the fact that the aromatic amino acids (AAA) are metabolized in the liver, while the BCAA are metabolized mainly in the muscle. AAA are precursors to various neurotransmitters, and compete with the BCAA for entry through the blood-brain barrier. Changes in the AAA/BCAA ratio have been implicated in the initiation of hepatic coma which is actually

Fig. 10. Effect of aflatoxicosis on the plasma concentrations of selected amino acids. Values are expressed as a percent change of the ad libitum control and are averages of the data from the chicks which received the basal ration (no supplementation) in the five separate feeding trials.



reversible by infusion of BCAA (48,49).

In the data presented in this report, the ratio of the sum of the BCAA, leucine, isoleucine, and valine, to the AAA, phenylalanine, and tyrosine, was reduced from 2.33 and 2.28 in the plasma of the ad libitum and pair fed controls respectively, to 1.27 in the aflatoxin groups. The decreased BCAA/AAA ratio was due primarily to the elevated plasma levels of AAA, and was very likely attributable to an aflatoxin induced impairment of hepatic function. A possible consequence of this altered ratio is that the competitive action of the branched chain amino acids on cerebral uptake of aromatic amino acids would be reduced (50). High levels of phenylalanine in plasma have been shown to raise cerebral levels of phenylalanine sufficiently to competitively inhibit neural tyrosine hydroxylase (51), and thus suppress the synthesis of catecholamines, while promoting the synthesis of false neurotransmitters (ea. octopamine, and phenylethanolamine), as is observed in hepatic insufficiency (52). Increased brain tryptophan would result in enhanced synthesis of serotonin (53). Although plasma tryptophan was not increased, elevations in brain tryptophan have been observed in hepatic syndromes (with no increase in plasma tryptophan) (48,49). Since serotonin containing neurons are associated with suppressed behavior, while neurons containing catecholamines are associated with arousal, the observed changes in BCAA/AAA ratios may be related to the behavioral changes such as anorexia. The ratio of tyrosine to phenylalanine (T/P) was also reduced from 1.28 and 1.33 in the ad libitum and pair fed controls respectively to 0.92 in the aflatoxin treatments. Anderson

has demonstrated that a correlation exists between a suppression in the T/P ratio and a decrease in feed intake (54). The 28% decrease in the T/P ratio in our data would be predicted to induce the 20% depression in feed intake observed in the aflatoxin groups by applying Anderson's model.

Taurine, which was also elevated in response to aflatoxin, is a component of the bile salt, taurocholic acid. Conjugation with taurocholic acid represents a major route for the excretion of aflatoxin and its metabolites in the chicken (34) and many other species (55). Dietary aflatoxin has been shown to reduce the concentration of bile salts in the bile, to but increase the volume of bile present in the gall bladder (45).

Elevated plasma levels of ammonia, glutamine, and perhaps BUN suggest a decreased ability to excrete nitrogen and/or an increased catabolism of protein or, other nitrogenous compounds. Although excretion of nitrogen as urea is a minor pathway in the chicken, the decreased plasma concentration of uric acid together with the elevated argininé, and ornithine, (urea cycle intermediates) and perhaps BUN, suggests an increased importance of urea production in the excretion of nitrogen during aflatoxicosis. This observation will be discussed in detail in Chapter V. Changes in plasma amino acid concentrations can have numerous physiological effects which will be discussed in more detail in Chapters V & VI. Some of the symptoms of aflatoxicosis may be directly attributable to these plasma abnormalities.

CHAPTER IV

EFFECT OF CHOLINE AND FOLATE SUPPLEMENTATION

Introduction

A major aspect of aflatoxicosis in livestock involves the nutritional status of the affected animal (47,56). Aflatoxin interacts with protein, lipid, and vitamin metabolism, and also increases most nutritional requirements (32,57). Concentrations of thiamine, riboflavin, vitamin B6, pantothenate, niacin, biotin and choline decrease in the plasma, bile and liver of chicks fed aflatoxin contaminated feed. Only the concentration of folate increases in plasma and bile (47). Dosages of aflatoxin too low to inhibit the growth of chicks influence lipid synthesis and transport (58). Low dosages of aflatoxin also diminish the concentration of bile secreted and pancreatic lipase activity is decreased in proportion to the dosage of aflatoxin. These two factors combine to produce steatorrhœa, a result of impaired digestion and absorption of lipophilic substances including the lipid soluable vitamins (45,46). The reduction in bile salts, which inhibit the growth of Gram positive bacteria, along with the alteration in the secretion of digestive enzymes, may result in the development of an atypical intestinal microflora (59) as well as hinder nutrient digestion or absorption. This suggestion is consistent with the observation that the administration of antibiotics, especially vitamin-antibiotic combinations to poultry with aflatoxicosis will improve feed conversion, weight gain, and mortality (56,60). These factors, together with aflatoxin induced anorexia, could be expected to produce nutritional deficiencies that may be responsive to dietary therapy.

A variety of vitamin and nutrient supplements or deficiencies have

been reported to interact with aflatoxicosis. Isocaloric diets formulated on a constant calorie/protein ratio and high in lipid, provide a mortality sparing effect independent of the degree of saturation, but if the lipid also contains a high level of unsaturated fatty acids, there is also a significant growth promoting effect (60). Dietary supplementation with vitamin B6 improves feed conversion of chicks with aflatoxicosis but increased pantothenate has no effect (unpublished results, Voigt, Wyatt, and Ayres). Supplemental choline combined with B12 provides a protective effect to quail with aflatoxicosis (61). Dietary deficiency of thiamine, or supplementation with folic acid combined with vitamin B6 and vitamin D have been reported to offer mediation to poultry consuming aflatoxin (56, 42). In addition, diets marginally deficient in choline, methionine, and folic acid enhance hepatocarcinogenesis and depress hepatic mixed-function oxidase activities in rats fed aflatoxin B1 (62), while folic acid added to swine rations containing mouldy corn has been reported to improve growth and feed conversion (63).

The object of the first two feeding trials was to evaluate the influence of dietary folate or dietary and intraperitoneal (IP) choline on the aflatoxin lesion. Choline was selected to be evaluated as a mediative agent for aflatoxicosis for the following reasons: (a) to determine if aflatoxicosis induces a nutritional deficiency of choline, since low levels of choline occur in plasma during aflatoxicosis (47), (b) to verify reports of positive effects from administration of various choline-containing dietary supplements

(61,62), and (c) to determine if the fatty liver syndrome induced by aflatoxicosis can be attributed to a deficiency of choline, since choline deficiency will also induce fatty liver (32). Choline was administered via IP injections, as well as through the diet, in an attempt to minimize the effects of any aflatoxin induced reduction in intestinal absorption of choline, although the enterohepatic circulation of bile prevents the total bypass of any effects attributable to malabsorption.

Like choline, folate functions in the biological reactions involving the transfer of single carbon units and is also a lipotropic factor. Similarities between the metabolic effects of folate deficiency and aflatoxicosis suggested exploring a dietary trial with folate, i.e. folacin is required for hemopoiesis, hemostasis and immune responses, all of which are affected by aflatoxin (65,66). Folate is also the only vitamin which is excreted at higher concentrations in the bile during aflatoxicosis (47).

#### Effects of choline on performance

The effect of dietary and IP choline supplements on the performance of chicks is shown in Table IV. Choline supplied in the diet, but not when given via IP injections, resulted in a 16% increase in the weights of birds stressed by aflatoxin or by pair-feeding, but only provided an 8% increase in the ad libitum controls. Dietary choline also produced an 11% increase in feed intake in the ad libitum controls, but no significant increase in the feed intake of the aflatoxin group. Supplemental choline is clearly not mediating the

Table IV. Effect of dietary and IP choline supplements on the performance of chicks with aflatoxicosis.

	Supplemental choline		
	a 0	b IP	b Diet
Weight Gain (g)			
Aflatoxin	521+23D	480+22D	603+44BC
Pair-fed	530+34D	499+22D	616+45BC
<u>Ad libitum</u>	638+60BC	587+36C	689+33A
Feed Intake (g)			
Aflatoxin	823+71BCD	728+61CDE	834+98BC
Pair-fed	796+67CDE	710+23E	792+82CDE
<u>Ad libitum</u>	928+65AB	875+78BC	1029+83A
Feed Conversion (Feed intake/weight gain)			
Aflatoxin	1.73+0.2A	1.66+0.2A	1.49+0.1AB
Pair-fed	1.63+0.1AB	1.55+0.1AB	1.38+0.1B
<u>Ad libitum</u>	1.56+0.1AB	1.60+0.1AB	1.90+0.2AB

Values (mean and standard deviations) for a parameter followed by the same letter are not significantly different at  $P \leq 0.05$ . Evaluation for the occurrence of interaction in the major effects is provided in Appendix Tables 1a-3a.

a No supplemental choline.

b Administration to achieve 175% of the NRC requirement (1977) of choline (basal ration provides 106%).

anorexic effects of aflatoxin, however the analysis of the effects of choline on aflatoxin toxicity is made difficult by the control response to choline supplementation. The 106% of the requirement of choline suggested by the NRC which was provided in the basal diet was not sufficient to provide maximal growth. This suggests that the NRC requirements for choline should be revised.

Effect of choline on hepatic lipid, and on selected biochemical constituents in plasma

The data on hepatic lipid content in Table V indicates that dietary choline significantly decreased hepatic lipid in the aflatoxin group, but not in either control group, while IP choline had no effect. As feed intake was actually increased by dietary choline in the aflatoxin group, the decreased hepatic lipid suggests a moderation of the toxic effects of aflatoxin.

Table VI compares the effect of IP and dietary choline supplements on the % change in selected plasma constituents between the aflatoxin and the pair fed controls. In all cases IP choline reduced the effect of aflatoxin, while dietary choline had little or no effect. \*

Similarly, IP choline reduced the effect of aflatoxin (see Table VII.) on glutamine, lysine, threonine, citrulline, ornithine, and tryptophan, while dietary choline had little or no effect. Arginine, tyrosine, phenylalanine, and serine were not affected by either supplement, while the increase in taurine was reduced by injected choline and further reduced by dietary choline. The tendency for choline, particularly IP choline, to normalize the plasma levels

Table V. Effect of dietary and IP choline on hepatic lipid content.

<u>Treatment</u>	<u>Hepatic lipid (% DWB)</u>		
	<u>No. choline</u>	<u>a IP choline</u>	<u>Dietary choline</u>
Aflatoxin	31.4+ 4.2A	32.1+ 4.2A	27.5+ 3.6B
Pair-fed	12.9+ 2.0C	14.3+ 2.8C	12.5+ 1.1C
<u>Ad libitum</u>	14.0+ 2.1C	14.4+ 1.4C	14.9+ 1.1C

Values (mean  $\pm$  standard deviations) for a parameter followed by the same letter are not significantly different at  $P \leq 0.05$ . Evaluation for the occurrence of interaction in the major effects is provided in Appendix Table 16a.

<sup>a</sup> Administration to achieve 175% of the NRC requirement (1977) of choline (basal ration provides 106%).

Table VI. Effect of dietary and IP choline on the percent change of selected plasma constituents between the chicks receiving aflatoxin and the pair-fed controls.

	<u>Aflatoxicosis vs. pair-feeding (% change)</u>		
	<sup>a</sup> No choline	<sup>b</sup> IP choline	<sup>b</sup> Dietary choline
Calcium	-31 *	-21 *	-36 *
Phosphorus	-18 *	-2	+8
Cholesterol	-57 *	-52 *	-66 *
Total protein	-57 *	-29 *	-57 *
Total transferin iron	-32 *	-8 *	-39 *
LDH	+73 *	+46 *	+71 *

<sup>a</sup> No supplemental choline

<sup>b</sup> Administration to achieve 175% of the NRC requirement (1977) of choline (basal ration provides 106%).

\* Percent change ([plasma constituent] in chicks receiving aflatoxin / [plasma constituent] in pair-fed control chicks X 100) is significant at  $P < 0.05$ . ie. The concentration of the given plasma constituent in chicks receiving aflatoxin is significantly different at  $P < 0.05$  from the same plasma constituent in the pair-fed controls. Raw data and evaluation for the occurrence of interaction in the major effects is provided in Appendix Tables 17 and 17a respectively.

Table VII. Effect of dietary and IP choline on the percent change of selected plasma amino acids between the chicks receiving aflatoxin and the pair-fed controls.

	Aflatoxicosis vs. pair-feeding (% change)		
	a 0	b IP	b DIETARY
Glutamine	+78 *	+42 *	+78 *
Lysine	-17	-4	-31 *
threonine	-31	-12	-12
citrulline	-14	+13	-40
ornithine	+210 *	+140 *	+391 *
arginine	+71 *	+85 *	+126 *
tyrosine	+63 *	+53 *	+30 *
phenylalanine	+167 *	+171 *	+202 *
tryptophan	+2	-18 *	+14
serine	-20 *	-24 *	-25 *
cystathionine	-20	-33 *	+14
taurine	+153 *	+107 *	-7

<sup>a</sup> No supplemental choline.

<sup>b</sup> Administration to achieve 175% of the NRC requirement (1977) of choline (basal ration provides 106%).

\* Percent change ([plasma constituent] in the aflatoxin chicks/[same constituent] in pair-fed controls X 100) is significant at P<0.05. i.e., The concentration of the given plasma constituent in chicks receiving aflatoxin is significantly different from the pair-fed controls.

of amino acids and other plasma constituents, suggests that nutrient absorption was improved by choline, perhaps by stimulating bile salt formation. The choline induced decreased in taurine is in agreement with increased taurocholic acid production.

#### Effects of folate

Data on the effect of supplemental folate on the performance and hepatic lipid content of chicks with aflatoxicosis are shown in Tables VIII and IX. Folate showed no protective action against aflatoxin. Similarly, folate supplementation had no significant effect on the plasma concentrations of the clinical parameters and amino acids measured (App. 18 and 22). However, it is interesting to note that it did produce a significant increase in 10 different amino acids (Table X) in the pair-fed controls which did not occur in the aflatoxin birds. It is known that certain antibiotics, and hormones of similar structure to aflatoxin, adversely affect the activity of folate. Perhaps aflatoxin has a similar effect.

Table VIII. Effect of supplemental folate on the performance of chicks with aflatoxicosis.

<u>Treatment</u>	Supplemental folate (% of NRC dietary requirement)		
	224	344	644
Aflatoxin	435+24EF	474+21E	412+12F
Pair-fed	517+19D	564+65C	552+45CD
<u>Ad libitum</u>	673+15A	583+13BC	617+22B
	Feed intake (g)		
Aflatoxin	881+73CD	834+32D	927+78C
Pair-fed	884+57CD	831+38D	920+59C
<u>Ad libitum</u>	1096+48B	1174+84AB	1202+63A
	Feed conversion (g)		
Aflatoxin	2.23+0.3B	1.91+0.2CD	2.48+0.3A
Pair-fed	1.84+0.2D	1.59+0.2E	1.79+0.1DE
<u>Ad libitum</u>	1.72+0.0DE	2.15+0.2B	2.07+0.1BC

Values (mean + standard deviations) for a parameter followed by the same letter are not significantly different at  $P \leq 0.05$ . Evaluation for the occurrence of interaction in the major effects is provided in Appendix Tables 4a-6a.

Table IX. Effect of supplemental folate on hepatic lipid.

<u>Treatment</u>	<u>Hepatic lipid (% dry weight basis)</u>		
	<u>Supplemental folate (% NRC dietary requirement)</u>	244	344
Aflatoxin		27.4 <sub>±</sub> 2.1A	28.9 <sub>±</sub> 2.8A
Pair-fed		15.9 <sub>±</sub> 3.2BC	17.5 <sub>±</sub> 1.1B
<u>Ad libitum</u>		14.1 <sub>±</sub> 1.2C	14.2 <sub>±</sub> 1.8C
			13.9 <sub>±</sub> 1.4C

Values (mean<sub>±</sub> standard deviations) for a parameter followed by the same letter are not significantly different at P<sub><</sub>0.05. Evaluation for the occurrence of interaction in the major effects is provided in Appendix Table 16a.

Table I. Percent change of selected plasma amino acids in chicks receiving a 16% increase in dietary folate.

	<u>Basal vs. supplemental folate (% change)</u>	
	<u>Aflatoxin</u>	<u>Pair-fed</u>
Asparagine	+2	+48*
Threonine	-9	+20*
Serine	+10	+37*
Glutamine	+4	+49*
Proline	+6	+18*
Glycine	+3	+25*
Alanine	-6	+24*
Valine	-2	+16*
Methionine	-8	+30*
Histidine	+11	+18*

\* Percent change ([plasma amino acid] in chicks receiving supplemental folate / [plasma amino acid] in chicks receiving the basal ration X 100) is significant at  $P \leq 0.05$ . ie. The plasma concentration of the given amino acid in plasma of the chicks receiving supplemental folate is significantly different from the concentration of the same plasma amino acid in chicks receiving the basal ration. Raw data and evaluation for the occurrence of interaction in the major effects is provided in Appendix Table 22 and 22a respectively.

CHAPTER V

EFFECT OF LYSINE AND THREONINE SUPPLEMENTATION

Introduction

As a result of its toxic and carcinogenic effects, aflatoxin has caused severe economic losses to the animal industries. Poultry are susceptible to the acute toxicity of aflatoxin, but resistant to its carcinogenic effects. The level of in vivo binding of metabolites of aflatoxin B1 to DNA appears to reflect the susceptibility of a species to the carcinogenic action of the toxin, while the level of protein binding reflects susceptibility to the acutely toxic action of aflatoxin B1 (67). Acute toxicity in poultry is characterized by poor performance (decreased feed consumption, weight gain, and feed conversion 68,69), accumulation of fat in the liver (70), impaired protein synthesis (71), and increased susceptibility to infection (72).

Data presented in Chapter III has shown that abnormal plasma concentrations of certain amino acids or amino acid ratios are consistently induced by administration of 2.5 ug aflatoxin/g diet to Hubbard/Hubbard broiler chicks for 24 days. These changes included marked elevation of plasma phenylalanine, tyrosine, arginine, and taurine, while lysine and threonine concentrations were reduced. Deficiencies, excesses and imbalances of amino acids can lead to similar changes in plasma free amino acids (73) and, like aflatoxin, may cause a marked reduction in feed consumption and weight gain (74). A rise in plasma phenylalanine roughly equivalent to that induced by dietary aflatoxin has been reported to result in impaired protein synthesis and a decrease in antibody response (75,76). Accumulation of fat in the liver has also been reported to result from

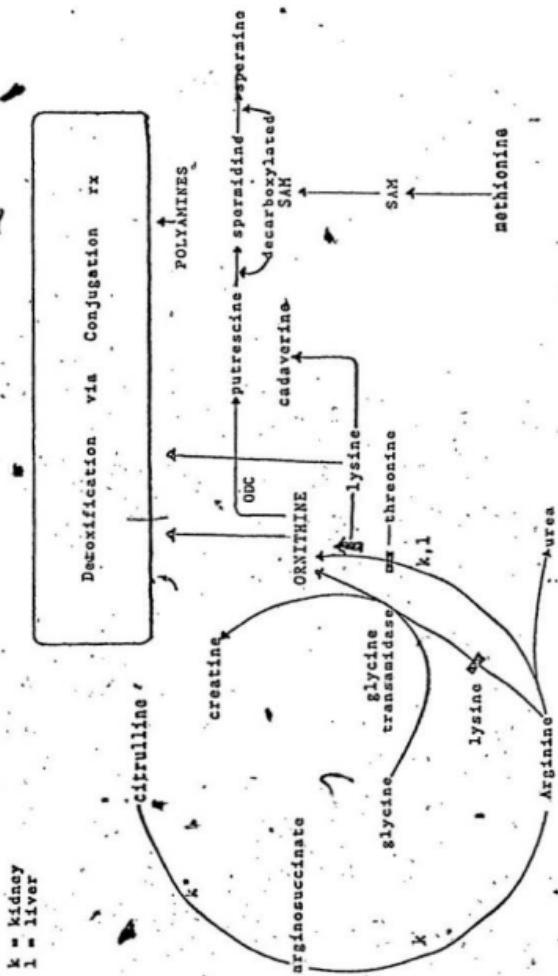
various plasma amino acid imbalances (77,78). If the effects of aflatoxin are related in part or in whole to these abnormal amino acid levels, the symptoms should be relieved if these concentrations could be normalized through appropriate dietary amino acid supplementation. Feeding trials 3 and 4 studied the effects of increasing increments of threonine or lysine on the performance and clinical biochemistry of chicks with aflatoxicosis.

Lysine and threonine were chosen because of the low concentrations of these amino acids in the plasma, and also because of the elevated levels of plasma arginine observed in the birds suffering from aflatoxicosis. Lysine competes with arginine for reabsorption in the renal tubules, and thus an excessive dietary intake of lysine can cause reduced arginine retention. In addition, lysine markedly increases renal arginase activity, thus stimulating the degradation of arginine to ornithine and urea (79,80). In contrast, threonine inhibits this reaction (80).

The resulting changes in ornithine production might also be expected to effect the symptoms of aflatoxicosis. Chickens do not reutilize ornithine for urea production as do ureatilic animals (Fig. 11). Instead ornithine is available to participate in detoxification through conjugation with a variety of xenobiotics, most notably benzoic acid. Ornithuric acid (dibenzoylornithine) is a major detoxification product in birds, snakes, and lizards, largely replacing the glycine conjugate hippuric acid which is excreted by mammals (81,82). Daily benzoic acid administration has been observed

Fig. 11. Role of ornithine in the formation of physiological nucleophiles

54



to increase urea nitrogen from 1% to 9% of the total nitrogen excreted by chickens. This rise in urea production was used to indicate the operation of the ornithine detoxification mechanism (82,83). Data presented in Chapter III showing an aflatoxin induced increase in plasma levels of ammonia, glutamine, possibly BUN, and the urea cycle intermediates, together with a decreased plasma concentration of uric acid, suggests a similar increased excretion of nitrogen as urea in response to daily aflatoxin administration, and thus suggests a similar reliance on the ornithine detoxification mechanism.

In addition, ornithine, together with lysine and methionine, is a precursor of the polyamines (putrescine, spermidine, spermine, and cadaverine). Polyamines are also known to be excreted bound to such compounds as plant alkaloids, antibiotics of microbial nature, and various carboxylic acids (84). If these highly nucleophilic compounds facilitate the excretion of aflatoxin or its metabolites through the formation of conjugates, then changes in the plasma concentrations of these compounds could be expected to effect the performance of chicks with aflatoxicosis. Metabolites of aflatoxin are known to be excreted in bile and urine as conjugates of compounds such as taurine, and glucuronic acid (34,55). The metabolism of aflatoxin B1 by chicken liver microsomes has been reported to result almost exclusively in the formation of the 2,3-dihydroxy-2,3-dihydro-aflatoxin B1 (DHD-B1) metabolite of aflatoxin B1 (previously misidentified as the B2a or hemiacetyl derivative), which binds extensively to microsomal protein at neutral pH (85). Protein binding

has also been observed in vivo with up to 63% of a given dose of aflatoxin reported to be concentrated in the muscle protein in broiler chicks (35), compared to 0.8% in piglets (33). It has been assumed that the dialdehydic phenolate ion of aflatoxin B1-dhd forms at physiological pH, and reacts with the primary amine groups of proteins to form a Schiff base (85). This same mechanism could apply to the formation of AFB1-dhd conjugates with the amine groups of free lysine, ornithine, arginine, and/or the polyamines. Thus the opposing effects of lysine and threonine on the production of ornithine and the polyamines provides a dual purpose for the choice of these two amino acids as dietary supplements to combine with dietary aflatoxin.

#### Contrasting effects of threonine and lysine on performance

Data on the effects of threonine and lysine supplementation on weight gains, feed conversion, and feed intake, are given in Tables XI and XII and XIII respectively. Lysine produced an increase in weight gain and feed conversion in the birds suffering from aflatoxicosis which was not observed in the pair-fed controls. Feed intake decreased in the controls, but remained unaffected in the aflatoxin treatments. In contrast, threonine failed to produce the beneficial effect on the body weight, feed conversion, and feed intake in the aflatoxin group, which is observed in the controls.

#### Lysine/arginine ratio

In Chapter III, plasma arginine concentration was shown to increase dramatically in response to aflatoxin, while plasma lysine

Table XI. Effect of graded levels of lysine or threonine on the final weight of chicks with aflatoxicosis.

		Final Weight (g)		
	Dietary supplement (% NRC requirement)	Aflatoxin	Pair-fed	Ad libitum
Lysine	102	367+32F	461+33C	545+47B
	122	411+25DE	446+13CD	586+22A
	146	402+20EF	448+11C	546+21B
Threonine	122	504+24EF	527+55DEF	620+39BC
	150	486+41F	547+17DE	628+24B
	175	519+36EF	573+59CD	729+40A

Values (means + standard deviations) for a parameter followed by the same letter are not significantly different at  $P \leq 0.05$ . Evaluation for the occurrence of interaction in major effects is provided in Appendix Tables 7a and 10a.

Table XIII. Effect of graded levels of lysine or threonine on feed conversion of chicks with aflatoxicosis:

		Feed Conversion (feed intake/weight gain)		
Dietary Supplement (% NRC requirement)	Aflatoxin	Pair Fed	Ad libitum	
Lysine	102	2.55±0.11A	1.94±0.17E	2.07±0.14CDE
	122	2.23±0.08BC	2.03±0.06DE	2.16±0.30BCD
	146	2.26±0.21B	1.99±0.08DE	1.93±0.11E
Threonine	122	1.84±0.13ABC	1.65±0.07C-F	1.72±0.15B-E
	150	1.90±0.14A	1.61±0.06DEF	1.73±0.04BCD
	175	1.79±0.13ABC	1.50±0.08F	1.57±0.08EF

Values (means ± standard deviations) for a parameter followed by the same letter are not significantly different at P<0.05. Evaluation for the occurrence of interaction in the major effects is provided in Appendix Tables 12a and 12b.

Table XIII. Effect of graded levels of lysine or threonine on feed intake of chicks with aflatoxicosis.

