Evidence for a Tradeoff Between Retention Time and Chewing Efficiency in Large Mammalian Herbivores

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Accessibility
Evidence for a tradeoff between retention time and chewing efficiency in large mammalian herbivores

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Running head: Particle size and digestion in herbivores
Abstract

Large body size is thought to produce a digestive advantage through different scaling effects of gut capacity and food intake, with supposedly longer digesta retention times in larger animals. However, empirical tests of this framework have remained equivocal, which we hypothesize is because previous comparative studies have not included digesta particle size. Larger particles require more time for digestion, and if digesta particle size increases with body mass, it could explain the lack of digestive advantage in larger herbivores. We combine data on body mass, food intake, digesta retention and digestibility with data on faecal particle size (as a proxy for digesta particle size) in 21 mammalian herbivore species. Multiple regression shows that fibre digestibility is independent of body mass but dependent on digesta retention and particle size; the resulting equation indicates that retention time and particle size can compensate for each other. Similarly, digestible food intake is independent of body mass, but dependent on food intake, digesta retention, and particle size. For mammalian herbivores, increasing digesta retention and decreasing digesta particle size are viable strategies to enhance digestive performance and energy intake. Because the strategy of increased digesta retention is usually linked to reduced food intake, the high selective pressure to evolve a more efficient dentition or a physiological particle separation mechanism that facilitates repeated mastication of digesta (rumination) becomes understandable.

Key words: herbivory, digestion, retention, mastication, metabolism, rumination
Introduction

Mammalian herbivores are thought to facilitate niche separation by the so-called Jarman-Bell-principle (Bell 1971; Geist 1974; Jarman 1974). This principle suggests that larger species can feed on diets of lesser quality (i.e., higher fibre content). The proposed mechanistic background of this concept is the fact that whereas metabolic requirements and hence food intake scales to body mass$^{0.75}$, gut capacity scales linearly to body mass; in other words, the amount of food ingested decreases per unit gut capacity, which should in theory lead to an increase in digesta retention time with increasing body mass (Parra 1978; Demment and Van Soest 1983; Demment and Van Soest 1985; Illius and Gordon 1992; Clauss et al. 2007a). Because digesta retention is a major determinant of digestibility, large body size has been suggested as a major digestive advantage and thus as one of the drivers of Cope’s rule in herbivores (Demment and Van Soest 1985). Additionally, the Jarman-Bell-principle has been suggested to facilitate intraspecific niche separation in sexually dimorphic ungulates (Barboza and Bowyer 2000).

Because of the perceived relevance of digesta retention, numerous studies have investigated this parameter, often in conjunction with digestibility measurements (reviewed in Clauss et al. 2007a). However, attempts to correlate digestive efficiency or digesta retention with body mass have remained unsatisfactory for at least three reasons. First, no systematic increase of either parameter with body mass could be demonstrated across species (Justice and Smith 1992; Pérez-Barberia et al. 2004; Clauss and Hummel 2005; Clauss et al. 2007a). Second, differences in digestive efficiency between sexes of dimorphic species are either absent or low (Gross et al. 1996; Pérez-Barberia et al. 2008). Third, it is felt that ruminants need to be considered separately from other herbivores due to their higher digestive efficiency (Illius and Gordon 1992), a preconception which prevents a unifying framework for herbivory.
While the concept that digesta retention increases with body size (Demment and Van Soest 1985; Illius and Gordon 1992) is only weakly supported by empirical evidence so far (Clauss et al. 2007a), the concept that chewing efficiency decreases (i.e., digesta particle size increases) with body mass (Pérez-Barbería and Gordon 1998) is clearly corroborated in comparative studies (Udén and Van Soest 1982; Clauss et al. 2002; Fritz et al. 2009). The relevance of reducing the particle size of ingested food is well understood, particularly in herbivores (Clauss and Hummel 2005); specifically, smaller food particles can be digested at a much faster rate. Therefore, many authors have speculated that an increase in chewing efficiency permits shorter digesta retention times, or that – vice versa – longer digesta retention can compensate for a reduced, or even lacking, chewing efficiency. This tradeoff has been evoked for comparisons between chewing and non-chewing dinosaurs (Farlow 1987; Sander and Clauss 2008), between reptiles and mammals (Karasov et al. 1986), between different large mammalian hindgut fermenters (Clauss et al. 2005), between ruminant and non-ruminant foregut fermenters (Schwarm et al. 2009), or between the sexes of a dimorphic ruminant species (Gross et al. 1995). However, a statistical demonstration of such a compensating effect across species has not been presented so far, most likely because data on digestibility, digesta retention and digesta particle size was not available for a sufficiently large dataset (Schwarm et al. 2009).

