Impacts of Timber Harvesting on Soil Organic Matter, Nitrogen, Productivity, and Health of Inland Northwest Forests

M.F. Jurgensen, A.E. Harvey, R.T. Graham, D.S. Page-Dumroese, J.R. Tonn, M.J. Larsen, and T.B. Jain

ABSTRACT. Soil organic components are important factors in the health and productivity of Inland Northwest forests. Timber harvesting and extensive site preparation (piling, windrowing, or scalping) reduces the amount of surface organic material (woody residues and forest floor layers) over large areas. Some wildfires and severe prescribed burns can have similar consequences. Such organic matter reductions can have important implications for soil chemical, biological and physical properties.

A number of studies have linked substantial reduction in mycorrhizae development and tree growth to high levels of soil disturbance, or removal of organic horizons. Timber harvesting also removes a large percentage of coarse woody debris, which has unknown ramifications on soil productivity. Current woody residue guidelines in this region recommend leaving <10 to 125 Mg ha^{-1} on site to replace woody materials lost during harvesting operations. Large amounts of soil nitrogen (>500 kg ha^{-1}) can also be lost from timber harvesting and site preparation, especially when using prescribed fire. The time required to replace this lost nitrogen may range from <10 to >275 yr, and depends on the severity of site treatments, presence or absence of nitrogen-fixing plants, and amounts of atmospheric deposition.

Maintaining adequate amounts of organic matter on some forest sites in the Inland Northwest may temporarily increase the risk of wildfire or favor the activity of certain insects or disease fungi. However, carefully planned prescribed burns and mechanical site preparation can be practiced on most sites with relatively low impacts on soil organic levels, while accomplishing the important forest management objectives of fuel reduction, seedbed preparation, and reducing competing vegetation. Organic matter management will be the most difficult on very dry sites, with their historically low soil organic and nitrogen content, and high fire potential. The maintenance of adequate soil organic matter levels is critical for sustaining forest health and productivity under the variable moisture and temperature conditions of this region. Thus, soil organic components will become more important in the future as ecosystem management systems are developed for western forests. For. Sci. 43(2):234–251.

Additional Key Words: Mounding, mycorrhizae, nitrogen fixation, scalping, site preparation, woody residue.

Timber harvesting and subsequent site preparation alter the cycling of aboveground forest organic materials and their incorporation into the soil. Soil organic matter is important to maintaining site productivity because of its roles in supporting soil nutrient availability, gas exchange, and water supply (Powers et al. 1990, Blake and Ruark 1992, Henderson 1995). Organic matter also is essential to soil microflora and microfauna active in nutrient cycling, soil aggregation, and disease incidence or prevention (Harvey et al. 1987a). In the past, wood removal was not considered detrimental to site productivity because harvesting old, mature stands left large amounts of residue. However, recent trends toward harvesting younger stands, coupled with total-tree utilization, raise concerns about how such

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management will impact soil processes, site productivity, global carbon sequestration, and forest biodiversity (McColl and Powers 1984, Harvey et al. 1989c, Harmon et al. 1990, Heilman 1990, Powers 1991, Johnson 1992).

Most environmental concerns regarding timber harvesting effects have focused on possible soil nutrient losses or changes in nutrient availability (Edmonds et al. 1989). Appreciable harvest-related losses of soil organic matter would be particularly important to nitrogen (N) cycling, since organic matter is the source of nearly all available soil N (McGill and Cole 1981). While the relationship of organic matter to soil nutrient cycling is a major concern, the importance of organic matter as a physical entity, apart from its nutrient content, also warrants consideration (Standish et al. 1988). Organic matter content has a great influence on many soil physical properties, such as water-holding capacity, aeration, drainage, and cation exchange. Changes in these soil properties following harvesting may be as significant as nutrient losses for subsequent forest growth on some sites.

Loss of soil organic matter during harvesting and site preparation greatly affects soil physical and biological properties, and may reduce overall site productivity, stability, and regeneration potential (Powers et al. 1990, Henderson 1995). Timber harvesting/soil organic matter information has come from many studies in different parts of the world. In Australia, losses of surface organic matter following timber harvesting caused site deterioration by lowering cation exchange capacity and reducing soil moisture retention (Farrell et al. 1986). Removing litter from pine stands decreased soil moisture levels and reduced tree growth in Canada, New Zealand, and the southeastern United States (Ginter et al. 1979, Ballard and Will 1981, Weber et al. 1985).

The majority of timber harvesting/site productivity studies in the Pacific Northwest have come from the moist, maritime forests on the west slopes of the Cascade and Sierra Mountains or in the Pacific Coast ranges (Perry et al. 1989). Much less information is available on the drier, continental forests of the Inland Northwest. However, studies in this region have reported substantial losses in site productivity 15 to 25 yr following clearcutting as a result of soil compaction and forest floor/surface soil displacement (Clayton et al. 1987, Bosworth and Studer 1991). Since soil organic matter removal and soil compaction often occur together, a national study has been established to try and separate the effects of these two harvest-related impacts on site productivity (Powers 1991). Two of these sites are located in the Inland Northwest (Idaho).

Recent changes in forest management on public lands to protect the spotted owl and other endangered species has caused significant timber harvest reductions in western Washington, Oregon, and northern California. Consequently, greater emphasis is being placed on the timber-producing forests of the Inland Northwest. Increased harvesting has also been advocated to alleviate some of the forest health problems in eastern Oregon and Washington caused by decades of fire control (Harvey 1994). This paper discusses the impact that this timber harvesting could have on soil organic matter components, and how this could affect soil N contents and fixation, mycorrhizal root development, seedling establishment and growth, forest health, and site productivity in this region.

The Inland Northwest

The Inland Northwest encompasses areas from the eas slopes of the Cascade Mountains in Washington and Oregon to the forests of Idaho and western Montana. It is bounded on the south by the Snake River in Idaho and the Continental Divide in Wyoming and Montana. Similar forest and soil characteristics occur in the northern portion of California east of the Cascades. Regional topography is characterized by massive mountains with deep V- and round-bottomed valleys. Elevations from 300 m to over 3,000 m dominate the forest zone. Precipitation ranges from 355 mm in the rain shadows of the Cascade and Bitterroot Mountains to over 1,525 mm on the east slopes of the Cascades and in northern Idaho. Snowfall accumulations over 6,500 mm are common with below-freezing temperatures occurring in any month at higher elevations.

Forests

Mixed-conifer forest types dominate forests of the Inland Northwest and comprise part of the Columbia Forest and Rocky Mountain ecoregions (Bailey 1980). These forests are highly variable, ranging from coastal to semi-arid conditions (Figure 1). The dry sites are characterized by widely spaced ponderosa pine (*Pinus ponderosa* Dougl. ex Laws.) or Douglas-fir (*Pseudostuga menziesii* [Mirb.] Franco) forests, intermixed with grasses (*Calamagrostis* spp.) and low shr (*Symphorocarpus* spp.) in the understory. These forests a often located in the rain shadow of major mountain ranges, such as the Bitterroots in Montana and the Cascades in Washington and Oregon. Because of many decades of fire suppression, large numbers of ponderosa pine and Douglas-fir seedlings and saplings are presently found on many of these sites. Down logs and coarse woody debris are abundant, but historically much of this material was consumed by frequent fires (Habeck and Mutch 1973, Arno 1980).

