# DEVELOPMENT OF A SUITABLE MODEL FOR ASCERTAINING THE GROWTH AND EGG PRODUCTION IN OUAILS 



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## DECLARATION

1 hereby declare that this thesis entitled "Development of a suitable model for ascertaining the growth and egg production in quails" is a bonafide record of research work done by me during the course of research work and the thesis has not previously formed the basis for the award to me of any degree, diploma, associateship, fellowship or other similar title of any other University or Society.

Vellanikkara,

## CERTIFICATE


#### Abstract

Certified that this thesis entitled "Development of a suitable model for ascertaining the growth and egg production in quails" is a record of research work done independently by Mr. John Thomas M. under my guidance and supervîsion and that it has not previously formed the basis for the award of any degree, fellowship or associateship to hîm.


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2)-7-91

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## CERTIFICATE

We, the undersigned members of the Advisory Committee of Mr. John Thomas, M., a candidate for the degree of Master of Science in Agricultural Statistics agree that the thesis entitled "Development of a suitable model for ascertaining the growth and egg production in quails" may be submitted by Mr. John Thomas, M., in partial fulfilment of the requirement for the degree.

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## ACKNOWLEDGEMENT

With great pleasure I express my heartfelt gratitude and obligation to Dr. K.C. George, Professor and Head, Department of Statistics, Kerala Agricultural University. As the Chairman of my Advisory Committee, he has given me the maximum help and encouragement by taking keen and lively interest throughout the course of this investigation. His deep academic interest, expert guidance helped me a lot that $I$ owe him beyond words.

I am happy to acknowledge with deep sense of gratitude, the valuable guidance received from Mr. K. I. Sunny, Associate Professor, Regional Agricultural Research Station, Pilicode.

Valuable suggestion received from Mr. S. Krishnan, Assistant Professor, Department of Agricultural Statistics contributed to the successful completion of the programme.

My profound gratitude is placed on record to Dr. G. Reghunathan Nair, Associate Professor, Centre of Excellence in Poultry Science for permitting me to make use of the existing facilities in the Poultry Farm for my thesis work and for his valuable guidance and suggestions.

I wish to place on record my sincere thanks to Mrs. Santha Bai, Junior Programmer, Department of Statistics for the help rendered by her during the analysis of the data.

I would like to express my sincere thanks to the Dean, College of Veterinary and Animal Sciences and the Associate Dean, College of Horticulture for providing necessary facilities for the study.

I will be failing my duty if $I$ do not express my extreme gratitude to the staff members of my department for their co-operation and sincere interest in my work.

I am privileged to acknowledge the encouragement rendered by my fellow colleagues especially Sukumaran, Anita, Sunil, Jayaraj, Chandrika, Seena, Biju and seniors.

I am grateful to my mother, brothers, sisters and relatives for their constant encouragement and co-operation which made me possible to complete the study.

I am also grateful to Mr. O.K. Ravindran, Peagles, Mannuthy for his unfailing patience, meticulous and neat typing.

I am grateful to the Kerala Agricultural University for the fellowship awarded to me during the course of research work.
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## CONTENTS



## INTRODUCTION

Coturnix Japanese quail belongs to the class Aves, family Phasianidae and genus coturnix and is named Coturnix coturnix Japonica. The Bobwhite quails belong to a different family. Coturnix.were either domesticated in Japan about the lith century or brought to Japan from China about that time. The male of this species is lighter than female and is identified by the cinnamon coloured feathers on the upper throat and lower breast region. The female is similar to male in colouration except that the feathers on the throat and upper breast are long, pointed and much lighter in cinnamon. Japanese quails are prolific layers and have showed that under tropical conditions quails mature by about 7 to 8 weeks of age and reach peak production by about 10 weeks. Under favourable environments (temperature, humidity, day light) they produce for long...periods and averaging 250 eggs per year. The weight of egg is approximately $10 \mathrm{~g}(6.7$ to 13.8 g$)$. The eggs are multicoloured ranging from dark brown-blue and white to buff each heavily mottled with black, brown and blue. Adult males weigh about 151 g (l lb to 190 g ) and females weigh about 166 g ( 130 to 200 g ) at lath week of maturity.

## Quails in general are easy to rear needing compara-

 tively smaller area for its rearing. Feeding is also very easy and less costly. It is generally disease free incomparison to poultry as a whole. Normally one quail egg cost about 50 paise in market. Quail egg and meat are considered to be of high medicinal value and hence it is more dear among the common people. By taking all these points into account it is very much economically viable for a lower middle class family to rear about 50 quails in their limited resources.

Very few studies were undertaken in the past with regards to body weight and egg production models in quails. The only available reference in literature at present are Laird (1965), Marks (1978), Ricklefs (1979), Kozaczynski (1985), Ricklefs (1985), Anthony ettal. (1986). Laird (1965) has fitted only a Gompertz curve for the growth pattern which he has not compared with any other model. Except Ricklefs (1979, 1985) none of them contributed much towards the development of suitable models for ascertaining the growth in quails. Since Ricklefs (1979) fitted logistic form for growth of Japanese quail and Ricklefs (1985) fitted Gompertz form, it was worth investigating the exact model for ascertaining the growth in quails. With regard to egg production models in quails no reference was available and hence it was also worth developing a suitable model for this purpose.

A study of growth curve may indicate the earliest age at which reliable prediction of adult weight may be made. Similarly the egg production also can be predicted through
suitable mathematical models. Under this situation it was thought useful to develop suitable models for ascertaining growth and egg production in quails with the 'following objectives.
I. To find a sui.table relationship between. age and body. weights.
2. To investigate the trend of egg production in quails through suitable mathematical models.
3. To study the impact of climatic elements (temperature, humidity) on egg prduction in quails.

With this study in view an experiment has been conducted on Japanese quails. Under this experiment 150 day old birds of same breed were hatched at the Kerala Agricultural University Poultry Farm. The birds were kept under homogenous conditions and were fed as per the package of practices recommended by KAU. Weekly body weights, daily egg production and daily climatological parameters were observed. This data has been used for development of suitable models.

Male


Quail eggs


Revien of Literature

## REVIEW OF LITERATURE

### 2.1 Growth studies

Growth curves reflect the life time inter relationship between an individual's inherent impulse to grow and mature in all body parts and environment in which these impulses are expressed. Knowledge of growth curves is important to all biologists regardless of specialisation, who are concerned with the effects of their research and recommendations on life time production efficiency. Development of the theory and techniques for fitting growth. curves may be traced both through time and scientific disciplines. In particular the theory and methodology of fitting growth curves owes much to the mathematicians, demographers and economists. A review of growth curve analyses in the chicks, ducks, quails and other avian species are presented here.
2.1.1 Growth studies (General)

Gompertz (1825) (See Winsor, 1932) developed a model $W_{t}=W_{\infty} \exp \left[-\left(I n W_{\infty}-\ln w_{t}{ }^{\prime}\right) \exp \left(-k\left(t-t^{\prime}\right)\right]\right.$ which is obtained by integrating the differential equation in terms of natural logarithm with respect to $t$

$$
d W_{t} / d t=k W_{t}\left(\ln W_{\infty}-W_{t}\right)
$$

where, $W_{\infty}=$ maximum weight, $W_{t}=$ weight at time $t$, K. = proportionality of growth rate constant

Ludwig (1929) pointed out that most of the equation representing growth can be reduced to four types. With $k$ standing for a constant, $t$ for time, $W$ for body weight, $A_{t}$ for time at completion of growth and $A_{w}$ for final weight, these four types can be expressed as follows:
$\phi=k\left(A_{t}-t\right)$ when rate of growth is considered proportional to the time remaining for completion of growth.
$\phi=k\left(A_{w}-w\right)$ when rate of growth is proportional to the weight to be gained.
$\phi=(k / t)$ when rate of growth is inversely proportional to the elapsed time.
$\phi=(k / w)$ when rate of growth is inversely proportional to the weight' already attained.

Bertalanffy (1938, 1949, 1957) outlined a general theory of organic growth. Bertalanffy's differential equation is given by

$$
d w / d t=a w^{m}-b w ; a, b, m \text { are constants. }
$$

Integrating, the function yielded the following growth curves

$$
W=\left(\frac{a}{b}-\left(\frac{a}{b}-w_{0}^{1-m}\right) \exp (-b(1-m) t)\right)^{1 / l-m}
$$

where, $W_{o}$ is weight at time. $t=0 \cdot$.
when $\quad m=0$;

$$
w=\frac{a}{b}-\left(\frac{a}{b}-w_{0}\right) \exp (-b t)
$$

which is monomolecular or modified exponential.
when

$$
\begin{aligned}
& m=2 \\
& W=1 /\left(\frac{a}{b}-\left(\frac{a}{b}-w_{0}\right) \exp (-b t)\right)
\end{aligned}
$$

which is autocatalytic or logistic curve.
when $m=1$,
the original differential equation gives
$W=w_{o} \exp [(a-b) t]$ which is exponential.

Under certain important assumption on constants and letting $m \rightarrow l$, this differential equation tends to Gompertz equation of the form

$$
W \rightarrow A \exp [B \exp (k t)]
$$

where, $A \rightarrow(a / b)^{1 / l-m}, \quad B=\log \left(A / w_{0}\right), \quad K \rightarrow b(m-1)$

Medawar (1940) pointed out that a growth curve would be of little interest to biologists if data representation were, its only function. The value of a growth curve lies in its potential for bringing out relationship which are not obvious from the data alone. This potential depends upon the correlation of the magnitude of growth curve parameters with significant experimental or biological conditions.

Brody (1945) defined two independent growth curves. One is based on the tendency for instantaneous rate of gain, prior to puberty or the point of inflection to be proportional to growth already made and is described by the differential equation

$$
\begin{equation*}
d W_{x} / d x=k W_{x} \tag{2.1.1}
\end{equation*}
$$

where, $k=$ proportionality or growth rate constant

$$
\mathrm{W}_{\mathrm{x}}=\text { weight of animal at time } \mathrm{x}
$$

Rearranging equation (2.1.1) and then integrating with respect to $x$ from $t$ ' to $t$ he obtained

$$
w_{t}=w_{\dot{0}} \exp (k t)
$$

where, $w_{o}$ is initial body weight

Following puberty the rate of gain tends to be proportional to the gain yet to be made and is described by the second equation

$$
\begin{equation*}
d w_{x} / d x i=-k \cdot\left(w_{\infty}-w_{x}\right) \tag{2.1.2}
\end{equation*}
$$

where, $k^{\prime}=$ growth rate constant and $W_{\infty}=$ maximum weight.

Rearranging equation. (2.1.2) and integrating with respect to $x$ from $t^{\prime}$ to $t_{0}$ he obtained

$$
w_{t}=w_{\infty}^{\prime}-\left(w_{\infty}-w_{o}\right) \exp \left(-k{ }^{\prime} t\right)
$$

when $\quad t^{\prime}=0$
where, $t^{\prime}$ is the initial time.

In 1838 , Verhulst (See Allee et al., 1949) developed. an equation to describe population growth and termed the function for this $S$-shaped curve the logistic function.

The equation for rate of gain, from which the logistic function was derived is,

$$
\begin{equation*}
d W_{x} / d x=k W_{x}\left(W_{\infty}-W_{x}\right) / W_{\infty} \tag{2.1.3}
\end{equation*}
$$

which indicates that the instantaneous rate of gain is a function of growth already made and potential for growth.

Rearranging equation (2.1.3), and then integrating, using partial fractions, between $t^{\prime}$ and $t$ with respect to $x$, we obtain

$$
\begin{equation*}
W_{t}=W_{\infty}\left(1+\left(\left(w_{\infty} / W_{t}\right)-1\right) \exp \left(-k\left(t-t^{\prime}\right)\right)^{-1}\right. \tag{2.1.4}
\end{equation*}
$$

Equation (2.1.4) relates weight, at a given time to a function of initial and final weights, growth rate constant and time.

Richards (1959) used an extended. form of lon Bertalanffy's growth function

$$
\begin{equation*}
\mathrm{W}=\left(\cdot \frac{\eta}{k}-\left(\frac{\eta}{k}-W_{0}^{l-m}\right) \exp (-(1-m) k t)\right)^{1 / 1-m} \tag{2.1.5}
\end{equation*}
$$

(which is sigmoid) to plant data for supplying an empirical fit. Here $W_{o}=$ weight at $t=0 ; ~ \sum$ (feta), k (kappa) are proportionality constants of anabolism and catabolism.
. $m=$ slope of Vo Bertalanffy's relation.

Equation (2.1.5)-can be abbreviated as

$$
\begin{equation*}
W^{l-m}=A^{l-m}-\beta \exp \left(-k^{\prime} t\right) \tag{2.1.6}
\end{equation*}
$$

where, $A^{1-m}=\eta / k ; \quad \beta=(\eta / k)-w_{0}^{l-m} ; k^{\prime}=(1-m) k$ are constants.

Therefore,

$$
\begin{aligned}
W^{1-m} & =A^{1-m}(1-b \exp (-k t)) \text { when } m<1-\ldots(2.1 .7) \\
& \left.=A^{1-m}(1+b \cdot \exp (-k t)) \text { when } m>1 \ldots-1.8\right)
\end{aligned}
$$

where, $b= \pm \beta^{m-1}$

```
when m = 0, equation (2.1.7) reduces to monomolecular
form W.= A(l-b exp(-kt)) and when m = 2 equation (2.1.8)
reduces to the autocatalytic form W = A (l+b exp(-kt) )
When m = l equation (2.1.6) is insoluble.
    When m lies between 0 and l the curves are transi-
tional in form between the monomolecular and Gompertz and when
m}\mathrm{ lies between l and 2 the curve lie between Gompertz and
autocatalytic. It was derived that as m->l equation (2.1.5)
represents the Gompertz equation
```

$$
W=A \exp (-b \exp (-k t))
$$

```
where, W = size at time t, A = ultimate limiting value,
    k = constant of catabolism (kappa)
```

    The absolute growth rate for (2.1.5) is given by
    
when $m=0,(2.1 .9)$ reduces to the growth rate of mono-
molecular function $k(A-W)$ and when $m=2$ equation (2.1.9)
reduces to growth rate of autocatalytic function kW (A-W)/A
Equation (2.1.9) becomes $\mathrm{kW} \log _{e}(A / W)$ which is growth
rate of Gompertz function. Here $k$ is the "rate constant"
which determines the spread of curve along time axis.

Nelder (1961) developed a logistic function
$W_{t}=W_{\infty}\left(1+\left(\left(W_{\infty}^{\prime} / W_{t}\right)^{1 / \theta_{-1}}\right) \exp \left(-k\left(t-t^{\prime}\right) / \theta\right)\right)^{-\theta}$
by integrating the differential equation

$$
\begin{equation*}
d W_{x} / d x=k W_{x}\left(I-\left(W_{t} / W_{\infty}\right)^{1 / \theta}\right. \tag{2.1.10}
\end{equation*}
$$

which is a generalization of logistic function given in differential equation form

$$
\begin{equation*}
d W_{x} / d x=k W_{x}\left(1-\left(W_{x} / W_{\infty}\right)\right) \tag{2.1.11}
\end{equation*}
$$

suggested by Verhulst (1838) (see Allee et al. 1949)

Here $\quad W_{\infty}=$ maximum weight, $W_{x}=$ weight of animal at time $x$

An advantage of Nelder's (1961) generalized logistic function is that when $u=1 / \theta \# 1$, the curve is asymmetrical and is more flexible.

Nelder (1962) (on reparameterization of Nelder (1961)) developed a logistic model of the type

$$
\left.W_{t}=w \dot{(i}+\left(\left(w_{\infty} / W_{t}\right)^{u}-1\right) \exp \left(-u k\left(t-t^{\prime}\right)\right)\right)^{-1 / u} \quad \text { which is obtained }
$$ by integrating the differential equation $d W_{x} / d x=k W_{x}\left(l-\left(W_{t} / W\right)^{u}\right)$ between $t$ and $\dot{t}$ with reference to $x$ and letting $u=1 / \theta$

Fabens (1965) gave properties and fitting of the Bertalanffy growth curve. He also developed a weight-growth curve,

$$
\mathrm{w}=(\mathrm{a}(1-\mathrm{b} \exp (-k t)))^{3}
$$

to recapture data as well as conventional age-size data. A computer programme is also presented to fit the curve,

$$
\text { where } a=q \eta / x, \quad b=1-\chi c / q \eta, k=\chi / 3
$$

$\eta=$ anobolic constant, $x=$ catabolic constant;
$q=$ constant characteristic of taxon,
$W=$ weight of animal
$c=a(1-b)$
The weight growth curve has an inflection point at $W=8 a^{3} / 27$.

Bhattacharya (1966) generalized the growth function suggested by Vo Bertalanffy as

$$
\left.y=(\alpha+\beta)^{t}\right)^{\delta}
$$

where, $\alpha, \beta, \gamma$ and $\delta$ are parameters.

The equation reduces to modified exponential when $\delta=1$; logistic equation when $\delta=-1$, Gompertz equation when $\delta \rightarrow \infty$

Laird et al. (1968) used a growth equation of the Gompertz type

$$
W=W_{0} \exp \left(\frac{A_{0}}{\alpha} \quad(1-\exp (-\alpha t))\right) ; A=A_{0} \exp (-\alpha t)
$$

where, $W=$ weight (or size in any other volumetric terms) at time $t, W_{o}=$ initial, weight at the start of the period of observation; $A_{0}$ and $A$ are specific growth rates at the starting time and at time $t$ respectively, $\alpha=$ rate of exponential decay of $A_{o}$ for representing the growth of individual parts of organism and of the whole organism.

Tallis (1968) suggested that growth and development can be regarded as a stochastic process in continuous time. Moreover in some situations of primary production, certain growth patterns may be more economical or otherwise more desirable than others.

Turner et al. (1976) introduced a general theory of growth which gave the following growth rate equation,

$$
x=\left(\beta / k^{n}\right) x^{I-n p}\left(k^{n}-x^{n}\right)^{l+p}
$$

where, $x=$ size of the system at time $t, k=$ size at $t=0$ and $\beta, n, p$ are parameters determining the particular characteristic of given curve.

Pruitt and Turner (1978), Turner and Pruitt (1978) have proved that general theory of growth is useful in numerical analysis of many and diverse biological and
biochemical processes. The range of applicability of the theory is illustrated by the fact that it yields

1. the logistic curve $(I+\exp (-\beta(t-\tau)))^{-1}$ with point of inflection $1 / 2$
2. the Gompertz $\exp (-\exp (-\beta-(t-\tau)))$ with point of inflection $1 / e$
3. Bertalanffy-Richards function $\left(1+\exp (-n \beta(t-\tau))^{-1 / n}\right.$
with point of inflection $(1+n)^{-1 / n}$. Here $\mathcal{Z}$ is constant of integration and is growth curve parameter.

Pruitt et al. (1979) developed the generic growth curve $\left.\left(1+(1+n p \beta(t-\tau))^{-l / p}\right)\right)^{-1 / n}$ whose point of inflection is given by $((1-n p) /(1+n))^{1 / n}$
where $n$ and $p$ are shape parameters, $\beta$ is maximum specific growth rate, $\tau$ is the constant of integration and reported that the analysis of growth in terms of the generic growth curve can be a powerful technique leading to relationships which may not be apparent from the growth data alone.
2.1.2 Growth studies in chicks

Lerner (1939) observed that the curve form $W=\mathrm{bt}^{\mathrm{a}}$ is not entirely satisfactory for representing the growth of a chick
but has been very widely used with many excellent results for limited portions of growth curve.

Here $W=$ weight at time $t, a$ and $b$ are constants.

The first derivative of this equation is
$d W / d t=u=a b t^{(a-1)}$

Sang (1962) used the logistic function,
$W_{t}=A /(1+\exp (c-k t))$
to analyse mean growth rate of selected lines of Brown Leghorns and found no sizeable differences among growth rates ( $k$ values) for six lines of chickens.

Roberts (1964) used a special case of the polynomial, a simple power function in time, to fit a linear portion of the early growth curve from hatching to seven weeks of age. He estimated the exponent of a function in the form

$$
W_{x}=a x^{k}
$$

Where $W_{x}=$ weight at time $x, a=$ initial weight, $k=$ early growth rate. The method was to take ratios of consecutive weights and solve for weekly k's in each individual using the equation'

$$
k=\log \left(w_{2} / w_{1}\right) / \log \left(x_{2} / x_{1}\right)
$$

Average values for $k$ in four different lines, over the seven weeks showed significant differences between lines and sexes when data were subjected to an analysis of variance.

Tanabe and Sacki,(1964)'working with pure and crossbred chicken from hatching to ten weeks of age, used the time function

$$
W_{x}=a x^{k}
$$

to estimate the values of $k$ (early. growth rate for each situ based on the log transformed function

$$
\log W_{x}=\log a+k \log x
$$

Here $W_{x}=$ weight at time $x, ~ a=i n i t i a l$ weight. In both the pure and crossbred lines, males and higher $k$ value than females of same breed.

$$
\begin{aligned}
& \text { Krause diet al. (1967) fitted the logistic function } \\
& W_{t}=\beta\left(1+\alpha \exp (-f t)^{-1}\right.
\end{aligned}
$$

to juvenile body weights of Athens-Canadian randombred chickens taken at 4 day intervals from 20 to 140 days of age.

Here $\quad \beta=w_{\infty}=$ maximum weight, $\alpha=\left(w_{\infty} / w_{t},\right)-1, w_{t},=$ weight at initial time $t^{\prime}, \gamma=k=$ proportionality or growth rate constant.

Pillai et al. (1969) while studying growth rate of chịckens from six different crosses found that simple exponential function $w=A \exp (k t)$ yielded a very good.fit. Here $A$, $k$ are constant, $w$ is the weight at time 't'.

Lilljedhl (1970) used a mathematical function of the Iogistic type

$$
Y=(A+B \exp (K X)) /(1+C \exp (K X))
$$

to give information about the growth of broiler chickens.

Here $\mathrm{y}=$ body weight, $\mathrm{X}=$ age.

