The effect of carbon fertilization on naturally regenerated and planted US forests

Over the last half century in the United States, the per-hectare volume of wood in trees has increased, but it is not clear whether this increase has been driven by forest management, forest recovery from past land uses, such as agriculture, or other environmental factors such as elevated carbon dioxide, nitrogen deposition, or climate change. This paper uses empirical analysis to estimate the effect of elevated carbon dioxide on aboveground wood volume in temperate forests of the United States. To accomplish this, we employ matching techniques that allow us to disentangle the effects of elevated carbon dioxide from other environmental factors affecting wood volume and to estimate the effects separately for planted and natural stands. We show that elevated carbon dioxide has had a strong and consistently positive effect on wood volume while other environmental factors yielded a mix of both positive and negative effects. This study, by enabling a better understanding of how elevated carbon dioxide and other anthropogenic factors are influencing forest stocks, can help policymakers and other stakeholders better account for the role of forests in Nationally Determined Contributions and global mitigation pathways to achieve a 1.5 degree Celsius target.

In the United States, over the last two decades, a large forest C-sink has sequestered 700–800 million tons of CO2 per year, which is roughly 10–11% of US gross CO2 emissions\(^1\). In addition, the per-hectare volume of trees has increased over the last 50 years\(^6\). Among the ten forest groups in this study, all except Aspen-Birch have increased their per-hectare wood volume from 1997 to 2017 (Fig. 1)\(^5\). It is not clear, however, whether the tree-volume and C-sink increase have been driven by forest recovery from past land uses, such as agriculture\(^7\), or other environmental factors such as elevated CO2, N deposition, or climate change\(^8\). Even less well understood is the role of forest management through planting, harvesting of secondary and old-growth forests, and forest management that is certified with sustainability criteria, even though these approaches are employed on an increasingly large share of the world’s forests\(^9\). As policymakers look to the future, where forests are expected to play a large role in Nationally Determined Contributions (NDCs) and global mitigation pathways to achieve a 1.5 °C target\(^10\), it is important to better understand how anthropogenic factors like elevated CO2 may influence current forest stocks and the outcomes of forest-expansion (e.g., afforestation) and conservation policies (e.g., avoided deforestation).

This analysis builds on the experimental and modeling literature that has examined the role of elevated CO2, like the FACE studies that controlled CO2 levels on plots within the same location and found that elevated CO2 increased net primary productivity (NPP)\(^14,15\). The effects these studies observed, however, might not scale up across ecosystems and over time with disturbances and other processes at play. Tree ring studies\(^16,17\) offer the potential to observe the effects of elevated CO2, but with limited observations and controls, they could not identify the effects of elevated CO2 separately from other factors, such as weather. Other research has shown that tree heights in Poland increased over time, although the role of CO2 concentration was not identified\(^18\), and that CO2 exposure generates smaller effects on tree-volume in settings that are colder and have more water stress\(^19\). A recent meta-analysis of experimental results estimated that each
100-ppm increase in CO2 increases aboveground volume in ecosystems by 8.2%\textsuperscript{20}. These experimental results underpin many dynamic global vegetation models (DGVMs), but they have not been replicated with observations from plots drawn randomly within ecosystems. This study addresses that gap by using quasi-experimental techniques\textsuperscript{21} and data drawn from tens of thousands of plots in the United States to identify, specifically and differentially, the role of elevated CO2 from other factors that also influence wood volume in forest stands.

We show that there has been a positive response in wood volume to elevated CO2 in ten temperate forest groups in the United States. Moreover, our estimated response in ecosystems is larger than that predicted by experimental results. Those estimates, however, included non-forested ecosystems, which do not appear to respond as strongly as forests to elevated CO2. For the forest groups in our study with both planted and naturally regenerated observations, we find that the proportional impact of elevated CO2 on planted stands is roughly equivalent to the impact in natural stands, but the results in cubic meters per hectare are mostly larger for planted stands because planted stands, on average, have more volume than naturally regenerated stands.

**Results**

This study estimates the impact of elevated CO2 on wood volume in the central stems of trees by comparing growing-stock volume on recently observed timber plots with matched plots observed decades prior using historical data from the US Forest Service (USFS) Forest Inventory and Analysis Program (FIA)\textsuperscript{22}. A key challenge in understanding how elevated CO2 affects the wood volume observed in forest plots is that CO2 concentrations change only temporally but not spatially. That means that all forests are exposed to the same CO2 concentration each year. Our research design overcomes this challenge by utilizing the fact that forests embed the historical CO2 concentrations to which they have been exposed. Thus, two plots of the same age and type of forest observed at two different time periods have effectively received two different CO2-exposure profiles. This is also true when two plots of the same type of forest are observed at the same time period, but their ages differ.

Since other factors such as climate, technology, pests, and tree stocking also affect the volume and vary over time, we use modern econometric techniques that combine matching with post-matching regression analysis, which includes time-fixed effects, to isolate the effects of elevated CO2 from other time-varying and time-invariant covariates that might also affect tree volumes\textsuperscript{23}. These matching methods\textsuperscript{24} systematically construct treatment and control groups of timber plots that are similar in observable characteristics, but that vary in their exposure to CO2 due to the observations having been taken in different decades.

Forests provide several advantages to facilitate empirical identification of the impact of elevated CO2 on tree wood volume. First, volume and management decisions on US forests have been systematically monitored and measured for many decades through the plot-level FIA database. Second, the FIA data indicate whether forest plots were regenerated naturally or through active planting. While the wood volume of several species of commercially planted conifers have been affected by changing seed technology over the last few decades, naturally regenerated forests—including most hardwoods—have not been influenced by such technological changes that would confound the identification of the impact of elevated CO2.

