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Observations

Enrichment of ¹³C and ¹⁵N against diet





Trophic level enrichment

$\delta^{13}C$	+0.4±0.12‰	(n=111)
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 δ^{15} N +2.0±0.20‰ (n=83)

 $\delta^{34}S$ $+0.4\pm0.52\%$ (n=13)

Dietary information (δ^{13} C) versus trophic level information ($\delta^{15}N$)

BUT: large spread of values

McCutchan et al. 2003 Oikos 102: 378-390

Food webs – trophic level enrichment in ¹³C and ¹⁵N





Arctic marine foodweb (NW Territories, Canada)

Foodweb base: POM, particulate organic matter (algae, bacteria, dead matter) and kelp/ice algae

Up to top predator: polar bear

Hobson & Welch 1992 MEPS 84:9

Food webs – trophic level enrichment in ¹⁵N

Number of trophic levels in an ecosystem calculated by dividing the full range of $\delta^{15}N$ by TLE (3‰ here)



Fig. 3. Ranges of δ¹⁵N values for marine organisms from Barrow Strait-Lancaster Sound and their associated trophic positions according to an isotopic model using a trophic enrichment value of + 3.8 ‰ (not applicable to marine birds)

Hobson & Welch 1992 MEPS 84:9

¹⁵N Amino acid enrichment – nonessential versus essential amino acids

δ^{15} N OF AMINO ACIDS IN PLANKTON

	First sampling		Second sampling	
$\delta^{15}N$ source	<i>T. suecica</i> (Food source)	B. plicatilis (Consumer)	<i>T. suecica</i> (Food source)	B. plicatilis (Consumer)
Bulk sample	-1.9	0.1	-1.7	-0.2
Amino Acids Alanine Aspartic acid Glutamic acid Glycine Isoleucine† Leucine† Lysine† Phenylalanine† Proline Serine	$\begin{array}{c} 0.5 \pm 0.7 \\ -0.5 \pm 0.3 \\ -0.4 \pm 0.2 \\ -4.7 \pm 0.4 \\ -0.8 \pm 0.5 \\ -0.4 \pm 0.8 \\ -1.1 \pm 0.4 \\ -5.3 \pm 0.2 \\ 0.1 \pm 0.3 \\ -7.8 \pm 0.6 \end{array}$	$\begin{array}{r} 4.8 \pm 0.1 \\ 4.6 \pm 0.2 \\ 6.5 \pm 0.4 \\ -3.2 \pm 0.4 \\ 3.3 \pm 0.1 \\ 2.8 \pm 0.1 \\ 0.7 \pm 0.4 \\ -4.2 \pm 0.2 \\ 4.3 \pm 0.2 \\ -7.1 \pm 0.2 \end{array}$	$\begin{array}{r} -1.6 \pm 0.1 \\ -0.6 \pm 0.4 \\ -0.1 \pm 0.3 \\ -4.4 \pm 0.3 \\ -1.3 \pm 0.9 \\ -1.5 \pm 0.3 \\ -1.8 \pm 0.2 \\ -4.3 \pm 0.4 \\ -0.4 \pm 0.2 \\ -7.8 \pm 0.3 \end{array}$	$\begin{array}{r} 4.0 \pm 0.4 \\ 3.2 \pm 0.4 \\ 6.4 \pm 0.3 \\ -4.1 \pm 0.8 \\ 2.4 \pm 0.4 \\ 2.0 \pm 0.3 \\ -0.4 \pm 1.2 \\ -4.8 \pm 1.0 \\ 3.3 \pm 0.3 \\ -6.9 \pm 0.3 \end{array}$
Threonine† Tyrosine Valine†	-2.4 ± 0.4 -2.2 ± 0.6 1.8 ± 0.3	-3.4 ± 0.4 -3.3 ± 0.6 4.5 ± 0.1	-2.3 ± 0.1 -2.5 ± 0.3 1.0 ± 1.5	-4.1 ± 0.5 -3.1 ± 0.1 5.6 ± 0.3

TABLE 1. Values of $\delta^{15}N$ of bulk samples and amino acids of *B. plicatilis* and *T. suecica*.

Notes: Samples were collected on two successive days after determining that bulk values were at a steady state. Values for bulk samples are from CFIRMS analysis (standard deviation is typically \pm 0.2%). Values for amino acids are means \pm 1 SE from three analyses of each sample by GCIRMS. Aspartic acid includes asparagine. Glutamic acid includes glutamine.

[†] Amino acids commonly required in animal diets (essential amino acids.)