All the four parameters $A, B, C$ and $K$ were significantly different from zero. For one of the forms in which the time difference between the early and late hatch of chicken tested, was so large that they represented two different stages of genetic improvement. Statistically significant difference between two hatches were found in all four parameters. By making second derivative of the body weight function equal to zero, some important growth characteristics such as co-ordinates of growth rates maximum, the corresponding inflection weight and proportion of body weight at slaughter (56 days) attained at the point of inflection (growth rate maximum) were derived. Also it was found that growth rate increased upto maximum of 29 g to 45 g per day - more in males than in females and it decreased subsequently. The maximum occurred between 36 and 48 days later in males than in females.

Zelenka (1970) while studying growth of chicken during the early period of post embryonal life used exponential function.

$$
w=a \exp (k t)
$$

where, $w=$ weight at time $t ; a, k$ are constants and the power function $y=a t^{b}$ to calculate growth from 2 to 22 days of age in 40 cockerals and 90 chicks of both sexes.

Tzeng and Becker (1981) used Gompertz model (Laird, 1965)

$$
W_{t}=w_{0} \exp \left(\frac{L}{k}(1-\exp (-k t))\right)
$$

in their study related to growth and found that it gave excellent fit to the live weight data as also the abdominal fat weight. Here $W_{t}=$ weight of the broiler or its part at time ' $t$ ', $d w / d t=$ L. $w_{t} \exp (-k t)=$ absolute growth rate. Other forms of non linear curves considered were logistic curve,

$$
W_{t}=A(1+\exp (-k t))^{-m} \text { and }
$$

Von Bertalanffy

$$
W_{t}=A(1-B \exp (-k t))^{3}
$$

where, $A, B, K$ and $m$ are parameters.

Grossman and Bohren (1982) in their study of "Comparison of proposed growth curve functions in chicken" contradicted the claim that growth in chicken is best described by any one of the growth functions but rather that under some set of environmental conditions one function should be able to describe accurately growth in chickens.

By describing overall body growth as a function of its components one should be able to have a better understanding of the growth characteristics and their relationships and be able to develop breeding plans to select for desirable aspects of the growth curve:

Jacob and Surendran (1984) observed that curves of the form

```
y = a exp(bx) (exponential)
y =ab c
```

were suitable in fitting body weights for 24 weeks. where ' $y$ ' is body weight at age $x$.

Grossman et al: (1985) used the logistic function model

$$
W_{t}=W_{\infty}\left(I+\left(\frac{W_{\infty}}{W_{\infty}}-1\right) \exp (-k t)\right)^{-1}
$$

for comparative purposes in two populations of chickens from hatching. through 45 week of age. Here $W_{t}=$ weight at time $t$, $k=$ growth rate constant, $w_{o}=$ initial weight, $w_{\infty}=$ maximum weight.

Grossman and Bohran (1985) used the logistic function

$$
W_{t}=(1+\exp (-a(t-\mu) / \sigma))^{-1}
$$

where $W_{t}=$ weight at time ' $t$ ', to determine whether two parameters of logistic growth function, growth rate constant

$$
\mathrm{K}=\frac{\mathrm{a}}{\sigma} \quad \text { where, } \mathrm{a}=\pi / \sqrt{2}
$$

and age at inflection point ( $F$ ) were inherited traits in chickens.

Knizetova et af. (1985) expressed the growth of chickens with widely different genetic growth. abilities by means of four parameters Richards function (Richards, 1959)

$$
\begin{aligned}
y_{t} & \left.=A(1 \pm b \exp (-k t))^{-1 / n}\right) n>-1 ; n \# 0 ; A, k>0 \\
Y_{t} & =A(1+b \exp (-k t))^{-1 / n} \text { for } n>0 \\
Y_{t} & =A(1-b \exp (-k t))^{-1 / n} \text { for } n<0
\end{aligned}
$$

The parameters estimated using the generalized least squares method are the following.
$Y_{t}=$ body weight (grams) at age $t$ (days)
$A=$ asymptotic value of size as $t \rightarrow \infty$; generally interpreted as average size at maturity independent of short term fluctuation of size in response to extraneous environmental effects.
$\mathrm{b}=$ integration constant, time scale parameter of no specific biological significance.
$k=$ rate at which logarithmic function of degree of maturity in body weight changes linearly per unit of time (maturity index), this rate estimates the maturation rate of curve (i.e. the relative rate at which A is reached).
$\mathrm{n}=$ shape parameter determining the position of the inflection point of the curve. In the original Richards function, it was designed as $m$ ( $m=n+1$ ) and by other authors as ( $M=-1 / n$ ), and established the degree of maturity in body weight at the point of inflection.

Tierce and Nordzkog (1985) analysed the body weight and shank length at 20 weeks of age and fitted an exponential equation of the type

$$
y=\alpha x^{\beta}
$$

where, $y=$ shank length, $x=$ body weight at 20 weeks of age, $\alpha, \beta$ are growth constants.

Indirabai et al. (1985) reported that growth curves of the form

$$
y=a+b x \text { (linear), } y=a \exp (b x) \text { (exponential) were }
$$

suitable for predicting the pattern of growth in broiler chicken.

Here $y$ is body weight at age $x$.

Ibe and Nwakalor (1987) fitted an allometric growth curve of the form $y=\alpha W^{\beta}$ in broiler chicken, where $w=$ body weight, $y=$ linear structural body parameters.

Grossman and Koops (1988) examined the growth of chickens with the help of multiphasic growth function based on a sum of logistic functions to describe mean body weight gain curves for four lines of chickens and to estimate number of phases and values of parameters within each phase. The function is of the form

$$
Y_{t}=\sum_{i=1}^{n}\left(a_{i}\left(l+\tanh \left(b_{i}\left(t-c_{i}\right)\right)\right)\right)
$$

with first derivative at age $t$ as

$$
y_{t}{ }^{\prime}=\sum_{i=1}^{n}\left(a_{i} b_{i}\left(1-\tan h^{2}\left(b_{i}\left(t-c_{i}\right)\right)\right)\right)
$$

where $y_{t}=$ mean weight (grams) at age $t, n=$ number of phases, tan $h=$ hyperbolic tangent, $a_{i}=$ half asymptotic weight, $b_{i}=$ growth rate relative to $a_{i}\left(\right.$ week $\left.^{-1}\right), c_{i}=$ age at maximum gain (weeks).

Study revealed that a diphasic function is appropriate to fit weight gain data for male and female chickens.
2.1.3 Growth studies in ducks

Susaki and Hamakawa (1965) while studying the growth of broiler ducks constructed growth curves from data on body weight of three broiler breeds and three crosses upto ten weeks of age. Curves of the type

$$
\begin{aligned}
& y=a x^{b} \\
& y=a+b x+c x^{2} \\
& y=a+b x+c(\log x)
\end{aligned}
$$

gave satisfactory fit to the data.

### 2.1.4 Growth studies in quails

Laird (1965) fitted the Gompertz equation to growth curves of several varieties of domestic chicken, turkey, goose, duck and quail.

Marks (1978) utilized four quail lines ( $P$, T, $S$ and $C$ which is maintained as a non selected control) to investigate growth patterns in quail. Body weight measurements suggested that the growth of all lines was best approximated by the logistic growth curve model

$$
y=A /\left(1+\exp \left(-k\left(x-T_{i}\right)\right)\right)
$$

where, $y=$ weight at agex,$k=$ constant proportional to overall growth rate, $\mathrm{T}_{\mathrm{i}}=$ age at infiection, $A=$ asymptotic weight

Ricklefs (1979) studied the patterns of growth development in Japanese quail, Common turn and the starling. It was observed that the growth rate varied inversely with functional maturity. The starling grows 4 times and the Turn two and a half time more rapidly than quail. Growth rate of each species was determined by fitting the growth curve by a logistic equation.

$$
W(t)=A /\left(1+\exp \left(-k\left(t-t_{i}\right)\right)\right)
$$

where $W(t)=$ weight at age $t, A=$ asymtote or weight plateau of growth curve, $k=$ growth rate constant, $t_{i}=$ age at inflection point of growth curve (the point of maximum growth rate).

Ino et al. (1985) observed that at 2, 4, 6, 8, 10 and $\%$ 12 week of age, the body weight of unspecified number of Japanese quails averaged 35.4, 75.3, 94.0, 96.7, 100.6 and 102.0 g respectively in male $\mathrm{V} / \mathrm{s} 36.1,77.9$, 109.0 , $118.6,122.3$ and 125.5 g in females. Age at sexual maturity averaged 35.9 and 44.8 days in male and female respectively.

Kozaczynski (1985) used the data obtained from 129 males and $15 i$ females Hungarians, 68 males and 101 females Pharroh $x$ Hungarian, 26 males and 33 females PAN, 92 males and 120 females BEK Japanese quails.

For the 4 breed types the body weight at 12 week of age averaged 152.4, 15I.1, 151.9 and 147.9 g for males and 187.8 ,
.181.5, 181.8 and 181.0 g for females respectively. Growth rate was highest from 7 to 21 days of age and lowest from 70 to 84 days. Correlation between body weight at various ages were mostly significant and ranged from 0.19 to 0.95 . Hungarian quails matured earlier than the other 3 types.

Ricklefs (1985) observed that growth curves of selected and unselected lines of broilers and Japanese quail show that the chicks respond to selection for 8 weeks or 4 week body mass respectively, by increasing the exponential growth rate during the first 2 weeks after hatching. The Gompertz equation used was of the form

$$
M(t)=A \exp (-\log (A / I) \exp (-k t))
$$

Here, $M(t)=$ Mass $(g)$ at age $t, A=$ mass plateau in grams (Asymptote), $K=$ rate of attainment of asymptote (unit $=1 /$ time), I = initial mass at age zero.

$$
\begin{aligned}
\log M(t):=\log A- & \log (A / I) \exp (-k t) \\
\frac{d}{d t}(\log M(t)) & =\log (A / I) K \exp (-k t) \\
& =k \log (A / M)=k(\log A-\log M)
\end{aligned}
$$

Anthony et al. (1986) have studied the growth curves of Japanese quails as modified by divergent solution of 4 week body weight of two weight selected lines and reported that the Gompertz curve is best for describing the growth of both the
lines. The logistic curve best fits the growth pattern of the low weight category. Also he observed that the pattern of growth of both the sexes were identical.

Sreenivasaih et al. (1987a) reported an initial average body weight of 100 Japanese quails (Monsoon hatch-August) as 5.74 g and the birds attained an average body weight of 117.27 g at 88 days ( 12.57 weeks) of age. The initial average body weight of 60 Japanese quails (WInter hatch-November) was 6.02 g and the birds attained an average body weight of 126.74 g at 88 days (12.57 weeks) of age.

## 2.2, Egg production studies

Mathematical models play a dominant role in poultry egg production. Models relating to egg production over time help in the prediction of egg production in certain time intervals or total egg production during any period. Also, from such models one can determine optimum time interval at which production is maximum.

Earlier workers (McNally, 1971; Timmermans, 1973) while trying to fit mathematical models for egg production employed the same models that were found to fit milk-yield at different lactation. One of the reasons for this is that milk yield and egg production both reach a peak and then onwards start declining.

A brief review of the works conducted are presented here.

Brody et al. (1923) employed the exponential curve $Y=a \exp (-b t)$
to describe average lactation course of a large number of cows of different breeds.

Here $y=$ average weekly or fortnightly milk yield, $t=$ time, $a, b$ are constants. They observed that the equation failed to fit the data remarkably well.

Sikka (1950) made an attempt to see whether the parabolic exponential equation

$$
y=a \exp \left(b t+c t^{2}\right)
$$

would represent the lactation curve better. Here $y=$ average weekly or fortnightly milk yield, and a, b, c are constants.

Narain (1962) studied three day egg laying of Drosophila melanogaster at various intervals throughout the life time. It was described that the decline in egg production of 0 . milanogaster from Nai-Basti (in India) by the model

$$
N(t)=66.56 \exp (-0.029 t)
$$

where $N(t)$ is daily egg production corresponding to day of egg laying.

Wood (1967) fitted a gamma type function

$$
y=a t^{b} \exp (-c t)
$$

to the lactation yield. The function accounted for 89.7 per cent of total variation in log daily yield. The function accounted for 89.7 per cent of total variation in log daily yield.

Tonkinson etal. (1969 b) suggested to use principal component analysis technique for the evaluation of egg production curves. The technique requires the computation of characteristic roots and vectors from the matrix of corrected sum of squares and cross products originating from the data matrix. The technique partitions the total variations of egg production curves with sets of independent derived responses. These responses are analysed by conventional ANOVA techniques for estimates of treatment, replicate and interaction effects. The curves can be reconstructed based upon the derived responses for visual interpretation of the analysis of variance results.

McMillan et al. (1970 a) used.the mathematical model
$N(T)=M\left(1-\exp \left(-\zeta\left(T-T_{0}\right)\right)\right) \exp (-\alpha T)$
to estimate the daily egg production of a Drosophila female whose production curve closely related to the curve of laying hens.

Here $N(T)=$ number of egg laid in period $T, M=$ the potential ${ }^{\text {Q }}$ maximum egg production per period, $T_{o}=$ the initial period of egg laying, $\xi=$ the rate of increase in egg laying, $\alpha=$ rate of decrease in egg laying.

McMillan et al. (1970 b) calculated parameters of egg production model
$N(t)=M\left(1-\exp \left(-\xi_{j}\left(t-t_{0}\right)\right)\right) \exp (-\alpha t)$
(Mc̣Milan et al. 1970 a)
of Drosophilla melanogaster From the model he also derived maximum egg production rate, time of this maximum, total egg production over 4-day intervals and total potential life time egg production. A feature of this model is that the parameters and derivations can be calculated withiout measuring daily egg productions throughout the entire life time of females.

$$
\begin{aligned}
& \text { McNally (1971) suggested that Wood model (1967) } \\
& \text { wheve } y_{t}=a t^{b} \exp (-c t) \\
& \text { and average daily yielded in the } i^{\text {th }} \text { week, } \\
& \text { a, b and } c \text { are constants }
\end{aligned}
$$

which has been found to fit lactation data in cattle can be taken as a basis of mathematical model for poultry egg production. He opined that the variation of number of eggs produced by a group of hens with time over a laying year has the same general form as that of milk yield over a lactation.

The addition of an extra term proportional to the square root of time to wood model improved the fit giving multiple correlation ranging from 0.936 to 0.994 . Hence Wood model may be written as

$$
Y_{t}=a t^{b} \exp \left(-c t+a t^{1 / 2}\right)
$$

The modified model was found to fit better than the Wood model with the highest values of $\mathrm{R}^{2}$ ranging from 93.6 to 99.4 per cent.

Gavora et al. (1971) verified that the egg production curve developed for egg production in D. melanogaster (McMillan, 1970) can be used for avian species also.

$$
N(t)=M\left(l-\exp \left(-\xi\left(t-t_{0}\right)\right)\right) \exp (-\alpha t)
$$

where $N(t)=$ number of egg laid on day $t ; M=$ potential maximum daily egg production, $t_{0}=$ initial day of egg laying, $\xi_{j}=$ rate of increase.in egg laying; ${ }_{n} \alpha=$ rate of decrease in egg laying.

The model was fitted for weekly, fortnightly and monthly time scale and also for all individuals as well as groups of hens.

Timmermans. (1973). used a mathematical model of egg production as suggested by McMillan (1970) and Gavora (1971) to a strain of white Leghorn bird (WB) and a strain composed of medium heavy breeds (MB). Both strains selected in a constant
environment. It was found that the function of McMillan fits in a high degree to the observed data. The prediction of the production has been investigated on hen housed basis. Later he proposed to use in further investigations the hen day data instead of hen housed data.

Schaeffer et ą. (1977) compared three methods 1. non linear method, 2. multiplicative method, 3. regression methiod to predict the 305 day milk and fat yields of Canadian Holstein and Jersey cattle.

The study found that the non linear method which consists of a non linear model

$$
Y_{i j}=A \exp \left(-\beta\left(i-t_{0}\right)\right)\left(1-\exp \left(-B\left(i-t_{0}\right)\right)\right) / B \exp \left(e_{i j}\right)
$$

was at least as accurate as either the multiplicative or regression methods because it requiresonly less computer storage for parameter estimates than other methods and could be implemented easily into a milk recording programme. Here $Y_{i j}=$ amount of milk given on the $i^{\text {th }}$ day of lactation of $j^{\text {th }}$ cow, $t_{o}=$ "lag time" parameter and may indicate when a cow's udder begins to lactate prior to calving, $B=$ slope of the lactation curve during the increasing production stage, $A=$ peak production, $\beta=$ slope during the decline in production after the peak, $\epsilon_{i j}=$ residual effect which was splitted into $\exp \left(\epsilon_{i j}\right)=$ $\exp (n \sin (i p)) \exp \left(e_{i j}\right)$ where $i \sin (i p)=$ period effect in a particular set of records.
$p$ is $2 \pi$ divided by the length of the period which could differ among lactation groups.

Ramachandra et al. (1979) fitted four models including the one suggested by McNally (1971) to identify the one which best fits the data and explain the minimum variation in egg production due to variation in the egg laying period in white cornish hens. The models are

1. $\quad \mu y \cdot t=\alpha+\beta t+t^{2}$
2. $\beta y \cdot t=\alpha+\beta \log t+\sqrt{ }(\log \cdot t)^{2}$
3. $\quad \boldsymbol{y} \cdot \mathrm{t}=\alpha \mathrm{t} \cdot \exp (-\mathrm{rt})$
4. $\quad$. y.t $=\alpha t \exp (-\gamma t+\delta \sqrt{t})$

In all the models $\boldsymbol{f} y . t$ represents the true (but unknown) average weekly egg production during laying period t. Under customary assumptions of the least square method $\mu \mathrm{y}$.t is measured as $y_{t}$ with a random error component which has zero expectation and constant variance.

The sample estimates of the unknown parameters in the case of each model were obtained by the application of the principle of least squares.

It was found that model 4 fitted to the sample data is an excellent one.

Adams and Bell (1980) used two equations

$$
y=\left(1.0 /\left(0.01+a r^{x-b}\right)\right)-c(x-d)
$$

where $x=$ age of flock, $y=$ per cent hen day production, $a, b$, $c, d$ and $r$ are constants, and

$$
y=a+b r^{x}
$$

where $x=$ age of flock, $y=$ weight per egg, $a, b, r$ are constants for predicting egg production and egg size as function of age.

Congleton et al. (1981) observed that prediction of laying hens using incomplete gamma curves
!

$$
y=A n^{b} \exp (-c n)
$$

is highly biased during most of the laying cycle. Here $y=$ predicted production for the $\mathrm{n}^{\text {th }}$ week of laying, $\mathrm{A}, \mathrm{b}, \mathrm{c}$ are parameters (specifying per cent production for each week in laying) which can be estimated by multiple linear expression.

An equation with an additional parameter (d)

$$
\hat{y}=A n^{b} \exp (-c n+d \sqrt{n})
$$

was also investigated and found that it did not improve the fit of the model.

McMillan (1981) derived an egg production model

$$
y_{t}=A\left(\exp \left(-k_{2} t\right)-\exp \left(-k_{1} t\right)\right)
$$

where $k_{1}$ and $k_{2}$ are instantaneous rates of increase and decrease in egg production, respectively and $A$ is the maximum potential egg production.

Gavora et al. (1982) fitted exponential models of McMillan et af. (1970 a, b)

$$
N_{p}=a(1-\exp (-c(t-d))) \exp (-b t)
$$

Wood (1967), $N_{p}=f t^{G} \exp (-h t)$ and a linear regression,

$$
N_{p}=m-k t
$$

to the results for individual hens, as well as to the mean results of groups of hens.

In all the three models, $N_{p}=$ the number of eggs laid in 28 day period $p, t=$ age of the birds in the middle of the 28 day period in days that is $t=$ (age at first egg $14+28$ p) and $a, b, c, d, f, g, h, k$ and $m$ are model parameters. The models were fitted to the data by the International Mathematics and 'Statistics Library Routine ZXSSQ. Overall the McMillan model gave the best results with mean $\mathrm{R}^{2}$ of 0.71 from fitting it to the results of 223 individuals with sixteen 28 day periods each and $R^{2}$ of 0.97 from fitting it to the period means for groups. The respective $R_{-}^{2}$ values were 0.68 and 0.95 for the wood model and 0.60 and 0.90 for the linear regression.

McMillan et al. (1986) compared three mathematical models of production curves with respect to their ability to predict 50 week egg production from actual egg production record lengths of 16,20 and 24 weeks.

1. Wood (1967) model $y=f t^{g} \exp (-h t)$
where $f, g$, $h$ are constants.
2. Compartmental model $\mathrm{y}=\mathrm{a}(1-\exp (-\mathrm{c}(\mathrm{t}-\mathrm{d}))) \exp (-\mathrm{bt})$
where $a, b, c, d$ are parameters which take different interpretations depending on the particular situations to which model is applied.
3. Regression model $y=m-k t$
$y=$ average total group egg production over time $t, m$ and $k$ are constants.

The study reported that when the model is intended only to predict full record egg production from past record, the linear model is better due to its simplicity and lower costs of fit. When the model is intended to serve several purposes, the compartmental model is preferable by virtue of the biological interpretation of its constants.

Johari et al. (1986) fitted McMillan model (1970 b) to annual egg production curves of white Leghiorn strain crosses to study their biological and statistical significance. The $R^{2}$ values were ranging from 80.31 to 97.87 per cent.

Johari et al. (1987) compared the fitting of five mathematical models.

Exponential (Brody et al. 1983) y $=\mathrm{a} \exp (-\mathrm{bt})$ Parabolic exponential (Sikka, 1950) $y_{t}=a \exp \left(b t+c t^{2}\right)$ Gamma type function (Wood, 1967) $y_{t}=a t^{b} \exp (-c t)$ Gamma type function (McNally, 1971) $y_{t}=a t^{b} \exp (-c t+d t)$ Inverse polynomial (Nelder, 1966). $y=t\left(a+b t+c t^{2}\right)^{-1}$
in the study on white leghorn strain cross entries.