		Feed intake (g)	
Dietary supplement (% NRC requirement)		Aflatoxin	Ad libitum
Lysine	102	839±61D	1050±22B
	122	830±52D	1180±69A
	146	821±56D	981±37C
Threonine	122	864±91B	1004±67A
	150	853±69B	1029±43A
	175	863±113B	1084±79A

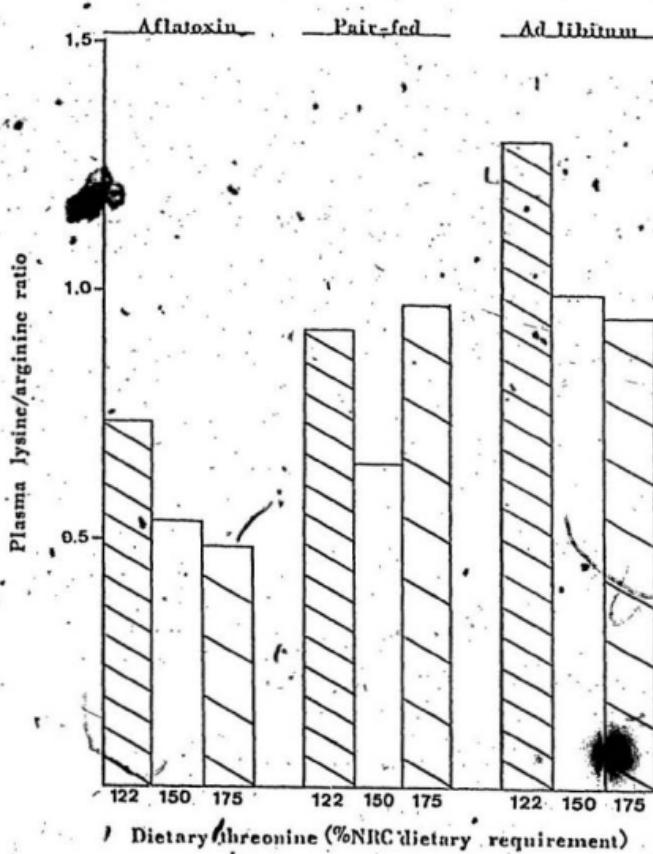
Values (means  $\pm$  standard deviations) for a parameter followed by the same letter, are not significantly different at P<0.05.

Evaluation for the occurrence of interaction in the major effects is provided in Appendix Tables 8a and 11a.

concentration decreased slightly. This aflatoxin induced decrease in plasma lysine/arginine ratio may be related to the negative effect of increased dietary threonine and the positive effect of increased dietary lysine on the performance of chicks receiving aflatoxin (Fig. 12 and 13). Fig. 13 shows that the 43% increase in dietary lysine increased the low plasma lysine/arginine ratio (0.48) in the birds suffering from aflatoxicosis to a more normal ratio (1.2). However, in the pair-fed and ad libitum controls, the ratios were increased to abnormally high ratios of 1.5, and 2.2, respectively, in response to increased dietary lysine. In contrast the already low lysine/arginine ratio (0.73) of the aflatoxin chicks was further reduced to 0.48 by their 40% increase in dietary threonine (Fig.12), while the control ratios were also decreased but remained within the normal range.

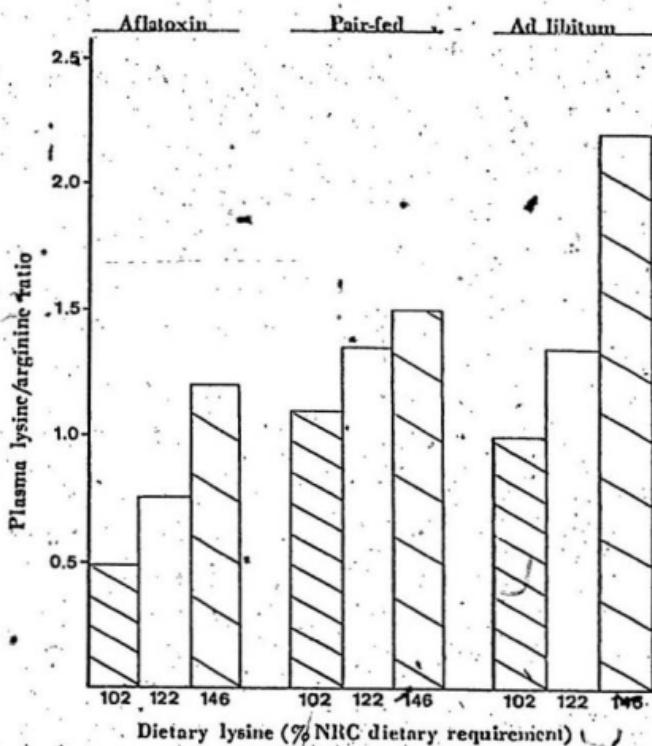
As previously mentioned, excess dietary lysine increases the degradation of arginine to ornithine and urea, while also decreasing arginine retention by competing for reabsorption by the renal tubules. This results in an increase in plasma lysine/arginine ratio and in dietary arginine requirement with a consequent decrease in performance which may be relieved by increased dietary arginine (79,80,86,87). Arginine requirement, and thus tolerance of excessive dietary lysine or arginine varies considerably from one strain of chicks to the next and even between families within strains (79,88). This variation appears to result from genetic differences in lysine metabolism (89). The selection of chicks with high arginine (H.A.), or low arginine (L.A.) requirement

Fig. 12. Effect of graded levels of threonine on plasma lysine/arginine ratio of chicks with aflatoxicosis.



62a

Fig. 13. Effect of graded levels of lysine on plasma lysine/arginine ratio of chicks with aflatoxicosis.



has produced two strains of chicks; Nesheim H.A and Nesheim L.A. (88).

Regardless of the type of diet fed, plasma lysine levels in the H.A. strain were nearly double those of the L.A. strain, while plasma arginine levels showed little variation.

Susceptibility to aflatoxicosis also varies considerably from one strain of chicks to the next or even between individuals within a strain (2). In light of the effects of aflatoxin on the plasma lysine/arginine ratio in the Hubbard/Hubbard strain, and the improved performance seen after normalization of this ratio, it would be interesting to compare the susceptibility of the H.A. and L.A. strains to aflatoxicosis. The H.A. strain, already having a high lysine/arginine ratio, could be expected to be less harmed by the decrease in the ratio produced by aflatoxin, than would the L.A. strain, which already have a low plasma lysine/arginine ratio. If these predictions are substantiated, this information could be of value in the selection of aflatoxin resistant chickens.

#### Contrasting effects of threonine and lysine on selected amino acids

Also of interest are the effects of lysine and threonine supplements on the plasma concentrations of ornithine and methionine (Table XIV). Predictably, the 43% increase in dietary threonine (inhibits arginase) caused a significant decrease in plasma ornithine concentration in both aflatoxin and pair-fed control groups. In contrast, the 43% increase in dietary lysine (stimulates arginase) caused an accumulation of ornithine in the pair-fed control birds, but in the aflatoxin group a decrease in ornithine concentration was

Table XIV. Effect of lysine and threonine supplements on plasma levels of selected amino acids in chicks with aflatoxicosis.

Amino Acid (nmole/ml of plasma)	Dietary Supplement (% NRC requirement)	Plasma amino acids (nmol/ml)		
		Aflatoxin	Pair-fed	Ad libitum
Arginine	102 lysine	505+31A	399+11B	390+21BC
	122 lysine	468+24A	357+50BCD	386+51BC
	146 lysine	391+25BC	323+25D	344+25CD
Ornithine	102 lysine	74.0+8ABC	62.8+19BC	81.4+13AB
	122 lysine	93.6+21A	53.3+14C	71.6+12ABC
	146 lysine	69.1+5.7BC	78.8+12AB	75.5+16ABC
Methionine	102 lysine	75.1+6.7B	76.3+7.3B	80.3+4.8AB
	122 lysine	62.9+10C	86.6+5.2A	80.4+1.4AB
	146 lysine	59.1+6.5C	78.5+4.2AB	81.4+10AB
Arginine	122 threonine	553+46A	439+67BC	351+68DE
	150 threonine	532+53A	348+16DE	422+26C
	175 threonine	502+24AB	348+62E	414+45CD
Ornithine	122 threonine	133+5.3A	100+21B	56+20DE
	150 threonine	95+12B	38+7.7E	70+14CD
	175 threonine	86+7.3BC	52+22DE	76+4.6C
Methionine	122 threonine	54+5.3CD	73+9.8A	60+9.8BC
	150 threonine	52+8.6CD	54+5.0CD	68+6.2AB
	175 threonine	46+2.0D	61+8.7BC	61+4.8BC

Values (means  $\pm$  standard deviations) for a parameter followed by the same letter are not significantly different at  $P \leq 0.05$ .

Evaluation for the occurrence of interaction in the major effects is provided in Appendix Tables 23a and 24a.

produced despite the large decrease in its precursor arginine which was observed in both the aflatoxin and control groups. This suggests that ornithine is being utilized more rapidly in the aflatoxin group. Methionine concentration was also decreased in response to increased lysine in the aflatoxin group but no change in methionine levels occurred in the control groups. Two alternate pathways for the utilization of ornithine and methionine include creatine synthesis and polyamine synthesis. Because lysine is known to inhibit hepatic glycine transamidase while increasing arginase activity (79,80), stimulation of creatinine synthesis in the presence of high concentrations of lysine is unlikely (Fig. 11, p. 54). The polyamines (putrescine, spermidine, spermine, and cadaverine) are required for many important biochemical processes including protein synthesis (90). Thus the beneficial effects of lysine on the performance of chicks with aflatoxicosis may be due, at least in part, to its role in the stimulation of ornithine and/or polyamine synthesis. Similarly, the inhibition of the production of these compounds by threonine may explain its negative effects on performance.

## CHAPTER VI

## EFFECT OF LYSINE/ARGININE SUPPLEMENTATION

Introduction

In the previous chapter, increased dietary lysine was shown to moderate the toxicity (as indicated by weight gain) but not the anorexic effects of aflatoxin, and it was suggested that the strain of chicken known as Nesheim H.A., having a comparatively high plasma lysine concentration, could be expected to be less susceptible to the effects of aflatoxin. As a follow up to this experiment, a fifth experiment was designed in which varying amounts of both lysine and arginine would be supplemented. In this way we hoped to relieve the lysine/arginine antagonism which was induced at the highest level of lysine supplementation in the previous experiment, and thus study the effects of increased plasma levels of the proposed detoxicants; lysine, arginine, ornithine, and the polyamines. Unexpectedly, the chicks used in this follow up experiment were found to differ quite markedly in their plasma levels of lysine, arginine, and lysine/arginine ratio (Table XV). The chicks from the lysine/arginine feeding trial appear to resemble quite closely the HA strain developed by Nesheim, which are characterized by high plasma lysine concentration and lysine/arginine ratio and hence a high dietary arginine (H.A.) requirement. In contrast the chicks in the former lysine experiment more closely resemble the LA strain developed by Nesheim, which are characterized by low plasma lysine concentration and lysine/arginine ratio, and hence a low dietary arginine (L.A.) requirement. The plasma lysine/arginine ratio of 0.99 reported by Nesheim (91) for the L.A. strain is remarkably

Table XV. Innate differences in the plasma concentrations of lysine, arginine, and lysine/arginine ratio in chicks from feeding trials 4 and 5.

	lysine experiment			lysine/arginine experiment		
	<sup>a</sup> Ad libitum Aflatoxin	Average Control	<sup>a</sup> Ad libitum Control	<sup>b</sup> Aflatoxin	Average Control	
lysine	401 <sup>±</sup> 74	242 <sup>±</sup> 11	415	600 <sup>±</sup> 145	425 <sup>±</sup> 58	633
arginine	390 <sup>±</sup> 21	505 <sup>±</sup> 31	395	316 <sup>±</sup> 37	501 <sup>±</sup> 96	340
lys/arg	1.03	0.48	1.05	1.90	0.85	1.86

a Chicks receiving feed Ad libitum with no supplementation.

b Average of all non-aflatoxin groups receiving no lysine supplementation (40 chicks in total).

c Chickens receiving aflatoxin (2.5ug/g diet) with no additional supplementation.

d Average of all non-aflatoxin groups (180 chicks in total).

By definition, the characteristics of the ad libitum controls receiving the basal ration (no supplementation) represent the innate characteristics of the entire group of chicks in each feeding trial. Thus the contrasting lysine/arginine ratios observed in the ad libitum controls from the two separate feeding trials indicates an innate difference in the two groups of chicks. This difference is also apparent when the aflatoxin groups from both feeding trials are compared, although both ratios have been reduced by aflatoxin.

In order to confirm that the twenty ad libitum control chicks are representative of the larger group of chicks in each feeding trial, the average control values, derived from a larger proportion of the total number of chicks, are also shown. These values correspond very closely with those of the ad libitum controls.

similar to the ratio of 1.03 observed in the ad libitum control chicks in the lysine feeding trial, while the ratio of 1.90 in the ad libitum control chicks from the lysine/arginine experiment more closely resemble the ratio of 3.06 reported by Nesheim for the H.A. strain.

Although unselected commercial broilers show great variation in arginine requirement, ranging from individuals which could be described as having low arginine requirement (L.A.) to those having high arginine requirement (H.A.), it is at first difficult to explain how two separate groups of chicks could be obtained from the same supplier, which differed so widely in this regard and yet showed relative uniformity within each group. However on closer examination, the finding that chicks of the H.A. strain hatch several hours sooner than chicks from the L.A. strain when eggs are treated the same during incubation; (92) provides a plausible explanation; it can easily be imagined that early and late hatching chicks, shipped immediately after hatched, would appear in separate shipments.

#### Relationship between plasma lysine/arginine ratio and performance

From this data, and the data presented in the previous chapter showing that elevated plasma lysine concentration reduced the toxic effects of aflatoxin, one might expect that the chicks used in the lysine/arginine experiment would be resistant to the toxic effects of aflatoxin. The feed intake and weight gain of the aflatoxin and control chicks are compared in Tables XVI. and XVII. respectively. As predicted, the chicks in the lysine/arginine experiment showed an anorexic effect in response to aflatoxin, but no toxic effect; weight

Table XVI. Effect of graded levels of lysine and arginine on feed intake of chicks receiving aflatoxin.

Treatment	Dietary Supplementation (% NRC requirement)			
	94 arg 102 lys	122 arg 102 lys	122 arg 122 lys	122 arg 146 lys
	Feed intake (g)			
Aflatoxin (2.5ug/g diet)	980+22B	898+56CD	930+104BCD	892+48CD
Ad libitum controls	1115+41A	1107+60A	1084+105A	1080+43A

Values (mean  $\pm$  standard deviations) followed by the same letter are not significantly different at  $P < 0.05$ . Evaluation for the occurrence of interaction in the major effects is provided in Appendix Table 13a.

Table XVII. Effect of graded levels of lysine and arginine on weight gain of chicks receiving aflatoxin.

Treatment	Dietary Supplementation (% NRC requirement)			
	94 ARG 102 LYS	122 ARG 102 LYS	122 ARG 122 LYS	122 ARG 146 LYS
Aflatoxin (2.5ug/g diet)	481+22DE	446+25EF	498+68CD	465+17DEF
Pair-fed controls	480+11DE	430+18F	482+21DE	439+22EF
Ad libitum controls	532+64BC	574+44AB	554+41AB	586+43A

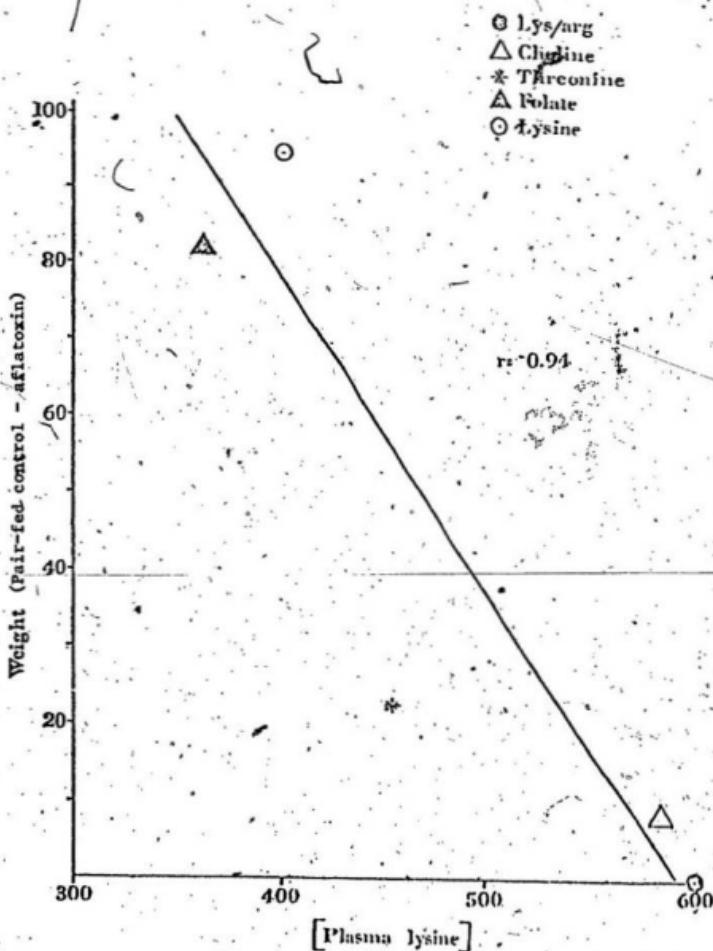
Values (mean  $\pm$  standard deviations) followed by the same letter are not significantly different at  $P \leq 0.05$ . Evaluation for the occurrence of interaction in the major effects is provided in Appendix Tables 14a.

gain is actually slightly higher in the aflatoxin group than in the pair fed controls!

Both plasma lysine concentration and the effect of aflatoxin on weight gain, varied considerably from one experiment to the next. Fig. 14 shows the relationship between these two factors. The difference in the final weight of the aflatoxin and pair fed controls receiving the basal ration, is plotted against the plasma lysine concentration in the ad libitum controls receiving the basal ration in each separate experiment. The correlation coefficient of -0.94 shows that there is a strong negative correlation between aflatoxin toxicity (as indicated by decreased weight gain over and above the decrease attributable to the anorexic effects of aflatoxin) and plasma lysine concentration, indicating that the Nesheim HA and genetically related chickens should be less susceptible to aflatoxin than the L<sub>A</sub> and related chickens.

73a

Fig. 14. Relationship between plasma lysine concentration and aflatoxin toxicity.



CHAPTER VII

SUMMARY AND CONCLUSIONS

---

Aflatoxin has been shown to produce a decrease in the plasma levels of a wide range of nutrients, in a pattern indicative of a malabsorption syndrome. Previous workers have shown that dietary aflatoxin reduces bile concentration and reduces pancreatic lipase activity. More research is necessary to show a causal relationship between these two findings. Supplementary choline, particularly when administrated through IP injections, significantly reduced the effects of aflatoxin on the majority of plasma constituents, and thus appears to relieve the aflatoxin induced malabsorption. Dietary choline also significantly reduced hepatic lipid and weight gain in the aflatoxin birds.

Although aflatoxin produced an overall reduction in plasma nutrient concentrations, specific plasma constituents were consistently increased in response to aflatoxin, most notably taurine, phenylalanine, and to a lesser extent tyrosine, together with several other urea cycle intermediates. Specific amino acid imbalances, particularly the decrease in tyrosine and BCAA/AAA may contribute to suppressed feed intake.

Previous researchers have observed daily benzoic acid administration to increase urea nitrogen from 1% to 9% of the total nitrogen excreted by chickens. This rise in urea production was used to indicate the operation of the ornithine detoxification mechanism. The aflatoxin induced increase in plasma levels of BUN, ammonia, glutamine and the urea cycle intermediates, together with the decreased plasma concentration of uric acid, suggests a similar increased excretion of nitrogen as urea in response to daily aflatoxin.

administration, and thus suggests a similar reliance on the ornithine detoxification mechanism.

The contrasting effects of lysine and threonine on the performance of chicks with aflatoxicosis may be related to their opposing effects on arginase, a key enzyme in the ornithine detoxification system. Regardless of the mechanism of the beneficial effects of lysine, genetic variations in plasma lysine concentration may be of value in the selection of aflatoxin resistant chickens.

77

REFERENCES

1. Davis N.D. & Diener U.L. (1970) Environmental factors affecting the production of aflatoxin. In: Proceedings of the First U.S.-Japan Conference on Toxic Microorganisms, pp.43-47, U.S. Dept. of the Interior, Washington D.C.
2. Smith, J.W. & Hamilton, P.B. (1970) Aflatoxicosis in the broiler chicken. *Poultry Sci.* 49, 207-215.
3. Patterson, D.S.P. (1977) Chemistry of mycotoxins: Biochemistry and physiology. In: Mycotoxic Fungi and Chemistry of Mycotoxins, (Wyllie, T.D. and Morehouse L.G., Ed.), Marcel Dekker Inc., New York, New York.
4. Patterson, D. S. P. & Roberts, B. A. (1979) Mycotoxins in animal feedstuffs: Sensitive thin layer chromatographic detection of aflatoxin, ochratoxin A, sterigmatocystin, zearalenone, and T-2 toxin. *JAOAC* 62:1265-1267.
5. Stubblefield, R. D. (1979) The rapid determination of aflatoxin M<sub>1</sub> in dairy products. *JAOCS* 56:800-802.
6. Shotwell, O.L., Hesseltinge, C.W., Stubblefield, R.D. & Sorenson, W.G. (1966) Production of aflatoxin on rice. *Appl. Microbiol.* 14:425-428.
7. West, S., Wyatt, R.D. & Hamilton, P.B. (1973) Improved yield of aflatoxin by incremental increases in temperature. *Appl. Microbiol.* 25:1018-1019.
8. Nabney, J. & Nesbitt, B.F. (1965) A spectrophotometric method of determining the aflatoxins. *Analyst* 90:155-160.
9. Wiseman, H.G. Jacobson, W.C. & Harmeyer, W.C. (1967) Note on removal of pigments from chloroform extracts of aflatoxin cultures with copper carbonate. *J. AOAC* 50:982-983.
10. Gitelman, H.J. (1967) An improved automated procedure for the determination of calcium in biological specimens. *Anal. Biochem.* 18:520-531.
11. Kraml, M. (1966) A semi-automated determination of phospholipids. *Clin. Chim. Acta* 13:442-448.

12. Persign, J.P., Vander Slik, W. & Riethorst, A. (1971) Determination of serum iron and latent iron-binding capacity (LIBC). *Clin. Chim. Acta* 35:91-98.
13. Neeley, W.E. (1972) Simple automated determination of serum or plasma glucose by hexokinase/glucose-6-phosphate dehydrogenase method. *Clin. Chem.* 18:509-515.
14. Sampson, E.J., Demers, L.M. & Krieg, A.F. (1975) Faster enzymatic procedure for serum triglycerides. *Clin. Chem.* 21:1983-1985.
15. Eggstein, M. & Kuhlman, E. (1974) Triglycerides and glycerol: Determination after alkaline hydrolysis. In: Methods in Enzymatic Analysis, (Bergmeyer, H.U., ed.), Vol.4, pp.1825-1831, Verlag Chemie, Weinheim/Academic Press Inc., New York.
16. Allain, C.C., Poon, L., Chan, S.G., Richard, W. & Fu, P. (1974) Enzymatic determination of total serum cholesterol. *Clin. Chem.* 20:470-475.
17. Goodwin, J., Baginski, E. & Zak, B. (1965) Simultaneous automated determination of serum albumin and total protein. In: Automation in Analytical Chemistry, (Skeggs, L.T., ed.), pp.563-568, Technicon Symposia, Mediad Incorporated, New York.
18. Doumas, B.T., Watson, W. & Briggs, H.G. (1971) Albumin standards and the measurement of serum albumin with bromcresol green. *Clin. Chim. Acta* 31:87-96.
19. Marsh, W.H., Fingerhut, B. & Miller, H. (1965) Automated and manual direct methods for the determination of blood urea. *Clin. Chem.* 11:624-627.
20. Sobrinho-Simoes, M. (1965) A sensitive method for the measurement of uric acid using hydroxylamine. *J. Lab. Clin. Med.* 65:665-668.
21. Gambino, S.R., & Schreiber, H. (1964) The measurement and fractionation of bilirubin on the autoanalyzer by the method of Jendrassik and Grof. In: Automation in Analytical Chemistry, (Skeggs, L.T., ed.) Technicon Symposia, Mediad Incorporated, New York.
22. Chasson, A.L., Grady, H.T. & Stanley, M.A. (1961) Determination of creatine by means of automatic chemical analysis. *Am. J. Clin. Pathol.* 35:83-88.

23. Morgenstern, S., Kessler, G., Auerbach, J., Flor, R. & Klein, B. (1965) An automated p-nitrophenyl phosphate serum alkaline phosphatase procedure for the autoanalyzer. Clin. Chem. 11, 876-888.
24. Morgenstern, S., Flor, R., Kessler, G. & Klein, B. (1966)-The automated determination of NAD-coupled enzymes, Part II. serum lactic dehydrogenase. Clin. Chem. 12: 274-281.
25. Scheidt, R.A., Nelson, V.A. & Levine, J.B. (1965) Automated determination of serum glutamic oxalacetic transaminase. In: Automation in Analytical Chemistry, (Skegg, L.T., ed.), pp.563-568, Technicon Symposia, Mediad Incorporated, New York
26. Henry, R.J., Chiamori, N., Golub, O.J. & Berkman, S. (1960) Revised spectrophotometric methods for the determination of glutamic-oxalacetic transaminase, glutamic-pyruvic transaminase and lactic acid dehydrogenase. Am. J. Clin. Path. 34:381-398.
27. Fraser, I.H., & Mookerjea, S. (1977) Purification of membrane-bound galactosyltransferase from rat liver microsomal fractions. Biochem. J. 164:541-547.
28. Folch, J., Lees, M. & Sloane-Stanley, G.H. (1957)- A simple method for the isolation and purification of total lipids from animal tissues. J. Biol. Chem. 226:497-509.
29. McIndoe, W. M., & Mitchell, G. G. (1978) Lactate dehydrogenase isozymes in the spermatozoa of the domestic fowl, Gallus domesticus and turkey, Meleagris gallopavo. Comp. Biochem. Physiol. 61B:433-437.
30. Blackburn, S. (1978) Destruction of amino acids. In: Amino Acid Determination Methods and Techniques, pp.10-15, Marcel Dekker, New York.
31. Fischer, J.E., Funovics, J.M., Aguirre, A., James, J.H., Keane, J.M., Wessdorp, R.I.C., Yoshimura, N. & Westman, T. (1975). The role of plasma amino acids in hepatic encephopathy. Surgery 78:276-290.
32. Hamilton, P.B., (1977) Interrelationships of mycotoxins with nutriton. Fed. Proc. 36:1899-1902.
33. Luthy, J., Zweifel, U. and Schlatter, U. (1980) Metabolism and tissue distribution of [<sup>14</sup>C] aflatoxin-B<sub>1</sub>, in pigs. Fd. Cosmet. Toxicol. 18:253-256.

