

## Age-Dependent Changes in Activity of Mallard Plasma Cholinesterases

Authors: Bennett, Richard S., and Bennett, Jewel K.

Source: Journal of Wildlife Diseases, 27(1): 116-118

Published By: Wildlife Disease Association

URL: https://doi.org/10.7589/0090-3558-27.1.116

BioOne Complete (complete.BioOne.org) is a full-text database of 200 subscribed and open-access titles in the biological, ecological, and environmental sciences published by nonprofit societies, associations, museums, institutions, and presses.

Your use of this PDF, the BioOne Complete website, and all posted and associated content indicates your acceptance of BioOne's Terms of Use, available at <a href="https://www.bioone.org/terms-of-use">www.bioone.org/terms-of-use</a>.

Usage of BioOne Complete content is strictly limited to personal, educational, and non - commercial use. Commercial inquiries or rights and permissions requests should be directed to the individual publisher as copyright holder.

BioOne sees sustainable scholarly publishing as an inherently collaborative enterprise connecting authors, nonprofit publishers, academic institutions, research libraries, and research funders in the common goal of maximizing access to critical research.

## Age-Dependent Changes in Activity of Mallard Plasma Cholinesterases

**Richard S. Bennett¹ and Jewel K. Bennett,²**¹ USEPA Environmental Research Laboratory, Corvallis, Oregon 97333, USA; ² NSI Technology Services Corporation, USEPA Environmental Research Laboratory, Corvallis, Oregon 97333, USA

ABSTRACT: Plasma acetylcholinesterase (AChE) and butyrylcholinesterase (BChE) activity was measured repeatedly in 27 mallard (Anas platyrhynchos) ducklings between 7 and 85 days of age to determine age-dependent changes in enzyme activity. Plasma AChE, BChE, and total cholinesterase (ChE) activity decreased significantly with age. The relative proportion of AChE in total ChE activity also decreased slightly with age. Since some anti-ChE chemicals can selectively inhibit AChE or BChE activity, characterization of age-dependent changes in the activity of each enzyme may be necessary to accurately identify the occurrence of pesticide exposure.

Key words: mallard, Anas platyrhynchos, cholinesterase, age-dependent changes, repeated measures.

Inhibition of brain and plasma cholinesterase (ChE) activity is commonly used for diagnosing wildlife exposure to organophosphorus and carbamate insecticides. The ChE activity of animals with suspected exposure to these insecticides can be compared to the activity of normal control animals of the same species to determine relative inhibition. Animal age also can be a critical factor for making valid comparisons. Brain ChE activity of European starlings (Sturnus vulgaris) increases several-fold between hatching and attainment of adult size (Grue and Hunter, 1984; Robinson et al., 1988), making it necessary to compare juveniles of the same age to determine appropriately the occurrence and degree of ChE inhibition. Agedependent increases in brain ChE activity have also been observed in nestling great egrets (Casmerodius albus), snowy egrets (Egretta thula), and black-crowned nightherons (Nycticorax nycticorax) (Custer and Ohlendorf, 1989). Precocial and semi-precocial species that have been studied differ from altricial species in the developmental pattern of brain ChE activity. Hoffman and Eastin (1981) found that brain ChE activity in mallards (*Anas platyrhynchos*) increases to adult levels during embryonic development. In other precocial and semi-precocial species, young and adults also have similar brain ChE activity levels (Hudson et al., 1972; Ludke et al., 1975; White et al., 1979; Grue and Hunter, 1984).

Plasma ChE activity, on the other hand, does vary with age in some precocial birds. Fairbrother et al. (1990) reported that plasma total ChE activity in normal mallards decreased by 43% between 5 and 58 days of age. Lyles et al. (1980) reported that plasma ChE activity decreased in developing chickens (Gallus domesticus), with a rapid decrease in acetylcholinesterase (AChE) activity after hatch, while butyrylcholinesterase (BChE) activity was relatively constant. These findings have been attributed to differences in the source of each enzyme. High levels of AChE at hatching appear to be associated with release of AChE from maturing muscle fibers, while BChE is synthesized and secreted by the liver (Lyles et al., 1980).

