Warming accelerates decomposition of decades-old carbon in forest soils

Francesca M. Hopkins\textsuperscript{a,b,1}, Margaret S. Torn\textsuperscript{c,d}, and Susan E. Trumbore\textsuperscript{a,b}

\textsuperscript{a}Department of Earth System Science, University of California, Irvine, CA 92697; \textsuperscript{b}Department of Biogeochemical Processes, Max Planck Institute for Biogeochemistry, 07745 Jena, Germany; \textsuperscript{c}Earth Sciences Division, Lawrence Berkeley National Laboratory, Berkeley, CA 94720; and \textsuperscript{d}Energy and Resources Group, University of California, Berkeley, CA 94720

Edited by William H. Schlesinger, Cary Institute of Ecosystem Studies, Millbrook, NY, and approved May 11, 2012 (received for review January 10, 2012)

Global climate carbon-cycle models predict acceleration of soil organic carbon losses to the atmosphere with warming, but the size of this feedback is poorly known. The temperature sensitivity of soil carbon decomposition is commonly determined by measuring changes in the rate of carbon dioxide (\textit{CO}_2) production under controlled laboratory conditions. We added measurements of carbon isotopes in respired \textit{CO}_2 to constrain the age of carbon substrates contributing to the temperature response of decomposition for surface soils from two temperate forest sites with very different overall rates of carbon cycling. Roughly one-third of the carbon respired at any temperature was fixed from the atmosphere more than 10 y ago, and the mean age of respired carbon reflected a mixture of substrates of varying ages. Consistent with global ecosystem model predictions, the temperature sensitivity of the carbon fixed more than a decade ago was the same as the temperature sensitivity for carbon fixed less than 10 y ago. However, we also observed an overall increase in the mean age of carbon respired at higher temperatures, even correcting for potential substrate limitation effects. The combination of several age constraints from carbon isotopes showed that warming had a similar effect on respiration of decades-old and younger (<10 y) carbon but a greater effect on decomposition of substrates of intermediate (between 7 and 13 y) age. Our results highlight the vulnerability of soil carbon to warming that is years-to-decades old, which makes up a large fraction of total soil carbon in forest soils globally.

The potential for carbon stored on land to become a source of carbon dioxide (\textit{CO}_2) to the atmosphere in the 21st century is a key uncertainty in predictions of future climate. Global warming increases the rate of decomposition of soil organic carbon (C\textsubscript{r}), a major loss pathway of C from the land surface to the atmosphere, thus contributing to the increase in atmospheric \textit{CO}_2 and hence, global temperatures. However, how much of the estimated 3,000 Pg C (1) stored in soils globally is vulnerable to enhanced decomposition with warming is highly uncertain and difficult to assess (2). In particular, the temperature sensitivity of C cycling on decadal timescales is a key uncertainty controlling the size of potential soil C responses to warming (3). Although there are no global estimates of decadal-aged C, it makes up the majority of C in mineral soils in temperate forests (4). We took advantage of a decade-long, whole-ecosystem C-isotope label to isolate the effect of warming on decomposition of decades-old C in a laboratory incubation experiment.

The temperature sensitivity of decades-old C is difficult to observe using traditional approaches, such as response of \textit{CO}_2 flux to experimental warming, because respiration is dominated by soil C cycling on fast timescales of 1 y or less. Previous studies using C isotopes to identify older C and assess its temperature sensitivity do not provide consistent results (recently reviewed in refs. 5 and 6). Most of these studies used a change in vegetation type (e.g., from C\textsubscript{3} to C\textsubscript{4} photosynthetic vegetation) as a means to distinguish old and young C. However, such vegetation shifts also change the amount and quality of C inputs to soil, affecting the decomposition process and potentially confounding measurements of temperature sensitivity. In addition, most studies took place in agricultural soils and may not be representative of less managed systems. Other methods to determine the temperature sensitivity of slower-cycling C also have significant drawbacks. Extended incubation periods to deplete the soil of fast-cycling C pools can change the decomposition process through substrate limitation (7). The response of slow-cycling soil C to warming is difficult to detect on the timescales of manipulative experiments, and it may be affected by covarying factors along natural temperature gradients (6). Model-derived predictions of temperature sensitivity of C pools cycling on different timescales are highly sensitive to assumptions in underlying model structures, such as which parameters are temperature-sensitive (8). Moreover, any inference of temperature sensitivity from bulk \textit{CO}_2 fluxes alone is difficult to relate to soil C destabilization processes, because respiration integrates across C pools stabilized by multiple interacting controls. Specifically, the mean residence time of different soil C pools is affected by both biology and physicochemical conditions, which are both likely to be temperature-sensitive (9). As a result, the effect of warming on the stability of soil C stocks is a topic of intense debate.

We investigated the temperature sensitivity of decades-old C by taking advantage of a whole-ecosystem C-isotope label in two temperate forest sites. Both sites had free air CO\textsubscript{2} enrichment (FACE) experiments, where atmospheric CO\textsubscript{2} concentrations in treatment plots are raised by fumigating with fossil-derived CO\textsubscript{2} that has a distinct C-isotope signature in \textit{δ}\textsuperscript{14}C and \textit{δ}\textsuperscript{13}C compared with background air [fumigation gas \textit{δ}\textsuperscript{14}C value of $\sim$1,000‰ compared with 50–100‰ for background air (10) and \textit{δ}\textsuperscript{13}C value of $\sim$43‰ compared with about $\sim$8‰ in background air (11, 12)]. Thus, C fixed by photosynthesis and incorporated into plant material and soil C under elevated CO\textsubscript{2} is isotopically distinguishable from previously existing soil C (FACE label). CO\textsubscript{2} enrichment began at both sites more than a decade before we sampled soils (Table 1), and therefore, the C-isotope label allows us to distinguish the contribution of decades-old C (pre-FACE C > 10 y) from the contribution of more recent C (FACE C < 10 y) to heterotrophic respiration during incubation using standard isotopic mixing models (13).

Author contributions: F.M.H., M.S.T., and S.E.T. designed research; F.M.H. performed research; F.M.H. and S.E.T. analyzed data; and F.M.H., M.S.T., and S.E.T. wrote the paper.

The authors declare no conflict of interest. This article is a PNAS Direct Submission.

Freely available online through the PNAS open access option.

\textsuperscript{1}To whom correspondence should be addressed. E-mail: fhopkins@uci.edu.

This article contains supporting information online at www.pnas.org/lookup/suppl/doi:10.1073/pnas.1120603109/-/DCSupplemental.
Although elevated CO₂ soils provide us with a unique opportunity to constrain how much decades-old C contributes to respiration by using the large difference in ¹⁴C and ¹³C between C fixed before and after FACE (Fig. 1), measurements of background levels of ¹³C in the nonenriched ambient CO₂ plots provide an additional age constraint. In ambient CO₂ plots, the ¹³C content of soil-respired CO₂ reflects the relative contribution of ¹⁴C fixed by photosynthesis into ecosystem C pools since aboveground nuclear weapons testing in the 1960s (Fig. 1). The atmospheric ¹⁴CO₂ signature has been declining by ~5% per y in recent years (14), and therefore, the mean age of respired C—the mean time elapsed since respired C was fixed from the atmosphere—can be determined using a time-dependent, steady-state model and the atmospheric history of ¹⁴C (15). With the bomb-derived ¹⁴C label, we can detect differences between C fixed from the atmosphere from 1 y to several decades before the date of sampling.

