Carbon storage, timber production, and biodiversity: comparing ecosystem services with multi-criteria decision analysis

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Abstract. Increasingly, land managers seek ways to manage forests for multiple ecosystem services and functions, yet considerable challenges exist in comparing disparate services and balancing trade-offs among them. We applied multi-criteria decision analysis (MCDA) and forest simulation models to simultaneously consider three objectives: (1) storing carbon, (2) producing timber and wood products, and (3) sustaining biodiversity. We used the Forest Vegetation Simulator (FVS) applied to 42 northern hardwood sites to simulate forest development over 100 years and to estimate carbon storage and timber production. We estimated biodiversity implications with occupancy models for 51 terrestrial bird species that were linked to FVS outputs. We simulated four alternative management prescriptions that spanned a range of harvesting intensities and forest structure retention. We found that silvicultural approaches emphasizing less frequent harvesting and greater structural retention could be expected to achieve the greatest net carbon storage but also produce less timber. More intensive prescriptions would enhance biodiversity because positive responses of early successional species exceeded negative responses of late successional species within the heavily forested study area. The combinations of weights assigned to objectives had a large influence on which prescriptions were scored as optimal. Overall, we found that a diversity of silvicultural approaches is likely to be preferable to any single approach, emphasizing the need for landscape-scale management to provide a full range of ecosystem goods and services. Our analytical framework that combined MCDA with forest simulation modeling was a powerful tool in understanding trade-offs among management objectives and how they can be simultaneously accommodated.

Key words: biodiversity; carbon sequestration and storage; forest ecosystem modeling; forest planning; multi-criteria analysis; multi-objective forest management; northern hardwood forests (USA); timber production.

INTRODUCTION

Growing concerns about global climate change and loss of biodiversity have intensified efforts to manage forests for multiple ecosystem services and functions. One critical function of forests is that they serve as large reservoirs of carbon within the global carbon cycle (Houghton 2007). How forests are managed can significantly affect whether they act as net carbon sources or sinks (Birdsey et al. 2006), and therefore forest management practices can play an important role in lessening the intensity of global climate change induced by rising atmospheric CO2 concentrations. Additionally, forests provide a diversity of important ecosystem goods and services, including timber. Not only do revenues from wood products provide substantial direct economic benefits, they can act as powerful incentives to sustain forests, habitats, and undeveloped space over the long term (Fischer et al. 2006, Ruddell et al. 2007). Moreover, forests support a large proportion of the world’s terrestrial species (Lindenmayer and Franklin 2002). Forest biodiversity provides humans with important food sources, recreational and aesthetic opportunities, and other benefits (Ehrlich and Ehrlich 1992, Daily et al. 1997).

Forest management practices vary greatly in their effects on carbon storage and temporal dynamics (Nunery and Keeton 2010), timber production, and provision of habitats for biodiversity (Lindenmayer and Franklin 2002). Challenges arise in managing these multiple ecosystem services because of trade-offs among the mix of services provided by a given management practice. A practice that maximizes one objective, such as timber production, is unlikely to simultaneously maximize all other ecosystem services. An additional challenge is quantifying the value of both market and
nonmarket goods and services provided by forests (Farley 2008), which may be expressed in disparate units that are not readily compared.

One approach for evaluating forest management alternatives is multi-criteria decision analysis (MCDA; also known as multiple criteria decision support, among other names). MCDA is a collection of formal methods used to select a justifiable course of action that explicitly accounts for multiple, conflicting objectives (Belton and Stewart 2002, Mendoza and Martins 2006). One procedure for MCDA is to begin by specifying management objectives and to weight the objectives based on value judgments (Fig. 1). For instance, in the simple case of deciding which management option to implement at a single site, a forest manager may value carbon storage, timber income, and biodiversity equally (with resulting weights of 0.333 for each objective), or may value one objective at twice the level of the others (with resulting weights of 0.50, 0.25, and 0.25). Given a set of standardized metrics and weights for each objective, a management prescription is assigned to the site and the expected effects of the chosen prescription are quantified. The quantities for the objectives are known as the “partial utilities” of a management decision and are then summed to yield the total utility of a management decision (Fig. 1). By repeating this process for other potential management prescriptions, the prescription that maximizes total utility can be identified and applied.

MCDA also allows a structured decision analysis for cases where different forest management prescriptions can be applied to different stands or compartments to maximize the total utility across compartments. In this case, a “portfolio” defines the management prescription assigned to each compartment and the total utility is the sum of utilities across compartments. For instance, if 10 compartments are available for management, the total utility may be maximized by assigning prescription A to compartments 1, 2, 4, 5, 8, and 9 and prescription B to compartments 3, 6, 7, and 10. In recent years, MCDA has been successfully applied to complex forest management scenarios (Kangas and Kangas 2005, Diaz-Balteiro and Romero 2008), although analyses to date have not combined the objectives of carbon storage and timber harvest with evaluation of an extensive set of species to represent biodiversity.

