Intestinal capacity of P-glycoprotein is higher in the juniper specialist, *Neotoma stephensi*, than the sympatric generalist, *Neotoma albigena*

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Abstract

Permeability-glycoprotein (Pgp) is a membrane-bound, ATP-dependent, transport protein that excludes many cytotoxic compounds including plant metabolites and pollutants from the barrier epithelia of many tissues including the small intestine. We hypothesized that intestinal Pgp capacity would be higher in *Neotoma stephensi*, a specialist on *Juniperus monosperma* known to be high in plant toxins, than the sympatric generalist, *Neotoma albigena*, which consumes juniper in the field, but is unable to tolerate a high juniper diet. We measured Pgp activity as the difference in accumulation of a known Pgp substrate, digoxin, between everted sections of small intestine exposed to ethanol vehicle control and a maximal level of a known competitive inhibitor of Pgp, cyclosporin A. We estimated intestinal capacity by averaging Pgp activity along the intestine and multiplying by total small intestine mass. These first measures of Pgp in wild mammals show a significant difference among species with the juniper specialist, *N. stephensi*, exhibiting a 2.4 fold higher capacity than the generalist, *N. albigena*. This result suggests that Pgp may play a role in the ability of *N. stephensi* to tolerate juniper.

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**Keywords**: P-glycoprotein; *Neotoma*; Woodrat; *Juniperus monosperma*; Multi-drug resistance

1. Introduction

Biochemical defenses against ingested toxins vary greatly among species and populations within species (Walker, 1980; Mead et al., 1985; Van Straalen, 1994; Fossi et al., 1995). A general hypothesis regarding this variation is that populations or species that are evolutionarily exposed to toxins and have developed a tolerance, should exhibit higher levels of biochemical defenses. Much of the evidence in support of this hypothesis comes from studies of the phase I and II biotransformation enzymes among major dietary groups such as omnivorous and piscivorous birds (Knight et al., 1981; Walker and Knight, 1981; Ronis and Walker, 1985; Fossi et al., 1995). Lower mixed function oxidase activity in piscivorous birds has been attributed to a lower evolutionary exposure to dietary toxins (Walker and Knight, 1981) and increased omnivory has been correlated with increased CYP 2B activity, which is responsible for detoxification of xenobiotics (Knight et al., 1981; Ronis and Walker, 1985; Fossi et al., 1995). None of these broad scale comparisons, however, accounted for phylogeny (Walker, 1980; Walker and Knight, 1981; Ronis and Walker, 1985). Furthermore, there have been no comparisons within other dietary groups (e.g., herbivores). Although we might expect dietary specialists and generalists to differ in defense mechanisms that act on a small range of compounds that differ in their diets, predicting the outcome for a more general defense mechanism that recognizes a broad range of substrates is more difficult. Without data to the contrary we cannot assume that a specialist and generalist differ in
overall toxin ingestion, which precludes formulating a
general hypothesis regarding dietary specialization.

As more general biochemical defenses have been
discovered and described in different species (Taipalusyy
et al., 2001; Scotto, 2003), their pattern of variation among
populations or species can be evaluated within the above
context. A recently described general biochemical defense
mechanism is Permeability-glycoprotein (Pgp), which has
affinity for numerous and varied compounds that include
toxins of plant origin (Ford and Hait, 1990). Pgp is a
membrane bound transporter that actively expels toxins that
diffuse into the cell membranes of barrier epithelial cells in
the small intestine, blood–brain barrier, blood testis barrier,
and in organs that routinely deal with toxins such as liver
and kidney (Gottesman and Pastan, 1993; Sarkar, 1995;
Hunter and Hirst, 1997; Brady et al., 2002).