The fitted models were compared using coefficients of determination ( $\mathrm{R}^{2}$ ). In all the models $\mathrm{a}, \mathrm{b}, \mathrm{c}, \mathrm{d}$ are constants. The study revealed that Nelder's inverse polynomial is the ideal one for hen housed egg production in layer type chickens. In this model ' 'a' is the rising extreme of the curve (depicting peak production) ' $b$ ' is the slope of curve (depicting the rate of increase of production) and 'c' is the decending phase (rate of decline of production).

Sreenivasaiah et al. (1987) reported an average egg weight of Japanese quails as $9.47 \pm 0.06$ of $g$ (Monsoon - August) $10.15 \pm 0.05 \mathrm{~g}$ (Winter - November).

Cason and Britton (1988) used weekly egg production data from six first cycle and l3 molted commercial layer flocks to compare: three non linear egg production models.

1. Compartmental or McMillan model

$$
P=\dot{a} \exp (-b t)(1-\exp (-c(t-d)))
$$

2. Adams-Bell model

$$
P=0.07\left(1 /\left(.01+a r^{t-b}\right)-c(t-d)\right)
$$

3. Compartmental type model based on a logistic growth curve $p=a \exp (-b t)(1 /(1+\exp (c+d t)))$
where $p=$ egg production in time period $t$, $t=$ age offlock in weeks, and $a, b, c, d, r$ are constants

Based on the comparisons the Adams Bell model was superior $\left(\mathrm{R}^{2}=.9938\right)$ to compartmental (McMillan model) $\left(\mathrm{R}^{2}=\right.$ .9523) in terms of goodness of fit or predictive ability for egg production in first cycle flocks and is as good as the compartmental model in molted flocks, for either hen housed or hen day egg production data.
, The logistic model ( $\mathrm{R}^{2}=.9930$ ) was superior to compartmental model ( $\mathrm{R}^{2}=.9423$ ) in terms of goodness of fit and predictive ability only for first cycle flocks. Here $\mathrm{R}^{2}$ is the mean coefficient of determination.

Predictions of total production based on 24 week of data were significantly more accurate for the Adams-Bell and logistic models than for compartmental model in terms of error or
percentage error in first cycle flocks, with no difference in molted flocks.

Yang et al. (1989) derived a new model (called as modified compartmental model)

$$
\begin{equation*}
y(t)=a \exp (-b t)(1+\exp (-c(t-d)))^{-1} \tag{2.2.1}
\end{equation*}
$$

for poultry egg production which not only has theoretical advantages over the compartmental model

$$
\begin{equation*}
y(t)=m(1-\exp (-p(t-q))(\exp (-n t) \tag{2.2.2}
\end{equation*}
$$

and wood model

$$
y(t)=f^{t} g \exp (-h t)
$$

but also appears better in respect of its goodness of fit to the data and its ability to predict.

In model (2.2.1) the parameters are 'a' = a scale parameter, b = rate of decrease in laying ability, $c=$ reciprocal indicator of the variation in sexual maturity, $d=$ mean age of sexual maturity of hens and $a e^{-b t}=$ exponential decay function.

In model (2.2.2) $\mathrm{m}, \mathrm{n}, \mathrm{p}$ and q are a scale parameters a measure of persistency of egg production, rate of sexual maturity and age at first egg respectively.

In model (2.2.3) $f, g$ and $h$ are parameters without reasonable interpretation.

Several criteria of goodness of fit, for different models including $R^{2}$ and errors of estimated annual egg numbers and estimated peaks of egg production curves are presented.

Results showed that modified compartmental model had higher $\mathrm{R}^{2}$, smaller errors of annual egg numbers and greater agreement of estimated peaks with actual peaks than the other two models.

Narayanankutty et al. (1989) in their study observed that for 40 eggs from 12 week old quails and 40 from 24 week old quails weight averaged $8.56 \pm 0.10$ and $9.95 \pm 0.13 \mathrm{~g}$ respectively.

### 2.3. Climatological studies

Yeates et al. (1941) and Rao et al. (1966), found detrimental efforts of hingh temperature on egg production in chicken.

Zannelli (1963) reported that higher weight gains was obsérved under low temperature regions.

Rao et al. (1966) reported that relative humidity had no effect on egg production in chicken.

Wilson et al. (1971) reported that 6 month old female Japanese quails kept in hot environment showed a higher egg production than those kept in cold environment.

Sreenivasaiah and Joshi (1987) observed that in Japanese quails mean temperature ( ${ }^{\circ} \mathrm{C}$ ) and vapour pressure (mm of Hg ) during the production period of two seasons (Monsoon - August, Winter - November) of hatch differed significantly. Egg production and egg weight of winter hatcched birds were significantly higher than the monsoon hatched birds.

Regression analysis indicated that regardless of season of hatch age had profound effect on both egg production and egg weight of monsoon hatched birds. Egg weight of winter hatched birds was greatly influenced by egg production. Definite detrimental effects due to high temperature and vapour pressure on egg production and egg weight were observed among monsoon hatched birds. Winter hatching was preferable and it was thought that values of climatic elements during production period of winter hatched birds were optimum for species. Both temperature and vapour pressure were found to influence egg production and egg weight. Also it was observed that relative humidity had . no effect on egg production in quails.

## MATEERIALS AND METHODS

### 3.1 Materials

The study was initiated using day old straight run birds of Japanese quails (Coturnix coturnix japonica.), one hundred and fifty in number from Kerala Agricultural University Poultry Farm, Mannuthy. The quails were hatched on lst February, 1989. They were serially numbered and wing banded.

On the 'day of hatching, the quails were placed in electrically operated thermostatically controlled battery type brooders. They were allocated to different compartments of the brooder at random. About thirty of chicks were allotted to each section of battery brooder. Commercial all-mash starter ration was fed ad libitum while the chicks were brooded in the batteries. Fresh water was made available at all time.

After few weeks the quails were divided into males and females and moved to individual cages. Necessary warmth was provided by infra-red bulbs. At this stage the birds were fairly well feathered and due to temperate weather only moderate heat was required. All the birds were fed on same feed formula and’all management practices were identical.

### 3.2 Methods

The body weight of each bird was recorded at weekly intervals until the birds attained an age of 12 weeks. At the
end of 12 th week weights were available on fifty eight (58) males and forty five (45) females. The weighing was stopped when females started laying. The weight of the birds were also recorded at. 27 th and 35 th experimental week.

The data so gathered were used for fitting appropriate functions of growth.

The choice of the appropriate curve to depict the growth pattern in any situation is not easy. on visual observation of data the pattern of growth appropriated to sigmoid curve upto l2th week. The following functions were considered.

| (i) Linear | $W_{t}$ | $=a+b t$ | (3.2.1) |
| ---: | :--- | ---: | :--- |
| (ii) Quadratic | $W_{t}$ | $=a+b_{1} t+b_{2} t^{2}$ | (3.2.2) |
| (iii) Exponential | $W_{t}=a \exp (b t)$ | (3.2.3) |  |
| (iv) Von Bertalanffy | $W_{t}=a[1-b \exp (k t)]^{3}$ | (3.2.4) |  |
| (v) Modified exponential | $W_{t}=k+a b t$ | (3.2.5) |  |
| (vi) Logistic | $W_{t}=a /[1+b \exp (-k t)]$ | (3.2.6) |  |
| (vii) Gompertz | $W_{t}=a \exp [-b \exp (-k t)](3.2 .7)$ |  |  |

Where $a, b, b_{1}, b_{2}$ and $k$ are constants and $W_{t}$ is the body weight at time $t$. The parameters of equations (3.2.1) to (3.2.4) were estimated using method of least squares and the parameters of equations (3.2.5) to (3.2.7) are estimated by method of partial sums (Nair, 1954).

The estimates of parameters and other related parameters are given as follows.

### 3.2.1 Linear

$$
w_{t}=a+b t
$$

The estimates of parameters are given as

$$
\begin{aligned}
& b=\left(N \Sigma t W_{t}-\Sigma t \Sigma W_{t}\right) /\left(N \Sigma t^{2}-(\Sigma t)^{2}\right. \\
& a=W-. b:
\end{aligned}
$$

Growth rate (GR) of an organism at any instant of time ( $t$ ) is defined as "the increase in organism material (body weight) of organism per unit of time.

$$
\text { i.e. } G R=d W_{t} / d t=b
$$

3.2.2 Quadratic

$$
w_{t}=a+b_{1} t+b_{2} t^{2}
$$

where, $a=D_{1} / D, b_{1}=D_{2} / D, b_{2}=D_{3} / D$

$$
\begin{aligned}
& D_{1}=\left|\begin{array}{ccc}
\Sigma W_{t} & \Sigma t & \Sigma t^{2} \\
\Sigma t W_{t} & \Sigma t^{2} & \Sigma t^{3} \\
\Sigma t^{2} W_{t} & \Sigma t^{3} & \Sigma t^{4}
\end{array}\right| \quad D_{2}=\left|\begin{array}{ccc}
n & \Sigma W_{t} & \Sigma t^{2} \\
\Sigma t & \Sigma t W_{t} & \Sigma t^{3} \\
\Sigma t^{2} & \Sigma t^{2} W_{t} & \Sigma t^{4}
\end{array}\right| \\
& D_{3}=\left|\begin{array}{ccc}
n & \Sigma t & \Sigma W_{t} \\
\Sigma t & \Sigma t^{2} & \Sigma t W_{t} \\
\Sigma t^{2} & \Sigma t^{3} & \Sigma t^{2} W_{t}
\end{array}\right| D=\left|\begin{array}{ccc}
n & \Sigma t & \Sigma t^{2} \\
\Sigma t & \Sigma t^{2} & \Sigma t^{3} \\
\Sigma t^{2} & \Sigma t^{3} & \Sigma t^{4}
\end{array}\right| \\
& \text { Growth rate }(G R)=b_{1}+2 b_{2} t
\end{aligned}
$$

### 3.2.3 Exponential

$$
W_{t}=a \exp (b t)
$$

where the constants $a$ and $b$ are estimated as $\hat{a}=\exp (A)$

$$
\begin{aligned}
& A=\bar{Z}_{t}-B \bar{t}, Z_{t}=\log W_{t} \\
& \hat{b}=B=\left(n \sum t z_{t}-\sum t \sum z_{t}\right) /\left[n \sum t^{2}-(\Sigma t)^{2}\right]
\end{aligned}
$$

When the curve is fitted the rate of growth at a particular period can be verified as the ratio of weight during the period to the weight during previous period minus one. In the case of exponential, the rate of growth is approximately given as $\exp (b)-1$.

### 3.2.4 Vo Bertalanffy

$$
W_{t}=a[1-b \exp (k t)]^{3}
$$

where 'a' is mature body weight which is known; $b$ and $k$ are constants.

$$
\begin{array}{ll}
\left(W_{t} / a\right)^{1 / 3} & =1-b \exp (k t) \\
b \exp (k t) & =1-\left(W_{t} / a\right)^{1 / 3} \\
\log b+k t & =\log \left[1-\left(W_{t} / a\right)^{1 / 3}\right] \\
B+k t & =z_{t}
\end{array}
$$

where $z_{t}=\log \left[1-\left(W_{t} / a\right)^{1 / 3]}\right.$

$$
\begin{aligned}
& \left.\mathrm{B}=\left(\Sigma z_{t} \sum t^{2}-\Sigma t \Sigma z_{t}\right) /\left(n \Sigma t^{2}-\bar{z} t\right)^{2}\right), \widehat{b}=\exp (B) \\
& \widehat{K}=\left(n \sum z_{t} t-\Sigma t \Sigma z_{t}\right) /\left(n \Sigma t^{2}-(\Sigma t)^{2}\right)
\end{aligned}
$$

The growth rate at a particular point is approximately given by
b $\exp (k t)[1-\exp (k)] /(l-b \exp (k t)]$
which depends on the value of $b \exp (K)$.
3.2.5 Modified exponential

$$
w_{t}=k+a b^{t}
$$

where the constants $a, b$, and $k$ are estimated as

$$
\begin{aligned}
& \hat{b}=\left[\left(s_{3}-s_{2}\right) /\left(s_{2}-s_{1}\right)\right]^{1 / n} \\
& \hat{a}=\left(s_{2}-s_{1}\right)(\hat{\mathrm{b}}-1) /\left(\hat{b}^{n}-1\right)^{2} \\
& \hat{k}=\left[s_{1}-\hat{a}\left(\hat{b}^{n}-1\right) /(\hat{b}-1)\right] / n
\end{aligned}
$$

Here $S_{1}, S_{2}, S_{3}$ are the sum of the $W_{t}$ values of three equal parts obtained from partial sums and 'n'is the number of observations in each part.

The growth rate at a particular period is approximately given by

$$
\begin{aligned}
& \left(a b^{t}+1-a b^{t}\right) /\left(k+a b^{t}\right) \\
& \approx a b^{t}(b-1) /\left(k+a b^{t}\right)
\end{aligned}
$$

3.2.6 Logistic

$$
w_{t}=a /(1+b \exp (-k t)
$$

which can be written as $Z_{t}=A+B C^{t}$

$$
\text { where, } \quad C=\exp (-K), A=1 / a, B=b / a, z_{t}=1 / w_{t}
$$

The estimates of parameters are given by,

$$
\begin{aligned}
& \widehat{K}=\log (1 / C) \text { where } c=\left(\left(S_{3}-S_{2}\right) /\left(S_{2}-s_{1}\right)\right)^{1-n} \\
& \hat{a}=1 / A \text { where } A=\left(S_{1}-\left(S_{2}-S_{1}\right) /\left(c^{n}-1\right) / n\right. \\
& \widehat{\mathrm{b}}=\mathrm{a} \cdot \mathrm{~B} \text { where } \mathrm{B}=\left(\mathrm{S}_{2}-\mathrm{S}_{1}\right)(\mathrm{c}-1) /\left(\mathrm{c}\left(\mathrm{c}^{\mathrm{n}}-1\right)^{2}\right) \\
& S_{1}=\sum_{t=0}^{3} z_{t^{\prime}}, \quad S_{2}=\sum_{t=4}^{7} z_{t}, \quad S_{3}=\sum_{t=8}^{l 1} z_{t}
\end{aligned}
$$

The growth rate at a particular period is given by

$$
\frac{1+b \exp (-k t)}{l+b \exp [-k t(t+l)]}-1
$$

3.2.7 Gompertz

$$
W_{t}=a \exp [-b \exp (-k t)]
$$

which can be written in the form

$$
z_{t}=A+B C^{t}
$$

where, $A=\log a, B=-b, c=\exp (-k)$, and $z_{t}=\log W_{t}$

The estimates of parameters are given by

$$
\begin{aligned}
& \hat{a}=\exp (A) \text { where } A=\left(S_{1}-\left(S_{2}-S_{1}\right) /\left(C^{n}-1\right)\right) / n \\
& \hat{b}=-B \text { where } B=\left(S_{2}-S_{1}\right)(C-1) /\left(C^{n}-1\right)^{2} \\
& \widehat{K}=\log _{e}(1 / C) \text { where } C=\left(\left(S_{3}-S_{2}\right) /\left(S_{2}-S_{1}\right)\right)^{1 / n} \\
& S_{I}=\sum_{t=0}^{3} Z_{t}, \quad S_{2}=\sum_{t=4}^{7} Z_{t}, \quad S_{3}=\sum_{t=8}^{11} Z_{t}
\end{aligned}
$$

The rate of growth at a particular period can be calculated as

$$
\frac{a \exp (-b \exp (-k(t+1)))}{a \exp (-b \exp (-k t))}-1
$$

### 3.3 Comparison of growth curves

In order to compare the relative efficiency of various growth curves and to select the one which best fit the observed data, two criteria are used, viz.
(i) Coefficient of determination ( $r^{2}$ ) and
(ii) Standard error of the estimate (s)

### 3.3.1 Coefficient of determination

It is calculated as the square of the correlation coefficient between the observed and predicted values. A large value of $r^{2}$ indicates best fit of the curve.

```
3.3.2 Standard error of the estimate
```

The standard error of the estimate measures the inadequacy of fit of the equation or of the error which is made in the estimation or prediction of $y$ from given values of x. The standard error of the estimate is calculated as

$$
s=\sqrt{\Sigma\left(y_{i}-\widehat{y}_{i}\right)^{2} /(n-2)}
$$

where $\hat{y}_{i}$ is the predicted value and ' $n$ ' is the number of observations. A small value of 'S' inđicates goodness of fit of the curve.

### 3.3.3 Comparison of rates of growth

Rao (1958) suggested a procedure for the comparison of rates of growth between different groups.

Let $Y_{t}$ denote the increase in body weight at time $t$ and $g_{t}$ is the mean of all $y_{t}$ 's in the experiment. Then $g_{t}$ is the time metameter. The difference in the values of $y_{t}$ are due to the time factor $\left(g_{t}\right)$, hence we may write

$$
Y_{t}=b g_{t}
$$

and the method of least squares leads to

$$
b=\left(\Sigma y_{t} g_{t}\right) /\left(\sum g_{t}^{2}\right)
$$

Thus obviously, comparison of difference in rates of growth between groups will be a comparison of b's. The 'b' values may be affected by initial body weight. Hence, a covariance analysis of the 'b' values taking initial values as concomitant variable can be adopted for comparing the growth rates of the groups.

### 3.4 Egg production

Daily egg production of quails were recorded from the end of March 1989 till the end of experiment. The data was converted to weekly, fortnightly, hen housed and hen day egg production basis.

McNally (1971) and Timmermans (1973) observed that the profile of egg production in poultry bears much resemblance to that one of the milk yield in dairy cattle. Hence the following milk production curves can be used to predict the egg production in quails.

$$
\begin{align*}
& \text { (i) Linear } \\
& Y_{t}=a+b t  \tag{3.5.1}\\
& \text { (ii) Exponential } \\
& \text { (Brody, 1923) } \\
& Y_{t}=a \exp (-b t)  \tag{3.5.2}\\
& \text { (iii) Parabolic }  \tag{3.5.3}\\
& \text { (iv) } \underset{\substack{\text { Inverse } \\
\text { (Nelynomial } \\
\text { Nelder } 1966)}}{ } \quad \mathrm{y}_{\mathrm{t}}=\mathrm{t}\left(\mathrm{a}+\mathrm{bt}+\mathrm{Ct}^{2}\right)^{-1} \\
& \text { (Sikka, 1950) } \\
& Y_{t}=a \exp \left(b t+C t^{2}\right)
\end{align*}
$$

```
    (v) Gamma function (Wood, 1967)
    (vi) Gamm type function }\quad\mp@subsup{Y}{t}{}=a\mp@subsup{t}{}{b}\operatorname{exp}(-ct+d\mp@subsup{t}{}{l/2}
    function
    (McNally, 1971)
```



```
    (Ramachandra
        et:al., 1979)
(viii) Quadratic
    function in }\quad\mp@subsup{Y}{t}{}=a+b(\mp@subsup{\operatorname{log}}{e}{}t)+c(\mp@subsup{\operatorname{log}}{e}{}t\mp@subsup{)}{}{2}(3.5.8
    (Ramachandra
                et a\ell.,1979)
    (ix) Quadratic-cum-
    log
        Yt}=a+bt+c\mp@subsup{t}{}{2}+d(\mp@subsup{\operatorname{log}}{e}{}t
        et a\ell.,1980)
    (x) Emperical }\quad\mp@subsup{Y}{t}{}=t/(a\operatorname{exp}(b\mp@subsup{t}{}{\prime})
(xi) Linear hyperbolic
\(\underset{\text { Sobrinho }}{\text { (Bianchini- }} \quad Y_{t}=a+b t+c / t\) et al., 1986)
```

Where $Y_{t}$ is the egg production in time scale (weekly, fortnightly, monthly hen housed, monthly hen day); $a, b, c, d$ are parameters.

For estimating the parameters of the above curves the following method suggested by Kendall et al. (1983) was used.

Let the linear model be

$$
\underline{Y}=\underset{\sim}{X} \underline{B}+\underline{U}
$$

where,

> | $\underline{Y}=$ | $n \times l$ vector of egg production |
| ---: | :--- |
| $\underline{X}=$ | $n \times k$ matrix of known coefficients |
| $\underline{B}=$ | $k \times 1$ vector of regression coefficients |
| $\underline{U}=$ | $n \times 1$ vector of error random variables with means |
|  | and dispersion matrix $E(\underline{U})=0$ |
| $V(\underline{U})=$ |  |

The vector of least square estimators of $B$ is given by $\underline{B}=\left(\underline{X}^{\prime} \underline{X}\right)^{-1} \quad \underline{X}^{\prime} \underline{y}$
and its dispersion matrix is given by

$$
v(\underline{B})=\sigma^{2}\left(\underline{x}^{\prime} \underset{\sim}{x}\right)^{-1}
$$

Unbiased estimator of $\sigma^{2}$ is $S^{2}$

$$
\text { Where, } \begin{aligned}
(n-k) s^{2} & =(\underline{Y}-\underline{X B})^{\prime}(\underline{Y}-X B) \\
& =\underline{Y}^{\prime} \underline{Y}-\underline{B}^{\prime} \underline{X}^{\prime} \underline{Y}
\end{aligned}
$$

### 3.5 Comparison of production curves

In order to compare the relative efficiency of various egg production curves and to select the one which best fit the observed data the following criteria are used.
3.5.1 Coefficient of determination
(As described in section 3.3.1)
3.5.2 Standard error of estimate
(As described in section 3.3.2)

### 3.5.3 Furnival index (I)

Furnival (1961) constructed an index (I) of fit as

$$
I=\left(\prod_{i=1}^{n} f^{\prime}\left(y_{i}\right)\right)^{-1 / n} \cdot s
$$

where, $f^{\prime}\left(y_{i}\right)$ is the derivative of some function of $f(y)$ of the dependent variable $Y$ with respect to y ; n is the number of data points, $s$ is the root mean square residual obtained from fitted regression.

A large value of $I$ indicates a poor fit and vice versa.

