

Genetic determination of the lysozyme activity levels in guineafowl egg white and serum*

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ABSTRACT

Guineafowl lysozyme levels in egg white and serum revealed considerable quantitative variations between varieties and between sexes. Electrophoretic variants were not detected. Inheritance studies indicated significant genetic influences. Heritability values were 0.31 ± 0.27 and 0.43 ± 0.29 , respectively, for lysozyme levels in serum and egg white. Moderate associations were observed for serum lysozyme with 4-week body weight, and for egg white lysozyme with age at first egg and albumin index.

Lysozyme (EC, 3.2.1.17), a widely distributed enzyme, is directly involved in antimicrobial defence mechanisms of body. Qualitative and quantitative variations of lysozyme activity in egg white and serum and its genetic control have been studied in chicken and other avian species (Baker *et al.* 1970, Manwell and Baker 1973, Sato and Watanabe 1976, Bessarabov and Krykanov 1985).

This article presents results of some observations on genetic control of quantitative and qualitative variations in guineafowl egg white and serum.

MATERIALS AND METHODS

Guncari guineafowls (370 individuals) belonging to 25 halfsib families maintained

at this Institute were used. Lysozyme levels were assayed in fresh serum and egg white samples (Lie 1980, Lie *et al.* 1986). Qualitative variations were investigated in acrylamide-gel electrophoresis (Sato and Watanabe 1976).

Effects of genetic and nongenetic factors on lysozyme levels were investigated using the following statistical model in which the effect of sire was taken as random, and those of sex, feathering phenotypes and variety were taken as fixed effects.

$$Y_{ijkl} = \mu + M_i + S_j + F_k + P_l + e_{ijkl}$$

Where,

Y_{ijkl} is the value of n th individual in $ijkl$ th cell,

μ is the overall mean of the population,

M_i is the effect of i th sire ($i = 1, 2, \dots, n$) (chosen at random),

S_j is the effect of the j th sex ($j = 1, 2$),

F_k is the effect of k th feathering phenotypes ($k = 1, 2$),

P_l is the effect of l th variety ($l = 1, \dots, 3$), and

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e_{ijkl} is the environmental deviations attributed to individuals assumed to be normally distributed with mean 0 and variance σ^2 .

RESULTS AND DISCUSSION

Existence of wide individual variations for lysozyme levels in egg white and other body fluids is well established among poultry species (Feeney and Allison 1969, Manwell and Baker 1973). In guineafowl serum lysozyme level at 4 weeks of age was 1.01 and 0.93 $\mu\text{g/ml}$, respectively, in male and female keets. A gradual increase at subsequent ages was seen in females. Significantly high lysozyme levels attained at maturity indicated higher cellular destruction taking place during the onset of laying (Fig. 1). Similar values were reported for chicken by Sato and Watanabe (1976). Differences between guineafowl varieties were significant, but effects of slow and rapid feathering phenotypes on lysozyme levels were not significant (Table 1). Population distribution pattern for lysozyme levels (8 weeks) revealed distinct

Table 1. Serum and egg white lysozyme least-square means for different varieties and feathering groups

	Lysozyme level	
	Serum ($\mu\text{g/ml}$)	Egg white (mg/ml)
<i>Varieties</i>		
White	1.14 \pm 0.07 ^a (68)	1.85 \pm 0.11 (40)
Lavender	1.26 \pm 0.06 ^b (88)	1.79 \pm 0.02 (36)
Pearl	1.18 \pm 0.05 ^c (114)	1.82 \pm 0.06 (114)
<i>Feathering type</i>		
Slow	1.15 \pm 0.05 (121)	1.78 \pm 0.07 (74)
Rapid	1.26 \pm 0.04 (149)	1.86 \pm 0.07 (116)
Overall	1.19 \pm 0.04 (270)	1.82 \pm 0.06 (190)

Figure in parenthesis is number of observations.

bimodality (Fig. 2).

Mean egg white lysozyme levels are comparable to the earlier reported values for guineafowl by Valdimirova (1974) and Sajjnar (1993). But these estimates are relatively much lower than the values reported by Feeney and Allison (1969) and

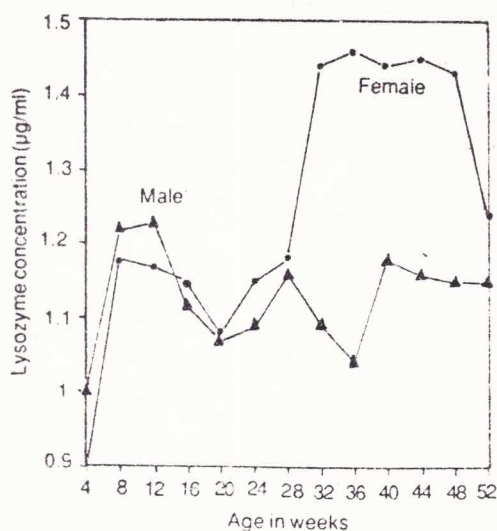


Fig. 1. Serum lysozyme levels in relation to age.