We sampled soils from both ambient and elevated CO₂ treatment plots at two FACE sites (Aspen FACE, Rhinelander, WI; Duke FACE, Durham, NC) after they had been exposed to elevated CO₂ for 11 y. Both sites are temperate forest plantations on old agricultural soils, but they differ with respect to species, lifeform, and stand age (Table 1). At Aspen FACE, deciduous aspen clones were planted in monoculture in 1997, and CO₂ enrichment was initiated the next growing season. At Duke FACE, evergreen loblolly pines were planted in 1983, and CO₂ enrichment began when the trees were already 13 y old. We incubated surface mineral soils (0- to 15-cm depth) at their site mean annual temperatures (MATs; 5 °C and 15 °C, respectively) and under two warming treatments (+10 °C and +20 °C). Respiration rates dropped with time (Fig. S1) in the elevated CO₂ treatment plots at both sites, with (i) a gradual increase in Δ¹⁴C of respiration from ambient CO₂ soils, reflecting relatively faster decomposition of 10-y-old C fixed during the CO₂ enrichment period, with slightly higher ¹⁴C content than the youngest C because of the gradual decline in atmospheric ¹⁴C in both CO₂ treatments over this time period.

### Results

#### Respiration Sensitivity to Warming

Warming consistently increased respiration rates from incubated surface soils for both CO₂ levels at the two sites (P < 0.0001 for temperature effect). Although respiration rates dropped with time (Fig. S1) in the Duke soils, the effect of temperature on respiration rate was consistent over the many months (up to 12 mo for Duke soils) of the experiment. The increase in respiration rates corresponded to a Q₁₀ of 1.5–1.9 for Duke and 2.9–3.1 for Aspen. The elevated atmospheric CO₂ treatment also significantly increased fluxes (P = 0.006) and interacted with the temperature effect (P = 0.044) at the Aspen site, but it had no statistically significant effect on fluxes at Duke.

#### Temperature Dependence of Decades-Old C (Pre-FACE C)

Isotopic signatures of the CO₂ respired in the incubations reflect the large influence of the isotopically depleted C fixed in the FACE experiments (Fig. 2). We used an isotopic mixing model to partition fluxes from the elevated CO₂ treatment into FACE-derived (<10 y) and pre-FACE (>10 y) pools using the FACE ¹⁴C label (SI Methods). In the FACE soils, roughly one-third of the C respired was fixed before the FACE experiment, regardless of temperature (Table 2). Warming increased the rate of losses from both pools, showing that decades-old C (>10 y) is vulnerable to immediate, enhanced losses on warming and has similar temperature sensitivity as younger FACE-derived (<10 y) C

### Table 1. Site information

<table>
<thead>
<tr>
<th>Site</th>
<th>Location</th>
<th>MAT (°C)</th>
<th>Mean annual precipitation (mm)</th>
<th>Planted (y)</th>
<th>+CO₂ (y)</th>
<th>Vegetation</th>
<th>Soil type</th>
<th>Soil texture</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aspen</td>
<td>45°N 89°W</td>
<td>4.9</td>
<td>810</td>
<td>1997</td>
<td>1998–2009</td>
<td><em>Populus tremuloides</em></td>
<td>Ultic Hapludalf</td>
<td>Sandy loam</td>
</tr>
</tbody>
</table>

#### Fig. 1. Atmospheric ¹⁴C content by year. The solid line represents background atmosphere Δ¹⁴C value since 1960, and the dashed and dotted lines represent atmosphere with FACE in elevated CO₂ plots at Aspen and Duke, respectively. Light gray lines represent the potential variability in ¹⁴C signature of the elevated CO₂ treatments based on the SD of CO₂ concentrations measured in elevated CO₂ plots (SI Methods).
Isotopic partitioning using the $\delta^{13}C$ label also supports the conclusion that both FACE and pre-FACE C are equally sensitive (Table 2).

We quantified the temperature effect on partitioned fluxes with an exponential model, where the temperature sensitivity coefficient $b$ defines the increase in flux per change in temperature and $A$ is a constant that represents the basal reaction rate (8). Within each site, we observed no significant differences in $b$ for C pools of different ages (Fig. 3 Inset). The flux of pre-FACE (>10 y) C was slightly more temperature-sensitive (but not statistically different) than the flux of FACE (<10 y) C. Values of $A$ were always higher for FACE C than pre-FACE C, confirming that the model separated pools with different overall cycling rates.

Increase in Substrate Availability with Temperature. Along with increased fluxes, we observed an immediate shift in the $\Delta^{14}C$ signature of respired $CO_2$ from warmed soils relative to the site MAT control soils (Fig. 2 A and C). Warming increased the mean age of respired C, which was shown by the significant increase in $\Delta^{14}C$ of respiration with incubation temperature from the ambient $CO_2$ treatment at both Aspen ($P = 0.0454$) and Duke ($P = 0.0058$). For the elevated $CO_2$ soils, $\Delta^{14}C$ of respiration also tended to increase with warming, although this difference was not statistically significant because of greater variability in $\Delta^{14}C$ of respiration among replicates (Table 3).

To test whether this pattern was caused by rapid depletion of fast-cycling C substrates, we normalized the isotopes of $CO_2$ flux data by amount of C lost (rather than by time). This normalization allows us to compare the sources of the equivalent amount of initial soil C.

### Table 2. Fraction of soil C stock (0- to 15-cm mineral soil) and respiration flux (first sampling) coming from pre-FACE (>10 y) C (±SEM) identified with $^{14}C$ and $^{13}C$ mixing models

<table>
<thead>
<tr>
<th>Stock ($f_{&gt;10} y$ by $^{13}C$)</th>
<th>Temperature (°C)</th>
<th>Respiration ($f_{&gt;10} y$ by $^{14}C$)</th>
<th>Respiration ($f_{&gt;10} y$ by $^{13}C$)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Duke</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>$CO_2$ ambient</td>
<td>15</td>
<td>0.33 (0.10)</td>
<td></td>
</tr>
<tr>
<td>$CO_2$ ambient</td>
<td>25</td>
<td>0.31 (0.15)</td>
<td></td>
</tr>
<tr>
<td>$CO_2$ ambient</td>
<td>35</td>
<td>0.27 (0.12)</td>
<td></td>
</tr>
<tr>
<td>$CO_2$ elevated</td>
<td>0.62 (0.03)</td>
<td>0.29 (0.16)</td>
<td>0.38 (0.15)</td>
</tr>
<tr>
<td>$CO_2$ elevated</td>
<td>15</td>
<td>0.24 (0.16)</td>
<td>0.28 (0.07)</td>
</tr>
<tr>
<td>$CO_2$ elevated</td>
<td>25</td>
<td>0.28 (0.22)</td>
<td>0.42 (0.06)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Aspen</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>$CO_2$ ambient</td>
<td>5</td>
<td>0.40 (0.31)</td>
<td></td>
</tr>
<tr>
<td>$CO_2$ ambient</td>
<td>15</td>
<td>0.34 (0.13)</td>
<td></td>
</tr>
<tr>
<td>$CO_2$ ambient</td>
<td>25</td>
<td>0.35 (0.12)</td>
<td></td>
</tr>
<tr>
<td>$CO_2$ elevated</td>
<td>0.68 (0.07)</td>
<td>0.28 (0.12)</td>
<td>0.43 (0.21)</td>
</tr>
<tr>
<td>$CO_2$ elevated</td>
<td>5</td>
<td>0.27 (0.08)</td>
<td>0.44 (0.09)</td>
</tr>
<tr>
<td>$CO_2$ elevated</td>
<td>15</td>
<td>0.28 (0.10)</td>
<td>0.37 (0.11)</td>
</tr>
<tr>
<td>$CO_2$ elevated</td>
<td>25</td>
<td>0.28 (0.10)</td>
<td>0.37 (0.11)</td>
</tr>
</tbody>
</table>
the size of the active pool (\(k_a\)) with warming, whereas the decay constant of the passive pool (\(k_p\)) for that pool stays relatively constant (Table S1) in the Duke soils (flux data from Aspen soils did not fit the model). This pattern has been found in many studies, and it has resulted in an ongoing debate about whether temperature dependence can be in both the pool size terms and the rate constant (18–20).

