

NOTE

Nitrogen isotopic composition of marine and freshwater invertebrates

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ABSTRACT: Stable nitrogen isotopes have customarily been used to delineate trophic position with only scant regard to source variability in isotopic composition. A compilation of literature data indicates, however, that marine invertebrates are enriched in ^{15}N relative to those inhabiting freshwaters. Estuarine mussels were also found to have intermediate $\delta^{15}\text{N}$ values which reflected their location along a freshwater-marine gradient. The use of invertebrate $\delta^{15}\text{N}$ as a measure of continental-marine coupling in complex coastal environments could therefore provide additional support for conclusions based on traditional $\delta^{13}\text{C}$ analysis.

KEY WORDS: $\delta^{15}\text{N}$ · Freshwater and marine invertebrates · Source enrichment

It is now widely accepted that a 3 to 4‰ fractionation in stable nitrogen isotopes occurs with food assimilation (Owens 1987). Some parallel work has indicated, however, that $\delta^{15}\text{N}$ (ratio of $^{15}\text{N}/^{14}\text{N}$ expressed as deviations from the recognized isotopic standard, in ‰) may also function as a source marker of material flow across ecotones. In oceans, for example, variability in $\delta^{15}\text{N}$ among animals depends upon process differences in internal cycling of 'old' and 'new' nitrogen (e.g. Rau 1981, Mullin et al. 1984, Checkley & Entzeroth 1985, Fisher et al. 1994). In estuarine and coastal environments, mixing of materials derived from terrestrial and oceanic sources has been assessed through $\delta^{15}\text{N}$ analysis of both suspended (Mariotti et al. 1984, Owens 1985, Croft et al. 1988) and deposited (Peters et al. 1978, Owens 1987) organic matter.

Broad-scale, continental-marine differences in $\delta^{15}\text{N}$ may also exist. The bone collagen of 64 mammals, 17 birds, and 17 fishes showed that marine animals have $\delta^{15}\text{N}$ values which average 6 to 10‰ higher (^{15}N enriched) than those for terrestrial or freshwater organisms (Schoeninger & DeNiro 1984). The low sample sizes (98 measurements) and narrow faunal

range (66 vertebrate species) of this previous study limits conclusions about the generality of broad-scale environmental differences in source variability of $\delta^{15}\text{N}$ [see Owens (1987) and Gearing (1988) for different interpretations]. To counter these limitations, the present investigation compiled literature data on over 400 measurements of $\delta^{15}\text{N}$, representing hundreds of different invertebrate species, to determine if marine animals are indeed enriched in ^{15}N relative to organisms inhabiting freshwaters.

Stable nitrogen values were obtained from tables and from figures with a digitizing reader. Data sources included Wada & Hattori (1976), Pang & Nriagu (1977), Macko et al. (1982), Checkley & Entzeroth (1985), Peterson & Howarth (1987), Yoshioka et al. (1989), Montoya et al. (1990), Rau et al. (1990, 1991), Toda & Wada (1990), and Mihuc & Toetz (1994), in addition to those studies listed in France (1995).

Invertebrate $\delta^{15}\text{N}$ was found to reflect both trophic-dietary and habitat-source fractionation. Over half of the organisms had $\delta^{15}\text{N}$ values within a single trophic position of one another (i.e. $\pm 2\%$) as centred about their respective modes in each data grouping (Figs. 1 & 2). The relative position of these modes, however, were found to be different for marine and freshwater invertebrates. Marine zooplankton (mode and mean = 10‰) and zoobenthos (mode and mean = 9‰) were on average enriched about 3 to 4‰ in $\delta^{15}\text{N}$ compared to freshwater zooplankton (mode = 6‰, mean = 7‰) and zoobenthos (mode and mean = 6‰). Therefore, Schoeninger & DeNiro's (1984) belief in broad-scale environmental differences in $\delta^{15}\text{N}$ between marine and freshwater animals, which they based on collagen samples of 17 fishes, is supported by the present compilation of whole-body samples for 443 invertebrates.