### 3.6 Climatological studies

Weekly egg production of thirty two (32) female Japanese quails hatched on first February 1989 and climatological parameters under which they were reared such as daily temperature and humidity (which was converted to weekly basis) were measured and utilised for this study. The birds started egg production from tenth week onwards and it was recorded daily up to the 34 th week. The egg production in any week depends on the previous two to three week climatological parameters. In order to correlate the effect of climatological parameters on egg production the previous three week average of those parameters were considered in correspondence with the present week's egg production, i.e. average of sixth,
seventh and eighth week climatological parameters were used to correlate with the nineth week egg production; average of seventh, eighth, nineth week climatological parameters were used to correlate the tenth week production etc.

In order to find a suitable relationship of temperature and humidity with that of weekly egg production, a multiple linear repression equation was fitted.

## RESULTS

The present investigation was undertaken mainly for the development of mathematical models for ascertaining growth and egg production in Japanese quails. Various mathematical models (described in sections $3.2,3.4$ ) were fitted to choose the best one for this purpose. The results obtained were presented in this chapter.

### 4.1 Average body weights

The average body weights along with standard error for male, female and birds irrespective of sex were presented in Table l. The body weights varied from 7.1172 g for first week to 157.6552 g during twelveth week in the case of males; 7.1622 g to 179.2000 g in the case of females; 7.1369 g to 167.0680 g in the case of birds irrespective of sex.

A graph with age (number of weeks) on $X$-axis and mean body weight of Japanese quails on Y -axis (Fig.I) was plotted.

Analysis of variance was conducted to test whether the difference in body weights of male and female Japanese quails were significant or not (Table 2.)

### 4.2 Growth study through mathematical models

The mathematical models utilised were described in section 3.2. The models were, fitted for fifty eight (58) male
and forty five (45) female Japanese quails using their body weights for first twelve weeks. The parameters of the fitted mathematical models namely linear, quadratic, exponential, Von Bertalanffy, modified exponential, logistic and Gompertz were presented in Table 3 to Table 9 for males and females along with the values of coefficient of determination $r^{2}$ and standard error of estiamte(s).

Models described in section 3.2 were also fitted for males, females and birds irrespective of sex using their average body weights over twelve weeks. The parameters of the fitted models were presented in Table 10 to 16 along with $r^{2}$ and $s$ values.

### 4.3 Rao's method

By the method of Rao (1958) the growth parameter 'b' was estimated for each of the 58 male and 45 female Japanese quails and were presented in Table 17. The growth parameter had a mean value of 0.009250 and 0.009741 for male and female Japanese quails respectively.

Analysis of covariance of 'b' values, taking initial body weight as concomitant variable was presented in Table 18.

### 4.4 Egg production study

Daily egg production data of thirty two (32) female Japanese quails reared for thirty four (34) weeks beginning
from first February 1989 to 30th September 1989 were utilised for the study. The birds started egg laying around loth week onwards. The data was coverted to total number of eggs produced on weekly, fortnightly and mean weekly egg production per bird basis (Table 19). Figure 6 and 7 shows the nature of weekly and fortnightly egg production. Hen housed and hen day egg production were worked out and presented in Table 20. A combined graph of hen housed and hen day egg production (y-axis) over various months (x-axis) were shown in Fig. 8.

Study was mainly conducted through mathematical models described in section 3.4. The models were fitted for four catagories namely total number of eggs produced on weekly, fortnightly and hen housed and hen day egg production data.

For comparison of fitted equations the coefficient of determination ( $r^{2}$ ), standard error of estimate(2) and Furnival index (I) were worked out.

In the case of $r^{2}$ value, it was assumed a linear trend of the variables. Hence the comparision under non linear models $r^{2}$ values will not be adequate. Therefore Furnival index (I) was taken as best criteria for comparison.

The parameters of the fitted mathematical models for the four categories (weekly, fortnightly, hen housed and hen day) were presented in Table 21 to 24 along with $r^{2}$, $s$ and $I$ values.

### 4.5 Climatological studies

- Under this study the data on temperature and humidity were collected upto 34 weeks. Data were converted to weekly basis in order to match with weekly egg production data. As mentioned in section 3.6 the climatological data of previous three weeks were correlated with present weeks average egg production per bird.

A multiple linear regression equation was fitted by taking previous three weeks average temperature $\left(z_{1}\right)$ and previous three weeks average humidity $\left(Z_{2}\right)$ as explanatory variables and present weekly average egg production per bird $(Y)$ as dependent variable (Table 25). The fitted equation was as given.

$$
Y=-41.7275+0.7687 * * z_{1}+0.3150 * * z_{2} \text { with } r^{2}=0.7654
$$

The ANOVA of fitted regression equation was given in Table 26.

Table 1. Mean and standard error of body weights (in g) of Japanese quails

| Week | Male | Female | Irrespective of ser |
| :---: | :---: | :---: | :---: |
| 1 | $7.1172 \pm 0.1591$ | $7.1622 \pm 0.1813$ | $7.1369 \pm 0.1698$ |
| 2 | $22.2362 \pm 2.1702$ | $22.5978 \pm 1.6934$ | $22.3942 \pm 1.9796$ |
| 3 | $46.6207 \pm 9.6479$ | $47.1556 \pm 9.3662$ | $46.8544 \pm 9.4831$ |
| 4 | $74.7241 \pm 13.4540$ | $76.8445 \pm 13.4400$ | $75.6505 \pm 13.4235$ |
| 5 | $95.5172 \pm 14.1089$ | $97.5111 \pm 12.9275$ | $96.3884 \pm 13.5764$ |
| 6 | $121.0690 \pm 14.9536$ | $123.3778 \pm 12.2980$ | $122.0777 \pm 13.8392$ |
| 7 | $131.4828 \pm 13.8136$ | $131.0222 \pm 10.6141$ | $131.2816 \pm 12.4613$ |
| 8 | $140.6552 \pm 12.1591$ | $142.4444 \pm 9.7318$ | $141.4369 \pm 11.1476$ |
| 9 | $147.2414 \pm 12.6426$ | $154.5333 \pm 14.2217$ | $150.4272 \pm 13.7540$ |
| 10 | $151.5862 \pm 14.8573$ | $166.8445 \pm 15.7450$ | $158.2524 \pm 20.1326$ |
| 11 | $152.1724 \pm 15.0708$ | $174.2667 \pm 15.7803$ | $161.8252 \pm 18.8578$ |
| 12 | $157.6552 \pm 14.5199$ | $179.2000 \pm 19.8055$ | $167.0680 \pm 20.0584$ |
| 27 | $172.4310 \pm 23.9862$ | $194.0000 \pm 17.7508$ | $181.8544 \pm 23.9376$ |
| 35 | $179.4483 \pm 22.1567$ | $192.4839 \pm 19.1841$ | $185.1456 \pm 21.8023$ |

Table 2. Mean sum of squares (M.S.S.) of body weights of Japanese quails over different weeks of age

| Source | DF | M.s.S. over various weeks |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |
| Between sexes | 1 | 0.0480 | 3.3010 | 7.2500 | 113.9370 | 100.6880 | 135.1250 | 5.3750 | 81.1250 |
| Within sexes | 101 | 0.0290 | 3.9070 | 90.7480 | 180.8470 | 185.1460 | 192.0820 | 156.7670 | 124.6960 |


| Source | DF | M.S.S. over various weeks |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 9 | 10 | 11 | 12 | 27 | 35 |
| Between sexes | 1 | 1347.5000** | 5899.5000** | 12369.7500** | 11762.2500** | 11788.616** | 4309.316** |
| Within sexes | 101 | 178.3140 | 232.5740 | 236.6630 | 289.8640 | 461.9624 | 437.3812 |

** Significant at 1 per cent level

Table 3. Fitting of linear form $W_{t}=a+b t$ using twelve weakb body weighte of Japanese quaile

| s1. | Male |  |  |  | Female |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | a |  | $z^{2}$ |  |  | b | ${ }^{2}$ | : |
| 1. | 12.1030 | 14.1406 | 0.8846 | 19.3184 | 8.9940 | 15.3266 | 0.9188 | 17.2295 |
| 2. | 11.1030 | 13.1406 | 0.9083 | 15.7899 | 8.0849 | 17.5434 | 0.9487 | 15.4337 |
| 3. | 28.0243 | 13.3476 | 0.8463 | 21.5140 | 13.8425 | 14.0678 | 0.9112 | 16.6119 |
| 4. | 7.9303 | 13.9248 | 0.9211 | 15:4079 | 5.7576 | 16.4168 | 0.9487 | 14.4324 |
| 5. | 7.0212 | 13.8339 | 0.9327 | 14.0563 | -1.5848 | 17.0297 | 0.9787 | 9.5069 |
| 6. | 4.3909 | 18.6437 | 0.9384 | 18.0709 | 7.3303 | 15.7556 | 0.9460 | 14.2361 |
| 7. | 23.5727 | 12.6927 | 0.8081 | 23.3900 | 1.4515 | 15.9164 | 0.9497 | 13.8561 |
| 8. | 16.2121 | 14.7238 | 0.8726 | 21.2791 | 9.1818 | 16.0105 | 0.8836 | 21.9778 |
| 9. | 1.3333 | 15.1154 | 0.9484 | 13.3230 | 9.2001 | 13.0692 | 0.9440 | 12.0345 |
| 10. | 21.8970 | 12.6287 | 0.8270 | 21.8438 | 25.8000 | 12.4000 | 0.7598 | 26.3621 |
| 11. | 24.5637 | 12.9364 | 0.8338 | 21.8431 | 13.6485 | 14.6797 | 0.9059 | 17.8960 |
| 12. | 17.0121 | 14.1622 | 0.8485 | 22.6296 | . 14.9378 | 14.9032 | 0.8639 | 22.3741 |
| 13. | 17.4227 | 13.3927 | 0.8523 | 21.0875 | 2.8061 | 17.0350 | 0.9287 | 17.8547 |
| 14. | -0.3757 | 14.9860 | 0.9475 | 13.3351 | 6.7394 | 14.5119 | 0.9099 | 17.2722 |
| 15. | 3.6485 | 13.7566 | 0.9327 | 13.9753 | 8.5151 | 16.3182 | 0.9150 | 18.8107 |
| 16. | 1.3455 | 14.2392 | 0.9282 | 14.9790 | 5.8182 | 17.2203 | 0.9669 | 12.0407 |
| 17. | 5.0727 | 14.8965 | 0.9003 | 18.7514 | 4.4848 | 14.7587 | 0.9321 | 15.0685 |
| 18. | 9.0121 | 16.3930 | 0.8977 | 20.9276 | -0.9333 | 19.1923 | 0.9717 | 12.3791 |
| 19. | 16.2727 | 14.8427 | 0.8789 | 20.8392 | -0. 5697 | 17.5209 | 0.9647 | 12.6716 |
| 20. | 10.5151 | 14.8566 | 0.8964 | 19.0997 | 19.4848 | 15.7741 | 0.8945 | 20.4844 |
| 21. | 19.7576 | 13.9476 | 0.8447 | 22.7197 | -8.9940 | 15.1350 | 0.9830 | 7.5259 |
| 22. | 15.3636 | 13.0594 | 0.8578 | 20.1091 | 7.0667 | 15.9462 | 0.9266 | 16.9687 |
| 23. | 18.4000 | 14.9615 | 0.8667 | 22.1884 | 7.4606 | 16.6035 | 0.9430 | 15.4341 |
| 24. | 11.8243 | 15.1014 | 0.8954 | 19.5168 | 1.8000 | 15.3154 | 0.9685 | 12.0464 |
| 25. | 19.9091 | 13.2217 | 0.8268 | 22.8860 | 10.5575 | 15.8399 | 0.9178 | 17.9220 |
| 26. | 13.3333 | 13.3615 | 0.8995 | 16.8888 | 12.3819 | 16.7336 | 0.8920 | 22.0164 |
| 27. | 18.6666 | 13.2077 | 0.8573 | 20.3765 | 5.0485 | 14.8874 | 0.9317 | 15.2428 |
| 28. | 5.6424 | 15.2678 | 0.9251 | 16.4291 | 1.2879 | 17.0955 | 0.9576 | 13.6083 |
| 29. | 11.4303 | 15.6595 | 0.8926 | 20.5456 | 10.0758 | 17.1025 | 0.8949 | 22.1602 |
| 30. | 13.3151 | 13.7490 | 0.9025 | 17.0910 | 11.6758 | 16.2717 | 0.8976 | 19.5079 |

Table 3. (Contd.)

| 51. <br> No. | Male |  |  |  | Female |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | a | b | $r^{2}$ | 8 | a | b | $r^{2}$ | 8 |
| 31. | 15.5272 | 14.2804 | 0.8703 | 20.8435 | 8.4848 | 15.3818 | 0.9270 | 16.3277 |
| 32. | 15.2000 | 14.2999 | 0.8654 | 21.3244 | 4.2970 | 17.0825 | 0.9539 | 14.2027 |
| 33. | -1.8303 | 12.9713 | 0.9749 | 7.8735 | 0.9334 | 17.5231 | 0.9674 | 12,1574 |
| 34. | 18.1091 | 14.2371 | 0.8593 | 21.7902 | 6.9757 | 15.6602 | 0.9601 | 12.0676 |
| 35. | -4.5878 | 18.7545 | 0.9748 | 11.4115 | 5.8848 | 15.6741 | 0.9521 | 13.2966 |
| 36. | 15.7152 | 16.1951 | 0.8640 | 24.2994 | -11.3151 | 15.7203 | 0.9663 | 11.1080 |
| 37. | 17.2879 | 12.8647 | 0.8600 | 19.6251 | -2.7001 | 17.4500 | 0.9598 | 13.5112 |
| 38. | 14.0697 | 13.3906 | 0.9057 | 16.3390 | 4.7545 | 17.9955 | 0.9453 | 16.3749 |
| 39. | 2.2364 | 12.8098 | 0.9563 | 10.3523 | -5.2819 | 18.9549 | 0.9627 | 14.1172 |
| 40. | 24.6224 | 11.9678 | 0.8164 | 21.4640 | 9.4151 | 16.1297 | 0.9381 | 15.6691 |
| 41. | 19.3090 | 15.5832 | 0.8543 | 24.3349 | 2.4454 | 13.8430 | 0.9050 | 16.9568 |
| 42: | 18.5940 | 11.6573 | 0.8205 | 20.6215 | 4.6303 | 16.1517 | 0.9526 | 13.6298 |
| 43. | 13.9879 | 14.0839 | 0.9041 | 17.3484 | 6.7212 | 17.4531 | 0.9269 | 18.5402 |
| 44. | 7.2999 | 16.6039 | 0.9100 | 19.7503 | -11.8604 | 18.1119 | 0.9907 | 6.6338 |
| 45. | 2.6151 | 12.8836 | 0.9658 | 9.1738 | -14.9212 | 19.3776 | 0.9907 | 7.1181 |
| 46. | -1.2485 | 16.9280 | 0.9445 | 15.5160 |  |  |  |  |
| 47. | 22.4182 | 12.8510 | 0.8181 | 22.9119 |  |  |  |  |
| 48. | 13.1455 | 14.2776 | 0.8704 | 20.8310 |  |  |  |  |
| 49. | 6.2667 | 16.7462 | 0.9204 | 18.6188 |  |  |  |  |
| 50. | 16.8728 | 15.3965 | 0.8853 | 20.9548 |  |  | - |  |
| 51. | 3.6303 | 13.7671 | 0.9494 | . 12.0218 |  |  |  |  |
| 52. | -1.1575 | 14.4524 | 0.9273 | 15.3079 |  |  |  |  |
| 53. | 8.4182 | 13.7049 | 0.8825 | 18.9088 |  |  |  |  |
| 54. | 19.2364 | 12.6559 | 0.8485 | 20.2226 |  |  |  |  |
| 55. | 0.1152 | 13.8797 | 0.9269 | 14.7359 |  |  |  |  |
| 56. | 12.9576 | 13.3706 | 0.8795 | 18.7137 |  |  |  |  |
| 57. | 1.6849 | 11.6434 | 0.9535 | 9.7236 |  |  | . |  |
| 58 | 7.8970 | 14.6364 | 0.9254 | 15.7180 |  |  |  |  |

Table 4. Fitting of quadratic form $H_{t}=a+b_{1} t+b_{2} t^{2}$ uging twelve weeks body weights of Japanese quails

| S1. | Male |  |  |  |  | Female |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Ho. | a | $\mathrm{b}_{1}$ | $\mathrm{b}_{2}$ | $5^{2}$ | 8 | a |  | $\mathrm{b}_{2}$ | ${ }^{2}$ | 5 |
| 1. | -35.2687 | 34.4426 | -1.5617 | 0.9853 | 7.2787 | -28.5590 | 31.4207 | -1.2380 | 0.9748 | 10.1265 |
| 2. | -28.7687 | 30.2284 | -1.3144 | 0.9931 | 4.5600 | -28.0590 | 33.0335 | -1.1916 | 0.9895 | 7.3560 |
| 3. | -31.8770 | 36.4481 | -1.7770 | 0.9862 | 6.7838 | -23.7406 | 30.1748 | -1.2390 | 0.9771 | 8.8861 |
| 4. | -29.4067 | 29.9264 | -1.2309 | 0.9883 | 6.2527 | -24.2731 | 29.2871 | -0.9900 | 0.9809 | 9.2780 |
| 5. | -26.7250 | 28.2966 | -1.1125 | 0.9870 | 5.9986 | -20.6795 | 25.2131 | -0.6295 | 0.9912 | 6.4544 |
| 6. | -39.6434 | 37.5155 | -1.4517 | 0.9915 | 7.0936 | -23.2342 | 28.8547 | -1.0076 | 0.9821 | 8.6381 |
| 7. | -34.3250 | 37.5060 | -1.9087 | 0.9787 | 8.2224 | -30.0978 | 29.4376 | -1.0401 | 0.9875 | 7.2731 |
| 8. | -37.8409 | 37.8894 | -1.7820 | 0.9918 | 5.6754 | -40.6589 | 37.3709 | -1.6431 | 0.9704 | 11.6760 |
| 9. | -27.7500 | 27.5797 | -0.9588 | 0.9841 | 7.8036 | -21.1406 | 26,0723 | -1.0002 | 0.9956 | 3.6429 |
| 10. | -33.8223 | 36.5083 | -1.8369 | 0.9903 | 5.4576 | -39.7455 | 40.4909 | -2.1608 | 0.9752 | 8.9300 |
| 11. | -30.4588 | 36.5174 | -1.8139 | 0.9868 | 6.4952 | -29.2001 | 33.0434 | -1.4126 | 0.9841 | 7.7420 |
| 12. | -39.8366 | 38.5259 | -1.8741 | $0.9872^{\circ}$ | 6.9370 | -39.6795 | 38.3106 | -1.8006 | 0.9815 | 8.6856 |
| 13. | -34.7705 | 35.7612 | -1.7207 | 0.9835 | 7.4187 | -37.9816 | 34.5154 | -1.3446 | 0.9827 | 9.2776 |
| 14. | -28.2998 | 26.9535 | -0.9206 | 0.9809 | 8.4799 | -34.5638 | 32.2132 | -1.3616 | 0.9846 | 7.5178 |
| 15. | -29.8364 | 28.1072 | -1.1039 | 0.9887 | 6.0242 | -38.2045 | 36.3409 | -1.5402 | 0.9911 | 6.4315 |
| 16. | -32.5181 | 28.7521 | -1.1164 | 0.9814 | 8,0297 | -21.5229 | 28.9378 | -0.9013 | 0.9916 | 6.3850 |
| 17. | -37.8818 | 33.3055 | -1.4161 | 0.9762 | 9.6593 | -31.0683 | 29.9957 | -1.1721 | 0.9869 | 6.9684 |
| 18. | -42.7000 | 38.5554 | -1.7048 | 0.9883 | 7.4585 | -23.1229 | 28.7021 | -0.7315 | 0.9849 | 9.5344 |
| 19. | -34.0682 | 36.4173 | -1.6596 | 0.9814 | 8.6069 | -30.2593 | 30.2451 | -0.9788 | 0.9928 | 6.0279 |
| 20. | -29.9774 | 32.2105 | -1.3349 | 0.9639 | 11.8769 | -25.0683 | 34.8683 | -1.4688 | 0.9669 | 12.0957 |
| 21. | -36.2501 | 37.9508 | -1.8464 | 0.9828 | 7.9328 | -14.4413 | 17.4695 | -0.1796 | 0.9843 | 7.6249 |
| 22. | -34.6591 | 34.4977 | -1.6491 | 0.9854 | 6.7828 | -31.0776 | 32.2937 | -1.2575 | 0.9804 | 9.2427 |
| 23. | -37.2592 | 38.8154 | -1.8349 | 0.9884 | 6.9080 | -25.4413 | 30.7043 | -1.0847 | 0.9806 | 9.4972 |
| 24. | -36.8046 | 35.9423 | -1.6031 | 0.9896 | 6.4875 | -26.2684 | 21.3447 | -0.9253 | 0.9912 | 5.8533 |
| 25. | -37.8407 | 37.9716 | -1.9038 | 0.9868 | 6.6673 | -34.0411 | 34.9536 | -1.4703 | 0.9916 | 6.0256 |
| 26. | -27.9319 | 31.0466 | -1.3604 | 0.9865 | 6.5171 | -42.7770 | 40.3731 | -1.8184 | 0.9903 | 6.9434 |
| 27. | -31.2045 | 35.0096 | -1.6771 | 0.9863 | 6.6515 | -29.5042 | 29.6957 | -1.1391 | 0.9826 | 8.1077 |
| 28. | -34.3047 | 32.5880 | -1.3169 | 0.9893 | 6.5351 | -29.1248 | 30.1294 | -1.0026 | 0.9883 | 7.5287 |
| 29. | -37.0321 | 36.4291 | -1.5977 | 0.9793 | 9.5128 | -42.2611 | 39.1040 | -1.6924 | 0.9767 | 10.9945 |
| 30. | -29.5414 | 32.1160 | -1.4128 | 0.9914 | 5.3415 | -34.6159 | 35.1110 | -1.5261 | 0.9812 | 8.8013 |

contd.