These results suggest that increased substrate availability may be the key to the initial stages of the warming response.

**Age of Respired C Substrates.** From the ambient CO\(_2\) treatment soils incubated at the site MAT, bomb \(^{14}\)C modeling estimates of the age of C respired were 2 y for Aspen (<1–5 y, 95% confidence interval) and 3 y for Duke (<1–6 y). Warming increased the mean age of C respired by 3–5 y at both sites relative to the MAT treatment (MAT + 10°C: +3 y at Aspen, +4 y at Duke; MAT + 20°C: +3.5 y at Aspen, +5 y at Duke). The young age of respired C agrees with the expectation that C with the fastest turnover time is metabolized early in incubation and that the youngest C dominates the heterotrophic respiration signal.

To confirm that additional substrate made available by warming was less than a decade old, we modified our original mixing model to include a warming-induced pool defined by the change in flux and \(^{14}\)C end members of substrate respired from warmed soils over the MAT control soils (Fig. 4). Using data for the same cumulative loss across temperatures, we assumed an equal contribution of <1 y C to fluxes in ambient CO\(_2\) soils and the same \(^{14}\)C end members at all temperatures, which enabled us to solve for the \(^{14}\)C value of the warming-induced substrates. We estimate this pool to have a mean age of 7–13 y in Aspen and 9–12 y in Duke.

**Discussion**

**Vulnerability of Decades-Old Soil C to Warming.** In these two temperate forest soils, warming increased respiration of soil C more than a decade old fixed before the FACE treatment. Such decades-old C is a major component of organic matter in these soils and temperate forests more broadly (4), implying that a large portion of soil organic C is vulnerable to increased decomposition with global warming. C more than a decade old made up 70% of mineral soil C in the 0- to 15-cm depth that we incubated, but ~30% of the C respired by 3 y from the warm soils, and the mean age of C respired by 3 y at Duke, +4 y at Duke; MAT + 20°C: +3.5 y at Aspen, +5 y at Duke)

**Temperature Sensitivity of Decades-Old Soil C Decomposition Is Robust Across Sites.** Warming increased decomposition rates of decades-old C at both sites, despite large differences in overall soil cycling rates (Fig. 3 and Table 3). Aspen had much slower...
respiration rates than Duke (Table 3) (23) and less contribution of decades-old C to the soil C stock, which was indicated by bomb-derived $^{14}$C in bulk ambient CO$_2$ soils ($\Delta^{14}$C values of 51% at Aspen and 73% at Duke). The total amount of soil C and its distribution among physical fractions also differed greatly between the sites (Table S2) (24, 25). Specifically, there was a greater proportion of mineral stabilized C at Aspen FACE relative to Duke FACE. Along with the $\Delta^{14}$C of bulk soils, this finding suggests a larger proportion of prebomb C and a larger overall pool of passive C not contributing to soil respiration at Aspen FACE than Duke FACE.

Nevertheless, the proportion of pre-FACE C in respiration and its response to warming was similar at both sites. For elevated CO$_2$ soils from Aspen and Duke, decades-old C comprised a surprisingly large 30% of respired CO$_2$ across all incubation temperatures. Based on the $\Delta^{14}$C of heterotrophically respired CO$_2$ from the ambient CO$_2$ soils, the estimated mean age of respired C was 2–3 y; however, additional information from the FACE-labeled soils indicates that this finding is averaging of very young C with almost one-third that is more than a decade old. Other studies with in situ isotope labels from agroecosystems show that up to 66% of respiration derived from decades-old C (>45 y old (26); other studies: 0–21%>40 y (27); 52%>14 y old (28); 45%>26 y old (26)).

The age distribution of respired C at both sites for the control temperature and warming treatments was consistent for both types of C-isotope tracers. Specifically, the offset between $\Delta^{14}$C–CO$_2$ respired in incubation and the $\Delta^{14}$C value of the atmosphere in the year of sampling was nearly the same at both sites, indicating similar residence time of respired C. Thus, respirable C was more uniform between soils than overall C stocks. Importantly, the similar response to warming—measured by the effect of warming on both proportional contribution of decade-old C to respiration fluxes and the increase in $\Delta^{14}$C respired—suggests that the sites have similar age distributions of C sources contributing to respiration and perhaps, similar mechanisms of temperature response.

**What Is Decadal-Aged C at These Sites?** We used the C-isotope signatures of soil C components (e.g., roots, microbes, and physical soil fractions) to identify sources of respired CO$_2$ and particularly, decades-old CO$_2$ (Fig. 5). C that was recently de-
posited by roots, either as exudate or litter, is probably the source of most C respired from these soils, although visible roots were removed before incubation. Roots are increasingly recognized as the primary source of C to microbes in A horizon soils (29). The age of roots coincides with Δ14C values of respired C at these sites; at Duke FACE, the mean age of roots was 4–6 y, with some roots >18 y old (30), and at Aspen FACE, the mean age of roots was 1–3 y old for <2-mm roots and 3–5 y old for >2-mm roots. Given these values, the time spent by C in structural root tissue was sufficient to give the respired CO2 age without significant additional time in soil C; the inferred age of respired C may be a function of time spent in structural root tissue rather than soil C pools.

Although decay of root tissues may be a major component of respired CO2, it is not the primary source of decades-old C to respiration. Although root tissue containing pre-FACE C may have been present at Duke FACE, roots at Aspen FACE are composed only of FACE-derived C, because those trees experienced an enriched CO2 atmosphere for their whole lives. The source of decades-old C to respiration is more likely to be C that is associated with minerals <250 µm in size, which were ~20% and ~30% of soil C at Aspen and Duke, respectively (24, 25). Temperature sensitivity of this fraction is consistent with the increase in respiration of both FACE-derived and pre-FACE C, because this fraction contains significant portions of both age classes of C and is the only soil pool with enough bomb-derived 14C to have caused the increase in Δ14C of respiration with warming. The larger-size fraction (>250 µm) of mineral-associated C at Duke FACE has very low 14C values in both elevated and ambient CO2 treatments, suggesting very slow turnover and negligible contribution to respiration. Although we do not have Δ14C measurements for physical soil fractions at Aspen FACE, incorporation of 13C-depleted C from the FACE label in mineral-associated size fractions suggests similar C turnover patterns as observed in the Duke FACE soils (24).

Microbial biomass has a similar C isotopic signature to its C sources and respiration (29), but it is too small of a C pool in itself to be solely responsible for the observed respiration flux. Up to 9% of total soil C at Duke was respired as active-pool C in the warmest soils compared with living microbial biomass that is, at most, 4–5% of total C in these soils (31).

Another explanation for the large release of decades-old C is a disturbance effect; however, this reason is unlikely to be a full explanation in our experiment. A major criticism of the incubation method is that preincubation sample preparation may change soil C decomposition rates. Particularly if soils are sieved, previously protected, decades-old soil C may be exposed to microbial attack and vulnerable to degradation. However, potential disturbance effects are unlikely to yield similar results for both sites, because Aspen soils were sieved, whereas Duke soils were not. In addition, soil aggregation is probably not the mechanism of soil C protection in Duke FACE soils (13).