Douglas-fir or grand fir (*Abies grandis* [Dougl. ex D. Don] Lindl.) forests occur in more mesic conditions, typically at elevations above dry Douglas-fir and climax ponderosa pine ecotypes. Species composition includes an admixture of these two species with lodgepole pine (*Pinus contorta* Dougl. ex Loud.), and western larch (*Larix occidentalis* Nutt.). In northern Idaho western white pine (*Pinus monticola* Dougl. ex D. Don.) is often a principal associate on moist sites. The understory is characterized by grasses (e.g., *Calamagrostis* spp.), forbs [e.g., *Clintonia uniflora* (Schult.) Kunth.], low shrubs (e.g., *Vaccinium* spp.) and tall shrubs (e.g., *Ceanothus* spp., *Physocarpus* spp., *Alnus* spp., *Acer* spp.). Because these forests are productive, they accumulate large amounts of coarse woody debris and tend to be densely stocked (Brown and See 1981, Graham 1988). Historically, ponderosa pin western larch, and Douglas-fir dominated most stands, es cially on drier sites, while western white pine was importan on moist sites (Habeck and Mutch 1973). The exclusion of

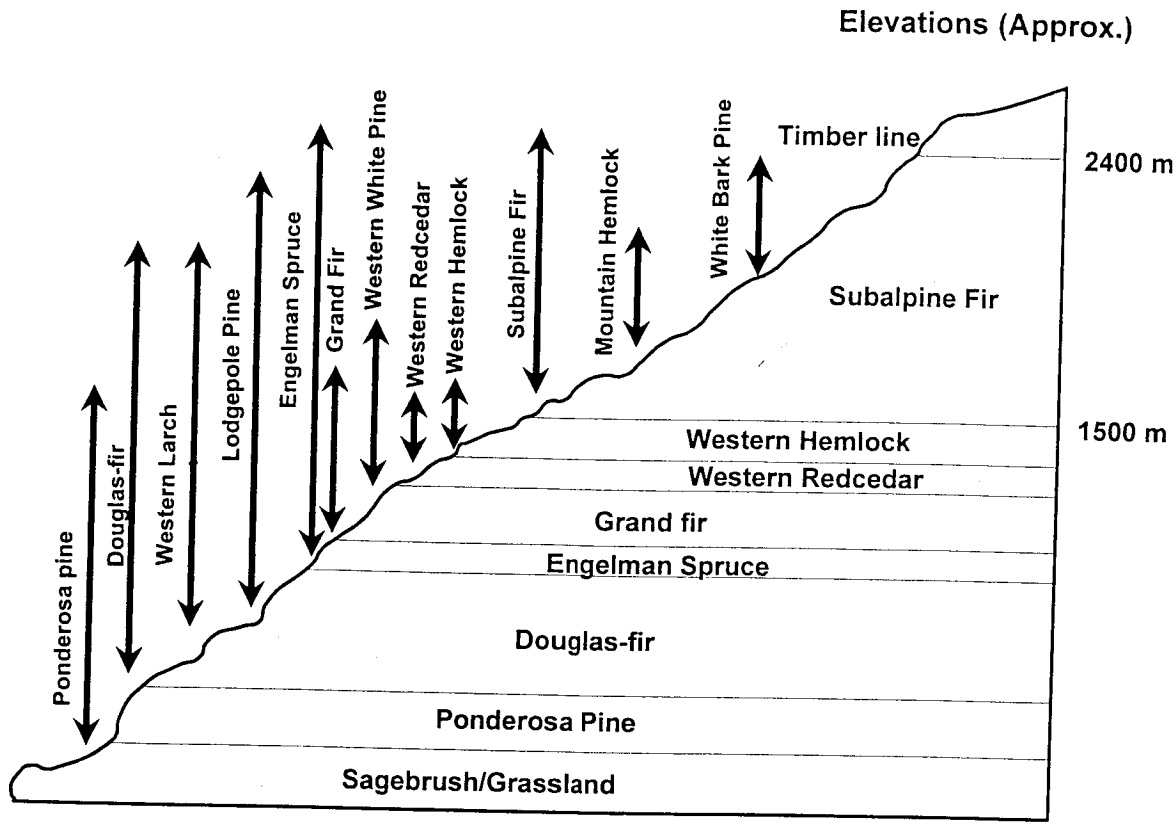

Figure 1. Generalized distribution of forest trees in the Inland Northwest. Horizontal bars indicate the major forest habitat type series found in the region with the vertical arrows indicating the relative elevation range of the species found in the habitat types (Pfister et al. 1977, Arno 1979, Steele et al. 1983, Cooper et al. 1991).

fire and the introduction of white pine blister rust (*Cronartium ribicola* J.C. Fisch.) in these forests has caused an invasion of shade-tolerant species, such as grand fir. These changes in stand composition and structure make these forests more susceptible to insect and disease attack, and to fires (Agee 1993, Hessburg et al. 1994).

Mixed western redcedar (*Thuja plicata* Donn ex D. Don), western hemlock (*Tsuga heterophylla* [Raf.] Sarg.), grand fir, and Douglas-fir forests dominate parts of northern Idaho, western Montana, and eastern Washington. These forests are the most productive in the Inland Northwest. They are characterized by high amounts of precipitation (1,300 mm yr^{-1}) and fertile volcanic ash soils (Graham 1990, Cooper et al. 1991). Before the occurrence of white pine blister rust, large expanses of western white pine covered much of this region, but now it is usually a minor stand component. The understory vegetation in closed canopy forests consists primarily of forbs (e.g., *Clintonia* spp., *Smilacina* spp., and *Pyrola* spp.). In openings, low shrubs such as huckleberry (*Vaccinium* spp.) and gooseberry (*Ribes* spp.), and tall shrubs such as alder (*Alnus* spp.) and Rocky Mountain maple (*Acer glabrum* Torr.) can occur. Large accumulations of coarse woody debris are common, especially where western white pine mortality from blister rust has been high (Brown and See 1981, Harvey et al. 1989c, Reinhardt et al. 1991).

The Engelmann spruce (*Picea engelmannii* Parry ex Engelm.)/subalpine fir (*Abies lasiocarpa* [Hook.] Nutt.) zone lies at elevations above the mixed fir forests and western redcedar–western hemlock forests (Steele et al. 1981, 1983, Cooper et al. 1991). This zone is characterized by Douglas-fir, western larch, and lodgepole pine in seral stages, and Engelmann spruce and subalpine fir in later stages of succession (Pfister et al. 1977, Cooper et al. 1991). Slow decomposition rates and long fire intervals (100 to 400 yr) in these forests cause large accumulations of coarse woody debris (Brown and See 1981). The understory is usually dominated by medium (*Vaccinium* spp.) and tall (*Menziesia ferruginea* Smith) shrubs.

On the east slopes of the Rocky and Cascade Mountains, and on many subalpine fir habitat types, lodgepole pine can be perpetuated by fire (Pfister et al. 1977). Dense, stagnated stands of pure lodgepole pine often develop, which are prone to bark beetle (*Dendroctonus* spp. and *Ips* spp.) attack and large stand-replacing fires. These sites are characterized by short growing seasons and cold temperatures. The amounts of coarse woody debris varies considerably and depends on stand fire history. Low shrubs (*Vaccinium* spp.) and grasses (*Calamagrostis* spp.) usually dominate the understory.

The subalpine forest types dominated by white bark pine (*Pinus albicaulus* Engelm.), mountain hemlock (*Tsuga mertensiana* [Bong.] Carr.) and subalpine fir are found at the highest elevations of the region. Trees in these forests are often widely spaced and have an understory of bear grass (*Xerophyllum tenax* [Pursh.] Nutt.), low shrubs (*Vaccinium* spp.), and sedges (*Carex* spp.). Cold temperatures and the short growing season limit forest productivity. These forests generally have little accumulation of coarse woody debris (Brown and See 1981).

Soils

Parent Material

Inland Northwest forests occupy a vast area of the western United States in which erosional processes, active volcanism, and weathering regimes have interacted to produce a diverse landscape. The resulting forest soils are equally diverse and have generally developed from three types of parent materials: (1) volcanic ash (including pumice), (2) glacial deposits, and (3) weathered bedrock. Nine of the eleven soil orders are present with Inceptisols, Alfisols, and Andisols being the most abundant (Geist and Cochran 1991, Meurisse et al. 1991, Harvey et al. 1994).

