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Tracing Biogenic Links of Natural Organic Substances at the Molecular Level with Stable Carbon Isotopes : *n*-Alkanes and *n*-Alkanoic Acids from Sediments

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Abstract: the $^{13}\text{C}/^{12}\text{C}$ compositions of *n*-alkanes and *n*-alkanoic acids from Eocene sediments are correlated in the 20-29 carbon number range.

n-Alkanes are an important class of natural compounds occurring almost ubiquitously in organic-rich sediments such as soils, petroleums and coals¹. Although extensively studied, the origin of sedimentary *n*-alkanes is still unclear, mainly because of the lack of structural information of these substances. They might derive from biological aliphatic precursors such as *n*-alkanes, *n*-alkohols, *n*-alkanoic acids or biopolymers¹⁻³. Recent studies suggest that $^{13}\text{C}/^{12}\text{C}$ compositions of molecular fossils could provide more information about their possible precursors in living organisms⁴. We would like now to report the correlation of *n*-alkanes and *n*-alkanoic acids from Eocene sediments by isotopic means.

The geological samples studied belong to the lacustrine Eocene formation (60×10^6 years) of Green River (Colorado, USA) and are located at depths varying between 300 m and 800 m. Separation procedures and isotopic analysis have been previously described^{4,5}. The following isotopic compositions were obtained (carbon number, δ ‰):

At 312 m depth: *n*-alkanes: (17, -36.8); (18, -31.7); (20, -31.9); (21, -30.2); (22, -32.0); (23, -30.0); (24, -31.0); (25, -30.5); (26, -29.9); (27, -29.8); (28, -29.6). *n*-Alkanoic acids: (18, -30.0); (19, -31.6); (20, -31.4); (21, -31.9); (22, -31.3); (23, -33.6); (24, -31.5); (25, -31.8); (26, -30.2); (27, -30.3); (28, -29.2); (29, -28.6).

At 778 m depth: *n*-alkanes: (17, -32.4); (18, -32.2); (20, -32.2); (21, -31.9); (22, -31.9); (23, -31.6); (24, -31.1); (25, -31.1); (26, -31.2); (27, -31.2); (28, -30.4). *n*-Alkanoic acids: (18, -29.2); (19, -32.6); (20, -33.2); (21, -32.6); (22, -32.5); (23, -32.6); (24, -32.3); (25, -32.1); (26, -32.2); (27, -32.1); (28, -32.3); (29, -32.1).

Fig. 1. Isotopic compositions of *n*-alkanes and *n*-alkanoic acids at 312 m (left) and 718 m depth (right).

In the 20-29 carbon number range, the isotopic compositions of *x* carbon-numbered *n*-alkanes and *x*+1 carbon-numbered *n*-alkanoic acids are close and display similar variations with increasing carbon number (Figure 1). Since various carbon sources with various isotopic compositions are available along the food chain, this correlation between two classes of organic substances suggests their biogenic link. Moreover, *n*-alkanes, which are seldom in living organisms, could derive from *n*-alkanoic acids occurring ubiquitously in eukaryotes and eubacteria as triglyceride esters. This correlation also suggests the formation of alkane *via* decarboxylation², accounting probably for the large amounts of carbon dioxide produced in sediments¹. Alternatively, the high isotopic composition difference observed between *n*-heptadecane and *n*-octadecanoic acid could arise from a different pathway of formation.

REFERENCES

1. Tissot, B. P., Welte, D. H. *Petroleum Formation and Occurrence*; Springer: Berlin. 1984.
2. Cooper, J. E., Bray, E. E. *Geochim. Cosmochim. Acta* **1963**, 27, 1113-1127. Behar, F. H., Albrecht, P. *Org. Geochem.* **1984**, 6, 597-604.
3. Tegelaar, E. W., Matthezing, R. M., Jansen, J. B. H., Horsfield, B., de Leeuw, J. W. *Nature* **1989**, 342, 529-531.
4. Hayes, J. M., Takigiku, R., Ocampo, R., Callot, H. J., Albrecht, P. *Nature* **1987**, 329, 48-51. Freeman, K. H., Hayes, J. M., Trendel, J. M., Albrecht P. *Nature* **1990**, 343, 254-256.
5. Lichtfouse, E., Freeman, K. H., Collister, J. W., Merritt, D. A., *J. Chromatogr.* **1991**, 585, 177-180 and refs. therein.