Here, we use the most comprehensive dataset from one single trial on food intake, digesta retention and digestive efficiency in large grazing mammals fed grass hay (Foose 1982), and add our own data on digesta particle size (from Fritz et al. 2009) determined for the same species by wet sieving analysis of faeces. We use conventional and phylogeny-based methods to investigate the scaling of digestive traits with body mass and to test two predictions. First, we predict that fibre digestibility is mainly influenced by digesta retention time and digesta particle size (and not by body mass). Second, we predict that energy intake is dependent on overall food intake, digesta retention, and digesta particle size (and not on body
We examine variation in a phylogenetic and statistical context that enable us to examine evolutionary change in two or more traits.

**Methods**

The dataset of Foose (1982) was used, which stems from feeding trials of non-domesticated herbivores held in captivity (Appendix). These trials were performed nearly forty years ago (1970-1980). We used data for a grass hay-only diet from species adapted to grazing. This resulted in exclusion of the tapirs, the giraffe, the black rhinoceros and the pygmy hippo from the original dataset, as browsing species have been reported to have difficulties in grass hay ingestion (Clauss et al. 2008a) and/or to produce larger faecal particles in captivity as compared to the wild (i.e., on their natural food) (Hummel et al. 2008). Thus, the data set includes ruminants and camels (“ruminants”, n=12), elephants, rhinoceroses and equids (“hindgut fermenters”, n=8) and the hippopotamus (“nonruminant foregut fermenter”, n=1).

The animals were adult and not lactating or pregnant beyond the first month (Foose 1982, p. 69). Data were available for body mass (kg), relative organic matter intake (g/kg^{0.75}/d), mean retention time (MRT, h), and the apparent digestibility of organic matter and neutral detergent fibre. Data on faecal (=digesta) particle size (mm) was gained from captive individuals of the same species kept in European zoos (from Fritz et al. 2009).

For all analyses, body mass and particle size were log-transformed to better meet the statistical assumptions, and we used two-tailed tests with a 5% significance level (α=0.05).

For non-phylogenetic analyses, data were analyzed by correlation analysis using SPSS 16.0.1 (SPSS Inc., Chicago, IL, USA). To analyze results in a phylogenetic context, we used phylogenetic generalized least squares (PGLS) (Pagel 1997; Pagel 1999). For this, we used the program BayesTraits (Pagel and Meade 2007) to calculate likelihood statistics under models of correlated or uncorrelated evolution. We calculated the parameter λ, which was used to assess whether traits show evidence for phylogenetic signal (Freckleton et al. 2002).
Values of $\lambda$ close to one indicate the existence of phylogenetic signal, and we used a likelihood ratio test to compare the likelihoods of models when $\lambda$ was estimated to models in which $\lambda$ was forced to be zero, as described in Freckleton et al. (2002). Forcing $\lambda$ to equal zero is equivalent to a non-phylogenetic test. We also used BayesTraits to implement a multiple regression model. Phylogenetic analyses were based on a recent estimate of mammalian phylogeny (Bininda-Emonds et al. 2007).

Results

Scaling with body mass

Results involving the phylogenetic scaling of four measures of digestive physiology with body mass are presented in Table 1. In all cases, we found significant phylogenetic signal ($\lambda$ close to 1 and significantly different from zero), and results from phylogenetic and non-phylogenetic analyses were consistent. We thus present bivariate plots and results of non-phylogenetic analyses in Figure 1 and statistical results from phylogenetic analyses in Table 1.