Extensive areas of soils with volcanic ash caps are found on a variety of parent materials, ranging from loess deposits and glacial tills to granitic and schist bedrock. Pumice soils predominate in northwestern California and central Oregon, whereas ash cap soils are most common in eastern Washington, northern Idaho, and northwestern Montana. Thickest ash caps occur on north-facing slopes where soil erosion has been the least. Soils developed from these volcanic deposits have relatively high inherent productivities, low bulk densities, and few coarse fragments (Geist et al. 1989). These properties also make them susceptible to mechanical displacement, especially when soil moisture is low (Geist and Cochran 1991). Ash and pumice soils are usually well drained with moderate to high infiltration rates and water holding capacities. However, once disturbed, damage to ash soils by compaction to depths of 30 cm can last for decades (Froehlich et al. 1985). Other site disturbances, such as ash displacement, remove nutrients, lower water-holding capacity, and also reduce site productivity (Geist and Cochran 1991).

Continental and alpine glaciers deposited various sized tills and outwash materials in northern Washington, Idaho, and Montana during the Pleistocene (Meurisse et al. 1991). Volcanic ash later covered many of these deposits. The soil material below the ash cap usually has a high percentage of rock fragments, and a low moisture and nutrient-holding capacity. If the ash cap is reduced, compacted, or removed, the productive capacity of the soil is significantly lowered (Cochran and Brock 1985).

Soils formed from weathered granite, gneiss, and schist bedrock are located primarily in southern Idaho and eastern Oregon. These skeletal soils are mostly Inceptisols and Alfisols. Low summer precipitation in these regions, co with low soil water and nutrient-holding capacities, gene. limit tree growth on these soils (Childs 1982, Clayton and Kennedy 1985).

Soil Organic Matter

The type and distribution of soil organic matter in Inland Northwest forests is variable and depends on stand age and location, tree species, and fire history. Large amounts of soil organic matter are found in surface organic layers (forest floor and woody residue) of old-growth soils, except in the fire-dominated lodgepole and ponderosa pine ecosystems (Table 1). Each of these soil organic components has a unique chemical and physical character based on the type of organic material present and the nature of underlying mineral soil (Quesnal and Lavkulich 1981, Prescott et al. 1989). Surface mineral horizons (< 30 cm deep) also contain large amounts of organic matter, especially if the soil has a significant volcanic ash content (Page-Dumroese et al. 1991, Harvey et al. 1994). Surface soil is the zone of greatest importance for site productivity, since root numbers, and presumably root activity, decrease rapidly below this depth (Kimmins and Hawkes 1978, Vogt et al. 1981, Strong and La Roi 1985, Little and Shainsky 1992). It is also the part of the soil most likely to be disrupted or destroyed by forest management activities. Considerable organic matter can also be four deeper soil horizons, which can be very important tc productivity on certain sites (Henderson 1995).

Total organic contents of Inland Northwest forest soils generally reflect site productivity and climate. Forest productivity is greatest in warm, moist cedar and hemlock habitat types in northern Idaho, and lowest on dry Douglas-fir and ponderosa pine sites (Table 1). Subalpine fir habitat types also have high soil organic matter contents, but productivity is generally lower than cedar and hemlock sites (Pfister et al.

Table 1. Soil organic matter and nitrogen content in representative old-growth forests of the Inland Northwest.[1]

	Cedar/ hemlock		Subalpine fir	Douglas-fir	Ponderosa pine	Lodgepole pine	
	Idaho	Montana	Montana	Montana	Montana	Wyoming	Oregon
Organic matter (Mg ha^{-1})							
Woody residue[2]	154	83	146	45	<20	2	38
Forest floor	71	101	72	63	9	24	18
Mineral soil[3]	201	145	153	133	160	162	52
Total	426	329	371	241	<189	188	108
Nitrogen (kg ha^{-1})							
Woody residue	231	125	219	68	<30	2	40
Forest floor	476	1,128	914	857	161	358	174
Mineral soil	3,045	1,729	1,686	2,183	3,433	4,030	1,152
Total	3,752	2,982	2,819	3,108	<3,624	4,390	1,366

[1] Values are from one stand in each of the forest cover types. Taken from Fahey (1983), Page-Dumroese et al. (1991), Busse 1994, J.B. Yavitt, unpublished data.
[2] Woody residue > 7.6 cm in diameter, except for lodgepole pine (Wyoming) where all down dead wood was measured.
[3] To a depth of 30 cm. These values do not include root weights.

1977, Steele et al. 1981, Cooper et al. 1991). Low temperatures on these higher elevation habitat types probably limit ›th organic matter decomposition and tree growth. Site ɔductivity is generally low on Douglas-fir and ponderosa pine sites, but organic levels in surface mineral layers may be quite high beneath certain grass communities (Nimlos and Tomer 1982).

While soil organic matter/productivity relationships are readily apparent among tree species, this relationship is much less evident when applied to a single species over a large area. Poor correlations of soil organic matter content with height growth have been reported for Douglas-fir and western hemlock in western Washington and Oregon (Edmonds and Chappell 1994), and for Douglas-fir in Idaho and western Montana (Monserud et al. 1990). Soil organic matter/tree growth responses would also be confounded by other site factors, especially precipitation and evapotranspiration. The large variability of soil organic matter content in response to individual site differences, such as slope, parent material, and soil depth, also make regional organic matter/productivity extrapolations difficult.

Woody Residues

Woody residue can be any size, but material larger than 7.5 cm in diameter has the potential to become a long-term component of soil ecosystems. Wood less than 7.5 cm in diameter usually decays quite rapidly or burns readily in both prescribed or wildfires (Edmonds 1991, Reinhardt et al. '). Studies have shown that woody material can have ɔrtant roles in tree seedling establishment, soil carbon cycling, nutrient and water storage, and animal activity (Sollins et al. 1980, 1987, Harmon 1987, Amaranthus et al. 1989, Arthur and Fahey 1992, Harmon et al. 1994, Tallmon and Mills 1994). Woody residue in many Inland Northwest forests may equal or exceed organic matter contents in the forest floor (Brown and See 1981). Generally, the levels of woody residue in undisturbed stands increases with stand age, as site moisture conditions improve, and as fire hazard decreases (Spies et al. 1988, Harcombe et al. 1990).

Woody residue decay is primarily a function of invertebrate activity, the colonization of wood by white-rot and brown-rot fungi, and soil surface microclimate (Larsen et al. 1980, Swift and Boddy 1984, Harmon et al. 1986, Edmonds 1991). As wood decay progresses, water and N content generally increase (Jurgensen et al. 1984, Harmon et al. 1986). Initial sapwood decay frequently is of the white-rot type, which eventually shifts to brown-rot. Heartwood of down-woody residue is usually decayed by brown-rot fungi. Typically, brown-rot fungi remove cellulose and modify lignin, while white-rot fungi remove similar amounts of both cellulose and lignin. However, the rates at which these materials are decomposed vary with fungal species (Setliff and Eudy 1979, Boddy 1991). Brown-rot fungi largely con- : decay patterns of large woody residue in the Inland ı .west (Larsen et al. 1980).

When decayed woody residues become incorporated into the forest floor or mineral soil, they can be termed *soil wood.* This advanced decayed wood is also called class IV and V logs in coarse, woody debris classification systems used in Washington and Oregon (Eubanks 1989). At this stage the wood often is covered by litter and not noticed as a soil component. Soil wood can comprise over 50% of the surface soil organic matter in old-growth stands of the Inland Northwest (Page-Dumroese et al. 1991). A large soil wood component has also been reported on sites in Oregon, Washington, and western Canada (Little and Ohmann 1988, Prescott et al. 1989). In some stands this accumulation is so pronounced that a separate forest floor designation (Lignomor) has been established (Green et al. 1993). The amount of soil wood present on a site generally increases as overall forest productivity increases. Soil wood retains more moisture than the forest floor during dry summer months and can be quite resistant to destruction by fire, especially on highly productive hemlock sites (Page-Dumroese et al. 1991, Reinhardt et al. 1991, 1994).

