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The Membrane-Bound Intestinal Enzymes of Waxwings and Thrushes: Adaptive and Functional Implications of Patterns of Enzyme Activity

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ABSTRACT
Cedar waxwings (Bombycilla cedrorum) feed predominantly on fruits that are rich in simple sugars and low in nitrogen, supplementing this diet with arthropod prey during the summer months as well as flowers and tree sap in springtime. In contrast, thrushes feed extensively on fatty, protein-rich invertebrate prey, supplemented with sugary and lipid-rich fruits. Simple sugars and fats are digested and/or absorbed by distinctly different physiological mechanisms, which suggests the possibility of contrasting digestive strategies in animals specialized to diets containing one of these two energy sources. In this study, we quantified enzymatic activity of three membrane-bound intestinal enzymes of cedar waxwings and five species of thrushes to explore this aspect of their digestive physiology. These enzymes catalyze the final steps in the digestion of carbohydrates (sucrase-isomaltase and maltase-glucoamylase) and protein (aminopeptidase-N). The two carbohydrases are homologous enzymes with overlapping functions; both enzymes catalyze the hydrolysis of maltase and isomaltase. The membrane-bound digestive enzyme systems that we described for cedar waxwings and thrushes can be explained by the particular nutrients contained within their respective natural diets. Consistent with previous work, cedar waxwings displayed intestinal sucrase activity, whereas thrushes did not. Correspondingly, cedar waxwings ate some foods containing sucrose, whereas thrushes do not. Sucrase-isomaltase conferred all maltase and isomaltase activity in cedar waxwings. In contrast, all maltase and isomaltase activity in thrushes was necessarily sucrase independent, which indicated the presence of maltase-glucoamylase. The absence of sucrase-independent maltase activity in cedar waxwings suggests that sucrase-isomaltase obviates the need for maltase-glucoamylase. Indeed, total maltase and isomaltase activities were much higher in cedar waxwings than in thrushes. Neither waxwings nor thrushes eat starchy foods; sucrase-isomaltase in waxwings and maltase-glucoamylase in thrushes probably function in digesting glycogen in animal foods. We suggest that digestive traits associated with specialization to monosaccharide-rich fruits (lack of a grinding gizzard) by frugivorous waxwings and thrushes may prevent utilization of starchy seeds. Total aminopeptidase-N activity in cedar waxwings was indistinguishable from the allometric pattern among thrush species, but the distribution of this enzyme along the intestines of waxwings and thrushes was distinctly different, which demonstrates that total enzyme activity can be insufficient as a descriptor of the functional activity of brush border enzymes. Aminopeptidase-N activity peaked in the anterior part of the intestines of thrushes and in the terminal portion of the intestines of waxwings, which suggests contrasting strategies for protein digestion from fatty versus sugary diets, respectively.

Introduction
Cedar waxwings (Bombycilla cedrorum) and thrushes show contrasting digestive strategies associated with specialization to sugary fruits and fatty, protein-rich invertebrates, respectively (Witmer 1996, 1998; Witmer and Van Soest 1998). Sugary fruits eaten by waxwings contain the simple sugars glucose and fructose, nutrients that are readily absorbed by active and passive mechanisms without enzymatic breakdown (Pappenheimer and Reiss 1987). During a brief period in the spring, cedar waxwings also regularly eat some foods that contain sucrose (plant sap, buds, flowers, scale insects; Witmer 1996). Thrushes, in contrast, consume predominantly arthropod prey, supplemented with both sugary (glucose and fructose) and fatty fruits throughout the year (Martin et al. 1951; Wheelwright 1986; Witmer 1996). Arthropods are typically rich in protein, fat, and glycogen (Redford and Dorea 1984; Bell 1990).

Carbohydrates and proteins are required on a daily basis by birds (Robbins 1993). Complex carbohydrates and proteins are broken down by a sequence of enzymatic reactions into their
Aminopeptidase-N has been reported in the cedar waxwing (McWilliams et al. 1999), but there are no comparable data for thrushes. If total activity of an intestinal enzyme is directly related to the quantity of its substrate in an animal’s natural diet, the relatively low protein content of the diet of the cedar waxwing (Witmer 1996) would predict relatively low aminopeptidase-N activity for this species.