34. Sawhney, D.S., Vadehra, D.V. & Baker, R.C. (1973) The metabolism of [<sup>14</sup>C] aflatoxins in laying hens. *Poultry Sci.* 52:1302-1309.
35. Mabee, M.S. & Chipley, J.R. (1973) Tissue distribution and metabolism of aflatoxin-B1 - [<sup>14</sup>C] in broiler chickens. *Appl. Microbiol.* 25: 763-769.
36. Garlich, J.D., Tung, H-T & Hamilton P.B. (1973) The effect of short term feeding of aflatoxin on egg production and some plasma constituents of the laying hen. *Poult. Sci.* 52:2206-2211.
37. Brown, J.M.M. & Abrams, L. (1965) Biochemical studies on aflatoxicosis. *Onderstepoort J. Vet. Res.* 32:119-146.
38. Clark, J.D., Jain, A.V., Hatch, R.C. & Mahaffey, E.A. (1980) Experimentally induced chronic aflatoxicosis in rabbits. *Am. J. Vet. Res.* 41:1841-1845.
39. Wyatt, R.D., Briggs, D.M. & Hamilton, P.B. (1973) The effect of dietary aflatoxin on mature broiler breeder males. *Poultry Sci.* 52:1119-1123.
40. Mohiddin, S.M., Mahendranath, D., Yadgiri, B. & Ahemed, S.R. (1981). Studies on the effects of aflatoxin on antibody synthesis against Ranikhet disease vaccine in chicks. *Indian J. Animal Sci.* 51:77-82.
41. Mattenheimer, H. (1971) Mattenheimer's Clinical Enzymology Principles and Applications. pp.109, Ann Arbor Science Publishers, Ann Arbor, Michigan.
42. Hamilton, P.B., Tung, H-T., Wyatt, R.D. & Donaldson, W.E. (1974) Interaction of dietary aflatoxin with some vitamin deficiencies. *Poultry Sci.* 53:871-877.
43. Lanza, G.M., Washburn, K.W., Wyatt, R.D. & Edwards, H.M. (1979) Depressed [<sup>59</sup>Fe] absorption due to dietary aflatoxin. *Poultry Sci.* 58:1439-1444.
44. Preston, J.A. (1971) Biochemical Profiling in Diagnostic Medicine, Volume 1, Technicon Instrument Corporation, Tarrytown, New York.
45. Osborne, D.J. & Hamilton, P.B. (1981) Decreased pancreatic digestive enzymes during aflatoxicosis. *Poultry Sci.* 60:1818-1821.

46. Osborne, D.J. & Hamilton, P.B. (1981) Steatorrhea during aflatoxicosis in chickens. *Poultry Sci.* 60:1398-1402.
47. Voigt, M.N., Wyatt, R.D., Ayres, J.G. & Koehler, F.E. (1980) Abnormal concentrations of B vitamins and amino acids in the plasma, bile and liver of chicks with aflatoxicosis. *Appl. Environ. Microbiol.* 40:870-875.
48. Rosen, H.M., Yoshimura, N., Hodgman, J.M. & Fischer, J.E. (1977) Plasma amino acid levels in hepatic encephalopathy of differing etiology. *Gastroenterology* 72:483-487.
49. Strombeck, D.R. & Rogers, Q. (1978) Plasma amino acid concentrations in dogs with hepatic disease. *J. Am. Vet. Med. Assoc.* 173:93-96.
50. Pardridge, W.M. (1977) Genetics of competitive inhibition of neural amino acid transport across the blood brain barrier. *J. Neurochem.* 28:103-108.
51. Ikeda, M., Levitt, M. & Udenfriend, S. (1967) Phenylalanine as substrate and inhibitor of tyrosine hydroxylase. *Arch. Biochem. Biophys.* 120:420-427.
52. Dodsworth, M.M., James, J.H., Cummings, M.C. & Fischer, J.E. (1974) Depletion of brain norepinephrine in acute hepatic coma. *Surgery* 75:811-820.
53. Fernstrom, J.D. (1978) The effects of tryptophan and diet on brain serotonin synthesis and release. In: *Depressive Disorders*, (Garattini, S. ed.), vol. 13. pp. 107-128, *Symposia Medica Hoechst*, F.K. Schattauer Verlag, New York.
54. Anderson, G.H. (1977) Regulation of protein intake by plasma amino acids. In: *Advances in Nutrition Research*, Vol 1, pp.145-166, Plenum Press, New York.
55. Bassir, O. & Osiyemi, F. (1967) Biliary excretion of aflatoxin in the rat after a single dose. *Nature (London)* 215:882.
56. Wilson, H.R., Douglas, C.R., Harms, R.A. & Edds, G.T. (1975) Reduction of aflatoxin effects on quail. *Poult. Sci.* 54:923-925.
57. Donaldson, W.E., Tung, H-T. & Hamilton, B.P.- (1972) Depression of fatty acid synthesis in chicken liver (*Gallus domesticus*) by aflatoxin. *Comp. Biochem. Physiol.* 41B:843-847.

58. Hamilton, P.B. (1975) Lipid and vitamin metabolism during aflatoxicosis. In: Microbiology, (Schlessinger, D., ed.), pp. 381-387, American Society for Microbiology, Washington D.C..
59. Hamilton, P.B. & Harris, J.R. (1971) Interaction of aflatoxin with Candida albicans infections and other stresses in chickens. *Poultry Sci.* 50:906-912.
60. Smith J.W., Hill C.H., & Hamilton P.B. (1971) The effect of dietary modifications on aflatoxicosis in the broiler chicken *Poultry Sci.* 50:768-774.
61. Knake, R.P., Rao, C.S. & Deyoe, C.W. (1973) Effects of feeding diets containing aflatoxin and added vitamins to *Coturnix* quail. *Poultry Sci.* 52:2050.
62. Campbell, T.C., Hayes, J.R. & Newberne, P.M. (1978) Dietary lipotropes, hepatic microsomal mixed function oxidase activities; and *in vivo* covalent binding of aflatoxin B1 in rats. *Cancer Res.* 38:4569-4573.
63. Anon. (1981) Folic acid shown to help pigs cope with moldy grain; gains improved. *Feedstuffs* 53:No.11:13.
64. Wogan, G.N., Edwards, G.S. & Shank, R.C. (1967) Excretion and tissue distribution of radioactivity from aflatoxin B1-[<sup>14</sup>C] in rats. *Cancer Res.* 27:1729-1736.
65. Tung, H.T., Cook, F.W., Wyatt, R.D. & P.B. Hamilton, (1976). The anemia caused by aflatoxin *Poultry Sci.* 54:1962-1969.
66. Cheville, N.F. (1979) Environmental factors affecting the immune response of birds - A review. *Avian Dis.* 23:308-314.
67. Ueno, I., Friedman, L., & Stone, C.L. (1980) Species differences in the binding of aflatoxin B1 to hepatic macromolecules. *Toxicol. Appl. Pharmacol.* 52:177-180.
68. Hamilton, P.B. & Garlich, J.D. (1972) Failure of vitamin supplementation to alter the fatty liver syndrome caused by aflatoxin. *Poultry Sci.* 51:688-692.
69. Park, L.E., Voigt, M.N., Fraser, I.H. & Davidson, W.S. (1983) Influence of dietary folate or dietary and interperitoneal administration of choline on free-amino acids and biochemical parameters in broiler chicks (Hubbard/hubbard) with aflatoxicosis. Submitted to: *Toxicol. Appl. Pharm.*

70. Tung, H-T., Donaldson, W.E. & Hamilton, P.B. (1972) Altered lipid transport during aflatoxicosis. *Toxicol. Appl. Pharmacol.* 22:97-104.
71. Hamilton, P.B. (1977) Interrelationships of mycotoxins with nutrition. *Fed. Proc.* 36:1899-1902.
72. Pier, A.C., Richard, J.L. & Thurstan, J.R. (1980) Effects of aflatoxin on the mechanisms of immunity and native resistance. *Med. Mycology, Zbl. Bakt. Suppl.* 8:301-309.
73. Rogers, Q.R. & Leung P.M.B. (1973) The influence of amino acids on the neuroregulation of food intake. *Fed. Proc.* 32:1709-1719.
74. Harper, A.E., Benevenga, N.J., & Wohlhueter, R.M. (1970) Effects of ingestion of disproportionate amounts of amino acids. *Physiol. Rev.* 50:428-558.
75. Ryan, W.L., Carver, M.J. (1964) Inhibition of antibody synthesis by L-phenylalanine. *Science* 143:479-480.
76. Roscoe, J.P., Eaton, M.D., & Gladys, C.C. (1968) Inhibition of protein synthesis in Kerbs 2 Ascites cells and cell-free systems by phenylalanine and its effect on leucine and lysine in the amino acid pool. *Biochem. J.* 109:507-515.
77. Best, C.H., Lucas, C.C., & Ridout, J.H. (1956) Vitamins and the protection of the liver. *British Med. Bull.* 12:9-14.
78. Harper, A.E. (1958) Nutritional fatty livers in rats. *Amer. J. Clin. Nutr.* 6:242-253.
79. Austic, R.E. & Nesheim, M.C. (1970) Role of kidney arginase in variations of the arginine requirement of chicks. *J. Nutr.* 100:855-868.
80. Austic, R.E. & Scott, R.L. (1975) Involvement of food intake in the lysine-arginine antagonism in chicks. *J. Nutr.* 105:1122-1131.
81. Tamir, H. & Ratner, S. (1963) A study of ornithine, citrulline and arginine synthesis in growing chicks. *Arch. Biochem. Biophys.* 102:259-269.
82. Sykes, A.H. (1971) Formation and composition of urine. In: *Physiology and Biochemistry of the Domestic Fowl*, Ed. 4, (Bell, D.J. and Freeman, B.M., ed.), Vol.1, pp. 274 Academic Press, New York.

83. Crowdle, J.H. & Sherwin, C.P. (1923) The chemical defence mechanism of the fowl. *J. Biol. Chem.* 55:15-31.
84. Seiler, N. (1979) Amide-bond-forming reactions of polyamines. In: *Polyamines in Biology and Medicine*, (Morris, D.R., & Marton, L.J., ed.), Marcel Dekker Inc., New York.
85. Neal, G.E., Judah, D.J. Stripe, F. & Patterson, D.S.P. (1981) The formation of 2,3-dihydroxy-2,3-dihydro-aflatoxin B1 by the metabolism of aflatoxin B1 by liver microsomes isolated from certain avian and mammalian species and the possible role of this metabolite in the acute toxicity of aflatoxin B1. *Tox. Appl. Pharmacol.* 58:431-437.
86. Jones, J.D. (1964) Lysine-arginine antagonism in the chick. *J. Nutr.* 84:313-321.
87. Jones, J.D., Petersburg, S.J., & Burnett, P.C. (1967) The mechanism of the lysine-arginine antagonism in the chick: Effect of lysine on digestion, kidney arginase, and liver transaminidinase. *J. Nutr.* 93:103-116.
88. Nesheim, M.C. (1968) Genetic variation in arginine and lysine utilization. *Fed. Proc.* 27:1210-1214.
89. Nesheim, M.C. (1967) Kidney arginase activity and lysine tolerance in strains of chicks selected for high or low requirement of arginine. *J. Nutr.* 95:79-87.
90. Loftfield, R.B., Eigner, E.A., & Pastuszyn, A. (1981) Polyamines and protein synthesis. In: *Polyamines in Biology and Medicine*, (Morris, D.R., and Marton, L.J. ed.) Marcel Dekker Inc, New York.
91. Nesheim, M.C. (1967) Kidney arginase activity and lysine tolerance in strains of chicks selected for high or low requirement of arginine. *J. Nutr.* 95:79-87.
92. Nesheim, M.C., Austic, R.E. & Wang S-H. (1971) Genetic factors in lysine and arginine metabolism of chicks. *Fed. Proc.* 30:121-126.

**APPENDIX A****RAW DATA TABLES 1--25**

App. 1. Effect of dietary or interperitoneal (IP) administration of choline on the weight gain of chicks with atelastinosis.

Day	Weight Gain (g/chick) <sup>a</sup>									
	Atelastin (1.5% w/w diet)			Restricted Feeding <sup>b</sup>			Control			
	0	IP	Dietary	0	IP	Dietary	0	IP	Dietary	0
1	40±1.56	39±0.56	42±1.04	41±1.74	40±1.56	41±1.74	41±2.04	41±1.74	29±1.04	41±2.04
2	48±1.04	46±1.24	47±1.24	44±1.04	44±1.04	45±1.44	43±1.44	43±1.44	43±1.44	43±1.44
3	62±1.44	62±1.44	62±1.44	62±1.14	62±1.14	62±1.44	62±1.44	62±1.44	62±1.44	62±1.44
4	72±1.44	71±2.14	70±1.34	72±2.14	72±2.14	72±2.14	72±2.14	72±2.14	72±2.14	72±2.14
5	87±2.34	85±2.74	90±2.34	87±2.74	87±2.74	86±2.34	86±2.34	86±2.34	86±2.34	86±2.34
6	101±2.58	101±2.58	108±1.68	102±2.58	102±2.58	110±2.68	110±2.68	105±2.74	99±2.24	112±2.68
7	119±1.04	117±2.24	117±2.24	117±2.24	117±2.24	112±2.24	112±2.24	112±2.24	112±2.24	112±2.24
8	141±4.86C	132±2.40B	141±2.14B	133±4.14C	130±2.38C	130±2.38C	144±2.44BNC	133±2.44BNC	130±2.44BNC	134±2.44B
9	151±2.34BC	149±2.34C	141±2.04BC	154±2.44BC	144±2.34BC	143±2.14BC	151±2.34BC	143±2.34BC	141±2.34BC	141±2.34BC
10	146±2.04BC	132±2.44C	149±2.44A	137±2.04C	137±2.04C	137±2.14C	140±2.44BC	137±2.44BC	140±2.44BC	140±2.44BC
11	185±2.04BC	185±2.40B	187±2.40A	181±2.44B	181±2.44B	175±2.44D	181±2.44B	181±2.44B	179±1.24B	181±2.44B
12	211±2.34AB	194±1.90	216±1.30A	205±2.54B	205±2.54B	216±1.84B	210±1.84B	210±1.84B	197±2.04B	212±2.44A
13	224±1.84A	213±1.14A	232±2.04A	223±1.64A	223±1.64A	219±1.64A	246±2.24B	217±2.04A	213±2.04A	234±2.24A
14	246±1.70E	241±1.04E	212±2.04A	234±1.54E	234±1.54E	234±1.54E	261±1.84C	261±1.84C	254±1.74E	272±1.94A
15	264±2.18	269±1.48	272±2.04B	262±1.64B	262±1.64B	269±2.14B	289±2.44B	289±2.44B	300±1.44A	291±1.64B
16	290±1.80	287±1.50	327±2.04B	208±1.70	209±1.70	295±1.64B	316±2.14B	322±1.64B	312±2.04A	312±2.04A
17	311±1.84C	311±1.84C	316±2.04B	286±2.74C	286±2.74C	286±2.74C	344±2.34BNC	344±2.34BNC	341±2.34BNC	341±2.34BNC
18	359±2.74C	360±1.70	360±1.84B	362±1.14C	362±1.14C	362±1.14C	362±1.50C	362±1.50C	362±1.50C	362±1.50C
19	379±1.80	362±1.50	366±2.54C	392±2.44C	392±2.44C	375±2.44B	407±2.44B	404±2.44B	420±2.44B	427±2.44A
20	439±2.74C	393±1.48	443±2.14B	424±1.04C	424±1.04C	408±2.34B	444±2.14B	441±2.14B	469±3.04C	513±2.24A
21	427±2.24EF	386±2.14F	471±1.90C	461±1.64E	461±1.64E	423±2.14F	508±2.84B	515±2.84B	492±2.84B	536±2.84A
22	451±2.60	468±1.36	494±1.24B	478±1.54C	478±1.54C	520±2.44D	540±2.44D	540±2.44D	507±2.84C	460±2.34A
23	493±2.44E	442±1.76	538±3.10C	503±2.04E	503±2.04E	491±2.44E	541±3.04C	542±2.44E	561±2.34C	442±2.34A
24	511±2.30	480±2.10D	603±2.84C	530±2.40	499±2.30	616±3.54C	589±3.04B	589±3.04B	587±2.34C	643±2.34A

<sup>a</sup> Values (mean ± standard deviation) for weight gain followed by the same letter are not significantly different at P < 0.05.

<sup>b</sup> Chicks fed to the feed intake of the corresponding atelastinosis group.

IP = supplemental choline.

Atelastinosis to achieve 175% of the NRC requirement (1977) of choline (choline ration provided 165%).

App. 1c. Major and interactive effects produced by dietary or intraperitoneal administration of cholera on weight gain of chicks with atlatonosis.

Day		Atlatonosis <sup>b</sup>	Effect of feeding		Effect of administration		ANOVA (FDr. 27)†	
			At latonosis <sup>b</sup>		Restricted Ad libitum		Feed	Admin
			Feed	Dietary <sup>c</sup>	Ipo	Dietary <sup>c</sup>	Feed	Admin
1	-	-	-A	-A	-A	-A	0.010(.897)	1.87(0.14)
2	-	-	-A	-A	-A	-A	1.71(0.20)	0.67(0.87)
3	-	-	-A	-A	-A	-A	1.08(0.36)	0.64(0.53)
4	-	-	-A	-A	-A	-A	1.09(0.35)	0.85(0.90)
5	-	-	-A	-A	-A	-A	0.40(0.87)	0.44(0.78)
6	-	-	-A	-A	-A	-A	0.55(0.38)	0.41(0.80)
7	-	-	-A	-A	-A	-A	0.010(.97)	0.42(0.40)
8	-	-	-A	-A	-A	-A	1.10(0.35)	1.84(0.001)
9	-	-	-A	-A	-A	-A	9.20(0.008)	0.81(0.30)
10	-	-	-A	-A	-A	-A	0.52(0.40)	0.11(0.38)
11	-	-	-A	-A	-A	-A	1.27(0.30)	1.17(0.002)
12	-	-	-A	-A	-A	-A	0.25(0.78)	7.08(0.004)
13	-	-	-A	-A	-A	-A	0.08(0.92)	5.54(0.005)
14	-	-	-A	-A	-A	-A	5.26(0.012)	0.26(0.40)
15	-	-	-A	-A	-A	-A	6.74(0.004)	0.44(0.41)
16	-	-	-A	-A	-A	-A	8.21(0.016)	0.71(0.39)
17	-	-	-A	-A	-A	-A	6.19(0.006)	0.37(0.44)
18	-	-	-A	-A	-A	-A	17.48(0.001)	0.48(0.73)
19	-	-	-A	-A	-A	-A	7.51(0.002)	0.12(0.97)
20	-	-	-A	-A	-A	-A	17.51(0.001)	0.12(0.97)
21	-	-	-A	-A	-A	-A	18.46(0.001)	0.84(0.65)
22	-	-	-A	-A	-A	-A	31.1(0.001)	0.11(0.98)
23	-	-	-A	-A	-A	-A	26.7(0.001)	0.11(0.98)
24	-	-	-A	-A	-A	-A	25.1(0.001)	0.01(0.41)
							26.7(0.001)	0.53(0.72)
							27.2(0.001)	0.27(0.89)

Means for the gain in weights within a major effect followed by the same letter are not significantly different at P<0.05.

Majority atlatonosis 2.35g per g of diet.

Calculated to the intake of the corresponding atlatonosis group.

No supplemental chlorine.

Administration to achieve 135% of the NRC requirement (1977) of chlorine (baud ration provided 100%).

Table - Major effect of administration. Feed = major effect of feeding. Feed x Admin = test for interaction.

App. 2. Effect of dietary or interperitoneal (IP) administration of choline on the feed intake of chicks with allantoisosis.

Day	Allantoin (2.5 % of diet) <sup>a</sup>			Feed intake (g/chick) <sup>b</sup>			Control	
	Restricted Feeding			Unrestricted Feeding				
	0 <sup>c</sup>	1/2 <sup>d</sup>	1 <sup>e</sup>	0	1/2	1		
1	0.000/0.04	0.000/0.04	0.000/0.04	0.000/0.04	0.000/0.04	0.000/0.04	0.000/0.04	
2	8.5±1.7 <sup>f</sup>	7.5±2.1 <sup>f</sup>	7.0±1.6 <sup>f</sup>	7.5±2.1 <sup>f</sup>	7.5±2.1 <sup>f</sup>	7.5±2.1 <sup>f</sup>	8.5±2.1 <sup>f</sup>	
3	215.2±4.6C	187.5±2.8C	191.6±2.8C	204.9±3.1A	184.9±2.8A	224.1±1.8B	132.1±1.7C	
4	267.3±3.9	204.9±3.9	218.9±3.1A	432.6±5.4	432.6±5.4	394.7±5.4	394.7±5.4	
5	244.4±3.6	47.9±4.9A	55.0±1.9A	63.9±2.2A	56.8±2.2A	52.0±1.0A	46.6±1.1A	
6	265.5±3.4	62.5±5.4A	76.2±3.4A	81.9±4.4A	73.9±4.4A	72.9±4.4A	64.6±5.4A	
7	100.2±3.4	87.5±5.4A	101.0±2.6A	101.0±2.6A	94.7±5.4A	94.7±5.4A	81.6±5.4A	
8	124.9±3.6A	114.9±5.0A	118.2±5.0A	124.9±1.6B	115.9±5.2B	120.9±5.4B	110.0±5.4B	
9	134.0±2.6	141.9±7.0A	136.2±9.0A	152.1±1.6A	147.9±2.0A	147.9±2.0A	130.0±2.0B	
10	183.9±7.6A	183.9±7.6A	184.5±7.6A	187.9±1.6C	187.9±1.6C	176.1±2.0A	183.9±7.6A	
11	217.9±5.6A	197.2±5.2B	215.2±5.2A	192.4±4.4A	192.4±4.4A	210.9±2.0B	226.1±7.9A	
12	248.0±5.4A	212.0±5.1B	241.5±5.1A	246.1±5.1A	220.6±5.1A	233.9±5.1B	248.0±5.4A	
13	289.1±4.2C	30.5±2.1BC	76.9±2.8BC	77.0±2.8BC	51.0±2.0C	39.9±2.8AB	50.2±1.4C	
14	320.1±4.4C	27.9±2.4C	31.1±4.1BC	31.1±4.1BC	29.4±2.1BC	33.5±2.0AB	37.7±2.1A	
15	360.4±4.6C	31.5±2.7B	35.5±4.7BCD	34.5±4.7BCD	31.4±2.5CD	33.9±2.5CD	38.4±2.3AB	
16	399.1±7.6CD	37.8±4.4CD	38.5±4.2CD	38.5±4.2CD	37.4±2.1CD	41.5±4.0BC	41.8±2.2A	
17	442.6±2.8CD	20.2±2.4B	44.0±5.7BCD	42.6±5.7BCD	39.7±2.9CD	41.6±2.8BC	47.4±2.2A	
18	492.2±8.0CD	43.5±2.8CD	49.0±5.4BCD	48.9±5.4BCD	42.7±2.1CD	44.5±3.0BC	44.1±2.3A	
19	545.2±8.0CD	47.2±2.2D	54.7±2.7ABC	51.9±2.0CD	48.9±2.0D	51.8±2.1CD	57.1±5.4BC	
20	601.2±9.0C	52.8±5.4D	60.1±2.9BC	57.1±2.9CD	50.8±2.4D	57.4±4.0CD	64.6±5.4BC	
21	632.4±4.6C	57.2±5.0D	63.4±5.2BC	62.8±4.3DC	56.3±2.4D	63.1±4.2CD	66.6±5.0AB	
22	704.5±5.2CD	67.2±5.0B	71.1±5.0C	67.9±5.1CD	60.5±2.7E	67.6±5.6BCD	74.9±5.8AB	
23	770.5±5.6CD	67.5±5.0B	73.0±5.0C	73.0±5.0C	61.4±2.9E	73.0±5.0C	85.2±5.4AB	
24	832.9±8.0CD	77.8±5.0E	83.4±5.2BC	79.8±5.2CD	71.0±2.2E	79.1±5.2CD	92.5±5.8A	

<sup>a</sup> Values (mean<sup>b</sup> standard deviation) for feed intake followed by the same letter are not significantly different at  $P < 0.05$ .

<sup>b</sup> Fed to the feed intake of the corresponding allantois group.

<sup>c</sup> No supplemental choline.

<sup>d</sup> Administration to achieve 175% of the BRC requirement (1977) of choline (heat ration provides 105%).

App. 2a. Major and interactive effects produced by dietary or intraperitoneal (IP) administration of choline on feed intake of chicks with efflaccitosis.

Day	Effect of feeding		Effect of supplemental choline		ANOVA ( $F_{(r \times f) \times f}$ ) <sup>a</sup>	
	Adolescent Reactive <sup>b</sup>	Adult <sup>b</sup>	Adolescent Reactive <sup>b</sup>	Adult <sup>b</sup>	Feed	Adolesc. x Admin
Weight gain (g/chick) <sup>c</sup>						
2	-A	-A	-A	-A	0.44(0.32)	0.75(0.48)
3	-A	-A	-A	-A	4.59(0.019)	2.31(0.040)
4	-A	-A	-A	-A	3.48(0.045)	2.52(0.059)
5	-A	-A	-A	-A	1.63(0.21)	1.07(0.39)
6	-A	-A	-A	-A	0.40(0.67)	1.72(0.18)
7	-A	-A	-A	-A	0.01(0.99)	1.56(0.22)
8	-A	-A	A	A	0.51(0.39)	2.38(0.11)
9	-A	-A	A	A	1.31(0.39)	1.77(0.16)
10	-A	-A	A	A	1.44(0.25)	2.34(0.12)
11	20.948	20.228	22.11A	18.64A	3.11(0.06)	0.47(0.76)
12	23.718	23.238	21.53A	19.88	3.16(0.059)	3.72(0.037)
13	27.658	26.228	29.81A	22.08	4.52(0.0091)	0.54(0.53)
14	30.318	29.88	34.04A	32.23A	5.05(0.0010)	1.44(0.14)
15	34.318	32.028	39.84A	32.48	4.40(0.0011)	4.81(0.016)
16	39.218	37.078	44.11A	40.15B	12.31(0.0001)	5.21(0.056)
17	42.518	41.028	49.16A	45.04A	14.10(0.0001)	4.91(0.035)
18	47.718	45.518	55.08A	50.08A	17.60(0.0001)	6.23(0.039)
19	51.318	50.118	61.24A	53.54A	20.10(0.0001)	0.81(0.53)
20	57.718	55.118	67.98A	63.13A	5.77A	7.91(0.0001)
21	62.818	60.718	72.94A	68.94A	21.26(0.0001)	0.64(0.63)
22	68.018	65.548	81.72A	72.84A	19.31(0.0001)	0.54(0.48)
23	73.718	70.918	87.56A	78.04A	23.40(0.0001)	8.75(0.001)
24	79.718	76.658	94.64A	85.04A	20.26(0.0001)	0.64(0.64)

<sup>a</sup>Means for feed intakes within a major effect followed by the same letter are not significantly different at P<0.05.