Virtually all of the soil wood in Inland Northwest forests is a product of brown-rot decay and comes from large residues with appreciable amounts of heartwood, especially pine species and Douglas-fir (Harvey et al. 1987a). Brown-rotted wood remains in the soil for hundreds of years (McFee and Stone 1966, Harvey et al. 1981), thus affecting soil properties for long periods. In contrast, white-rotted soil wood, originating from conifer sapwood or hardwood sapwood/ heartwood, rapidly loses its structural integrity once it becomes part of the forest floor. However, it has a lower extractive content, a higher potential for nonsymbiotic N fixation, and more rapid nutrient fluxes than brown-rotted wood (Larsen et al. 1980, Harmon et al. 1986, Jurgensen et al. 1989).

Timber Harvesting Impacts

Timber harvesting and subsequent site preparation techniques can have a major impact on soil properties and productivity. Historically, clearcutting followed by burning or mechanical site preparation (e.g., soil scarification, scalping) has been favored in the Pacific Northwest to manage for early succession species, such as Douglas-fir or western larch (Graham et al. 1989a, Ferguson and Carlson 1991, Arno and Fischer 1995). Site preparation can improve early seedling growth and survival on many sites in this region where competition from other plants is high (Ross et al. 1986, Thomson and McMinn 1989, Packee 1990, Messier et al. 1995). However, these treatments, which are often accompanied by soil organic matter losses, soil compaction, and surface mineral soil displacement, can also reduce tree growth (Atzet et al. 1989, Powers 1991). The degree of soil disturbance will vary according to: (1) intensity of site treatment, (2) type of equipment used, (3) time of year harvesting occurs, and (4) amount of organic matter on the soil surface (Ross and Walsted 1986, Minore and Weatherly 1988, Clayton 1990, Graham et al. 1991b, Page-Dumroese 1993). Soils having higher amounts of organic matter in forest floor and soil wood would be more affected by these disturbances than soils having higher organic contents in surface mineral layers.

Site preparation methods that increase organic matter content in surface soil layers (e.g., mounding or bedding)

Table 2. Soil properties, western white pine seedling growth, and amount of competition on two sites in northern Idaho 3 yr after site preparation.[1]

	Soil properties				Seedling characteristics			
Site treatment	Bulk density (mg m^{-3})	Organic matter (kg kg^{-1})	Total N (g kg^{-1})	Available N (mg kg^{-1})	Height (cm)	Root depth (cm)	Total weight (g)	Compe vegeta (kg ha
Grand fir habitat type								
Mounded	0.70	0.17	2.9	62	26	28	7	9,192
Mounded-herbicide[3]	0.68	0.17	3.0	58	39	34	26	280
Scalped	0.92	0.08	1.3	18	32	22	15	684
None	0.76	0.12	2.6	32	36	24	10	3,760
Western hemlock habitat type								
Mounded	0.58	0.24	2.8	37	31	21	12	1,711
Mounded-herbicide	0.58	0.18	2.8	52	29	29	13	<50
Scalped	0.85	0.11	1.5	19	24	20	8	<50
None	0.65	0.27	2.2	66	28	21	8	<50

1 Values from Page-Dumroese et al. (1997).
2 Values from Graham et al. (1989a).
3 Herbicide applied—1.7 kg ha^{-1} glyphosate.

improve soil physical and chemical properties, and can result in greater seedling growth (Table 2, Sutton 1993). In contrast, site treatments that compact the soil and significantly lower surface soil organic and nutrient levels, such as scalping, can have negative impacts on seedling development. Scalping is a common method of site preparation in the Inland Northwest and can benefit seedling establishment and growth by eliminating competing vegetation from the planting site and raising soil temperature on cold sites (Table 3, Sutton 1985, Thomson and McMinn 1989,

Table 3. Seedling response to scalping and herbicide treatments in the Inland Northwest.[1]

					Plantation growth index[3]	
Species/treatment	Age[2]	Survival (%)	Height (cm)	Diameter (cm)	(Index)	(T/C[4])
Montana						
Ponderosa pine	6					
Control		31	64.7	13.2	3,453	—
Scalping		60	65.1	14.1	7,786	2.2
Herbicide		83	97.8	23.0	50,631	14.7
Lodgepole pine	6					
Control		31	53.7	9.4	1,526	—
Scalping		72	58.2	11.5	7,553	5.0
Herbicide		75	78.1	16.1	15,025	9.8
Washington						
Douglas-fir	3					
Control		75	34.2	5.2	813	—
Scalping		74	31.1	5.3	1,143	1.4
Herbicide		78	44.1	7.5	2,010	2.5
Idaho						
Douglas-fir	6					
Control		46	52.9	7.7	1,463	—
Scalping		74	64.7	11.5	7,704	5.3
Herbicide		46	66.7	11.6	4,111	2.8
Lodgepole pine	6					
Control		27	84.2	13.9	4,627	—
Scalping		83	118.6	24.8	79,852	17.3
Herbicide		31	100.5	16.0	10,950	2.4

1 Values from Boyd (1986). Herbicide data shown for Velpar. If different rates, formulations, or method of application used, results were averaged.
2 Number of growing seasons since planting.
3 Plantation Growth Index is the product of survival, mean height, and mean diameter^2—a value proportional to the amount of stemwood produced/100 planted trees.
4 T/C = the ratio of site treatment to control.

McNabb et al. 1993, Fleming et al. 1994, Powers 1997). However, as shown in Table 2 and in other studies (Miller and Breuer 1984, Graham et al. 1989a), scalping can sometimes have little benefit and occasionally reduces tree growth. An alternative site treatment to scalping is the application of herbicides, which reduce competition but maintain surface organic matter layers intact. In many instances, seedlings grow better after herbicide treatment when compared to scalping or grubbing (Table 3). These positive growth responses to herbicides occur even when plant competition is equal to, or greater than, scalped treatments. Consequently, scalping should only be used on sites where environmental concerns restrict herbicide application or mineral soil is required for seed germination.

A key issue of site preparation effects on soil properties is the length of time such treatments impact site productivity. Several studies have shown that tree growth is influenced by site preparation to at least the sapling stage (Cole and Schmidt 1986, Clayton et al. 1987, Graham et al. 1989a). One method to estimate site preparation effects on future tree development is to use a stand projection model such as PROGNOSIS (Wykoff et al. 1982). By using height increment and physical site information, this model estimates the impacts of site treatments on long-term productivity. A short-term scenario assumes that early growth responses to site preparation have stopped when the trees reach a diameter of 7.5–12.5 cm, while a long-term response assumes growth differences continue throughout a rotation (100 yr). Using the more conservative stand projection (short-term effects), scalping reduced total volume of western larch on two sites by 4–6% (Table 4). If a long-term response is assumed, scalping caused a 20–27% reduction in volume. In contrast, if soil is mounded and competing vegetation is controlled, a 5–12% increase is projected for the shortterm. If these site preparation benefits are extended over the 100 yr life of the stand, a 63–70% increased yield is projected. Similar results were found with Douglas-fir (Graham et al. 1991a). While other sites and stands may react differently, the potential impact of differing site preparation techniques on final tree volume is apparent.

Soil Organic Matter

Short-term losses of organic matter (1 to 5 yr) after clearcutting and site preparation have been determined by measuring woody residue and soil organic matter levels before and after harvest, or on adjacent cut and uncut stands. Results from these studies indicated organic matter losses from woody residues and the forest floor in western conifer forests ranged from 2 to over 150 Mg ha^{-1}, with higher losses found after prescribed burns (Cromack et al. 1979, DeByle 1980, Feller 1988, Reinhardt et al. 1991, Johnson 1992). Harvesting operations also incorporate forest floor material and harvest residues into the mineral soil, which can increase organic content in the surface soil for the first few years after harvest (Packer and Williams 1976, Cromack et al. 1979). However, these higher levels of organic matter in the mineral soil generally decrease as the new stand develops (Kraemer and Hermann 1979, Durgin 1980).