Woodrats of the Neotoma genus provide an excellent
opportunity to investigate a general defense mechanism like
Pgp in a wild mammalian herbivore system. Both dietary
generalists and specialists that consume toxin laden plants,
exist within Neotoma (Atsatt and Ingram, 1983; Dial, 1988;
Edwards et al., 2001). Neotoma albigula, a generalist, and
N. stephensi, a juniper specialist, are sympatric and differ in
their dietary inclusion (Dial, 1988) and tolerance (Dearing et
al., 2000) of Juniperus monosperma leaves. Although we
lack a complete understanding of the chemical composition
in either the specialist or generalist diet, there is evidence
indicating a difference in overall toxin load. J. monosperma
is high in numerous terpenoids and phenolic compounds
(Adams et al., 1981; Adams, 1994), and the dominant
monoterpenes, alpha-pinene, is known to cause toxicity in
mammals (Sperling et al., 1967; Savolainen and Pfaller,
1978; Dearing et al., 2000). Although N. albigula also
consumes J. monosperma (24% of diet), 29% of the diet is
made up of Yucca baileyi (Dial, 1988). There are no data on
the toxicity of Y. baileyi, so we must rely on data from a
congener. Yucca shidigera is used as a feeding supplement
in domestic mammals (i.e. pigs, cattle, rabbits) and results
indicate only positive effects including mass gain and
increased digestive efficiency (Hussain and Cheeke, 1995;-
Hussain et al., 1996; Killeen et al., 1998; Min et al., 2001;
Hong et al., 2001). Added to the dietary data are physiological comparisons between N. stephensi and N.
albigula gavaged with alpha-pinene that indicate N.
stephensi exhibits a lower plasma concentration of alpha-
pinene (Sorensen and Dearing, 2003), and an increased
excretion of unchanged alpha-pinene in the feces (Sorensen
et al., 2004). The authors implicated a difference in
intestinal absorption possibly mediated by Pgp. Together
these data lead us to hypothesize that N. stephensi will
exhibit higher Pgp activity than N. albigula.

To test this hypothesis, we measured Pgp activity in the
small intestine, which is the first Pgp expressing organ that
could serve as a barrier to absorption of ingested toxins. To
strengthen our comparison within an evolutionary/phyloge-
etic context, we included the more distantly related
congener (Edwards et al., 2001; Edwards and Bradley,
2002). N. cinerea, that also consumes a generalist diet
(Johnson and Hansen, 1979; Haufier and Nagy, 1984; Frase
and Sera, 1993). Among the three species we compared
intestinal Pgp capacity, which we define as the activity of
Pgp over the entire small intestine.

2. Materials and methods

2.1. Animals

Specialist (Neotoma stephensi) (2 male, 3 female) and
generalist (N. albigula) (4 male, 3 female) woodrats were
trapped outside the south border of Wupatki National Park,
45 km NE of Flagstaff, AZ (35°30' N 111°27' W) from the
same area with equal access to J. monosperma. The diets of
these species were described in an extensive study by Dial
(1988). Neotoma cinerea (2 male, 3 female) were trapped at
a private residence near Heber City, Summit, UT. N. cinerea
is a dietary generalist (Johnson and Hansen, 1979; Haufier
and Nagy, 1984; Frase and Sera, 1993). The caches of the N.
cinerea suggested they were foraging on Opuntia cladles,
sagebrush (Artemiseta tridentata) and various non-native
garden plants. Animals were transported to the University of
Utah Animal Facility and were housed individually in shoe
box cages (48×27×20 cm) with bedding and cotton batting
at 20 °C on a 12L:12D photoperiod. All animals were fed
Teklad ground rabbit chow (formula 2120) and water ad
libitum. Animals were in captivity for 6–12 months prior to
experiments. All experiments on woodrats conformed to
University of Utah IACUC protocol #04-02012.

2.2. Chemicals

[3H]-digoxin and [14C]-polyethylene glycol were pur-
chased from PerkinElmer Life and Analytical Sciences
(Boston, MA, USA). [14C]-D-glucose and [3H]-L-glucose
were purchased from Moravek (Brea, CA, USA). Phloridzin
was purchased from Sigma Aldrich (St. Louis, MO, USA)
and cyclosporin A (CsA) was purchased from Qbiogene
(Carlsbad, CA, USA).