We compared total hydrolytic activity of sucrase-isomaltase, maltase-glucoamylase, and aminopeptidase-N, as well as the distribution of enzyme activities throughout the intestines of cedar waxwings and five species of thrushes. We undertook this work as an exploratory, descriptive study of this facet of the digestive physiology of two groups of fruit-eating birds that differ in the nutrient composition of their natural diets. By comparing cedar waxwings to thrushes varying in body size, we were able to make quantitative comparisons that controlled for effects due to body size. We depict enzyme activity as a function of body size, comparing values for cedar waxwings to the regression that describes the allometric relationship among thrush species. Because phylogenetic relatedness confounds the use of species as units of replication, we also compared enzyme activity in cedar waxwings to other members of the Muscicapa- poidea, the monophyletic clade that includes both cedar waxwings and thrushes, using phylogenetically independent contrasts (Felsenstein 1985). We make inferences about the adaptive and functional significance of patterns of intestinal enzyme activity in waxwings and thrushes.

Material and Methods

We compared the activities of three membrane-bound enzymes (sucrase-isomaltase, maltase-glucoamylase, and aminopeptidase-N) in cedar waxwings (n = 4) to those in five species of thrushes (American robin, Turdus migratorius, n = 3; wood thrush, Hylocichla mustelina, n = 3; hermit thrush, Catharus guttatus, n = 1; gray-cheeked thrush, Catharus minimus, n = 1; and Swainson’s thrush, Catharus ustulatus, n = 1). Birds were captured near Ithaca, New York, and maintained in captivity for more than 2 yr before this study. The maintenance diet was moistened Eukanuba Small Bites puppy chow (Iams, Dayton, Ohio) supplemented with a variety of natural fruits. Bird diets were supplemented with crab apple fruits (Malus spp.) for the 4 d before tissue collection. We maintained birds under similar conditions of captivity and diet for an extended period to control for potentially confounding environmental influences on the activity of digestive enzymes in our comparative study. The quantitative expression of digestive enzymes can be modulated by dietary nutrient content (Afik et al. 1995; Martínez del Rio et al. 1995; Karasov and Hume 1997; Sabat et al. 1998; Levey et al. 1999).

For collection of gut tissues, birds were fasted overnight, killed with an isofluorane overdose, and their intestines were immediately excised and placed in ice-cold 1.02% saline. At
least three, but up to 11, samples of intestinal tissue approximately 1 cm in length were taken throughout the intestine, slit longitudinally to open the gut tube. For two individuals (cedar waxwing and wood thrush), the entire intestine was sampled. Position of each section (distance from pylorus) and the length and width of each intestinal segment were recorded to calculate intestinal surface area (nominal area). Tissues were rinsed with clean ice-cold saline, blotted, weighed, and stored in liquid N₂.

Tissue samples were prepared for enzyme analysis by thawing at 4°C and homogenizing in 350 mM mannitol in 1 mM Hepes/KOH, pH 7.5, for 30 s in an OMNI 5000 homogenizer at setting 6. Disaccharidase activities were measured by the rate at which stock solutions of each sugar substrate (sucrose, maltose, isomaltose) incubated with tissue homogenates yielded glucose. We refer to rates of maltose and isomaltose hydrolysis as maltase and isomaltase activity, respectively, regardless of the enzyme(s) responsible for catalytic activity (sucrase-isomaltase and maltase-glucoamylase). Glucose concentrations of incubated solutions were assayed colorimetrically using Glucose-Trinder 500 reagent (Sigma Chemical). These methods are modified from Dahlqvist (1984), as described in Martínez del Río (1990) and Martínez del Río et al. (1995). Tissue homogenates (100 µL) were incubated at 40°C (Prinzinger et al. 1991) with 100 µL of 56 mM sugar solutions in 0.1 M maleate/NaOH buffer, pH 6.5, for 10 min. Reactions were then arrested by adding 3 mL of the Glucose-Trinder stop/develop reagent, and absorbance was measured at 505 nm.

We used l-alanine-p-nitroanilide as a substrate for aminopeptidase-N. Hydrolysis of the peptide bond yields alanine and p-nitroaniline; the concentration of the latter reaction product was determined colorimetrically at 384 nm. We added 10 µL of homogenate to 1 mL of assay mix (2.0 mM l-alanine-p-nitroanilide in 0.2 M NaH₂PO₄/Na₂HPO₄ buffer, pH 7.0), incubated this solution at 40°C for 10 min, and arrested the reaction with 3 mL of ice-cold 2 N acetic acid. See Martínez del Río (1990) and Martínez del Río et al. (1995) for justification of the use of intestine homogenates and reaction rates relative to Vₘₐₓ under these assay conditions. We measured the pH optima for each enzyme to validate our assay protocol (M. C. Witmer and C. Martínez del Río, unpublished data); pH optima for these enzymes in cedar waxwings are reported in McWilliams et al. (1999).