Increased Vulnerability of Intermediate-Aged (7–13 y) C with Warming. The central question of our experiment was whether decades-old C had different temperature sensitivity than faster-cycling C, because the feedback between soil respiration and climate warming in earth system models is particularly sensitive to this premise (32, 33). The FACE C-isotope label (both 13C and 14C) allowed us to unequivocally determine that decades-old C fixed before FACE had similar temperature sensitivity to C fixed during the last 10 y. However, integrating the bomb-derived 14C label into the analysis, we identified a subtle difference in the age of C respired with increasing temperature. Specifically, we observed a parallel increase in Δ13C of CO2 respired with warming in both ambient and elevated CO2 soils that was not caused by exhaustion of fast-cycling C. The similar increase in Δ14C in both CO2 treatments suggests that warming increased the contribution of C fixed earlier in the decade since the FACE treatment began, reflecting the ~60% decline in atmospheric Δ13C over the period of the FACE experiment in ambient CO2 plots and an ~40% decline under enriched CO2, where the decline in background atmosphere Δ13C was diluted by addition of FACE label C during CO2 enrichment.

Our study is not unique in the finding that the 14C content of respiration increased with warming. Two other incubations of forest soils with background levels of 14C inferred higher temperature sensitivity of soil C with a similar age as the age in our study. In boreal forest soils, increased 14C of respiration with warming corresponded to higher temperature sensitivity of decadal cycling C compared with annually and centennially cycling C pools (34). In a temperate forest soil, 25 °C of warming increased the 14C-derived mean residence time of respiration by up to 4 y from 7.9 to 11.9 y (35). These findings show that warming increases the respiration of C up to several decades old (fixed during the postbomb period; i.e., post-1960).

With the additional time constraint of the FACE label, we can eliminate the possibility that C much older than 10 y was more temperature-sensitive than older ages of C. If this finding were the case, we would expect to observe a much more rapid increase in 14C respired by elevated CO2 soils than ambient CO2 soils. From the similar rate of change in both types of isotope labels, we conclude that some portion of soil organic C aged 7–13 y responded disproportionately to warming. Different temperature sensitivity of this age of C does not contradict our finding of similar temperature sensitivity of decades-old and younger C; in fact, C of this age would be partitioned into both the FACE and pre-FACE C pools in the mixing model.

Higher temperature sensitivity of intermediate-aged C provides a potential explanation for inconsistencies between conclusions of previous studies that used C-isotope labels in soil to infer temperature sensitivities of different ages of C. Isotope label studies differ in the length of the labeling period before sampling, resulting in varying definitions of older C. If C with a similarly disproportionate temperature sensitivity and age is present in the soils of these studies, then inconsistent conclusions for temperature sensitivity of older C may depend on whether this C was included as part of the older or younger C pool. In shorter experiments, temperature sensitivity of this intermediate C is likely to be categorized as older C, and therefore, its disproportionate response to warming gives the appearance that older C is more temperature-sensitive [e.g., soils sampled after 5 y of label (36) or 14 y of label (29)]. In contrast, studies with a longer labeling period [e.g., labels of 26 and 45 y (26) or 33 y (16)] find equal temperature sensitivity between age classes. This finding suggests that, when this intermediate-aged C with higher temperature sensitivity is categorized as younger C, its response cannot be resolved from the temperature response of the majority of respiratory C substrate, resulting in equal apparent temperature sensitivity of the two age classes. The finding of equal sensitivity with longer label times suggests that the contribution of disproportionately temperature-sensitive C to the total flux is relatively minor.

The combination of the FACE isotope label and the bomb 14C signal allowed us to identify the effects of warming on three different timescales of C cycling and avoid some confounding factors present in previous studies. The age constraint of the FACE label improved age estimates over those estimates from bomb-derived 14C alone. Also, measurement of 14C has advantages over the 13C label used in most studies. Although Δ13C data reported here are corrected for mass-dependent fractionation, the 13C isotope may be affected by temperature-dependent kinetic fractionation by microbial respiration (37) or preferential use of 13C-depleted substrate (38). Other confounding factors include differential substrate depletion between temperature treatments (39), differences in substrate conditions because of
seasonal effects (40), and differences in C cycling under C3 and C4 vegetation. In FACE experiments, manipulation of CO2 concentrations may have altered decomposition rates, resulting in differences between CO2 treatments at these sites (41, 42); however, this manipulation is unlikely to affect our results. Although the Δ13C mixing model assumes similar decomposition rates of pre-FACE C between CO2 treatments, model results are not sensitive to this term. In addition, the Δ13C mixing model gave similar estimates of the fraction of pre-FACE C and does not require the assumption that C cycling rates are similar between the two treatments.

What Potential Mechanisms Underlie the Observed Temperature Response? The similarity in temperature sensitivity of the two broad age classes suggests a common suite of mechanisms of soil C response to warming. This finding is consistent with the conceptual framework emerging from recent synthesis efforts (6, 9), which emphasizes different temperature controls over microbial respiration and supply of soil C to microbial respiration. In the short term, the temperature sensitivity of microbial respiration is the primary control of the temperature dependence of soil respiration. The constant proportion of decay of older and younger C respired across temperatures is likely determined by their fractional contributions to microbially assimilable C. Hence, the warming response could simply reflect faster respiration of assimilable C by microbes. Alternatively, it could mean that the availability of younger and older C sources was controlled by the same process or that their respective controls were similarly temperature-sensitive.

Although the temperature sensitivity of microbial respiration has been well-established, much less is known about temperature dependence of substrate supply to microbial respiration, which controls C availability in the long term. Multiple lines of evidence suggest that warming increased the supply of C of both age classes to microbes, including consistently higher flux rates over the whole incubation period, larger pools of actively cycling C, and increase in the mean age of respiration substrate at higher temperature. Previous incubation and litter decomposition studies also suggest that warming increases the fraction of soil C that is assimilable by microbes (43, 19, respectively).

Various potentially temperature-sensitive processes could influence substrate supply or cause an apparent change in supply in an incubation, such as shift in microbial community composition (44), change in microbial efficiency (45), increased turnover of microbial biomass (46), change in biochemical composition of soil C substrates respired (47), increased desorption of mineral-adsorbed organic C (6), and increased diffusion. In our study, increased substrate availability coincided with an increase in respiration of soil organic C with a mean age of 7–13 y, suggesting that a greater proportion of C of this age became available with warming. Some of these processes can be ruled out, because they would increase assimilation of substrates without a change in substrate age, such as increased diffusion, change in microbial efficiency, or increased turnover of microbial biomass.

Other mechanisms may be consistent with a change in the age of respired CO2. A warming-induced shift in microbial community or enzyme production could change the use of C of different ages (48); however, it is unlikely that such a shift would happen within the relatively short time period over which we collected CO2 from these soils (7). Chemical kinetic theory, also known as the carbon quality temperature hypothesis (8, 49), provides a potential explanation for an increased contribution of slower turnover compounds because of higher temperature sensitivity of compounds with greater total bond strength, which is often associated with compounds that are more structurally complex (i.e., more chemical bonds) (47). If substrates with greater complexity are also retained in soils longer (i.e., become older) and warming disproportionately promotes their decomposition, we would expect to see an increase in the mean age of respired CO2 with increased temperature. However, the radiocarbon age of soil C is not necessarily indicative of biochemical stability (48), and we do not know the extent to which biochemical stability or activation energy of compounds per se controls C decomposability in mineral soils (50).

It is difficult to tease apart mechanisms in incubations such as this incubation or field respiration studies; heterotrophic respiration integrates over multiple C sources and reflects overlapping mechanisms of soil C stabilization. In addition, extended incubation periods have been criticized for their departure from in situ conditions (51, 52). Incubation isolates soils from sources of C input and results in rapid onset of substrate limitation to decomposers, which can modify the apparent response of respiration to warming (53). Substrate depletion was eventually observed in the incubation of Duke FACE soils at all temperatures, suggesting that the increase in amount of assimilable C under warming may not be sustained over time. It remains an open question whether increased substrate availability observed with warming in incubations would be sustained in a field setting or is the product of a finite, exhaustible pool as some studies suggest (54).