2.3. Western blot analysis for Pgp

Pgp has previously been found in numerous tissues
including the small intestine of Sprague–Dawley rat (i.e.
Ambudkar et al., 1999). We conducted Western blot analysis
to verify that a Pgp homolog was present in the small
intestine of the three species of woodrat in this study.
Brush border membrane vesicles (BBMV) for analysis of
membrane-bound Pgp were prepared from frozen tissues as
previously described (Barnes, 2001). Frozen tissues were
homogenized with a SDT 1810 Tekmar Tissumizer (Cin-
cinnati, OH, USA) in ice-cold homogenization buffer (300
mM mannitol, 12 mM Tris/HCl, 5 mM EGTA, pH 7.1–7.2
sterile filtered, with 2 μg/mL aprotinin, 2 μg/mL leupeptin, 1mM PMSF). The homogenate was centrifuged at 1000×g for 5 min at 4°C. The supernatant was made 10 mM with MgCl₂, placed on ice for 15 min, then centrifuged at 23000×g for 15 min at 4°C. Membrane vesicles were isolated by centrifugation at 23,000×g for 30 min at 4°C, re-suspended in storage buffer (300mM mannitol, 20 mM HEPES, 10 mM Tris/HCl, 4 mM MgCl₂, pH 7.4, sterile filtered), aliquoted, and stored at -70°C until analyzed.

For the immunodetection of Pgp, 10–30 μg of solubilized BBMV protein were electrophoresed on vertical 7% polyacrylamide mini-gels and Western blotting of the separated proteins was performed using standard protocols. We used rabbit polyclonal primary antibodies raised against human Pgp (C-219, Signet Pathology Systems, Dedham, MA, USA) followed by anti-rabbit IgG conjugated with horse-radish peroxidase (HRP). Pgp is highly conserved among a diversity of species and the C-219 antibody has been used to detect Pgp homologs in mammals (Georges et al., 1990; Kamath and Morris, 1998; Veau et al., 2002), birds (Barnes, 2001), fish (Doi et al., 2001), and even a mollusk (Kepper and Ringwood, 2001). Immunoreactive proteins were visualized using photographic film. Membrane vesicles prepared from the rat hepatoma cell line, H-4-II-E, were run with the samples from Neotoma species to verify the molecular weight of Pgp. We have used this method previously to detect Pgp homologues in a variety of species including Japanese quail (Coturnix japonica), chicken, and American robin (Turdus migratorius) (unpublished data). We used the gels only to verify presence of a Pgp homolog, not to quantify protein expression.

2.4. Measures of Pgp activity using tissue accumulation of digoxin

For tissue preparation and mounting, we followed closely the procedures described in (Karason and Diamond, 1983; Karason et al., 1985). Briefly, animals were euthanized and the intestine from the stomach to the proximal end of the caecal attachment was quickly removed (4–6 min) and flushed with ice-cold Ringer solution. Solution composition (in mM) was 50 mannitol, 100 NaCl, 4.7 KCl, 2.5 CaCl₂, 1.2 KH₂PO₄, 1.2 MgSO₄, and 20 NaHCO₃ gassed with 95% O₂ and 5% CO₂. Osmolarity (mosM) was 290. After measuring intestine length, the intestine was everted, and short sleeves (1.5 cm long) were cut and then mounted on stainless steel rods by tying the tissue down over grooves exactly 1-cm apart. During this preparatory stage, mounted and un-mounted tissues were kept in cold Ringer gassed with 95% O₂ and 5% CO₂.