Enzyme activities were standardized to nominal area of the intestine (Martínez del Río 1990; Martínez del Río et al. 1995). We statistically evaluated enzyme activity as a function of intestinal position by repeated-measures ANOVA, with bird species as a factor. For this analysis, we grouped the three Catharus spp. thrushes together, comparing among genera. Total (summed) enzyme activity for each bird was calculated by adding the activities of assayed gut sections and extrapolating activities over segments that were not included in enzyme assays. We assumed linear changes in enzyme activity between measured segments, approximating the pattern that we found for two individuals for whom we examined the entire intestine (cedar waxwing and wood thrush; Figs. 1, 2). We compared total maltase activity and aminopeptidase-N activity in cedar waxwings to the allometric patterns that we report among the five thrush species. We computed phylogenetically independent contrasts with the Macintosh program of Purvis and Rambaut.
Digestive Enzymes of Waxwings and Thrushes

To evaluate whether enzyme activity in cedar waxwings differs from the allometric pattern among other members of the Muscicapoidea, including data for European starlings (*Sturnus vulgaris*) and gray catbirds (*Dumetella carolinensis*) from Martínez del Río et al. (1995). The phylogeny and branch lengths used to structure and standardize our contrasts, respectively (Garland and Adolph 1994), were estimated from DNA hybridization (Sibley and Ahlquist 1990). For cedar waxwings, the only species that showed significant sucrase activity, we evaluated maltase and isomaltase hydrolysis capacity as a function of sucrase activity of individual intestinal segments by regressing hydrolysis rates of these two disaccharides against sucrase activity (Martínez del Río 1990). We used ANCOVA, with individual bird as a factor, to evaluate how maltase and isomaltase hydrolysis changed as a function of sucrase activity.

**Results**

***Patterns of Hydrolase Activity along the Intestine***

Enzyme activities for the single individuals of each of the three *Catharus* spp. thrushes were very similar. Variation among these individuals was no greater than interindividual variation within the other three bird species that we examined (Figs. 1–3). Thus, we suspect that the *Catharus* spp. thrushes are similar in intestinal enzyme activity and that our data for these three individuals are reasonable approximations of the mean values of these measurements for these species under our experimental conditions.

Hydrolysis rates of carbohydrates (illustrated by maltase activity) in both cedar waxwings and thrushes peaked in the anterior intestine and declined toward zero activity in the posterior intestine (repeated-measures ANOVA, using data for the three anterior tissue samples: $F_{3,18} = 11.6, P < 0.001$; Fig. 1). This is a typical pattern for intestinal membrane-bound enzymes (Alpers 1987; Biviano et al. 1993; Martínez del Río et al. 1995). Cedar waxwings showed maltase activity per unit nominal area of intestine approximately four times higher compared with the five thrush species (repeated-measures ANOVA: $F_{5,18} = 13.9, P = 0.001$; Fisher’s protected least significant difference between *Bombycilla* and each of the thrush genera: $P < 0.001$; Fig. 1).

The pattern of aminopeptidase-N activity along the intestine was strikingly different between cedar waxwings and thrushes (repeated-measures ANOVA, genus × gut position interaction: $F_{5,18} = 8.2, P < 0.001$; Fig. 2). For thrushes, aminopeptidase-N activity tended to decline distally. American robins showed a slightly different pattern than the other thrush species, with the highest aminopeptidase-N activity in the terminal portion of the small intestine. In contrast to the general pattern for all thrush species, aminopeptidase-N activity was highest in the terminal portion of the intestine of cedar waxwings, with relatively high enzyme activity in the distal ileum and proximal rectum. Our data are consistent with McWilliams et al.’s (1999) report that cedar waxwings showed comparable levels of aminopeptidase-N activity in the posterior and anterior halves of the intestine.
Comparisons of Total Enzyme Activity

Cedar waxwings showed sucrase, maltase, and isomaltase activity, whereas thrushes showed very low levels of sucrase and isomaltase activity (Table 1). Total maltase activity in cedar waxwings was higher than the allometric relationship among thrush species (approximately fourfold), whereas total aminopeptidase-N activity in cedar waxwings was similar to the allometric pattern among thrush species (Table 1; Fig. 3). The independent contrasts of total enzyme activity against body mass (Fig. 4) corroborated our preceding results for the traditional allometric comparisons. Maltase activity for the Bombycillidae versus Muscicapidae/Sturnidae contrast was significantly different from the other Muscicapoidea contrasts (the maltase contrast involving the Bombycillidae was negative because, by the convention of the program, the values for the clade with the smaller value for the independent variable [body mass] are subtracted from the corresponding values for the clade with the larger value for the independent variable). The contrast of aminopeptidase-N activity against body mass for cedar waxwings was indistinguishable from the pattern among other Muscicapoidea standardized contrasts.