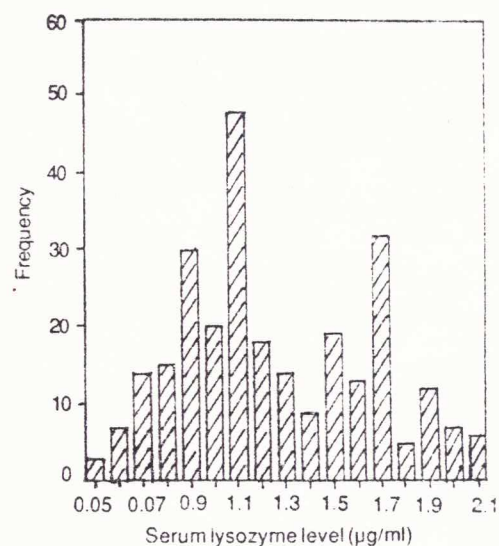


Fig. 2. Serum lysozyme distribution of GUNCARI population at 8 weeks.

Arora *et al.* (1974) for chicken. During electrophoresis, guineafowl egg white lysozyme resolved as a single discrete band between the quail and chicken lysozyme components (Fig. 3), but no variations were identified in this species.

Guineafowl serum and egg white lysozyme heritability estimates 0.311 ± 0.27 and 0.428 ± 0.294 , respectively, are comparable to moderate heritability values reported for chicken (Bessarabov and Krykanov 1985). Inheritance of lysozyme levels among the progeny from matings between parents with high and low lysozyme levels indicated a dominance trend for high type over low type (Table 2). These observations imply existence of some major gene influences besides the limited additive gene control. Similar genetic control was reported for serum alkaline phosphatase (Tamaki 1975), and lysozyme levels (Lie *et al.* 1980). Low positive phenotypic and genetic correlations were obtained between serum lysozyme levels and body weights at hatch and 4 weeks of age (0.08 , 0.02 ; 0.049 ± 0.368 , 0.329 ± 0.459 respectively). Egg white lysozyme correlations with albumin index (0.59 , 0.727 ± 0.0214) were positive but negative with age at first egg (-0.014 , -0.683 ± 0.278). In spite of the high standard errors obtained in a few cases, these associations indicated the

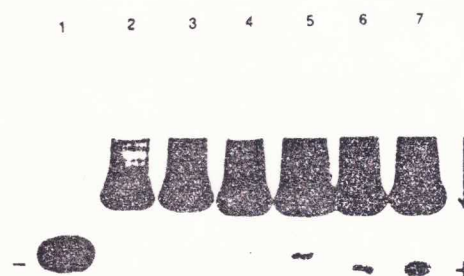


Fig. 3. Zymogram of egg white lysozyme 1, lysozyme standard, 2-3, guineafowl, 4-5 chicken, 6-7, quail.

possible trends and also suggested caution and further varification.

Lysozyme defined as 1,4- β -acetyl muramidase lyses bacteria by splitting the β (1-4) linkage between N-acetylmuramic acid (NAG) N-acetyl-glucosamine component of its cell-wall. Released from blood platelets, lysozyme is an important nonspecific bacteriolytic agent in blood and other biological fluids. The sires arbitrarily classified as high, medium and low lysozyme level types revealed, respectively, 1.4, 4.5 and 6.2% mortality among their progeny. This negative association between lysozyme level of sire and progeny mortality pattern, positive correlations for body weights, and the negative correlation with age at first egg indicated a better health status for birds with higher lysozyme levels. These observations are in agreement

Table 2. Results for matings between parents with different serum lysozyme level (high-1.75, medium-1.4—1.2, low-1.0 $\mu\text{gm/ml}$)

Sire	Dam	N	Offspring (%)			Overall progeny average
			High	Medium	Low	
High	High	18	44.4	33.3	22.3	1.85
High	Low	23	21.7	43.4	34.9	1.31
Low	Low	28	10.7	32.1	57.2	0.95

with the report of a positive correlation between dam's lysozyme level and nonspecific general resistance of progeny (Bassarabov and Krykanov 1985). Efficient functioning of this highly evolved nonspecific defence system in egg white also seems crucial for resistance to microbial invasions of embryos (Sajjanar 1993). Observations of very low lysozyme levels in egg white of penguins and related birds as compared to high lysozyme levels observed among other avian species from relatively dirty habitats suggest existence of an evolutionary association with level of environmental pollution (Manwell and Baker 1972).

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