Modeling the Temperature Response of Soil C Decomposition. In many soil C models, temperature sensitivity is expressed exclusively in the rate constants of linear, donor-controlled soil C pools (55). When we modeled our data with this model structure, increased respiration was best simulated with an increase in the size of the active pool rather than a change in the rate constants. Indeed, including the effect of warming on substrate availability in current model structures would require a highly temperature-sensitive pool to rapidly transfer previously slow-cycling C to the fast pool. If this new warming-induced supply is rapidly depleted, then the flux from this highly temperature-sensitive pool may be transitory—a case we cannot determine with an incubation, because substrate limitation is observed at all temperatures. In that case, however, inferring the changes in respiration rates using only a temperature-sensitive rate constant may overstate the warming effect on the soil C stock.

If chemical bond strength (activation energy) were the fundamental limit to decomposition rates, then the Arrhenius equation of chemical kinetic theory (8) can be used to quantify increases in respiration substrate availability with warming. However, recent attempts to model this effect either explicitly (16) or implicitly (49) assumed that respiratory substrate stays constant under warming by parameterizing temperature sensitivity in the rate constants that directly control respiration rates. This approach predicts a more rapid loss of active pool C in warmed soils than soils at the MAT control temperature, which is counter to our findings. In contrast, our data suggest that the change in respiration rate with warming is more strongly controlled by substrate availability than temperature. As a result, caution must be taken in deriving parameter estimates in models from measurements of warming on respiration in incubations or field studies.

Earth system models are designed to predict future climate, but they still lack a predictive understanding of how much soil C is vulnerable on timescales of the next century. In this timeframe, the most important C response will come from C cycling on decadal timescales. Older soils (centuries to millennia) are also an important component of global soil C stocks (15), but their very long turnover times (and correspondingly slow decomposition rates) indicate that they will not have much effect on feedbacks in the 21st century and cannot be measured in incubation experiments in any case (56).

Our results indicate that large amounts of C (1,750–4,700 kg m−2) in the top 15 cm of mineral soils at these two temperate
forest sites) were vulnerable to increased decomposition losses with warming. The fact that we saw similar results at the two sites, despite differences in soil C stabilization therein, suggests that the pattern we observed may apply more broadly. The importance of decadal-aged C to the large amount of C in forest soils globally suggests that soil C could become a source of atmospheric CO₂ under global warming.

A continuing challenge for models is to understand the unresolved mechanisms where C of different ages and stability can have the same temperature sensitivity. Although more research is needed to better incorporate soil C decomposition processes into models, our results suggest that we need models and experiments that explicitly separate the temperature sensitivity of microbial metabolism and the temperature sensitivity of substrate supply rather than parameterizing the temperature sensitivity of any particular compound or fraction.

Methods

We sampled the top 0–15 cm mineral soil from the Duke and Aspen FACE sites, which have been documented extensively elsewhere (97). These FACE experiments have a similar design, consisting of replicate 30-m diameter forested plots, one-half of which receive CO₂ fumigation (elevated CO₂ experiments that explicitly separate the temperature sensitivity of microbial metabolism and the temperature sensitivity of substrate supply rather than parameterizing the temperature sensitivity of any particular compound or fraction.

We report error as the SEM of experimental replicates or by propagating the error in isotope calculations (60). Reported P-values are from comparisons of treatment means in ANOVA done using PROC GLM (unless t test was indicated), and exponential fits to data are done by PROC NLIN in SAS 9.2.

ACKNOWLEDGMENTS. We thank Xiaomei Xu and the staff of the W. M. Keck Carbon Cycle Accelerator Mass Spectrometer, University of California at Irvine for radiocarbon analyses. We thank Tim Filley and Sara Top of Purdue University for collection of soils from Aspen FACE and John Lichter at Bowdoin College for providing density and size fractions of soil from Duke FACE. We also thank to the principle investigators and staff of the Duke and Aspen FACE experiments for access, Carlos Sierra for discussion, and Claudia Czimczik and two anonymous reviewers for insightful comments on the manuscript. The Duke FACE experiment was funded by US Department of Energy’s Office of Science (DOE-BER) Grant DE-FG02-95ER62083. The Aspen FACE experiment was funded by DOE-BER, with additional support from the US Forest Service (USFS) Global Change Program, Michigan Technological University, the Canadian Forest Service, and the USFS Northern Research Station. F.M.H. was supported by a National Science Foundation Graduate Research Fellowship, an Achievement Rewards for College Scientists Foundation Scholarship, and a Ralph and Carol Cicerone Graduate Fellowship. M.S.T. was supported by DOE-BER Contract DE-AC02-05CH11231.


Supporting Information

Hopkins et al. 10.1073/pnas.1120603109

SI Methods

1.1 Site Description. Litter and soil were collected at two free air CO2 enrichment (FACE) experiments: Duke Forest FACE near Chapel Hill, North Carolina and Aspen FACE near Rhinelander, Wisconsin.

1.1.1 Duke FACE. The Duke FACE site, under forest cover since at least the 1940s, was burned and planted with loblolly pine (Pinus taeda L.) in 1983. Year-round CO2 enrichment began in 1996 in three of four replicate plots. A prototype plot that served as the fourth replicate was additionally fumigated during the summers of 1994 and 1995 (1). There was no difference in C-isotope values of soils from the prototype plot and the values of the other three replicates. Loblolly pine dominates the aboveground biomass, with deciduous trees in the understory. The soils are low-fertility acidic clay loam Hapludalfs in the Enon Series, and there are poorly drained during wet periods. In 2005, the top 15 cm mineral soil had an average C content of 2.16 kg C m⁻² (±0.15) and 2.10 kg C m⁻² (±0.18) and δ¹³C signature of −27.40 (±0.04) and −30.86 (±0.32) in ambient and elevated CO2 plots, respectively (2). During the period of the FACE experiment, C was accumulating in the soils of both the ambient and elevated CO2 plots because of recovery from disturbance (3).

1.1.2 Aspen FACE. The Aspen FACE site was agricultural land for at least 50 yr up to 1972, when it was purchased by the US Department of Agriculture Forest Service and subsequently used for short-rotation forestry. The site was cleared and disked in 1996 and 1997, and Aspen clones (Populus tremuloides Michx.) were planted on one-half of each FACE plot in June of 1997. The remaining one-half of each plot was planted as one-quarter aspen with paper birch and one-quarter aspen with sugar maple, but for this experiment, only soils from the aspen monoculture were used. CO2 fumigation began in 1998. Soils are Pandus clay loam: mixed, frigid, coarse loamy Al Histosols. Soil C was measured as 5.12 (+1.13) kg C m⁻² in ambient CO2 plots and 3.27 (+0.8) kg C m⁻² in elevated CO2 plots for the top 20 cm mineral soil, and soil C in elevated CO2 plots was −31.1% in δ¹³C (4).

1.2 FACE Experiments. Both experiments enriched CO2 levels by 200 µmol mol⁻¹ for the whole stands in each 30-m diameter treatment plot (1). The added CO2 was derived from natural gas, which is very depleted in ¹³C and ¹⁴C relative to the background atmosphere. Fumigation gas was −43‰ in δ¹³C and −1,000‰ in δ¹⁴C (5–7) compared with −8‰ in δ¹³C and 40–110‰ in δ¹⁴C for the background atmosphere over the period of the FACE experiments (8, 9). The δ¹³C of roots and leaves in elevated CO2 plots confirms fixation of CO2 with the expected isotopic admixture of fumigation gas and background air; root ingrowth cores measured root inputs as −39‰ at Duke, and roots picked from soil cores the year before our sampling (2008) measured root inputs as −39‰ at Aspen (10, 4, respectively).