Dietary efflaccitosis = 2.5g per g of diet.

Efflaccitosis = the state of the corresponding efflaccitosis group.

No supplemental choline.

Administration to achieve 17% of the NRC requirement (1971) of choline (water solution provided IP).

Adolesc. = major effect of administration. Feed = major effect of feeding. Admin = test for interaction.

App. 3. Effect of dietary or interpetitonal (IP) administration of choline on the feed conversion of chicks with ascites.

Day	Feed conversion ratio <sup>a</sup>			Control	
	Restricted Feeding <sup>b</sup>				
	0	IP	Dietary		
1	1.2921-1.14	1.5450-1.11A	0.8112-0.08	3.0942-0.08	
2	1.0000-0.20A	0.4820-1.18A	1.1410-1.04A	0.8102-0.08	
3	1.2920-1.29A	0.9710-1.06A	1.2850-1.23A	0.8040-0.15A	
4	1.2640-1.28A	0.9710-1.06A	1.4320-1.20A	0.8040-0.15A	
5	1.2780-1.12A	1.0160-1.09A	1.3200-1.21A	1.2660-1.21A	
6	1.2650-1.12A	1.0750-1.17A	1.2650-1.21A	1.2250-1.19A	
7	1.2720-1.14	1.2050-1.20A	1.3050-1.10A	1.2050-1.13A	
8	1.2320-0.98A	1.2050-1.12A	1.2640-1.14A	1.2050-1.12A	
9	1.2520-1.25A	1.2520-1.25A	1.2850-1.18A	1.2520-1.25A	
10	1.2460-1.17A	1.4020-0.93AB	1.2050-1.11B	1.4520-0.97AB	
11	1.2320-1.22A	1.5520-1.08A	1.2050-1.04A	1.5110-0.97A	
12	1.2420-0.94B	1.4520-1.22A	1.2050-1.04A	1.4420-0.92AB	
13	1.5720-1.17AC	1.5250-1.34BC	1.3710-0.78BC	1.4740-0.98BC	
14	1.2640-1.35A	1.3840-0.97BCD	1.3520-1.11CD	1.4520-0.96AD	
15	1.5460-1.28A	1.3730-0.93BCD	1.4320-1.32C-C	1.4420-0.97AC-C	
16	1.4020-1.17A	1.5250-1.22AB	1.2910-1.22AB	1.4710-0.98AB	
17	1.5860-1.24A	1.4440-1.12A	1.5110-1.21A	1.4460-0.94A	
18	1.5320-1.04A	1.4520-1.21A	1.4450-1.10A	1.4450-0.97A	
19	1.4140-1.04A	1.4710-1.18A	1.5520-1.04A	1.4740-0.97A	
20	1.5220-1.24A	1.5020-1.09A	1.5020-1.04A	1.4740-0.96A	
21	1.4820-1.23A	1.6110-1.04A	1.5320-1.04A	1.4910-1.04A	
22	1.2780-1.17A	1.4630-1.04A	1.5730-1.04C	1.5320-1.04NC	
23	1.5700-1.18A	1.5860-1.10AB	1.5460-1.08AB	1.6000-1.10AB	
24	1.2720-1.18A	1.4660-1.04A	1.4950-1.04A	1.6250-0.98AB	

<sup>a</sup> Values (mean  $\pm$  standard deviation) for feed conversion (feed intake/weight gain) followed by the same letter are not significantly different at  $P \leq 0.05$ .

<sup>b</sup> pair fed to the feed intake of the corresponding ascites group.

c No supplemental choline.

d Administration to achieve 175% of the NRC requirement (1977) of choline (based ration provides 108%)

App. 2a. Major and interactive effect produced by dietary or intraperitoneal (IP) administration of choline on the feed conversion ratio of chicks with ateliosis.

Day	Feed Conversion ratio <sup>a</sup>						ANOVA ( $F_{(r_e > r_f)^b$ )	
	Effect of feeding			Effect of administration				
	Ateliosis <sup>c</sup>	Restricted Ad libitum	IP	Ateliosis	IP	Dietary		
2	-A	-A	-A	-A	-A	-A	0.20(0.82)	
3	-A	-A	-A	-A	-A	-A	0.69(0.51)	
4	-B	-A	-A	-A	-A	-A	2.21(0.68)	
5	-B	-A	-A	-A	-A	-A	1.08(0.38)	
6	-B	-A	-A	-A	-A	-A	1.70(0.20)	
7	-B	-A	-A	-A	-A	-A	1.08(0.35)	
8	-B	-A	-A	-A	-A	-A	0.50(0.61)	
9	-B	-A	-A	-A	-A	-A	1.25(0.31)	
10	-B	-A	-A	-A	-A	-A	0.05(0.91)	
11	-B	-A	-A	-A	-A	-A	0.12(0.89)	
12	-B	-A	-A	-A	-A	-A	1.84(0.19)	
13	-B	-A	-A	-A	-A	-A	1.01(0.42)	
14	-B	-A	-A	-A	-A	-A	2.12(0.14)	
15	-B	-A	-A	-A	-A	-A	0.91(0.43)	
16	-B	-A	-A	-A	-A	-A	0.88(0.47)	
17	-B	-A	-A	-A	-A	-A	2.57(0.65)	
18	-B	-A	-A	-A	-A	-A	1.58(0.34)	
19	-B	-A	-A	-A	-A	-A	1.40(0.29)	
20	-B	-A	-A	-A	-A	-A	4.85(0.18)	
21	-B	-A	-A	-A	-A	-A	1.56(0.70)	
22	-B	-A	-A	-A	-A	-A	0.66(0.52)	
23	-B	-A	-A	-A	-A	-A	2.70(0.04)	
24	-B	-A	-A	-A	-A	-A	1.35(0.28)	
25	-B	-A	-A	-A	-A	-A	3.41(0.08)	
26	-B	-A	-A	-A	-A	-A	0.79(0.54)	
27	-B	-A	-A	-A	-A	-A	0.18(0.92)	
28	-B	-A	-A	-A	-A	-A	4.81(0.16)	
29	-B	-A	-A	-A	-A	-A	4.33(0.02)	
30	-B	-A	-A	-A	-A	-A	7.14(0.04)	
31	-B	-A	-A	-A	-A	-A	1.28(0.30)	
32	-B	-A	-A	-A	-A	-A	3.32(0.04)	
33	-B	-A	-A	-A	-A	-A	2.40(0.11)	
34	-B	-A	-A	-A	-A	-A	1.88(0.14)	
35	-B	-A	-A	-A	-A	-A	0.84(0.51)	
36	-B	-A	-A	-A	-A	-A	2.91(0.02)	
37	-B	-A	-A	-A	-A	-A	0.71(0.52)	
38	-B	-A	-A	-A	-A	-A	1.34(0.28)	
39	-B	-A	-A	-A	-A	-A	0.10(0.90)	
40	-B	-A	-A	-A	-A	-A	1.63(0.21)	
41	-B	-A	-A	-A	-A	-A	1.34(0.28)	
42	-B	-A	-A	-A	-A	-A	0.10(0.90)	
43	-B	-A	-A	-A	-A	-A	1.08(0.35)	
44	-B	-A	-A	-A	-A	-A	1.08(0.35)	
45	-B	-A	-A	-A	-A	-A	1.39(0.23)	
46	-B	-A	-A	-A	-A	-A	0.50(0.74)	
47	-B	-A	-A	-A	-A	-A	2.77(0.01)	
48	-B	-A	-A	-A	-A	-A	0.60(0.53)	
49	-B	-A	-A	-A	-A	-A	0.60(0.67)	
50	-B	-A	-A	-A	-A	-A	1.47(0.25)	
51	-B	-A	-A	-A	-A	-A	0.85(0.63)	
52	-B	-A	-A	-A	-A	-A	1.77(0.17)	
53	-B	-A	-A	-A	-A	-A	0.58(0.57)	
54	-B	-A	-A	-A	-A	-A	2.23(0.04)	
55	-B	-A	-A	-A	-A	-A	0.22(0.53)	
56	-B	-A	-A	-A	-A	-A	3.64(0.04)	
57	-B	-A	-A	-A	-A	-A	1.23(0.32)	

None of feed conversion ratios (feed intake/weight gain) within a major effect followed by the same letter are not significantly different at  $P \leq 0.05$ .

Ateliosis at day 2.5% per g of diet.

Pair-wise to the isolate of the corresponding ateliosis group.

No supplemental choline.

Administration to achieve 175% of the NRC requirement (1977) of choline (basal ration provided 106%).

Feed = major effect of feeding. Feed x Admin = test for interaction.

App. 4. Effect of graded levels of dietary folate acid on the weight gain of chicks with atlantosis.

Day	Weight Gain (g/chick) <sup>a</sup>			General		
	Atlanthin (2.5% of diet)	344c	644c	Restrictive Feeding	244	644
0	391±.68	391±.36	391±.76	380±.19A	380±.79A	380±.36A
1	493±.28	493±.08	493±.24	473±.48A	473±.68A	481±.56A
2	575±.74	575±.26	575±.38	575±.16A	575±.16A	562±.94A
3	685±.24	715±.24	685±.04	664±.44A	693±.86A	682±.54A
4	795±.48	765±.08	772±.64A	765±.08A	812±.36A	812±.46A
5	935±.44A	885±.74	893±.54A	945±.67A	945±.50A	924±.52A
6	1047±.30A	1015±.78	1012±.48B	1084±.36AB	1115±.26AB	1092±.04AB
7	1197±.16A	1192±.86C	1202±.26C	1256±.04A	1252±.26C	1256±.78AB
8	1424±.04A	1364±.18C	1323±.60	1484±.94A	1475±.34	1462±.84A
9	1550±.46C	1527±.08C	1495±.46C	1825±.68B	1622±.24B	1605±.60A
10	1714±.30C	1679±.46C	1622±.60	1802±.59ABC	1829±.48BC	1825±.48AB
11	1915±.26C	1895±.46C	1784±.50	2082±.72AB	2119±.38B	1991±.24A
12	2024±.26D	2068±.26C	1928±.20	2315±.08AB	2395±.56A	2092±.04AB
13	2175±.29	2215±.10	2032±.68	2469±.18BC	2541±.24ABC	2447±.78
14	2325±.26E	2380±.39	2165±.52	2682±.05C	2715±.36C	2664±.1A
15	2509±.2C	2324±.70	2315±.45C	3032±.73	3325±.78	3197±.38
16	2535±.26	2815±.20	2515±.46C	3305±.38C	3435±.98C	3425±.4AC
17	2684±.72C	3064±.4C	2715±.70	3625±.98	3615±.58	3625±.48B
18	3045±.70E	3255±.60	2885±.27	3825±.70CD	3775±.21C	4005±.25A
19	3264±.28E	3154±.70	3115±.95	4165±.68C	4045±.28C	4335±.28A
20	341±.67	3735±.94	3825±.57	4325±.30CD	4175±.18B	4175±.21A
21	365±.78F	395±.26	3558±.69	4475±.61C	4325±.9D	4495±.25B
22	391±.68	420±.22F	3744±.56C	4675±.62	4525±.18	5615±.26A
23	405±.56G	457±.22F	415±.20	503±.40K	4995±.78	5385±.25D
24	435±.48F	474±.21E	412±.27	517±.90	564±.65C	583±.15C

<sup>a</sup> Values (means ± standard deviations) for weight gain followed by the same letter are not significantly different at  $P \leq 0.05$ .

Pair fed to the feed intake of the corresponding atlantia group.

% Percentage of the NRC dietary requirement (4977) of folic acid (basal ration provides 2444).

App. 4a. Major and interactive effects produced by graded levels of dietary folic acid on the weight gain of chicks with effluviation.

Day	Effect of feeding ATTENUS® Restricted Ad libitum	Effect of administration <sup>a</sup>			ANIMAL (Litter #)			Feed x Admin <sup>b</sup>	
		244		344	644		Admin		
		Feed	Admin	Feed	Admin	Feed	Admin		
0	-	-A	-A	-A	-A	-A	-A	0.310(0.59)	
1	-	-A	-A	-A	-A	-A	-A	0.310(0.53)	
2	-	-A	-A	-A	-A	-A	-A	1.200(0.32)	
3	-	-A	-A	-A	-A	-A	-A	0.460(0.64)	
4	-	-A	-A	-A	-A	-A	-A	0.740(0.57)	
5	-	-A	-A	-A	-A	-A	-A	0.640(0.54)	
6	-	-A	-A	-A	-A	-A	-A	0.470(0.75)	
7	-	-A	-A	-A	-A	-A	-A	0.030(0.39)	
8	-	-A	-A	-A	-A	-A	-A	0.010(0.39)	
9	-	-A	-A	-A	-A	-A	-A	0.010(0.39)	
10	-	-A	-A	-A	-A	-A	-A	0.510(0.53)	
11	1858	1036	1084	1228	1238	1258	1284	1.780(0.19)	
12	2128	1284	12648	1284	1238	1258	1284	7.070(0.03)	
13	2128	2514	2514	2514	2514	2514	2514	4.250(0.02)	
14	2128	1437	1434	1434	1434	1434	1434	5.480(0.014)	
15	2128	1634	1634	1634	1634	1634	1634	9.450(0.008)	
16	2688	2828	2828	2828	2828	2828	2828	0.500(0.61)	
17	2876	2128	2084	2084	2084	2084	2084	12.610(0.001)	
18	3046	2324	2324	2324	2324	2324	2324	28.910(0.001)	
19	3106	2514	2514	2514	2514	2514	2514	44.810(0.001)	
20	3196	2688	2884	2884	2884	2884	2884	55.4(0.001)	
21	3196	3046	3196	3196	3196	3196	3196	0.220(0.81)	
22	3246	3328	3484	3484	3484	3484	3484	94.760(0.001)	
23	3246	3238	3238	3238	3238	3238	3238	1.640(0.33)	
24	3246	2818	2884	2884	2884	2884	2884	92.000(0.001)	
25	3246	3484	3484	3484	3484	3484	3484	0.640(0.34)	
26	3446	3484	3636	3636	3636	3636	3636	6.540(0.004)	
27	3596	4184	4184	4184	4184	4184	4184	80.310(0.001)	
28	4416	5488	5234	5234	5234	5234	5234	1.08(0.25)	
29	4416	5128	6164	4116	4116	4116	4116	97.310(0.001)	
30	4416	5488	5234	5234	5234	5234	5234	5.310(0.001)	
31	4416	5488	5234	5234	5234	5234	5234	1.24(0.30)	
32	4416	5488	5234	5234	5234	5234	5234	2.67(0.048)	
33	4416	5128	6164	4116	4116	4116	4116	7.851(0.001)	
34	4416	5488	5234	5234	5234	5234	5234	4.931(0.013)	
35	4416	5488	5234	5234	5234	5234	5234	14.3(0.001)	
36	4416	5488	5234	5234	5234	5234	5234	12.44(0.001)	
37	4416	5488	5234	5234	5234	5234	5234	6.591(0.001)	
38	4416	5488	5234	5234	5234	5234	5234	5.54(0.001)	
39	4416	5488	5234	5234	5234	5234	5234	19.44(0.001)	
40	4416	5488	5234	5234	5234	5234	5234	5.961(0.001)	
41	4416	5488	5234	5234	5234	5234	5234	1.04(0.27)	
42	4416	5488	5234	5234	5234	5234	5234	9.961(0.001)	
43	4416	5488	5234	5234	5234	5234	5234	7.26(0.004)	

<sup>a</sup>Reason for the gain weights within a major effect followed by the same letter are not significantly different at P<0.05.

Dietary effluviation = 2.5% per & diet.

Opalinized to the intake of the corresponding (1977) of folic acid (basal ration provided 2412).

Opalinized to the NRC dietary requirement (1977) of folic acid (basal ration provided 2412).

Effluviation = major effect of administration. Feed = major effect of feeding. Feed x Admin = test for interaction.

App. 5. Effect of graded levels of dietary folic acid on the feed intake of chicks with adrenostenosis.

Day	Feed intake (g chick <sup>-1</sup> ) <sup>a</sup>		Feed intake (g chick <sup>-1</sup> ) <sup>b</sup>		Control	
	Alloxazine (2.5 µg of diet)	SAc	Restricted feeding	SAc	Control	SAc
1	1000.67A	9.049±0.68A	9.050±0.58A	8.150±0.59C	7.540±0.72C	8.051±0.28C
2	2149.70B	20.23±0.96C	21.51±2.08	17.51±0.70	18.51±1.20	16.61±1.62D
3	2121.14A	22.52±0.85C	22.52±1.28A	20.51±1.70	20.21±1.20	20.51±1.70
4	5.52±3.68A	5.25±0.86C	5.75±1.18AB	4.65±1.18	4.92±0.26B	4.15±0.68C
5	7929.08E	72.19±2.06C	82.59±1.58	64.52±1.20	69.52±1.20	70.54±1.48C
6	10299.09ECD	91.59±1.02C	102.60±1.20C	71.59±1.10	99.51±1.02D	97.54±1.48C
7	12121.28C	118.69±1.52C	141.51±1.88A	119.58±1.80C	98.21±1.10B	100.94±1.02CD
8	14665.02C	169.84±1.00C	154.50±1.44A	144.51±1.20	135.51±1.30C	130.51±1.20C
9	2084.21ECD	18.21±1.08C	20.02±0.84C	20.02±0.80C	17.75±1.14	21.51±1.08C
10	2812.28ECD	22.02±1.18	28.95±3.04C	24.62±2.02D	23.16±2.00	28.45±2.08B
11	2884.28ECD	25.71±1.30	31.84±0.68A	27.51±1.20	30.52±2.00C	24.61±1.20
12	3212.23ECD	27.96±1.40	32.24±2.44A	32.25±2.00D	32.25±2.00	33.44±2.00A
13	3825.44ECD	37.95±2.00D	42.75±2.28AB	37.25±1.80C	39.25±2.00D	39.51±2.00A
14	6215.54ECD	30.45±2.35C	44.35±2.44A	42.15±2.30C	44.55±2.00D	45.05±2.20A
15	4445.57ECD	43.25±2.46C	47.15±2.48C	43.95±2.70C	47.75±2.58AB	48.45±2.20A
16	5932.59ECD	47.45±2.18C	53.55±2.74AB	51.55±2.30C	49.55±2.58AB	50.25±2.34
17	5605.60ECD	51.15±2.78	61.15±2.78	55.55±3.58C	50.95±4.16C	54.45±3.06C
18	5915.60ECD	54.65±2.80C	63.05±3.58A	59.05±3.40C	54.55±4.26C	59.05±3.40C
19	6315.65ECD	58.85±2.80C	69.35±2.80C	63.55±2.80C	68.55±4.20	72.05±3.58
20	6815.65ECD	63.65±2.80	75.55±2.80C	61.55±1.00	67.45±2.00	71.55±2.80C
21	7115.65ECD	67.45±2.80	78.45±2.80C	72.55±3.40C	67.55±2.80	70.55±3.58C
22	7705.67ECD	72.25±2.80	89.55±2.76	78.55±2.00C	77.55±2.40	89.55±2.90C
23	8305.70ECD	77.15±2.80	87.65±2.76C	81.55±4.60C	76.95±2.30	86.55±4.06C
24	8815.75ECD	83.45±2.10	92.75±2.76C	84.55±7.00D	83.55±2.80	109.55±4.48B

<sup>a</sup> Values (mean ± standard deviation) for feed intake followed by the same letter are not significantly different at  $P \leq 0.05$ .

<sup>b</sup> Pair fed to the feed intake of the corresponding adrenostenosis group.

c Percentage of the NRC dietary requirement (13.77) of folic acid (folic acid provides 24.83%).

Table 5a. Major and interactive effects produced by graded levels of dietary folate and on the feed intake of chicks with *afattoinosis*.

Day	Feed intake (g/day)*		Effects of administration			ADULT HEN ( $P > 0.1$ ) <sup>b</sup>	
	Effect of Fattening / Aftatinosis <sup>c</sup>	Restricted Ad libitum	244	344	644	Feed	Feed x Admin
1	9.08A	7.88B	9.61A	-	-	14.8(0.0001), 1.44(0.26)	1.37(0.27)
2	20.56A	17.68	21.1A	-	-	20.76(0.0001), 0.52(0.49)	0.87(0.48)
3	36.5A	31.3B	37.5A	-	-	26.5(0.0001), 0.47(0.53)	1.38(0.27)
4	53.6A	47.5B	58.1A	-	-	18.7(0.0001), 0.62(0.55)	2.09(0.11)
5	77.1B	69.6C	84.8A	-	-	12.7(0.0001), 0.74(0.49)	2.98(0.037)
6	100B	92.6B	110C	-	-	9.3(0.0003), 1.62(0.32)	5.05(0.33)
7	130B	122B	143A	127B	129B	141A	6.55(0.0013), 4.25(0.023)
8	167B	156B	167A	161B	161B	180A	6.02(0.0018), 5.55(0.0017)
9	203B	197B	221A	202B	198B	222A	6.10(0.0017), 5.25(0.0012)
10	245B	236B	270A	241B	240B	271A	6.25(0.0015), 5.45(0.0014)
11	288B	260B	316A	260B	282B	321A	6.35(0.0013), 5.54(0.0013)
12	333B	326B	353B	327B	376A	376A	6.15(0.0003), 5.81(0.0003)
13	387B	376B	423B	373B	375B	434A	5.06(0.014), 5.18(0.009)
14	432B	427B	450A	437B	428B	494A	6.99(0.0046), 8.45(0.0014)
15	488B	478B	550A	481B	481B	549A	7.46(0.0016), 7.27(0.0020)
16	533B	518B	608A	536B	528B	591A	10.56(0.0004), 9.77(0.0012)
17	561B	554B	662A	511B	572B	653A	13.51(0.0001), 5.81(0.0001)
18	598B	581B	715A	614B	615B	679A	21.76(0.0001), 5.81(0.0079)
19	640B	633B	782A	633B	663B	730A	27.31(0.0001), 5.48(0.0077)
20	681B	676B	850A	716B	769B	873A	40.11(0.0001), 6.55(0.0043)
21	722B	723B	920A	770B	763B	872A	50.31(0.0001), 6.67(0.0044)
22	773B	771B	932A	826B	819B	892A	60.51(0.0001), 6.11(0.0053)
23	822B	819B	1073A	835B	874B	937A	74.7(0.0001), 7.14(0.0032)
24	861B	872B	1157A	954B	946B	1012A	42.31(0.0001), 4.47(0.012)

\*Mean for feed intake within a row effect followed by the same letter are not significantly different at P<0.05.

<sup>a</sup>Dietary aftatin = 215μg per g of diet.

<sup>b</sup>Refers to the intake of the corresponding afattoin group.

<sup>c</sup>Percentage of the inc. dietary requirement (1972) of folic acid (basal ration provided 244t).

Admin = major effect of administration. Feed = major effect of feeding. Feed x Admin = test for interaction.

App. 6. Effect of graded levels of dietary folic acid on the feed conversion of weanling rats.

Dose	Feed conversion ratio <sup>a</sup>		Control	
	Unrestricted feeding	Restricted feeding		
24C	24A	6A	6A	
1	1.0109±.17A	0.9900±.020A	0.9800±.020A	0.9310±.07A
2	1.1650±.06A	1.1150±.07A	1.0500±.06A	1.0450±.06A
3	1.2650±.02A	1.1650±.03A	1.1250±.02A	1.0450±.00A
4	1.4150±.12A	1.3250±.13A	1.4650±.08A	1.3250±.11A
5	1.4550±.14A	1.3250±.13A	1.4650±.08A	1.3250±.13A
6	1.5850±.17A	1.4250±.02AC	1.6550±.09A	1.3250±.18
7	1.6250±.18C	1.4350±.02AC	1.6950±.21A	1.2850±.16D
8	1.6350±.18C	1.4350±.02AC	1.7450±.21A	1.3050±.02D
9	1.7250±.20A	1.5150±.04C	1.8150±.21A	1.3650±.12D
10	1.8850±.23A	1.5750±.07AC	2.0250±.20A	1.4850±.19C
11	1.9850±.23DC	1.6950±.07C	2.2150±.22A	1.6150±.17DC
12	2.0450±.21DC	1.7350±.07DC	2.4250±.28A	1.6350±.07DF
13	2.1450±.21A	1.8450±.10DC	2.5850±.30A	1.7650±.17DC
14	2.2350±.20DC	1.9250±.14DC	2.7150±.33A	1.8450±.19DC
15	2.3750±.21A	1.9750±.21DC	2.7750±.35A	1.8250±.16DF
16	2.3750±.21A	1.9550±.24DC	2.7050±.32A	1.7950±.03DF
17	2.3750±.24A	1.9250±.23DC	2.6250±.31A	1.7150±.12DF
18	2.2340±.23A	1.8150±.23DC	2.4650±.27A	1.7150±.09DF
19	2.2950±.23A	1.8450±.21B	2.3450±.34A	1.7150±.09DF
20	2.2350±.23A	1.8150±.17DC	2.3250±.28A	1.7050±.07DF
21	2.1950±.20A	1.8150±.17DC	2.4850±.27A	1.7150±.09DF
22	2.1950±.20A	1.8150±.17DC	2.4150±.24A	1.7150±.09DF
23	2.2750±.23A	1.8150±.15DC	2.3750±.24A	1.7450±.08E
24	2.2750±.23A	1.8150±.16DC	2.4850±.27A	1.6450±.15D

<sup>a</sup> Values (means ± standard deviation) for feed conversion (feed intake/weight gain) followed by the same letter are not significantly different at P < 0.05.

b Ratio fed to the feed intake of the corresponding astatin group.

c Percentage of the PRC dietary requirement (1977) of folic acid (based ration provides 2442).