Just prior to a flux measurement, a rod with a mounted tissue was pre-incubated in 37°C Ringer solution bubbled with 95% O₂/5% CO₂ for 2 min (Pgp assays) or 6 min (D-glucose uptake). Pre-incubation solutions also contained either test compounds or ethanol vehicle control to maximize tissue exposure to our treatments. The tissue was then transferred to a flat-bottomed water-jacketed test tube containing 8 mL gassed Ringer with radiolabeled probe and marker solutes at 37°C. The tube contained a stir bar spinning rapidly (approx. 1000 rpm) to control the unstirred layer thickness (Karason and Diamond, 1983). Probe molecules used were a specific substrate of Pgp, [³H]-digoxin (9.1 kBq/mL), for measuring Pgp activity (Sababi et al., 2001; Stephens et al., 2001), or [¹⁴C]-D-glucose (1.1 kBq/mL) for measuring D-glucose uptake activity. To correct for non-absorbed probe in adherent mucosal fluid, we used tracer concentrations of a membrane-impermeable marker [¹⁴C]-polyethylene glycol (PEG; MW=4000; 1.1 kBq/mL), or in the case of [¹⁴C]-D-glucose uptake we used [³H]-L-glucose (31.4 kBq/mL) to correct for both non-absorbed probe in adherent mucosal fluid and diffusive flux (Karason and Diamond, 1983). After 12 min (Pgp assay) or 4 min (D-glucose uptake) the incubation was terminated. We used longer incubation times with [³H]-digoxin (12 min) than are generally used for measuring sugar and amino acid uptake (generally 1–4 min) because digoxin is passively absorbed relatively slowly, and because the target of our measurements was not the accumulation in tissue per se, but a difference in digoxin accumulation in the presence and absence of a competitor, which would reflect digoxin excluded by Pgp (Green et al., in press). After incubation, tissues were then either simply blotted (Pgp assay) or rinsed in stirred cold Ringer and then blotted (D-glucose uptake). The mounted tissue was cut from between the two grooves with a razor blade, weighed, incubated in 1 mL of tissue solubilizer (Soluene-350, Packard, Meriden, CT, USA), and counted in 10 mL of scintillation cocktail (Ecolume, Packard) with 0.5% by volume acetic acid on a Beckman LS 5801 (Beckman Coulter) with counting channels set to minimize spill (counts of the alternate isotope appearing in the same counting channel) and programmed with a standard curve to correct for background quench. Calculation of digoxin accumulation followed (Karason and Diamond, 1983). For simplicity of expression, we assumed a digoxin concentration of 1 fmol/μL incubation solution (it was actually approximately 0.75 fmol/μL), yielding units of fmol digoxin accumulated per mg tissue. Total intestine mass was estimated as the product of the summed mass of the 16, 1-cm tissue samples and the ratio of total intestine length: 16 cm.

All experiments were conducted between April 11, 2003 and May 7, 2003 within which we blocked for time and randomized species within each time block. One experiment was conducted each day between 12:00 noon and 16:00 h. Two types of trials, CsA dose response and tissue viability, were used to optimize the method in woodrats. They are described in detail below.

2.5. CsA dose response

First, we measured [³H]-digoxin uptake in the absence and presence of increasing concentration of CsA (1, 5, 10
μM, and saturated (~15 μM) CsA), a known competitive inhibitor of Pgp (Lan et al., 1996). This was done because digoxin accumulation in the tissue is a balance between passive diffusion into and Pgp-mediated export out of the cells. The desired measure of Pgp activity index was the difference in digoxin accumulation per mg tissue between a tissue with Pgp maximally inhibited by the competitor (mimicking a tissue without Pgp) and the accumulation in an adjacent tissue without the competitor (ethanol vehicle control). This estimates the amount of digoxin that would have diffused into the tissue if Pgp were not present. In these trials, we used adjacent slices of intestine from the mid-intestinal region. The preincubation was 2 min in control (0.15% ethanol) or CsA-containing Ringer solution and the incubation was 12 min in the same solutions with [3H]-digoxin.

2.6. Determination of tissue viability and damage

Second, as checks on the structural and functional integrity of the intestinal sleeves, we histologically inspected adjacent tissues in two individuals per species for signs of villus damage (Starck et al., 2000). Also, we measured mediated D-glucose uptake over the same combined pre-incubation and incubation time, using adjacent tissues exposed to CsA (15 μM), vehicle control (0.15% ethanol), and phlorizin (1 mM), a known inhibitor of the intestinal glucose transporter SGLT-1. Histology preparations indicated that woodrat intestinal tissue was not damaged during the experimental procedure for any of the three species. Villi in experimental sleeves were shorter than in sleeves that were not exposed to the experimental procedure, but remained intact (data not shown). Glucose uptake data are presented in Results.