We examine ten temperate forest-type groups in the United States whose range in the FIA database is shown in Fig. 2. First, we focus only on observations of naturally regenerated (unmanaged) forest plots because natural stands arguably have not been affected by the advances in tree planting and seed selection that have improved the quality of planted stands over the same time period. To address the potential for bias in our data, we follow recent advances in quasi-experimental econometrics and “trim” the data using matching methods\textsuperscript{25} to create a dataset of one-to-one matches between a control group comprised of observations taken by the USFS during the pre-1990 (low CO2) period and a treatment group whose observations were made in the post-2000 (high CO2) period, while controlling for multiple salient covariates that are correlated with time and/or could affect yield, such as age and stocking conditions (Supplementary Tables 1–6 and Supplementary Data 1, 2). A variety of alternative functional forms are tested (Supplementary Data 3–10), with results robust across specifications. The effect of elevated CO2 is identified by the natural log of lifetime CO2, which is the logged sum of the annual CO2 concentrations experienced by each stand-up to its age class at the time the plot is measured. Climate-related effects on wood volume are captured with seasonal temperature and precipitation variables.
Fig. 2 | Geographic range of forest groups based on observations taken by USFS. Note: This figure for each forest group details (in green) all counties in which the US Forest Service Forest Inventory and Analysis (USFS-FIA) database has recorded the forest group’s presence between 1968 and 2018. This is based on its annual resource inventories and is limited to observations of stands between 1 and 100 years of age.
To strengthen the identification of elevated CO$_2$ other environmental factors correlated with time, such as nitrogen deposition, large-scale disturbances associated with invasive species (e.g., Emerald Ash Borer, Agrilus planipennis Fairmaire), and widespread phenomena like Sudden Aspen Death, are captured by including time-dummy variables (fixed effects). For Table 1, the dummy variable is designed to capture changes between the pre-1970 and the post-2000 periods that are not tied to the gradually increasing CO$_2$ trendline (e.g., the impact of nitrogen deposition). The approach is detailed in Model (1) of Supplementary Data 11 and the multivariate-regression results are presented as Model (1) in Supplementary Data 12–22. In six of the ten forest groups, two-sided t-tests indicate that the time-dummy coefficients are significant with two forest groups positively impacted and four negatively (Table 1). Aspen/Birch appears to have experienced the largest decline over time (−18.0%; P < 0.01; 99% CI from −30.8 to −5.2%; t = −3.62; df = 16,416), a result potentially due to aspen decline. Emerald Ash/Cottonwood also decreases (−11.8%; P < 0.01; 99% CI from −23.1 to −0.6%; t = −2.7; df = 7,562), a result that might be the result of the Emerald Ash Borer that has plagued Ash trees in the Eastern US from 2002 to the present. White/Red/Jack pine shows the largest increase (21.2%; P < 0.01; 99% CI from 0.1 to 42.2%; t = 2.6; df = 12,458) while Spruce/Fir experienced (9.0% CI from 1.1 to 17.2; df = 15,726). Aspen/Birch also increases (P = 0.01; 99% CI from 0.0 to 18.9%; t = 2.7; df = 16,416) due to the spruce budworm (Choristoneura fumiferana) outbreaks in the 1960s and early 1980s. Overall, across all forests, these large-scale changes from episodic phenomena have reduced wood volume by an average of 8.9% (P < 0.01; 99% CI from −10.7 to −7.0; t = −12.4; df = 123,145).

Given that the Western US faces different threats and disturbances than the Eastern US, we repeat the process used to create Table 1 with only plots from the 33 states the USFS considers part of the East. Specifically, this comprises all states fully east of the 100th meridian plus Texas and Oklahoma. Balance statistics are provided in Supplementary Data 23. Supplementary Data 11 details the models used for each forest group individually (Model 3) and the combined sample (Model 4). Regression results are presented in Supplementary Data 24, 25. To enable ease of comparison with Table 1, a revised version for the Eastern US is also included (Supplementary Table 7). Results for the four forest groups whose original match results had no observations from the Western US, of course, are identical. Among the remaining six, there are slight changes in the impact of elevated CO$_2$. At age 75, Elm/Ash/Cottonwood increases from 10.0% (16.0 m$^3$ ha$^{-1}$) for all observations to 11.0% (18.0 m$^3$ ha$^{-1}$); P < 0.01; 99% CI from 7.4 to 17.2%; t = 4.1; df = 6,724) for observations in the Eastern US. Oak/Pine increases from 12.6% (25.1 m$^3$ ha$^{-1}$) to 13.1% (26.7 m$^3$ ha$^{-1}$); P < 0.01; 99% CI from 7.4 to 18.7%; t = 5.4; df = 9,914), and Aspen/Birch increases from 12.1% (17.6 m$^3$ ha$^{-1}$) to 12.4% (18.1 m$^3$ ha$^{-1}$); P < 0.01; 99% CI from 8.0 to 16.8%; t = 6.6; df = 16,348). Maple/Beech/Birch increases from 9.9% (15.6 m$^3$ ha$^{-1}$) to 10.0% (15.8 m$^3$ ha$^{-1}$); P < 0.01; 99% CI from 5.4 to 14.7%; t = 5.0; df = 17,340). Spruce/Fir decreases from 10.6% (19.3 m$^3$ ha$^{-1}$) to 10.3% (18.6 m$^3$ ha$^{-1}$); P = 0.01; 99% CI from 5.9 to 14.9%; t = 3.0; df = 19,493). Elm/Ash/Cottonwood, the largest increase over time (−30.8 to −28.9%), has a 33.1% (51.1 m$^3$ ha$^{-1}$) impact of elevated CO$_2$ exposure parameter indicates that each 1% increase in lifetime CO$_2$ exposure, which is an additional 3.2–3.3 ppm CO$_2$. For each additional 1% increase in CO$_2$ exposure from 1970 to 2015, cumulative lifetime CO$_2$ increased 11.1% for 75-year-old stands, whereas for 25-year-old stands, it increased 18.9%. For each 1% increase in lifetime CO$_2$ exposure from 1970 to 2015, cumulative lifetime CO$_2$ increased 11.1% for 75-year-old stands, whereas for 25-year-old stands, it increased 18.9% (Supplementary Data 26, 27 and the impacts are similar in sign and magnitude to those shown in Table 1 (Supplementary Tables 8, 9).

We next examine the impact of elevated CO$_2$ separately for naturally regenerated and planted stands for the three forest groups where sufficient data of both types is available (Supplementary Data 11: Model 7). Since many replanted stands are harvested in relatively short timber rotations, meaning there are few stands older than age 50, we limit the data for both to ages less than or equal to 50 years to better compare the different stands. Because of this change in age classes, we create new matches for both naturally regenerated and planted stands (Supplementary Data 28–30) and then run new post-matching regressions (Supplementary Data 31–33). Results indicate that elevated CO$_2$ has positive effects (P < 0.01) on wood volume for each forest type examined on both natural and planted stands. Based on these regressions, the predicted change in volume due to elevated CO$_2$ at 25 years of age for planted stands in 1970 versus 2015 (Table 2) for Slash/Longleaf pine is +15.5 m$^3$ ha$^{-1}$ (+21.7%; P < 0.05; 99% CI from +1.8 to +29.3 m$^3$ ha$^{-1}$; t = 1.9; df = 859). For Loblolly/Shortleaf pine, the change is +34.6 m$^3$ ha$^{-1}$ (+27.4%; P < 0.05; 99% CI from +17.4 to +51.8 m$^3$ ha$^{-1}$; t = 3.9; df = 2,757), and for White/Red/Jack pine, the change is +16.0 m$^3$ ha$^{-1}$ (+42.8%; P < 0.05; 99% CI from +2.1 to +29.8 m$^3$ ha$^{-1}$; t = 2.3; df = 671). For naturally regenerated stands, the gains are smaller. For Slash/Longleaf pine, the change is +12.5 m$^3$ ha$^{-1}$ (+21.7%; P < 0.05; 99% CI from +4.3 to +20.7 m$^3$ ha$^{-1}$; t = 2.5; df = 1,981), for Loblolly/Shortleaf pine, the change is +21.4 m$^3$ ha$^{-1}$ (+28.8%; P < 0.05; 99% CI from +15.9 to +26.9 m$^3$ ha$^{-1}$; t = 7.6; df = 10,170), and for White/Red/Jack pine, the change is +15.3 m$^3$ ha$^{-1}$ (+29.4%; P < 0.05; 99% CI from +4.5 to +26.1 m$^3$ ha$^{-1}$; t = 2.8; df = 1,280). The results for natural stands are consistent with the results presented in Table 1 that are derived from all observations aged 1 to 100.