App. 6a. Major and interactive effects produced by graded levels of dietary folic acid on the feed conversion of chicks with aflatoxinosis.

Day	Feed conversion ratio*		Effects of administration		ANOVA (F <sub>1,16</sub> > F <sub>1,15</sub> )	
	244 344		644		Feed	
	Aflatoxin/Restricted Ad libitum		Feed A Admin		Feed A Admin	
1	0.928	0.818	0.918	0.74	0.74	3.16(0.05)
2	1.111	0.898	1.098	1.4	1.4	2.12(0.14)
3	1.190	1.028	1.298	1.4	1.4	0.86(0.43)
4	1.414	1.118	1.538	1.4	1.4	0.91(0.91)
5	1.516	1.238	1.538	1.4	1.4	2.25(0.0034)
6	1.578	1.258	1.578	1.4	1.4	0.31(0.73)
7	1.558	1.258	1.618	1.4	1.4	0.97(0.39)
8	1.478	1.428	1.708	1.50	1.568	1.726
9	1.798	1.578	1.798	1.688	1.658	1.826
10	1.908	1.638	1.888	1.738	1.758	1.926
11	1.598	1.618	1.668	1.718	1.728	1.908
12	2.068	1.658	1.898	1.778	1.778	2.068
13	2.198	1.768	1.938	1.878	1.858	2.188
14	2.398	1.858	2.008	1.938	1.928	2.288
15	2.348	1.798	1.978	1.978	1.948	2.284
16	2.348	1.758	1.888	1.938	1.938	2.204
17	2.278	1.726	1.918	1.868	1.868	2.158
18	2.258	1.706	1.928	1.868	1.868	2.118
19	2.218	1.686	1.898	1.838	1.858	2.088
20	2.238	1.676	1.908	1.878	1.898	2.084
21	2.198	1.726	1.918	2.118	1.776	1.908
22	2.188	1.708	1.908	1.898	1.948	2.078
23	2.168	1.736	1.898	1.898	1.968	2.038
24	2.218	1.746	1.988	1.938	1.898	2.118

Means of feed conversion ration (feed intake/weight gain) within a major effect followed by the same letter are not significantly different at P<0.05.

Supplementary aflatoxin = 2.0 mg per kg of diet.

o - related to the intake of the corresponding aflatoxin group.

Difference of the NRC dietary requirement (1972) of folic acid (base) ration provided 2.04%.

Admin = major effect of administration. Feed = Major effect of feeding. (Feed x Admin = test for interaction).

App. 7. Effect of graded levels of dietary threonine on the weight gain of chicks with afflatoxins.

Day	Afflatoxin (G.5. U/gm of diet)		Weight Gain (g/Chick)*		Control			
	0	120C	135C	170C		120	135	170
0	260.96A	260.60AB	291.6A	341.6B	362.8AB	252.23B	252.23B	262.4AB
3	3461.5E	5521.7D	6251.7B	4021.4BCD	6725.5A	6051.18e-D	6325.29e-C	5723.4
6	9126.5E	9216.5E	10151.2A	9941.7AB	10725.5A	10425.5AB	10425.5AB	10251.7AB
9	14023.8H	14241.2H	15648.5AB	14549.4AB	13525.5AB	13525.5AB	13525.5AB	13462.5A
12	19392.5J	19641.2J	21512.2CD	19321.20	21029.1CD	21725.1CD	22292.5CA	24411.6AB
15	21526.3P	22021.7AP	21512.5ABP	18481.1AF	21029.4CD	21525.4CD	21029.4CD	21122.5AB
18	31151.2D	31032.7D	31142.9	31615.1D	34615.1C9	33151.2D	36792.9C	36151.1B
22	44392.2D	43024.0D	45123.2CD	44644.6D	49721.5CD	49351.6BC	53952.1B	53251.2B
23	48452.2F	46521.0F	50051.2AF	50252.3AF	53725.4D	53725.4D	61151.47A	60192.1CB
24	50452.4F	48651.1F	51812.8AF	52251.2AF	54721.7BC	57251.5CD	62052.9CA	62851.2AB

\* Values (mean  $\pm$  standard deviation) for weight gain followed by the same letter are not significantly different at  $P \leq 0.05$ .

b/c = ratio of the feed intake of the corresponding afflatoxin group.

c Percentage of the NRC dietary requirement (1977) of threonine (basal ration provides 12.8%).

App. 7a. Major and interactive effects produced by graded levels of dietary threonine on the weight gain of chicks with Aflatoxinosis.

Day	Effect of Aflatoxin Aflatoxin <sup>a</sup> Restricted Ad libitum	Weight gain (g/chick) <sup>b</sup>			ANOVAs (F <sub>r</sub> > F <sub>t</sub> ) <sup>c</sup>		
		Effect of administration <sup>d</sup>			Feed x Admin		
		12B	1355	179	Feed	Admin	Feed x Admin
0		26.7A	24.9B	25.4AB	24.9B	25.4AB	26.8A
3		57.1A	62.2A	59.8AB	57.1A	62.2A	59.8AB
6		96.5A	104A	102A	96.5A	102AB	103A
9		147B	153AB	162A	147B	153AB	161A
12		201B	208B	243A	205B	217B	229A
15		232B	270B	331A	248B	287AB	299A
18		323B	331B	402A	335B	347B	373A
22		440C	471B	569A	473B	478AB	552A
23		483C	532B	639A	532A	535B	586A
24		505C	549B	659A	515A	554B	607A
					49.71(0.001)	49.71(0.001)	7.78(0.002)

<sup>a</sup>Mean for the gain in weights within a major effect followed by the same letter are not significantly different at P<0.05.

<sup>b</sup>Dietary aflatoxin = 2.5% per g of diet.

<sup>c</sup>Attributed to the intake of the corresponding aflatoxin group.

<sup>d</sup>Percentage of the NRC dietary requirement (1977) of the threonine (basal ration provided 12%).

<sup>e</sup>Admin = major effect of administration. Feed = major effect of feeding. Feed x Admin = test for interaction.

App. 8. Effect of graded levels of dietary threonine on the feed intake of chicks with aflatoxins.

Day	Feed intake (g/chick) <sup>a</sup>			Control
	Aflatoxin (1.5 mg/kg diet)	1.0% restricted feeding <sup>b</sup>	1.0% unrestricted feeding <sup>b</sup>	
1	128 <sup>c</sup>	128 <sup>c</sup>	128 <sup>c</sup>	128
2	154 <sup>c</sup> , 1CD	140 <sup>c</sup> , 1CD	163 <sup>c</sup> , 1CD	155 <sup>c</sup>
3	255 <sup>c</sup> , 1CD	255 <sup>c</sup> , 1CD	262 <sup>c</sup> , 1.0A	155 <sup>c</sup>
4	295 <sup>c</sup> , 1CD	389 <sup>c</sup> , 1.0D	424 <sup>c</sup> , 1.0C	179
5	544 <sup>c</sup> , 1CD	535 <sup>c</sup> , 1.0E	672 <sup>c</sup> , 1.0B	128
6	744 <sup>c</sup> , 1CD	734 <sup>c</sup> , 1.0C	675 <sup>c</sup> , 1.0A	128
7	964 <sup>c</sup> , 1CD	964 <sup>c</sup> , 1.0C	1021 <sup>c</sup> , 1.0C	135 <sup>c</sup>
8	1205 <sup>c</sup> , 1CD	1169 <sup>c</sup> , 1.0B	1249 <sup>c</sup> , 1.0A	132 <sup>c</sup> , 2.0A
9	1485 <sup>c</sup> , 1.0B	1472 <sup>c</sup> , 1.0B	1482 <sup>c</sup> , 1.0B	134 <sup>c</sup> , 1.0A
10	1725 <sup>c</sup> , 1CD	1725 <sup>c</sup> , 1CD	1725 <sup>c</sup> , 1CD	1725 <sup>c</sup> , 1CD
11	2001 <sup>c</sup> , 1CD	2001 <sup>c</sup> , 1CD	2001 <sup>c</sup> , 1CD	2012 <sup>c</sup> , 1CD
12	2352 <sup>c</sup> , 1CD	2431 <sup>c</sup> , 1CD	2671 <sup>c</sup> , 1.0D	2389 <sup>c</sup> , 1CD
13	2662 <sup>c</sup> , 1CD	2562 <sup>c</sup> , 1CD	2632 <sup>c</sup> , 1CD	2667 <sup>c</sup> , 1CD
14	3152 <sup>c</sup> , 1CD	2984 <sup>c</sup> , 1CD	3032 <sup>c</sup> , 1CD	3120 <sup>c</sup> , 1CD
15	3525 <sup>c</sup> , 1CD	3372 <sup>c</sup> , 1CD	3415 <sup>c</sup> , 1CD	3524 <sup>c</sup> , 1CD
16	4014 <sup>c</sup> , 1CD	4232 <sup>c</sup> , 1CD	4327 <sup>c</sup> , 1CD	4094 <sup>c</sup> , 1CD
17	4395 <sup>c</sup> , 1C	4343 <sup>c</sup> , 1C	4317 <sup>c</sup> , 1C	4285 <sup>c</sup> , 1CD
18	4902 <sup>c</sup> , 1C	4142 <sup>c</sup> , 1C	5144 <sup>c</sup> , 1C	4385 <sup>c</sup> , 1CD
19	5264 <sup>c</sup> , 1C	5272 <sup>c</sup> , 1C	5665 <sup>c</sup> , 1C	5095 <sup>c</sup> , 1CD
20	6032 <sup>c</sup> , 1C	5853 <sup>c</sup> , 1C	6185 <sup>c</sup> , 1CD	5847 <sup>c</sup> , 1CD
21	6652 <sup>c</sup> , 1B	6454 <sup>c</sup> , 1B	6745 <sup>c</sup> , 1B	6234 <sup>c</sup> , 1B
22	7365 <sup>c</sup> , 1A	7194 <sup>c</sup> , 1A	7375 <sup>c</sup> , 1A	6829 <sup>c</sup> , 1A
23	7920 <sup>c</sup> , 1B	7757 <sup>c</sup> , 1B	7932 <sup>c</sup> , 1B	7612 <sup>c</sup> , 1B
24	8445 <sup>c</sup> , 1B	8324 <sup>c</sup> , 1B	8632 <sup>c</sup> , 1B	8059 <sup>c</sup> , 1B

<sup>a</sup> Values (mean  $\pm$  standard deviations) for feed intake followed by the same letter are not significantly different at  $P \leq 0.05$ .

<sup>b</sup> Feed fed to the feed intake of the corresponding aflatoxin group.

<sup>c</sup> Percentage of the NRC dietary requirement (1977) of threonine (basal ration provides 128%).

App. 8a. Major and interactive effects produced by graded levels of dietary threonine on the feed intake of chicks with *afflatenosis*.

Day	Effect of feeding Afflatenosis Restricted Ad Libitum	Feed Intake (g/day)		Effect of administration		SOMA (FF(Fe > F)) <sup>a</sup>		
		125 235		179		Feed		
		Feed	Admin	Feed	Admin	Feed	Admin	
1	5.648	7.728	-A	-A	-A	12.910 (0.001)	0.020 (0.51)	
2	15.038	17.94	-A	-A	-A	10.310 (0.005)	0.740 (0.48)	
3	25.50	30.04	-B	-A	-A	19.010 (0.001)	1.510 (0.23)	
4	40.00	46.04	-A	-A	-A	19.410 (0.63)	2.310 (0.039)	
5	56.30	67.84	-A	-A	-A	13.310 (0.001)	0.130 (0.48)	
6	76.80	84.16	B2-A	-A	-A	21.210 (0.001)	0.210 (0.56)	
7	99.60	103.8	102A	102A	102A	5.210 (0.912)	2.410 (0.074)	
8.	125B	128B	132A	125B	125AB	4.260 (0.025)	2.780 (0.080)	
9.	154B	153B	167A	153B	154B	2.6710 (0.689)	3.1810 (0.598)	
10	181B	183B	201A	181B	188AB	6.210 (0.008)	3.8210 (0.034)	
11	211B	210B	212A	210B	210A	7.4610 (0.008)	3.9110 (0.322)	
12	248B	241B	281A	210B	210A	9.3110 (0.007)	3.4410 (0.040)	
13	279B	268B	275A	278B	278A	12.3510 (0.001)	3.5110 (0.644)	
14	317B	308B	381A	-A	-A	14.410 (0.001)	0.3010 (0.12)	
15	358B	346B	432A	-A	-A	18.3510 (0.001)	1.9210 (0.16)	
16	408B	393B	491A	-A	-A	20.510 (0.001)	1.6810 (0.20)	
17	-	445B	438B	539A	-A	-A	24.210 (0.001)	1.9510 (0.16)
18	-	493B	473B	592A	-A	-A	18.110 (0.001)	0.2810 (0.89)
19	-	545B	523B	660A	-A	-A	20.210 (0.001)	1.8910 (0.17)
20	-	602B	575B	730A	-A	-A	20.810 (0.001)	1.3350 (0.28)
21	-	662B	631B	690A	-A	-A	22.210 (0.001)	0.9910 (0.38)
22	-	731B	692B	888A	-A	-A	23.810 (0.001)	0.2710 (0.49)
23	-	787B	761B	932A	-A	-A	23.4410 (0.001)	0.6110 (0.55)
24	-	860B	815B	1039A	-A	-A	26.3110 (0.001)	0.3510 (0.84)

Reasons for feed intakes within a major effect followed by the same letter are not significantly different at P<0.05.

Isoprotein afflatenosis = 2.75% per g of diet.

Spliced to the intake of the corresponding afflatenosis group.

Difference of the NRC dietary requirement (1977) of threonine (base) - required 1.88%.

Feed = major effect of feeding.

Admin = major effect of administration.

App. 9. Effect of graded levels of dietary threonine on the feed conversion of chicks with allostrotinosis

Dose	Feed conversion ratio <sup>a</sup>					
	Allostrotin (2.5 g/lb of diet)	155	179	172	155	179
	Restricted feeding <sup>b</sup>					
3	1.4249-0.06A	1.350-0.15AB	1.1029-0.12CD	1.1729-0.11ACD	1.069-0.07D	1.1359-0.07CD
6	1.3449-0.08A	1.3029-0.13A	1.2469-0.09A	1.2259-0.07A	1.1869-0.01A	1.2929-0.25A
9	1.4249-0.09A	1.4149-0.23A	1.4029-0.08A	1.3529-0.08A	1.2669-0.05A	1.3029-0.08A
12	1.5049-0.16A	1.5329-0.12A	1.5169-0.11A	1.4649-0.08A	1.4369-0.08A	1.4259-0.15A
15	1.6849-0.21A	1.5929-0.22AB	1.7129-0.15A	1.5529-0.07AB	1.4419-0.03E	1.4749-0.08AB
18	1.7249-0.19AB	1.6569-0.17AB	1.7469-0.20AB	1.6329-0.08AB	1.5329-0.07AB	1.6569-0.14AB
22	1.8149-0.19A	1.8469-0.15A	1.7529-0.17AB	1.6749-0.17AB	1.5949-0.07BC	1.5249-0.07BC
23	1.7249-0.16AB	1.4529-0.14A	1.7129-0.12ABC	1.6129-0.10B-C	1.5329-0.07BC	1.4649-0.07E
24	1.8449-0.17AB	1.9049-0.14A	1.7929-0.13ABC	1.6529-0.07C-F	1.6169-0.06EF	1.7249-0.13B-E

<sup>a</sup> Values (mean  $\pm$  standard deviation) for weight gain followed by the same letter are not significantly different at  $P \leq 0.05$ .

<sup>b</sup> Pairs fed to the feed intake of the corresponding allostrotin group.

<sup>c</sup> Percentage of the NRC dietary requirement (1977) of threonine (basal ration provides 12.8%).

App. 9a. Major and interactive effects produced by graded levels of dietary threonine on the feed conversion of chicks with elastotensis.

Day		Feed conversion ratio*			ANOVA (F <sub>128</sub> , 155 > F <sub>1, D</sub> ) <sup>b</sup>	
		Effect of diet				
		Effect of elastotensis	Effect of administration <sup>c</sup>	Feed x Adnin		
3		1.129A	1.118B	-A	5.86(0.00813)	
6		1.126 <sup>d</sup>	-A	-A	1.05(0.326)	
9		1.126 <sup>d</sup>	-A	-A	2.53(0.412)	
12		1.121A	1.118AB	-A	0.34(0.791)	
15		1.124A	1.120B	-A	0.34(0.792)	
18		1.124A	1.120B	-A	0.67(0.432)	
22		1.127A	1.123B	-A	1.17(0.233)	
23		1.125A	1.120B	-A	0.43(0.661)	
26		1.125A	1.120B	-A	2.37(0.019)	

\*Means of feed conversion ratios (feed intake/weight gain) within a major effect followed by the same letter are not significantly different at P<0.05.  
†Protein elastotensis = 2.5% per g of diet.

<sup>a</sup>Refers to the feed intake of the corresponding elastotensis group.

<sup>b</sup>Dependent of the NRC dietary requirement (1972) of threonine (basal ration provided).

<sup>c</sup>Adnin = major effect of administration. Feed = major effect of feeding. Feed x Adnin = test for interaction.

App. 13. Effect of graded levels of dietary lysine on the weight gain of chicks with dl-alanotaurine.

Dose	Weight gain (g chick) <sup>a</sup>			Control
	Alanyltaurine (1.1% w/w of diet)	Metathiolated Feeding <sup>b</sup>	102	
0	37.469 ± .584	38.152 ± .654	37.750 ± .504	38.247 ± .526
3	51.462 ± .648	51.451 ± .524	52.023 ± .461	50.745 ± .548
6	74.442 ± .116	80.454 ± .486	81.154 ± .286	84.296 ± .648
9	113.2 ± .06	124.5 ± .28	124.5 ± .48	130.0 ± .048
12	159.2 ± .81	171.5 ± .20	172.9 ± .540	194.7 ± .348
15	203.9 ± .29	228.5 ± .150	227.0 ± .116	277.5 ± .048
18	252.1 ± .46	272.5 ± .092	277.4 ± .030	332.9 ± .649
22	301.4 ± .86	346.4 ± .246	388.5 ± .040	457.9 ± .356
24	367.2 ± .37	411.0 ± .258	402.0 ± .087	461.0 ± .356

<sup>a</sup> Values (means ± standard deviations) for weight gain followed by the same letter are not significantly different at P < 0.05.<sup>b</sup> pair fed to the feed intake of the corresponding alanyltaurine group.<sup>c</sup> Percentage of the NRC dietary requirement (1977) of lysine (base ration provides 102%).

App. 10a. Major and interactive effects produced by graded levels of dietary lysine on the weight gain of chicks with arlatoxins.

Day		Weight gain (g/chick) <sup>a</sup>		ANOVA ( $F_{1,12} > F_{1,1}$ ) <sup>b</sup>	
		Effect of feeding Arlatoxin <sup>c</sup> restricted Ad libitum		Effect of administration <sup>d</sup>	
		102	122	146	122
0		52.7AB	50.6B	53.1A	51.0B
3		76.7B	60.3AB	63.7A	78.4B
6		120B	129A	132A	128AB
9		167B	150A	155A	170B
12		219C	268B	280A	-A
15		237C	356B	354A	-A
18		308C	446B	532A	442B
21		393C	452B	559A	-A
24					-A

<sup>a</sup>Mean for the gain in weights within a major effect followed by the same letter are not significantly different at P<0.05.

<sup>b</sup>Dietary arlatoxin = 2.5% per g of diet.

<sup>c</sup>Pair-fed to the intake of the corresponding arlatoxin group.

<sup>d</sup>Percentage of the NRC dietary requirement (1977) of lysine (basal ration provided 102%).

<sup>e</sup>Admin = major effect of administration. Feed = major effect of feeding. Feed x Admin = test for interaction.

App. 11 Effect of graded levels of dietary lysine on the feed intake of chicks with effluviosis.

Day	Aflatoxin (1.5 mg/kg of diet)	Feed Intake (g/chick) <sup>a</sup>			General
		Restraint	Feeding	102	
1	102 <sup>c</sup>	132 <sup>c</sup>	146 <sup>c</sup>	102	146
2	10-24-7A 13-25-2B 20-25-3BC	7.930-7.728 6.156-6.138 7.943-7.827	11.125-11.126 11.125-11.126 7.562-7.468	13.125-13.126 13.125-13.126 7.562-7.468	8.943-8.846 8.569-8.569 8.472-8.474
3	33-124-3A 33-124-3A	23.934-27.802 23.934-27.802	23.125-24.467 23.125-24.467	23.125-24.467 23.125-24.467	18.029-20.818 17.725-19.840
4	53.009-0.048 41-459-3C	35.652-34.488 49-749-4C	35.652-34.488 49-749-4C	35.652-34.488 49-749-4C	29.422-4.488 35.725-35.725
5	73-249-4AB 82-021-3C	60-105-1CD 71-145-1C	60-105-1CD 71-145-1C	60-105-1CD 71-145-1C	44.85-51.740 82-115-1CA
6	91-253-3AB	71-145-1C	71-145-1C	71-145-1C	83-32.5-46.4 104.8-104.8
7	126-148-8A	109.5-17C	109.5-17C	109.5-17C	145.0-13A 112-12BC
8	138-151-6AB	144.2-18BCD	134.5-21C	128.9-20	130.6-18C 132.5-20C
9	200-219-8C	181.5-18BC	179.5-20C	160.9-22C	18.845-13C 18.669-13A
10	240-234-8	220-231-BC	210-149-3C	201-141-1C	218.6-15BC 216.1-16B
11	282-232-2BC	251-231-BC	241-159-3C	240-151-1C	216.6-15BC 212.0-17B
12	296-238-BC	287-231-BC	289-238-BC	273.2-11C	290-238-BC 300-230-1C
13	349-238-BC	328-231-1CD	333-231-1CD	309.5-42B	325-231-1CD 373-231-1CD
14	375-231-1C	364-231-1C	364-231-1C	351-14C	418.5-14B 418.5-14B
15	413-231-1C	403-232-1C	409-231-1C	391-14C	407.5-13C 407.5-13C
16	449-141-1CD	442-152-3CD	449-152-3D	448.1-1CD	450.5-13B 525.6-10B
17	487-246-CD	481-124-1B	485-146-1CD	472-152-3D	492-246-CD 492-246-CD
18	533-250-1B	523-250-1B	521-155-3B	532-151-1B	532-151-1B 648.5-12B
19	573-247-1B	568-252-1B	563-252-1B	561-155-1D	575.5-11D 71.26-14B
20	634-254-1B	620-252-1B	614-149-1D	621-151-1D	604-251-1B 784-151-1B
21	675-254-1B	623-253-1B	623-253-1B	623-253-1B	623-253-1B 784-253-1B
22	712-257-1B	722-257-1B	720-256-1B	700-151-1D	702-251-1D 702-251-1D
23	792-245-1B	783-244-1B	774-152-1B	754-151-1B	710-241-1B 705-151-1D
24	839-161-1D	830-252-1B	821-152-1B	817-149-1B	818-151-1D 1030-22B

<sup>a</sup> Values (mean + standard deviation) for feed intake followed by the same letter are not significantly different at P  $\leq 0.05$ .

b pair fed to the feed intake of the corresponding aflatoxin group.

c percentage of the NRC dietary requirement (1977) of lysine (base ration provides 10.2%).

App. IIa. Major and interactive effects produced by graded levels of dietary lysine on the feed intake of chicks with ascitesosis.

Day	Effect of Lysine Administration Additive <sup>a</sup>	Effect of administration Additive <sup>b</sup>		ANOVA ( $F(Fe > Fe)$ ) <sup>c</sup>	
		Feed intake (g/chick)		Feed	Admin
		102	122	146	146
1	-	9.02A	6.36B	6.07B	2.16(0.11)
2	13.95	12.95	16.6A	16.36	1.82(0.045)
3	26.68	24.08	20.1A	-A	5.42(0.0019)
4	42.28	38.98	50.1A	-A	7.18(0.0023)
5	61.18	58.08	69.5A	63.44B	8.50(0.0014)
6	83.88	77.38	90.9A	85.2A	5.50(0.0090)
7	111.28	101C	123A	114AB	1.05A
8	145.8	130C	158A	152A	10.5(0.0004)
9	187.8	163C	201A	185AB	1.75A
10	221.8	207C	248A	226AB	2.27A
11	256.8	246B	280A	-A	8.34(0.0015)
12	291.8	276B	320A	-A	4.42(0.0037)
13	334.8	311C	383A	357A	1.53A
14	-	368.8	355A	379AB	4.00A
15	409.8	405B	428A	420B	2.71A
16	443.8	446B	547A	475B	508A
17	-	483.8	486B	517B	552A
18	-	528.8	527B	660A	546AB
19	-	568.8	567B	723A	652A
20	-	619.8	609B	791A	670B
21	-	671.8	662B	859A	723B
22	-	783.8	769B	930A	781B
23	-	783.8	764B	1004A	844B
24	-	830.8	821B	1069A	946A

Means for feed intakes within a major effect followed by the same letter are not significantly different at  $P \leq 0.05$ .

<sup>a</sup>Protein ascites = 2.5% per g of diet.

<sup>b</sup>Related to the intake of the corresponding ascites group.

<sup>c</sup>Percentage of the FFC dietary requirement (1972) of lysine (total ration provided 10.2%).

<sup>d</sup>Abuse = major effect of administration; Feed = major effect of feeding; Feed  $\times$  Admin = test for interaction.

Effect of graded levels of dietary lysine on the feed conversion of chicks with aflatoxinosis.

Age days	Feed conversion (g chick) <sup>a</sup>		Control
	Aflatoxin (2.5 mg/kg of diet) <sup>b</sup>	Unrestricted feeding <sup>c</sup>	
3	102 <sup>e</sup>	122 <sup>c</sup>	146 <sup>c</sup>
6	2.43±0.26A	1.76±0.48ABC	1.39±0.06C
9	2.45±0.32A	2.10±0.17BC	1.64±0.26C
12	2.23±0.15A	2.16±0.16BC	1.69±0.20C
15	2.46±0.10A	2.17±0.16AB	2.12±0.27BCD
18	2.49±0.09A	2.32±0.13B	2.18±0.16AB
21	2.52±0.11A	2.21±0.11AB	1.82±0.23AB
24	2.55±0.11A	2.20±0.11AB	1.82±0.17B
			2.15±0.54ABC

<sup>a</sup> Values (means ± standard deviations) for feed conversion (feed intake/weight gain) followed by the same letter are not significantly different at P ≤ 0.05.<sup>b</sup> Pair fed to the feed intake of the corresponding aflatoxin group.<sup>c</sup> Percentage of the NRC dietary requirement (1977) of lysine (mean ration provided "025").