1.3 Sampling Procedure. We sampled the top 0–15 cm mineral soil plus the overlying litter layer in each of the elevated and ambient CO2 plots. We treated each FACE ring or experimental plot as the level of replication (Duke n = 4, Aspen n = 3). Organic soil (litter layer) and mineral soils were collected from the Duke FACE on July 7, 2008. We sampled the litter layer by cutting a 100-cm² rectangle of all organic material found above the mineral soil surface. Mineral soils were sampled under the cleared area down to 15-cm depth using a 5-cm diameter slide hammer corer. Litter and mineral soils were collected at Aspen FACE on July 1, 2009. After surface litter removal, mineral soils were sampled with a 5-cm diameter impact corer in increments of 0–5 and then 5–15 cm.

1.4 Incubation Procedure. Soils were transported to the laboratory on ice and refrigerated before incubation. For both soils, rocks and roots were removed before incubation. For Duke soils, whole soil cores (minus roots) were placed in interior containers inside of 2-L Mason jars at field moisture content, and a subsample was taken after incubation to determine water content. For the Aspen site, mineral soils were additionally sieved to 4 mm, composited by ring, and then split using a cone and quarter method. About 140 g soil were used for incubation, with a subsample taken to determine soil moisture content before incubation. Because Aspen soils dried out during sieving, deionized water was added to soils after they were in incubation jars to return them to field moisture level.

Soils were kept at field moisture over the incubation period by high relative humidity inside of incubation jars. Soils were incubated continuously at site mean annual temperature, +10 °C, and +20 °C of warming, (Aspen: 5 °C, 15 °C, and 25 °C; Duke: 15 °C, 25 °C, and 25 °C; ±1 °C) for around 1 d before CO2 fluxes were measured for the first time. Temperature treatments were monitored by a Stowaway Tidbit Temperature Logger (Onset Corporation) kept alongside soil samples during incubation and sampling. Soils were removed briefly from their temperature treatments for flux sampling (less than 1 h per sampling). Jars were removed from their temperature treatments and allowed to come to room temperature (24 °C) before CO2 was removed from the jar.

1.5 CO2 flux and isotope sampling. For soil CO2 flux and isotope measurements, jars were capped with modified Mason jar lids fitted with stopcock-type sampling ports. Effective jar volume and potential leakage were checked by measuring the pressure change on expansion of the jar headspace into an evacuated known volume (11). At the beginning of the flux measurement period, jar headspace was purged with CO2-free air at 1 L min⁻¹ until the jar’s volume had been completely purged at least three times before closing the jar and allowing CO2 to accumulate in the headspace. Headspace CO2 concentrations were measured in 2-mL syringe samples injected upstream of a CO2 scrubber and air pump in line with a Licor 6252 Infrared Gas Analyzer (12). Fluxes were calculated as the CO2 evolved for the effective jar volume for a given incubation time.

For each flux sampling period, we measured δ¹³C–CO2 on air sampled directly from the jar headspace. Syringe samples were injected into He-filled vials for measurement of δ¹³C–CO2 by continuous flow isotope ratio MS (Thermo Finnigan Gas Bench coupled to Delta Plus). δ¹³C values are reported relative to the Pee Dee Belemnite standard.

For measurement of Δ¹⁴C–CO2, jar headspace CO2 was allowed to build up to 0.3–2.5% and then was collected using an evacuated container. CO2 was purified cryogenically on a vacuum line and then reduced to graphite using the methods in the work by Xu et al. (13). Radiocarbon content of samples was measured at University of California at Irvine’s W. M. Keck Carbon Cycle Accelerator Mass Spectrometer facility (14). Data are reported as Δ¹³C, the per mil difference in radiocarbon content relative to 95% of the activity of the oxalic acid I standard. Δ¹⁴C was corrected for fractionation by normalizing to
-25% for samples from the ambient CO2 plots (15). In enriched CO2 atmospheres, the C-isotope signature of C fixed in photosynthesis does not just reflect C-isotope fractionation but rather, fractionation plus the mixing of different source gases (i.e., fumigation gas and atmosphere). Therefore, a different fractionation correction for Δ14C must be used. Using the method from the work of Torn and Southon (16), we assumed that fractionation of C isotopes in photosynthesis was similar between CO2 treatments, and we used the δ13C values for ambient CO2 soils to correct for mass-dependent fractionation in elevated CO2 soils.

Error is reported as the SEM of samples from replicate plots, reflecting spatial heterogeneity among plots within the experiment sites. Error was propagated in isotopic mixing calculations using the procedure outlined in the work by Phillips and Gregg (17). Precision of all measurements was better than the SEM of replicate plots.

1.6 Postincubation. After incubation, Duke soils were sieved to 2 mm to remove any remaining rocks, and subsampled for soil moisture determination. No differences in soil moisture were found between temperature treatments. Both Duke and Aspen soils were dried at 60 °C and ground for analysis of bulk solid sample C and C-isotope content. Percent C and δ13C were measured on aliquots of each sample using an NA 1500 NC elemental analyzer (Fisons Instruments) coupled to isotope ratio MS (as previously described).

1.7 Bomb 14C Modeling. We used the record of Northern Hemisphere atmospheric 14C content (9) and a steady-state, one-pool model of 14C to determine the real mean residence time of CO2 respired during incubation (18). The model works on annual time steps, assuming that all C input to soil has the 14C signature of that year’s atmosphere. The decomposition rate, k, is inversely proportional to the turnover time of the modeled soil C pool, which is, in turn, closely related to that pool’s mean residence time (assuming no lags in living tissue). If we consider the respired CO2 to be coming from one homogenous pool, k can be determined from Δ14C of CO2 respired. The model is solved independently for each treatment by iteratively adjusting k until model 14C content equals the 14C signature of that year’s atmosphere. The turnover time, estimated from 1/k, represents the mean age of carbon being respired assuming a homogenous C pool that contributes to microbial respiration. This mean age includes time spent by C in living tissues (e.g., roots or twigs) as well as the mean time for decomposition.

1.8 Isotopic Mixing Model. 1.8.1 14C mixing model. We used an isotopic mixing model to take advantage of the decade-long FACE label. The principle of the model is similar to the principle of the 13C mixing models used extensively at the FACE sites; we use the difference between 14C content of preexisting soil C and FACE label C added since initiation of CO2 fumigation. First, we calculate the 14C signature of the FACE atmosphere (elevated CO2 plots) for the mixture of background air and fossil-derived (-1,000‰) fumigation gas (Eq. S1):

\[ \Delta^{14}C_{\text{FACE}} = \Delta^{14}C_{\text{new}} \frac{[\text{CO}_2]_{\text{amb}}}{[\text{CO}_2]_{\text{FACE}}} - 1000\% \cdot \frac{[\text{CO}_2]_{\text{FACE}} - [\text{CO}_2]_{\text{amb}}}{[\text{CO}_2]_{\text{FACE}}} . \]  

We used the daytime CO2 concentration records (7:00 AM to 7:00 PM) for experimental plots from the website of the two experiments (Duke, http://face.env.duke.edu; Aspen, http://aspenface.mtu.edu) to define the average CO2 concentration of ambient and elevated CO2 plots from the beginning of the CO2 experiment to the year before the sampling. [CO2]amb, the atmospheric CO2 content in ambient plots (and the background CO2 endmember for the elevated CO2 plots), is defined as the average (±SD) CO2 concentration in ambient CO2 plots (Duke, 379 ± 7, Aspen, 370 ± 18), and [CO2]preFACE is the measured CO2 concentration in elevated CO2 plots (Duke, 534 ± 16; Aspen, 531 ± 13). We assumed the Δ14Cnew was equal to the Δ14C of the atmosphere in the year of sampling (Duke, 45‰; Aspen, 40‰), with a 5‰ error bar reflecting the seasonal cycle of Northern Hemisphere atmospheric Δ14C (9).