Because thrushes lacked sucrase-isomaltase, maltose and isomaltose hydrolysis was necessarily sucrase independent, which indicates the presence of maltase-glucosamylase (Table 1). For cedar waxwings, we estimated sucrase-independent maltase/iso- maltase activity from the y-intercept of the regression of maltase/isomaltase activity on sucrase activity (Martínez del Rio 1990). The relationship between both maltase and isomaltase activity and sucrase activity of individual intestinal segments did not differ among individual waxwings (ANCOVA, bird x enzyme activity interaction: maltase $F_{1,16} = 0.85$, $P = 0.49$; isomaltase $F_{1,16} = 1.39$, $P = 0.28$) and showed a tight positive relationship (ANCOVA: maltase $F_{1,16} = 391.2$, $P < 0.001$; isomaltase $F_{1,16} = 675.5$, $P < 0.001$; Fig. 5). The y-intercepts for both regressions...
were not significantly different from zero (Fig. 5), which suggests that maltase and isomaltase activities in cedar waxwings were due solely to the presence of sucrase-isomaltase.

Discussion

Our results corroborate previous work showing intestinal sucrase activity in cedar waxwings (Martínez del Río et al. 1989; McWilliams et al. 1999) but not in thrushes (Karasov and Levey 1990; Martínez del Río 1990). The presence of sucrase-isomaltase in cedar waxwings is probably associated with this species’ regular seasonal consumption of foods containing sucrose, such as flowers, buds, sap, and scale insects (Wittmer 1996 and references therein). Rufous-tailed plant cutters (Phytotoma rara), which ingest young leaves and shoots in addition to fruit, also exhibit high levels of sucrase-isomaltase (Meynard et al. 1999; López-Calleja and Bozinovic 2000). Unlike thrushes, cedar waxwings did not show sucrase-independent amylase activity that would indicate the presence of maltase-glucoamylase (Martínez del Río 1990). This result conflicts with McWilliams et al.’s (1999) report that 29%–39% of maltase activity in cedar waxwings was sucrase independent, a discrepancy that calls for further study. Consistent with our result, the absence of sucrase-independent maltase activity has been reported for two other passerines, the frugivorous yellow-rumped warbler (Dendroica coronata; Afik et al. 1995) and the frugivorous/folivorous rufous-tailed plant cutter (Meynard et al. 1999). The presence of sucrase-isomaltase confers relatively high maltase and isomaltase activities in waxwings, which presumably eliminates the need for maltase-glucoamylase. Relatively high amylase activity in cedar waxwings, instead of representing a functional trait, may simply be a consequence of conserved functional similarity between sucrase-isomaltase and maltase-glucoamylase. These homologous enzymes share identical catalytic sites and are members of the same enzyme family (Nichols et al. 1998).

Thrushes showed only trace levels of sucrose activity, and these birds do not consume sucrose-rich foods. The traces of sucrose activity may be the result of minor microbial contamination of intestinal samples. In contrast to cedar waxwings, thrushes showed sucrase-independent maltase and isomaltase activities, which indicates the presence of maltase-glucoamylase. Despite possessing intestinal enzymes that catalyze the final products of starch digestion in both cedar waxwings (sucrase-isomaltase) and thrushes (maltase-glucoamylase), these frugivores do not eat starchy seeds. Intestinal amylase activity in these birds probably functions in digesting glycogen in the animal foods that they eat. We suggest that other features of their digestive systems associated with specialization to frugivory and seed dispersal may preclude the utilization of starchy seeds, like the absence of the powerful seed-grinding gizzard that is found in granivores (McLelland 1975).