McClelland & Montoya 2002 Ecology 83:2173

¹⁵N Amino acid enrichment – nonessential versus essential amino acids



FIG. 1. Differences in stable N isotope values ($\Delta\delta^{15}$ N) of bulk material and individual amino acids between consumer (*B. plicatilis*) and food source (*T. suecica*). Points are means from two sampling dates, and error bars reflect the range of values from *B. plicatilis* (on points) and *T. suecica* (on zero line). Amino acids are grouped into those that show changes greater than the bulk material and those that change less than the bulk material. Abbreviations are: Ala, alanine; Asp, aspartic acid; Glu, glutamic acid; Ile, isoleucine; Leu, leucine; Pro, proline; Val, valine; Gly, glycine; Lys, lysine; Phe, phenylalanine; Ser, serine; Thr, threonine; Tyr, tyrosine. ¹⁵N enrichment in "non-essential" amino acids between 4 and 7 permil (metabolic N recycling)

No such enrichment in "essential" amino acids (isotopic signature of food source)

 $\Delta\delta^{15}$ N Glu – Phe approx. 7 permil per trophic level i.e. a measure of trophic level enrichment without knowing the exact food source and its isotopic composition

> McClelland & Montoya 2002 Ecology 83:2173

Animal diet – two-isotope mixing models to calculate fractions of mixed diets

Partitioning of wolf diets



C₃ versus C₄ diets – Browsers and grazers



Fig. 3 Estimated diets for modern *Loxodonta* and *Elephas* compared to the estimated diets for fossil proboscideans. Modern data from Figs. 1, 2. Fossil data for Africa from Table 2; fossil data from Pakistan from Stern et al. (1994), Morgan et al. (1994), and from this study

Cerling et al. 1999 Oecologia 120: 364



Fig. 1 Histograms showing δ^{13} C for modern plants, and estimated diets calculated from tooth enamel or collagen for modern elephants and for modern African herbivores using fractionation as described in the text. Data sources are this study, and from Sukumar et al. (1987), van der Merwe et al. (1988, 1990), Vogel et al. (1990a, b), Sukumar et al. (1987), Sukumar and Ramesh (1992, 1995), Koch et al. (1995), and Bocherens et al. (1996b)

Isotopic differences between tissues



Fir, bones, enamel, and horn are isotopically distinct from diet

Roth & Hobson 2000 Can. J. Zool. 78: 848

Tissue turnover and time integration of dietary isotope signals

Different dietary time integration of tissues such as blood, liver, muscles, bone collagen, hair keratin of gerbil (<u>Wüstenrennmaus</u>)

Gerbil Tissue Isotope Shift After Diet Change -10 -12 Starting Diet (Corn) Musck Brain Hair Hair $(t_{1/2} = 47 \text{ d})$ Brain (28 d) -20 -22 Muscle (27 d) -24 Liver (6 d) -26 Fat (15 d) 0 20 40 60 80 100 120 140 160 Days Since Diet Change Figure 4. When a gerbil's diet is switched to food with a different 513C value, its tissues change over time to reflect this. This happens as carbon in the tissues is replaced. Some tissues show this change more rapidly than others. For example, liver tissue achieves the 513C value of the new diet in less than 80 days, whereas hair takes well over twice as long.



Herbivore diet and N losses



Deamination/transamination remove ¹⁴N preferentially that is excreted in urine and enriches ¹⁵N in body protein (key enzymatic step unknown)

Herbivore diet and diet protein content



Figure 2. Box plot showing δ^{15} N values for the alfalfa diet and six mammalian herbivore species that had been eating this diet. The range in trophic enrichment is between 2.8‰ and 6.4‰. Note the difference between the alpacas and rabbits is 3.6‰, more than enough to be attributed to a shift in trophic level. The hindgut fermenting rabbits have lower δ^{15} N values than all of the foregut fermenters, possibly because they digest a lower percentage of their microflora.



Figure 3. Box plot showing diet-hair fractionation (Δ_{d-h}) for herbivores eating high-protein alfalfa diets and low-protein coastal bermudagrass (CBG) diets. Note the consistently higher diet-hair fractionation for animals eating the high-protein alfalfa.

> Alfalfa 19% crude protein Bermuda grass 9% crude protein

> > Sponheimer et al. (2003) Int. J. Osteoarchaeol. 13: 80

Herbivore ¹⁵N enrichment with low and high protein diets





You are what you eat, plus a few permilles?

Mr. T.E. Cerling goes west (from USA to mongolia)..... Changing food - changing beard C isotope signatures

or You are what you eat ?

Hair as a chronological archive

+ continuous growth (0.4 mm/day)

- + biologically inactive after formation
- + resistant to degradation



Figure 1: This figure plots the carbon isotopic composition of beard hair (expressed as δ^{13} C) vs date that the hair was collected. The kinks in the Mongolian time lines correspond to the six-day time lag between hair formation and its eruption through the skin surface.

Human diet – The human food web



FIGURE 8. The distribution of carbon and nitrogen isotopes in a human food web. Open circles: Japanese foods. (From Minagawa, M. et al., *Chikyuu-Kagaku*, 2,79, 1986. With permission.) Closed circles: American foods. (Adapted from Schceller, D. A. et al., *Ecol. Food Nutrit.*, 18, 159, 1986.)

Herbivorous versus omnivorous humans -Hair isotopic signatures



Fig. 2. Mean isotopic analyses for each dietary preference group.

Macko et al. (1999) Phil. Trans. R. Soc. London B 354: 65