App. 12a. Major and interactive effects produced by graded levels of dietary lysine on the feed conversion of chicks with *allatoxins*.

Day	Feed conversion (g/chick) <sup>a</sup>														
	Effect of feeding				Effect of administration <sup>d</sup>				ANCOVA (Feed x F1) <sup>e</sup>						
	Allatoxin <sup>b</sup>		Restricted Ad <sup>c</sup>		102		122		146		Feed		Admin		Feed x Admin
Day	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
3	2.08A	1.84B	1.98AB	2.22A	1.80A	1.86A	-	-	-	0.46(0.64)	2.77(0.081)	2.69(0.033)	-	-	
6	2.12A	1.79B	2.15A	2.12A	1.91B	1.88B	-	-	-	3.11(G,0.61)	3.62(0.041)	3.27(0.0021)	-	-	
9	2.12A	1.79B	2.15A	2.07AB	1.97B	1.97B	-	-	-	16.81(0.0001)	2.91(0.072)	2.90(0.031)	-	-	
12	2.27A	1.82C	2.03B	-A	-A	-A	-	-	-	20.90(0.0001)	1.65(0.21)	4.81(0.0046)	-	-	
15	2.27A	1.86C	2.03B	-A	-A	-A	-	-	-	24.26(0.0001)	0.95(0.43)	3.41(0.018)	-	-	
18	2.21A	1.76C	2.03B	-A	-A	-A	-	-	-	24.70(0.0001)	-0.84(0.44)	4.78(0.0048)	-	-	
22	2.21A	1.86C	2.03B	2.13B	2.06B	2.01A	-	-	-	24.70(0.0001)	-2.55(0.097)	4.54(0.0032)	-	-	
24	2.25A	1.99E	2.03B	2.13B	2.06B	2.01A	-	-	-	25.30(0.0001)	2.98(0.069)	4.14(0.0098)	-	-	

Percentages of feed conversion (g/g) (gained/weight gain) within a pair effect followed by the same letter are not significantly different at P<0.05.

Dietary allatoxin = 2.5% per diet.

Restricted ad = the feed intake of chicks not receiving allatoxin group.

Opencage of the NRC dietary requirement (1977) of lysine (base provided 102).

Major = major effect of administration. Feed = major effect of feeding. Feed x Admin = test for interaction.

APPLIED POLYMER SYMPOSIA, VOL. 13, PAGES 179-192 (1970)

Year	Period	Estimated Exports (in \$ millions)												Estimated Imports (in \$ millions)												
		Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec	
2005	Jan	16.5	17.0	17.5	18.0	18.5	19.0	19.5	20.0	20.5	21.0	21.5	22.0	22.5	23.0	23.5	24.0	24.5	25.0	25.5	26.0	26.5	27.0	27.5	28.0	
2005	Feb	16.8	17.2	17.6	18.0	18.4	18.8	19.2	19.6	20.0	20.4	20.8	21.2	21.6	22.0	22.4	22.8	23.2	23.6	24.0	24.4	24.8	25.2	25.6	26.0	
2005	Mar	17.1	17.5	17.9	18.3	18.7	19.1	19.5	19.9	20.3	20.7	21.1	21.5	21.9	22.3	22.7	23.1	23.5	23.9	24.3	24.7	25.1	25.5	25.9	26.3	
2005	Apr	17.4	17.8	18.2	18.6	19.0	19.4	19.8	20.2	20.6	21.0	21.4	21.8	22.2	22.6	23.0	23.4	23.8	24.2	24.6	25.0	25.4	25.8	26.2	26.6	
2005	May	17.7	18.1	18.5	18.9	19.3	19.7	20.1	20.5	20.9	21.3	21.7	22.1	22.5	22.9	23.3	23.7	24.1	24.5	24.9	25.3	25.7	26.1	26.5	26.9	
2005	Jun	18.0	18.4	18.8	19.2	19.6	20.0	20.4	20.8	21.2	21.6	22.0	22.4	22.8	23.2	23.6	24.0	24.4	24.8	25.2	25.6	26.0	26.4	26.8	27.2	
2005	Jul	18.3	18.7	19.1	19.5	19.9	20.3	20.7	21.1	21.5	21.9	22.3	22.7	23.1	23.5	23.9	24.3	24.7	25.1	25.5	25.9	26.3	26.7	27.1	27.5	
2005	Aug	18.6	19.0	19.4	19.8	20.2	20.6	21.0	21.4	21.8	22.2	22.6	23.0	23.4	23.8	24.2	24.6	25.0	25.4	25.8	26.2	26.6	27.0	27.4	27.8	
2005	Sep	18.9	19.3	19.7	20.1	20.5	20.9	21.3	21.7	22.1	22.5	22.9	23.3	23.7	24.1	24.5	24.9	25.3	25.7	26.1	26.5	26.9	27.3	27.7	28.1	
2005	Oct	19.2	19.6	20.0	20.4	20.8	21.2	21.6	22.0	22.4	22.8	23.2	23.6	24.0	24.4	24.8	25.2	25.6	26.0	26.4	26.8	27.2	27.6	28.0	28.4	
2005	Nov	19.5	19.9	20.3	20.7	21.1	21.5	21.9	22.3	22.7	23.1	23.5	23.9	24.3	24.7	25.1	25.5	25.9	26.3	26.7	27.1	27.5	27.9	28.3	28.7	
2005	Dec	19.8	20.2	20.6	21.0	21.4	21.8	22.2	22.6	23.0	23.4	23.8	24.2	24.6	25.0	25.4	25.8	26.2	26.6	27.0	27.4	27.8	28.2	28.6	29.0	
2006	Jan	19.1	19.5	19.9	20.3	20.7	21.1	21.5	21.9	22.3	22.7	23.1	23.5	23.9	24.3	24.7	25.1	25.5	25.9	26.3	26.7	27.1	27.5	27.9	28.3	28.7
2006	Feb	19.4	19.8	20.2	20.6	21.0	21.4	21.8	22.2	22.6	23.0	23.4	23.8	24.2	24.6	25.0	25.4	25.8	26.2	26.6	27.0	27.4	27.8	28.2	28.6	29.0
2006	Mar	19.7	20.1	20.5	20.9	21.3	21.7	22.1	22.5	22.9	23.3	23.7	24.1	24.5	24.9	25.3	25.7	26.1	26.5	26.9	27.3	27.7	28.1	28.5	28.9	29.3
2006	Apr	20.0	20.4	20.8	21.2	21.6	22.0	22.4	22.8	23.2	23.6	24.0	24.4	24.8	25.2	25.6	26.0	26.4	26.8	27.2	27.6	28.0	28.4	28.8	29.2	29.6
2006	May	20.3	20.7	21.1	21.5	21.9	22.3	22.7	23.1	23.5	23.9	24.3	24.7	25.1	25.5	25.9	26.3	26.7	27.1	27.5	27.9	28.3	28.7	29.1	29.5	29.9
2006	Jun	20.6	21.0	21.4	21.8	22.2	22.6	23.0	23.4	23.8	24.2	24.6	25.0	25.4	25.8	26.2	26.6	27.0	27.4	27.8	28.2	28.6	29.0	29.4	29.8	30.2
2006	Jul	20.9	21.3	21.7	22.1	22.5	22.9	23.3	23.7	24.1	24.5	24.9	25.3	25.7	26.1	26.5	26.9	27.3	27.7	28.1	28.5	28.9	29.3	29.7	30.1	30.5
2006	Aug	21.2	21.6	22.0	22.4	22.8	23.2	23.6	24.0	24.4	24.8	25.2	25.6	26.0	26.4	26.8	27.2	27.6	28.0	28.4	28.8	29.2	29.6	30.0	30.4	30.8
2006	Sep	21.5	21.9	22.3	22.7	23.1	23.5	23.9	24.3	24.7	25.1	25.5	25.9	26.3	26.7	27.1	27.5	27.9	28.3	28.7	29.1	29.5	29.9	30.3	30.7	31.1
2006	Oct	21.8	22.2	22.6	23.0	23.4	23.8	24.2	24.6	25.0	25.4	25.8	26.2	26.6	27.0	27.4	27.8	28.2	28.6	29.0	29.4	29.8	30.2	30.6	31.0	31.4
2006	Nov	22.1	22.5	22.9	23.3	23.7	24.1	24.5	24.9	25.3	25.7	26.1	26.5	26.9	27.3	27.7	28.1	28.5	28.9	29.3	29.7	30.1	30.5	30.9	31.3	31.7
2006	Dec	22.4	22.8	23.2	23.6	24.0	24.4	24.8	25.2	25.6	26.0	26.4	26.8	27.2	27.6	28.0	28.4	28.8	29.2	29.6	30.0	30.4	30.8	31.2	31.6	32.0
2007	Jan	22.7	23.1	23.5	23.9	24.3	24.7	25.1	25.5	25.9	26.3	26.7	27.1	27.5	27.9	28.3	28.7	29.1	29.5	29.9	30.3	30.7	31.1	31.5	31.9	32.3
2007	Feb	23.0	23.4	23.8	24.2	24.6	25.0	25.4	25.8	26.2	26.6	27.0	27.4	27.8	28.2	28.6	29.0	29.4	29.8	30.2	30.6	31.0	31.4	31.8	32.2	32.6
2007	Mar	23.3	23.7	24.1	24.5	24.9	25.3	25.7	26.1	26.5	26.9	27.3	27.7	28.1	28.5	28.9	29.3	29.7	30.1	30.5	30.9	31.3	31.7	32.1	32.5	32.9
2007	Apr	23.6	24.0	24.4	24.8	25.2	25.6	26.0	26.4	26.8	27.2	27.6	28.0	28.4	28.8	29.2	29.6	30.0	30.4	30.8	31.2	31.6	32.0	32.4	32.8	33.2
2007	May	23.9	24.3	24.7	25.1	25.5	25.9	26.3	26.7	27.1	27.5	27.9	28.3	28.7	29.1	29.5	29.9	30.3	30.7	31.1	31.5	31.9	32.3	32.7	33.1	33.5
2007	Jun	24.2	24.6	25.0	25.4	25.8	26.2	26.6	27.0	27.4	27.8	28.2	28.6	29.0	29.4	29.8	30.2	30.6	31.0	31.4	31.8	32.2	32.6	33.0	33.4	33.8
2007	Jul	24.5	24.9	25.3	25.7	26.1	26.5	26.9	27.3	27.7	28.1	28.5	28.9	29.3	29.7	30.1	30.5	30.9	31.3	31.7	32.1	32.5	32.9	33.3	33.7	34.1
2007	Aug	24.8	25.2	25.6	26.0	26.4	26.8	27.2	27.6	28.0	28.4	28.8	29.2	29.6	30.0	30.4	30.8	31.2	31.6	32.0	32.4	32.8	33.2	33.6	34.0	34.4
2007	Sep	25.1	25.5	25.9	26.3	26.7	27.1	27.5	27.9	28.3	28.7	29.1	29.5	29.9	30.3	30.7	31.1	31.5	31.9	32.3	32.7	33.1	33.5	33.9	34.3	34.7
2007	Oct	25.4	25.8	26.2	26.6	27.0	27.4	27.8	28.2	28.6	29.0	29.4	29.8	30.2	30.6	31.0	31.4	31.8	32.2	32.6	33.0	33.4	33.8	34.2	34.6	35.0
2007	Nov	25.7	26.1	26.5	26.9	27.3	27.7	28.1	28.5	28.9	29.3	29.7	30.1	30.5	30.9	31.3	31.7	32.1	32.5	32.9	33.3	33.7	34.1	34.5	34.9	35.3
2007	Dec	26.0	26.4	26.8	27.2	27.6	28.0	28.4	28.8	29.2	29.6	30.0	30.4	30.8	31.2	31.6	32.0	32.4	32.8	33.2	33.6	34.0	34.4	34.8	35.2	35.6
2008	Jan	26.3	26.7	27.1	27.5	27.9	28.3	28.7	29.1	29.5	29.9	30.3	30.7	31.1	31.5	31.9	32.3	32.7	33.1	33.5	33.9	34.3	34.7	35.1	35.5	35.9
2008	Feb	26.6	27.0	27.4	27.8	28.2	28.6	29.0	29.4	29.8	30.2	30.6	31.0	31.4	31.8	32.2	32.6	33.0	33.4	33.8	34.2	34.6	35.0	35.4	35.8	36.2
2008	Mar	26.9	27.3	27.7	28.1	28.5	28.9	29.3	29.7	30.1	30.5	30.9	31.3	31.7	32.1	32.5	32.9	33.3	33.7	34.1	34.5	34.9	35.3	35.7	36.1	36.5
2008	Apr	27.2	27.6	28.0	28.4	28.8	29.2	29.6	30.0	30.4	30.8	31.2	31.6	32.0	32.4	32.8	33.2	33.6	34.0	34.4	34.8	35.2	35.6	36.0	36.4	36.8
2008	May	27.5	27.9	28.3	28.7	29.1	29.5	29.9	30.3	30.7	31.1	31.5	31.9	32.3	32.7	33.1	33.5	33.9	34.3	34.7	35.1	35.5	35.9	36.3	36.7	37.1
2008	Jun	27.8	28.2	28.6	29.0	29.4	29.8	30.2	30.6	31.0	31.4	31.8	32.2	32.6	33.0	33.4	33.8	34.2	34.6	35.0	35.4	35.8	36.2	36.6	37.0	37.4
2008	Jul	28.1	28.5	28.9	29.3	29.7	30.1	30.5	30.9	31.3	31.7	32.1	32.5	32.9	33.3	33.7	34.1	34.5	34.9	35.3	35.7	36.1	36.5	36.9	37.3	37.7
2008	Aug	28.4	28.8	29.2	29.6	30.0	30.4	30.8	31.2	31.6	32.0	32.4	32.8	33.2	33.6	34.0	34.4	34.8	35.2	35.6	36.0	36.4	36.8	37.2	37.6	38.0
2008	Sep	28.7	29.1	29.5	29.9	30.3	30.7	31.1	31.5	31.9	32.3	32.7	33.1	33.5	33.9	34.3	34.7	35.1	35.5	35.9	36.3	36.7	37.1	37.5	37.9	38.3
2008	Oct	29.0	29.4	29.8	30.2	30.6	31.0	31.4	31.8	32.2	32.6	33.0	33.4	33.8	34.2	34.6	35.0	35.4	35.8	36.2	36.6	37.0	37.4	37.8	38.2	38.6
2008	Nov	29.3	29.7	30.1	30.5	30.9	31.3	31.7	32.1	32.5	32.9	33.3	33.7	34.1	34.5	34.9	35.3	35.7	36.1	36.5	36.9	37.3	37.7	38.1	38.5	38.9
2008	Dec	29.6	30.0	30.4	30.8	31.2	31.6	32.0	32.4	32.8	33.2	33.6	34.0	34.4	34.8	35.2	35.6	36.0								

Incubation period of 21 days; mortality rate of 10%.

Under the new rules, the percentage of the EEC statutory requirement (1971) of 15 per cent of capital or equivalent (based ratios provide 10% and 10%, respectively).

App. 13a. Major and interactive effects produced by graded levels of dietary lysine and arginine on feed intake of chicks with enterotoxosis.

Day	Effect of feeding	Effect of administration				Feed intake (g/chick)	ANOVA ( $F/F_r > F_{0.05}$ )		
		Affatoxin D Restricted Cd Litter		Share 1010g/w					
		122Arg101Lys	122Arg122Lys	122Arg144Lys	122Arg154Lys				
2	-A	-A	-A	-A	-A	1.34(0.23)	1.48(0.24)	1.27(0.30)	
3	-A	-A	-A	-A	-A	2.31(0.14)	1.95(0.30)	2.91(0.09)	
4	-A	-A	-A	-A	-A	1.11(0.34)	1.45(0.24)	1.53(0.20)	
5	-A	-A	-A	-A	-A	1.79(0.18)	1.56(0.22)	2.15(0.07)	
6	95.18	94.78	101.14	94.78	94.78	2.80(0.039)	1.46(0.21)	1.12(0.37)	
7	223.87	148.89	128.64	128.64	128.64	6.12(0.0031)	1.93(0.29)	0.74(0.82)	
8	147.99	153.8	153.8	153.8	153.8	7.43(0.0026)	6.42(0.24)	0.81(0.72)	
9	172.8	170.8	183.6	183.6	183.6	5.21(0.0035)	0.16(0.52)	0.68(0.62)	
10	200.8	200.8	217.4	217.4	217.4	8.71(0.0003)	0.27(0.64)	0.25(0.61)	
11	234.8	228.8	234.4	234.4	234.4	15.01(0.001)	0.48(0.62)	1.04(0.42)	
12	265.8	261.8	252.4	252.4	252.4	19.81(0.0001)	0.35(0.66)	0.89(0.66)	
13	297.8	289.8	333.4	333.4	333.4	29.10(0.0001)	0.38(0.63)	0.92(0.49)	
14	323.8	323.8	377.4	377.4	377.4	38.61(0.0001)	0.35(0.62)	0.94(0.48)	
15	346.8	359.8	420.6	420.6	420.6	42.01(0.0001)	0.27(0.54)	0.91(0.47)	
16	411.8	393.0	473.2	473.2	473.2	53.61(0.0001)	0.74(0.53)	1.01(0.44)	
17	455.8	438.8	526.4	526.4	526.4	54.51(0.0001)	1.02(0.40)	0.81(0.73)	
18	503.8	482.0	586.4	586.4	586.4	83.91(0.0001)	1.20(0.33)	0.82(0.71)	
19	544.8	528.8	642.4	642.4	642.4	62.11(0.0001)	0.91(0.45)	0.81(0.72)	
20	590.8	524.8	693.4	693.4	693.4	74.01(0.0001)	1.00(0.40)	0.86(0.62)	
21	640.8	618.8	760.6	760.6	760.6	71.14(0.0001)	1.44(0.20)	0.71(0.44)	
22	701.8	648.0	831.4	758.4	719.8	726.8*	2.50(0.03)	0.84(0.44)	
23	757.8	729.8	894.4	821.4	779.8	777.8*	82.41(0.0001)	2.88(0.049)	
24	813.8	785.6	946.4	850.4	841.6	836.8	62.51(0.0001)	3.02(0.047)	
25	874.8	845.6	1019.6	958.4	902.8	910.8	5.26(0.032)	0.56(0.76)	
26	923.8	902.8	1098.4	1015.6	957.6	972.8	40.21(0.0001)	1.38(0.023)	

\*Shows for feed intake within a major effect followed by the same letter are not significantly different at  $P \leq 0.05$ .

Obscure enterotoxosis = 2-hp per g diet.

†Refers to the feed intake of the corresponding enterotoxosis group.

‡Percentage of the NRC dietary requirement (1977) of lysine or arginine (meal ratio provided 102% and 94%, respectively).

§Shows a major effect of administration. Feed x Admin = test for interaction.

Fig. 2 Effect of graded levels of lysine and methionine sulphate with uridine triphosphate.

Whichever [one] it is, for which you have by the time [letter] are not specifically directed at you, [letter]

Applies to the first batch of the corresponding materials produced.

<sup>11</sup> The following section draws on the discussion of the post-industrial transition in Chapter 11 of *Industrial Decline and Post-industrial Transition* (1997) by John G. Turner and Richard J. Williams.

App. 14a. Major and interactive effects produced by graded levels of dietary lysine and arginine on weight gain of chicks with allatoxins.

Day	Effect of feeding			Effect of administration <sup>d</sup>			ANOVA ( $F(Pr > F)$ ) <sup>e</sup>	
	Allatoxin <sup>f</sup> Restricted - Ad libitum	94kg <sup>g</sup> /102kg <sup>h</sup>	122kg <sup>g</sup> /102kg <sup>h</sup>	122kg <sup>g</sup> /122kg <sup>h</sup>	122kg <sup>g</sup> /144kg <sup>h</sup>	Feed	Ad libitum	Feed x Ad libitum
0	-A	-A	-A	-A	-A	-A	1.28(0.27)	0.14(0.94) 0.95(0.53)
2	-A	-A	-A	-A	-A	-A	1.48(0.24)	1.71(0.18)
6	102A	97.1B	98.8B	104A	100B	96C	101AB	2.25(0.002) 2.31(0.002)
9	146A	160B	150B	144A	142A	135B	146A	4.45(0.019)
12	153A	184B	194A	-A	-A	-A	-A	6.18(0.0017) 2.71(0.018)
15	239B	224C	236A	-A	-A	-A	-A	3.91(0.029)
18	294B	287B	325A	-A	-A	-A	-A	17.1(0.0001) 0.93(0.43) 3.31(0.011)
21	353B	355B	417A	-A	-A	-A	-A	32.0(0.0001) 0.38(0.77) 2.34(0.026)
24	24.5B	427B	502A	-A	-A	-A	-A	44.4(0.0001) 0.63(0.61) 2.31(0.040)
25	451B	433B	535A	-A	-A	-A	-A	28.7(0.0001) 1.49(0.24) 2.15(0.071)
26	472B	457B	561A	-A	-A	-A	-A	44.1(0.0001) 1.34(0.38) 2.35(0.027)
								38.4(0.0001) 1.17(0.34) 2.16(0.010)

<sup>a</sup>Mean for the gain in weight within a major effect followed by the same letter are not significantly different at  $P \leq 0.05$ .

<sup>b</sup>Dietary allatoxin 2.5% per g of diet.

<sup>c</sup>Pair-fed to the feed intake of the corresponding allatoxin group.

<sup>d</sup>Percentage of the NRC dietary requirement (1977) of lysine or arginine (basal ration provided 102% and 98% respectively).

<sup>e</sup>Ad libitum = major effect of administration. Feed = major effect of feeding. Feed x Ad libitum = test for interaction.

App. 15. Effect of graded levels of lysine and arginine on feed conversion of chicks with effluvium.

(9.9%). Initial number of 21 chicks per treatment was reduced to 20 chicks at 7 days.

JOURNAL OF POLYMER SCIENCE: PART A: POLYMERS IN ADVANCED TECHNOLOGY

App. 15a. Major and interactive effects produced by graded levels of dietary lysine and arginine on feed conversion of chicks with sitotrichosis.

Day	Effect of feeding		Effect of administration		Feed conversion (g/chick) <sup>a</sup>		ANOVA ( $F(1, r - 1)$ ) <sup>b</sup>		
	Arginine b Restricted c Ad libitum	Arginine d Ad libitum	Arginine e Ad libitum	Arginine f Ad libitum	122Kg <sup>g</sup> /125Lysine	122Kg <sup>g</sup> /145Lysine	Feed	Aditin	Feed x Adtin
3	1.239	1.355A	1.422	1.358	1.308	1.455A	1.328	5.39(0.009)	4.38(0.009) 2.96(0.019)
6	1.63C	1.76B	1.88A	1.68C	1.778	1.91A	1.66C	19.16(0.001)	10.86(0.001) 5.54(0.004)
9	1.72B	1.82B	1.94A	1.78B	1.798	1.99A	1.74B	9.44(0.0003)	7.21(0.003) 3.20(0.003)
12	1.74C	1.88A	1.97A	1.86AB	1.87AB	1.96A	1.81B	9.54(0.0005)	7.21(0.004) 4.20(0.0013)
15	1.89B	1.97A	2.00A	1.93AB	1.95AB	2.02A	1.89B	4.92(0.013)	3.11(0.038) 5.25(0.006)
18	-A	-A	-A	-A	-A	-A	-A	0.37(0.69)	1.45(0.24) 4.46(0.018)
21	-A	-A	2.13A	2.05AB	2.05AB	2.05AB	2.00B	0.80(0.46)	3.27(0.011) 3.40(0.028) 3.27(0.011)
24	2.16A	2.05B	2.11AB	2.21A	2.14A	2.05B	2.02B	4.07(0.023)	7.47(0.0003) 1.74(0.14)
33	-A	-A	A	2.22A	2.19A	2.10B	2.09B	0.83(0.43)	4.34(0.010) 2.51(0.038)
26	-A	-A	-A	2.26A	2.20AB	2.11B	2.10B	0.89(0.42)	3.98(0.015) 1.90(0.11)

<sup>a</sup>Mean of feed conversion rates (feed intake/weight gain) within a major effect followed by the same letter are not significantly different at P<0.05.  
<sup>b</sup>Dietary sitotrichosis = 2.5% part B of diet.

<sup>c</sup>Restricted to the feed intake of the corresponding sitotrichosis group.

<sup>d</sup>Percentage of the NRC dietary requirement (10%) of lysine or arginine (basal ration provided 1021 and 942 respectively).

<sup>e</sup>Aditin = major effect of administration. Feed = major effect of feeding.  
<sup>f</sup>Aditin = test for interaction.

App. 16. Effect of administrations of nutrients and aflatoxins on mortality and also on hepatic weight, enolase and lipid.

values ( $\text{mean} \pm \text{SD}$ ) for a parameter followed by the same letter are not significantly different at  $p < 0.05$ .

\* Fair fed to the seed intake of the corresponding aitoxia group.

**App. 16a.** Major and interactive effects produced by administrations of nutrients and aflatoxin on hepatic weight, moisture and lipid.