Our results suggest that protein digestion is fundamentally different in cedar waxwings and thrushes. This may be a consequence of different strategies associated with digestion of protein from lipid- versus monosaccharide-rich diets. The distribution of aminopeptidase-N activity along the intestines of thrushes suggests that protein digestion takes place principally in the duodenum, where emulsification and digestion of fats typically occurs (Place 1992; Place and Stiles 1992). Relatively high aminopeptidase-N activity in the hindgut of cedar waxwings suggests that the final stages of protein digestion occur in the terminal portion of the intestine. A similar pattern of aminopeptidase-N activity along the intestine has been reported in the folivorous rufous-tailed plant cutter (Meynard et al. 1999), which suggests a common trait for birds specialized to sugary, low-protein foods. For sugary fruits that can be rapidly processed through the gut because simple sugars are readily absorbed, the distribution of aminopeptidase-N in cedar waxwings may maximize exposure of dietary protein to acid and enzymatic hydrolysis before the final stages of digestion and absorption. Meynard et al. (1999) proposed a similar hypothesis for rufous-tailed plant cutters. This trait may also enhance the ability of birds specialized to foods with a low nitrogen-to-carbohydrate ratio to recover precious nitrogen from digestive secretions and excretory products. The lower intestine of birds is an important site of water and electrolyte absorption (Skadhauge 1982; Thomas 1982; Anderson and Braun 1985), and for cedar waxwings, this terminal region appears to be important in nutrient digestion and absorption as well. In the distal ileum and proximal rectum of cedar waxwings, the villi become papillate (M. C. Wittmer and D. McClearn, unpublished data).
Figure 4. Phylogenetically independent contrasts of enzyme activity against body mass for cedar waxwings, four thrush species, gray catbirds (maltase only), and European starlings, closely related frugivores in the superfamily Muscicapoidea (Sibley and Ahlquist 1990). The phylogeny is from Sibley and Ahlquist’s (1990) DNA-hybridization study, with nodes indicated by capital letters, followed by $\Delta T_{m,H}$ values (the difference in melting temperatures of homoduplex and heteroduplex DNA hybrids). We used $\Delta T_{m,H}$ values as estimates of evolutionary branch lengths to standardize our contrasts (Felsenstein 1985; Garland and Adolph 1994; Purvis and Rambaut 1995). Maltase activity was significantly different for the contrast including cedar waxwings compared with other Muscicapoidea contrasts, whereas the corresponding contrast for aminopeptidase-N activity was indistinguishable from other Muscicapoidea contrasts.
Figure 5. Maltase and isomaltase activities plotted against sucrase activity for cedar waxwing intestinal segments to estimate sucrase-independent hydrolysis of maltose and isomaltose. Symbols distinguish individual birds. Lines show least squares regressions (±95% confidence interval) for all data. Both relationships intercept the origin, which suggests that hydrolysis of maltose and isomaltose in cedar waxwings is solely due to sucrase-isomaltase.

carrier-mediated glucose transport remains elevated, and the luminal contents are mixed by retroperistaltic refluxing (Levey and Duke 1992). These traits may be associated with digestive specialization to a diet rich in simple sugars.

Our finding that total aminopeptidase-N activity was the same for both cedar waxwings and thrushes did not support the prediction that proteinase activity would be correlated with the proportion of protein-rich animal foods in a species' overall diet. This result may be partly a consequence of the fact that the scale at which digestive systems operate is the nutrient mixture of gut contents rather than the overall diet. Indeed, the guts of fruit-eating birds tend to contain single food types that reflect their most recent meal (fruit or animal foods; Witmer 1996; C. Martinez del Rio and S. Helmy, unpublished data). Furthermore, our results showing different patterns of aminopeptidase-N activity along the intestines of waxwings and thrushes, but similar levels of total enzyme activity, suggest that total enzyme activity can be insufficient in describing functional differences in the digestive enzymes of animals. Using a single number to estimate digestive capacity of enzymes can obscure important differences among species in the locale and mode of nutrient digestion.

The membrane-bound hydrolases of cedar waxwings and thrushes correspond well with the nutrients in their natural diets. The extent to which the digestive traits of these frugivores influence diet selection will require nutritional studies that measure the utilization of nutrients from experimental diets that vary in nutrient type and concentration. For instance, the lack of sucrase precludes the utilization of sucrose, an abundant plant sugar, strongly influencing the preferences of birds for sucrose-rich foods (Martinez del Rio et al. 1988; Martinez del Rio and Stevens 1989; Martinez del Rio et al. 1990; Malcarney et al. 1994). The ability of frugivorous birds to modulate the amounts and distributions of carbohydrazes and aminopeptidase-N in response to diet composition is not well documented (see Afik et al. 1995). Passerine birds appear not to modulate disaccharidases to the nutritional composition of the diet, but dipeptidase activity often varies with the level of dietary protein (Afik et al. 1995; Karasov 1996; Sabat et al. 1998; but see Levey et al. 1999). Thus, for frugivorous passerines, results for amnepptidase-N are likely to depend on the dietary regimen of experimental subjects. Because nutrient utilization is a complex process that is determined by a large suite of morphological and physiological traits (Alpers 1987), as well as interactions among dietary components, determining the functional limits to diet breadth of any particular trait will require detailed nutritional and physiological studies.

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