The size of the coefficient on the natural log of lifetime CO$_2$ for natural stands is larger though for the analysis using forests aged 1 to 50 than the analysis for Table 1 using forests aged 1 to 100. For Loblolly/Shortleaf, using only observations aged 1 to 50, the parameter
Table 1 | Change in predicted wood volume due to elevated CO₂ at ages 25/50/75 and due to other episodic phenomena from 1970 to 2015 using observations of naturally regenerated stands aged 1–100

| Forest Type     | Carbon Fertilization | Episodic Phenomena Mean Impact | Number of Observations |
|-----------------|----------------------|--------------------------------|------------------------|
|                 | 25 years old         | 50 years old                   | 75 years old           | Post-2000 v. Pre-1990 |
|                 | Mean | CI | Mean | CI | Mean | CI |                      |                        |
| White/Red/Δ (%) | 23.6 | (9.7, 37.6) | 18.0 | (4.0, 31.9) | 14.4 | (0.4, 28.3) | Δ (%) 21.2          | 2362                   |
| Jack Pine Δ (m³/ha) | 12.9 | (5.3, 20.6) | 18.3 | (4.1, 32.5) | 26.3 | (0.7, 51.9) | Sig. **              |                        |
| Spruce/Δ (%)    | 17.4 | (5.0, 29.9) | 13.2 | (0.8, 25.7) | 10.6 | (1.8, 19.4) | Δ (%) 19.2          | 4980                   |
| Fir Δ (m³/ha)   | 5.8  | (1.6, 9.9)  | 9.0  | (0.5, 17.5) | 10.0 | (1.7, 18.4) | Sig. *              |                        |
| Slath/Δ (%)     | 21.8 | (10.3, 33.3) | 16.6 | (5.0, 28.1) | 13.2 | (1.7, 24.8) | Δ (%) 9.8           | 3582                   |
| Longleaf Pine Δ (m³/ha) | 11.2 | (5.3, 17.2) | 17.3 | (6.3, 29.3) | 22.5 | (2.9, 42.1) | Sig. **              | 15,188                 |
| Lobolly/Δ (%)   | 21.9 | (17.1, 26.8) | 16.7 | (11.8, 21.5) | 13.3 | (8.5, 18.2) | Δ (%) 15.3         | 268                    |
| Shortleaf Pine Δ (m³/ha) | 17.5 | (13.6, 21.3) | 31.1 | (22.1, 40.1) | 40.1 | (25.5, 54.6) | Sig.                |                        |
| Oak/Δ (%)       | 20.7 | (15.2, 26.3) | 15.8 | (10.2, 21.3) | 12.6 | (7.1, 18.1) | Δ (%) 12.5         | 9,780                  |
| Pine Δ (m³/ha)  | 12.2 | (8.0, 15.5)  | 18.4 | (11.6, 26.2) | 25.1 | (14.1, 36.2) | Sig.                |                        |
| Oak/Δ (%)       | 19.0 | (16.4, 21.5) | 14.4 | (11.9, 16.9) | 11.5 | (9.0, 14.1) | Δ (%) 6.2          | 39,268                 |
| Hickory Δ (m³/ha) | 11.2 | (9.7, 12.6)  | 16.2 | (13.3, 18.9) | 19.2 | (15.4, 24)  | Sig. ***            | 11,446                 |
| Oak/Gum/Δ (%)   | 22.4 | (16.6, 28.1) | 17.0 | (11.2, 22.8) | 13.6 | (7.8, 19.4) | Δ (%) 2.9          | 7,618                  |
| Cypress Δ (m³/ha) | 17.7 | (13.2, 22.3) | 25.3 | (16.7, 33.8) | 33.0 | (19.0, 47.0) | Sig.                |                        |
| Elm/Ash/Δ (%)   | 16.5 | (10.6, 22.4) | 12.5 | (6.6, 18.4)  | 10.0 | (4.1, 15.9) | Δ (%) 6.8          | 12,514                 |
| Cottonwood Δ (m³/ha) | 10.3 | (6.6, 14.0)  | 14.4 | (7.6, 21.2)  | 16.0 | (6.6, 25.4) | Sig. ***            | 7,618                  |
| Maple/Beech/Δ (%) | 16.4 | (11.7, 21.0) | 12.4 | (7.8, 17.1)  | 9.9  | (5.3, 16.4) | Δ (%) 1.6          | 12,514                 |
| Birch Δ (m³/ha)  | 10.2 | (7.3, 13.1)  | 13.7 | (8.6, 18.8)  | 16.8 | (8.4, 22.9) | Sig.                |                        |
| Aspen/Δ (%)     | 19.9 | (15.5, 24.2) | 15.1 | (10.7, 19.4) | 12.1 | (7.7, 16.4) | Δ (%) 18.0         | 16,472                 |
| Birch Δ (m³/ha)  | 9.7  | (7.6, 11.8)  | 14.6 | (10.4, 18.8) | 17.6 | (11.2, 23.9) | Sig. ***            |                        |
| All Forest Δ (%) | 20.3 | (15.9, 24.6) | 15.4 | (11.0, 19.8) | 12.3 | (7.9, 16.7) | Δ (%) 8.9          | 123,210                |
| Groups Δ (m³/ha) | 11.8 | (9.2, 14.4)  | 18.4 | (13.2, 23.7) | 23.0 | (14.8, 31.2) | Sig. ***            |                        |

Matches were created using observations of naturally regenerated plots with the control being observations from 1968–90 and the treatment being observations from 2000–18. Post-matching, full multivariate regression analysis was performed and the p value for the CO₂ variable was less than 0.01 for each forest group individually and also collectively. Then, the carbon fertilization effect from 1970 to 2015 was analyzed using a one-sided t-test to assess the significance of the change in CO₂ exposure from 1970 to 2015 on wood volume at 25/50/75 years of age (i.e., a 25-year-old stand in 1970 received the sum of yearly exposure values from 1946 to 1970 and a 25-year-old stand in 2015 received the sum of yearly exposure values from 1991 to 2015). Climate variables were held at their 1970 levels. The impact of the other episodic phenomena was captured by the dummy variable comparing observations pre-1990 with those post-2000. Using a one-sided t-test to test the hypothesis that volume in 2015 had increased significantly from the volume in 1970, Table 1 displays a 99% confidence interval for all but Spruce/Fir at 75 years of age, where a 95% CI is shown. Model (1) results (Supplementary Data 12–22) were used as inputs in these calculations. ***P < 0.01, **P < 0.05, and *P < 0.10.
on the carbon-fertilization variable is 1.66 (P < 0.01; 99% CI from 1.59 to 1.73; t = 43.8), whereas, when using observations aged 1 to 100, the parameter is smaller at 1.29 (P < 0.01; 99% CI from 1.25 to 1.33; t = 59.9) (Supplementary Table 10). Likewise, for White/Red/Jack pine, the values are 1.70 (P < 0.01; 99% CI from 1.49 to 1.90; t = 16.1) and 1.38 (P < 0.01; 99% CI from 1.26 to 1.50; t = 21.9) (Supplementary Table 10), respectively. These larger parameters, when using just the observations of younger stands, suggest that the effect of carbon fertilization could attenuate over time for some forest groups. For Slash/Longleaf though, the parameter values are relatively consistent at 1.25 (P < 0.01; 99% CI from 1.08 to 1.42; t = 14.5) and 1.27 (P < 0.01; 99% CI from 1.16 to 1.37; t = 23.9) (Supplementary Table 10), respectively.