2.7. Total Pgp capacity

Along with a measure of Pgp activity in the mid-gut, we included measures in the proximal and distal regions. To estimate the intestine’s total capacity, we computed a weighted average of Pgp activity per mg intestine (0.25×proximal+0.5×mid+0.25×distal) and multiplied that by the small intestine mass. We tested differences among species by first comparing intestinal Pgp capacity using body mass as a covariate after examining the interaction between body mass and Pgp capacity. Then, because many physiological processes are scaled to body mass, we also scaled Pgp capacity to body mass using (1) kg0.54, because this is how food (and hence toxin) intake scale with mass in woodrat species (Smith, 1995), and (2) kg0.75, a generally accepted metabolic mass scalar.

2.8. Statistical analyses

When possible, we analyzed treatment effect with repeated measures, two-factor ANOVA with treatment and species as factors. When all treatments or concentrations were not represented in all animals we determined treatment effect with a one-way, two-factor ANOVA with treatment and species as the two factors. We also tested differences in intestinal Pgp capacity using ANCOVA with log [intestinal Pgp capacity] vs log [body mass] as the covariate after examining the interaction, and using one-way ANOVA on intestinal Pgp capacity scaled to body mass. We used Fisher’s LSD for pair-wise comparisons when global tests resulted in P<0.05. Plots of residuals were visually inspected to verify a random distribution. A P<0.05 was considered significant and 0.05<P<0.10 a trend. All data are reported as mean±S.E., n=sample size.

3. Results

Western blot analysis confirmed the presence of a Pgp homolog in the small intestine of all three woodrat species (Fig. 1) as expected based on our previous results with other species (unpublished data). The main difference seen among the species was that Pgp from N. stephensi migrated further down the blot than either N. albigena or N. cinerea and we will discuss this subsequently. We removed one N. cinerea from the study due to consistently low values for mediated D-glucose uptake indicating a loss of functional integrity in intestinal sleeves.

Accumulation of tracer [3H]-digoxin, corrected for adherent label, increased sublinearly with increasing concentration of CsA in the solution for intestinal sleeves from all three species (Fig. 2). Cyclosporin A precipitated from solution at around 15 μM, but [3H]-digoxin accumulation at ~15 μM was not significantly higher than at 10 μM. Therefore, in subsequent experiments, we used 10 μM CsA as the maximal inhibitory concentration, and calculated the index of Pgp activity as the difference between accumulation of [3H]-digoxin in the presence and absence of 10 μM

![Fig. 1. Western blot analysis of Pgp homologs using protein from brush border membrane vesicles concentrated from mid-intestinal tissue of N. albigena, N. cinerea, and N. stephensi. Membrane vesicles from the rat (Rattus norvegicus) hepatoma cell line, H-4-II-E, were used as positive control and to verify molecular weight of the Pgp homolog. Numbers in parentheses are μg of total protein loaded in the lane.](image-url)
Fig. 2. (a) Digoxin accumulation (fmol/mg tissue) in mid-intestinal sleeves exposed to cyclosporin A (1, 5, 10, ~15 μM) or vehicle control (0 μM CsA; 0.15% ethanol) and (b) the difference in digoxin accumulation (fmol/mg tissue) between CsA treated and control tissues during 2 min pre-incubations and 12 min incubations in Neotoma albigna (n=4–7 sleeves/concentration), N. cinerea (n=3–4 sleeves/concentration), and N. stephensi (n=3–5 sleeves/concentration). All three species show sublinear increase in digoxin accumulation and in the difference between tissues exposed to CsA and control, and the relationships reach an asymptote by 10 μM (repeated measures ANOVA within species: digoxin accumulation at 10 μM CsA>0 μM CsA, P<0.05; digoxin accumulation at 10 and 15 μM CsA not significantly different, P=0.1). In the box above the plots, significant differences (P<0.05) from control within species are indicated by (*) based on paired Student’s t-tests in (a) or one sample Student’s t-tests in (b). Points are mean±S.E.