Using the atmosphere’s Δ14C value as the new (<10 y) Δ14C endmember assumes that all flux is coming from pools either <1 or >10 y old. However, the model is very insensitive to this endmember value because of the large isotopic difference between the CO2 in FACE elevated CO2 plots and ambient CO2 plots. We tested this assumption by increasing Δ14Cnew by about 10‰ to simulate a mix of substrates with a 3 y mean residence time. This endmember changed the estimated proportion of flux from the pre-FACE pool by 0.01 (4% of original estimate).

Then, we partition fluxes from both ambient and elevated CO2 plots into >10 and <10 y components by writing two mass balance equations for the flux (Eq. S2):

\[ \text{CO}_2^{\text{lev}} = \text{CO}_2^{>10} + \text{CO}_2^{<10,\text{lev}} \]  

and (Eq. S3)

\[ \text{CO}_2^{\text{amb}} = \text{CO}_2^{>10} + \text{CO}_2^{<10,\text{amb}} . \]  

We also wrote two equations for mass balance of the isotopes of flux (Eq. S4):

\[ \Delta^{14}C_{\text{levelev}} = \Delta^{14}C^{>10}\text{CO}_2^{>10} + \Delta^{14}C_{\text{FACE}}\text{CO}_2^{<10,\text{lev}} \]  

and (Eq. S5)

\[ \Delta^{14}C_{\text{ambamb}} = \Delta^{14}C^{>10}\text{CO}_2^{>10} + \Delta^{14}C_{\text{new}}\text{CO}_2^{<10,\text{amb}} . \]  

If we assume the flux and Δ14C content of the >10 y pool are the same for both CO2 treatments (i.e., CO2 fumigation had no effect on decomposition rates of previously existing soil C), then the system of equations can be solved for the fluxes from the <10 y pool of the elevated CO2 treatment and ambient CO2 treatment. In the original model (Fig. 3), we partitioned fluxes separately for each temperature treatment and site, deriving an independent Δ14C_y for each incubation temperature.

1.8.2 Modified 14C mixing model. We modified the CO2 flux mixing model for the warming treatments by adding another CO2 source pool to account for changes in the isotopic signature of respiration substrate with warming. First, we selected fluxes that represent the same amount of total C respired for each temperature treatment: 2.68% of soil carbon at Duke and 0.11% of soil carbon at Aspen. Second, we partitioned the fluxes from the elevated CO2 treatment into FACE and pre-FACE C using the previous approach but with isotopic endmembers that did not change with temperature. In contrast to the original mixing model, where we determined Δ14C_y separately for each temperature, we fixed Δ14C_y to the value determined from the mean annual temperature control treatment. This change had little effect on the fraction of >10 y C contributing to overall flux across temperatures (Fig. 4).

From the increase in Δ14C of CO2 respired from the ambient CO2 treatment with warming, we know that the increased flux was of an intermediate age: >3 y because the higher Δ14C values indicate carbon fixed earlier in the bomb period but <10 y because of the proportion of pre-FACE carbon remains the same. We determined the age of additional substrates respired under warming using CO2 fluxes from the ambient CO2 treatment as...
previously, we assumed that the flux of $>10$ y C was the same for a given temperature between CO2 treatments. In addition, we assumed that the contribution of C with a mean residence time of less than 1 y (i.e., C with the $\Delta ^{13}C$ signature of that year’s atmosphere) was constant across temperatures, and therefore, any additional flux with warming after subtracting out the increase in $>10$ y C was attributed to the warming-induced pool. We then used flux and isotopic mass balance equations to solve for the $\Delta ^{13}C$ value of the warming-induced substrate, which was 7–13 y at Aspen and 9–12 y at Duke.

1.7.3 $^13C$ mixing model. We also used $\delta ^{13}C$ of respiration fluxes and overall soil C stock to distinguish FACE C from pre-FACE C. We solve for fraction pre-FACE C (Eq. $S_6$): 

\[
f_{\text{pre-FACE}} = \frac{\delta ^{13}C_{\text{flux}} - \delta ^{13}C_{\text{new}}}{\delta ^{13}C_{\text{amb}} - \delta ^{13}C_{\text{new}}}, \tag{S6}
\]

where $\delta ^{13}C_{\text{flux}}$ is the $\delta ^{13}C$ signature of respired CO2 (or soil C) from elevated CO2 soils, $\delta ^{13}C_{\text{amb}}$ is the $\delta ^{13}C$ signature of respired CO2 (or soil C) from ambient CO2 soils, and $\delta ^{13}C_{\text{new}}$ is the $\delta ^{13}C$ signature of new photosynthate under CO2 enrichment [Duke, $-39\%$ (10); Aspen, $-39\%$ (4)]. The model is very sensitive to the selection of the $\delta ^{13}C_{\text{new}}$ endmember. The $\delta ^{13}C$ of microbial respiration has been observed to change with both temperature and time in incubation (11, 19). A 2% change in the value of $\delta ^{13}C_{\text{new}}$ results in a 20–50% change in the calculated proportion of pre-FACE C contributing to respiration. Instead, we report the error on the fraction of pre-FACE C in Table 2 as the SEM $\delta ^{13}C$ values of mixing model components, which was propagated by the method in the work by Phillips and Gregg (17).

1.9 Cumulative Respiration Calculations. To correct for potential substrate limitation, we compared the same amount of C respired across temperatures (20). We chose a target amount of total C respired for which the isotopic signature could be consistently estimated across temperature treatments (2.68% of initial C respired for Duke and 0.11% of initial C respired for Aspen). We used periodic measurements of respiration and isotopes and linear interpolation between these measurements to determine cumulative amount of C respired and $\Delta ^{14}C$ values when required.

1.9 Modeling of Data. All nonlinear model fits to data were performed using the Marquardt method in PROC NLIN of SAS version 9.2.

1.9.1 Modeling pool sizes and decay rates using fluxes over time. We fit a two-pool exponential decay model to fluxes from each temperature and CO2 treatment combination in the Duke FACE soils, which was similar to the method described in the work by Paul et al. (21) (Eq. $S7$):

\[
R_t = k_pk_{at}e^{-k_pt} + k_d(1-C_{at})e^{-k_dt}. \tag{S7}
\]

Fluxes ($R_t$; units of $\mu g$ $C_{active}$ g $C_{col}^{-1}$ d$^{-1}$) were measured nine times over the course of the 373 d incubation for each of the four replicates, and these data were used to fit $k_p$ (decay rate of active pool; units of d$^{-1}$), $k_d$ (decay rate of slow pool; units of d$^{-1}$), and $C_{at}$ (size of active pool; units of $\mu g$ $C_{active}$ g $C_{col}^{-1}$).

No model was fit to Aspen data; there was no trend in fluxes over time (linear regression slope of fluxes against time was not significantly different from zero).

1.9.2 Modeling the temperature effect on partitioned fluxes. To describe the exponential dependence of flux on temperature, we fit an exponential model (11) to FACE label-partitioned fluxes [$R_{>10}$ $S_9(T)$, $R_{<10}$ $S_9(T)$] for each site and CO2 treatment (Eq. $S8$): 

\[
R_{>10}(T) = A_{>10}e^{b_{>10}T} \tag{S8}
\]

and (Eq. $S9$) 

\[
R_{<10}(T) = A_{<10}e^{b_{<10}T}. \tag{S9}
\]

Temperature dependence can be quantified as (Eq. $S10$)

\[
\frac{dr}{dT} = b \ast R. \tag{S10}
\]

where $b$ is the temperature sensitivity coefficient. We fit $A$ and $b$ separately for pre-FACE flux ($R_{>10}$) and FACE flux ($R_{<10}$). We report best fits and SE estimates from the nonlinear fit procedure.