Hepatic Parameter	Effect of feeding				Effect of Administration <sup>d</sup>				ANOVA [F(Fr)] <sup>e</sup>									
	Effect of Aflatoxin <sup>b</sup>		Effect of Ad libitum		A		B		C		D		Feed	Admin	Feed x Admin			
	Restricted	Unrestricted	Ad libitum	A	B	C	D	A	B	C	D	F	F	F				
Choline <sup>c</sup>	21.0A	16.3C	16.2C	A	A	A	A	A	A	A	A	18.3(0.0001)	3.00(0.006)	0.41(0.80)				
Weight (g)	A	A	A	A	A	A	A	A	A	A	A	1.98(0.16)	0.36(0.70)	1.01(0.42)				
Moisture (%)	30.3A	13.2B	14.4B	A	A	A	A	A	A	A	A	1.44(0.0001)	1.54(0.23)	1.13(0.36)				
Lipid (% DBW)																		
Folate <sup>c</sup>																		
Weight (g)	A	A	A	A	A	A	A	A	A	A	A	2.26(0.12)	0.07(0.93)	4.11(0.0007)				
Moisture (%)	72.7B	72.6B	74.0A	A	A	A	A	A	A	A	A	7.58(0.0024)	11.01(0.00316)	2.10(0.0007)				
Lipid (% DBW)	28.2A	16.2B	14.1C	A	A	A	A	A	A	A	A	165.0(0.0001)	1.07(0.36)	0.35(0.84)				
Threonine <sup>c</sup>																		
Weight (g)	20.0A	16.4B	16.7B	A	A	A	A	A	A	A	A	65.8(0.0001)	0.41(0.58)	4.50(0.0064)				
Moisture (%)	75.3A	73.3C	74.5B	A	A	A	A	A	A	A	A	49.5(0.0001)	1.61(0.22)	1.77(0.16)				
Lipid (% DBW)	21.0A	15.1B	14.3B	A	A	A	A	A	A	A	A	57.3(0.0001)	0.66(0.53)	7.89(0.0002)				
Lysine <sup>c</sup>																		
Weight (g)	17.5A	12.8C	15.7B	A	A	A	A	A	A	A	A	42.4(0.0001)	0.51(0.61)	0.62(0.65)				
Moisture (%)	A	A	A	A	A	A	A	A	A	A	A	2.43(0.11)	0.84(0.44)	1.87(0.15)				
Lipid (% DBW)	27.5A	14.9B	15.7B	A	A	A	A	A	A	A	A	181.0(0.0001)	0.28(0.76)	1.15(0.35)				
Lysine + arginine <sup>c</sup>																		
Lipid (% DBW)	30.5A	14.1B	15.0B	A	A	A	A	A	A	A	A	17.98	17.78	21.9A	18.7B	272(0.0001)	6.70(0.033)	2.76(0.14)

**Means of a parameter within a major effect followed by the same letter are not significantly different at  $p \leq 0.05$ .**

Basal ratios provided the lowest %NRC values, but 106% of the NRC in the choline study.

App. 17. Effect of dietary or enteropetional administration (IP) of choline on plasma components in plasma of chicks with allatostatin.

Parameter (units)	Allatostatin (1.5 μg/kg of diet)			Restricted feeding			Ad libitum control		
	0	IP	Dietary	0	IP	Dietary	0	IP	Dietary
Gallbladder	4.480-480	5.260-580	4.760-580	7.0-0	5.0-0	5.260-580	6.740-750	7.260-780	7.160-780
Phosphorus, inorganic (mg %)	5.450-550	5.450-510	5.450-510	5.350-580	5.350-580	5.350-580	6.450-630	6.450-630	6.450-630
Total iron (μg %)	7.540-700	8.410-700	6.445-700	10.950-580	11.750-580	9.950-580	11.650-1100	11.750-1100	12.150-800
TIBC <sup>a</sup> /Total	11.950-120	14.650-120	11.950-120	9.750-710	16.650-1100	11.950-1100	13.950-170	13.950-170	16.950-130
Saturated transferin(%)	55.620-520	71.53-500	67.560-700	72.95-700	72.95-700	72.95-700	73.94-730C	73.94-730C	80.95-710B
Glycogen (mg %)	24.05-140	34.71-150	21.90-140	35.25-140	35.25-140	35.25-140	34.85-130	34.85-130	34.85-130
Cholesterol (mg %)	85.440-300	81.850-300	81.850-300	54.150-100	61.250-100	54.150-100	11.050-50	11.050-50	11.050-50
Triglycerides (mg %)	49.650-110A	44.000-120A	44.000-120A	25.950-160B	25.950-160B	25.950-160B	6.450-50	6.450-50	6.450-50
Glycerol (μg %)	1.340-100	2.000-150	1.340-100	3.00-0	0.00-0	1.720-170	2.450-100	2.450-100	3.150-100
Total protein (g %)	0.530-0.580	0.270-0.580	0.530-0.580	1.450-0.948	1.450-0.948	1.450-0.948	1.250-1.120	1.250-1.120	1.450-0.948
Albumin (g %)	0.320-0.350	1.250-1.400	0.320-0.350	1.050-1.180	1.050-1.180	1.050-1.180	1.150-1.100	1.150-1.100	1.150-1.100
Globulins/globulin ratio	0.7450-0.6700	0.5650-0.6700	0.6480-0.1400	0.8500-0.1000	0.9250-0.1100	0.7150-0.0980	0.8180-0.0840	0.8450-0.0840	0.8450-0.0840
Urine urea nitrogen (mg %)	1.5400-0.3800	2.8400-0.5600	2.2400-0.5400	1.5400-0.5800	1.5400-0.5800	1.5400-0.5800	2.000-0.000	2.000-0.000	1.6400-0.1000
Uric acid (mg %)	5.460-0.080	5.460-0.080	5.460-0.080	4.20-0.70C	3.851-1.1C	3.851-1.1C	2.540-0.6C	2.540-0.6C	2.540-0.6C
Total Nitrogen (g %)	0.1200-0.06	0.1200-0.06	0.1200-0.06	0.100-0.0	0.100-0.0	0.100-0.0	0.180-0.05A	0.180-0.05A	0.180-0.05A
Allatostatin phosphate (0/0.320/0.0)	347.250/248A	346.250/232A	350.000/200A	310.000/0.0	310.000/0.0	310.000/0.0	218.000/0.0A	218.000/0.0A	218.000/0.0A
LNT (U/L)	88.150-140	83.950-170A	78.950-160A	50.850-50	61.050-50	44.450-50	44.250-170	44.250-170	44.650-180
AST (U/L)	16.850-117	13.440-120C	13.150-120C	20.650-170C	22.950-120C	21.650-140	18.750-100E	18.750-100E	20.350-78C
AlT (U/L)	6.0	6.0	6.0	80	80	80	80	80	80
SGOT (μmol/l/h) <sup>b</sup>	4.480-110	6.440-90A	4.500-130	6.300-33A	6.340-55A	6.340-55A	6.440-30A	6.440-30A	6.440-30A
SGPT (μmol/l/h) <sup>b</sup>	2.450-70A	2.650-70A	2.950-110	3.050-70B	6.250-13A	6.450-13A	4.250-10B	4.250-10B	6.150-13A
Prolactin (μg/ml)	21.450-11A	19.350-11A	19.350-11A	17.450-11A	17.450-11A	17.450-11A	22.750-50A	18.850-11A	21.050-11A
Choline (μg/ml)	26.050-76	27.480-110B	25.524-41C	36.820-110C	51.540-140B	51.540-140B	46.550-130C	61.250-110A	39.550-110D

\* Values (SDN) for a clinical parameter followed by the same letter are not significantly different at P<0.05.

<sup>a</sup> Creatinine was not detected (< 0.10 μM).

<sup>b</sup> Pair fed to the feed intake of the corresponding allatostatin group.

c No supplemental choline.

d Allatostatin to achieve 133% of the NRC requirement (1977) of choline (basal ration peptides 100%).

e TIBC = total iron binding capacity, LTH = lactate dehydrogenase, gparaparate malonosterate, ALT = alanine aminotransferase, SGOT = aspartate aminotransferase, Gal = galactosyltransferase.

f Not detected at less than 0.1 U/L.

App. 17a. Major and interactive effects produced by dietary or intraperitoneal (IP) of choline on various components in plasma of chicks with effluations.

Parameter (units)	Effect of feeding		Effect of administration		ANOVA ( $F$ [ $F_c > F$ ]) <sup>a</sup>	Feed $\times$ Admin
	Alfalfa restricted Ad libitum	IP choline	IP choline	Admin		
Calcium (mg/l)	4.888	6.976	7.174	A	84.710 (0.001)	1.77 (0.17)
Phosphorus, inorganic (mg/l)	5.18C	5.67B	6.45A	A	39.110 (0.001)	0.310 (0.73)
Total iron (µg/l)	73.4C	107B	94.10A	92.10A	5.310 (0.032)	1.84 (0.15)
TIBC <sup>b</sup> (µg/l)	112B	168A	157A	153B	17.710 (0.008)	3.340 (0.023)
Saturated transferin (%)	65.0B	70.2B	79.2A	75.0A	65.3B	4.49 (0.031)
Fatty acid (mg/ml)	A	A	A	A	18.810 (0.006)	1.47 (0.29)
Glycose (mg/l)	249B	251A	252A	A	38.110 (0.001)	0.40 (0.87)
Cholesterol (mg/l)	33.9C	130A	113B	A	33.010 (0.001)	1.20 (0.12)
Triglycerides (mg/l)	798A	592C	645AB	677AB	705A	3.19 (0.023)
Glycerol (µg/l)	4.41A	2.32B	1.87B	A	31.710 (0.001)	1.29 (0.12)
Total protein (g/l)	1.52C	2.92B	20.2A	2.42B	2.60A	3.46 (0.013)
Albumin (g/l)	0.339B	1.23A	1.38A	1.08B	1.15A	1.07 (0.12)
Globulin (g/l)	0.938B	1.58A	1.64A	1.23B	1.45A	1.37 (0.12)
Albumin/globulin ratio	0.478	0.85A	0.84A	A	18.410 (0.001)	0.72 (0.45)
Blood urea nitrogen (mg/l)	2.17A	1.42B	1.92A	A	5.910 (0.075)	1.97 (0.16)
Uric acid (µg/l)	5.50B	3.91C	2.43A	A	14.010 (0.001)	0.54 (0.43)
Total bilirubin (µg/l)	A	A	A	A	5.50 (0.01)	0.34 (0.45)
Alkaline phosphatase (U/L)	A	A	A	A	2.00 (0.16)	0.50 (0.74)
Lact (µV/L)	831A	521B	438C	634A	605A	0.24 (0.79)
AST <sup>c</sup> (U/L)	171C	216A	194B	187B	205A	15.110 (0.001)
Choline (µg/ml)	26.4B	48.1A	49.4A	36.4B	44.3A	5.310 (0.008)
					24.105 (0.001)	5.910 (0.007)
						2.84 (0.04)

<sup>a</sup>Name for a clinical parameter within a major effect followed by the same letter is not significantly different at P<0.05.

<sup>b</sup>Pair fed to the feed intake of the corresponding effluation group.

<sup>c</sup>No supplemental choline

TIBC = total iron binding capacity, AST = aspartate aminotransferase,

Administration to achieve NRC requirement (1977) of choline (basal ration provided).

Admin = major effect of administration. Feed = major effect of feeding. Feed  $\times$  Admin = test for interaction.

Effluation = 2.5 % per of diet.

App. 18. Effect of graded levels of dietary folic acid on various' components in plasma of chicks with allatostatin.

Parameter (units)	Plasma parameter <sup>a</sup>		
	Allatostatin (2.5, nM/L of diet) <sup>b</sup>	Allatostatin (5, nM/L of diet) <sup>b</sup>	Allatostatin Control <sup>b</sup>
Calcium	7.444	7.444	7.444
Phosphorus, inorganic (mg %)	7.440-4.662	7.890-3.956	7.540-5.102
Total iron (μg %)	6.090-1.962	5.890-1.887	6.150-3.700
Total (mg %)	7897.50	7959.44	11255.44
Saturated triglycerin (mg %)	1114.312	1725.2946	12541.2486
Cholesterol (mg %)	7291.398	6931.46	6952.648
Cholesterol ratio	4737.0C	4556.4C	4156.4C
Triglyceride (mg %)	69.4420A	75.1425A	11165.5A
Glycogen (μg/dL)	4161.058	4912.031	3295.006
Total protein (g %)	1.750-2.26	1.750-1.50	1.500-1.24
Albumin (g %)	0.950-1.17C	0.880-1.05C	1.550-0.96C
Globulin (g %)	1.000-0.88	1.000-1.18	0.930-1.08
Albumin/globulin ratio	0.450-1.53	0.480-0.72	1.450-1.23
Blood urea nitrogen (mg/dL)	2.550-5.58	2.000-0.020	0.860-0.08
Uric acid (mg %)	5.040-4.89E	5.160-4.62CD	4.250-4.1E
Total bilirubin (mg %)	0.3250-0.05A	0.2800-0.02AB	0.2500-0.02AB
Creatinine (mg %)	80	80	80
Lipid profile <sup>c</sup> (mg %)	383007238C 393954977A 340154693C	383020320AC 41329511A8	4005112753AB 445275AB 31351128A
Lipop (U/L)	857591AB	891523AA	465527CD
AST (U/L)	1818.50	21251.5A	182451.5B
ALT (U/L)	1.851-1.08CD	4.010-0.82A	1.251-0.72C
SALD (μmol/L/dL)	5.150-7.28A	6.650-3.94BC	4.250-7.10CE
GALD (μmol/L/dL)	67.155.5A	55.652.7BC	60.957.6A
Whole blood plasma (μM)	4.1652.2A	15.856.4B	15.052.1A
Folic acid (μM/L)	1715158C	3592606A	3462528AB 43.45±0C

<sup>a</sup> Values (mean ± SD) for a clinical parameter followed by the same letter are not significantly different at P<0.05.<sup>b</sup> Fed to the feed intake of the corresponding allatostatin group.<sup>c</sup> Percentage of the NRC dietary requirement (1977) of folic acid ration provided 245.42%.<sup>d</sup> TAC = total iron binding capacity, TBC = lactate dehydrogenase, AST = aspartate aminotransferase, ALT = alanine aminotransferase, GAL = galactosyltransferase, GALC = galactosyltransferase.<sup>e</sup> ND = not detected at less than 0.01 mg %.<sup>f</sup> ND = not detected.

App. 1B Major and interactive effects produced by graded levels of dietary folic acid on various components in plasma of chicks with arachidonitis.

Parameter (units)	Plasma parameter <sup>a</sup>		Effect of administration <sup>b</sup>	Alpha [(Prc-F)] <sup>c</sup>		
	Effect of feeding	Aflatoxin/B restricted Ad libitum		Feed	Adnin	Feed vs Adnin
Calcium (mg%)	7.578	9.07A	9.23A	.A	.A	.A
Phosphorus, Integ. (mg%)	5.95C	7.30A	6.55B	.A	.A	.A
Total iron (ug%)	10.9A	10.7A	10.7A	.A	.A	.A
TIBC (ug%)	119.8	127.6B	130.8A	.A	.A	.A
Saturated transferin (%)	70.1C	79.8B	86.8A	.A	.A	.A
Glucone (mg%)						
Cholesterol (mg%)	28.8B	29.1A	28.2B	.A	.A	.A
44.3C	111.0A	89.0B				
Triglycerides (mg%)	A	A	A	.A	.A	.A
Glycerol (ug%)	44.5A	33.90A	29.5B	.A	.A	.A
Total protein (g%)	1.60B	2.63A	2.75A	2.50A	2.36AB	2.33B
Albumin (g%)	0.53B	1.38A	1.31A	.A	.A	.A
Globulin (g%)	0.50B	1.45A	1.44A	1.34A	1.26AB	1.25AB
Albumin/globulin ratio	0.94B	0.59A	0.59A	.A	.A	.A
Blood urea nitrogen (ug%)	.A	.A	.A	.A	.A	.A
Uric acid (mg%)	4.80C	6.23A	5.92B	.A	.A	.A
Total bilirubin (ug%)						
Unconjugated bilirubin (ug/L)	2652B	4239AB	4862A	.A	.A	.A
Conjugated bilirubin (ug/L)	848A	491B	455B	.A	.A	.A
AST (U/L)						
ALT (U/L)	212A	215A	195B	.A	.A	.A
Fatty acid, plasma (ug/ml)						
Palmitic acid, s/bt (ug/ml)	12.3A	6.59B	13.1A	.A	.A	.A
Stearic acid (ug/ml)	292A	51.4C	194B	101B	202AB	234A

<sup>a</sup>Means for a clinical parameter within a major effect followed by the same letter are not significantly different at P<0.05.

<sup>b</sup>Significant effect of the intake of the corresponding Aflatoxin group.

<sup>c</sup>Significant effect of the intake of the corresponding Aflatoxin group.

Alpha = total iron binding capacity, IBC = lactate dehydrogenase, AST = aspartate aminotransferase, ALT = alanine aminotransferase, the NRC dietary requirement (1972) of feline basal ration provided 2441.

Major = major effect of administration, Feed = major effect of feeding feed & Adnin = test for interaction.

App. 19. Effect of graded levels of dietary thiomine on the plasma biochemistry of chicks with astatocontics.

Parameter (units)	Astatoxin (2.5 mg/kg diet)			Restricted feeding			Control		
	128	135	179	128	135	179	128	135	179
Calcium	4.240-2.262	4.240-3.375	4.440-3.375	6.240-3.388	6.240-3.388	6.540-3.388	6.640-4.288	6.640-4.288	6.440-3.288
Phosphorus, inorganic (mg/dl)	4.140-1.182	4.140-1.182	4.140-1.182	4.460-1.300	4.460-1.300	4.460-1.300	5.110-1.324	5.110-1.324	4.460-1.324
Total iron (μg/dl)	80.85±14.5	51.85±16.0	62.25±15.0	110.95±18.0	110.95±18.0	110.95±18.0	110.95±18.0	110.95±18.0	110.95±18.0
TIBC	18.95±4.82	11.95±3.00	13.65±2.00	17.71±2.120	17.71±2.120	17.71±2.120	16.65±2.140	16.65±2.140	17.25±2.140
Saturated triterpenoids (μg/dl)	66.85±1.48	43.65±1.60	76.62±3.60	66.85±1.48-C	66.85±1.48-C	66.85±1.48-C	62.51±1.68	62.51±1.68	62.51±1.68
Cleicase (ug/dl)	165.74±3.20	238.95±3.40	239.80±3.40	231.56±5.68	231.56±5.68	231.56±5.68	236.54±3.44	236.54±3.44	232.50±4.48
Cholesterol (mg/dl)	50.55±4.00	42.55±2.50	47.55±6.50	13.25±6.68	14.95±5.58	13.25±5.58	10.95±4.95	12.75±4.18	7.05±0.50
Triglycerides (mg/dl)	77.45±2.10	108.65±6.6	111.56±1.16	86.05±4.84	119.51±2.0	121.51±1.84	88.24±6.84	88.54±6.84	4.45±0.50
Glyceral (mg/dl)	379.51±6.6A	253.95±10.8BC	315.95±14.3AB	79.55±6.2D	109.51±10.2CD	82.51±9.50	82.05±9.50	188.91±10.9BC	25.51D
Total protein (g/dl)	1.360-2.95	1.360-2.45C	1.440-2.95C	2.940-1.88B	2.840-1.04A	2.840-1.04A	2.940-1.04A	2.940-1.04A	2.240-4.18
Albumin/globulin ratio	0.4010±0.12C	0.3545±0.10C	0.3485±0.10C	1.1610±0.08A	1.1610±0.08A	1.1610±0.08A	1.1610±0.08A	1.1610±0.08A	1.1610±0.08A
Blood urea nitrogen (mg/dl)	0.4450±0.08	0.3920±0.08	0.3920±0.08	0.6160±0.08A	0.6160±0.08A	0.6160±0.08A	0.6160±0.08A	0.6160±0.08A	0.6160±0.08A
Uric acid (mg/dl)	1.09±0.82A	ND	1.09±0.84A	0.59±0.10A	ND	0.59±0.10A	ND	0.59±0.10A	ND
Total bilirubin (mg/dl)	6.449±0.83A	5.461±0.86C	5.749±0.77BC	7.251±2.4	4.749±0.94C	4.651±0.78B	6.281±0.66C	6.281±0.66C	7.450±0.67A
Alkaline phosphatase (USS/100ml)	0.050±0.05A	0.050±0.05A	0.050±0.05A	0.050±0.05A	0.050±0.05A	0.050±0.05A	0.050±0.05A	0.050±0.05A	ND
Lecithin/β-alanide (mg/ml)	320.95±0.0	320.95±0.0	320.95±0.0	320.95±0.0	348.95±0.0	348.95±0.0	350.95±0.0	350.95±0.0	344.95±0.0
ATP (U/ml)	771.45±1.5A	394.55±1.5A	432.55±1.5A	448.95±1.5A	448.95±1.5A	448.95±1.5A	333.45±1.5A	333.45±1.5A	417.95±1.5A
ALT (U/ml)	160.95±1.5A	132.55±1.5C	160.95±1.5A	123.25±1.0C	110.95±1.5C	129.55±1.5C	134.95±1.5C	134.95±1.5C	160.95±1.5A
AST (U/ml)	ND	ND	ND	ND	ND	ND	ND	ND	ND
GSH (μmol/ml)	5.65-5.84B	5.15-5.65AB	5.15-5.65AB	5.17-5.74BC	5.05-5.65AB	4.75-5.44C	5.87-5.84A	5.72-5.82ABC	5.140-5.35A
Packed cell volume (%)	31.51±3.0D	30.74±2.20	32.46±5.5C	32.46±4.5CD	31.51±4.5CD	31.62±4.5CD	35.02±3.2B	35.12±3.2C	37.25±4.1A
Folate acid (μg/dl)	8.99±1.6A	5.16±0.56A	5.17±0.51A	5.05±1.59A	5.05±1.59A	5.05±1.59A	11.45±1.4A	5.93±0.53A	10.04±0.82A

Values (mean ± standard deviation) for a climated parameter followed by the same letter are not significantly different<sup>a,b,c,d</sup>, at  $P \leq 0.05$ .

b/c = fed to the feed intake of the corresponding astatoxin group.

c/d = percentage of the NMC dietary requirement of thiomine (NaCl ration provides 125%).

d/e = total iron binding capacity, TBC = lactate dehydrogenase, AST = aspartate aminotransferase, ALT = alanine aminotransferase, GSH = γ-glutamyltransferase;

e/f = Not Detected.

App. 19a Major and interactive effects produced by graded levels of dietary threonine on the plasma biochemistry of chicks with aflatoxinosis.

Parameter [units]	Plasma parameter				Alpha [F (P>F)] <sup>a</sup>
	Effect of feeding		Effect of administration <sup>b</sup>		
Methionine <sup>c</sup> Restricted Ad libitum	120	155	179	Feed x Admin	
Calcium (mg%)	4.256	6.694	6.458	A	A
Phosphorus, inorganic (mg%)	4.125	4.244	4.084	A	A
Total iron (ug%)	50.38	55.148	109A	A	A
TIBC (ug%)	111B	175A	142B	A	A
Saturated transferrin (%)	47.06 <sup>d</sup>	66.5A	66.5A	A	A
Packed cell volume (%)	31.98	32.38	35.6A	A	A
Glycose (ug%)	250C	283A	271B	A	A
Cholesterol (ppm)	46.9C	132A	93.9B	106A	92.4B
Triglycerides (ug%)	-	A	A	A	A
Glyceral (ug%)	316A	68.8B	104B	A	A
Total protein (g%)	1.200	2.88A	2.78A	A	A
Albumin (g%)	0.308	1.13A	1.08A	A	A
Globulin (g%)	0.310	1.75A	1.70A	A	A
Albumin/globulin ratio	0.486	0.65A	0.65A	A	A
Blood urea nitrogen (ug%)	A	A	A	A	A
Uric acid (ug%)	7A	A	A	6.88A	5.54B
Total bilirubin (ug%)	A	A	A	A	6.71A
Alkaline phosphatase (U/L)	A	A	A	A	1.1510 (0.33)
Lip <sup>e</sup> (U/L)	178BA	443B	401B	3500A	3360A
AST (U/L)	-	162A	125C	-	A
AST <sup>f</sup> (U/L)	-	A	A	A	A
Fatty acid (ug/ml)	5.23A	9.10A	5.68B	B, 5.6AB	6.92D
Insulin (ug/ml)	16.8A	5.44B	14.9A	A	A
Glutathione (nmol/ugbw)	306B	222C	472A	377A	228B
					418A 19.4 (0.0001) 98.2 (0.0006) 12.0 (0.0001)

<sup>a</sup>Means for a clinical parameter within a major effect followed by the same letter are not significantly different at P<0.05.

<sup>b</sup>Unilateral aflatoxin = 2.5 mg per g of diet.

<sup>c</sup>Refers to the level of the corresponding aflatoxin group.

<sup>d</sup>Percentage of the dietary requirement (1971) of lysine (basal ration provided 128%).

<sup>e</sup>TIBC = total iron binding capacity, U/L = lactate dehydrogenase, AST = aspartate aminotransferase, ALAT = alanine aminotransferase.

<sup>f</sup>Major effect of administration. Feed = major effect of feeding. Feed x Admin = test for interaction.

App. 20. Effect of graded levels of dietary lysine on the plasma biochemistry and hepatic glutathione of chicks with elatostatinosis.

Parameter (units)	Elatostatin (2.5 µg/g of diet)		Control	
	Untreated	Restricted Feeding <sup>b</sup>	Untreated	Restricted Feeding <sup>b</sup>
Ca:Cr (eq 2)	1.02 <sup>c</sup>	3.22 <sup>c</sup>	1.02	1.22
Phosphorus, %P <sub>2</sub> O <sub>5</sub> (eq 2)	4.81±0.256	4.50±0.220	6.33±0.410	6.25±0.310
Total iron (mg/g)	74.35±2.92	64.22±0.68	53.90±0.238	57.00±0.144
TIBC (mg/g)	109.22±6.76	71.35±7.50	61.25±9.30	12.67±0.748
Saturated transferrin (%T)	67.54±2.32	67.35±1.28	70.61±1.96	77.11±1.58
Packed cell volume (%)	25.84±1.58	26.51±1.38	27.49±2.58	29.00±2.58
Glucose (mg/dl)	57.61±4.90	53.54±1.90	48.01±4.50	48.06±1.14
Cholesterol (mg/dl)	67.05±2.20	84.46±2.20	109.61±0.60	120.69±0.60
Total triglycerides (mg/dl)	45.31±1.66	41.65±0.88	48.14±2.54	101.04±4.46
Total protein (g/dl)	1.43±0.198	1.18±0.130	1.11.13±0.210	2.43±0.154
Albumin (g/dl)	0.38±0.035	0.58±0.100	0.50±0.035	1.15±0.038
Globulin (g/dl)	0.88±0.138	0.60±0.086	0.63±0.132	1.20±0.084
Albumin/globulin ratio	0.67±0.030	0.96±0.015	0.80±0.016	1.35±0.015
Blood urea nitrogen (mM)	1.83±0.574	1.12±0.540	1.50±0.368	2.05±0.490
Uric acid (mg/dl)	5.48±0.548	5.48±1.948	4.80±0.548	5.93±1.448
Total bilirubin (mg/dl)	0.20±0.004	0.20±0.004	0.18±0.004	0.18±0.004
Alkaline phosphatase (U/l)	3889±1450	4460±1100	2300±943	3200±390
Lact <sup>d</sup> (U/l)	70±19.8	75±7.8	45±2.2	42±5.2
AST (U/l)	137±13.5	148±1.5	133±5.30	205±1.36
ALT (U/l)	0.09±0.08	7.5±9.6A	0.01±0.08	2.5±5.0A
Fatty acid (µg/ml)	9.6±0.258	8.6±0.141	8.6±0.141	5.7±0.562
Insulin (µU/ml)	17.9±2.5A	16.0±2.5A	16.4±2.5A	13.1±2.5A
Glucagon (ng/ml)	31.9±1.5B	26.3±1.9B	23.5±1.1B	21.5±1.1A
Glutathione (nmol/g web)	320±16.8	343±3.8B	320±16.8	275±12.8C

<sup>a</sup> Values (mean ± standard deviation) for a clinical parameter followed by the same letter are not significantly different.