Longer term organic matter changes following harvest can be evaluated by comparing soil organic content in stands having different ages since cutting. Because preharvest soil conditions usually are not known, sites of similar stand history, species composition, topography, etc., are selected to limit the inherent variability in organic matter contents among soils. Unfortunately, only a few studies of this type have been conducted in western forests. Clayton and Kennedy (1985) estimated that it may take more than 50 yr to restore soil organic matter and nutrients to previous levels in disturbed forest ecosystems in central Idaho. Results from less controlled studies in the Inland Northwest also indicate reductions in the thickness of surface organic layers for a considerable time after

Table 4. Western larch growth projections to age 100 yr after site preparation on two sites in northern Idaho.[1]

	Short-term calibration[2]				Long-term calibration[3]		
Site treatment[4]	Seedling ht[5] (cm)	Mean diam (cm)	Mean ht (m)	Total vol (m^3 ha^{-1})	Mean diam (cm)	Mean ht (m)	Total vol (m^3 ha^{-1})
Grand fir habitat type							
Mounded	162	30.7	35.1	203	30.7	37.8	203
Mounded-herbicide	201	33.3	37.2	262	42.2	35.1	525
Scalped	113	31.5	34.8	224	32.0	36.0	235
None	140	32.5	34.4	233	35.8	40.5	323
Western hemlock habitat type							
Mounded	128	36.8	52.4	326	38.4	40.2	384
Mounded-herbicide	192	36.6	41.2	325	40.1	41.5	433
Scalped	107	35.6	37.5	289	31.5	37.5	201
None	128	36.6	38.4	309	34.0	38.7	254

1 Values from Graham et al. (1995).
2 The growth rate established by the 5 yr height increment attenuated to the average growth rate for the site by the time the trees reach 7.6 cm to 12.7 cm in diameter.
3 The growth rate established by the 5 yr height increment is maintained for the 100 yr.
4 Site preparations described by Page-Dumroese et al. (1997).
5 Seedling height at the end of eight growing seasons after planting.

harvesting or fire (Harvey et al. 1986). Most of this information was obtained from sites which were clearcut and had fairly intensive site preparation. However, the current emphasis on ecosystem management in western forests will likely reduce the use of these silvicultural practices in the future (O'Hara et al. 1994), thus alleviating some detrimental soil impacts associated with timber harvesting.

Losses of organic matter from the forest floor and mineral soil following timber harvesting are generally a result of increased organic matter decomposition by soil microorganisms (Hendrickson et al. 1982). Increased soil moisture, temperature, and alkalinity after harvesting, especially if fire is used for slash disposal, favors increased microbial activity (Hungerford 1980, Jurgensen et al. 1981, 1982). Mechanical site preparation mixes the forest floor into mineral soil, which generally increases organic matter decomposition rates (Salonius 1983), and improves the growth of subsequent regeneration (Graham et al. 1989a, Thomson and McMinn 1989). However, on some sites with thick H horizons, soil mixing can reduce microbial activity and organic matter decomposition (Messier et al. 1995).

Considerable interest has developed on leaving adequate amounts of woody residue after timber harvesting. It is generally recognized that some wood should be left on site to increase biodiversity, and this may also be important for long-term site productivity (Heilman 1990, Swanson and Franklin 1992). However, the optimal amount of woody residue to leave will depend on habitat type, fire hazard, regeneration method, and preharvest soil organic matter levels. The site preparation method used will also be a major factor (see Table 7). A minimum of 22 to 36 Mg ha^{-1} of residual woody material has been recommended for moist habitat types of northwestern Montana and northern Idaho (Harvey et al. 1987a). Leaving greater amounts of woody residue may further benefit stand productivity on many sites, but may also increase fire risk. After considering fire hazard and other site preparation objectives, Reinhardt et al. (1991) established a fairly wide range of allowable woody residue loadings (22–125 Mg ha^{-1}) for mixed conifer forests in the northern Rocky Mountains. However, on some of the drier sites in the region (e.g., ponderosa pine, lodgepole pine, and Douglas-fir), leaving residue loadings of less than 10 Mg ha^{-1} may be adequate to maintain soil organic matter levels (Graham et al. 1994).

How much organic matter to leave on a site after harvesting will also be influenced by whether the new stand is to be established by planting or by relying on natural regeneration. The removal of large amounts of surface organic matter by machine piling, windrowing, or prescribed burning to expose mineral soil is desirable when establishing regeneration from seed, especially western larch and ponderosa pine (Haase 1986, Shearer and Stickney 1991). However, seedbed preparation runs the risk of depleting soil organic matter reserves needed for subsequent seedling growth and development (Graham et al. 1989b, Minore and Weatherley 1990, Miller 1991). Such extensive soil disturbance and organic matter removals are normally not required if a site is planted with nursery stock (Harvey et al. 1987b).

Soil Nitrogen

Nitrogen is required for tree growth in greater amounts than any other mineral nutrient, and is usually the nutrient most limiting in western forest soils (Binkley 1991, Miller et al. 1991, Moore et al. 1994). Since nearly all soi' forest soils is present as organic forms, soil N cont Inland Northwest soils generally parallels that of soil organic matter (Table 1). Consequently, logging operations in western forests that result in large organic matter losses also remove considerable soil N (Cromack et al. 1979, Stark 1979, Jurgensen et al. 1981, Feller and Kimmins 1984). Increased residue removal, shorter stand rotation age, and total tree harvesting may further increase N losses (Wollum and Davey 1975, Jurgensen et al. 1980). If prescribed fire is also used as a site preparation tool, soil N losses can exceed 500 kg ha^{-1} (Feller 1988, Little and Ohmann 1988). Nitrogen lost in fires is directly proportional to the amount of organic matter consumed and temperatures produced (Mroz et al. 1980, Hungerford et al. 1991). The significance of such N depletion to site productivity depends on total N capital present in the soil and the magnitude of N losses, especially from surface organic layers (Jurgensen et al. 1980, Edmonds and Bigger 1984).

Replacement of soil N lost from timber harvesting or fire would come from precipitation and dry deposition (dust, pollen, etc.), biological nitrogen fixation, and forest fertilization. Nitrogen fertilizer applications are economically feasible on many sites in the Inland Northwest (Moore et al.1994), but may increase risk of endemic root disease or animal damage in some stands (Nelson 1989, Entry 1991, Miller et al. 1991). In areas with high precipitatic large urban and industrial centers, such as the northeastern United States, atmospheric N inputs can add > 20 kg N ha^{-1} yr^{-1} (Kimmins et al. 1985). However, in the relatively dry Inland Northwest with its low population density, N additions to forested sites normally average between 0.5 and 2.5 kg ha^{-1} yr^{-1} (Tiedemann et al. 1978, Clayton and Kennedy 1985, Fahey et al. 1988, NADP/NTN 1992). Nitrogen additions from biological N fixation come from two sources: (1) microorganisms living in specialized nodules on plant roots (symbiotic N-fixation), and (2) free-living soil microorganisms (nonsymbiotic N-fixation). The contribution of each varies considerably depending on site location, elevation, soil type, stand seral stage, and cover type. Most of the estimates of N fixation in western forests were obtained using the acetylene reduction technique (Silvester 1983). While many questions have been raised about the suitability of this technique, it does give a relative measure of N-fixing potential among various ecosystems and site treatments.