Table 4. (Contd.)

| 92. | Hale |  |  |  |  |  | Female |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| No. | a | $\mathrm{b}_{1}$ | $\mathrm{b}_{2}$ |  | $r^{2}$ | * | a | $b_{1}$ | $b_{2}$ | $r^{2}$ | B |
| 31. | -37.1774 | 36.8681 | -1.7375 |  | 0.9906 | 5.9180 | -27.8184 | 30.9404 | -1.1968 | 0.9793 | 9.1543 |
| 32. | -37.7769 | 37.0044 | -1.7465 |  | 0.9859 | 7.2745 | -21.6727 | 28.2124 | -0.8561 | 0.9763 | 10.7439 |
| 33. | -17.5046 | 19.6888 | -0.5167 |  | 0.9893 | 5.4108 | -21.2637 | 27.0360 | -0.7318 | 0.9832 | 9.2091 |
| 34. | -36.7317 | 37.7402 | -1.8079 ${ }^{\circ}$ | ; | 0.9886 | 6.5452 | -19.2819 | 26.9134 | -0.8656 | 0.9875 | 7.1192 |
| 35. | -30.2772 | 29.7642 | -0.8469 |  | 0.9933 | 6.1906 | -24.8272 | 28.8364 | -1.0125 | 0.9892 | 6.6648 |
| 36. | -45.2769 | 42.3346 | -2.0107 |  | 0.9883 | 7.5168 | -23.1634 | 20.7981 | -0.3906 | 0.9718 | 10.6992 |
| 37. | -32.1704 | 34.0611 | -1.6305 |  | 0.9890 | 5.8039 | -30.5297 | 29.3770 | -0.9175 | 0.9845 | 8.8322 |
| 38. | $-25.8430$ | 30.4960 | -1.3158 |  | 0.9873 | 6.3145 | -34.4389 | 34.7926 | -1.2921 | 0.9908 | 7.0955 |
| 39. | -21.7182 | 23.0760 | -0.7897 |  | 0.9902 | 5.1569 | -33.6573 | 31.1157 | -0.9354 | 0.9845 | 9.5738 |
| 40. | -29.282 | 35.0783 | -1.7778 |  | 0.9845 | 6,5747 | -22.3846 | 29.7581 | -1.0483 | 0.9751 | 10.4790 |
| 41. | -40.3272 | 41.1416 | -1.9660 |  | 0.9812 | 9.2075 | -31.3389 | 28.3220 | $-1.1138$ | 0.9597 | 11.6412 |
| 42. | -29.2090 | 32.1443 | -1.5759 |  | 0.9604 | 10.2078 | -18.1046 | 25.8952 | -0.7495 | 0.9717 | 11.0948 |
| 43. | -28.3451 | 32.2266 | -1.3956 |  | 0.9869 | 6.7508 | -26.1728 | 31.5506 | -1.0844 | 0.9603 | 14.4064 |
| 44. | -40.4842 | 37.0828 | -1.5753 |  | 0.9864 | 8.0873 | -17.6181 | 20.5794 | -0.1898 | 0.9917 | 6.5991 |
| 45. | -18.3936 | 21.9730 | -0.6992 |  | 0.9923 | 4.5822 | -22.1180 | 22.4619 | -0.2373 | 0.9920 | 6.9242 |
| 46. | -35.7859 | 31.7298 | -1.1386 |  | 0.9844 | 8.6746 |  |  |  |  |  |
| 47. | -35.3769 | 37.6204 | -1.9053 |  | 0.9860 | 6.7026 |  |  |  |  |  |
| 48. | -38.3770 | 36.3587 | -1.6985 |  | 0.9854 | 7.3690 |  |  |  |  | , |
| 49. | -38.6042 | 35.9766 | -1.4793 |  | 0.9875 | 7.7893 |  |  |  |  |  |
| 50. | -33.9225 | 37.1659 | -1.6746 |  | 0.9831 | 8.4877 |  |  |  |  |  |
| 51. | -24.2408 | 25.7119 | -0.9188 |  | 0.9889 | 5.9484 |  |  |  |  |  |
| 52. | -32.6046 | 27.9297 | -1.0367 |  | 0.9718 | 10.0489 |  | - |  |  |  |
| 53. | -36.3547 | 32.8932 | -1.4760 |  | 0.9781 | 8.6126 |  |  |  |  |  |
| 54. | -31.7182 | 34.4936 | -1. 6798 |  | 0.9880 | 5.9943 |  |  |  | - |  |
| 55. | -31.5819 | 27.4641 | -1.0450 |  | 0.9760 | 8.9075 |  |  |  |  |  |
| 56. | -31.7090 | 32.5134 | -1.4725 |  | 0.9791 | 8.2193 |  |  |  |  |  |
| 57. | -17.3910 | 19.8187 | -0.6289 |  | 0.9795 | 6.8117 |  |  |  |  |  |
| 58. | -30.2998 | 31.0064 | -1.2592 |  | 0.9893 | 6.2734 |  |  |  |  |  |

Table 5. Fitting of exponential form $H_{t}=a \exp (b t)$ using twelve weeks body weighte of Japanese quails

| Sl. <br> No. | Male |  |  |  | Female |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | a | $\exp (\mathrm{b})$ | $r^{2}$ | * | a | $\exp (\mathrm{b})$ | $\mathrm{r}^{2}$ | 8 |
| 1. | 18.6083 | 1.2504 | 0.6260 | 52.3858 | 18.5951 | 1.2574 | 0.6916 | 52.3038 |
| 2. | 18.0813 | 1.2440 | 0.6639 | 45.5260 | 19.4494 | . 1.2684 | 0.7357 | 58.9079 |
| 3. | 21.0599 | 1.2375 | 0.5919 | 53.9838 | 19.3081 | 1.2468 | 0.6969 | 48.5182 |
| 4. | 17.3570 | 1.2532 | 0.6840 | 47.0061 | 18.1009 | 1.2669 | 0.7515 | 52.4613 |
| 5. | 17.0077 | 1.2541 | 0.7067 | 45.2047 | 16.9860 | 1.2728 | 0.8019 | 48.0468 |
| 6. | 18.3764 | 1.2812 | 0.6892 | 66.8291 | 18.3391 | 1.2613 | 0.7466 | 50.4252 |
| 7. | 20.7171 | 1.2356 | 0.5462 | 54.4006 | 16.2102 | 1.2733 | 0.7257 | 51.4704 |
| 8. | 20.3614 | 1.2475 | 0.6059 | 57.0104 | 18.8951 | 1.2608 | 0.6193 | 59.6094 |
| 9. | 16.5810 | 1.2624 | 0.7257 | 46.4843 | 17.9044 | 1.2423 | 0.7342 | 40:5268 |
| 10. | 21.0496 | 1.2309 | 0.5658 | 51.4690 | 22.0419 | 1.2263 | 0.4856 | 55.4780 |
| $\Perp$. | 22.1088 | 1.2299 | 0.5818 | 52.7820 | 19.6843 | 1.2492 | 0.6744 | 52.0692 |
| 12. | 19.9029 | 1.2467 | 0.5761 | 56.8957 | 19.1449 | 1.2563 | 0.5975 | 59.2909 |
| 13. | 19.4909 | 1.2433 | 0.5913 | 52.9997 | 17.6269 | 1.2724 | 0.6795 | 58.5227 |
| 14. | 15.4677 | 1.2686 | 0.7280 | 46.0798 | 16.7217 | 1.2624 | 0.6574 | 51.4847 |
| 15. | 15.3594 | 1.2632 | 0.6917 | 45.6180 | 18.7422 | 1.2642 | 0.6584 | 58.9429 |
| 16. | 14.7895 | 1.2701 | 0.6814 | 47.6338 | 19.4832 | 1.2632 | 0.7784 | 52,4081 |
| 17. | 15.9056 | 1.2709 | 0.6418 | 54.2924 | 16.3689 | 1.2650 | 0.6958 | 49.4599 |
| 18. | 17.9022 | 1.2718 | 0.6262 | 63.2113 | 18.2429 | 1.2807 | 0.7987 | 56.9942 |
| 19. | 20.5943 | 1.2470 | 0.6301 | 55.8939 | 17.1711 | 1.2766 | 0.7471 | 54.8400 |
| 20. | 18.8433 | 1.2530 | 0.6723 | 51.6142 | 22.1127 | 1.2478 | 0.6782 | 57.8483 |
| 21. | 20.3256 | 1.2451 | 0.5836 | 56.8721 | 12.8356 | 1.2846 | 0.8459 | 36.8402 |
| 22. | 18.8204 | 1.2427 | 0.5959 | 50.5028 | 17.7309 | 1.2674 | 0.7002 | 55.0064 |
| 23. | 20.7950 | 1.2489 | 0.6068 | 59.1939 | 18.5973 | 1. 2666 | 0.7407 | 54.6617 |
| 24. | 19.0667 | 1.2551 | 0.6368 | 55.8689 | 16.4780 | 1.2660 | 0.7468 | 46.9878 |
| 25. | 20.5753 | 1.2369 | 0.5567 | 54.0915 | 19.0171 | 1.2605 | 0.6711 | 56.9433 |
| 6. | 19.0710 | 1.2413 | 0.6652 | 46.6798 | 19.5356 | 1.2657 | 0.6200 | 65.3159 |
| 7. | 20.6730 | 1.2361 | 0.6079 | 51.2181 | 17.2060 | 1.2598 | 0.6971 | 49.0024 |
| 8. | 17.0109. | 1.2657 | 0.6755 | 53.1318 | 17.3909 | 1.2733 | 0.7403 | 54.2300 |
| 29. | 18.4408 | 1.2642 | 0.6398 | 59.3534 | 19.1383 | 1.2690 | 0.6357 | 64.9277 |
| 0. | $18.9738^{\circ}$ | 1.2457 | 0.6602 | 49.0949 | 19.1871 | 1.2555 | 0.6454 | 55.8726 |

Table 5. (Contd.)

| S1. Male |  |  |  |  | Female |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| No. | a | $\exp (\mathrm{b})$ | $\mathrm{r}^{2}$ | $s$ | a | $\exp (b)$ | $r^{2}$ | 8 |
| 31. | 19.4966 | 1.2491 | 0.6045 | 55.5889 | 17.8601 | 1.2630 | 0.7044 | 52.7448 |
| 32. | 18.6062 | 1.2555 | 0.5978 | 57.2052 | 18.6890 | 1.2661 | 0.7676 | 52.2408 |
| 33. | 13.4374 | 1.2658 | 0.7951 | 35.1019 | 17.8025 | 1.2730 | 0.7902 | 51.7521 |
| 34. | 19.4266 | 1.2522 | 0.5918 | 58.1758 | 18.9281 | 1.2556 | 0.7762 | 46.8500 |
| 35. | 15.9855 | 1.2927 | 0.7659 | 59.0629 | 18.3825 | 1.2586 | 0.7445 | 48.9226 |
| 36. | 19.2399 | 1.2670 | 0.5811 | 67.9310 | 12.6966 | 1.2885 | 0.7795 | 41.9944 |
| 37. | 19.5613 | 1.2379 | 0.6048 | 49.4563 | 16.4697 | 1,2793 | 0.7426 | 54.0776 |
| 38. | 19.3353 | 1.2406 | 0.6758 | 46.4911 | 18.5644 | 1.2750 | 0.7126 | 61.5585 |
| 39. | 15.0793 | 1.2538 | 0.7495 | 37.2738 | 16.2829 | 1.2905 | 0.7451 | 59.9064 |
| 40. | 21.7750 | 1.2234 | 0.5666 | 49.0667 | 19.3650 | 1.2589 | 0.7438 | 52.1628 |
| 41. | 21.2913 | 1.2515 | 0.5906 | 63.2767 | 15.3883 | 1.2615 | 0.6653 | 45.9360 |
| 42. | 19.3210 | 1.2293 | 0.5806 | 45.5530 | 17.8463 | 1.2653 | 0.7838 | 48.3728 |
| 43. | 19.6009 | 1.2449 | 0.6704 | 49.7207 | 18.7307 | 1.2711 | 0.7191 | 59.0194 |
| 44. | 18.0937 | 1.2701 | 0.6495 | 61.0716 | 15.0095 | 1.2871 | 0.8409 | 44.7818 |
| 45. | 15.8084 | 1.2489 | 0.7761 | 35.5877 | 14.4959 | 1.2994 | 0.8309 | 50.2210 |
| 46. | 15.7104 | 1.2834 | 0.7055 | 56.5855 |  |  |  |  |
| 47. | 20.8017 | 1.2351 | 0.5518 | 54.0037 |  |  |  |  |
| 48. | 18.3704 | 1.2546 | 0.6035 | 55.5098 |  |  |  | , |
| 49. | 18.1195 | 1.2701 | 0.6643 | 60.1168 |  |  |  |  |
| 50. | 19.9951 | 1. 2565 | 0.6362 | 59.9892 |  |  |  |  |
| 51. | 15.7974 | 1,2594 | 0.7364 | 42.6064 |  |  |  |  |
| 52. | 14.3508 | 1.2730 | 0.6826 | 47.2688 |  |  |  |  |
| 53. | 16.5308 | 1.2583 | 0.6193 | 50.8118 |  |  |  |  |
| 54. | 19.5489 | 1.2384 | 0.5896 | 50.5878 |  |  |  |  |
| 55. | 14.1945 | 1.2705 | 0.6841 | 45.5747 | i |  |  |  |
| 56. | 18.7336 | 1.2433 | 0.6343 | 48.3947 |  |  |  |  |
| 57. | 14.4528 | 1.2453 | 0.7627 | 31.5465 |  |  |  |  |
| 58. | 18.1124 | 1.2537 | 0.6887 | 49.0423 |  |  |  |  |
|  |  |  |  |  |  | - |  |  |

Table 6. Fitting of Von-Bertalanffy frof $H_{t}=a[1-b \text { exp(xt) }]^{3}$ using twelve weeks body woights of Japanese quails

| Sl. | Male ( ${ }^{\text {a }}$ = 210 g ) |  |  |  | Female ( $\mathrm{a}=230 \mathrm{~g}$ ) |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| No. | b | k | $\mathrm{r}^{2}$ | 0 | b | k | $r^{2}$ | 8 |
| 1. | 0.6408 | -0.1831 | 0.9441 | 13.7832 | 0.6879 | -0.1843 | 0.9542 | 12.9738 |
| 2. | 0.6369 | -0.1643 | 0.9509 | 11.7185 | 0.7812 | -0.2386 | 0.9845 | 8.5417 |
| 3. | 0.5754 | -0.1751 | 0.9156 | 16.2999 | 0.6357 | -0.1625 | 0.9404 | 13.7063 |
| 4. | 0.6694 | -0.1771 | 0.9625 | 10.8239 | 0.7449 | -0.2063 | 0.9725 | 10.6347 |
| 5. | 0.6778 | -0.1757 | 0.9676 | 9.9103 | 0.8090 | -0.2145 | 0.9895 | 6.7366 |
| 6. | 1.0482 | -0.3473 | 0.9966 | 4.3183 | 0.7115 | -0.1919 | 0.9702 | 10.6442 |
| 7. | 0.5540 | -0.1620 | 0.8775 | 19.0038 | 0.7343 | -0.1867 | 0.9783 | 9.2312 |
| 8. | 0.6298 | -0,2006 | 0.9494 | 13.9593 | 0.6816 | -0.1940 | 0.9416 | 15.7477 |
| 9. | 0.7391 | -0. 1984 | 0.9830 | 7.8234 | 0.6467 | -0.1442 | 0.9626 | 9.9326 |
| 10. | 0.5607 | -0.1596 | 0.8936 | 17.4203 | 0.5445 | -0.1398 | 0.8196 | 22.9441 |
| 11. | 0.5551 | -0.1685 | 0.9043 | 16.9174 | 0.6392 | -0.1713 | 0.9463 | 13.6855 |
| 12. | 0.6094 | -0.1871 | 0.9255 | 16.3635 | 0.6256 | $-0.1735$ | 0.9215 | 17.2914 |
| 13. | 0.5987 | -0.1719 | 0.9157 | 16.2273 | 0.7415 | -0. 2085 | 0.9757 | 10.7214 |
| 14. | 0.7485 | -0.1937 | 0.9795 | 8.5130 | 0.6719 | -0.1636 | 0.9464 | 13.4739 |
| 15. | 0.6966 | -0.1716 | 0.9666 | 10.0045 | 0.6896 | -0.1971 | 0.9687 | 11.8086 |
| 16. | 0.7179 | -0.1789 | 0.9653 | 10.5893 | 0.7805 | -0.2278 | 0.9866 | 8.7840 |
| 17. | 0.7022 | -0.1937 | 0.9526 | 13.2450 | 0.6925 | -0.1674 | 0.9623 | 11.3534 |
| 18. | 0.7253 | -0.2367 | 0.9742 | 11.2970 | 1.3003 | -0.3528 | 0.9698 | 18.0415 |
| 19. | 0.6373 | -0.2046 | 0.9496 | 13.9069 | 0.7984 | -0.2213 | 0.9935 | 5.5094 |
| 20. | 0.6867 | -0.2038 | 0.9467 | 13.8003 | 0.6457 | -0.2022 | 0.9427 | 15.2226 |
| 21. | 0.5936 | -0.1851 | 0.9197 | 16.7332 | 0.8103 | -0.1734 | 0.9836 | 7.4176 |
| 22. | 0.6046 | -0.1639 | 0.9148 | 15.8026 | 0.7071 | -0.1926 | 0.9630 | 12.1684 |
| 23. | 0.6271 | -0.2092 | 0.9487 | 14.4187 | 0.7464 | -0.2134 | 0.9715 | 10.9862 |
| 24. | 0.6678 | -0.2056 | 0.9620 | 12.2443 | 0.7218 | -0.1767 | 0.9821 | 8.0268 |
| 25. | 0.5794 | -0.1703 | 0.8998 | 17.7348 | 0.6746 | -0.1904 | 0.9659 | 11.8295 |
| 26. | 0.6280 | -0.1700 | 0.9450 | 12.6784 | 0.6778 | -0.2099 | 0.9614 | 13.6960 |
| 27. | 0.5853 | -0.1702 | 0.9191 | 15.6414 | 0.6939 | 0.1713 | 0.9631 | 11.2776 |
| 28. | 0.7138 | -0.2043 | 0.9755 | 9.7315 | 0.7703 | -0.2125 | 0.9865 | 7.7652 |
| 29. | 0.6948 | $\therefore 0.2213$ | 0.9598 | 12.9903 | 0.7073 | -0.2191 | 0.9585 | 14.2337 |
| 30. | 0.6340 | -0.1774 | 0.9527 | 12.1551 | 0.6571 | -0.1808 | 0.9460 | 14.3384 |

Contd

Table 6. (Contd.)

| $\begin{aligned} & \text { S1. } \\ & \text { No. } \end{aligned}$ | Male ( $\mathrm{a}=210 \mathrm{~g}$ ) |  |  |  | Female ( $\mathrm{a}=230 \mathrm{~g}$ ) |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | b | k | $r^{2}$ | a | b | k | $\mathrm{r}^{2}$ | 5 |
| 31. | 0.6234 | -0.1888 | 0.9406 | 14.5642 | 0.6873 | -0.1824 | 0.9583 | 12.4632 |
| 32. | 0.6279 | -0.1892 | 0.9347 | 15.2810 | 0.7867 | -0.2244 | 0.9709 | 11.3698 |
| 33. | 0.7336 | -0.1581 | 0.9841 | 6.3162 | 0.8283 | -0.2322 | 0.9784 | 10.0605 |
| 34. | 0.6123 | -0.1908 | 0.9331 | 15.4928 | 0.7128 | -0.1903 | 0.9781 | 9.0214 |
| 35. | 1.2237 | -0.3506 | 0.9896 | 8.9333 | 0.7092 | -0.1873 | 0.9787 | 8.9344 |
| 36. | 0.6778 | -0.2408 | 0.9612 | 14.0067 | 0.8154 | -0.1772 | 0.9830 | 7.9689 |
| 37. | 0.5927 | -0.1617 | 0.9159 | 15.4530 | 0.8031 | -0.2166 | 0.9871 | 7.7156 |
| 38. | 0.6268 | -0.1718 | 0.9496 | 12.1137 | 0.7941 | -0.2408 | 0.9884 | 7.6628 |
| 39. | 0.6948 | - -0.1553 | 0.9757 | 7.8093 | 0.9019 | -0.2564 | 0.9900 | 7.3125 |
| 40. | 0.5419 | $-0.1498$ | 0.8765 | 17.8171 | 0.7158 | -0.2030 | 0.9632 | 12.1426 |
| 41. | 0.6382 | -0.2282 | 0.9488 | 15.2913 | 0.6960 | -0.1536 | 0.9311 | 14.4841 |
| 42. | 0.5738 | -0.1420 | 0.8635 | 18.0756 | 0.7615 | -0.2046 | 0.9636 | 12.1104 |
| 43. | 0.6405 | -0.1863 | 0.9555 | 12.0456 | 0.7807 | -0.2352 | 0.9544 | 14.6611 |
| 44. | 0.7502 | -0.2425 | 0.9792 | 10.1079 | 0.9174 | -0.2311 | 0.9894 | 7.3058 |
| 45. | 0.6931 | -0.1573 | 0.9819 | 6.7420 | 1.0377 | -0.2670 | 0.9896 | 8.1224 |
| 46. | 0.8286 | -0.2425 | 0.9895 | 7.0487 |  |  |  |  |
| 47. | 0.5620 | -0.1646 | 0.8892 | 18.2026 |  |  |  |  |
| 48. | 0.6365 | -0.1863 | 0.9364 | 15.0076 |  |  |  |  |
| 49. | 0.7642 | -0.2462 | 0.9843 | 8.8511 |  |  |  |  |
| 50. | 0.6665 | -0.2237 | 0.9588 | 12.9626 |  |  |  |  |
| 51. | 0.7027 | -0.1734 | 0.9750 | 8.5524 |  |  |  |  |
| 52. | 0.7363 | -0.1812 | 0.9638 | 10.9674 |  |  |  |  |
| 53. | 0.6576 | -0.1721 | 0.9327 | 14.5268 |  |  |  |  |
| 54. | 0.5819 | -0.1596 | 0.9057 | 16.1697 |  |  |  |  |
| 55. | 0.7195 | -0.1713 | 0.9601 | 11.0457 |  |  |  |  |
| 56. | 0.6248 | -0.1687. | 0.9299 | 14.4928 |  |  |  |  |
| 57. | 0.6864 | -0.1374 | 0.9663 | 8.8203 |  |  |  |  |
| 58. | 0.6849 | $-0.1923$ | 0.9715 | 9.9683 |  |  |  |  |