The present study was conducted to document age-dependent changes in AChE and BChE in growing mallards. Since some anti-ChE chemicals can selectively inhibit AChE or BChE activity (Aldridge, 1953; Silver, 1974), characterization of age-dependent changes in the activity of each enzyme and their relative contribution to total ChE activity may be necessary to accurately identify the occurrence of pesticide exposure.

Day-old mallard ducklings, purchased from Whistling Wings, Inc. (Hanover, Illinois 61041, USA), were housed in brooders maintained at 39 C and provided with Purina® (St. Louis, Missouri 63166, USA) gamebird starter and water ad libitum.

_		Enzyme activity		_
Age (days)	<b>AChE</b>	BChE	Total ChE	Percent AChE

TABLE 1. Mean (±SE) and range for plasma acetylcholinesterase (AChE), butyrylcholinesterase (BChE),

	Enzyme activity			
Age (days)	AChE	BChE	Total ChE	Percent AChE <sup>b</sup>
7	$302 \pm 14$ $161,440$	$1,045 \pm 55^{\circ}$ $645, 1,524$	$1,343 \pm 64^{\circ}$ 913, 1,885	22.4° 15, 37
14	$276 \pm 16$ $120, 456$	$1,014 \pm 51$ $495, 1,407$	$1,290 \pm 61$ $616, 1,711$	21.8 15, 35
29	$286 \pm 17$ $172,538$	$938 \pm 24$ $706, 1,154$	$1,224 \pm 28$ 878, 1,445	21.4 15, 39
50	208 ± 13 90, 361	$784 \pm 35$ $465, 1,276$	$992 \pm 40$ 556, 1,547	20.3 14, 36
85	$184 \pm 12^{\circ}$ $109, 338$	$744 \pm 32^{\circ}$ 489, 1,071	$923 \pm 33$ $680, 1,244$	18.4° 12, 37

<sup>\*</sup> Enzyme activity is expressed as micromoles of acetylthiocholine hydrolyzed/min/l plasma.

Brooder temperatures were reduced to 35 C at 7 days of age. Ducklings were moved to outdoor ground pens  $(2.5 \times 5.0 \text{ m})$  at 21 days of age. Blood samples (2 ml) were collected from each of 27 ducklings at 7. 14, 29, 50, and 85 days of age by jugular venipuncture. The blood was collected in blood collection tubes (Vacutainer®, Bectin Dickson, Rutherford, New Jersey 07070, USA) containing sodium heparin and kept on wet ice until centrifugation. Each sample was centrifuged for 10 min at 2 to 5 C at 3,000 × g. Plasma was collected in plastic vials (Sardstedt®, Princeton, New Jersey 08540, USA) and stored at -75 C until analysis. After the last collection period, all samples were analyzed at the same time in random order. Methods for conducting ChE assays are described in Fairbrother et al. (1990). AChE activity was determined by incubating samples in the presence of  $1 \times 10^{-5}$  M iso-OMPA (tetraisopropylpyrophosphoramide, Sigma Chemical Co., St. Louis, Missouri 63178, USA) (Aldridge, 1953). BChE was calculated as the remainder of total ChE minus AChE.

The mean activity for AChE, BChE, and total ChE decreased with increasing age in mallard ducklings (Table 1). The temporal changes in ChE activity were examined by a repeated measures analysis of variance. Significant  $(P \le 0.0001)$  linear trends over time were observed for each measurement. There was no evidence (P > 0.08) of curvilinear trends. A test for serial correlation using the Durbin-Watson statistic (Durbin and Watson, 1971) did not detect serial correlation in total ChE and BChE data, but did for AChE. Linear regression of enzyme activity on age was used to describe the linear trends as follows: 1) AChE =  $(308.8 \pm 11.0, SE) - (1.6)$  $\pm$  0.2) Days,  $R^2 = 0.24$ ; 2) BChE = (1056.8  $\pm 30.6$ ) -  $(4.1 \pm 0.7)$  Days,  $R^2 = 0.23$ , and 3) total ChE =  $(1364.3 \pm 35.3) - (5.7)$  $\pm$  0.8) Days,  $R^2 = 0.30$ . The percent of AChE in total ChE activity decreased slightly with age (Table 1). However, percent of AChE varied much more among individuals than within individuals, as observed in the ranges for percent AChE (Table 1). The mean percent AChE for each bird ranged from 15 to 37%. Variation between individuals in the relative proportions of AChE and BChE activity may help explain variability in individual sensitivity to pesticide effects and illustrates the advantage of using repeated measurements of ChE activity as a means of separating possible pesticide effects from individual variability (Fairbrother et al., 1989).

