relation between terrestrial C and atmospheric CO₂. Assessment of modern terrestrial inventories alone will not provide sufficient information to predict potential future sinks and sources for atmospheric CO_2 . One must also base such predictions on time-series data and process studies to determine the effects of landscape alterations, including those caused by human activities (2).

REFERENCES AND NOTES

- W. H. Schlesinger, *Nature* **348**, 232 (1990).
 R. A. Houghton, J. E. Hobbie, J. M. Melillo, *Ecol.* Monoar. 53, 235 (1983)
- 3. P. P. Tans, I. Y. Fung, T. Takahashi, Science 247, 1431 (1990).
- N. J. Shackleton, in The Fate of Fossil Fuel CO2 in the Oceans, N. R. Andersen and A. Malahoff, Eds. (Plenum, New York, 1977), pp. 401-427.
- J. N. Adams, H. Faure, L. Faure-Denard, J. M. 5 McGlade, F. I. Woodward, Nature 348, 711 (1990).
- 6. H. Jenny, The Soil Resource: Origin and Behavior (Springer-Verlag, New York, 1980).
- 7 D. S. Jenkinson, Philos. Trans. R. Soc. London 329, 361 (1990)
- W. J. Parton, D. S. Schimel, C. V. Cole, D. S. 8 Ojima, Soil Sci. Soc. Am. J. 51, 1173 (1987).
- We use the term accumulation for site-scale data with units of grams per square meter per year and the term sequestration for areally extrapolated data with units of grams per year. 10. A. S. Dyke and V. K. Prest, *Geog. Phys. Quat.* **41**,
- 237 (1987).
- National Átlas of Canada (Canada Department of Energy and Resources: Surveys and Mapping Branch, Ottawa, 1973).
- Soil Conservation Service, in National Atlas of the 12 United States (U.S. Geological Survey, Reston, VA, 1987), pp. 42–43.
- 13. H. Jenny, S. P. Gessel, F. T. Bingham, Soil Sci. 68, 419 (1949).
- 14. A soil chronosequence is a series of soil profiles from sites that have similar factors such as land use, climate, vegetation, slope, and original material but differ in age of the landform (37). Each chronosequence in our study was fit to a curve of the form of Eqs. 1 and 2 with the use of the Marquardt-Levenberg nonlinear regression implementation of Sigma Plot Version 4.0 (Jandel Scientific, Corte Madera, CA, 1990). Chronose-quences were selected where geologic ages of deposits represented age of geomorphic surface and inception of soil development. Chronosequence data for soils other than peats apply to mineral soil below organic leaf-litter layers. 15. J. M. Kimble, H. Eswaran, T. Cook, in International
- Congress of Soil Science (International Congress of Soil Science, Kyoto, 1990), pp. 248-253.
- 16. P. W. Birkeland, Arct. Alp. Res. 10, 733 (1978)
- K. R. Everett, ibid. 3, 1 (1971). 17
- C. Tarnocai, in Global Change, Canadian Wet-18. lands Study, H. I. Schiff and L. A. Barrie, Eds. (Canadian Institute for Research in Atmospheric Chemistry, York University, North York, U.K., 1988), pp. 21–25.
- 19. L. Ovenden, Quat. Res. 33, 377 (1990).
- 20. W. D. Billings, *Quat. Sci. Rev.* **6**, 165 (1987). 21. S. C. Zoltai and D. H. Vitt, *Quat. Res.* **33**, 231 (1990)
- D. Schimel, M. A. Stillwell, R. G. Woodmansee, 22. Ecology 66, 276 (1985)
- 23. S. Trumbore, *Ecol. Appl.*, in press. 24. J. Brown, *Soil Sci. Soc. Am. Proc.* **31**, 686 (1967).
- E. Gorham, Ecol. Appl. 1, 182 (1991). 25.
- D. P. Franzmeier and E. P. Whiteside, Mich. Agric Exp. Stn. Q. Bull. 46, 1 (1963); L. R. Barrett and R. J. Schaetzl, Can. J. Soil Sci., in press.
- 27. F. C. Ugolini, in Soil Development and Ecological Succession in a Deglaciated Area of Muir Inlet,

Southeast Alaska, R. P. Goldthwaite, F. Loewe, F. C. Ugolini, H. F. Decker, D. M. DeLong, Eds. (Ohio State University, Columbus, 1974), pp. 29-56.