In this paper we apply MCDA to support an overall goal of managing forests for multiple ecological services and functions, as represented by three management objectives: (1) storing carbon, (2) producing wood products, and (3) sustaining biodiversity using a set of wildlife species (Fig. 1). We tested the analysis using a representative forest ecosystem in the northeastern United States for a set of specific metrics over a 100-year time period based on the output of forest development simulation modeling. We considered four alternative forest management prescriptions that spanned a range of harvesting intensities and frequencies. Questions we investigated included the following: (1) To what extent can all three objectives be simultaneously accommodated, and what trade-offs are necessary? (2) How do different management prescriptions compare in their capacity to support individual objectives and the overall goal of multifunctional forest ecosystems? (3) What are the implications for management of forests across multiple sites or large areas?

METHODS

Study area, sites, and field sampling

Our study investigated northern hardwood–conifer forests in the state of Vermont, USA, which is predominantly forested and in which Acer saccharum (sugar maple), Fagus grandifolia (American beech), Tsuga canadensis (eastern hemlock), and Betula alleghaniensis (yellow birch) form the major late-successional species. Within the study area, we measured breeding bird biodiversity and forest characteristics at 533 randomly located sites that were predominantly forested (including early successional forests); see Mitchell and Donovan (2008) for an overview of sampling procedures. Elevation of sites ranged from 29 to 877 m (mean 424 m) and data were collected in 2003 or 2004. At each site, the species identity, diameter at breast height (dbh, 1.37 m height), and live/dead status were recorded for trees at least 2.54 cm in diameter selected in a variable radius plot with a metric 2.3-factor prism. Bird observations consisted of three single-observer 10-minute point counts, separated by 2-minute intervals, conducted at each site during the breeding season. All bird species observed within 75 m were recorded.

Our analysis involved simulating forest development responses to a range of possible management activities. From the 533 sites, which included a variety of forest types and successional stages, we chose a subset of 42 sites (Appendix A; see Plate 1) representative of mature northern hardwood composition (basal area ≥ 25 m²/ha, quadratic mean diameter ≥ 20 cm, and elevation < 600 m). These sites contained sufficient stocking to implement forest management at the commencement of forest simulation, which maximized the effect of management.

Forest simulation model

We used the Forest Vegetation Simulator (FVS) to simulate forest changes at each site over 100-year model runs in 5-year time steps and to estimate carbon (C) storage and timber production at each time step. FVS is a distance-independent, individual tree-based model of forest growth designed for even- and uneven-aged stands with simple- to mixed-species composition (Crookston and Dixon 2005). Although many forest vegetation simulation models are available, each with strengths and weaknesses (Wolfslehner and Seidl 2010), we chose FVS for its ability to simulate forest management activities and the availability of a model variant calibrated for northern hardwoods (Ray et al. 2009a). We used the Northeast Variant (NE-FVS), which uses growth and
yield equations from NE-TWIGS (Hilt and Teck 1989) and embedded height equations and bark ratios specific to northeastern species. We also used regeneration data specific to the study area (Nunery and Keeton 2010). Regional validation studies of NE-FVS have shown adequate predictions of forest growth in northern hardwood forests, with modeled volume predictions within 10–15% of actual volumes (Yauussy 2000). While the absolute numerical predictions generated by FVS carry uncertainty, the model as used here is appropriate for comparing relative differences among management prescriptions (Ray et al. 2009a, Nunery and Keeton 2010). Site-specific individual tree data (species, dbh, live or dead) and environmental characteristics (elevation, slope, aspect) for the 42 sites served as inputs into FVS.

We implemented four alternative silvicultural prescriptions in FVS to determine their relative influence on C storage, timber production, and forest stand structural characteristics. In situ C storage was calculated based on aboveground biomass (live and dead) estimated using species group-specific allometric equations from Jenkins et al. (2003). For simulations involving timber harvesting, FVS also tracked C fluxes to and among wood products pools from production to disposal using life cycle data from Smith et al. (2006). Carbon pools resulting from harvest, each with their own rates of release of C to the atmosphere or flux to another C pool, included coarse woody debris (remaining on site), wood products burned for energy, wood products in use, and landfilled wood and paper. FVS does not currently estimate soil C stocks because they are highly variable and do not respond to management actions in a uniform manner (Hoover and Rehain 2011). FVS does track coarse root biomass as a ratio relative to aboveground biomass, but because other pools including soil C and fine roots are not modeled we did not include
Individual tree selection (ITS) low (30-year entry cycle) harvest individual trees with target basal area of live trees was field measurements described previously with NED-2 for the landscape context of the simulation sites. The covariates improved occupancy estimates by accounting occupancy (Schwenk and Donovan 2011). The latter we previously demonstrated to be important to bird birds (e.g., Hobson and Schieck 1999, MacFaden and structure metrics expected to influence occurrence of predictive covariates were used (Table 2): (1) forest and ages and therefore is considerably more informative than an assessment of a single species or proxy of biodiversity. To predict how forest management practices would affect bird populations, we modeled the relationship between FVS forest structure outputs and occupancy ($\psi$), the probability that a site was occupied by a species. The species set consisted of all territorial, terrestrial birds with relatively small home ranges and sufficient detections (>10 sites) for modeling (common and scientific names of the 51 species are provided in Appendix B). We implemented single-season occupancy models (MacKenzie et al. 2002) in MARK 5.1 (White and Burnham 1999) using data from all 533 original sites, which allowed assessment of bird responses to a range of vegetation conditions. For all species, two types of predictive covariates were used (Table 2): (1) forest structure metrics expected to influence occurrence of birds (e.g., Hobson and Schieck 1999, MacFaden and Capen 2002), and (2) patch and landscape metrics that we previously demonstrated to be important to bird occupancy (Schwenk and Donovan 2011). The latter covariates improved occupancy estimates by accounting for the landscape context of the simulation sites. The four forest structure covariates were calculated from field measurements described previously with NED-2 software (Twery et al. 2005). Basal area of live trees was selected as a metric of how forest stand structure affects bird occurrence. Basal area of live conifers was selected based on numerous studies documenting the importance of conifer trees (or their absence) to habitat selection. Basal area of standing dead trees primarily indicated snag availability, recognizing the importance of dead trees to cavity-nesting, insectivorous, and other birds, but also indicated structural development more generally. Quadratic mean diameter (QMD) of live trees, which is the diameter of the tree of average basal area, was selected to integrate information about both mean diameter and tree size distribution. The patch and landscape covariates (Table 2) reflected landscape condition at multiple scales. A categorical variable indicated whether the site was mid- to late-successional forest (1) or in early-successional stages (0). Distance to edge indicated distance to the nearest major land cover type, such as agricultural land. Percentage forest and road density within 1 km reflected major human influences on the surrounding landscape including conversion of forests and development. Using these occupancy covariates, we generated 16 models for each species. The model set consisted of all possible combinations of the four forest structure covariates (including a model with no covariates). All models included the patch and landscape occupancy covariates. The equation for the model containing all occupancy covariates was as follows:

$$
\logit(\psi) = \beta_0 + \beta_1(\text{basal area of live trees}) + \beta_2(\text{basal area of live trees})^2 + \beta_3(\text{basal area of conifers}) + \beta_4(\text{basal area of conifers})^2 + \beta_5(\text{basal area of dead trees}) + \beta_6(\text{QMD}) + \beta_7(\text{QMD})^2 + \beta_8(\text{mature forest})^2 + \beta_9(\text{distance to edge}) + \beta_{10}(\text{percentage forest within 1 km}) + \beta_{11}(\text{percentage forest within 1 km})^2 + \beta_{12}(\text{road density})
$$

The quadratic terms, which were always paired with their corresponding unsquared terms, allowed assess-

<table>
<thead>
<tr>
<th>Prescription</th>
<th>Harvesting frequency</th>
<th>Structural retention</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Clearcut</td>
<td>high (80 years)</td>
<td>none</td>
<td>clearcut: 2004, 2084; remove slash from site; commercial thin when stand fully stocked</td>
</tr>
<tr>
<td>Shelterwood</td>
<td>low (&gt;100 years)</td>
<td>medium</td>
<td>residual basal area 14 m²/ha; removal cut in 2024 of trees ≥15 cm diameter, retain 6 legacy trees/ha; leave slash on site; commercial thin when stand fully stocked</td>
</tr>
<tr>
<td>Individual tree selection (ITS)</td>
<td>low (30-year entry cycle)</td>
<td>high</td>
<td>harvest individual trees with target q-factor of 1.3; residual basal area 19 m²/ha; retain 12 legacy trees/ha (average diameter 41 cm)</td>
</tr>
<tr>
<td>No management</td>
<td>none</td>
<td>high</td>
<td>no active management simulated</td>
</tr>
</tbody>
</table>
ment of a curvilinear relationship between the covariates and occupancy. We included quadratic terms because we expected that for some species occupancy might be greatest at intermediate values of basal area, QMD, or percentage forest in the landscape. We evaluated the importance and effect of covariates on occupancy in a multimodel inference framework with Akaike’s Information Criterion (AIC; Burnham and Anderson 2002), which uses information from all models in the set to draw statistical inferences. For each species, relative importance of a covariate was calculated as the sum of AIC weights for the models containing the covariate. We considered a covariate to receive substantial support if its relative importance was $>0.5$; the sum of weights for all models $= 1$.

**Linking bird occupancy models to FVS simulations**

Combining the FVS simulation outputs with the beta coefficient values from the occupancy models allowed us to predict occupancy for all bird species at each simulation time step (see also Table 2). Values of basal area and QMD were obtained from FVS for each site at each 5-year interval. Covariate values for percentage forest within 1 km and road density were assumed to remain constant at each site throughout the simulation. Clear-cut and shelterwood harvesting (but not ITS) were assumed to affect covariate values for mature forest and distance to edge as follows. Harvesting changed sites from mid-/late-successional to early-successional forest for 20 years after harvest; i.e., the categorical variable for mature forest was “no” during the first four time steps after harvest (4, 9, 14, and 19 years), and “yes” during all other time steps. This time period was based on studies showing that some early-successional bird species utilize harvested areas at least 15–20 years after harvesting (Keller et al. 2003, Schlossberg and King 2009). During time steps when mature forest was no, distance to edge was reduced to 20 m to be representative of bird occupancy in small early-successional patches within a larger forest matrix. We chose this distance because harvesting in the study region generally occurs on a small scale and because the range of distances to edge for early successional sites recorded in our data set was small (0–50 m, mean 18.6 m). At all other time steps the actual value of distance to edge obtained for each site was used. We calculated a weighted average occupancy for each site/prescription/time step/species combination using the top-ranked models representing $>95\%$ AIC weight for a given species.