CsA in adjacent intestinal tissues (i.e., digoxin excluded per mg wet tissue mass).

Although we assumed that the enhancement of digoxin accumulation by CsA is due to competitive inhibition of digoxin’s exclusion by Pgp, it is also conceivable that CsA acts by depressing ATP generation, which powers the expulsion of compounds by Pgp (Ambudkar et al., 1999). To test for this nonspecific effect, we compared meditated uptake of 1 mM D-glucose, which is also powered by ATP generation, in the presence and absence of 15 μM CsA (in 0.15% ethanol as vehicle). We also included a measure of D-glucose uptake in the presence of phloridzin (1 mM), a potent inhibitor of the D-glucose transporter SGLT-1, to

Fig. 3. D-Glucose uptake in mid-intestinal sleeves exposed to vehicle control (0.15% ethanol), CsA (15 μM), and phloridzin (1 mM) during 10 min pre-incubation and 4 min incubation in N. albigna, N. cinerea, and N. stephensi. (+) indicates significant differences (P<0.01) from control after significant repeated measures ANOVA (F_{2,14}=12.217, P=0.001) with no significant interactions.

Fig. 4. Variation in Pgp activity (fmol digoxin excluded/mg tissue) along the intestinal tract of N. albigna (n=7), N. cinerea (n=4), and N. stephensi (n=5) as determined by the difference in digoxin accumulation in intestinal sleeves exposed to vehicle control (ethanol) or 10 μM CsA during 2 min pre-incubations and 12 min incubations. N. albigna and N. cinerea, but not N. stephensi exhibit significant differences in Pgp activity from proximal to distal sections of the intestine (repeated measures ANOVA within species, P=0.005, P=0.013, P=0.013, respectively). Different letters (a, b, c) indicate significant differences (pair wise comparisons, P<0.05) among intestinal sections within each species. Points are mean±S.E.
confirm that our measure was indeed of active D-glucose uptake. D-glucose uptake was significantly depressed by phlorizin (P=0.004), but not by CsA (P=0.148) (Fig. 3; repeated measures ANOVA, F2,13=12.217, P=0.001, followed by pair wise comparisons).

* N. albigna and N. cinerea, but not N. stephensi exhibited significant differences in Pgp activity from proximal to distal sections of the intestine (Fig. 4; repeated measures ANOVA within species: P=0.005, P=0.013, P=0.091, respectively). Data on body mass, small intestine length and mass, and intestinal Pgp capacity are summarized in Table 1. We calculated intestinal Pgp capacity by multiplying the weighted average of Pgp activity along the intestine, by intestine mass. Because N. cinerea is significantly larger than either N. stephensi or N. albigna we included log [body mass] as a covariate in an ANCOVA of log [intestinal Pgp capacity]. The scaling (i.e. mass exponent) for the covariate, however, was much higher (1.31) than expected (0.54–0.75). Therefore, we also tested the difference by scaling intestinal Pgp capacity to body mass (kg\(^{-0.54}\)) based on the scaling of food intake in three Neotoma species (Smith, 1995), and to body mass (kg\(^{0.75}\)), a generally recognized scaling factor for metabolic processes. By whatever comparative analysis used, N. stephensi exhibited a significantly higher intestinal Pgp capacity than N. albigna (Table 1).

4. Discussion

In this study, we made the first measures of Pgp activity in wild mammals and the first comparison of intestinal Pgp capacity among mammalian herbivores that differ in dietary specialization. Our data support the hypothesis that the juniper specialist N. stephensi has greater Pgp activity along the small intestine (intestinal Pgp capacity) than the sympatric generalist N. albigna. First we discuss the validity of the method in Neotoma, then the possible reasons for the observed differences in Pgp and potential future research.