Table 7. Fitting of modified exponential form $W_{t}=k+a b^{t}$ using twelve weeks body weights of Japanese quails

|  | Male |  |  |  |  | Female |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| No. | k | a | b | $r^{2}$ | B | k | a | b | $\mathrm{r}^{2}$ | 3 |
| 1. | 163.9384 | -245.0447 | 0.7521 | 0.9565 | 12.3864 | 200.9154 | -245.1225 | 0.8 .366 | 0.9708 | 10.334 |
| 2. | 165.3366 | -215.8630- | 0.8052 | 0.9810 | 7.4207 | 226.1211 | -282.1480 | 0.8327 | 0.9877 | 7.704 |
| 3. | 159.8471 | -241.6212 | 0.7212 | 0.9766 | 8.7057 | 173.3957 | -228.8422 | 0.7918 | 0.9780 | 8.400 |
| 4. | 167.7558 | -229.2746 | 0.7948 | 0.9721 | 9.5396 | 210.0499 | -262.0112 | 0.8346 | 0.9776 | 9.609 |
| 5. | 170.6955 | -223.6132 | 0.8117 | 0.9782 | 8.1923 | 266.6588 | -301.7841 | 0.8948 | 0.9880 | 7.146 |
| 6. | 225.6341 | -304.7634 | 0.8108 | 0.9782 | 11.3062 | 212.4865 | -255.0987 | 0.8489 | 0.9823 | 8.207 |
| 7. | 150.5409 | -249.2208 | 0.6839 | 0.9588 | 11.3223 | 198.8199 | -261.1647 | 0.8280 | 0.9752 | 10.275 |
| 8. | 177.8341 | -253.6819 | 0.7646 | 0.9724 | 10.3710 | 213.3607 | -267.4097 | 0.8356 | 0.9562 | 13.717 |
| 9. | 230.7521 | - 266.5086 | 0.8846 | 0.9785 | 8.7578 | 165.0680 | -209.8250 | 0.8164 | 0.9885 | 5.626 |
| 10. | 147.0304 | -261.4604 | 0.6656 | 0.9465 | 13.3823 | 157.7321 | -231.4638 | 0.7316 | $0.9404^{\circ}$. | 13.391 |
| 11. | 156.3837 | -239.9615 | 0.7081 | 0.9773 | 8.4033 | 173.8451 | -246.6636 | 0.7667 | 0.9742 | 9.741 |
| 12. | 166.1128 | -259.1395 | 0.7300 | 0.9572 | 12.7911 | 167.9366 | -284.7830 | 0.7061 | 0.9440 | 15.749 |
| 13. | 157.1927 | -241.4964 | 0.7282 | 0.9609 | 11.3147 | 227.8141 | -285.5292 | 0.8478 | 0.9732 | 11.380 |
| 14. | 195.7738 | -246.9884 | 0.8476 | 0.9692 | 10.5027 | 168.9433 | -242.5633 | 0.7795 | 0.9615 | 11.753 |
| 15. | 174.6588 | -225.9999 | 0.8288 | 0.9772 | 8.4693 | 194.5022 | -271.6741 | 0,7895 | 0.9730 | 11.155 |
| 16. | 177.3641 | -234.0547 | 0.8265 | 0.9658 | 10.6913 | 283.5470 | -311.4117 | 0.8982 | 0.9928 | 5.623 |
| 17. | 165.2859 | -263.4979 | 0.7487 | 0.9373 | 16.0256 | 178.1192 | -241.5617 | 0.8069 | 0.9715 | 10.161 |
| 18. | 187.1813 | -285.4888 | 0.7574 | 0.9616 | 13.7961 | 300.0216 | -338.6324 | 0.8942 | 0.9842 | 9.270 |
| 19. | 172.9213 | -270.0449 | 0.7326 | 0.9574 | 13.4977 | 260.1619 | -303.5030 | 0.8816 | 0.9895 | 7.072 |
| 20. | 171.9506 | -250.8732 | 0.7636 | 0.9454 | 14.2928 | 210.5899 | -252.5438 | 0.8234 | 0.9754 | 9.894 |
| 21. | 158.8989 | - -285.4201 | 0.6728 | 0.9446 | 15.1265 | 274.9760 | -305.4132 | 0.9205 | 0.9765 | 8.877 |
| 22. | 148.5922 | -249.4521 | 0.7023 | 0.9418 | 13.9106 | 190.9590 | -262.9399 | 0.7967 | 0.9704 | 11,293 |
| 23. | 173.4858 | -273.2112 | 0.7216 | 0.9675 | 11.7267 | 215.7255 | -266.6726 | 0.8365 | 0.9806 | 9.099 |
| 24. | 176.1137 | -261.9190 | 0.7587 | 0.9659 | 11.9466 | 204.0956 | -250.1310 | 0.8525 | 0.9841 | $\checkmark 7.593$ |
| 25. | 157.1719 | -246.0018 | 0.7185 | 0.9556 | 12.1497 | 195.6006 | -255.1274 | 0.8069 | 0.9837 | 8.057 |
| 26. | 160.5914 | -227.1659 | 0.7693 | 0.9738 | 9.1244 | 203.2485 | -278.3002 | 0.7900 | 0.9763 | 10.630 |
| 27. | 154.0206 | -251.9375 | 0.7009 | 0.9635 | 11.2577 | 212.9009 | -249.7476 | 0.8668 | 0.9764 | 9.019 |
| 28. | 187.5848 | -249.0773 | 0.8130 | 0.9756 | 9.6615 | 296.0054 | -323.4373 | 0.9090 | 0.9857 | 7.904 |
| 29. | 178.0179 | -275.6724 | 0.7439 | 0.9568 | 13.8691 | 218.0186 | -282.6589 | 0.8188 | 0.9687 | 12.363 |
| 30. | 166.2498 | -227.3428 | 0.7794 | 0.9803 | 7.8729 | 212.9579 | -256.4950 | 0.8467 | 0.9730 | 10.222 |

Table 7. (Contd.)

| 51. | Male |  |  |  |  | Female |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | k | a | b | $r^{2}$ | B | k | a | b | $\mathrm{r}^{2}$ | B |
| 31. | 166.8358 | -253.6160 | 0.7398 | 0.9655 | 11.3732 | 209.9149 | -247.3254 | 0.8523 | 0.9797 | 8.6113 |
| 32. | 165.1522 | -258.3229 | 0.7305 | 0.9619 | 12.0600 | 277.1983 | -305.4186 | 0.8974 | 0.9771 | 10.0037 |
| 33. | 205.5454 | -233.7702 | 0.8962 | 0.9860 | 5.9429 | 276.8853 | -310.3356 | 0.8948 | 0.9830 | 8.8016 |
| 34. | 167.3957 | -253.4505 | 0.7336 | 0.9730 | 9.9598 | 225.0890 | -257.4897 | 0.8696 | 0.9883 | 6.5315 |
| 35. | 273.3392 | -320.4421 | 0.8827 | 0.9892 | 7.4939 | 229.6618 | -265.2423 | 0.8726 | 0.9880 | 6.7279 |
| 36. | 188.9175 | -285.9516 | 0.7461 | 0.9679 | 12.422 | 359.7891 | -389.3833 | 0.9415 | 0.9699 | 10.5803 |
| 37. | 148.8030 | -242.4947 | 0.7065 | 0.9576 | 11.6661 | 310.7318 | -346.4321 | 0.9119 | 0.9820 | 9.3285 |
| 38. | 171.2174 | -216.9931 | 0.8077 | 0.9842 | 6.7630 | 235.7042 | -291.6541 | 0.8435 | 0.9867 | 8.1902 |
| 39. | 174.7937 | -213.3276 | 0.8559 | 0.9825 | 6.8333 | 276.7466 | -332.1031 | 0.8796 | 0.9794 | 10.9740 |
| 40. | 143.9125 | -238.9098 | 0.6778 | 0.9660 | 9.8901 | 220.2350 | -259.5834 | 0.8514 | 0.9783 | 9.2791 |
| 41. | 180.2350 | -289.7511 | 0.7155 | 0.9578 | 14.0100 | 175.2133 | -228.8705 | 0.8292 | 0.9443 | 13.2978 |
| 42. | 131.1208 | -268.3412 | 0.6225 | 0.8936 | 17.8562 | 227.1223 | -260.4118 | 0.8694 | 0.9717 | 10.5774 |
| 43. | 168.0497 | -231.2896 | 0.7726 | 0.9767 | 8.6608 | 308.6038 | -332.1158 | 0.9090 | 0.9587 | 13.9283 |
| 44. | 198.4853 | -279.3443 | 0.7921 | 0.9699 | 12.1818 | 955.1025 | -975.2821 | 0.9785 | 0.9916 | 6.4043 |
| 45. | 189.2563 | -218.9779 | 0.8762 | 0.9885 | 5.394 | 485.2807 | -518.2831 | 0.9487 | 0.9898 | 7.5135 |
| 46. | 206.7318 | -276.0309 | 0.8256 | 0.9682 | 12.1911 |  |  |  |  |  |
| 47. | 153.3342 | -246.7610 | 0.7003 | 0.9613 | 11.1568 |  |  |  |  |  |
| 46. | 162.4482 | -264.3253 | 0.7237 | 0.9501 | 14.1011 |  |  |  |  |  |
| 49. | 204.0732 | -272.3026 | 0.8099 | 0.9727 | 11.1508 |  |  |  |  |  |
| 50. | 182.0986 | -260.9365 | 0.7564 | 0.9777 | . 9.4760 |  |  |  |  | - |
| 51. | 177.8614 | -222.1326 | 0.8390 | 0.9813 | 7.4297 |  |  |  |  |  |
| 52. | 190.2545 | -239.0581 | 0.8512 | 0.9587 | 11.7102 |  |  |  |  |  |
| 53. | 155.3202 | -238.1950 | 0.7506 | 0.9431 | 13.7223 |  |  |  |  |  |
| 54. | 151.6216 | -224.6323 | 0.7329 | 0.9730 | 8.7740 |  |  |  |  |  |
| 55. | 164.9066 | -229.6761 | 0.8087 | 0.9512 | 12,599 |  |  |  |  |  |
| 56. | 151.1785 | -245.8327 | 0.7186 | 0.9402 | 14.1243 |  |  |  |  |  |
| 57. | 175.5328 | -201.3312 | 0.8830 | 0.9739 | 7.3288 |  |  |  |  |  |
| 58. | 182.0564 | -242.539 | 0.8094 | 0.9765 | 9.3833 |  |  |  |  |  |
|  |  | . |  |  | - |  |  |  |  |  |

Table 8. Fitting of logistic form $W_{t} \quad a / l+b$ exp(-kt) using twelve weeks body weights of Japanese qualls

| s1. | Male |  |  |  |  | Female |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | a | b | k | $r^{2}$ | 5 | $a$ | b | k | $\mathrm{r}^{2}$ | E |
| 1. | 152.5710 | 38.8391 | 0.8162 | 0.9917 | 6.4079 | 163.7900 | 38.4549 | 0.7658 | 0.9555 | 14.1010 |
| 2. | 144.6990 | 33.7999 | 0.7619 | 0.9888 | 6.9623 | 185.3642 | 46.6043 | 0.8088 | 0.9586 | 15.6354 |
| 3. | 152.1891 | 43.3044 | 0.9037 | 0.9742 | 10.7228 | 153.4382 | 39.2967 | 0.8245 | 0.9552 | 13.1737 |
| 4. | 148.6116 | 36.0033 | 0.7704 | 0.9926 | 5.7522 | 170.5569 | 41.9428 | 0.7885 | 0.9501 | 15.2617 |
| 5. | 147.1801 | 35.1431 | 0.7568 | 0.9872 | 7.0510 | 174.7130 | 38.3427 | 0.7111 | 0.9616 | 23.5972 |
| 6. | 193.1123 | 51.5837 | 0.8250 | 0.9851 | 10.2302 | 167.6904 | 40.0934 | 0.7664 | 0.9527 | 14.8025 |
| 7. | $145 \% 9866$ | 45.6057 | 0.9513 | 0.9755 | 9.7540 | 164.3494 | 41.4814 | 0.7698 | 0.9848 | 8.1877 |
| 8. | 163.4043 | 41.9336 | 0.8407 | $0: 9858$ | 8.8767 | 174.2610 | 40.6608 | 0.7670 | 0.9710 | 12.1802 |
| 9. | 162.1092 | 33.4109 | 0.6765 | 0.9920 | 6.4594 | 141.3042 | 31.8986 | 0.7478 | 0.9825 | 7.8714 |
| 10. | 143.4711 | 42.6213 | 0.9423 | 0.9911 | 6.2749 | 147.7484 | 40.1381 | 0.8968 | 0.9458 | 13.8536 |
| 11. | 150.0159 | 42.4876 | 0.9195 | 0.9721 | 10.9997 | 158.8080 | 40.4362 | 0.8374 | 0.9765 | 10.1852 |
| 12. | 156.8975 | '42.6880 | 0.8754 | 0.9886 | 7.3226 | 160.7172 | 48.0852 | 0.9147 | 0.9918 | 10.1852 6.2218 |
| 13. | 148.9376 | 40.3527 | 0.8628 | 0.9819 | 8.6864 | 180.4389 | 41.0787 | 0.7462 | 0.9898 | 7.9240 |
| 14. | 155.6389 | 34.3841 | 0.7017 | 0.9926 | 5.3995 | 152.4851 | 38.7752 | 0.7891 | 0.9943 | 7.9240 5.0401 |
| 15. | 145.3980 | 35.1438 | 0.7382 | 0.9931 | 5.0736 | 173.0641 | 43.8345 | 0.8140 |  | 5.0401 |
| 16. | 148.1724 | 36.1794 | 0.7338 | 0.9955 | 4.0477 | 185.3612 | 43.8345 39.9989 | 0.8140 0.7321 | 0.9939 0.9500 | 6.4892 17.0383 |
| 17. | 153.2709 | 43.7801 | 0.8403 | 0.9964 | 3.6507 | 154.1474 | 38.7047 | 0.7737 | 0.9500 | 17.0383 |
| 18. | 172.3861 | 49.3458 | 0.8662 | 0.9943 | 5.8306 | 196.6278 | 43.5528 | 0.7737 0.7426 | 0.9908 | 6.1210 |
| 19. | 162.7998 | 44.6126 | 0.8839 | 0.9829 | 9.0639 | 196.6278 | 43.5528 | 0.7426 | 0.9414 | 19.0055 |
| 20. | 156.2952 | 40.3950 | 0.8332 | 0.9674 | .9 .0639 11.3178 | 182.7520 | 39.6691 | 0.7194 | 0.9813 | 10.3071 |
| 21. | 154.3252 | 49.4482 |  | 0.9674 | 11.3178 | 178.1444 | 42.5364 | 0.8221 | 0.9216 | 20.5125 |
| 22. |  | 49.4482 | 0.9680 | 0.9912 | 6.4013 | 150.3265 | 34.1394 | 0.6559 | 0.9567 | 12.0654 |
| 22. | 142.8177 | 40.3136 | 0.8786 | 0.9926 | 5.4400 | 167.0787 | 44.6193 | 0.8236 | 0.9727 | 11.1325 |
| 23. | 164.7700 | 45.8276 | 0.9039 | 0.9861 | 8.5766 | 174.9730 |  |  |  | 11.1325 |
| 24. | 162.2651 | 41.7339 | 0.8399 | 0.9914 |  |  | 43.9420 | 0.7982 | . 0.9490 | 15.9868 |
| 25. | 149.4700 | 39.9554 |  | 0.9914 | 6.5969 | 159.8176 | 36.0041 | 0.7203 | 0.9874 | 7.4966 |
| 26. | 146.7926 | 35.3372 | 0.8826 | 0.9801 | 8.8259 | 170.1170 | 41.8489 | 0.7960 | 0.9814 | 10.3575 |
| 27. | 147.9313 | 35.3372 41.4027 | 0.8078 | 0.9848 | 7.6555 | 180.8682 | 47.7206 | 0.8361 | 0.9836 | 10.4336 |
| 28. | 147.9313 161.0933 | 41.4027 | 0.9143 | 0.9861 | 7.4762 | 160.0292 | 34.9574 | 0.7105 | 0.9776 | 9.9324 |
| 29. | 161.0933 | 39.5184 | 0.7682 | 0.9925 | 6.3009 | 182.7188 | 37.5701 | 0.6864 | 0.9751 | 12.1881 |
| . | 165.5757 | 48.4897 | 0.8857 | 0.9825 | 8.9757 | 184.4013 |  |  |  |  |
| 30. | 150:6449 | 37.0269 | 0.8005 |  |  | 184.4013 | 46.2166 | 0.8109 | 0.9690 | 13.4583 |
|  |  |  | 0.6005 | 0.9851 | 8.3204 | 170.4015 | 38.9918 | 0.7632 | 0.9638 | 13.4687 |

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Table 8, (contd.)

| $\begin{aligned} & \text { si. } \\ & \text { No. } \end{aligned}$ | Male |  |  |  |  | Female |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | a | $b$ | k | $r^{2}$ | 8 | a | b | k | $r^{2}$ | 5 |
| 31. | 156.6387 | 42.5693 | 0.8612 | 0.9906 | 7.0740 | 165.4490 | 39.0798 | 0.7589 | 0.9525 | 14.6862 |
| 32. | 155.9276 | 46.2486 | 0.8868 | 0.9877 | 7.6311 | 180.1149 | 39.0989 | 0.7311 | 0.9335 | 18.5582 |
| 33. | 135.3267 | 30.3695 | 0.6587 | 0.9754 | 8.0874 | 181.3279 | 40.2197 | 0.7326 | 0.9394 | 17.7085 |
| 34. | 157.9105 | 47.0649 | 0.8974 | 0.9795 | .9.9371 | 167.0109 | 35.7250 | 0.7319 | 0.9512 | 14.9132 |
| 35. | 189.3867 | 45.9464 | 0.7392 | 0.9751 | 12.1992 | 1688429 | 36.0780 | 0.7227 | 0.9694 | 12.1634 |
| 36. | 176.0976 | 53.7911 | 0.9041 | 0.9826 | 10.2178 | 164.4605 | 32.1779 | 0.5881 . | 09948 | 4.5155 |
| 37. | 142.7670 | 39.0415 | 0.8815 | 0.9911 | 6.0080 | 185.1556 | 38.3856 | 0.6846 | 0.9797 | 10.7926 |
| 38. | 149.5566 | 33.7501 | 0.7666 | 0.9729 | 10.5430 | 1884689 | 44.2608 | 0.7780 | 0.9735 | 127843 |
| 39. | 136.2337 | 29.4012 | 0.6829 | 0.9932 | 4.6018 | 193.5117 | 44.5576 | 0.7389 | 0.9756 | 12.0826 |
| 40. | 139.9199 | 41.2120 | 0.9386 | 0.9796 | 8. 6642 | 172.4566 | 39.4060 | 0.7753 | 0.9327 | 17.8678 |
| 41. | 171.6660 | 48.9980 | 0.9173 | 0.9826 | 9.8546 | 144.5802 | 32.7734 | 0.7193 | 0.9728 | 9.1852 |
| 42. | 128.6274 | 41.4450 | 0.9866 | 0.9827 | 6.5552 | 166.9997 | 38.6369 | 0.7454 | 0.9289 | 17.9847 |
| 43. | 153.2663 | 36.9524 | 0.8068 | 0.9781 | 9.8691 | 189.5505 | 42.6814 | 0.7320 | 0.9151 | 22.2938 |
| 44. | 175.5572 | 44.7244 | 0.8180 | 0.9902 | 7.4456 | 188.7253 | 35.8378 | 0.6188 | 0.9778 | 10.8188 |
| 45. | 138.6245 | $26.7885$ | 0.6486 | $0.9877$ | 6.2181 | 195.4401 | 39.6018 | 0.6436 | 0.9821 | 10.1162 |
| 46. | 171.9819 | 43.3034 | 0.7638 | $0.9943$ | $5.3873$ |  |  |  |  | 10.1162 |
| 47. | 147.4760 | 43.2312 | 0.9199 | 0.9787 | 9.2910 |  |  |  |  |  |
| 48. | 153.9172 | 44.8359 | 0.8859 | 0.9954 | 4.6554 |  |  |  |  |  |
| 49. | 176.3955 | 42.9709 | 0.7785 | 0.9911 | 7.6518 |  |  |  |  |  |
| 50. | 168.1246 | 48.3309 | 0.8870 | 09680 | 12.8064 |  |  |  |  |  |
| 51. | 144.7622 | 33.3731 | 0.7191 | 0.9848 | 7.2827 |  |  |  |  |  |
| 52. | 150.4717 | 34.0455 | 0.6833 | 0.9962 | 3.9147 |  |  |  |  |  |
| 53. | 144.5246 | 38.1363 | 0.8059 | 0.9926 | 5,2550 | ' |  |  |  |  |
| 54. | 143.4013 | 39.4547 | 0.8665 | 0.9758 | 9.6590 |  |  |  |  |  |
| 55. | 142.4020 | 35.4530 | 0.7357 | 0.9982 | 2.3833 |  |  |  |  |  |
| 56. | 143.7593 | 37.6537 | 0.8505 | 0.9908 | 5.7188 |  |  |  |  |  |
| 57. | 126.3302 | 23.9437 | 0.6091 | 0.9920 | 4.5280 |  |  |  |  |  |
| 56. | 157.3121 | 36.5884 | 0.7695 | 0.9896 | 6.9291 | . |  |  |  |  |

Table 9. Fitting of Gompertz form $W_{t}=a \exp [-b \exp (-k t)]$ uing twelve weeks body weights of Japanese quail


Contd.