As an aid in presenting and interpreting results, we grouped species into clusters with similar patterns of occupancy, and therefore similar associations with predictor covariates, based on the full set of 533 study sites. For this analysis, we used Proc Distance and Proc Cluster (SAS Institute 2008) to generate a dendrogram of occupancy relationships based on hierarchical agglomerative clustering on the species set (predicted occupancy of each species at each site). We then examined the dendrogram to select a small number of species clusters.

### Table 2. Covariates used in modeling occupancy for 51 species of landbirds in Vermont (field data collected in 2003 and 2004).

<table>
<thead>
<tr>
<th>Covariates</th>
<th>Data source</th>
<th>Accompanied by quadratic term?</th>
<th>Data set range</th>
<th>Data set median</th>
<th>Change at sites during forest simulations?</th>
</tr>
</thead>
<tbody>
<tr>
<td>Forest structure variables tested in model set</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Basal area of live trees, $\geq 2.5$ cm dbh (m$^2$/ha)</td>
<td>field measurement</td>
<td>yes</td>
<td>0.0–59.7</td>
<td>25.3</td>
<td>yes</td>
</tr>
<tr>
<td>Basal area of live conifer trees, $\geq 2.5$ cm dbh (m$^2$/ha)</td>
<td>field measurement</td>
<td>yes</td>
<td>0.0–50.7</td>
<td>2.2</td>
<td>yes</td>
</tr>
<tr>
<td>Basal area of standing dead trees, $\geq 2.5$ cm dbh (m$^2$/ha)</td>
<td>field measurement</td>
<td>no</td>
<td>0.0–18.4</td>
<td>2.3</td>
<td>yes</td>
</tr>
<tr>
<td>Quadratic mean diameter (QMD) of live trees, $\geq 2.5$ cm dbh (cm)</td>
<td>field measurement</td>
<td>yes</td>
<td>0.0–67.0</td>
<td>19.2</td>
<td>yes</td>
</tr>
<tr>
<td>Mature forest (or early successional) within 25 m</td>
<td>field assessment</td>
<td>no</td>
<td>†</td>
<td>N/A</td>
<td>yes: forest changed to early successional for 19 yr following harvest</td>
</tr>
<tr>
<td>Patch and landscape variables appearing in all models:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Distance to edge of nearest other land cover type (m)</td>
<td>NLCD (2001)$§$ supplemented by aerial and field photographs</td>
<td>no</td>
<td>0–2030</td>
<td>150</td>
<td>yes: edge distance reduced to 20 m when mature forest = “no”</td>
</tr>
<tr>
<td>Percentage forest within 1 km</td>
<td>NLCD 2001</td>
<td>yes</td>
<td>5.7–100</td>
<td>94</td>
<td>no</td>
</tr>
<tr>
<td>Road density within 1 km (km/km$^2$)</td>
<td>Vermont e911 roads data</td>
<td>no</td>
<td>0.0–9.7</td>
<td>0.6</td>
<td>no</td>
</tr>
</tbody>
</table>

**Notes:** Variable radius plot field measurements (tree species and diameter) were used to calculate basal areas and QMD for all 533 sites used to develop occupancy models; future values of these covariates were simulated in FVS for the 42 simulation sites.

† Range of values for 533 sites used to develop occupancy models.

‡ Categorical: yes, no.

§ National Land Cover Database (MRLC 2001).
Multi-criteria decision analysis

We implemented the MCDA in three parts: (1) estimating partial utilities of objectives when a single prescription was assigned to all sites (single-prescription portfolio); (2) estimating the total utility under different weighting scenarios for single-prescription portfolios; and (3) allowing the assigned prescription to vary among sites (multi-prescription portfolios) to identify the combination of prescriptions that maximized total utility. For part 1, we computed partial utilities for carbon (C), timber (T), and biodiversity (B) for each site (i) and time step given a management prescription (j). We scaled the partial utilities between 0% and 100%, such that 0 represented the lowest possible scoring and 100 represented the maximum possible scoring for any given objective. The partial utility for C storage, \( U_{C,i,j} \), was calculated as the mean annual C stored during the 100-year simulation (at site i for prescription j), divided by the mean annual C for the site-prescription combination with the maximum C storage (considering all four prescriptions). Similarly, the partial utility \( U_{T,i,j} \) for timber production was calculated as the mean annual volume of merchantable wood produced during the simulation (at site i for prescription j) divided by the mean annual volume produced for the site and prescription with the maximum timber production.

We calculated the corresponding partial utility for biodiversity for each site and prescription, \( U_{B,i,j} \), in several steps. After estimating average occupancy for each bird species during the simulated time period, we rescaled occupancy estimates so that species with greater estimated occupancy (more common species) did not have a disproportionate influence on the utility function. The rescaled occupancy estimates were summed across the 51 species:

\[
B_{i,j} = \sum_{k=1}^{51} \frac{\psi_{i,j,k}}{\psi_{\text{max},i,j,k}}
\]

where \( \psi_{i,j,k} \) was the mean occupancy for an entire simulation (site i, prescription j, species k) and \( \psi_{\text{max},i,j,k} \) was the maximum of these occupancy values. The final partial utility converted the bird occupancy scores to the 0–100% scale:

\[
U_{B,i,j} = \frac{B_{i,j}}{B_{\text{max},i,j}}
\]

where \( B_{\text{max},i,j} \) was the maximum bird occupancy score among all site–prescription combinations.