- 28. The timing of peat land expansion in northern Asia, around 5 ka or later (25), and the size of peat land in north central Asia (38) were comparable to those of the Laurentide region. The Laurentide Ice Sheet represents about 20% of deglaciated land (39, 40). Peats are the most important C sink. The calculation implies about 3 \times 10¹² m² of peat land in deglaciated regions outside of the Laurentide region. Total world peat inventories are 3×10^{12} to 4×10^{12} m² (25), so our extrapolation overestimates C sequestration by deglacial peats. The exclusion from our inventories of soils formed on continuous permafrost helps to compensate for this overestimate. Carbon is sequestered in many of these soils because of the lack of drainage and limited decomposition. Field studies and better maps would help improve this extrapolation.
- 29. W. H. Schlesinger, Biogeochemistry, An Analysis of Global Change (Academic Press, San Diego, CA. 1992). Preindustrial rates include those of peats and low rates for accumulation for non-peat soils
- 30. S. C. Porter, in Americans Before Columbus: Ice Age Origins, R. C. Carlisle, Ed. (Department of Anthropology, Pittsburgh University, Pittsburgh, PA, 1988), pp. 1-24.

- 31. J. L. Sarmiento and E. Sundquist, Nature 356, 589 (1992)
- A. Neftel, H. Oeschger, J. Schwander, R. Stauffer, R. Zumbrunn, *ibid*. **295**, 220 (1982). 32.
- 33. W. S. Broecker and T. H. Peng, Tracers in the Sea (Lamont Doherty Geological Observatory, Palisades, NY, 1982).
- J. G. Bockheim, Arct. Alp. Res. 11, 289 (1979). 34
- L. J. Evans and B. H. Cameron, Can. J. Soil Sci. 35 59, 203 (1979).
- E. J. Ciolkosz, in Quaternary Deposits and Soils of the Central Susquehanna Valley of Pennsylvania, D. Marchand, E. J. Ciolkosz, M. E. Bucek, G. H. Crowl, Eds. (Agronomy Department, Pennsylva-nia State University, University Park, 1978).
- H. Jenny, Factors of Soil Formation (McGraw-Hill, 37. New York, 1941).
- 38 Food and Agriculture Organization, United Nations Educational, Scientific, and Cultural Organization, Soil Map of the World (1978).
- 39 K. C. Prentice and I. Y. Fung, Nature 346, 48 (1990).
- G. H. Denton and T. Hughes, The Last Great Ice 40. Sheets (Wiley-Interscience, New York, 1981).
- 41. L. E. Jackson, in Quaternary Geology of Greenland and Canada, R. J. Fulton, Ed. (Geological Survey of Canada, Ottawa, 1989), pp. 667-697.
- We thank P. Showalter for work on Geologic 42 Informations Systems and W. Schlesinger, S Trumbore, and C. Tarnocai for reviews

8 July 1992; accepted 2 November 1992

Evidence from the Lamarck Granodiorite for Rapid Late Cretaceous Crust Formation in California

Drew S. Coleman,*† Thomas P. Frost, Allen F. Glazner

Strontium and neodymium isotopic data for rocks from the voluminous 90-million-year-old Lamarck intrusive suite in the Sierra Nevada batholith, California, show little variation across a compositional range from gabbro to granite. Data for three different gabbro intrusions within the suite are identical within analytical error and are consistent with derivation from an enriched mantle source. Recognition of local involvement of enriched mantle during generation of the Sierran batholith modifies estimates of crustal growth rates in the United States. These data indicate that parts of the Sierra Nevada batholith may consist almost entirely of juvenile crust added during Cretaceous magmatism.

Present models for the origin of the Sierra Nevada batholith developed from the strontium isotope studies of Kistler and Peterman (1) and indicate that the isotope systematics of the batholith can be reproduced through mixing between depleted mantle and Proterozoic upper crust (2-5), or by mixing of crustally contaminated mafic magmas and siliceous melts of Proterozoic lower crust (6). These models assume that the regular east/west change in the isotopic composition of granitic rocks in the batholith reflects an eastward increase in the contribution of old continen-

tTo whom correspondence should be addressed.

SCIENCE • VOL. 258 • 18 DECEMBER 1992

tal crust. In particular, the ${}^{87}\text{Sr}/{}^{86}\text{Sr} = 0.706$ isopleth (1), or the ${}^{87}\text{Sr}/{}^{86}\text{Sr} = 0.708$, $\varepsilon_{\text{Nd}} =$ -7 isopleths (3, 7), commonly are interpreted to reveal the approximate western limit of the continental crust under the United States. DePaolo et al. (8) reached similar conclusions regarding the origin of the batholith using Nd model ages for granitic rocks.