Digesta retention time was not significantly related to body mass (Fig. 1a), but faecal particle size increased with body mass (Fig. 1b). Given these two first findings, one would expect fibre digestibility to decrease with body mass, but no significant association was found (Fig. 1c); instead, fibre digestibility appears to be higher in ruminants. Body size was also neither correlated significantly with relative organic matter intake (Fig. 1d) nor with relative digestible organic matter intake (Fig. 1e). In our dataset, the intake of digestible organic matter (i.e., the product of organic matter intake and organic matter digestibility) can be considered as a good proxy for energy intake. Because we limited our dataset to only species that readily accepted the offered diet (grass hay), and because no weight loss was evident during the trials in these species (Foose 1982), this relative digestible organic matter intake represents relative maintenance energy requirements (expressed on a metabolic body weight-
basis). Fig. 1e therefore indicates that differences in maintenance energy requirements and hence metabolic rate do occur between different large herbivore species, similar to variation in metabolic rates reported in smaller mammals (McNab 2008).

**Determinants of fibre digestion**

Fibre digestibility increases with digesta retention time (likelihood ratio test: $\chi^2=10.6$, $p=0.001$, $\lambda=0.9$; Fig. 2a), again at generally higher levels in ruminants. Across all species, fibre digestibility decreases with increasing faecal particle size (Fig. 2b), but this result was not significant after controlling for phylogeny (likelihood ratio test: $\chi^2=2.00$, $p=0.16$, $\lambda=1.0$).

We also used PGLS to run a multiple regression analysis with fibre digestibility as the dependent variable and digesta retention time, faecal particle size, body mass and relative organic matter intake as independent variables. The model explained a large proportion of the variation in fiber digestibility ($R^2=0.89$), and the maximum likelihood estimate of $\lambda$ was 0 (equivalent to a non-phylogenetic test). The model produced significant effects for only digesta retention time and particle size (Table 2). As the regression coefficients for these two variables had opposite signs, this is consistent with a compensating effect of these two variables. Based on the high $\lambda$’s in the bivariate tests, we re-ran the multiple regression with $\lambda$ set to 1 ($R^2=0.58$). The results again demonstrated significant (and opposite) effects for retention time and particle size (Table 2).

**Determinants of energy intake**

In order to test for the relevant factors that influence energy intake, and allow a higher relative metabolic rate in a species, we ran a second multiple regression analysis, with relative digestible organic matter intake as the dependent variable, and body mass, relative food intake, retention time and particle size as independent variables. Although the result could be considered self-evident because the dependent variable (relative intake of digestible material)
is a product of the independent variables (relative intake and the factors shown to determine digestibility), this analysis is important because it is the overall intake of digestible material (i.e. energy) that is the currency relevant for the energy budget of the organism, not digestibility itself; actually, the goal to achieve a high digestibility may set a constraint on intake (Clauss et al. 2007b).

As in the previous multiple regression model, the maximum likelihood estimate of $\lambda$ was 0, and a high proportion of the variation was accounted for by the model ($R^2=0.97$). Relative organic matter intake was a highly significant predictor of relative digestible organic matter intake. Body mass was not statistically significant (Table 3). Digesta retention time was also statistically significant, and faecal particle size approached significance. We repeated the analysis with $\lambda=1$ ($R^2=0.90$). In this analysis, only relative food intake was statistically significant, although digesta retention approached significance (Table 3).