In part 2 of the MCDA, we estimated total utility by including weights for management objectives while retaining the assumption that all sites receive the same management prescription. Weights represented preferences or values assigned to the management objectives (\( w_C = \) C weight, \( w_T = \) timber weight, and \( w_B = \) biodiversity weight) under the constraint that the weights summed to 1. We explored a full range of weights to examine how they affected the expected utility for the four prescriptions. For any given scenario, the assigned weights were multiplied by the site-specific partial utilities described in part 1. The total utility of a management prescription \( j \) was the following linear additive model (averaged across the 42 sites):

\[
U_j = \sum_{i=1}^{42} w_C \times U_{C,i,j} + w_T \times U_{T,i,j} + w_B \times U_{B,i,j}
\]

In part 3 of the MCDA, we added the ability to differentially manage sites and identify the portfolio \( p \) that maximized the total utility across sites. The total utility \( (U_p) \) was expressed as in the previous equation, with the addition of a choice vector to identify which prescription \( j \) was assigned to each site. For the four prescriptions \( j_1-4 \), values were binary (0 or 1) and only one prescription could be chosen per site for the duration of the simulation, resulting in the following total utility for the entire portfolio:

\[
U_p = \frac{1}{42} \sum_{j=1}^{42} \left[ w_C U_{C,j-1} + w_T U_{T,j-1} + w_B U_{B,j-1} \right]
\]

Because the partial utilities were scaled between 0% and 100% and weights ranged from 0 to 1, \( U_p \) was also scaled between 0% and 100%.

We considered four weighting scenarios that represented a range of preferences. In one scenario the weights for the objectives were equal; in the other three, one objective was weighted 0.60 and two were weighted 0.20. For a weighting scenario, the portfolio with the maximum utility was the one where each site was assigned the prescription that maximized the utility at that site. We identified the portfolio with maximum utility under each weighting scenario, and tallied the number of prescriptions of each type that had been assigned.

**Results**

**Starting condition of forest characteristics and bird occupancy**

At the beginning of the simulation, live basal area at the 42 sites (Appendix A) ranged from 23.0 to 50.5 m²/ha (mean 29.6) and QMD ranged from 20.8 to 48.0 cm (mean 28.5 cm). Percentage of live basal area composed of conifers ranged from 0% to 54.9%, with a mean of 7.7% or 2.6 m²/ha.

All four forest structure covariates were important to occupancy estimates for multiple species (Table 3), with conifer basal area being important for the most species. A variety of linear and nonlinear associations between covariates and occupancy were apparent. For example, the association between conifer basal area and occupancy was positive for Blackburnian Warbler (Setophaga fusca), negative for Veery (Catharus fusescens), and
Simulations of changes in forest characteristics over time

Values of FVS outputs (basal area, total live aboveground biomass, and QMD) showed considerable fluctuation over time and substantial differences among prescriptions (Fig. 2). Harvesting under all three active management prescriptions resulted in substantial decreases in average live, conifer, and dead basal area compared to no management. On average, live basal area (Fig. 2a) and aboveground live biomass (Fig. 2e) were greatest for the no-management prescription and least for the clear-cut prescription. For the first half of the simulation period, live conifer basal area (Fig. 2b) was greatest for the no-management prescription, but subsequently became greater under the ITS and clear-cut prescriptions as a result of model regeneration inputs. Basal area of dead trees (Fig. 2c) was consistently greatest in the no-management prescription. Temporal patterns for QMD (Fig. 2d) were stable for the no-management prescription due to compensatory effects of in-growth of small trees and increases in dominant canopy tree sizes. Harvesting decreased QMD except in the case of the shelterwood prescription, where the initial cut resulted in an increase in mean tree diameter, followed by a large decrease after the subsequent overstory removal. Clear-cut QMD was much reduced as an average over two rotations compared to no management, as expected for even-aged management. ITS QMD values were intermediate due to the removal of some canopy trees at each harvest coupled with regeneration and stem development across a range of tree sizes.

Simulations of carbon storage and timber production

The forest simulations resulted in a clear pattern of increasing C storage with decreasing intensity of forest management (Fig. 3). Mean C storage for the simulation under the no-management prescription (147.3 Mg C/ha) was more than triple that of the clearcut prescription (46.3 Mg C/ha). More intensive forest management resulted in greater storage of C in wood products, but these quantities did not make up for concomitant large reductions in aboveground live biomass (Figs. 2f and 3). Under the simulations, a considerable portion of harvested biomass did not become part of wood products, including wood that remained on site (where it decomposed), was discarded during processing (bark and limbs), or was burned for energy. The decline in C stored in wood products over time (Fig. 2f) reflected relatively short half-lives (<15 yr) projected for certain products, such as wooden pallets and railroad ties. ITS most closely approached the no-management prescription, with mean C storage (112.6 Mg C/ha) 76.4% of the no-management prescription, of which 12.5% (14.1 Mg C/ha) constituted long-term storage in harvested wood products. No management and lower intensity management had higher recruitment rates for snags, which led to an increase in the downed log C pool over time.

Annual timber production and volume removals were greatest in the clearcut prescription (2.5 ± 0.068 m³/ha, mean and SE for 42 sites), with less harvested in the ITS (2.0 ± 0.084 m³/ha) and shelterwood prescriptions (1.5 ± 0.075 m³/ha). The no-management prescription did not include harvests.