All of these models assume a depleted composition for the mantle end member. However, isotope studies in the Basin and Range province of the western United States suggest that parts of eastern California, Nevada, and Utah were underlain by enriched lithospheric mantle during Cenozoic (9–13) and Mesozoic (14) magmatism. Limited data from mantle xenoliths and alkalic lavas also suggest the presence of an enriched mantle component as far west as the central Sierra Nevada [Fig. 1 (15, 16)]. If an enriched mantle was involved in Sierra Nevada batholith formation, estimates of crust for-

D. S. Coleman and A. F. Glazner, Department of Geology, University of North Carolina, Chapel Hill, NC 27599

T. P. Frost, U.S. Geological Survey, West 904 Riverside Avenue Spokane, WA 99201

^{*}Present address: Massachusetts Institute of Technology, Department of Earth, Atmospheric, and Planetary Sciences, 80 Ames Street 54-1016, Cambridge, MA 02139



Fig. 1. Location of the Lamarck intrusive suite. Inset shows location of main figure. Ruled area on inset outlines the approximate limit of the area characterized by Cenozoic igneous rocks with enriched Sr isotopic compositions [modified from the Sierran Province of Menzies *et al.* (*11*)]. Patterned area on the main figure is the outline of the Sierra Nevada batholith.

mation rates during Mesozoic arc magmatism in the western United States could increase significantly over earlier estimates (8) because the isotopic composition of enriched mantle (EM) may be variable and significantly different from that of depleted mantle (DM) [for example, $\epsilon_{\rm Nd}$ (EM) 0 to -11 (11, 12); $\epsilon_{\rm Nd}$ (DM) +8.5 (8)]. No isotopic data have been available for mafic plutonic rocks from the eastern Sierra Nevada batholith to test this possibility.

Here we report Sr and Nd isotope data (17) from the voluminous, compositionally heterogeneous, 90-million-year-old Lamarck intrusive suite in the central Sierra Nevada batholith (Fig. 1). The Lamarck lies at the western limit of the Sierran enriched mantle province of Menzies *et al.* (11), between the Cenozoic Big Pine volcanic field, which contains lavas with an enriched mantle signature (9), and the Big Creek pipe, where Domenick *et al.* (16) collected a mantle xenolith with



Fig. 2. (a) Initial Nd versus Sr for the Lamarck intrusive suite. The isotopic compositions of the mafic rocks are interpreted to be representative of their mantle source whereas the granodiorites are interpreted to reflect derivation from the mafic rocks with variable and apparently minor contamination by older crustal rocks. (b) Nd versus silica concentration. The gabbros and diorites show a tight cluster of Nd isotopic compositions despite a variation of almost 5% SiO₂ by weight. With the exception of one sample, the rocks show a weak positive relation between Nd isotopic composition and SiO₂ concentration. This trend is opposite that predicted by simple mixing of depleted mantle and Proterozoic crust. Solid squares, gabbros and diorites; open circles, granodiorites and granites; solid circles, hybrid inclusions; open squares, cumulus schliere. Error bars are discussed in (17).

enriched mantle characteristics. This area also lies in a gap in the existing Nd data base for the Sierra Nevada batholith between 36°30' and 37°00'N latitude. Field relations and geochemistry of the Lamarck intrusive suite were reported by Frost (18) and Frost and Mahood (19), who concluded that much of the chemical variation among rocks with silica contents between 49 and 64% by weight could be explained by mixing between mafic and felsic end members coupled with concomitant fractional crystallization of both endmember magmas.

Field, petrographic, and chemical data indicate that many mafic schlieren in the Lamarck suite represent cognate accumulations of mafic minerals from the host granodiorite magma similar to those described elsewhere in the Sierra Nevada batholith (20). However, the gabbro and diorite intrusions and inclusions in the suite are not cumulates, but were intruded as magmas that interacted with the felsic magmas of the Lamarck (19). Only one sample (76c) that we analyzed is considered to be a cumulate on the basis of field and petrographic data, and it is clearly chemically distinct from those samples considered to represent magmatic or hybrid compositions [(18, 19) Table 1].