Discussion

In contrast to the common assumption in the literature (Demment and Van Soest 1985; Illius and Gordon 1992), we found no evidence that an increase in body mass confers a digestive advantage. The absence of an effect is unlikely to be due to insufficient variation in body mass, as our dataset included species that ranged from 133 to 3402 kg. Instead, the results suggest that to increase digestive efficiency, herbivores either increase digesta retention, or enhance chewing efficiency, or both (or select a diet of higher digestibility, an option not open for most larger-sized herbivores). In comparison to earlier herbivore digestion models (Demment and Van Soest 1985; Illius and Gordon 1992), digesta particle size thus becomes an important variable for understanding digestive adaptations in herbivores. Actually, the lack of consideration that digesta particle size received in earlier concepts might explain why they remain unsatisfactory when applied to mammalian data. The strong phylogenetic signal detected in our analyses indicates that the parameters investigated – mainly digesta retention
and particle size – have played a fundamental role in the evolution of different mammalian large herbivore groups; actually, these results underline the impression already evident from the graphical depiction of the data (Fig. 1-3) that differences in digestive strategy are at the core of the phylogenetic differentiation of large herbivores. The fact that both digesta retention time and faecal particle size show a strong phylogenetic signal, but only particle size shows a significant correlation with body mass after accounting for phylogeny (Table 1), could indicate different constraints on these two parameters – namely that whereas animal lineages might evolve retention times independent of their body size (Clauss et al. 2007a), evolutionary strategies used so far could not completely liberate digesta particle size from the constraining effect of body mass (Fritz et al. 2009). A likely explanation for this interpretation is that tooth size and chewing frequency are both allometrically correlated with body mass (Shipley et al. 1994; Pérez-Barberia and Gordon 1998).

In order to evolve the potential for a higher metabolism, i.e. a higher energy intake, herbivore species should, above all, increase food intake, but should also increase digesta retention in the gut, and reduce digesta particle size. Because an increase in digesta retention is, among large mammals, not the automatic result of an increase in body mass, there appear to be two major strategies to increase digesta retention: an increase of gut capacity or a reduction in food intake (Clauss et al. 2007a). Both of these strategies imply conceptual disadvantages that might limit the scope of adaptation that can be derived from an increase in digesta retention time.

Increasing the relative capacity of the gut might constrain, by volume displacement, the function of other organs. For example, Clauss et al. (2003) and Mortolaa and Lanthier (2005) independently speculated that the high water content in the faeces of large cattle-like ruminants (defecating in ‘pies’), or the observed unusually high breathing frequency in this group of ruminants, could be the result of a space competition between organs in the body cavity, with the particularly voluminous forestomach in these animals reducing the space
available for the organs of water-reabsorption from digesta (colon) or air exchange (lung), respectively. Additionally, increasing gut capacity might ultimately limit the agility of the animal, and therefore, particularly high gut capacities might only be an option for animals that are, due to their ecology or body size, relatively immune to predation, such as hippos or sloths. Yet, gut capacity might be, across vertebrate herbivores, more flexible than one would expect based on mammal data alone: in herbivorous dinosaurs such as stegosaurs, ankylosaurs, or sauropods, relative gut capacities exceeding the ones in mammalian herbivores have been suggested to facilitate long digesta retention to compensate for a lack of chewing mechanisms (Bakker 1986; Coe et al. 1987; Franz et al. 2009).

Food intake was negatively associated with digesta retention in our dataset (Fig. 3). A negative association between food intake and digesta retention follows the common-sense logic that an increased input into a tube will result into an increased output and a shorter passage time; this association has been found both within and between species (Clauss et al. 2007a; Clauss et al. 2007b). Among primates, this relationship was also demonstrated using phylogenetically independent contrasts (Clauss et al. 2008b). In our dataset, however, this relationship was not significant in a phylogenetic test (likelihood ratio test: $\chi^2=0.1$, $p=0.75$). This can be explained by the taxonomic clustering of data along these dimensions (see Figure 3): while artiodactyls (hippopotamus and ruminants) cluster at the low-intake, long-retention end, perissodactyls and elephants cluster at the high-intake, short-retention end of the spectrum. This finding again emphasizes that alternative digestive strategies were a major determinant of lineage diversification in large mammalian herbivores. Nevertheless, the evolutionary option to increase energy gain by increasing digesta retention is potentially constrained by the consecutive, necessary reduction in overall food intake.