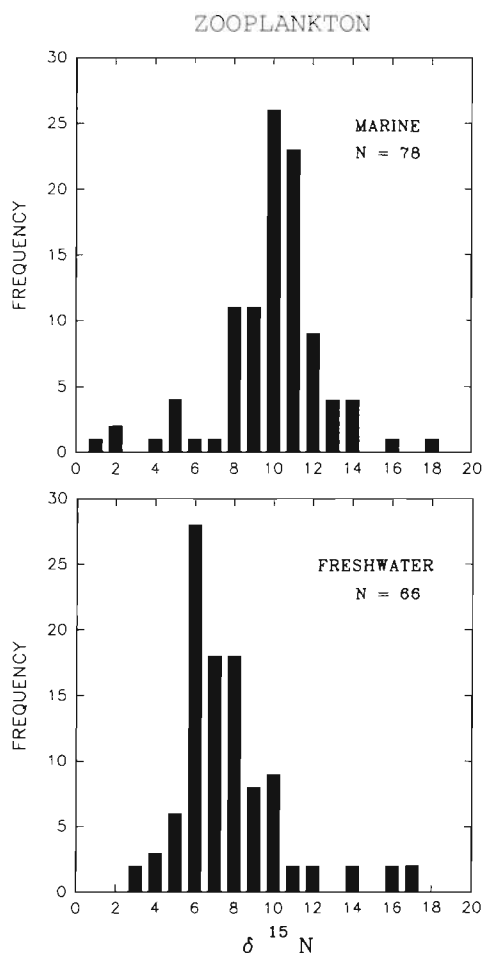


Fig. 1. Percentage frequency distributions of stable nitrogen isotope ratios for marine and freshwater zooplankton

Within a single species, analysis of $\delta^{15}\text{N}$ can be used to measure the dietary proportion of food originating from different environmental sources. For example, a reanalysis of mussel data from Peterson et al. (1985) indicates that the $\delta^{15}\text{N}$ of estuarine invertebrates reflects the degree of terrestrial-oceanic mixing (Fig. 3). This substantiates Schoeninger & DeNiro's (1984) finding of intermediate $\delta^{15}\text{N}$ values for organisms such as migratory birds and anadromous fishes which spend part of their life cycles feeding in both freshwater and marine environments.

The use of invertebrate $\delta^{15}\text{N}$ as a marker of ecotonal coupling in coastal environments could therefore provide additional support for conclusions based on the traditional analysis (e.g. Incze et al. 1982, Stephenson & Lyon 1982) of $\delta^{13}\text{C}$ alone.

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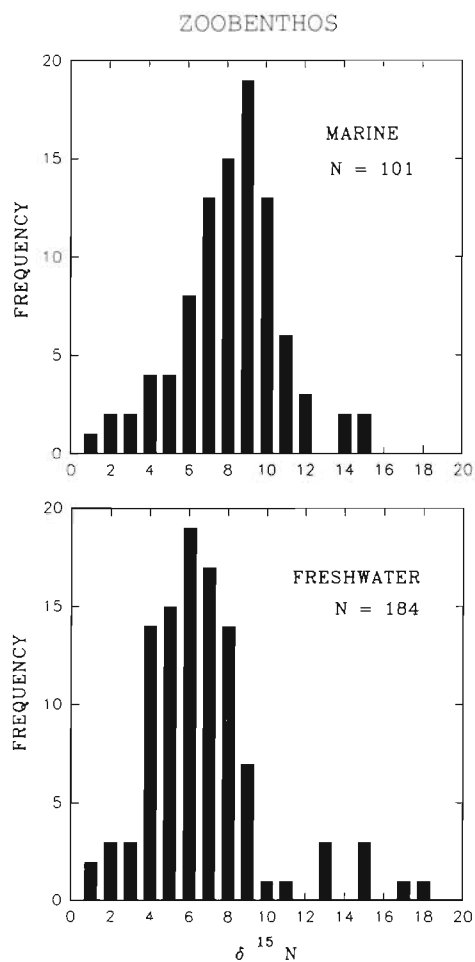


Fig. 2. Percentage frequency distributions of stable nitrogen isotope ratios for marine and freshwater zoobenthic invertebrates

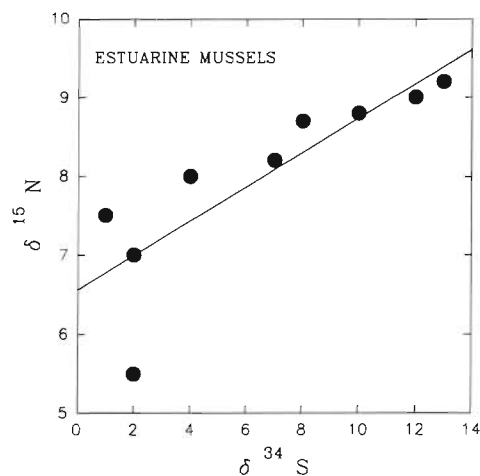


Fig. 3. Relationship between stable isotopes of nitrogen and sulfur for mussels (*Geukensia demissa*) in an estuary (re-analysis of data from Peterson et al. 1985). The sulfur isotopic signature for marine sediments is about 15‰ while that for freshwater sediments is below 0

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