Because age-dependent changes in plasma AChE and BChE activity exist, similar

<sup>&</sup>lt;sup>b</sup> Median value and range for percent AChE activity of total ChE.

<sup>6</sup> Means based on sample size of 26 ducklings.

age birds are necessary when evaluating exposure of young mallards to ChE inhibitors. It is not known if the relationships observed in mallards and chickens apply to other precocial or altricial young, but given the amount of change observed in plasma ChE activity during growth, age should be a critical factor when evaluating exposure to ChE inhibitors. More research is needed to determine how changes in brain and plasma ChE activity during growth relate to age-related differences in pesticide sensitivity.

We thank Susan Schiller for assistance with ChE analyses, Lisa Ganio for statistical advice, and Anne Fairbrother, Brad Marden, and Mark Meyers for review of the manuscript. Mention of trade names or commercial products does not constitute endorsement or recommendation for use.

## LITERATURE CITED

- ALDRIDGE, W. N. 1953. The differentiation of true and pseudo cholinesterase by organophosphorus compounds. Biochemical Journal 53: 62-67.
- CUSTER, T. W., AND H. M. OHLENDORF. 1989. Brain cholinesterase activity of nestling great egrets, snowy egrets and black-crowned night-herons. Journal of Wildlife Diseases 25: 359-363.
- DURBIN, J., AND G. S. WATSON. 1971. Testing for serial correlation in least squares regression III. Biometrica 58: 1-19.
- FAIRBROTHER, A., R. S. BENNETT, AND J. K. BENNETT. 1989. Sequential sampling of plasma cholinesterase in mallards (Anas platyrhynchos) as an indicator of exposure to cholinesterase inhibitors. Environmental Toxicology and Chemistry 8: 117-122.
- , M. A. Craig, K. Walker, and D. O'Lough-

- LIN. 1990. Changes in mallard (Anas platy-rhynchos) serum chemistry due to age, sex, and reproductive condition. Journal of Wildlife Diseases 26: 67–77.
- GRUE, C. E., AND C. C. HUNTER. 1984. Brain cholinesterase activity in fledgling starlings: Implications for monitoring exposure of songbirds to ChE inhibitors. Bulletin of Environmental Contamination and Toxicology 32: 282-289.
- HOFFMAN, D. J., AND W. C. EASTIN, JR. 1981. Effects of malathion, diazinon, and parathion on mallard embryo development and cholinesterase activity. Environmental Research 26: 472-485.
- HUDSON, R. H., R. K. TUCKER, AND M. A. HAEGELE. 1972. Effect of age on sensitivity: Acute oral toxicity of 14 pesticides to mallard ducks of several ages. Toxicology and Applied Pharmacology 22: 556-561.
- LUDKE, J. L., E. F. HILL, AND M. P. DIETER. 1975. Cholinesterase (ChE) response and related mortality among birds fed ChE inhibitors. Archives of Environmental Contamination and Toxicology 3: 1-21.
- Lyles, J. M., E. A. Barnard, and I. Silman. 1980. Changes in the levels and forms of cholinesterases in blood plasma of normal and dystrophic chickens. Journal of Neurochemistry 34: 978–987.
- ROBINSON, S. C., R. J. KENDALL, R. ROBINSON, C. J. DRIVER, AND T. E. LACHER, JR. 1988. Effects of agricultural spraying of methyl parathion on cholinesterase activity and reproductive success in wild starlings (Sturnus vulgaris). Environmental Toxicology and Chemistry 7: 343–349.
- SILVER, A. 1974. The biology of cholinesterases. Frontiers of Biology, Vol. 36, A. Neuberger and E. L. Tatum (eds.). North Holland Publishing Company, New York, New York, pp. 449–488.
- WHITE, D. H., K. A. KING, C. A. MITCHELL, E. F. HILL, AND T. G. LAMONT. 1979. Parathion causes secondary poisoning in a laughing gull breeding colony. Bulletin of Environmental Contamination and Toxicology 23: 281–284.

Received for publication 22 December 1989.