Isotope data for rocks from the Lamarck intrusive suite show little variation despite a wide variation in bulk chemistry, and variation among the mafic rocks is considerably smaller even though analyses were obtained from four different intrusions separated by a total distance of 40 km (Table 1 and Fig. 2). The variation among samples probably reflects open-system interaction of silicic magmas with their wall rocks. Interestingly, there is a general trend (with the exception of one sample) toward decreasing 87 Sr/ 86 Sr ratio and increasing ϵ_{Nd} with increasing silica (Fig. 2).

Table 1. Sr and Nd isotopic compositions of rocks from the Lamarck intrusive suite. Sample names, unit designations and oxide concentrations are from Frost (18) and Frost and Mahood (19). Trace element

concentrations were determined by isotope dilution techniques except for starred (*) samples which have Rb and Sr data from Frost (18). Analytical procedures are described in (17).

Sample	Unit	SiO ₂ (%)	MgO (%)	K ₂ O (%)	Rb (ppm)	Sr (ppm)	(⁸⁷ Sr/ ⁸⁶ Sr),	Sm (ppm)	Nd (ppm)	$\epsilon_{Nd}(t)$
80fG	Mount Gilbert gabbro	50.67	5.69	1.75	66.30	735.98	0.70641	4.98	25.75	-4.20
20b*	Mafic inclusion in 20a	50.97	4.66	2.36	190	400	0.70656	5.36	25.17	-4.70
80c*	Mount Gilbert gabbro	51.63	5.51	1.63	48	850	0.70631	4.66	21.70	-4.72
6c	Mafic inclusion in 6a	52.15	4.87	2.13	110.35	756.71	0.70656	5.89	29.99	-5.63
SAB-1b	Lake Sabrina diorite	52.45	4.68	2.00	100.13	712.04	0.70638	6.79	37.71	-4.75
76c	Schliere in 76a	52.78	7.19	2.35	109.43	251.25	0.70641	17.30	86.68	-4.54
87a	Piute Creek diorite	53.39	5.38	1.93	101.86	582.33	0.70633	4.01	20.27	-4.33
UB-1b*	Upper Basin gabbro	53.55	4.73	1.73	74	630	0.70646	4.13	18.57	-4.94
UB-5	Upper Basin gabbro	54.94	3.12	2.03	55.17	847.46	0.70648	5.20	26.72	-4.79
83c	Mount Gilbert diorite	55.03	3.66	1.96	76.53	809.13	0.70640	7.70	40.64	-4.34
80fL	Lamarck granodiorite	61.99	2.86	1.58	65.17	698.80	0.70636	5.47	32.68	-3.87
76a	Lamarck granodiorite	65.34	2.23	3.70	114.25	466.81	0.70652	4.03	21.37	-4.35
20a*	Lamarck granodiorite	65.39	2.01	3.47	130	445	0.70658	3.67	18.78	-4.62
15a	Lamarck granodiorite	67.95	1.54	3.82	141.45	463.26	0.70679	3.28	19.87	-6.55
6a	Lamarck granodiorite	69.82	1.11	4.50	130.48	379.66	0.70639	3.26	19.98	-3.84
UB-1a	Lamarck granodiorite	70.53	0.91	3.55	102.81	391.05	0.70619	4.08	24.09	-2.91

Fig. 3. Nd isotopic composition versus Nd concentration for mafic and felsic rocks of the Lamarck intrusive suite, with representative mixing models. Data symbols are the same as in Fig. 2. We consider that a mixing origin for the trend of the data is unlikely (see text for discussion). Inset shows trends generated by open and closed system effects on the main diagram. Labeled increments along model curves show weight percent crustal component in the mixture. Model 1 uses the arc tholeiite mantle source of Hill et al. (4) mixed with a crust with $\varepsilon_{Nd} = -15$ and 50 ppm Nd. The concentration of Nd chosen for this contaminant is unusually high, but an extreme



value was used in order to minimize the amount of crustal component necessary to generate the gabbros (~20%). Model 2 uses the mantle and crustal parameters of DePaolo et al. (8) and requires approximately 40% recycled crustal materials in the rocks.

These trends are opposite those predicted by mixing between depleted mantle and Proterozoic crust.

The simplest interpretation of these data is that the isotopic compositions of the mafic rocks reflect the composition of their mantle source, a source that is similar in isotope and trace-element characteristics to that inferred for nearby Cenozoic basalts and xenoliths (9, 15, 16). If this is correct, then the more silicic members of the Lamarck series could be produced by commonly invoked petrologic processes [for example, coupled assimilation-fractional crystallization, or partial melting of the mafic members of the series (13, 21)]. The lack of a significant ancient crustal component in the Lamarck implied by this interpretation is supported by the apparent lack of inherited zircons in the granodiorite (22).