### Discussion

Our estimated response of wood volume in ecosystems to elevated CO2 is stronger than that predicted by experimental results. Over the period 1970 to 2015, CO2 concentrations increased by 75 ppm, and we find that this increase in CO2 stimulated an increase in wood volume in naturally regenerated 75-year-old forests in the United States by 12.3% (7.5 and 17.1%; 99% CI). This is larger than the 8.2% effect found in experimental studies20, although those estimates included non-forested ecosystems, which do not appear to respond as strongly as wood volume to elevated CO2. Also, our results are for temperate forest groups that exist largely in parts of the Eastern US where the climate is mostly warm and wet. This is important given the recent finding that, in settings that are colder and have more water stress, CO2 exposure generates smaller effects on tree-volume20.

For the southern pine forest groups, our estimates indicate that the proportional response of wood volume in planted stands is roughly equivalent to the response in natural stands, but the results in cubic meters per hectare are larger for planted stands because these planted stands have more volume on average than naturally regenerated stands. For White/Red/Jack pine, our estimates indicate a different pattern. The proportional response of wood volume in planted stands is far larger than the response in natural stands, but the results in cubic meters per hectare are smaller for planted stands early in life. Potentially, there are some management activities associated with the replanting of White/Red/Jack pine, such as competition management, that initially reduce biomass. Using the results for 25-year-old stands (Table 2), the 75-ppm increase in CO2 between 1970 and 2015 generated a 28.8% (21.4 m3 ha−1) increase in wood volume in 25-year-old, naturally regenerated Loblolly/Shortleaf stands and a 27.4% (34.6 m3 ha−1) increase in planted stands of the same age. This suggests that from 1970 to 2015, wood volume on natural Loblolly/Shortleaf stands increased by about 0.5 m3 ha−1 yr−1 due to elevated CO2, while volume increased by 0.8 m3 ha−1 yr−1 on planted stands. The parameter on logged CO2 for Loblolly/Shortleaf and White/Red/Jack pine is smaller for the regressions using forests aged 1 to 100 than those using only forests aged 1 to 50 (Supplementary Table 10). This suggests that for these two forest groups, carbon fertilization may attenuate over time—that is, the proportional response in wood volume to elevated CO2, when measured with data including older stands, is smaller than when measured with data including only younger stands. Although the proportional effect of elevated CO2 appears to decline for older stands, the additional accumulation of volume in older stands due to elevated CO2 remains substantial nonetheless because older stands have more wood volume.

The methods in this study explicitly control for other time-linked phenomena. Climate change is controlled directly through our experimental design by matching observations before 1990 with those after 2000 and then including polynomial functions of seasonal climate variables in our post-matching estimations. Other episodic factors, such as nitrogen deposition and invasive species, are controlled with dummies variables that account for the year the plot was observed. While we cannot observe these specific factors, the results indicate that, on average, these episodic factors have caused wood volume in naturally regenerated stands to decline. In fact, across all forest groups, these time-related factors reduced volume by 8.9% (P < 0.01) between 1970 and 2015 (Table 1). Only two of the forest groups showed an increasing trend in wood volume that is not attributable to climate change or elevated CO2: White/Red/Jack pine (P < 0.01) and Spruce/Fir (P < 0.10). We hypothesize that these results are both tied to recovery from past natural disturbances.

For planted stands aged 1 to 50, we find a significant negative trend in wood volume for Loblolly/Shortleaf and White/Red/Jack (but not for Longleaf/Slash) pine that is attributed to episodic temporal factors unrelated to elevated CO2 and climate change, but for similarly aged naturally regenerated stands, the effect is smaller or not significant (Supplementary Data 31–33). This outcome suggests that on planted stands, temporal factors unrelated to elevated CO2 and climate change have had a negative influence on wood volume in the last 30 to 40 years. Although we control for stocking conditions in the matching process, increased thinning activity or lower investments in fertilizing after 2006, as timber prices declined and fertilizer prices
increased during the Great Recession, could explain this result. Forest managers, as well, may have shifted management objectives from being largely focused on increasing volume to increasing value.

Overall, our results present evidence that there has been a positive response in wood volume to elevated CO$_2$ in ten temperate forest groups in the United States. Our matching approach, which allows us to identify the effect of elevated CO$_2$, does not enable us to assess how elevated CO$_2$ interacts with its resulting climate impacts, such as changes in temperature, precipitation, forest composition, and range. The pooled modeling approach we include in the robustness analysis (Supplementary Data 3, 4) may provide an opportunity to test these interactions. Our results hint that the effects of elevated CO$_2$ could attenuate in older forests, but we cannot directly test for attenuation with our approach. Future research should address this question.

**Methods**

**Materials**

Information on wood volume and the physical environment of the plots were obtained from the US Forest Service Forest Inventory and Analysis (USFS-FIA)$^{22}$. The FIA database categorizes each plot into one of 33 forest groups, but 23 groups do not have sufficient data in the control period (before 1990) to enable robust matching and so were dropped from this study. As a result, several western forest groups (e.g., Douglas-fir) were not included in our study. The following ten forest groups [(1) Loblolly/Shortleaf Pine, (2) Slash/Shortleaf Pine, (3) White/Red/Jack Pine, (4) Spruce/Fir, (5) Elm/Ash/Cottonwood, (6) Maple/Beech/Birch, (7) Oak/Hickory, (8) Oak/Gum/Cypress, (9) Aspen/Birch, and (10) Oak/Pine] all had more than 5000 observations and large numbers of observations both from before 1990 and from 2000 on. Data for the 48 conterminous states from evaluation years between 1968 and 2018 were included in the study. We limited our analysis to plots with trees from 1 to 100 years of age, resulting in trees that had been planted somewhere between 1869 and 2018—a period during which atmospheric CO$_2$ increased from roughly 287 to more than 406 ppm$^{25,26}$. The geographic distribution of the ten forest groups presented in Fig. 2 shows in orange all counties in which the USFS recorded in at least one year between 1968 and 2018 the presence of a plot of the respective forest group that meet the age requirements for inclusion in this study. Precipitation and temperature data were obtained from the PRISM Climate Group$^{42}$.

**Methods**

Results in Tables 1 and 2 are based on estimated exponential tree-volume functions of the generalized form shown in Eq. 1. The left-hand side is the natural log of the volume per hectare in the central stem of trees on each plot in cubic meters. Volume is assumed to be a function of age, the logged cumulative lifetime concentration of CO$_2$, and other variables, including plot-specific variables that vary across plots but not time ($X_i$), weather variables that vary across plots and time ($W_{it}$), and time-specific fixed effects that vary across time but not plots ($E_{it}$).

$$\ln \left( \frac{\text{Volume}}{\text{Hectare}} \right)_{it} = \alpha + \beta_0 + \frac{1}{\text{Age}_{it}} + \beta_1 \ln\left(\text{CumCO2}[\text{Life},i]\right) + \beta_2 X_i + \beta_3 W_{it} + \beta_4 E_{it} + e_{it}$$ (1)

The nonparametric smearing estimate method was used to transform logged-volume results into a volume in cubic meters per hectare$^{22}$. The climate variables, obtained from the PRISM Climate Group$^{42}$ and described in Supplementary Table 1, enter as cubic polynomials of the lifetime seasonal temperature and precipitation averages that a plot of a given age at a given time experienced.