<sup>b</sup> At 2.0%.

<sup>c</sup> Paired fed to the feed intake of the corresponding elatostatin group.

<sup>d</sup> Percentage of the NRC dietary requirement (1977) of lysine (basal ration provides 101%).

LDH = total serum lactate dehydrogenase, ALT = aspartate aminotransferase, AST = alanine aminotransferase.

<sup>e</sup> At 2.0%.

<sup>f</sup> ...

Fig. 204. Major and interactive effects produced by graded levels of dietary lysine on the plasma biochemistry of chicks with glutathionuria.

Parameter (units)	Plasma parameters				Age (days)	Feed ( $\text{g}/\text{kg}^{-1}$ )	
	Effect of feeding	Effect of distribution	Effect of diet	Effect of time			
Affiliation (kg $^{1/2}$ )	4.370	6.884	-	-	68.50 (0.001)	0.340 (0.1)	0.370 (0.1)
Phosphorus, non- nitrogenous (mg $^{1/2}$ )	6.174	5.534	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Protein, non- nitrogenous (mg $^{1/2}$ )	69.124	121.4	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen (mg $^{1/2}$ )	79.418	82.486	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	82.486	82.486	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	27.408	27.408	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	30.034	30.034	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	53.4C	53.4C	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	12.6A	9.5B	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	9.5B	9.5B	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	4.4A	4.4A	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	22.0B	17.0B	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	2.44A	2.44A	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	1.12A	1.12A	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	1.12A	1.12A	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	0.70B	1.2A	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Urea nitrogen, diet content (mg $^{1/2}$ )	1.2A	1.2A	-	-	4.70 (0.001)	0.890 (0.4)	2.11 (0.4)
Albumin/globulin ratio	1.52A	0.43C	1.00B	-	2.34 (0.12)	0.740 (0.079)	2.460 (0.069)
Total urea nitrogen (mg $^{1/2}$ )	5.25B	6.18B	7.28A	-	1.26 (0.31)	1.24 (0.30)	1.31 (0.29)
Total bilirubin (mg $^{1/2}$ )	70.1A	44.8B	39.7C	-	20.20 (0.001)	0.00 (0.10)	0.40 (0.61)
Urea nitrogen, phosphate (U/L)	166C	198A	174A	-	8.05 (0.001)	0.00 (0.10)	0.25 (0.51)
Urea nitrogen, phosphate (U/L)	166C	198A	163B	-	1.71 (0.19)	0.25 (0.19)	0.58 (0.21)
Urea nitrogen, phosphate (U/L)	166C	198A	174A	-	0.35 (0.48)	0.11 (0.34)	0.58 (0.21)
Urea nitrogen, phosphate (U/L)	166C	198A	163B	-	96.10 (0.001)	0.47 (0.43)	1.71 (0.20)
Urea nitrogen, phosphate (U/L)	166C	198A	174A	-	1.63 (0.001)	0.47 (0.43)	1.41 (0.20)
Urea nitrogen, phosphate (U/L)	166C	198A	163B	-	8.94A	0.760 (0.51)	0.550 (0.19)
Urea nitrogen, phosphate (U/L)	166C	198A	174A	-	8.94A	0.760 (0.51)	0.550 (0.19)
Urea nitrogen, phosphate (U/L)	166C	198A	163B	-	8.94A	0.760 (0.51)	0.550 (0.19)

Glutathione (mol/g wet) 3068 222C effect followed for 1-2 days

P < 0.05;

proteolytic *aflatoxin* = 2.5ug per g of diet.

pair-set to the intake of the corseid

percentage of the NRC dietary requirement

• total mean binding capacity, LDI.

### **Amidotransferase**

Administer a major effect of administration.

卷之三

卷之三

卷之三

卷之三

• 4

卷之三

卷之三

App. II. Effect of dietary or interperitoneal administration (IP) of colchicine on concentrations of free amino acids (FAAs) and other sulphur positive substances (MRS) in plasma of chicks with alastrism.

**Table 2.** Values (mean  $\pm$  SEM) for PAA or VPS followed by the same letter are not significantly different at  $P < 0.05$ .

feed intake of the corresponding lactation group.

e no suplemento 1 chalilat.

Administration to achieve 175% of the WMC requirement (1977) of chlorine bleach ration 106.7 kg/2000 kg total chlorine available. It was decided at the time that the chlorine bleach ration would be 9.4 metric tons per day.

ESTATE PLANNING

Fig. 2b. Major and interactive effects produced by dietary or intraperitoneal administration of choline on concentrations of free-fatty acids (FFA) and other elutroposin positive substances (EP) in plasma of chicks with affluvitis.

	FFA or EP (nmol/dl) <sup>a</sup>		Effect of administration of system		Effect of diet		Effect of diet & administration of system	
	Effect of diet	Effect of administration of system	Feed	Abuse	Feed & Abuse	Abuse (FFA > EP)		
Affluvitis	553.8	442.0	617.6	-	12.7 (0.0001)	6.4 (0.533)	2.5 (0.042)	
Choline+Metabolic acid	15.48	25.56	14.38	15.76	16.36	23.6 (0.0001)	15.4 (0.0001)	
Ammonium	260.0	235.9	232.6	238.0	237.6	2.4 (0.023)	2.4 (0.023)	
Ammonium	51.64	24.88	48.52	43.86	40.07	2.2 (0.033)	2.2 (0.033)	
Aspartic acid	143.8	76.46	281.6	193.6	193.6	7.4 (0.012)	7.4 (0.012)	
Aspartic acid	161.44	63.13	45.38	55.18	70.76	2.0 (0.0001)	5.3 (0.001)	
Citric acid	-	-	-	6.186	4.428	4.238	6.2 (0.0001)	
Citric acid	10.39	10.39	13.74	10.48	12.36	9.73 (0.0001)	5.5 (0.014)	
Cysteathione	6.37 (C)	34.18	18.76	23.22	20.18	23.4 (0.0001)	6.3 (0.0004)	
Cysteathione	6.37 (C)	34.18	18.76	23.22	20.18	23.4 (0.0001)	6.3 (0.0004)	
Glutamic acid	62.46	71.21	90.24	-	-	41.0 (0.0001)	6.4 (0.023)	
Glutathione	20.54	6.818	10.39	10.39	10.39	14.2 (0.0001)	2.4 (0.033)	
Glutathione	115.9	117.8	138.4	126.8	126.8	11.9 (0.0001)	4.2 (0.013)	
Glycine	162.14	61.72	81.58	-	-	21.5 (0.0001)	2.4 (0.013)	
Glycine	118.8	200.0	81.84	-	-	22.4 (0.0001)	2.9 (0.012)	
Histidine	151.6	65.49	132.4	-	-	12.9 (0.0001)	6.1 (0.004)	
Histidine	-	-	-	7.5 (0.0001)	7.5 (0.0001)	1.2 (0.211)	6.1 (0.0006)	
Hydroxyproline	112.0	121.9	138.4	-	-	6.7 (0.0001)	6.7 (0.0002)	
Hydroxyproline	-	-	-	12.2 (0.0001)	12.2 (0.0001)	2.1 (0.014)	0.1 (0.951)	
Isoleucine	124.8	107.0	121.6	123.8	123.8	1.0 (0.0001)	2.8 (0.004)	
Lysine	138.6	139.0	132.4	-	-	4.8 (0.0001)	4.8 (0.0001)	
Lysine	-	-	-	4.2 (0.0001)	4.2 (0.0001)	2.2 (0.0001)	2.2 (0.0001)	
methionine	65.09	82.26	92.46	64.26	67.26	81.4 (0.0001)	2.3 (0.0001)	
methionine	-	-	-	10.18	10.18	2.4 (0.0001)	4.4 (0.004)	
3-methylbutyrate	17.76	14.70	14.56	14.60	15.39	14.3 (0.0001)	5.2 (0.0001)	
Phenylalanine	89.78	79.26	112.6	-	-	14.1 (0.0001)	2.0 (0.015)	
Proline	27.04	96.46	14.89	-	-	40.5 (0.0001)	0.1 (0.976)	
Proline	25.38	23.72	4.372	-	-	13.0 (0.0001)	0.4 (0.932)	
Arginine	6.20C	8.92	7.718	6.238	7.718	13.2 (0.0001)	6.3 (0.010)	
Arginine	23.14	14.88	14.58	-	-	5.2 (0.0001)	1.1 (0.023)	
Threonine	56.20	57.88	95.16	74.56	65.63	5.0 (0.0001)	1.1 (0.027)	
Threonine	22.24	11.12 <sup>c</sup>	16.82	-	-	43.4 (0.0001)	1.0 (0.003)	
Threonine	21.88	20.62	24.64	27.84 <sup>b</sup>	20.04	11.9 (0.0001)	4.2 (0.003)	

Note: FFA or EP within a major effect followed by the same letter are not significantly different at P<0.05.

<sup>a</sup>Significant difference = 2.297 nmol/dl.

<sup>b</sup>Excluded from the feed batch of the corresponding elutroposin group.

<sup>c</sup>No significant difference.

Administration to achieve 151% of BMC requirement (1972) of choline (chick ration provided (1981)).

Table = Major effect of administration. Feed = major effect of feeding. Feed & Abuse = test for interaction.

App. 22. Effect of graded levels of dietary folic acid on concentrations of free-methionine (FMA) and other sulphuric acid substances<sup>a</sup> in plasma of chicks with ascites.

FMA or MPS	Free-methionine substances (mg/mℓ of plasma)			Control
	Atmospheric level of diet	544C level of diet	Restricted feeding	
Alanine	336±56C	484±56C	735±56A	735±56A
γ-Aminobutyric acid	2.15±.24	4.05±.24	2.45±.14	2.44
α-Aminoglyclic acid	9.44±.68Y	8.15±.37	3.52±.34	6.51±.34
Ammonia	23.52±4.48B	29.25±4.44A	23.52±3.24B	23.52±3.24B
Aspartic acid	1015±9C	1395±58B	1079±48B	1055±44B
Cysteine	1129±12A	1129±11A	49±4.2D	211±6.4AB
Glutamine	5.01±.48CD	5.00±.70ACD	5.06±.48A	5.05±.31A
Glutathione	5.39±.18	5.39±.18	4.35±.18	4.35±.18
Optic acid/phosphoarginine	1.52±.14A	1.52±.14A	1.52±.14A	1.52±.14A
β-Alanine	2.25±.22A	2.15±.21A	2.15±.21A	2.15±.21A
β-Hydroxybutyrate	1.67±.23A	1.67±.23A	1.67±.23A	1.67±.23A
β-Hydroxyvaleric acid	1.15±.18A	1.15±.18A	1.15±.18A	1.15±.18A
Choline	1125±11A	987±11A	1125±11A	1125±11A
Glycine	72.52±13B	70.52±10B	72.52±13B	72.52±13B
Methionine	12.85±.52C	12.85±.52C	12.85±.52C	12.85±.52C
Nicotinamide	3.96±.21AB	3.15±.16AC	4.00±.06A	2.46±.43C
Hydroxyproline	16.61±1.8D	17.06±2.0D	20.57±.98D	20.57±.58B
Taurine	18.01±1.1C	18.51±1.4C	9.69±.4C	12.15±.8A
Lysine	19.01±1.6CD	21.27±2.5CD	18.71±.26	13.03±.54A
Arginine	20.63±4.6C	21.73±3.9C	16.67±.50	16.67±.50
Homocysteine	11.75±.23A	11.75±.23A	11.75±.23A	11.75±.23A
Homocysteine	11.75±.23A	11.75±.23A	11.75±.23A	11.75±.23A
Ornithine	12.85±.52C	12.85±.52C	12.85±.52C	12.85±.52C
Phospholamban	1.66±.20	1.44±.48CD	1.51±.47CD	1.44±.44AB
Prolidine	4.53±.1CD	7.15±.5A	4.05±.92	6.51±.43B
Serine	20.63±4.6C	24.52±2.1A	20.63±4.6B	11.54±.7C
Taurine	21.92±2.0C	49.94±0.7	34.62±2.0B	9.25±0.9A
Threonine	56.32±0.8F	47.94±0.7	61.63±1.8C	88.65±6.6B
Tryptophane	33.63±1.0D	32.52±1.0D	30.95±2.0D	10.93±1.2C
Tyrosine	54.43±1.0C	66.53±.59	50.94±.5C	8.62±.4A
Valine	22.84±1.9A	19.92±2.0A	22.82±1.9C	12.97±.4D
	23.25±2.0AC	23.25±2.0AC	23.25±2.0AC	23.25±2.0AC

<sup>a</sup> Values (means ± SD) for FMA or MPS followed by the same letter are not significantly different at P < 0.05. BD = Not detectable at levels less than of 0.1 mℓ folic acid.

<sup>b</sup> Determined by a radioisotope assay based on the method of Hsu et al. (1961), modified by Kornblith et al. (1962).

<sup>c</sup> Percentage of the NRC dietary requirement (1959)<sup>d</sup> of folic acid ration provided 2443D.

The first and earliest effects produced by *Streptomyces* spores on *Neurospora* seedlings were described by Friesenbeck and colleagues (1948) and Schärer

These two files are very similar, except that different file names are used.

Intensity of lesions = 1.50 per 1 ml. dose.

卷之三

THE JOURNAL OF CLIMATE

Effect of purified levulinic acid on concentrations of free-sulfuric acid (TSA) and other aldehydes in positive substances

Wavelengths (nm & standard deviations) for FIA or WPS followed by the same letter are not significantly different at P<0.05.  
N.D.: Not detectable at levels less than 0.1 nmol/L. a: sodium acid, anerine, y-anisidine, sulfonate, benzoquinone, methionine sulfoxide, carnosine, diacetylcarnosine, glycosaminoglycans, heparan sulfate, heparin, homocysteine, methionine.

J. Neurosci., November 1, 2006 • 26(44):11875–11886 • 11885

Percentages of the NRC dietary requirement (1977) of thiamine that rat provides 1387

This major and minor  $\beta$ -agonist effect produced by graded levels of the diet on growth rate of *Fusco-nation* male (FM) and other alpaca is positive substance (PS) in plasma of chicks with ascites.

Mean fat PDI at 60° within a main effect followed by the same letter are not significantly different at  $P < 0.01$ .

10

THE JOURNAL OF CLIMATE

Table 2 gives the effect of whole wheat bran feed on major effect of feeding. Feed is whole or bran for lactation

App. 2a. Effect of graded levels of dietary lysine on concentrations of free-amino acids (FAA) and other nitrogen positive substances (MRS) in plasma of chicks with ascites.

FAA or MRS	Proteinase, acid or alkali-catalyzed hydrolysis		Unrestricted Feeding		Control	
	10% 100% 1000%	12% 100% 1000%	10% 100% 1000%	12% 100% 1000%	10% 100% 1000%	12% 100% 1000%
Alanine	45196.588 46615.038	39951.508 44691.558	54941.163 53468.848	54941.163 53468.848	59495.254 59495.254	59495.254 59495.254
α-Aketo-β-hydroxy acid	16,479.561 16,479.561	16,479.561 16,479.561	16,479.561 16,479.561	16,479.561 16,479.561	16,479.561 16,479.561	16,479.561 16,479.561
Arginine	3025.114 4487.544	3025.114 4487.544	3025.114 4487.544	3025.114 4487.544	3025.114 4487.544	3025.114 4487.544
Asparagine	21662.114 17452.118	18493.142 14923.142	69,112.23 10641.082	69,112.23 10641.082	13291.062 91,157.176	10572.827 14193.020
Aspartic Acid	62,249.538 62,249.538	48,112.448 48,112.448	21,729.588 18,048.778	21,729.588 18,048.778	14,952.534 13,849.014	18,011.814 18,011.814
Citrulline	8,435.514 8,435.514	6,290.546 6,290.546	6,126.548 4,126.548	6,126.548 4,126.548	5,729.524 5,729.524	4,492.946 4,492.946
Cysteine	9,489.174 9,489.174	8,449.524 8,449.524	6,149.524 4,149.524	6,149.524 4,149.524	5,159.514 5,159.514	4,121.154 4,121.154
Cysteic acid	4,976.524 4,976.524	4,976.524 4,976.524	4,976.524 4,976.524	4,976.524 4,976.524	4,976.524 4,976.524	4,976.524 4,976.524
Glutamic acid	13593.646 13593.646	13593.646 13593.646	13593.646 13593.646	13593.646 13593.646	13593.646 13593.646	13593.646 13593.646
Glutamine	28,454.724 28,454.724	16,951.534 16,951.534	27,129.524 27,129.524	20,251.524 20,251.524	22,512.524 22,512.524	20,251.524 20,251.524
Glycine	67295.268 67295.268	64259.648 64259.648	61259.538 61259.538	61259.538 61259.538	61259.538 61259.538	61259.538 61259.538
Histidine	15591.664 15591.664	12159.524 10159.524	10159.524 10159.524	10159.524 10159.524	13465.544 13465.544	14151.514 14151.514
Hydroxyproline	2,060.554 2,060.554	2,870.554 2,870.554	2,650.554 2,650.554	2,650.554 2,650.554	2,350.554 2,350.554	2,085.544 2,085.544
Isoleucine	18275.820 18275.820	14075.820 14075.820	13575.820 13575.820	13575.820 13575.820	18159.524 18159.524	22105.524 22105.524
Leucine	11385.544 11385.544	11151.544 11151.544	11061.544 11061.544	11061.544 11061.544	13151.544 13151.544	14351.544 14351.544
Lysine	22125.520 22125.520	19175.520 19175.520	22125.520 22125.520	22125.520 22125.520	26251.524 26251.524	26251.524 26251.524
Methionine	24411.119 24411.119	32001.662 32001.662	42701.016 42701.016	42701.016 42701.016	47295.524 47295.524	49329.524 49329.524
Phenylalanine	73,149.524 73,149.524	62,950.264 62,950.264	59,158.524 59,158.524	78,297.28 78,297.28	86,453.284 86,453.284	80,254.524 80,254.524
1-Histidine	21,145.524 21,145.524	14,495.524 14,495.524	18,172.524 18,172.524	18,172.524 18,172.524	14,495.524 14,495.524	17,449.524 17,449.524
2-Hydroxybutyrate	16,455.524 16,455.524	13,455.524 13,455.524	21,525.524 21,525.524	21,525.524 21,525.524	18,455.524 18,455.524	17,750.524 17,750.524
Ornithine	76,145.524 76,145.524	69,453.524 69,453.524	69,153.524 69,153.524	62,891.524 62,891.524	78,461.528 81,461.528	75,561.524 75,561.524
Proline	32652.528 32652.528	35952.528 35952.528	100542.520 100542.520	111625.520 111625.520	127621.020 13051.020	13671.020 13761.020
Taurine	28485.524 28485.524	55951.524 55951.524	42870.524 42870.524	48891.524 48891.524	51899.524 52991.014	52195.524 53010.004
Threonine	40251.524 40251.524	42259.524 42259.524	31457.524 31457.524	47215.524 47215.524	61599.524 61599.524	61599.524 61599.524
Tryptophan	18,455.524 18,455.524	3,125.524 3,125.524	6,000.524 6,000.524	6,000.524 6,000.524	6,000.524 6,000.524	10,000.524 10,000.524
Tyrosine	27727.524 27727.524	23124.524 23124.524	28621.524 28621.524	18495.524 18495.524	13597.524 13597.524	11034.524 11413.524
Valine	28129.524 28129.524	24925.524 24925.524	28624.524 28624.524	29094.524 29094.524	31259.524 31259.524	31259.524 31259.524

a Values (means  $\pm$  standard deviations) for FAA or MRS followed by the same letter are not significantly different at  $P \leq 0.05$ .

b Ratio fed to the feed intake of the corresponding amino acid group.

c Percentage of the NRC dietary requirement (1977) of lysine (based on protein digestibility coefficient), methionine, homocysteine, methionine sulfide or

d Not detectable at levels less than 0.01 mg/g diet—methionine, methionine sulfide, homocysteine, methionine sulfide.

e Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

f Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

g Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

h Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

i Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

j Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

k Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

l Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

m Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

n Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

o Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

p Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

q Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

r Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

s Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

t Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

u Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

v Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

w Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

x Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

y Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

z Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

aa Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

bb Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

cc Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

dd Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ee Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ff Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

gg Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

hh Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ii Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

jj Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

kk Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ll Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

mm Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

nn Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

oo Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

pp Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

qq Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

rr Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ss Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

tt Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

uu Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

vv Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ww Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

xx Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

yy Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

zz Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

aa Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

bb Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

cc Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

dd Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ee Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ff Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

gg Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

hh Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ii Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

jj Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

kk Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ll Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

mm Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

nn Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

oo Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

pp Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

qq Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

rr Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ss Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

tt Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

uu Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

vv Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ww Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

xx Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

yy Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

zz Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

aa Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

bb Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

cc Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

dd Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ee Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ff Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

gg Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

hh Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

ii Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

jj Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

kk Determined at levels less than 0.01 mg/g diet—ascorbate, α-ketoglutamate, α-ketoglutarate, α-ketobutyrate, α-ketothiolate.

App. 26a. Major and interactive effects produced by graded levels of dietary lysine on concentrations of free-sulfuric acid (TSA) and other sulphur-positive substances (TSS) in plasma of chicks with atelocentosis.

TSA or TSS	Effect of feeding Atelocentosis Restricted Ad libitum	Effect of administration			ANOVA [F (rep. f) <sup>a</sup> ]	
		102	122	140	Feed	Ad lib. Feed x Ad lib.
Alanine	407.0	523.0	571.0	-A	-A	4.10(0.018)
γ-nicotinoylbutyric acid	16.70	15.04	14.78	17.2A	13.2B	6.24(0.019)
Ammonia	43.53	21.30	34.18	23.2A	25.0A	8.12(0.017)
Arginine	43.54	36.08	37.38	43.1A	40.8A	15.0(0.001)
Asparagine	50.10	54.43	14.64	51.8A	79.1B	10.9A
Aspartic acid	103.8	114.0	127.8	128.8	147.6	21.2(0.001)
Citrulline	7.28A	5.00B	5.65B	-A	-A	7.65(0.023)
Cystathione	-A	-A	-A	-A	-A	3.28(0.017)
Cysteic acid/phosphocysteine	15.4A	10.0B	10.8B	-A	-A	1.08(0.18)
Glutamine	107.4	93.6B	111.6	-A	-A	0.25(0.013)
Glutamate	-A	-A	-A	-A	-A	2.74(0.012)
Glycine	155.8	140.0	161.6	-A	-A	1.39(0.13)
Glycine	-A	-A	-A	-A	-A	1.34(0.13)
Glycine	-A	-A	-A	-A	-A	2.37(0.11)
Histidine	-A	-A	-A	-A	-A	2.20(0.13)
Hydroxylysine	-A	-A	-A	-A	-A	0.77(0.17)
Hydroxyproline	13.6C	17.7B	22.8A	-A	-A	0.12(0.89)
Isotocine	11.8B	13.8A	13.2A	-A	-A	1.81(0.18)
Lysine	20.6C	23.1B	26.3A	21.7B	21.7A	3.28(0.013)
Tyrosine	30.31C	44.82B	53.58B	41.8B	58.9A	13.5(0.001)
Nicotinamide	63.7B	80.4A	80.7A	-A	-A	1.7(0.001)
1-methylhistidine	-A	-A	-A	-A	-A	1.35(0.17)
3-methylhistidine	-A	-A	-A	-A	-A	1.47(0.11)
Ornithine	70.7A	65.0B	76.1AB	-A	-A	3.28(0.011)
Phenylalanine	23.8A	101C	13.1B	-A	-A	6.6(0.010)
Proline	-A	-A	-A	-A	-A	1.27(0.22)
Serine	53.2B	62.0A	63.0A	-A	-A	6.55(0.007)
Taurine	23.8A	15.9B	14.3B	-A	-A	1.47(0.17)
Theanine	* 206C	231B	68.1A	-A	-A	20.1(0.001)
Tyrosine	23.8A	17.9B	18.2B	-A	-A	22.2(0.001)
Valine	23.5B	29.8A	31.0A	-A	-A	5.97(0.013)
Tryptophan	9.38A	0.02B	0.02B	-A	-A	2.20(0.023)

Means for TSA or TSS within a major effect followed by the same letter are not significantly different at P<0.05.

Majority atelocentosis = 25% per g of diet.

Corrected to the feed intake of the corresponding atelocentosis group.

Percentage of the NRC dietary requirement (1977) of lysine (hept) ration provided (1022).

Ad lib. = major effect of administration. Feed = minor effect of feeding. Feed x Ad lib. = test for interaction.

App. 25. Effect of graded levels of lysine and arginine on the concentrations of plasma free amino acids (FAA) and other aliphatic positive substances (RPS) in plasma of chicks with aflatoxicosis.

Williams ( $\pm$  standard deviation) for PAA or PAA<sub>1</sub> followed by the same letter are not significantly different at  $P < 0.05$ . Not detectable at levels less than 0.1 mmol/mol creatinine, ascorbic acid,  $\alpha$ -ketoglutaric acid,  $\beta$ -hydroxybutyric acid, creatine kinase, lactate dehydrogenase, glucose-6-phosphate dehydrogenase, hemopexin, methionine sulfone or sulfate.

App. 23a. **Holter and interactive effects produced by graded levels of lysine and arginine on the concentrations of plasma free amino acids (FAA) and other labile nitrogen positive substances (LNP) in plasma of chicks with efflavinosis.**

A HISTORY OF THE CHURCH IN AMERICA

THE JOURNAL OF CLIMATE





