S_{obs} is the 1.40.4072/m s 4042007400

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Fig. S1. Isotopic profiles of POM (filled circles) of zooplankton (open symbols) in Paul (Left) and Crampton (Right) lakes. In Paul L., only Daphnia (open circles) is shown because Daphnia and Holopedium were indistinguishable at this scale. In Crampton L., Holopedium (open circles) and Leptodiaptomus (▽) are both shown. Error bars at each represent the temporal SD of samples taken four times during 2009. In Paul L., both zooplankton taxa were measured at all depths on all four dates. In Crampton L., we did not measure both taxa at all depths or on all dates because in some cases, taxa were too rare to obtain sufficient mass for isotope analyses. Thus, samples without SD are from single measures. The approximate position of terrestrial isotope values is shown for each isotope; the approximate value for the δ^2 H of phytoplankton is shown in Upper.

Fig. S2. Results of fitting dietary water in IsoSource. Upper shows the distributions of dietary water, fitted by IsoSource in models that included all three isotopes (C, N, and H) and four end members as possible sources (terrestrial, surface phytoplankton, deep phytoplankton, and benthic algae). Lower shows the distribution of the mean values of dietary water, fitted by IsoSource, for all of the three isotope models shown in Fig. 3.

Table S1. Reported values of allochthony for zooplankton in freshwater systems

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The table is divided into two parts: Qualitative estimates and Quantitative estimates, and then ordered chronologically. "Qualitative" means that the author provided some written assessment of allochthony as large or small but did not report a numerical value. "Support" means that either the qualitative or quantitative results of the study support (Yes) or do not support (No) the hypothesis that zooplankton are subsidized by terrestrial organic matter.

1. del Giorgio PA, France RL (1996) Ecosystem-specific patterns in the relationship between zooplankton and POM or microplankton delta C-13. Limnol Oceanogr 41:359–365.

- 2. Sobczak WV, Cloern JE, Jassby AD, Müller-Solger AB (2002) Bioavailability of organic matter in a highly disturbed estuary: The role of detrital and algal resources. Proc Natl Acad Sci USA 99:8101–8105.
- 3. Karlsson J, Jonsson A, Meili M, Jansson M (2004) delta N-15 of zooplankton species subarctic lakes in northern Sweden: Effects in s of diet and trophic fractionation. Freshw Biol 49: 526–534.
- 4. Perga ME, Bec A, Anneville O (2009) Origins of carbon sustaining the growth of whitefish Coregonus lavaretus early larval stages in Lake Annecy: Insights from fatty-acid biomarkers. J Fish Biol 74:2–17.
- 5. Meili M, et al. (1996) Sources and partitioning of organic matter in a pelagic microbial food web inferred from the isotopic composition δ¹³ C and δ¹⁵ N) of zooplankton species. Arch Hydrobiol Spec Issues Advanc Limnol 48:53–61.
- 6. Jones RI, Grey J, Sleep D, Quarmby C (1998) An assessment, using stable isotopes, of the importance of allochthonous organic carbon source to the pelagic food web in Loch Ness. Proc Biol Sci 265:105–111.
- 7. Cole JJ, Carpenter SR, Kitchell JF, Pace ML (2002) Pathways of organic carbon utilization in small lakes: Results from a whole-lake C-13 addition and coupled model. Limnol Oceanogr 47: 1664–1675.

8. Cole JJ, et al. (2006) Differential support of lake food webs by three types of terrestrial organic carbon. Ecol Lett 9:558-568.

- 9. Karlsson J, Jonsson A, Meili M, Jansson M (2003) Control of zooplankton dependence on allochthonous organic carbon in humic and clear-water lakes in northern Sweden. Limnol Oceanogr 48:269–276.
- 10. Carpenter SR, et al. (2005) Ecosystem subsidies: Terrestrial support of aquatic food webs from C-13 addition to contrasting lakes. Ecology 86:2737–2750.

11. Matthews B, Mazumder A (2006) Habitat specialization and the exploitation of allochthonous carbon by zooplankton. Ecology 87:2800–2812.

12. Pace ML, et al. (2007) Does terrestrial organic carbon subsidize the planktonic food web in a clear-water lake? Limnol Oceanogr 52:2177–2189.