The only non-ruminant foregut fermenter in this dataset, the hippopotamus, is a good example of the strategy of particularly long retention times due to a low food intake and an enormous gut capacity (Clauss et al. 2003; Clauss et al. 2004; Clauss et al. 2007b). Due to the
obligatory low food intake on this long-retention strategy, hippos are characterized by remarkably low maintenance energy requirements (Schwarm et al. 2006). Apparently, the range of adaptation possible due to increased digesta retention is limited to comparatively low energy requirements and low metabolic rates. Among other mammals, this strategy is common among non-ruminant foregut fermenters and some small hindgut fermenters (Clauss et al. 2007a; Clauss et al. 2008b; Munn et al. 2008).

On the other hand, there are two major strategies to increase chewing efficiency and thus reduce digesta particle size: by evolving a more efficient dental design, or by increasing the time spent chewing per unit digesta (i.e., rumination). The prerequisite for efficiently increasing the time spent chewing per unit digesta is a sorting mechanism that separates smaller from larger particles (Fritz et al. 2009; Schwarm et al. 2009). Rumination sets a constraint on food intake, because it represents a relevant proportion of the activity budget that can therefore not be used for feeding (Van Soest 1994). Due to this strategy of repeated mastication and moderately long digesta retention, ruminants are thus limited in the amount of food they can ingest (Fig. 1d); but due to the exceptional small digesta particle sizes they achieve (Fig. 1b), they can attain disproportionately high digestibilities for their digesta retention (Fig. 2a). The equids of our dataset represent the strategy of a particularly sophisticated dental design (Jernvall et al. 1996; Fritz et al. 2009) that allows a high degree of digesta particle size reduction (Fig. 1b) without a constraint on food intake. Thanks to their efficient teeth, equids can afford a high food intake and still respectable digestive efficiencies, which potentially allows them higher intakes of digestible matter and energy than ruminants (Foose 1982; Duncan et al. 1990).

Increasing digesta retention will increase digestive efficiency; however, it will, in varying degrees between species, also limit food intake. Increasing chewing efficiency therefore appears as an attractive alternative to enhance energy uptake. This implication explains the high selective pressure on mammals to acquire more efficient dental designs if they were to...
fuel organisms of increasing metabolic scope (Reilly et al. 2001) – because the adoption of a
more efficient dental design is ultimately the only strategy to enhance digestive efficiency
without compromising food intake. In order to fully understand the ecophysiological
diversification of herbivores, not only gut capacity, food intake, and digesta retention, but also
ingesta particle size reduction must be taken into consideration.

Acknowledgements

The core data used in this study stems from the study of Thomas J. Foose, who passed away
in 2006, before we could ask him to co-author this paper. This is contribution no. 54 of the
DFG Research Unit 533 “The Biology of Sauropod Dinosaurs”.

284
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<th>Species</th>
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<th>Organic matter intake²</th>
<th>Mean retention time³</th>
<th>Organic matter digestibility⁴</th>
<th>Fibre (NDF) digestibility⁵</th>
<th>Mean particle size²</th>
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<td>43.44</td>
<td>7.285</td>
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¹data from Foose (1982)
²data from Fritz et al. (2009)
References


Schwarm, A., Ortmann, S., Wolf, C., Streich, W.J., Clauss, M., 2009. More efficient mastication allows increasing intake without compromising digestibility or necessitating a larger gut: comparative feeding trials in banteng (Bos javanicus) and pygmy hippopotamus (Hexaprotodon liberiensis). Comparative Biochemistry and Physiology A 152, 504-512.