1.11 Estimates of Decades-Old C. To determine how much of the overall 0–15 cm mineral soil C pool is vulnerable at both sites, we related the isotopic identity of fluxes to measurable C pools in the soil. Respiration fluxes consist of annually to decadal cycling C (Table 2), and therefore, we have to determine how much of the soil C stock is cycling on these timescales. We used different methods to assess the fraction of soil C stocks vulnerable to decomposition at Duke and Aspen sites because of differences in the data available and inherent soil C age structure between the two sites. Modeling C pools at both the sites using bomb radiocarbon incorporation is complicated by the relatively recent plowing history and young stand ages. Aspen is farther from steady state than Duke, because site disturbance is more recent.

1.11.1 Duke. Using the rate of addition of FACE-derived C into density and size fractions, the work by Lichter et al. (2) estimated the turnover time of pools in the Duke FACE soil. To relate these experimentally determined pools to the $\Delta ^{14}C$ values of respiration, we measured the $\Delta ^{14}C$ values of size and density fractions of 0–5 cm soils taken in July of 2008 from the Duke FACE experiment plots (Fig. 5). All fractions at the Duke FACE site are dominated by bomb-derived C (i.e., decadal or younger) with the exception of the >250-µm size fraction, which is dominated by prebomb C (i.e., millennial C). We estimated the size of the decadal soil carbon pool by subtracting carbon in the >250-µm size fraction for 0- to 15-cm depth from the total soil C stock at this depth.

This estimate is in accord with a bomb radiocarbon model for soil at the Duke site, consisting of 25% of C with a mean residence time of 3,700 y and 75% of C with a mean residence time of 25 y (22).

1.11.2 Aspen. Because of its recent disturbance history and young stand age, the Aspen site is far from steady state, which complicates turnover time calculations for soil C fractions. All size fractions experienced an increase in FACE-derived C over 4 y of the elevated CO2 treatment, meaning that none of them are completely inert (23). However, stability of pools can be inferred from changes in the amount of pre-FACE C; the work by Hofmockel et al. (23) showed a significant decrease in the amount of pre-FACE C in the fine particulate organic matter (fPOM; 53–250 µm) and mineral-associated organic matter (MAOM; 53–3µm) size fractions, but no change in the amount of pre-FACE C in the coarse particulate organic matter (>250 µm) fraction.

The work by Hofmockel et al. (23) also showed no time trend in total fPOM or MAOM stocks, suggesting that the pre-FACE components of C in the fPOM and MAOM fractions are indeed single homogeneous pools at steady state with respect to the total carbon. This finding means that we can assess the relative stability of these pools by dividing the stock by the flux to confirm that these pools are turning over on the order of decades (14 y for fPOM and 40 y for MAOM). If the pre-FACE fraction of C in coarse particulate organic matter soil size class represents

[38x4]Hopkins et al. www.pnas.org/cgi/content/short/1120603109

3 of 5
a stable pool, then we can subtract it from the total soil C to get an estimate of the maximum vulnerability of soil carbon at Aspen FACE for 0–15 cm soils: 94%.

Given the very low flux rates at Aspen FACE compared with Duke FACE, we can use another approach to estimate the minimum amount of decades-old C in the Aspen FACE soils. In this study, we partitioned CO₂ fluxes into FACE and pre-FACE components using the Δ₁⁴C of respired CO₂. We also partitioned the bulk soil C stock into FACE and pre-FACE components using δ¹³C, showing that C predating the FACE treatment makes up 70% of the soil C pool at the 0- to 15-cm depth but only 30% of the flux.

Differences in the cycling rates of the fractions show that it is unlikely that all pre-FACE soil C is a homogeneous reservoir (23). We can assume that only a small part of total soil pre-FACE C is contributing to CO₂ flux and that the minimum turnover time for that component of pre-FACE C is 10 y (in keeping with the fact that this pool is >10 y old and ignoring any lags in plant tissue). We then multiply the measured flux of pre-FACE C by this turnover time to get an estimate for the smallest possible fraction of pre-FACE C that is contributing to respiration. Added to the stock of FACE C (which we assume is all decadal or younger), we estimate a minimum of 53% of the Aspen stock cycling on decades or shorter timescales.


Fig. S1. CO₂ fluxes (micrograms C grams soil⁻¹ day⁻¹) over 373 d of incubation of Duke FACE soils. Solid symbols represent fluxes from the ambient CO₂ treatment, and open symbols represent fluxes from the elevated CO₂ treatment. Colors (blue, 15 °C; yellow, 25 °C; red, 35 °C) denote incubation temperature. Error bars represent the SEM of samples from four replicate plots for each treatment.
Fig. S2. Schematic representation of cumulative C respired (percent of initial soil C) by time into incubation. An equivalent amount of respired C was selected across temperature treatments, which is represented by the dashed line, and the isotopes of respiration were compared for the time period for which this C amount was respired.

Table S1. Two-pool model fit to flux time series data (Duke FACE only)

<table>
<thead>
<tr>
<th>Temperature (°C)</th>
<th>$k_a$</th>
<th>$k_s$</th>
<th>$C_a$</th>
<th>Temperature (°C)</th>
<th>$k_a$</th>
<th>$k_s$</th>
<th>$C_a$</th>
</tr>
</thead>
<tbody>
<tr>
<td>15</td>
<td>17 (2)</td>
<td>0.09 (0.01)</td>
<td>18 (1)</td>
<td>15</td>
<td>16 (3)</td>
<td>0.09 (0.01)</td>
<td>25 (3)</td>
</tr>
<tr>
<td>25</td>
<td>11 (1)</td>
<td>0.14 (0.01)</td>
<td>44 (4)</td>
<td>25</td>
<td>7 (3)</td>
<td>0.14 (0.04)</td>
<td>64 (21)</td>
</tr>
<tr>
<td>35</td>
<td>13 (1)</td>
<td>0.20 (0.02)</td>
<td>90 (6)</td>
<td>35</td>
<td>11 (3)</td>
<td>0.17 (0.05)</td>
<td>86 (16)</td>
</tr>
</tbody>
</table>

$k_a$ (year$^{-1}$) is the decay constant of the active pool, $k_s$ (year$^{-1}$) is the decay constant of the slow pool, and $C_a$ (milligrams C active grams C soil$^{-1}$) is the size of the active pool. Parameters were estimated for four replicate samples (±SEM).

Table S2. Soil carbon fractions (density and size separated) from Aspen and Duke FACE sites

<table>
<thead>
<tr>
<th>Fraction</th>
<th>C amount (g C m$^{-2}$)</th>
<th>$f_{&gt;10 \mu m}$ by $^{13}$C</th>
<th>C amount (g C m$^{-2}$)</th>
<th>$f_{&gt;10 \mu m}$ by $^{13}$C</th>
</tr>
</thead>
<tbody>
<tr>
<td>Light fraction</td>
<td>Ambient CO$_2$</td>
<td>Elevated CO$_2$</td>
<td>1,431</td>
<td>Elevated CO$_2$</td>
</tr>
<tr>
<td>&gt;250 μm</td>
<td>480</td>
<td>460</td>
<td>0.50</td>
<td>473</td>
</tr>
<tr>
<td>53–250 μm</td>
<td>510</td>
<td>520</td>
<td>0.55</td>
<td>371</td>
</tr>
<tr>
<td>&lt;53 μm</td>
<td>2,900</td>
<td>2,800</td>
<td>0.61</td>
<td>326</td>
</tr>
</tbody>
</table>

Data from the works by Hofmockel et al. (23) and Lichter et al. (3).