The variable for atmospheric carbon was constructed as the logarithmic transformation of the sum of yearly atmospheric CO$_2$ exposures over the lifetime of the stand. Other site-specific covariates were obtained from the FIA data (Supplementary Table 2), such as the availability of water, the quality of the soil, the photoperiod of the plot, whether disturbances had impacted the land, and whether the land was publicly or privately owned$^{33,34}$. The time-specific fixed effects ($E_i$) in the model control for episodic factors like nitrogen deposition and invasive species, which are correlated with time but cannot be observed over space for the whole time period. These time-dummy variables account for underlying, unobservable systematic differences between the 21st-century period when atmospheric CO$_2$ was higher and the pre-period when levels were much lower. Controlling for these factors aids the identification of the impact of elevated CO$_2$, which varies annually.

A potential concern is that wood volume changes over time could be related to an increased number of trees per hectare rather than increased wood volume of the trees. To assess whether controls for the stocking condition were needed, we examined data on the number of trees per acre of each forest type. First, we looked at a group of southern states (Supplementary Table 3) and found double-digit percentage changes in tree stocking between 1974 and 2017 for seven of the nine forest groups. However, the changes were mixed, with four having increased tree density and five decreasing tree density. The FIA data do not record the Aspen/Birch forest group as present in these southern states in these evaluations.

The examination of a group of northern states involved a comparison of the average stocking conditions around 1985 with those in 2017. The changes in tree density for these forest types (Supplementary Table 4) were also split with four showing increased stocking and five having less dense stocking. The change for Loblolly/Shortleaf pine was relatively large, with stocking density increasing by 27.2%. Slash/Longleaf was not recorded as present in these states in these evaluations.

Next, we analyzed changes, over the period from around 1985 to 2017, in all states east of the 100th meridian, as those states comprised the bulk of the data in our study (Supplementary Table 5). Results for seven of the ten forest groups showed a less dense composition. Loblolly/Shortleaf pine again was shown to have become more densely stocked, with an increase of 13.2%.

The last check included all of the 48 conterminous states and compared changes in stocking conditions from years around 1985 to 2017 (Supplementary Table 6). Seven of the ten forest groups showed decreased stocking density over time. Not surprisingly (because most Loblolly/Shortleaf is located in the Eastern US), the change in Loblolly/Shortleaf pine density is the same for this check as was shown in the results in Supplementary Table 5. Based on the results from all these comparisons and given that stocking density has changed over time, we controlled for it both in the matching and in the multivariate-regression analysis.

Genetic matching (GM), the primary approach used for this analysis, combines propensity score matching and Mahalanobis matching techniques$^{35}$. The choice of GM was made after initially considering other approaches, such as nearest-neighbor propensity score matching with replacement and a non-matching, pooled regression approach. Three options were tested on the samples for Loblolly/Shortleaf pine and Oak/Hickory, and the regression results are presented in Supplementary Data 3-4.

The results across these different approaches were quite similar, suggesting that the results are not strongly driven by methodological choice. We focused on matching rather than a pooled regression approach to help reduce bias and provide estimates closer to those that would be obtained in a randomized controlled trial. When choosing the specific matching approach, we considered that standard matching methods are equal percent bias reducing (EPBR) only in the unlikely case that the covariate distributions are all roughly normal$^{36}$ and that EPBR may not be desirable, as in the case where one of two covariates has a nonlinear relationship with the dependent variable$^{37}$. We also noted that GM is a matching algorithm that at each step
minimizes the largest bias distance of the covariates and that GM has been shown to be a more efficient estimator than other methods like the inverse probability of treatment weighting and one-to-one greedy nearest-neighbor matching. Additionally, when the distributions of covariates are non-ellipsoidal, this nonparametric method has been shown to minimize bias that may not be captured by simple minimization of mean differences. Lastly, as sample size increases, this approach will converge to a solution that reduces imbalance more than techniques like full or greedy matching. Given the support that this choice has in the literature, we decided to employ GM to create all the matched data used in this study using R software.

Artificial regeneration of forest stands, noted as planting throughout the text is used as the main proxy for the impact of forest management. The other indicator of management activity is what can be described as interventions, which are a range of human on-site activities that the USFS details. We define unmanaged land as stands with natural regeneration and where no interventions occurred on the plot.

To create Table 1, we first excluded all plots on which there had been either planting activities or some type of human intervention. Then, we created treatment and control groups by forming two time periods separated by an intervening period of ten years to ensure a more than a marginal difference between the groups in terms of lifetime exposure to atmospheric CO2. The control period used forest plot data sampled between 1968 and 1990, and the treatment period used forest plots sampled between 2000 and 2018. Note that even though the earlier period contains more years, there are fewer overall observations.

Matches were then made to balance the treatment and control groups based on the following observable covariates: (1) Seasonal Temperature, (2) Seasonal Precipitation, (3) Stocking Condition, (4) Aspect, (5) Age, (6) Physiographic Class, and (7) Site Class. The propensity score was defined as a logit function of the above covariates to generate estimates of the probability of treatment. Callipers with widths less than or equal to 0.2 standard deviations of the propensity score were also employed to remove at least 98% of bias.

Balance statistics for the primary covariates are presented in Supplementary Data 1–2 and show a strong balance for all covariates across all forest groups. Thus for each forest group, our sample of plots includes control plots (pre-1990) and treatment plots (post-2000) that are comparable (balanced) in climate and other biophysical attributes.

After trimming our sample using this matching process and obtaining strongly balanced matches, we turned to regression analysis, where we employed Stata software. To confirm that we had the most appropriate model structure, tests of the climate and atmospheric carbon variables were undertaken using various polynomial forms, and the main variable of interest, atmospheric carbon, was tested both with and without transformation of that variable. Results (Supplementary Data 5–10) show that the climate variables were not improved with complexity beyond cubic form. Moreover, selection tools, like the Akaike and Bayesian information criterion, favored the cubic choice, and so we utilized the cubic formulation throughout this study. Results for the CO2 variable were similar in both sign and significance for the linear and logged form. We use the logged form as it allows easier interpretation of the effect, suppresses heteroscedasticity, and removes the assumption that each unit increase in CO2 exposure will have a linear (constant) effect on volume.

The estimated effect of CO2 exposure for each forest group (Supplementary Data 12–21) was estimated using alternate specifications of the independent variables included in Eq. 1. For each forest type, the Model (1) specification (Eq. 2) is the basis for the results presented in Table 1. The β0 coefficient details the impact on the volume of the main variable of interest, atmospheric carbon.

\[
\ln(\text{volume per hectare}) = a + \beta_0 \ln(\text{LifetimeCO2}) + \beta_1 \frac{1}{\text{Age}} + \beta_2 \text{Site Class} + \beta_3 \text{Seasonal Temperature} + \beta_4 \text{Seasonal Temp}^2 + \beta_5 \text{Seasonal Temp}^3 + \beta_6 \text{Seasonal Precipitation} + \beta_7 \text{Seasonal Precip}^2 + \beta_8 \text{Seasonal Precip}^3 + \beta_9 \text{Stocking} + \beta_{10} \text{Disturbances} + \beta_{11} \text{Physiographic Class} + \beta_{12} \text{Aspect} + \beta_{13} \text{Slope} + \beta_{14} \text{Elevation} + \beta_{15} \text{Latitude} + \beta_{16} \text{Longitude} + \beta_{17} \text{Ownership} + \beta_{18} \text{Time Dummies} + \beta_{19} \text{Seasonal Vapor Pressure Deficit} + \beta_{20} \text{Length of Growing Season} + \epsilon
\]

After estimating Eq. 2 for each forest type individually (Supplementary Data 12–21), all plots were pooled across forest groups, with additional forest-group dummy variables, to estimate a general tree-volume function (Supplementary Data 22).