Table 9. (Contd.)

| S1. | Male |  |  |  |  | Female |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| N | 8 | b | k | $\mathrm{r}^{2}$ | 8- | a | $b$ | k | $\mathrm{r}^{2}$ | B |
| 31. | 160.1809 | 5.2446 | 0.5071 | 0.9968 | 3.3076 | 176.3127 | 4.4848 | 0.3919 | 0.9763 | 9.5574 |
| 32. | 159.0620 | 5.5511 | 0.5262 | 0.9957 | 3.8149 | 199.1793 | 4.2359 | 0.3489 | 0.9654 | 12.5675 |
| 33. | 150.1221 | 4.2419 | 0.3320 | 0.9899 | 5.0056 | . 200.2744 | 4.3847 | 0.3527 | 0.9716 | 21.5022 |
| 34. | 161.0165 | 5.4534 | 0.5274 | 0.9958. | 3.8129 | 180.9018 | 4.1817 | 0.3663 | 0.9785 | 9.0938 |
| 35. | 207.9940 | 4.7461 | 0.3612 | 0.9929 | 6.1260 | 183.2219 | 4.2544 | 0.3611 | 0.9870 | 7.1370 |
| 36. | 180.1215 | 5.7087 | - 0.5188 | 0.9958 | 4.3037 | 195.3182 | 4.2907 | 0.2755 | 0.9887 | 6.4459 |
| 37. | 144.9853 | 5.3604 | 0.5448 | 0.9965 | 3.1733 | 208.2546 | 4.4067 | 0.3253 | 0.9888 | 7.3534 |
| 38. | 156.1814 | 4.3610 | 0.4221 | 0.9876 | 6.1229 | 200.5710 | 4.7938 | 0.4021 | 0.9914 | 6.5949 |
| 39. | 146.4951 | 4.2989 | 0.3649 | 0.9976 | 2.4517 | 211.4276 | 4.8930 | 0.3677 | 0.9894 | 7.6041 |
| 40. | 141.3684 | 5.4142 | 0.5905 | 0.9953 | 3.4448 | 183.9129 | 4.4695 | 0.3984 | 0.9657 | 11.8911 |
| 41. | 174.6436 | 5.6516 | 0.5468 | 0.9920 | 5.7268 | 153.4808 | 4.6819 | 0.3985 | 0.9715 | 9.3145 |
| 42. | 129.6722 | 6.3541 | 0.6635 | 0.9741 | 8.0884 | 181.2520 | 4.3246 | 0.3727 | 0.9589 | 12.9123 |
| 43. | 158.3134 | 4.6927 | 0.4611 | 0.9907 | 5.5262 | 210.8006 | 4.2564 | 0.3399 | 0.9478 | 16.2553 |
| 44. | 182.3380 | 5.2599 | 0.4557 | 0.9963 | 4.0778 | 234.6334 | 4.1579 | 0.2616 | 0.9934 | 5.6635 |
| 45. | 151.5491 | 3.9899 | 0.3367 | 0.9951 | 3.4884 | 232.5885 | 4.4501 | 0.2888 | 0.9952 | 5.1286 |
| 46. | 182.0225 | 5.0885 | 0.4110 | 0.9966 | 3.9775 |  |  |  |  |  |
| 47. | 149.5542 | 5.4500 | 0.5631 | 0.9913 | 5.0249 |  |  |  |  |  |
| 48. | 156.8388 | 5.6686 | 0.5329 | 0.9966 | 3.5834 |  |  |  |  |  |
| 49. | 185.1975 | 4.8652 | 0.4206 | 0.9948 | 4.8067 |  |  |  |  |  |
| 50. | 172.4804 | 5.2958 | 0.5048 | 0.9896 | 6.4176 |  |  |  |  |  |
| 51. | 154:2059 | 4.4497 | 0.3861 | 0.9934 | 4.3798 |  |  |  |  |  |
| 52. | 162.2282 | 4.5628 | 0.3635 | 0.9880 | 6.2778 |  |  |  |  |  |
| 53. | 148.5103 | 5.2240 | 0.4814 | 0.9891 | 5.9308 |  |  |  |  |  |
| 54. | 146.2260 | 5.0748 | 0.5176 | 0.9916 | 4.8664 |  |  |  |  |  |
| 55. | 149.4904 | 4.9931 | 0.4172 | 0.9921 | 5.0622 |  |  |  |  |  |
| 56. | 146.574 | 5.2999 | 0.5229 | 0.9882 | 5.9668 |  |  |  |  |  |
| 57. | 139.4108 | 3.8540 | 0.3173 | 0.9892 | 4.7057 |  |  |  |  |  |
| 58. | 164.7302 | 4.6989 | 0.4235 | 0.9953 | 4.0006 |  |  |  |  |  |

Fitting of models using average body weights of Japanese qualls over twelve weeks

| sex | a | b | $x^{2}$ | B |
| :---: | :---: | :---: | :---: | :---: |
| Male | 11.5402 | 14.2256 | 0.9039 | 17.5456 |
| Female | 5.1070 | 16.1753 | 0.9538 | 13.4568 |
| Irrespective of sex | 8.7296 | 15.0774 | 0.9295 | 15.6984 |

Table 11. Quadratic $W_{t}=a+b_{1} t+b_{2} t^{2}$

| Sex | $a$ | $b$ | $b_{2}$ | $x^{2}$ | b |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Male | -32.7086 | 33.1894 | -1.4588 | 0.9926 | 5.1460 |
| Female | -28.3067 | 30.4955 | -1.1015 | 0.9951 | 4.6110 |
| Irrespective | -20.7853 | 32.0124 | -1.3027 | 0.9943 | 4.7077 |
| of sex |  |  |  |  |  |

Table 12. Exponential $W_{t}=a \exp (b t)$

| Sex | A | $\exp (b)$ | $\mathrm{r}^{2}$ | $B$ |
| :---: | :---: | :---: | :---: | :---: |
| Male | 18.4335 | 1.2520 | 0.6534 | 51.3204 |
| Female | 17.9536 | 1.2655 | 0.7336 | 52.3387 |
| Irrespective of gex | 18.2102 | 1.2581 | 0.6916 | 51.7545 |

Table 14. Modified exponential $W_{t}=k+a b^{t}$

| Sex | $k$ | $a$ | $b$ | $r^{2}$ | $s$ |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Male | 168.6513 | -240.1105 | 0.7718 | 0.9754 | 9.3476 |
| Female | 219.5777 | -265.1914 | 0.8532 | 0.9921 | 5.7300 |
| Irrespective | 187.1863 | -245.3882 | 0.8106 | 0.9858 | 7.3496 |
| of sex |  |  |  |  |  |

Table 15. Logistic $a / 1+b \exp (-k t)$

| Sex | a | $b$ | $k$ | $r^{2}$ | $s$ |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Male | 153.7328 | 39.1829 | 0.8161 | 0.9918 | 6.4281 |
| Fernale | 171.6814 | 39.2857 | 0.7496 | 0.9778 | 10.7745 |
| Irrespective <br> of sex | 161.5433 | 39.0390 | 0.7829 | 0.9863 | 8.3039 |

Table 16. Gompertz $\mathcal{H}_{t}=a \exp [-b \exp (-k t)]$

| Sex | $a$ | $b$ | $k$ | $r^{2}$ | $s$ |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: |
| -158.6057 | 4.9667 | 0.4677 | 0.9991 | 1.6903 |  |
| Male | 183.8152 | 4.5238 | 0.3846 | 0.9949 | 4.6230 |
| Female | 169.1025 | 4.7301 | 0.4267 | 0.9982 | 2.6054 |
| Irrespective <br> Of Bex |  |  |  |  |  |

Table 13. Von-Bertalanffy $W_{t}=a[1-b \exp (k t)]^{3}$

| Sex | b | k | $\mathrm{r}^{2}$ | $s$ |
| :---: | :---: | :---: | :---: | :---: |
| Male $(a=210 \mathrm{~g})$ | 0.6515 | -0.1854 | 0.9580 | 11.9131 |
| $\begin{aligned} & \text { Female } \\ & (\mathrm{a}=230 \mathrm{~g}) \end{aligned}$ | 0.7221 | -0.1954 | 0.9851 | 7.8073 |
| Irrespective of sex <br> (a) 220 g ) | 0.6813 | $-0.1880$ | 0.9724 | 10.0626 |

Table 17. Initial body weights $\left(W_{0}\right)$ and 'b' values of male and female Japanese quails for Rao's method

| Sl. | Male |  | Female |  |
| :---: | :---: | :---: | :---: | :---: |
| No. | $W_{0}$ | b | $\mathrm{W}_{0}$ | b |
| 1. | 7.0 | 0.009532 | 7.0 | 0.009383 |
| 2. | 7.0 | 0.008268 | 7.0 | 0.010811 |
| 3. | 7.0 | 0.009923 | 7.0 | 0.009777 |
| 4. | 6.9 | 0.009056 | 7.2 | 0.010352 |
| 5. | 6.9 | 0.008853 | 6.9 | 0.009385 |
| 6. | 6.9 | 0.010741 | 6.9 | 0.010228 |
| 7. | 6.9 | 0.009980 | 6.9 | 0.009567 |
| 8. | 7.0 | 0.010210 | 7.0 | 0.010455 |
| 9. | 7.0 | 0.008240 | 7.0 | 0.008020 |
| 10. | 7.0 | 0.009208 | 7.2 | 0.009408 |
| 11. | 7.0 | 0.009595 | 7.2 | 0.010195 |
| 12. | 7.2 | 0.010136 | 7.2 | 0.010521 |
| 13. | 7.2 | 0.009998 | 7.2 | 0.010385 |
| 14. | 7.2 | 0.008936 | 7.2 | 0.009359 |
| 15. | 7.2 | 0.008137 | 7.0 | 0.010283 |
| 16. | 7.2 | 0.008195 | 7.0 | . 0.010525 |
| 17. | 7.2 | 0.009277 | 7.0 | 0.009125 |
| 18. | 7.2 | 0.010391 | 7.4 | 0.011679 |
| 19. | 7.0 | 0.010215 | 7.4 | 0.010113 |
| 20. | 7.0 | 0.009807 | 7.4 | 0.011260 |
| 21. | 7.0 | 0.010021 | 7.0 | 0.007931 |
| 22. | 7.0 | 0.009146 | 7.0 | 0.010455 |
| 23. | 7.4 | 0.010568 | 7.0 | 0.010825 |
| 24. | 7.4 | 0.009834 | 7.0 | 0.009019 |
| 25. | 7.4 | 0.009552 | 7.0 | 0.010535 |
| 26. | 7.4 | 0.009063 | 7.0 | 0.011331 |
| 27. | 7.4 | 0.009406 | 7.0 | 0.008658 |
| 28. | 7.0 | 0.009573 | 7.3 | 0.010202 |
| 29. | 7.0 | 0.010816 | 7.3 | 0.011204 |

Table 17. (Contd.)

| Sl. | Male |  | Female |  |
| :---: | :---: | :---: | :---: | :---: |
|  | $\mathrm{W}_{0}$ | b | $W_{0}$ | b |
| 30. | 7.0 | 0.009374 | 7.3 | 0.010065 |
| 31. | 7.0 | 0.010039 | 7.2 | 0.010431 |
| 32. | 7.0 | 0.010150 | 7.2 | 0.010385 |
| 33. | 7.0 | 0.007082 | 7.2 | 0.010676 |
| 34. | 7.0 | 0.010326 | 7.2 | 0.009972 |
| 35. | 7.0 | 0.010861 | 7.2 | 0.009921 |
| 36. | 7.0 | 0.011209 | 7.2 | 0.007862 |
| 37. | 7.3 | 0.009264 | 7.5 | 0.010127 |
| 38. | 7.3 | 0.009258 | 7.5 | 0.011036 |
| 39. | 7.2 | 0.007353 | 7.5 | 0.010767 |
| 40. | 7.2 | 0.009300 | 7.5 | 0.010512 |
| 41. | 7.2 | 0.011744 | 7.5 | 0.008152 |
| 42. | 7.2 | 0.008783 | 7.0 | 0.010316 |
| 43. | 7.2 | 0.009587 | 7.2 | 0.011618 |
| 44. | 7.5 | 0.010677 | 7.2 | 0.009040 |
| 45. | 7.5 | 0.007571 | 7.2 | 0.010074 |
| 46. | 7.0 | 0.009984 |  |  |
| 47. | 7.0 | 0.009986 |  |  |
| 48. | 7.0 | 0.009821 |  |  |
| 49 | 7.0 | 0.010506 |  |  |
| 50. | 7.0 | 0.010801 |  |  |
| 51. | 7.0 | 0.008665 |  |  |
| 52. | 7.0 | . 0.007907 |  |  |
| 53. | 7.2 | 0.009109 |  |  |
| 54. | 7.2 | 0.009312 |  |  |
| 55. | 7.2 | 0.007680 |  |  |
| 56. | 7.2 | 0.009384 |  |  |
| 57. | 7.2 | 0.006609 |  |  |
| 58. | 7.2 | 0.009038 |  |  |

Table 18. Analysis of covariance of initial body weights ( $W_{0}$ ) and 'b' values by Rao's method

| Source | D.F. | SS(X) | SS(Y) | SP(XY) | $\begin{aligned} & \text { ADJ } \\ & \mathrm{D.F} \end{aligned}$ | ADJ M.S.S | F | Mean of <br> 'b' values | ADJ Values $w_{i o}-\left(x_{i o}-6.241748\right)$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Between groups | 1 | 579.6616 | 0.001085 | 0.794905 | 1 | 7.586867E-06 | 9.318347 | 0.009539 | 0.009250 |
| Within group | 101 | 2.75 | 0.000082 | 0.000902 | 100 | 8.141859E-07 |  | 0.0100432 | 0.000741 |
| Total | 102 | 582.4116 | 0.001176 | 0.795807 | 101 |  |  |  |  |

Regression estimate $=0.000328$
To test ' $b$ ' $\left.=\frac{[S P(X Y)}{S S(X)}\right]^{2}$, ADJ M.S.E.
$=0.3631<1$ (not significant)

Table i9. Total weekly fortnightly and mean weekly egg production per bird of Japanese quails

| Weeks | Total weekly egg production | Total fortnightly egg production | Mean weekly egg production |
| :---: | :---: | :---: | :---: |
| 1 | 23 |  | 0.7188 |
| 2 | 69 | 92 | 2.1563 |
| 3 | 119 |  | 3.7188 |
| 4 | 134 | 253 | 4.1875 |
| 5 | 153 |  | 4.7813 |
| 6 | 147 | 300 | 4.5938 |
| 7 | 138 |  | 4.3125 |
| 8 | 178 | 316 | 5.5625 |
| 9 | 196 |  | 6.1250 |
| 10 | 180 | 376 | 5.6250 |
| 11 | 185 |  | 5.7813 |
| 12 | 166 | 351 | 5.1875 |
| 13 | 170 |  | 5.3125 |
| 14 | 168 | 338 | 5.2500 |
| 15 | 141 |  | 4.4063 |
| 16 | 121 | 262 | 3.7813 |
| 17 | 117 |  | 3.6563 |
| 18 | 154 | 271 | 4.8125 |
| 19 | 158 |  | 4.9375 |
| 20 | 168 | 326 | 5.2500 |
| 21 | 179 |  | 5.5938 |
| 22 | 180 | 359 | 5.6250 |
| 23 | 164 |  | 5.1250 |
| 24 | 145 | 309 | 4.5313 |
| 25 | 146 |  | 4.5625 |
| 26 | 139 | 285 | 4.3438 |

Table 20. Hen housed and hen day egg production of Japanese quails during the period l.l.l989 to 30.9.1989

| Months | Number of layers at the begining of month | Number of days in each month | Total hen days | Total <br> number <br> of eggs <br> produced | Hen <br> housed <br> egg <br> production | Number of birds expired | Number of days absent by dead birds during the month | Hen <br> days <br> of <br> survi- <br> vors | Hen <br> day <br> egg <br> product- <br> ion |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| (I) | (2) | (3) | $\begin{gathered} (4) \\ (=2 \times 3) \end{gathered}$ | (5) | $\begin{aligned} & \left(\frac{5}{4}\right)^{6} \times 100 \end{aligned}$ | (7) | (8) | $\begin{gathered} (9) \\ (4)-(8) \end{gathered}$ | $\begin{aligned} & (10) \\ & (5) \times 100 \end{aligned}$ |
| February | 62 | 28 | 1736 | 0 | 0 | 0 | -- | 1736 | 0 |
| March | 62 | 31 | 1922 | 38 | 1.9771 | 0 | -- | 1922 | 1.9771 |
| April | 62 | 30 | 1860 | 885 | 47.5806 | 0 | -- | 1850 | 47.5806 |
| May | 62 | 31 | 1922 | 1304 | 67.8460 | 3 | 41 | 1881 | 69.3248 |
| June | 59 | 30 | 1770 | 1294 | 73.1073 | 3 | 12 | 1758 | 73.6064 |
| July | 56 | 31 | 1736 | 976 | 56.2212 | 5 | 29 | 1707 | 57.1763 |
| August | 51 | 31 | 1581 | 1003 | 63.4409 | 8 | 195 | 1386 | 72.3665 |
| September | 43 | 30 | 1290 | 722 | 55.9690 | 2 | 41 | 124.9 | 57.8063 |

Table 21. Fitting of mathematical models for, weekly egg production of Japanese quails

| Models | a | b | c | d | $r^{2}$ | s | I |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Linear | 121.4000 | 1.9419 | -- | -- | 0.1601 | 34.7162 | 34.7162 |
| Exponential | 101.1137 | 0.0237. | -- | -- | 0.1264 | 38.1350 | 12228.2900 |
| Parabolic exponential | 57.0929 | 0.1462 | -4.5362 | -- | 0.4850 | 0.3165 | 101.4965 |
| Inverse polynomial | 0.0205 | 0.0032 | 0.0001 | -- | 0.9213 | 0.0145 | 280.6831 |
| Gamma function | 35.5322 | 1.0571 | -0.0833 | -- | 0.8223 | 0.1860 | 59.6288 |
| Gamma type function | 1049.1900 | 3.6236 | 0.2729 | -4.0810 | 0.9223 | 0.1258 | 40.3254 |
| Quadratic function | 69.0193 | 13.1663 | -0.4157 | -- | 0.4889 | 27.6640 | 27.6640 |
| Quadratic function log scale | 13.4228 | 124.6064 | -25.6003 | -- | 0.7471 | 19.4584 | 19.4584 |
| Quadratic-cum-log | 31.6676 | -17.2984 | 0.2521 | 122.8046 | 0.7602 | 19.3758 | 19.3758 |
| Emperical | 0.0266 | -0.0775 | -- | -- | 0.6480 | 23.4910 | 7532.5820 |
| Linear hyperbolic | 192.6841 | -1.3239 | -183.4538 | -- | 0.7194 | 20.4970 | 20.4970 |

Table 22. Fitting of mathematical models for fortnightly egg production of Japanese quails

| Models | a | b | c | d | $r^{2}$ | s | I |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Linear | 241.1923 | 7.7198 | -- | -- | 0.1749 | 68.1949 | 68.1949 |
| Exponential | 210.2932 | 0.0420 | -- | -- | 0.1426 | 73.1730 | 47549.3000 |
| Parabolic exponential | 119.7881 | 0.2671 | -0.0161 | -- | 0.5443 | 0.2650 | 172.2323 |
| Inverse polynomial | 0.0063 | 0.0013 | 0.0001 | -- | 0.9374 | 0.0034 | 269.1243 |
| Gamma function | 129.8996 | 1.1039 | -0.1627 | -- | 0.8166 | 0.1681 | 109.2483 |
| Gamma type function | 90987.1600 | 4.8825 | 0.7049 | -7.5894 | 0.9240 | 0.1141 | 74.1584 |
| Quadratic function | 126.0351 | 53.7827 | -3.2902 | -- | 0.5245 | 54.2986 | 54.2986 |
| Quadratic function log scale | 97.1038 | 268.8874 | -75.5807 | -- | 0.8018 | 35.0586 | 35.0586 |
| Quadratic-cum-log | 195.6462 | -107:5919 | 3.3230 | 370.8704 | 0.8190 | 35.3130 | 35.3130 |
| Emperical | 0.0074 | -0.1435 | -- | -- | 0.6793 | 43.3137 | 28146.1400 |
| Linear hyperbolic | 450.5895 | -10.0145 | 348.5215 | -- | 0.8052 | 34.7535 | 34.7535 |

Table 23. Fitting of mathematical models for hen housed egg production of Japanese quails

| Models | a | b | c | d | $\mathrm{r}^{2}$ | s | I |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Linear | 26.2958 | 6.5026 | -- | -- | 0.3500 | 20.9682 | 20.9882 |
| Exponential | 8.3317 | 0.3720 | -- | -- | 0.1428 | 38.8248 | 3298.7340 |
| Parabolic exponential | 0.6035 | 2.1221 | -0.2188 | -- | 0.7807 | 0.7445 | 63.2596 |
| Inverse polynomial | 0.6405 | -0.2770 | 0.0302 | -- | 0.6786 | 0.1150 | 156.6152 |
| Gamm function | 8.4609 | 5.2193 | -1. 2209 | -- | 0.9371 | 0.3987 | 33.8736 |
| Gamma type function | $1.109345 \mathrm{E}+19$ | 25.4443 | 5.7197 | -48.8799 | 0.9963 | 0.1123 | 9.5416 |
| Quadratic function | -27.2613 | 42.2073 | -4.4631 | -- | 0.8448 | 11.4576 | 11.4576 |
| Quadratic function log scale | I. 2589 | 96.6707 | -35.3734 | -- | 0.9571 | 6.0220 | 6.0220 |
| Quadratic-cum-log | 50.4287 | -51.2273 | 1.9775 | 137.3186 | 0.9609 | 6.6385 | 6.6385 |
| Emperical | 0.1197 | 0.0668 | --- | -- | 0.2687 | 30.4821 | 2589.9000 |
| Linear hyperbolic | 127.2179 | -7.7595 | -118.4475 | -- | 0.9533 | 6.2830 | 6.2830 |