Table 1. Phylogenetic signal and scaling with body mass of mammalian digestive physiology parameters

| Parameter                      | Phylogenetic signal | | | Correlated evolution | | | | | | |
|-------------------------------|---------------------|-----------------|-----------------|-----------------|-----------------|-----------------|
|                               | $\lambda$ | $L_h(\lambda)$ | $L_h(\lambda=0)$ | P-value | $L_h(\lambda,r=0)$ | P-value |
| Digesta retention             | 0.95     | 21.8            | 8.85            | <0.0001 | 21.3            | 0.32             |
| Mean particle size            | 0.97     | -4.3            | -16.1           | <0.0001 | -10.0           | 0.0007           |
| Fibre digestibility           | 0.99     | 34.9            | 20.3            | <0.0001 | 34.6            | 0.44             |
| Relative organic matter intake| 1.02     | 17              | -0.6            | <0.0001 | 16.5            | 0.32             |
| Relative digestible organic matter intake | 1.00 | 15.1 | 1.85 | <0.0001 | 14.3 | 0.21 |

Notes: Tests of phylogenetic signal compare likelihoods ($L_h$) for a model in which $\lambda$ is estimated to a model in which $\lambda$ was forced to equal zero; in both models, we estimated the correlation between traits. For tests of correlated evolution (last two columns), we further developed a model in which $\lambda$ was estimated and the covariance between traits was forced to equal zero. In this case, we compared the model to the likelihood score from $\lambda$ estimated, i.e., $L_h(\lambda)$, in a likelihood ratio test.
Table 2. Multiple linear regression according to Fibre digestibility (%) = a + b logBody mass + c Relative organic matter intake + d Digesta retention time + e logFaecal particle size. A PGLS model with $\lambda=0$ is equivalent to a non-phylogenetic test.

<table>
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<tr>
<td>Faecal particle size</td>
<td>-0.07</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Table 3. Multiple linear regression according to Relative digestible organic matter intake = a + b logBody mass + c Relative organic matter intake + d Digesta retention time + e logFaecal particle size. A PGLS model with $\lambda=0$ is equivalent to a non-phylogenetic test.

<table>
<thead>
<tr>
<th>Factor</th>
<th>PGLS ($\lambda=0$)</th>
<th>PGLS ($\lambda=1$)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Beta</td>
<td>p</td>
</tr>
<tr>
<td>Body mass</td>
<td>0.018</td>
<td>0.57</td>
</tr>
<tr>
<td>Relative organic matter intake</td>
<td>1.03</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Digesta retention time</td>
<td>0.181</td>
<td>0.001</td>
</tr>
<tr>
<td>Faecal particle size</td>
<td>-0.045</td>
<td>0.06</td>
</tr>
</tbody>
</table>
Fig. 1. Correlations between a) body mass and digesta retention time ($R=0.18$, $p=0.429$); b) body mass and faecal particle size ($R=0.82$, $p<0.001$); c) body mass and fibre digestibility ($R=-0.20$, $p=0.394$); d) body mass and relative organic matter intake ($R=-0.06$, $p=0.801$); e) body mass and relative digestible organic matter intake ($R=-0.13$, $p=0.563$) in large mammalian herbivores (ruminants: open triangles = true ruminants, grey triangles = camelids; nonruminant foregut fermenter: black square = hippopotamus; hindgut fermenters: open circles = equids, grey circles = rhinoceroses, black circles = elephants; statistics for raw data; results of maximum likelihood [PGLS] methods in Table 1).
Fig. 2. Correlations between a) digesta retention time and fibre digestibility ($R=0.77$, $p<0.001$); b) faecal particle size and fibre digestibility ($R=-0.56$, $p=0.009$) in large mammalian herbivores (ruminants: open triangles = true ruminants, grey triangles = camelids; nonruminant foregut fermenter: black square = hippopotamus; hindgut fermenters: open circles = equids, grey circles = rhinoceroses, black circles = elephants; statistics for raw data; results of maximum likelihood [PGLS] methods in text)

Fig. 3. Correlation between organic matter intake and digesta retention time ($R=-0.65$, $p=0.002$) in large mammalian herbivores (ruminants: open triangles = true ruminants, grey triangles = camelids; nonruminant foregut fermenter: black square = hippopotamus; hindgut fermenters: open circles = equids, grey circles = rhinoceroses, black circles = elephants; statistics for raw data; results of maximum likelihood [PGLS] methods in text)