Our main Model (1) results are provided in Supplementary Data 12–22, along with three additional models that assess the robustness of the elevated CO2 effect to different specifications. The simplest specification, Model (4), included only stand age, CO2 exposure, and a time-dummy variable. Model (3) took the Model (4) base and added in an array of site-specific variables, including those for the climate. Model (2) was similar to Model (1) in that it included the impact of vapor pressure deficit and the length of the growing season on the variables included in Model (3), but it differed from Model (1) in that it tested an alternate approach to capturing the impact of underlying, unobservable systematic differences like nitrogen deposition.

Using the estimated coefficients from the preferred Model (column 1) specification (Eq. 2), the estimated change in growing-stock volume between two CO2 exposure scenarios was calculated at ages 25, 50, and 75. The first scenario examined CO2 exposure up to 1970 (that is, when calculating growing-stock volume for a 25-year-old stand, the CO2 exposure would have the summation of the yearly values for the years from 1946 to 1970 [310 to 326 ppm CO2]). The second scenario examined CO2 exposure up to 2015 (that is, when calculating growing-stock volume for a 25-year-old stand, the CO2 exposure was the summation of the yearly values for the years from 1991 to 2015 [347 to 401 ppm CO2]). In both scenarios, climate variables were maintained at their 1970 exposure levels, covering the same historical years (e.g., for a 25-year-old stand, 1946 to 1970 were the years of interest), while using seasonal, not annual values and calculating average values, not lifetime summations.

Forest dynamics in the Western US differ from those in the East (e.g., generally drier conditions; greater incidence of large wildfires) and as most of the observations for this study are of forest groups located in the 33 states that the USFS labels as comprising the Eastern US, robustness tests were conducted to assess whether results would differ were only eastern observations utilized. Three forest groups [(1) Lobolly/Shortleaf pine, (2) Oak/Gum/Cypress, and (3) Slash/Longleaf pine] have no observations in the Western US. A fourth, White/Red/Jack Pine, has a slight presence in a few Western states, but no western observations were selected in the original matching process (Supplementary Data 2). For the other six forest groups, all observations from Western US states were dropped. As can be seen from Fig. 2, this had the biggest impact on Aspen/Birch and Elm/Ash/Cottonwood. With this data removed, the GM matching algorithm was again used. Balance statistics are presented in Supplementary Data 23 and again show a strong balance for all covariates across all forest groups. With matches made, the average treatment effect on the treated was estimated using the Model (1) specification used to create Table 1. Regression results are presented in Supplementary Data 24, 25, and a revised version of Table 1 for just the observations from the Eastern US is presented as Supplementary Table 7.

As an additional robustness check on the results in Table 1, we tested an alternative functional form of the volume function. This
alternative volume function is shown in Eq. 3. It has a similar shape as
the function used for the main results in the paper, however, this
equation cannot be linearized with logs in a similar way. Thus, it was
estimated with nonlinear least squares, using the matched samples of
naturally regenerated forests for individual forest groups, as well as the
aggregated sample.

\[
\text{Volume} = \frac{a}{(b + \exp(-c \cdot \text{Age}))}
\]  

(3)

We began by estimating two separate growth functions, one for
the pre-1990 (low CO₂) period and one for the post-2000 (high CO₂)
period using Eq. 3. That is, observations from the pre-1990 (low CO₂)
control period and from the post-2000 (high CO₂) treatment period
were handled in separate regressions. For this initial analysis with
the nonlinear volume function, we did not control for CO₂ concentra-
tion or other factors that could influence volume across sites (e.g., weather,
soils, slope, aspect), and thus, results likely show the cumulative
impact of these various factors. Using the regression results (Supple-
mentary Data 26), we calculated the predicted volume for the pre-1990
and post-2000 periods and compared the predicted volumes (Sup-
plementary Data 26), we calculated the predicted volume for the pre-1990
and post-2000 periods and compared the predicted volumes (Sup-
plementary Table 8).

Next, we tested this yield function on the combined sample
(assuming both control and treatment observations) and all forest
groups. Here the model was expanded to better identify the impact of
elevated CO₂ by including all covariates. Instead of using a dummy
variable for each forest group, though, a single dummy variable was
used to differentiate hardwoods from softwoods. Once again, the
equation was logarithmically transformed for ease of comparison with
the results presented in Table 1. All covariates were originally input, but
those which were not significant were removed. That process yielded
the functional form shown in Eq. 4. Results for the regression are
presented in Supplementary Data 27. The predicted change in volume
due to CO₂ fertilization from 1970 to 2015 is shown in Supplementary
Table 9.

\[
\text{Volume} = \left[ (0+\text{a \cdot \text{Time Dummy} + \text{a2 \cdot \text{Ln(Lifetime CO2)} + \text{a3 \cdot \text{Ln(Seasonal Temperature)}}} \right. \\
\left. + \text{a4 \cdot \text{Ln(Seasonal Precipitation)}} + \text{a5 \cdot \text{Site Class}}} \\
\left. + \text{a6 \cdot \text{Physiographic Dummy + a7 \cdot \text{Aspect Dummy + a8 \cdot \text{Stacking Code}}} \\
\left. + \text{a9 \cdot \text{Disturbances + a10 \cdot \text{Hardwood/Softwood Dummy)}} + \text{b0 \cdot \text{b1 \cdot \text{Time Dummy}}} \\
\left. + \text{b2 \cdot \text{Ln(Lifetime CO2)} + \text{b3 \cdot \text{Ln(Seasonal Temperature)}}} \\
\left. + \text{b4 \cdot \text{Ln(Seasonal Precipitation)}} + \text{b5 \cdot \text{Site Class}}} \\
\left. + \text{b6 \cdot \text{Physiographic Dummy + b7 \cdot \text{Aspect Dummy + b8 \cdot \text{Stacking Code}}} \\
\left. + \text{b9 \cdot \text{Disturbances + b10 \cdot \text{Hardwood/Softwood Dummy}}} + \text{exp}(-\text{c0 + c1 \cdot \text{Time Dummy + c2 \cdot \text{Lifetime CO2}}} \\
\left. + \text{c3 \cdot \text{Ln(Seasonal Precipitation)}} + \text{c4 \cdot \text{Ln(Seasonal Temperature)}} + \text{c5 \cdot \text{Site Class}}} \\
\left. + \text{c6 \cdot \text{Physiographic Dummy + c7 \cdot \text{Aspect Dummy + c8 \cdot \text{Stacking Code}}} \\
\left. + \text{c9 \cdot \text{Disturbances + c10 \cdot \text{Hardwood/Softwood Dummy}} + \text{Ag)}} \\
\right)
\]  

(4)

As the results using the nonlinear volume functions were similar
in sign and magnitude to the multivariate-regression results and as the
practice of matching and then running a multivariate-regression
represents a doubly robust econometric approach that has been
shown to yield results that are robust to misspecification in either the
matching or the regression model[28-32,35-37], the main text results are based
on estimations utilizing multivariate-regression analysis post-
matching.