Simulations of changes in bird occupancy over time

Changes in forest covariates (Fig. 2), plus accompanying changes to distance to edge following harvests, resulted in changes in predicted bird occupancy over time within management prescriptions and differences among management prescriptions (Fig. 4). We grouped the species into five clusters with similar patterns of occupancy (cluster assignments for all species are presented in Appendix B). Species in cluster 1 (which we termed "edge and early successional species"), such
as Song Sparrow (*Melospiza melodia*), showed substantial increases in occupancy when sites were in an early successional state (Fig. 4a, b). They also were associated with relatively unforested landscapes. Cluster 2 species (“species of intermediate forest landscapes”), such as Veery, tended to occur most frequently in landscapes with an intermediate percentage of forest within 1 km and were not particularly sensitive to local forest management. Cluster 3 species (“forest interior species”), such as Black-throated Green Warbler (*Setophaga virens*), decreased in occurrence following intensive harvesting and were more likely to occupy sites with greater values of percentage forest within 1 km. Like forest interior species, species of cluster 4 (“coniferous forest”) were associated with highly forested landscapes, but occupancy was greater in forests with higher conifer basal areas. In general, Blackburnian Warbler and other coniferous forest species did not respond as negatively to harvesting as forest interior species. Cluster 5 species tended to occur near edges and in early successional sites within landscapes that otherwise were predominantly forested, so we termed them “early successional/forest
matrix species.” Chestnut-sided Warbler (*Setophaga pensylvanica*) is representative of this cluster.

Considering all species collectively, the most substantial changes over time occurred in the clearcut prescription, reflecting its intensive harvest regime (Fig. 4a). For 20 years following clearcuts, occupancy predictions for early successional species (clusters 1 and 5) increased while they decreased for forest interior species. Increasing basal area of conifers as regeneration proceeded positively affected coniferous forest species until the second clearcut. In the shelterwood prescription (Fig. 4b), early-successional and forest interior species exhibited similar responses to harvesting as in the clearcut prescription, except that fluctuations were more muted and of longer duration, reflecting the less intensive harvest regime implemented over a longer time period. By maintaining greater stand structural complexity as measured by vegetation covariates, the ITS prescription (Fig. 4c) resulted in occupancy patterns similar to the no-management prescription (Fig. 4d). For both of these prescriptions, occupancy of early-successional species was suppressed relative to the more intensive prescriptions and was much more stable over time. Conversely, forest interior and coniferous forest species were associated with more mature forests and collectively showed greater occupancy under the ITS and no-management prescriptions.

**Fig. 3.** Predicted carbon stocks (mean ± SE), including stocks stored in harvested wood products, under different management prescriptions applied to 42 northern hardwood forest sites (average for 100-year simulations).

**Fig. 4.** Average predicted occupancy summed for all 51 forest birds species, grouped in clusters, for the four management prescriptions: (a) clearcut with short harvest interval (clearcuts in 2004 and 2084), (b) shelterwood with long harvest interval (partial harvest in 2004, removal cut in 2024), (c) individual tree selection, 30-year entry cycle, and (d) no management.
Multi-criteria decision analysis

In part 1 of the MCDA, where portfolios consisted of a single prescription assigned to all sites, the partial utilities for an objective were clearly related to management prescription (Fig. 5). The no-management prescription utility was greatest for C storage but least for timber production and biodiversity; the clearcut prescription utility had the reverse pattern. Although the ITS prescription utility was not the greatest for any objective, it was consistently high for all objectives (Fig. 5).

Sites varied more in estimated annual timber production, given the same management prescription, than they did for C storage or biodiversity (Table 4). Timber production was especially variable for the shelterwood prescription, with the most productive site estimated to yield greater than three times the volume of the least productive site. C storage and biodiversity values varied across sites by less than a factor of 2 when all sites received the same prescription.

In the part 2 of the MCDA, we identified the prescription that maximized total utility when all sites received the same prescription and when weighting factors were applied to the three objectives. Which prescription maximized total utility depended on the assigned weights. The no-management and clearcut prescriptions varied strongly depending on the assigned weights, with total relative utilities ranging from 0 (for no management if timber weight = 1.0) and 0.3 (for clearcut if C weight = 1.0) to 1. The ITS prescription frequently was the second-ranked prescription and consistently had a high total relative utility that was 75–80% of the maximum possible utility. The shelterwood prescription was never the highest ranked prescription.

In the case where sites could be managed differentially (MCDA part 3), multi-prescription portfolios achieved higher total utilities than single-prescription portfolios under the combinations of weights we investigated (Fig. 7). The dominant prescription when optimizing allocation was consistent with the pattern observed for single-prescription portfolios (Figs. 6 and 7). Although multi-portfolio prescriptions were favored, in all cases the utility of at least one single-prescription portfolio nearly equaled the maximum utility (Fig. 7). In fact, utility of the all-ITS portfolio was ≥90% of the maximum utility for all weights we tested. The all-clearcut portfolio also attained 90% or more of the maximum utility whenever the weight for C storage was ≤0.33. On the other hand, the all-shelterwood portfolio utility did not exceed 90% of maximum utility for any weighting scenario, and none of the maximum utility portfolios included shelterwood prescriptions. When weights for C storage were large, the utility of the portfolio exclusively consisting of
no management nearly equaled the maximum possible utility.