Table 24. Fitting of mathematical models for hen day egg production of Japanese quails

| Models | a | b | $c$ | d | $r^{2}$ | s | I |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Linear | 24.9896 | 7.3183 | -- | -- | 0.4022 | 21.1125 | 21.1125 |
| Exponential | 8.1594 | 0.3847 | -- | -- | 0.1748 | 40.4675 | 3542.5890 |
| Parabolic exponential | 0.5927 | 2.1329 | -0.2185 | -- | 0.7884 | 0.7398 | 64.7593 |
| Inverse Folynomial | 0.6414 | -0.2772 | 0.0301 | -- | 0.6838 | 0.1146 | 165.6879 |
| Gamm function | 8.2854 | 5.1986 | -1.2019 | -- | 0.9378 | 0.4011 | 35.1081 |
| Gamma type function | $4.855436 \mathrm{E}+18$ | 25.0332 | 5.6049 | -47.9365 | 0.9934 | 0.1505 | 13.1725 |
| Quadratic function | -28.5834 | 43.0336 | -4.4644 | -- | 0.8513 | 11.7737. | 11.7737 |
| Quadratic function log scale | 1.2386 | 95.4551 | -33.5394 | -- | 0.9382 | 7.5873 | 7.5873 |
| Quadratic-cum-1og | 42.7397 | -42.7434 | 1.4483 | 126.064 | 0.9401 | 8.6287 | 8.6287 |
| Emperical | 0.1222 | 0.0795 | -- | -- | 0.3046 | 31.5089 | 2758.3370 |
| Linear hyperbolic | 124.5305 | -6.7486 | -116.8264 | -- | 0.9346 | 7.8062 | 7.8062 |

Table 25. Weekly mean temperature and humidity with weekly average egg production per bird

| Average weekly egg production Y | Previous three weeks average |  |
| :---: | :---: | :---: |
|  | Temperature $\mathrm{Z}_{1}$ | $\begin{gathered} \text { Humidity } \\ \mathrm{Z}_{2} \end{gathered}$ |
| 0.72 | 29.00 | 65.80 |
| 2.16 | 29.00 | 68.80 |
| 3.72 | 29.70 | 69.90 |
| 4.19 | 29.90 | 70.70 |
| 4.78 | 29.80 | 74.40 |
| 4.59 | 30.00 | 75.90 |
| 4.31 | 30.00 | 75.70 |
| 5.56 | 30.20 | 75.50 |
| 6.13 | 29.60 | 77.10 |
| 5.63 | 29.10 | 79.00 |
| 5.78 | 28.50 | 80.10 |
| 5.19 | 27.50 | 82.50 |
| 5.31 | 26.80 | 84.00 |
| 5.25 | 26.00 | 85.90 |
| 4.41 | 26.30 | 84.50 |
| 3.78 | 26.60 | 82.30 |
| 3.66 | 27.00 | 80.70 |
| 4.81 | 26.50 | 81.80 |
| 4.94 | 26.10 | 84.00 |
| 5.25 | 25.70 | 84.40 |
| 5.59 | 25.90 | 83.70 |
| 5.63 | 26.00 | 84.80 |
| 5.13 | 26.20 | 84.10 |
| 4.53 | 26.60 | 84.00 |
| 4.56 | 26.90 | 81.80 |
| 4.34 | 26.60 | 82.60 |

Table 26. ANOVA

| Source | DF | M.S.S. |
| :--- | :---: | :---: |
| Regression | 2 | $12.8714^{* *}$ |
| Error | 23 | 0.3430 |

## fig.l AGE v/e MEAN BODY WEIGHT OF QUAIL8



FIG. 2 GOMPERTZ CURVE FIT TED TO JAPANE8E QUAILS
(MALE) UBING AMERAGE WEEKLY BODY WEIGHT


## FIG. 3 QUADRATIC CURVE FITTED TO JAP.QUAIL8 (FEMALE)USING AU.WEEKLY BODY WEIGHT



## fig. 4 GOMPERTZ CURVE FITTED TO JAPANE8E QUAIL8 (FEmALE)UBING A.WEEKLY BODY WEIGHT



## FIG. 5 GOMPERTZ CURVE FITTED TO JAPANE\&E QUAIL8 USING AMERAGE WEEKLY BODY WEIGHT (IRREAPECTIVE OF 3EX)



## FIG. 6 WEEKLY EGG PRODUCTION OF JAPANEBE QUAIL8 (COTURNIX COTURNIX JAPONICA)



- number of egre

FIG. 7 FORTNIGHTLY EGG PRODUCTION OF JAPANESE QUAILS (COTURNIX COTURNIX JAPONICA)


## FIg. 8 HEN HOUSED \& HEN DAY EGG PRODN. JAP.QUALLS(COTURNIX COTURNIX JAPONICA)



- HEN HOUSED EGG PROD.
T- HEN DAY EGG PRODN


## FIG. 9 EXPECTED WEEKLY EGC PRODN.OETANED FROM QUADRATIC-CUM-LOE \& GUADRATIC FNJN LOG BCALE ALONG WITH OBSERYED EGG PRODN






## DISCUSSION

The results of the present investigations were already given in chapter 4. Most of the results obtained were having a reasonable comparison with the results obtained by other research workers in this field with few exception.

In the case of egg production in quails practically no work have been done. Hence could not have a comparative study of this aspect. The discussion of the result obtained were given in this chapter.

### 5.1 Growth study

The average hatching weight of one hundred and three (103) Japanese quails (58 males and 45 females) were found to be $7.1369 \pm 0.1698$ g. This was slightly higher than the average hatching weight (6.50 g) given by'Ricklefs (1979) but falls in the range of $5.12-8.05 \mathrm{~g}$ for Japanese quails as hatching weight given by Ricklefs (1979). The average hatching weight in the present study was also higher than the average hatching weight given by Sreenivasaiah et al. (1987a) who reported an average initial body weight as 5.74 g (Monsoon hatch - August) and 6.02 g (Winter hatch - November).

It was observed that the average hatching weight for males and females were found to be $7.1172 \pm 0.1591 \mathrm{~g}$ and $7.1622 \pm 0.1813 \mathrm{~g}$ respectively and females weighed more.

The body weight steadily increased for males, females and birds irrespective of sex (Table 1) upto l2th week. In all the cases females exceeded males with respect to body weights. The same phenomenon was observed by Ino et al. (1985) for males and females eventhough the body weights were not in agreement with the present study.

The average body weight during the 12 th weeks were found to be $157.6552 \pm 14.5199 \mathrm{~g}$ and $179.2000 \pm 19.8055 \mathrm{~g}$ for males and females respectively. This was found to be in agreement with the results obtained by Kozaczynski (1985.).

Analysis of variance (Table 2), showed that there was no significant difference in body weights between males and females upto 8th week. But from 9th week to 12 th week there was significant difference in body weights between male and female Japanese quails. Fig.l further substantiate the claim.

In order to study the trend in body weights further body weights were taken at 27 th and 35 th week for males and females.

During the 27 th week the average body weights again shown an increasing trend with females weighing more (194 g) than males (172.4310 g) and the analysis of variance have shown that there was significant difference in body weights between males and females.

At the 35 th week there was a slight decline in average body weights for females (192.4839 g) while the average body weights for males (179.4483 g) again shown an increasing trend and the analysis of variance again showed that there was significant difference in body weights between males and females. The average body weights for birds irrespective of sex were also worked out for 27 th and 35 th week and it was found to be $181.8544 \pm 23.9376 \mathrm{~g}$ and $185.1456 \pm 21.8023 \mathrm{~g}$ respectively. In general it could be concluded that the body weights of quails showed an increasing trend even upto 35 th week.

Various mathematical models were fitted and Gompertz curve has emerged as best one for ascertaining growth in quails over twelve weeks having higher ' $r$ ' values and lower 's' values. This was found to be in agreement with the curve suggested by Laird (1965) and Ricklefs (1985).

It was observed that for the development of suitable models for ascertaining growth in quails using average body weights over twelve weeks, Gompertz curve emerged as the best one followed by quadratic and logistic in the case of males and birds irrespective of sex while quadratic curve (Fig.3) emerged as best one followed by Gompertz and modified exponential in the case of females.

In general Gompertz curve was found to be most suitable for the development of models for ascertaining growth in Japanese quails having highest ' $r^{2}$ ' and ${ }_{\wedge}$ 's's' values. Fig.2, 4, 5 showing observed and expected body weights over twelve weeks further substantiate the claim.

The Gompertz form fitted to the average body weights over twelve weeks were of the form.

$$
\begin{aligned}
& W_{t}=158.6057 \exp (-4.9667 \exp (-0.4677 \mathrm{t})) \quad \text { (Male) } \\
& \mathrm{r}^{2}=0.9991, \mathrm{~s}=1.6903 \\
& \mathrm{~W}_{\mathrm{t}}=183.8152 \exp (-4.5238 \exp (-0.3846 \mathrm{t})) \quad \text { (Female) } \\
& \mathrm{r}^{2}=0.9949, \mathrm{~s}^{\prime}=4.6230 \\
& \mathrm{~W}_{\mathrm{t}}=169.1025 \exp (-4.7301 \exp (-0.4267 \mathrm{t})) \\
& \mathrm{r}^{2}=0.9982, \mathrm{~s}=2.6054^{\circ} \text { (Birds irrespective of.sex) }
\end{aligned}
$$

Rao's method have shown that the initial body weight had no significant effect on growth rate of Japanese quails.

### 5.2 Egg production study

The average age at sexual maturity for 32 femalo Japanese quails was approximately 9 to 10 weeks or $69.41 \pm$ 9.18 days. This was found to be higher than the reported age at sexual maturity by Ino et al. (1985) as 35.9 days for males and 44.8 days for females.

The average weight of egg was found to be $12.20 \pm 1.05 \mathrm{~g}$ which was also found to be higher than the reported egg weight by Sreenivasaiah et al. (1987b) and Narayanankutty et al. (1989).

With regard to egg production study scientists fitted a number of mathematical models in chicken, ducks, turkey, goose, turn, starling etc.

Still none of these scientists havc developed a suitable model for ascertaining the egg production in Japanese quails which was evident from the available literature.

In the present study a number of mathematical models tried in milk production in cattle were used to predict the egg production in quails. The comparisons were made on the value of Furnival index (I).

It was observed that for predicting weekly egg production quadratic-cum-log function and quadratic function in log scale emerged as first and second best having respective $I$ values $19.3758,19.4584$ and $r^{2}$ values 0.7602 , 0.7471 ( Table 21). It is evident that there is not much difference between respective $I$ values and $r^{2}$ values. Figure 9 further shows that there is not much variation between expected egg productions in comparison with observed egg production in the case of these two curves.

With regard to prediction of fortnightly egg production linear hyperbolic function and quadratic function in logarithmic scale emerged as first and second best having respective I valụes 34.7535 and 35.0586 and $r^{2}$ values 0.8052 and 0.8018 (Table 22). Figure 10 further shows that there is not much variation between expected egg production in comparison with observed egg production of these two curves.

It was observed that hen housed and hen day egg production were highest in the month of June (73.1073\% and 73.6064\% respectively) followed by August (Table 20).

In the case of prediction of hen housed and hen day egg production, quadratic function in logarithmic scale has emerged as best one having least $I$ values and comparatively good $r^{2}$ values (Tables 23 \& 24 for hen housed and hen day egg production respectively). Figure 11 and 12 (for hen housed and hen day egg production respectively) between observed and expected egg production. shows a good fit for quadratic function in logarithmic scale.

In general, quadratic function in logarithmic scale can be considered as suitable model for ascertaining egg production in quails with regard to weekly, fortnightly, hen housed and hen day egg production. The form of the fitted quadratic function in logarithmic scale was

$$
\begin{aligned}
& Y=13.4228+124.6064\left(\log _{e} t\right)-25.60003\left(\log _{e} t\right)^{2} \\
& I=19.4584, r^{2}=0.7471, s=19.4584(\text { weekly) } \\
& Y=97.1038+268.8874\left(\log _{e} t\right)-75.5807\left(\log _{e} t\right)^{2} \\
& I=35.0586, r^{2}=0.8018, s=35.0586(\text { Fortnightly) } \\
& Y=1.2589+96.6707\left(\log _{e} t\right)-35.3734\left(\log _{e} t\right)^{2} \\
& I=6.0220, r^{2}=0.9571, s=6.0220(\text { Hen housed) } \\
& Y=1.2386+95.4551\left(\log _{e} t\right)-33.5394\left(\log _{e} t\right)^{2} \\
& I=7.5873, r^{2}=0.9382 ; s=7.5873(\text { Hen day) }
\end{aligned}
$$

### 5.3 Climatological studies

The average weekly temperature and average weekly humidity under which the experiment was conducted were found to be $27.8 \pm 1.6^{\circ} \mathrm{C}$ and $79.4 \pm 5.7 \%$ respectively.
(i.e. $26.2^{\circ} \mathrm{C}$ to $29.4^{\circ} \mathrm{C}$ - temperature 73.7\% to $85.1 \%$ - humidity)

The multiple linear regression equation fitted by taking previous three weeks temperature $\left(Z_{1}\right)$ and previous three weeks average humidity $\left(z_{2}\right)$ and the present weekIy average egg production per bird (Table 25) showed an $r^{2}$ value of $76.5 \%$ which gives good fit. The analysis of variance of this regression equation (Table 26) showed a highly significant value for regression equation. Further analysis of the coefficients of equation showed highly significant values of
coefficients of $z_{1}$ and $z_{2}$ which indicated that the average temperature and humidity were contributing factors for the egg production in quails.

It was evident from the study of previous workers (Wilson et al., 1971; Sreenivasaiah and Joshi, $1987 \mathrm{a}, \mathrm{b}$ ) that temperature was having significant effect on production. Accordingly multiple regression equation was fitted by taking average temperature and humidity of the previous three weeks as explanatory variable and the present weekly average egg production per bird as dependent variable.

## Summary

With a view to develop suitable models for ascertaining growth and egg production in quails an experiment was initiated on first February, 1989 at the Kerala Agricultural University Poultry Farm, Mannuthy. One hundred and fifty (150) day old quails have been used in this experiment. Forty seven (47) birds died during the period of twelve weeks due to some reason or other. The data on the remaining birds consisting of 58 males and 45 females were considered for the present study.

The body weights of these birds were recorded continuously upto 12 weeks. The initial average body weights were $7.1172 \pm 0.1591 \mathrm{~g}, 7.1622 \pm 0.1813 \mathrm{~g}$ and $7.1369 \pm 0.1698 \mathrm{~g}$ for males, females and birds irrespective of sex respectively.

At the l2th week the average body weights of males and females were $157.6552 \pm 14.5199 \mathrm{~g}$ and $179.2500 \pm 19.8055 \mathrm{~g}$ respectively.

Since the body weights have shown an increasing trend (Fig.l) for males and females and females had begun egg production (females started egg production from around loth week onwards and they have to be kept undisturbed for uninterrupted egg production) body weights were not taken upto 26 th
week. Further body weights were taken only at 27 th and 35 th week to ascertain the trend by which time the males and females had reached an average body weights of
$172.43 \mathrm{~g} ; 19.4 .00 \mathrm{~g}$ ( 27 th week)
$179.45 \mathrm{~g} ; 192.00 \mathrm{~g}$ (35th week) respectively which again showed an increasing trend with a slight decline in the case of average body weights of females.

Analysis of variance was conducted for the body weights of 58 males and 45 females birds which showed that there was significant difference in body weights between males and females from 9 to 12 weeks as well as on 27 th and 35 th weeks.

Mathematical models such as linear, quadratic, exponential, Von-Bertalanffy, modified exponential, logistic and Gompertz were fitted and were compared using coefficient of determination ( $r^{2}$ ) and standard error of estimate(s) values. By comparison Gompertz curve $W_{t}=a \exp [-b \exp (-k t)]$ - was chosen as the best one for ascertaining growth in Japanese quails from 1 to 12 week. It was also concluded that Gompertz curve of the following forms were most suitable for ascertaining growth in Japanese quails using average body weights over 12 weeks. The forms were

$$
\begin{aligned}
& W_{t}=158.6057 \exp [-4.9667 \exp (-0.4677 t)] \quad(\text { Male }) \\
& W_{t}=183.8152 \exp [-4.5238 \exp (-0.3846 t)] \text { (Female) } \\
& W_{t}=169.1025 \exp [-4.7301 \exp (-0.4267 t)]
\end{aligned}
$$

(Birds irrespective of sex)

Figures of Gompertz curves showing observed and expected body weights were also drawn to support the claim.

Rao's method was tried and it was observed that initial body weight had no significant effect on growth rate of Japanese quails.

Mathematical models such as linear, exponential, parabolic exponential, inverse polynomial, Gamma function, Gamma type function, quadratic function, quadratic function in logarithmic scale, quadratic-cum-log function, emperical and linear hyperbolic functions were tried for predicting egg production. The fitted models were compared using Furnival index (I). The $r^{2}$ and $s$. values were used only as a second criteria for comparison. Before fitting, the egg production -data was converted to weekly and fortnightly basis. Hen housed and hen day egg production were also computed. The models were fitted to weekly, fortnightly, hen housed and hen day egg production data. It was concluded that quadratic function in logarithmic scale was mot suitable for ascertaining egg production in quails. The forms of the models were

$$
\begin{aligned}
& Y=13.4228+124.6064\left(\log _{e} t\right)-25.6003\left(\log _{e} t\right)^{2}(\text { weekly) } \\
& Y=97.1038+268.8874\left(\log _{e} t\right)-75.5807\left(\log _{e} t\right)^{2}(\text { fortnight }) \\
& Y=1.2589+96.6707\left(\log _{e} t\right)-35.3734\left(\log _{e} t\right)^{2} \\
& Y=1.2386+95.4551\left(\log _{e} t\right)-35.5394\left(\log _{e} t\right)^{2}(\text { hen housed }) \\
& Y=0 y)
\end{aligned}
$$

Figures of quadratic function in logarithmic scale showing observed and expected egg production were also drawn to support the claim.

Climatological parameters (temperature and humidity) recorded daily from the beginning of the experiment till the end, was utilised to study the impact of these parameters on egg production. The parameters were converted to weekly basis. As the present average weekly egg production per bird is correlated with the previous three weeks climatological parameters, a multiple linear regression equation of the form

$$
\mathrm{Y}=-41.7275+0.7687 * * \mathrm{Z}_{1}+0.3150 * * \mathrm{z}_{2} \text { with } \mathrm{r}^{2}=0.7654
$$ wàs fitted with average weekly egg production per bird ( $Y$ ) as dependent variable and weekly average temperature $\left(Z_{1}\right)$ and weekly average humidity $\left(Z_{2}\right)$ as explanatory variable. It was found that the weekly average temperature and weekly average humidity have significant effect on the average egg production.

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# DEVELOPMENT OF A SUITABLE MODEL FOR ASCERTAINING THE GROWTH AND EGG PRODUCTION IN QUAILS 

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# ABSTRACT OF A THESIS <br> Submitted in partial fulfilment of the requirement for the degree of <br> flatier of Scimice (Agritultural Statistics) 

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#### Abstract

An investigation was carried out into the growth and egg production aspect of Japanese quails at the Kerala Agricultural University Poultry Farm, Mannuthy on lst February, 1989 with the following objectives. 1. t.o fird a suitable relationship between age and body weight. 2. to investigate the trend of egg production in quails through suitable mathematical models. 3. to study the impact of climate parameters (temperature, humidity) on egg prodưction in quails.

The birds were reared under uniform feed formula and identical management practices (recommended by Kerala Agricultural University Package of Practices). The investigation mainly depended on data consisting of weekly body weights of individual birds, daily e'gg production of birds (beginning from age at sexual maturity) and daily climatological parameters (temperature and humidity) from beginning till the end of experiment of 30 th September, 1989.

Mathematical models such as linear, quadratic, exponential, Von-Bertalanffy, modified exponential, logistic and Gompertz were fitted for the purpose using body weights of


individual birds as well as average body weights over twelve weeks and the fitted models were compared using coefficient of determination ( $r^{2}$ ) and standard error of estimate(s).

Mathematical models such as linear, exponential, parabolic exponential, inverse polynomial, Gamma function, Gamma-type functicn, quadratic function, quadratic function in logarithmic scale, quadratic-cum-log, emperical and linear hyperbolic functions were fitted for the development of suitable models for ascer̈taining egg production using total weekly, fortnightly egg production, hen housed and hen day egg production and fitted models were compared using Furnival index, $r^{2}$ and $s$.

Multiple linear regression equation was fitted using average weekly egg production per bird as dependent variable and weekly temperature and humidity as explanatory variable to study the impact of climatological parameters on egg production in quails.

The investigation has the following, salient features.
(i) The hatching weight of Japanese quails were 7.1369 g .
(ii) The females weighed more than the males during the entire period of experiment and the body weights have shown an increasing trend. At the end of 12 th week the average body weights of males and females were 157.6552 g and 179.2500 g respectively.
(iii) Rao's method jus'tified that initial body weights had no significant effect on growth rate.
(iv) Gompertz curve $W_{t}=a \exp [-b \exp (-k t)]$ was most suitable for ascertaining growth in quails on individual basis as well as on the basis of average body weights over twelve weeks.
(v) Averace age at sexual maturity (females) was found to be approximately 10 weeks and on an average the eggs weighed 12.20 g .
(vi) Quadratic function in logarithmic scale $W_{t}=a+b\left(\log _{e}{ }^{t}\right)+c\left(\log _{e}{ }^{t}\right)^{2}$ was most suitable for ascertaining egg production in quails (weekly, fortnightly, he'n housed and hen day, production basis).
(vii) Climatic parameters had significant impact on egg production in quails.