To develop Table 2, which compares naturally regenerated stands
with planted stands, we used the same general approach as was used to
create Table 1. The analysis and comparison of planted and naturally
regenerated stands was conducted only for stands with enough
observations of both to make a comparison: White/Red/Jack, Slash/
Longleaf, and Loblolly/Shortleaf pine. We followed the same matching
and regression procedures as above, but conducted the matching
separately for naturally regenerated and planted stands. We also
limited the data to stands less than or equal to 50 years of age, as there
are few planted stands of older ages due to the economics of rotational forestry[38-40]. Balance statistics for the matched samples are presented
in Supplementary Data 28–30. Again, the matching process resulted in
a good balance in observable plot characteristics, which implies that
we achieved comparable treatment and control plots.

Using the matched data, we estimated the same regression as in
Eq. 2. Estimation results, which use the Model (2) specification from
Supplementary Data 19–21 that was used with the data for these three
forest groups from ages 1–100, are presented in Supplementary
Data 30–32. A comparison of the parameter estimates on the natural
log of lifetime CO₂ exposure between the results for ages 1–50 (from
Supplementary Tables 31–33) and those for ages 1–100 (from Sup-
plementary Data 19–21) is presented in Supplementary Table 10.

Reporting summary
Further information on research design is available in the Nature
Research Reporting Summary linked to this article.

Data availability
The compiled data generated from the raw data in the USFS-FIA and
PRISM group databases have been deposited in the Ohio State Uni-
versity repository, which is a public database that does not require
special permission to access. The data can be accessed using: Click
Link [https://u.osu.edu/forest/co2-fertilization/]. Source data are pro-
vided with this paper.

Code availability
The code files have also been deposited in the Ohio State University
repository, which is a public database that does not require special
permission to access. The code can be accessed using: Click
Link [https://www.dropbox.com/sf/cbi8tdsp4nutw6u/AABF_J-YFyfrkKt2-
vqM3za/FIA_CO2FertilizationData?dl=0&subfolder_nav_tracking=1].