13. Taipale S, Kankaala P, Tiirola M, Jones RI (2008) Whole-lake dissolved inorganic 13C additions reveal seasonal shifts in zooplankton diet. Ecology 89:463–474.

14. Mohamed MN, Taylor WD (2009) Relative contribution of autochthonous and allochthonous carbon to limnetic zooplankton: A new cross-system approach. Fund Appl Limnol 175: 113–124.

15. Berggren M, et al. (2010) Lake secondary production fueled by rapid transfer of low molecular weight organic carbon from terrestrial sources to aquatic consumers. Ecol Lett 13: 870–880.

16. Caraco N, Bauer JE, Cole JJ, Petsch S, Raymond P (2010) Millennial-aged organic carbon subsidies to a modern river food web. Ecology 91:2385–2393.

Table S2. Limnologic conditions in Paul and Crampton Lakes, Vilas County, WI

Color is a measure of light absorption at 400 nM. DOC, dissolved organic C; POC, particulate organic C. Z_{mix} is the depth of the upper mixed layer; Z_{oxic} is the depth at which anoxic water starts. Paul Lake is entirely feed by ground water and Crampton mostly so, except for an intermittent stream. The watersheds of both lakes are entirely forested and undeveloped; watershed areas from Cardille et al. (1). For both lakes, the terrestrial C can enter these lakes as DOC in groundwater, as Aeolian-transported POC, and POC can be formed in the lake from DOC by flocculation.

1. Cardille JA, et al. (2007) Carbon and water cycling in lake-rich landscapes: Landscape connections, lake hydrology, and biogeochemistry. J Geophys Res-Biogeo 112:G202031.

The terrestrial end members represent averages of leaves of the dominant trees in these watersheds and were measured directly. Benthic algal isotopes were obtained by deploying clean tiles for 1–2 wk in each lake and sampling the algae that grew there and measured directly. Phytoplankton isotopes were obtained by a unique method. Using dilution regrowth experiments in each lake, we obtained the δ^2 H of phytoplankton. The contrast between the δ^2 H of phytoplankton and the water it grew in (ε_Η,) was large, and nearly identical (between 150 and 160 del units) between the lakes and in other systems so far studied (1–4). Unlike the case for C, there is no known physiological effect on ε_H of bulk organic matter from either growth rate or cell size. Thus, it is possible to predict the δ^2 H of phytoplankton from that of water in principal under any growth conditions. Using the measured δ^2 H of water at each depth and time, we calculated the δ^2 H of phytoplankton by using the mean $\varepsilon_{\rm H}$. We assume that POM is a mixture of phytoplankton and terrestrial material. Because we have measured values of the δ^2 H, δ^{13} C, and δ^{15} N of terrestrial material, we can solve algebraically for the 13 C and 15 N of phytoplankton. Solving Eq. 1 for φ_T gives us the fraction of POM that is terrestrial. We can then solve Eq. 2 for the δ^{13} C (or a similar equation for δ^{15} N) of phytoplankton. Eq. 1: δ^2 H $_{\rm POM}$ = $\phi_{\rm T}\times\delta^2$ H $_{\rm T}$ + (1 – $\phi_{\rm T})$ δ²H_P (all are knowns except ϕ_T) Eq. 2: δ¹³C_{POM} = ϕ_T × δ¹³C_T+ (1 – ϕ_T) × δ¹³C_p (now all are knowns except δ¹³C_P), where the subscripts "POM" and "P" denote particulate organic matter and phytoplankton, respectively, and ϕ_T is the fraction of POM that is terrestrial.

1. Caraco N, Bauer JE, Cole JJ, Petsch S, Raymond P (2010) Millennial-aged organic carbon subsidies to a modern river food web. Ecology 91:2385–2393.

2. Doucett RR, Marks JC, Blinn DW, Caron M, Hungate BA (2007) Measuring terrestrial subsidies to aquatic food webs using stable isotopes of hydrogen. Ecology 88:1587–1592.

3. Finlay JC, Doucett RR, McNeely C (2010) Tracing energy flow in stream food webs using stable isotopes of hydrogen. Freshwat Biol 55:941–951.

4. Solomon CT, et al. (2009) The influence of environmental water on the hydrogen stable isotope ratio in aquatic consumers. Oecologia 161:313–324.

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