Symbiotic Nitrogen Fixation

Nitrogen-fixing plants in forests of the Inland Northwest can be grouped into two major categories: (1) plants in the family Leguminosae—9 genera, and (2) nonleguminous plants—5 genera from 4 families (Table 5). The distrib patterns of these plants generally reflect soil moisture perature conditions and stand successional stage. Legu are more widely distributed than nonleguminous N-fixing

Table 5. Occurrence of nitrogen-fixing plants in forest climax series of Idaho, Western Montana, and Western Wyoming.[1]

	Pinus flexilis (58)[2]	*Pinus ponderosa* (180)	*Pseudo-tsuga* (1,124)	*Abies grandis* (411)	*Thuja* (311)	*Tsuga hetero-phylla* (189)	*Picea* (219)	*Tsuga merten-siana* (103)	*Abies lasio-carpa* (1,645)	*Pinus albicaulis* (53)
	Percent of stands in which genus found									
Nonlegumes										
Alnus	0	0	9	4	4	4	10	3	8	0
Ceanothus	3	9	11	10	4	2	0	0	1	0
Cercocarpus	5	2	3	0	0	0	0	0	0	0
Purshia	7	31	8	<1	0	0	0	0	<1	6
Shepherdia	19	6	14	4	1	5	35	0	17	23
Legumes										
Astragalus	53	16	19	4	3	0	22	0	7	28
Hedysarum	21	5	4	<1	<1	0	11	2	5	7
Lathyrus	0	8	2	10	6	<1	3	0	1	0
Lotus	0	4	0	0	0	0	0	0	0	0
Lupinus	12	41	24	5	1	2	9	2	18	23
Oxytropis	7	2	<1	0	0	0	0	0	0	2
Thermopsis	0	1	<1	13	9	0	0	2	1	0
Trifolium	0	5	2	6	3	<1	3	0	2	2
Vicia	2	17	4	6	4	2	5	0	1	0

[1] Data are from late seral and climax forest stands (Pfister et al. 1977, Steele et al. 1981, 1983, Cooper et al. 1991).
[2] Total number of stands examined within each habitat series.

plants in older stands throughout the region. *Lupinus* spp. is the most common N-fixing genus across all habitat types, while buffaloberry (*Shepherdia canadensis* [L.] Nutt.) is the most frequently occurring nonleguminous N-fixing plant. Bitterbrush (*Purshia tridentata* [Pursh. DC.]) is restricted to drier sites, while *Alnus* spp. is found in cooler, wetter sites. *Ceanothus* spp. shows only a scattered distribution in these older stands. Ponderosa pine and Douglas-fir sites have the greatest occurrence of N-fixing plants, which likely reflects their more open stand structure.

Although species of at least 1 N-fixing plant occurred in all but 6 of the 115 habitat types examined in Montana, Idaho and Wyoming, these plants are not major understory components in most older Inland Northwest forests. Even when several N-fixing plants are found on the same site, the combined coverage of these species rarely exceeds 10% (Jurgensen et al. 1979). In contrast, early seral forests resulting from timber harvesting or fire usually have greater understory development and higher N-fixing plant frequencies than older stands (Bormann and Gordon 1989).

Most ecological studies on N-fixing plants have been very site specific and usually detail early successional development after a disturbance, such as timber harvesting or fire. *Ceanothus* spp. has received particular attention, since it often becomes abundant after prescribed burns or wildfires. *Ceanothus* spp. seed, which can remain viable in the soil for up to 200 yr, requires a heat treatment to break dormancy (Noste and Bushey 1987). Reports of *Ceanothus* spp. percent coverages on burned sites in Idaho and Montana have ranged from less than 5% to over 80% (Jurgensen et al. 1991). Generally, the hotter the burn, the greater the development of *Ceanothus* spp. (Orme and Leege 1976, Noste and Bushey 1987).

The occurrence of several other N-fixing plants also increases after site disturbance. *Trifolium* spp. has been found on disturbed and burned sites in northern Idaho (Mueggler 1965, Stickney 1986), and *Hedysarum* spp. responded strongly to burning and mechanical site preparation on two habitat types in western Montana (Arno et al. 1985). *Alnus* spp. was either more or less common after timber harvesting or fire in various habitat types of Idaho and Montana (Mueggler 1965, Wittinger et al. 1977, Stickney 1980, 1986). Conflicting results have also been reported for *Lupinus* spp. and *Astragalus* spp. (Zamora 1975, Lyon and Stickney 1976, Arno et al. 1985, Stickney 1986, Steele and Geier-Hayes 1987, 1989, Brown and DeByle 1989). In contrast, coverage of both *Purshia* sp. and *Shepherdia* sp. was reduced after fire (Wagstaff 1980, Brown and DeByle 1989). The scattered and incomplete nature of these studies indicates that much more information is needed on the successional roles of N-fixing plants in this region.

Little information is available on the actual contribution of N-fixing plants to the N economy of Inland Northwest forests after harvesting. Studies on moist, highly productive sites in western Oregon and Washington have reported additions over 100 kg of N ha^{-1} yr^{-1} from *Ceanothus* spp. and *Alnus* spp. (Conard et al. 1985, Hibbs and Cromack 1990, Binkley et al. 1992). Nitrogen gains from N-fixing plants are likely lower on the generally drier sites in the Inland Northwest. Estimated N fixation by snowbrush (*Ceanothus velutinus* Dougl. ex Hook.) in central Oregon ponderosa pine stands range from 7 to 72 kg N ha^{-1} yr^{-1}, depending on stand age and snowbrush coverage (Youngberg and Wollum 1976, McNabb et al. 1979, Busse 1992). In contrast, Hendrickson and Burgess (1989) reported that *Lupinus* spp. and *Shepherdia* sp. added only 2.8 kg N ha^{-1} yr^{-1} to a cutover lodgepole pine site in southern British Columbia. Other estimates of N fixation by *Lupinus* spp. range from 0.1 kg N ha^{-1} yr^{-1} in older, southeastern Wyoming lodgepole pine stands (Fahey et al. 1985) to 0.6 kg N ha^{-1} yr^{-1} in Utah aspen stands (Skujins et al. 1987). Nitrogen additions from *Purshia* sp. in ponderosa and lodgepole pine stands of central Oregon can vary from

Table 6. Nonsymbiotic nitrogen fixation in old-growth stands of the Inland Northwest.[1]

	Subalpine fir (Montana)		Cedar/hemlock (Montana)		Douglas-fir (Montana)		Cedar/hemlock (Idaho)	
Soil component	(g N)	(% N fix)	(g N)	(% N fix)	(g N)	(% N fix)	(g N)	(% N fix)
Wood residue	515[2]	35[3]	230	27	159	20	1,428	49
Forest floor	328	22	192	23	101	13	88	4
Soil wood	250	17	91	11	95	12	178	6
Mineral soil[4]	379	26	326	39	442	55	1,197	41
Total	1,472		839		797		2,891	

1 Values from Jurgensen et al. (1991).
2 Total N fixed/ha over a 180 day period in 1977 as measured by the acetylene reduction technique using a 3:1 ethylene to N conversion ratio.
3 Percentage of total soil N fixed in that soil component.
4 Mineral soil sampled to a depth of 30 cm.

less than 0.1 kg N ha^{-1} yr^{-1} to 8 kg N ha^{-1} yr^{-1} (Dalton and Zobel 1977, Busse 1992).

There has been worldwide interest in artificially establishing N-fixing plants after harvesting, especially under intensive plantation management, or after fire (Turvey and Smethurst 1983). Several studies on seeding N-fixing legumes on cut and burned sites have been conducted in the Inland Northwest (Everett et al. 1991, Trowbridge and Holl 1992). Early results have been encouraging. On some dry sites, N-fixing plants may enhance natural regeneration by providing shade during the critical seed germination and seedling establishment period (Geier-Hayes 1994, Jones 1995). However, on many other sites N-fixing plant competition for light and water may outweigh the possible N benefits to tree seedlings (Stewart et al. 1984, Petersen et al. 1988, McDonald and Fiddler 1989, Loucks and Harrington 1991). Additional studies are needed to determine the management potential for such N-fixing plant treatments.