**DISCUSSION**

Our findings offer new insights for sustainable forest management, demonstrating the utility of analytical approaches that combine forest simulation modeling with MCDA. These approaches have great potential for evaluating trade-offs among multiple management objectives. The framework we present can be readily modified to incorporate (1) alternative models or (2) additional modules to consider factors such as natural disturbance and climate change. Previous researchers have noted the usefulness of MCDA (Kangas and Kangas 2005, Wolfslehner and Seidl 2010), but ours is the first to integrate carbon and timber objectives with biodiversity objectives represented by a large set of bird species. An emerging theme from our study was that variation in the effects of management prescriptions, site characteristics, and weights placed on management objectives meant that no single management approach had the greatest utility in all circumstances. Therefore, a diversity of silvicultural approaches is likely to be preferable and simultaneously accommodating all three objectives requires compromises among them. Providing a full range of ecosystem goods and services requires holistic, landscape-scale management in which a diversity of silvicultural and conservation approaches are applied in tandem (Lindenmayer and Franklin 2002, Keeton 2007).

With the recent expansion of both voluntary and compliance carbon market systems, demand is increasing for forest carbon management approaches that can be integrated with other objectives, including timber, non-timber forest products, wildlife, water, aesthetics, and recreation (Jackson et al. 2005, Ray et al. 2009b). Our analysis framework demonstrated an intuitively understandable approach to integrating these objectives. Our findings are consistent with a number of recent studies showing that either no management or silvicultural approaches emphasizing less frequent harvesting and greater structural retention result in the greatest net carbon storage (Harmon and Marks 2002, Seidl et al. 2007, Swanson 2009, Nunery and Keeton 2010). Our simulations indicated that considerable carbon can be stored in wood products, but because of losses of wood

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**TABLE 4.** Comparison of site variability among 42 sites for C storage, timber production, and bird occurrence (annual averages for 100-year simulation, summarized for cases when all sites received the same prescription).

<table>
<thead>
<tr>
<th>Prescription</th>
<th>C storage (Mg/ha)</th>
<th>Timber production (m³/ha)</th>
<th>Probability of occupancy (sum of 51 species)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
<td>CV†</td>
</tr>
<tr>
<td>Clearcut</td>
<td>46</td>
<td>6</td>
<td>12</td>
</tr>
<tr>
<td>Shelterwood</td>
<td>86</td>
<td>11</td>
<td>12</td>
</tr>
<tr>
<td>ITS</td>
<td>113</td>
<td>9</td>
<td>8</td>
</tr>
<tr>
<td>No management</td>
<td>147</td>
<td>16</td>
<td>11</td>
</tr>
</tbody>
</table>

† Coefficient of variation ([standard deviation/mean] × 100; %). † The no-management prescription did not entail timber harvests.

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**FIG. 6.** Management prescriptions with the highest relative utilities given all possible combinations of values (weights) assigned to the objectives of carbon storage, timber production, and biodiversity. For example, the illustrated point occurs at a carbon storage weight of 0.1, a timber production weight of 0.4, and a biodiversity weight of 0.5. Given this combination of weights, the clearcut management prescription has the highest total utility.
during production processes, the relatively short (e.g., 1–50 years) life cycle of many wood products (Smith et al. 2006), and decay following eventual disposal, the wood products pool (under intensive wood management) is insufficient to compensate for carbon storage in less intensively managed forests. Our findings also indicated that the less timber harvested, the greater the total (in situ + wood products) amount of carbon likely to be stored over the long term as an annual average. But the results also clearly showed that there are intermediate approaches, particularly when silvicultural prescriptions are coordinated across multiple sites, that could be employed to yield a mix of carbon storage and timber production. Most likely applied forest carbon management will involve some combination of more and less intensive silvicultural approaches depending on the economic incentives offered by carbon markets and the overall management objectives of a given landowner (Ryan et al. 2010, Keeton et al. 2011).

We should point out that our conclusions on carbon implications depend upon the approaches and assumptions we used, including timing of harvests and fate of wood products. Two considerations with potential to modify our conclusions are soil carbon and substitution of wood for more fossil-fuel-intensive alternatives (termed “substitution effects”). Soil carbon represents a significant C pool in forests, but responses of soil C to management are highly variable and uncertain, and consequently are not tracked by FVS (Hoover and Rebain 2011). Some studies have found that intensive management can result in large losses of soil C, whereas others have reported much more modest effects; the intensity of soil scarification, the methods of harvesting and site preparation, and soil type are important in determining soil C fluxes (Yanai et al. 2003, Jandl et al. 2007, Nave et al. 2010). If our analyses had assumed that harvesting induced substantial losses of soil C, then the differences in C storage among the management prescriptions we examined would have been even more pronounced. By contrast, consideration of substitution effects might have had the opposite effect. Some studies that have incorporated assumptions about the offset of emissions achieved by replacing more energy-intensive building materials with wood have concluded that intensified harvesting can reduce net emissions (Perez-Garcia et al. 2005, Eriksson et al. 2007). Given the host of factors necessary for a comprehensive life cycle analysis, such as carbon emissions in harvesting and

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**Fig. 7.** Utilities for portfolios given four combinations of weights (a, b, c, d) for the objectives of carbon storage, timber production, and biodiversity. In each panel, the bar to the left of the vertical dashed line represents the portfolio that maximizes total utility (total utility = 100%), allowing for sites to be assigned different management prescriptions (segment heights represent the number of sites assigned to each prescription; ITS, individual tree selection). The four bars on the right of each panel represent portfolios consisting of all sites assigned the same management prescription, with bar heights equal to total utility as a percentage of the maximum total utility.
transporting wood and cradle-to-grave carbon implications of alternatives, we did not attempt to incorporate substitution effects into our analysis. Both soil carbon and substitution effects can be readily incorporated into MCDA, however, if adequate data are available.