References
1. Pan, Y. et al. A large and persistent carbon sink in the world’s for-
ests. Science 333, 988–993 (2011).
2. Harris, N. L. et al. Global maps of twenty-first century forest carbon
fluxes. Nat. Clim. Change 11, 234–240 (2021).
3. U.S. EPA. Inventory of U.S. Greenhouse Gas Emissions and Sinks.
EPA, 430-R-19-001 (2019).
4. Oswalt, S. N. & Smith, W. B. U.S. forest resource facts and historical
trends. FS-1035. USFS (2012).
5. Smith, W. B. Forest Resources of the United States, 1997. (North
Central Research Station, Forest Service, US Department of Agri-
culture, 2001).
6. Oswalt, S. N., Smith, W. B., Miles, P. D. & Pugh, S. A. Forest
Resources of the United States, 2017: A Technical Document Sup-
porting the Forest Service 2020 RPA Assessment. Gen. Tech. Rep.
WO-97. (US Department of Agriculture, Forest Service, 2019).
7. Pugh, T. A. et al. Role of forest regrowth in global carbon sink
dynamics. Proc. Natl Acad. Sci. USA 116, 4382–4387 (2019).
8. Schimel, D., Stephens, B. B. & Fisher, J. B. Effect of increasing CO₂
on the terrestrial carbon cycle. Proc. Natl Acad. Sci. USA 112,
436–441 (2015).
9. FAO. Global Forest Resources Assessment 2020: Main Report
(FAO, 2020).
10. Forrell, N. et al. Assessing the INDCs’ land use, land use change,
and forest emission projections. Carb Bal. Mgmt. 11, 1–17 (2016).
11. Grassi, G. et al. The key role of forests in meeting climate targets
requires science for credible mitigation. Nat. Clim. Change 7, 220–226
(2017).
12. Roe, S. et al. Contribution of the land sector to a 1.5 C world. Nat.
Clim. Change 9, 817–828 (2019).
13. Masson-Delmotte, V. et al. Global warming of 1.5°C. IPCC Spec. Rep. Impacts Glob. Warming. 1, 1–9 (2018).
14. Norby, R. J. et al. Forest response to elevated CO2 is conserved across a broad range of productivity. Proc. Natl Acad. Sci. USA 102, 18052–18056 (2005).
15. Norby, R. J. & Zak, D. R. Ecological lessons from free-air CO2 enrichment (FACE) experiments. Ann. Rev. Ecol. Evol. Syst. 42, 181–203 (2011).
16. Gedalof, Z. & Berg, A. A. Tree ring evidence for limited direct CO2 fertilization of forests over the 20th century. Glob. Biogeochem. Cycles. 24 (2010).
17. Silva, L. C. & Anand, M. Probing for the influence of atmospheric CO2 and climate change on forest ecosystems across biomes. Glob. Ecol. Biogeogr. 22, 83–92 (2013).
18. Socha, J., Solberg, S., Tymina-Czabańska, L., Tornpalski, P. & Vallet, P. Height growth rate of Scots pine in Central Europe increased by 29% between 1900 and 2000 due to changes in site productivity. For. Ecol. Mgmt. 490, 119102 (2021).
19. Cabon, A. et al. Cross-biome synthesis of source vs. sink limits to tree growth. Science 376, 758–761 (2022).
20. Song, J. et al. A meta-analysis of 1,119 manipulative experiments on terrestrial carbon-cycling responses to global change. Nat. Ecol. Evol. 3, 1309–1320 (2019).
21. Butsic, V., Lewis, D.J., Radeloff, V.C., Baumann, M. & Kuemerle, T. Quasi-experimental methods enable stronger inferences from observational data in ecology. Basic Appl. Ecol. 19, 1–10 (2017).
22. Bursell, E. S. et al. The Forest Inventory and Analysis Database Description and User Manual Version 8.0 for Phase 2. (USFS, 2018).
23. Ferraro, P. J. & Miranda, J. J. Panel data designs and estimators as substitutes for randomized controlled trials in the evaluation of public programs. J. Assoc. Environ. Econ. 4, 281–317 (2017).
24. Sekhon, J. S. Multivariate and propensity score matching software with automated balance optimization: the matching package for R. J.Stat. Softw. 42 (2011).
25. Bartos, D. L. & Campbell, R. B. Decline of quaking aspen in the Interior West—examples from Utah. Rangelands 20, 17–24 (1998).
26. Bartos, D. L. Fading Gold: The Decline of Aspen in the West. (USFS, 2001a).
27. Woolley, S. C., Walker, S., Vernon, J. & Lindroth, R. L. Aspen decline, aspen chemistry, and elk herbivory: are they linked? Rangelands 30, 17–21 (2008).
28. Moser, W. K. et al. Paper Birch (Wiigwaas) of the Lake States, 1980-2010. Gen. Tech. Rep. NRS-149. 1–37 (USFS, 2015).
29. Herms, D. S. & McCullough, D. G. Emerald ash borer invasion of North America: history, biology, ecology, impacts, and management. Ann. Rev. Entomol. 59, 13–30 (2014).
30. Flower, C. E., Knight, K. S. & Gonzalez-Meler, M. S. Impacts of the emerald ash borer (Agrilus planipennis Fairmaire) induced ash (Fraxinus spp.) mortality on forest carbon cycling and successional dynamics in the eastern United States. Biol. Invasions 15, 931–944 (2013).
31. Gunn, J. S., Ducey, M. J., Buchholz, T. & Belair, E. P. Forest carbon resilience of eastern spruce budworm (Choristoneura fumiferana) salvage harvesting in the Northeastern United States. Front. For. Glob. Change 3, 14 (2020).
32. Keeling, C. D. et al. Atmospheric carbon dioxide variations at Mauna Loa observatory, Hawaii. Tellus B 28, 538–551 (1976).
33. Thoning, K. W., Tans, P. P. & Komhyr, W. D. Atmospheric carbon dioxide at Mauna Loa Observatory: 2. J. Geophys. Res. Atmos. 94, 8549–8565 (1989).
34. NCEI. NOAA National Centers for Environmental Information. (2020).
35. Noormets, A. et al. The role of harvest residue in rotation cycle carbon balance in loblolly pine plantations. Glob. Change Biol. 18, 3186–3201 (2012).
36. Maier, C. S., Albaugh, T. J., Allen, H. L. & Dougherty, P. M. Respiratory carbon use and carbon storage in mid-rotation loblolly pine (Pinus taeda L.) plantations: the effect of site resources on the stand carbon balance. Glob. Change Biol. 10, 1335–1350 (2004).
37. Love-Ouress, K. R., Clark, A. III, Schimleck, L. R., Jokela, E. J. & Daniels, R. F. Specific gravity responses of slash and loblolly pine following mid-rotation fertilization. For. Ecol. Mgmt. 257, 2342–2349 (2009).
38. Seymour, R. S. & Smith, D. M. A new stocking guide formulation applied to eastern white pine. For. Sci. 33, 469–484 (1987).
39. Benzie, J. W. Manager’s Handbook for Red Pine in the North-Central States. General Technical Report NC-33. (USFS, 1977).
40. Brais, S., Paré, D. & Roch, O. Impacts of wild fire severity and salvage harvesting on the nutrient balance of jack pine and black spruce boreal stands. For. Ecol. Mgmt. 137, 231–243 (2000).
41. PRISM Climate Group. Oregon State University. http://prism. oregonstate.edu (2019).
42. Duan, N. Smearing estimate: a nonparametric retransformation method. J. Am. Stat. Assoc. 78, 605–610 (1983).
43. Mären, I. E., Karki, S., Prajapati, C., Yadav, R. K. & Shrestha, B. B. Facing north or south: does slope aspect impact forest stand characteristics and soil properties in a semiarid trans-Himalayan valley? J. Arid Environ. 112, 112–123 (2015).
44. Newingham, B. S. et al. No cumulative effect of 10 years of elevated [CO2] on perennial plant volume components in the Mojave Desert. Glob. Change Biol. 19, 2168–2181 (2013).
45. Diamond, A. & Sekhon, J. S. Genetic matching for estimating causal effects: A general multivariate matching method for achieving balance in observational studies. Rev. Econ. Stat. 95, 932–945 (2013).
46. Rubin, D. B. & Stuart, E. S. Affinely invariant matching methods with discriminant mixtures of proportional ellipsoidal symmetric distributions. Ann. Stats. 34, 1814–1826 (2006).
47. Colson, K. E. et al. Optimizing matching and analysis combinations for estimating causal effects. Sci. Rep. 6, 1–11. (2016).
48. Holmes, W. M. Using Propensity Scores in Quasi-Experimental Designs (Sage Publications, 2013).
49. Rosenbaum, P. R. & Rubin, D. B. Constructing a control group using multivariate matched sampling methods that incorporate the propensity score. Am. Stat. 39, 33–38 (1985).
50. Sekhon, J. & Grieve, R. A new non-parametric matching method for bias adjustment with applications to economic evaluations. (2008).
51. Vose, M. D. Modeling simple genetic algorithms. Found. Genet. Algorithms 2, 63–73 (1993).
52. Nix, A. E. & Vose, M. D. Modeling genetic algorithms with Markov chains. Ann. Math. Artif. Intel. 5, 79–98 (1992).
53. The Forest Inventory and Analysis Database Description and User Manual Version 8.0 for Phase 2. USFS, 2018.
54. Colson, K. E. et al. Optimizing matching and analysis combinations for estimating causal effects. Sci. Rep. 6, 1–11. (2016).
55. Holmes, W. M. Using Propensity Scores in Quasi-Experimental Designs (Sage Publications, 2013).
56. Rosenbaum, P. R. & Rubin, D. B. Constructing a control group using multivariate matched sampling methods that incorporate the propensity score. Am. Stat. 39, 33–38 (1985).
57. Sekhon, J. & Grieve, R. A new non-parametric matching method for bias adjustment with applications to economic evaluations. (2008).
58. Vose, M. D. Modeling simple genetic algorithms. Found. Genet. Algorithms 2, 63–73 (1993).
59. Nix, A. E. & Vose, M. D. Modeling genetic algorithms with Markov chains. Ann. Math. Artif. Intel. 5, 79–98 (1992).
60. R Core Team. R: a language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. https://www.R-project.org/ (2022).
61. StataCorp. Stata Statistical Software: Release 14. (StataCorp LP, 2015).
62. Imbens, G. W. & Wooldridge, J. M. Recent developments in the econometrics of program evaluation. J. Econ. Lit. 47, 5–86 (2009).
63. Stuart, E. S. Matching methods for causal inference: a review and a look forward. Stat. Sci. 25, 1 (2010).
64. Ho, D. E., Imai, K., King, G. & Stuart, E. S. Matching as a nonparametric preprocessing for reducing model dependence in parametric causal inference. Political Anal. 15, 199–236 (2007).

Acknowledgements

Funding for this project was obtained from the USDA National Institute for Food and Agriculture Grant 20176702326275. The findings and conclusions in this publication are those of the author(s) and should not be construed to represent any official USDA or US Government determination or policy. This research was supported [in part] by the US Department of Agriculture, Economic Research Service.
Author contributions
Conceptualization, Methodology, and Writing—review and editing: B.S., E.C.D., and D.J.L. Data curation: E.C.D. Investigation: E.C.D. Visualization: E.C.D. and B.S. Funding acquisition: B.S. and D.J.L. Project administration: B.S. Software: E.C.D. Supervision: B.S. Writing—original draft: E.C.D. and B.S.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary information The online version contains supplementary material available at https://doi.org/10.1038/s41467-022-33196-x.

Correspondence and requests for materials should be addressed to Eric C. Davis.

Peer review information Nature Communications thanks Weisheng Zeng and the other, anonymous, reviewer(s) for their contribution to the peer review of this work. Peer reviewer reports are available.

Reprints and permission information is available at http://www.nature.com/reprints

Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/.

This is a U.S. Government work and not under copyright protection in the US; foreign copyright protection may apply 2022