These data on Pgp activity are the first in wild mammals, so a concern is the validity of the method and viability of woodrat intestinal tissue throughout the experimental procedure. Evidence that suggests our method was successful in measuring Pgp activity in these woodrats include: (1) A Pgp homolog is found in woodrat intestinal tissue as indicated by Western blot analysis. (2) We observed the expected sublinear digoxin accumulation with an increasing dose of the known Pgp competitive inhibitor, CsA, that reached an asymptote by 10 \(\mu\)M in each of the three species. These results match our previous results in chicken and Sprague-Dawley rat (Green et al., in press). (3) Tissues withstood the experimental procedure both structurally, based on histological preparations that show intact villi, and functionally, based on active D-glucose uptake over the experimental time period with or without CsA. An indication of a loss in functional integrity allows us to remove a study animal as was done with one N. cinerea in this study.

Pgp expression and activity in rodents can be induced by substrates administered in the diet (Salphati and Benet, 1998; Sandstrom and Lennernas, 1999), and Pgp induction may play a role in how Neotoma woodrats deal with dietary toxins. We feel that our data represent basal levels of Pgp activity as all animals were in captivity for 6–12 months prior to the experiment, feeding on the same chow diet. Of interest would be whether there are differences in the ability of each species to induce Pgp when exposed to dietary substrates.

Our finding that the juniper specialist, N. stephensi, has significantly higher intestinal Pgp capacity than N. albigna adds to the mounting evidence that these two species differ in their treatment of ingested toxins. Dearing et al. (2000) found that N. albigna produces a greater volume of more acidic urine compared to N. stephensi when on the same diet high in alpha-pinene, the dominant monoterpene in J. monosperma. This indicates a difference in the detoxification pathway. Supporting this assertion are preliminary data from S. Haley (unpublished data) suggesting a difference in the activity levels of phase I and II enzymes in liver tissue of these two species. Sorensen and Dearing (2003), using a pharmacokinetic approach, compared the in vivo absorption of alpha-pinene in N. stephensi and N. albigna. Their results suggested less absorption of alpha-pinene in the specialist than the generalist. Adding to these results is a recent finding that N. stephensi excretes more unchanged alpha-pinene in the feces than N. albigna (Sorensen et al., 2004). These latter two studies implicated Pgp as a potential mechanism that decreases intestinal absorption of alpha-pinene to a greater extent in the specialist than the generalist.

Table 1

<table>
<thead>
<tr>
<th>Species</th>
<th>Body mass (g)</th>
<th>Small intestine length (cm)</th>
<th>Small intestine mass (g)</th>
<th>Intestinal Pgp capacity (fmol digoxin/kg(^{0.54}))</th>
<th>Intestinal Pgp capacity (fmol digoxin/kg(^{0.75}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>N. albigna (n=7)</td>
<td>192.8±7.3</td>
<td>38.6±1.6</td>
<td>3.2±0.3</td>
<td>414±82</td>
<td>993±247</td>
</tr>
<tr>
<td>N. cinerea (n=4)</td>
<td>239.9±36.1</td>
<td>42.2±9.4</td>
<td>5.2±0.4</td>
<td>803±49</td>
<td>1563±327</td>
</tr>
<tr>
<td>N. stephensi (n=5)</td>
<td>182.0±19.1</td>
<td>53.5±2.7</td>
<td>7.5±2.6</td>
<td>955±186</td>
<td>2404±293</td>
</tr>
</tbody>
</table>

Statistics:
- \(F_{2,13}=12.4, P<0.001\)
- \(F_{2,13}=19.3, P=0.001\)
- \(F_{2,13}=13.0, P=0.001\)
- \(F_{2,13}=6.2, P=0.014\)
- \(F_{2,13}=6.8, P=0.010\)
- \(F_{2,13}=6.9, P=0.009\)

Different letters in parentheses (a or b) indicate significant differences among species (\(P<0.05\)) within each column from pair wise comparisons following significant one-way ANOVA, unless otherwise noted.