Our study is one of the first to link projection of future forest condition to empirically derived models of wildlife occurrence in the context of multifunctional forest management. It therefore establishes a useful framework for future analyses applicable where diversity of species and natural communities are to be considered, as is required for the U.S. National Forest System. Most prior multi-objective studies, including the few studies that have simultaneously considered carbon storage, timber production, and biodiversity (e.g., Seely et al. 2004, Fürstenau et al. 2007, Briceño-Elizondo et al. 2008), have represented biodiversity by a few species or by indirect proxies of biodiversity (such as density of deadwood). By assessing the probability of occupancy for multiple species simultaneously, our framework presents an alternative to analytical approaches heavily dependent on indicator species for assessing biodiversity.

We found that bird species responded most strongly to the more intensive harvesting practices (clearcut, shelterwood prescriptions). Given the heavily forested landscape in which sites were located and the small patch size of harvests, the predicted positive response of species following intensive harvesting tended to be of a greater magnitude than the negative response of species most common under late successional conditions. The net result was that more intensive prescriptions received larger biodiversity utility values. Results would differ for large clearcuts or fragmented landscapes. The patterns of predicted bird occurrence we observed were generally consistent with studies that have examined how birds respond to silvicultural practices in the northeastern United States (Thompson and Capen 1988, Germaine et al. 1997, Hagan et al. 1997, Costello et al. 2000, Keller et al. 2003, Goodale et al. 2009). Dense stands of seedlings and saplings with low basal area and QMD, which are characteristic of initial years following harvests in an even-aged system, provide high quality habitat for birds that forage on the ground and in low foliage (Keller et al. 2003). Examples of such species from our study are Common Yellowthroat (Geothlypis trichas; cluster 1) and White-throated Sparrow (Zonotrichia albicollis; cluster 5). More mature or less intensively managed stands, which have greater basal area and QMD than recently harvested even-aged systems, offer new niches and greater vertical complexity for species that use upper canopy foliage, bark, and cavities, but provide fewer resources for species of low foliage (Keller et al. 2003). Red-eyed Vireo (Vireo olivaceus; cluster 3) and Brown Creeper (Certhia americana; cluster 4) are examples of species from our study characteristic of more mature and complex forests. Species less sensitive to management appeared to use a wide range of forest types or occur more frequently in forests of intermediate

**Plate 1.** Example of one of 42 northern hardwood forest sites used in simulation of forest management and comparison of predicted future ecosystem services. A white measuring tape used in field sampling is visible in the lower central portion of the image. Photo credit: Charley Eiseman.
Such forests may provide services including watershed protection, riparian function, soil retention, genetic resources, and aesthetic experiences that we did not consider (de Groot et al. 2002, Keeton et al. 2007). They may also harbor species of conservation concern not found in simpler or younger forests (Fischer et al. 2006). Such considerations can readily be addressed within the MCDA framework. Second, landscape context matters (Fischer et al. 2006, Kupfer et al. 2006). Our focal study sites were located in a mostly unfragmented forested landscape, which has important implications for the biodiversity results. In such landscapes, our findings that moderate levels of harvesting may have beneficial effects on biodiversity appear reasonable because adverse effects to forest interior species are not expected to be large (Rodewald and Yahner 2001, Thompson et al. 2002). They also are consistent with arguments that early successional species are of high conservation concern in eastern North America and can benefit from regeneration induced by silvicultural practices (Hunter et al. 2001, DeGraaf and Yamasaki 2003). However, our findings are unlikely to translate well to landscapes where forests are already substantially fragmented. In such landscapes, populations of late successional species may be under stress due to lack of habitat and increased risks from edge predators and nest parasites (Donovan et al. 1995, Fischer and Lindenmayer 2007). A final consideration involves uncertainty, which is inherent in multi-criteria analyses involving models of future conditions (Wolfslehner and Seidl 2010). The uncertainty most readily addressed is the current and future preferences of forest management stakeholders, because our analyses presented results involving a spectrum of weights for objectives. More difficult to quantify are uncertainties related to the FVS model and bird occurrence models. Recognizing that we did not undertake a formal model uncertainty analysis, we caution readers not to assume that small differences between portfolios of silvicultural prescriptions are conclusive evidence of superiority of one portfolio or prescription over another.

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LITERATURE CITED

occupancy rates when detection probabilities are less than one. Ecology 83:2248–2255.


Supplemental Material

Appendix A

Characteristics of 42 northern hardwood forest sites used in simulation modeling (Ecological Archives A022-085-A1).

Appendix B

Bird species and model results for species used in occupancy estimation (Ecological Archives A022-085-A2).