Alternatively, isotope data for the mafic rocks could be explained by cryptic contamination (23) of mafic magma derived from depleted mantle. Figure 3 shows two possible mixing scenarios, in which depleted mantle magmas are contaminated by mafic gneisses with variable Nd concentrations and isotope ratios. In either scenario, mixing of depleted mantle magma with 20 to 40% contaminant by weight, followed by closed system fractional crystallization, could reproduce the Nd concentration and isotope ratio data.

We view such an origin for the Lamarck suite as plausible but unlikely for the following reasons. First, for this model to explain the entire data set, a huge volume of mafic magma (enough to generate all of the rocks in the Lamarck intrusive suite) must be emplaced into the crust and experience significant and uniform contamination by a component with a very negative $\epsilon_{\rm Nd}$ (24). The magma must then undergo almost no further crustal contamination: formation of the silicic end of the series would have to occur by closed system fractional crystallization alone (unless the contaminant fortuitously had isotopic compositions identical to those of the contaminated mafic magma). Second, for the fractional crystallization model to work requires that

 $D^{\rm Nd} << 1$ in order to form the Nd-enriched mafic to intermediate magmas, followed by further closed system differentiation with $D^{\rm Nd}$ \geq 1 to form the lower Nd silicic members of the suite (25). This scenario is plausible but fails to account for other geochemical variations; in particular, in the gabbros and diorites, Nd concentration varies by a factor of 2, but the concentration of K_2O , which would have D << 1 during fractional crystallization, is essentially constant in these rocks (Table 1). Furthermore, as discussed above, field and petrographic data argue against a cumulus origin for the mafic intrusions (19).

If the Lamarck suite was derived from an enriched mantle with $\epsilon_{Nd}\approx-4.5$ and an $^{87}Sr/^{86}Sr$ ratio of 0.7064, then the amount of juvenile material it represents may approach 100%. This possibility could locally double estimates of the amount of new crust generated during Mesozoic magmatism over earlier estimates (8). If further studies support widespread involvement of enriched mantle in generation of the Sierra Nevada batholith, then the Mesozoic (in particular, the Late Cretaceous) may have been a time of exceedingly rapid crust formation in the western United States.

REFERENCES AND NOTES

- R. W. Kistler and Z. E. Peterman, *Geol. Soc. Am.* Bull. 84, 3489 (1973); U.S. Geol. Surv. Prof. Pap 1071 (1978).
- D. J. DePaolo, J. Geophys. Res 86, 10470 (1981) З. G. L. Farmer and D. J. DePaolo, ibid. 88, 3379
- (1983)M. Hill, J. R. O'Neil, H. Noyes, F. A. Frey, D R. 4.
- Wones, Am. J. Sci. 288-A, 213 (1988)
- J H. Chen and G R. Tilton, Bull. Geol. Soc. Am 5. 103, 439 (1991) 6.
- R. W. Kistler, B. W. Chappell, D. L. Peck, P. C Bateman, Contrib. Mineral. Petrol. 94, 205 (1986). 7
- Nd isotope ratios are presented in the ϵ notation (2), where $\epsilon_{Nd}t$ is defined as $[^{143}Nd/^{144}Nd_{sample}(t)]/[^{143}Nd/^{144}Nd_{CHUR}(t)] 1\} \times 10000$, where t is the age of the rock, and assuming that present-day values for CHUR (chondritic uniform reservoir) are ¹⁴³Nd/¹⁴⁴Nd_{CHUR} = 0.512638 and ¹⁴⁷Sm/¹⁴⁴ Nd_{CHUR} = 0.1967.
 B. J. DePaolo, A. M. Linn, G. Schubert, *J. Geo-*
- phys. Res. 96, 2071 (1991)
- W. P. Leeman, Geochim. Cosmochim. Acta 34, 857 (1970); W P. Leeman, Bull. Geol. Soc Am. 93, 487 (1982); D. S. Ormerod, C. J. Hawkesworth, N. W.