Nonsymbiotic Nitrogen Fixation

Reports of prolific development of *Ceanothus* spp. in certain habitat types after timber harvesting or fire should not obscure the fact that this and other N-fixing plants are lacking or of low frequency on many harvested sites in the Inland Northwest. On these sites, the replacement of N losses due to timber removal would have to come from nonsymbiotic N fixation and atmospheric deposition. Nitrogen additions from nonsymbiotic N fixation are highest in moist northern Idaho cedar or hemlock habitat types, and lowest on dry Douglas-fir habitat types in western Montana (Table 6). Surface organic materials contributed well over half of the nonsymbiotic N fixed in these soils, except for the dry, low organic matter Douglas-fir site. Nonsymbiotic N fixation rates show considerable annual variation, which relate to yearly differences in soil temperature and moisture (Jurgensen et al. 1992).

Coarse woody residue is an especially important source of N fixation in Inland Northwest forests and can account for up to 50% of the total N fixed (Table 6). The greater the woody residue loadings on a site, the greater the N gains from nonsymbiotic N fixation. On warm, dry sites, the accumulation of woody residue and N fixation is lower than on w more productive sites. While residue weights for these s (see Table 1) are higher than regional averages, they are nowhere near the maximum for these forest types (Jurgensen et al. 1987). Consequently, on sites with heavy residue loadings, such as in overmature stands on moist sites, gains from N fixation could be much higher.

Nonsymbiotic N fixation is especially susceptible to harvesting impacts due to its dependence on adequate organic matter supplies (Jurgensen and Davey 1970, Sprent and

Table 7. Nonsymbiotic nitrogen fixation as affected by clearcutting and site preparation treatments on a cedar/hemlock site in northern Idaho.[1]

	Clearcut — residue treatment									
	None		Prescribed burned		Intensive removal		Heavy slash[2]		Uncut	
Soil component	(Mg[3])	(g N[4])	(Mg)	(g N)	(Mg)	(g N)	(Mg)	(g N)	(Mg)	(g N)
Wood residue[5]	146.0	984	57.9	177	0.6	111	249.8	1,483	154.3	1,428
Forest floor	16.7	110	5.5	26	13.3	73	34.5	326	23.2	88
Soil wood	50.9	109	22.4	47	51.6	195	50.3	430	47.9	178
Mineral soil	—	1,218	—	826	—	1,125	—	1,608	—	1,197
Total	213.6	2,421	85.8	1,076	75.5	1,504	334.6	3,847	225.4	2,891

1 Values from Jurgensen et al. (1992).
2 Residue from the intensive removal plots added to these plots.
3 Dry mass ha^{-1} of organic material on top of mineral soil.
4 Estimate of total N fixed ha^{-1} over a 180 day period.
5 Woody residue > 7.6 cm in diameter.

Sprent 1990). The greater the amounts of organic matter removed or destroyed by harvesting and site treatments, the greater the reduction in N fixation. Clearcutting and subsequent prescribed burning lowered N fixation on a cedar/hemlock site in northern Idaho by 63%, compared to an adjacent uncut stand (Table 7). Clearcutting and slash removal by dozer (intensive removal) lowered N fixation by 48%, while clearcutting without any site preparation reduced N fixation by only 16%. In contrast, increasing woody residue and forest floor weights above preharvest conditions increased N fixation by 33%.

Clearcutting and subsequent site preparation greatly reduces the amount of woody residue on harvested old-growth sites (Table 7). The contribution of woody residues to soil N inputs after harvesting depends on both the amount and the decay stage of woody material left. Nitrogen fixation rates generally increase as wood decay progresses (Jurgensen et al. 1987, Griffiths et al. 1993). Most woody residue in old-growth stands are large decaying logs, many of which are destroyed during harvest. They are replaced by smaller amounts of less decayed wood from cut trees, which have much lower rates of N fixation (Jurgensen et al. 1992). Soil wood generally is less disturbed by harvesting and site treatments, and consequently becomes a more important site of N fixation.

While annual soil N gains from nonsymbiotic N fixation after harvesting are small, they become significant as new stands age. For example, in northwestern Montana only 110 kg of soil N was lost by clearcutting a subalpine fir stand, followed by a low intensity prescribed fire (Jurgensen et al. 1981). Assuming an average N fixation gain of 0.6 kg N ha^{-1} yr^{-1} for this site (Jurgensen et al. 1992), and 1.5 kg N ha^{-1} yr^{-1} is added in precipitation (NADP/NTN 1992), these N losses would be replaced in 52 yr. When N removed in bolewood is also included (175–200 kg N ha^{-1}), the length of time to replace this N loss would increase to 135–150 yr. In contrast, soil N loss from clearcutting and prescribed burning an old-growth cedar/hemlock site in Idaho was 640–700 kg N ha^{-1} (Jurgensen et al. 1991). Using an N fixation rate of 1.1 kg ha^{-1} yr^{-1} (Table 7), and 1.5 kg N ha^{-1} yr^{-1} added in precipitation, these N losses would be replaced in 250–270 yr. Such a slow return to original soil N levels could have a considerable impact on subsequent stand growth.

As discussed earlier, N-fixing plants may become established in many habitat types after harvesting. If this occurred on either of the two sites discussed above, soil N losses would be replaced much more rapidly than by nonsymbiotic N fixation alone. Assuming symbiotic N fixation rates could range from a low of 3 kg N ha^{-1} yr^{-1} (Hendrickson and Burgess 1989) to a high of 72 kg N ha^{-1} yr^{-1} (Youngberg and Wollum 1976), the recovery time to preharvest soil N levels could vary from 10 to 125 yr on the cedar/hemlock site, and only 4 to 60 yr on the subalpine fir site. However, competition from these N-fixing plants for light and water could greatly reduce tree seedling growth response.

Other sources of N fixation not accounted for in these calculations could also add appreciable amounts of N to the soil and reduce the time required for site recovery. Nonsymbiotic N fixation in tree stumps, cull trees left for snags after harvesting, and large dead roots can add up to 1 kg ha^{-1} yr^{-1} (Granhall and Lindberg 1980, Harvey et al. 1989b). Increased development of N-fixing algae and lichens on the soil surface after harvest can add N, but this is likely very low due to generally dry conditions in this region throughout the summer. Nitrogen-fixing bacteria are also active in the rhizosphere of developing tree seedlings and shrubs (Amaranthus et al. 1990) and in mycorrhizae (Li et al. 1992), but how much N they add to the soil is unknown.

Mycorrhizal Root Development

Nearly all of the important tree species in the Inland Northwest are ectomycorrhizal, except western red cedar, which has vasicular-arbuscular mycorrhizae. Ectomycorrhizal root development is strongly related to stand productivity in many habitat types of this region (Harvey et al. 1979). Studies in old-growth stands have shown a strong relationship of ectomycorrhizal root tips with soil organic layers (Table 8). Even though surface organic matter comprised only 6 to 33% of the surface soil volume on these sites, they contained > 70% of the active ectomycorrhizal root tips except for very dry ponderosa pine. Even when surface organic layers are reduced in thickness due to harvesting, silvicultural operations, or fire, ectomycorrhizae still predominate in these layers (Harvey et al. 1986).