* ANCOVA of log [intestinal Pgp capacity] with log [body mass] as the covariate.
There are two issues that should be considered when extrapolating our data to the observed difference in juniper tolerance. First, extrapolation of these data presupposes juniper toxins are recognized by Pgp. Establishing a causal link between Pgp and alpha-pinene absorption, for example, requires that we determine whether Pgp recognizes alpha-pinene as a substrate, which is the subject of ongoing research. Second, although intestinal Pgp capacity as we defined it is significantly higher in *N. stephensi*, this does not necessarily translate into decreased toxin absorption at the whole animal level because there may be other transporters that influence the absorption and distribution of toxins. For example, multidrug resistance-associated proteins (MRPs) or MRP-like organic anion transporting polypeptides (OATPs) could be present in intestinal cells (Walters et al., 2000; Cattori et al., 2001) and transport digoxin (Kullak-Ublick and Becker, 2003; Mikkaichii et al., 2004) and thus influence digoxin accumulation. This might explain why the baseline intracellular digoxin levels of the specialist *N. stephensi* were not significantly different from those of the generalist *N. albiculata* (see Fig. 2a), rather than being lower as might be expected if Pgp expression were the only factor influencing those levels. Although other factors might influence the baseline digoxin levels in our experiments, to the best of our knowledge the CsA inhabitable component of digoxin uptake is most specific to Pgp, which is why we think that the specialist exhibits higher Pgp activity. But, the diversity of toxin transporting and biorienting polypeptides underscores how complicated it can be to investigate biochemical defense against ingested toxins. New techniques that can simultaneously measure activities of numerous genes or gene products, such as DNA micro-arrays, may prove to be more powerful in comparative studies than comparisons of only one enzyme or transporter among species.

Identifying the underlying mechanism for the observed difference in Pgp activity requires more study. We found the Pgp homologs among woodrat species migrated differently down the gel (Fig. 1). This could be due to post-translational modification of Pgp such as glycosylation that is not known to affect function (Ambudkar et al., 2003), or differences in isoforms. Future studies should look more closely at specific differences in the Pgp protein in *Neotoma* and how they relate to function. We did not draw any conclusions from relative blot intensity among the species because we had a small sample size limited to mid-intestinal tissue that was not consistently collected from the same region as our activity measures.

An ancillary goal of this research was to continue acquiring data within the *Neotoma* genus to better elucidate the evolutionary trajectory of biochemical defense mechanisms. The intermediate level of intestinal Pgp capacity in the more distantly related generalist *N. cinerea*, however, makes it difficult to determine the ancestral trait in *Neotoma*. It may be that the generalist diets of *N. cinerea* and *N. albiculata* are not equivalent in overall toxin load. Clearly, more data on dietary toxin load and Pgp activity are needed before we can confidently predict Pgp levels based on dietary specialization. Fortunately, there are a number of *Neotoma* species for which we have data on wild diet (i.e. Thompson, 1982; Atsatt and Ingram, 1983; Karasov, 1989; Post, 1992; McMurry et al., 1993), that can be tested to address this relationship.

Variation in the tolerance of plant defensive chemicals among mammals is complicated, probably reflecting the diversity of ingested plant compounds and the diversity of the herbivores’ physiological and biochemical adaptations. Studies of the variation among species in phase I and II detoxifying enzymes have thus far dominated the literature (i.e. Walker, 1980; Knight et al., 1981; Ronis and Walker, 1985; Walker and Ronis, 1989; Van Straalen, 1994). Our study underscores that interspecies variation in Pgp activity adds to this complexity. Also, to date there have been 10 ABC-superfamily transport proteins shown to confer cellular resistance to toxins (Taipalensuu et al., 2001; Scotto, 2003). Research on these transporters has been limited to their role in human physiology, so we lack data on non-laboratory animals. The similarities in regulation of the ABC-transporters (Scotto, 2003) and the potential link between Pgp and phase I detoxifying enzymes (Cummins and Benet, 2001; Schuetz and Strom, 2001) suggest a coordinated response that may prove to be very complex. The everted sleeve technique has proven useful to determine the function of Pgp and may be modified to investigate other transporters, providing a tissue-level check of data from in vivo, cell culture, or DNA micro-array techniques.

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