SCIENCE • VOL. 258 • 18 DECEMBER 1992

Rogers, W. P. Leeman, M. A. Menzies, Nature 333. 349 (1988)

- C. E. Hedge and D. C. Noble, Bull. Geol. Soc. Am 10 82, 3503 (1971); J. G. Fitton, D. James, W. P Leeman, J. Geophys. Res. 96, 13693 (1991); J. G. Fitton, D. James, P. D. Kempton, D. S. Ormerod, W P. Leeman, J. Petrol, 29, 331 (1988); D. S. Coleman and J. D. Walker, Mem. Geol. Soc. Am. 176, 391 (1990); J. D. Walker and D. S. Coleman, Geology 19, 971 (1991).
- 11. M. A Menzies, W. P Leeman, C J Hawkesworth, Nature 303, 205 (1983)
- G. L. Farmer et al., J. Geophys. Res. 94, 7885 12 (1989)
- 13. D S. Coleman and J D Walker, ibid. 97, 11011 (1992)14
- J. E. Wright and J L Wooden, Geology 19, 457 (1991)
- 15. G. Van Kooten, Contrib. Mineral. Petrol. 76, 378 (1981); , W. P. Leeman, M. A Menzies, Eos 66, 414 (1985)
- M. A. Domenick, R. W. Kistler, F. C. W. Dodge, M. 16. Tatsumoto, Geol Soc. Am. Bull. 94, 713 (1983).
- Ages used for correction of isotopic compositions 17. are 90 million years ago for all samples. All isotopic analyses were performed at the University of North Carolina (UNC). Sample preparation was performed according to procedures outlined in Frost (18). Samples were dissolved at approximately 150°C in a sealed beaker or at 180°C in a Teflon bomb using an HF-HNO3 mixture. Separation of Rb, Sr, and the rare earth elements was accomplished using standard cation exchange techniques Sm and Nd were separated on anion columns using the HNO₃-CH₃OH method. All iso-topic analyses were performed on a VG Sector 54 thermal ionization mass spectrometer Sr and Nd were analyzed using dynamic-multicollector mode with $^{88}{\rm Sr}$ = 4 V and $^{144}{\rm Nd}$ = 1 V. Rb and Sm were analyzed using static-multicollector mode with ^{87}Rb = 200 mV and ^{147}Sm = 500 mV. Analytical blanks for all elements reported are less than 100 pg. On the basis of multiple analyses of standard BCR-1, analytical precision for Sr and Nd (shown as error bars on Fig. 2) is better than 20 ppm ($^{87}Sr/^{86}Sr$ = ±0.000014 and $^{143}Nd/$ $^{144}Nd = \pm 0.000010$). Comparison of trace element concentrations determined by isotope dilution and energy-dispersive x-ray fluorescence spectrometry (18) show excellent agreement. Sr data are normalized to a 86Sr/88Sr ratio of 0.1194, and replicate UNC analyses of NBS-987 yield a 87 Sr/ 86 Sr ratio of 0.710250 \pm 0 000016 (number of analyses, n = 169) Nd data are normalized to a 146Nd/144Nd ratio of 0.7219, and replicate UNC analyses of La Jolla Nd yield a 143Nd/144Nd ratio of 0.511854 ± 0.000007 (n = 49).
- 18. T P. Frost, U.S. Geol. Surv Open-File Rep. 87-193 (1987)
- _____ and G A. Mahood, *Geol. Soc Am. Bull.* 99, 272 (1987) 19.
- 20. F C. W. Dodge and R. W. Kistler, J Geophys. Res. 95, 17841 (1990)
- G. A Mahood and A. N. Halliday, Contrib. Mineral. 21. Petrol. 100, 183 (1988), W. Hildreth, A. N. Hallıday, R L Christiansen, J Petrol. 32, 63 (1991); C. M. Johnson, J. Geophys. Res. 96, 13485 (1991)
- T. W Stern, P. C. Bateman, B. A Morgan, M. F 22 Newell, D. L. Peck, U.S. Geol Surv. Prof Pap. 1185, 17 p. (1981).
- A. F. Glazner, G. L. Farmer, W. T. Hughes, J. L. Wooden, W. Pickthorn, *J. Geophys. Res.* 96, 23. 13673 (1991); A. F. Glazner and G. L. Farmer, Science 255, 72 (1992)
- 24. W. Hildreth and S. Moorbath, Contrib. Mineral. Petrol. 98, 455 (1988).
- 25. DNd is defined as the distribution coefficient for Nd ($C_{\text{fractionating solids}}/C_{\text{magma}}$), where C is concentration in weight units
- Supported by National Science Foundation grants 26 EAR-8816941 and EAR-8917300 (A F.G.) We appreciate helpful reviews by C. M. Johnson, J. S. Miller, J J. W. Rogers, J. D. Walker, and an anonymous reviewer.

12 June 1992; accepted 22 October 1992

1926