Ectomycorrhizae show their best development in the forest floor, but brown-rotted soil wood is also a good site for root development, especially in cedar and hemlock forests (Page-Dumroese et al. 1994). Studies in other forest ecosystems have reported similar results (Bernstein 1960, Day and Duffy 1963, Knapp and Smith 1982, Geier-Hayes 1987, Webb 1988, Perala and Alm 1989). In periods of adequate moisture, the forest floor supports the highest level of ectomycorrhizae, but during dry summer condi-

Table 8. Ectomycorrhizal root tips in the surface soil of old-growth stands in the Inland Northwest.[1]

	Soil volume[2]		Ectomycorrhizae	
Stand[1]	Surface organic[3]	Mineral soil	Surface organic	Mineral soil
	(%)			
Montana				
Western hemlock	32	68	89	11
Subalpine fir	33	67	93	7
Douglas-fir	21	79	77	23
Idaho				
Western hemlock	17	83	80	20
Western white pine	17	83	93	7
Washington				
Ponderosa pine	14	86	16	84
Wyoming				
Lodgepole pine	6	94	71	29

[1] Stands sampled in late spring/early summer (values from Harvey et al. 1986).
[2] % of soil volume to a 30 cm depth.
[3] Forest floor and soil wood.

Table 9. Ectomycorrhizal root tips, soil moisture content, and soil temperature during the growing season in an old-growth subalpine fir stand in western Montana.[1]

	May/June			July/August			September/October		
	Myco[2]	H_2O (%)	Temp (C°)	Myco	H_2O (%)	Temp (C°)	Myco	H_2O (%)	Temp (C°)
Forest floor	369	130	11.8	75	74	14.1	59	142	8.0
Soil wood	75	205	11.0	109	118	12.2	28	244	7.2
Mineral soil[3]	14	37	10.0	1	28	11.4	3	41	7.0

1 Values from Harvey et al. (1978).
2 Numbers of ectomycorrhizal root tips per liter of soil volume.
3 Sampled to a depth of 30 cm.

tions prevalent in this region, soil wood is the most active site (Table 9). Soil wood usually is wetter and cooler than the surrounding forest floor and is affected less by daily or seasonal moisture and temperature changes (Hungerford 1980, Reinhardt et al. 1991). However, on sites where the soil already is cool and moist (e.g., Alaska), soil wood may be detrimental to root growth (Loopstra et al. 1988).

An organic matter-ectomycorrhizae relationship is also seen in the establishment and growth of young conifer seedlings. In old-growth stands, natural regeneration generally becomes established on microsites with deeper than average surface organic horizons (Table 10). This is especially evident in cool, moist subalpine fir and western hemlock stands. In contrast, organic matter is not beneficial to early seedling establishment on ponderosa pine sites, where mineral soil provides necessary conditions for seed germination. However, once seedlings become established on these dry sites, ectomycorrhizal root development is favored in the forest floor and soil wood (Harvey et al. 1989a).

The successful establishment of tree seedlings after timber harvesting also depends on adequate ectomycorrhizae development (Harvey et al. 1986, Perry et al. 1987, Kropp and Langlois 1990). Maintaining adequate amounts of organic matter on the soil surface after harvesting is important for ectomycorrhizae development (Harvey et al. 1987a). Forest floor and soil wood can be favorable seedbeds for the establishment of natural regeneration in recently harvested stands (Harvey et al. 1986, Geier-Hayes 1987, 1994, Tonn and Graham 1991). Reduced ectomycorrhizae formation occurs on some sites after timber harvesting, particularly where fire is used as site preparation (Harvey et al. 1980a and b, Perry et al. 1982, Parke et al. 1984).

Table 10. Soil organic matter depths supporting natural conifer seedlings in 4 old-growth stands in western Montana.

	Depth of surface organic matter[1]	
Stand type	Stand average (cm)	Seedling[2] (cm)
Ponderosa pine		
Forest floor	1.0	0.1
Soil wood	0.2	0.0
Total	1.2	0.1
Douglas-fir		
Forest floor	1.4	1.2
Soil wood	0.8	1.2
Total	2.2	2.4
Subalpine fir		
Forest floor	1.9	2.9
Soil wood	1.7	4.5
Total	3.6	7.4
Western hemlock		
Forest floor	1.8	4.2
Soil wood	1.9	1.6
Total	3.7	5.8

1 Values from Harvey et al. (1987b, 1989a).
2 Soil horizon depths beneath 100 randomly selected seedlings/stand.

Ectomycorrhizae have pronounced effects on the quality and quantity of carbon allocated to tree seedling root systems (Vogt et al. 1991, Rygiewicz and Anderson 1994). While most studies have shown beneficial effects of ectomycorrhizae on seedling growth, they occasionally may have little or even a negative effect (Bledsoe et al. 1982, Gagnon and Langlois 1987, Shaw et al. 1987, Harvey et al. 1996). In these situations, the energy costs for ectomycorrhizal seedlin maintain fungal hyphae and greater numbers of fine outweigh the benefits derived from increased water or nutrient uptake. Ectomycorrhizae-induced growth reductions most likely occur under harsh site conditions, such as drought or heavy competition, but can vary by tree species (Perry et al. 1987, Harvey et al. 1996). As shown earlier, growth-limiting moisture and/or nutrient levels are more common in soils with low organic matter contents. However, growth reductions caused by mycorrhizal fungi under these high-stress soil conditions are more than offset by increased seedling survival (Harvey et al. 1986).

Forest Health

Forest diseases and insects are often assumed to have only negative influences on stand development. However, in terms of site productivity, they can be very important to the functioning of certain forest ecosystems. These organisms have integral roles in the trophic dynamics of carbon and N cycling, habitat development, and landscape diversity (Harvey et al. 1992, Harvey 1994). The connection between forest stress and forest health is especially important in forests of the Inland Northwest, which are often limited by low soil moisture and nutrient levels (Geist and Cochran 1991)

Soil organic content, insect and disease activities, an incidence are strongly interrelated. This is especially t this region, where organic matter is generally produced at a higher rate than it can be cycled through the decay process

(Harvey 1994). Current forest health problems in these ecosystems are driven primarily by accumulations of both live nd dead organic matter, accompanied by high stand densi-ies and changes in tree species composition (Monnig and Byler 1992, Hessburg et al. 1994, Lehmkuhl et al. 1994). These problems may also be aggravated by soil nutrient deficiencies (Moore et al. 1994). This regional organic matter buildup is mostly due to fire suppression and tree mortality from white pine blister rust (Covington et al. 1994, Harvey 1994). In addition to causing insect and disease problems, high levels of surface organic matter also immobilize considerable amounts of soil nutrients in woody biomass, and increase the incidence of stand-replacing fires. Consequently, controlling organic matter accumulation will be as important to future forest health and productivity as maintaining adequate organic reserves (Oliver et al. 1994).

Harvesting and site preparation practices can have a major impact on forest disease incidence and insect populations (Martin 1988, Harvey et al. 1992, Lehmkuhl et al. 1994). Of major importance is the amount and type of organic materials left on site after harvesting operations. In some ecosystems the accumulation of surface organic matter can increase the incidence of insect and disease attack (Fellin 1980, Thies and Russell 1984). Large fuel accumulation after timber harvesting operations also creates the potential for hot, stand-replacing fires, and large losses of soil organic matter (Harvey 1994). Even when organic matter buildup does not occur, ber harvesting can aggravate endemic root rot and insect blems by increasing stress in the subsequent stand (Fellin 1980, McDonald et al. 1987, Stoszek 1988). Some sites in northern Idaho have experienced multiple harvests since the late 1800s, and they are now losing as much as 50% of their potential productivity to root rots. This has been caused by improved fire control (favoring more susceptible hosts), off-site planting, inappropriate seed sources, lower soil productivity, and increased fungal inoculum from infected roots and residue (Harvey et al. 1989a).

Maintaining historically appropriate organic matter reserves after timber harvesting and site preparation is important for maintaining forest health and soil productivity (Everett et al. 1994). Interactions between forest management prescriptions, insects, and disease fungi will be major considerations for most forests in the Inland Northwest (Harvey 1994). The relationship of forest health to soil organic matter will vary considerably among different ecosystems, depending on tree age, stand structure, and fire susceptibility. However, minimum organic contents on each site should be defined and maintained within specified target ranges (Reinhardt et al. 1991, Oliver et al. 1994).

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