Influences of Wind and Snow on Northern Tree-Line Environments at Churchill, Manitoba, Canada

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ABSTRACT. The influences of wind and snow distribution at the tree line near Churchill, Manitoba, were studied. Wind profiles above the snow surface, snow crystal size and quantity were examined during abrasion experiments with white spruce branchlets. For white spruce, the abrasion zone was evidently within 80 cm above the snow surface, and needle removal by abrasion was strongly influenced by branch age. Removal by abrasion of similar-aged needles was highest in new branches and declined with branch age up to 7 or 8 years, and then increased again with older branches. Trees that grew 80 cm in height in 7 or 8 years were successful in passing through the abrasion zone.

In forest-tundra environments the matted forms of trees were snow covered early in winter and received little abrasion. Sparsely needled islands of trees were largely covered with snow at the base. More densely needled trees and some trees within woodlands were less exposed to abrasion. The blockage of air flow resulted in high-speed jets, which cleared out snow within a "horseshoe-shaped vortex" around the base of the tree. Both in much of the woodlands and in the open forest, snow abrasion was minimal or non-existent and snow load on the branches (qali) becomes the prevalent stress.

During winter, a large proportion of the snowfall was initially blown off the exposed surfaces of Hudson Bay and the coastal tundra regions, into the woodlands, and eventually across the tree line and into the open forest. As the woodlands filled up with snow in mid-winter, the rate of snow accumulation in the forest increased from double to triple the snowfall. Variations in the rate of accumulation occurred with wind speed and direction.

During May, snowmelt began on exposed tundra first and usually ran off the frozen surface. Snowmelt occurred about three weeks later in the open forest and was characterized there by less runoff, as the water more readily permeated the thawing peat. The late snow beds are characteristic of the forest-tundra woodlands and were usually gone by mid-July. The woodlands were snow free for 1.5-2 months during the year, while the open forest was snow free for about 3 months and the tundra was largely snow free for 6 months or more.

Key words: tree line, wind, snow, white spruce, abrasion

INTRODUCTION

In tree-line ecosystems, the importance of wind in winter is demonstrated by its effects: desiccation by wind (Warlde, 1981; Frey, 1983), abrasion by wind-driven snow particles (Hadley and Smith, 1987, 1989), and friction from strong winds (Grier, 1988), which can severely damage overwintering trees. The redistribution of snow by wind may result in the protection of trees from negative wind effects during winter (Marr, 1977; Hadley and Smith, 1983, 1987) but also can cause severe damage during snowmelt (Payette and Filion, 1985; Lavoie and Payette, 1992). Within the tree line, trees have reached sufficient density to buffer the direct wind effects and subsequently modify their environment. Beyond the tree line, trees are more scattered and different stresses are prevalent. One distinction is the recognition of different (apparently wind-generated) crown forms (Yoshino, 1973; Payette, 1974; Scott et al., 1987; Stevens and Fox, 1991). Unique tree responses to wind stress indicate variations in physical environments in the area of the tree line and that a classical ecotone or gradient approach to snow-loaded forests and wind-abraded islands of tree shrubs may be an oversimplification of separate and discrete systems. Identifying
the processes that lead to different crown forms is fundamental to understanding the development and subsequent stability of the open forest and forest-tundra systems around the tree line.

This study reports a series of experiments that complement descriptions of wind-generated processes occurring during winter in a tree-line region. First, the relevance of wind to the development of the zone of abrasion and the impact of that abrasion on needle and branch survival in white spruce (*Picea glauca* [Moench] Voss) are examined. Then the wind-speed and snow-depth profiles around different crown forms of spruce in their natural environment are evaluated. Finally, snow redistribution relative to the presence or absence of spruce individually, in islands and woodlands and within the open forest is compared and discussed relative to the snow-free period.

**METHODS**

**Regional Description**

The tree-line region around Churchill, Manitoba (58°45′N, 94°04′W) has been delineated using white spruce crown forms (Scott et al., 1987). An area close to Hudson Bay where trees are absent or stemless mats exist in low abundance is defined as tundra. The resultant boundary is similar to that from defining tundra by the extreme limit of trees in other studies (Tranquillini, 1979; Payette, 1983; Kullman, 1990). The forest-tundra is composed of wind-abraded islands of trees and occasional woodlands that have varying crown forms. The region where greater than 95% full-crowned trees are prevalent is defined as the open forest. The line between the open forest and the forest-tundra defines the tree line in this study.

**Tree Abrasion Zone**

The abrasion zone was defined by measuring two separate profiles of wind velocity above the snow surface. Two anemometers were wired to an encased CR21x (Campbell Scientific Instruments, Edmonton, Alberta) datalogger that was strapped to a komatik (sled) and heated with thermochemical Hot Shots (Johnson & Johnson, Peterborough, Ontario). The reference anemometer measured wind speed at 1.6 m above the snow surface. The second anemometer measured wind speed at 25 cm intervals above the snow surface. The CR21x was programmed to integrate the wind speed over 20 second intervals and then average three such measurements. For each profile the procedure was repeated over a minimum of three minutes. The average wind speeds at each height interval were then converted to percentages of the reference wind speeds.

To examine the character of snow in the abrasion zone, ice crystals were captured and measured within the first metre above the snow surface. Glass vials were taped onto a grill at 1 cm intervals for the first 20 cm, then 4 cm intervals up to 1 m. The grill was placed in an exposed area and aligned so that the trapping vials were facing obliquely to the wind for a 12 hour period with wind speeds averaging approximately 30 km·h⁻¹ and exceeding 40 km·h⁻¹. In a sheltered, unheated room, a minimum of 20 ice crystals were sampled from each vial, placed on a calibrated slide and examined under a microscope. The longest two axes were measured to the nearest 0.01 mm so that the cross-sectional area of the largest surface could be determined.

To relate the rate of vertical growth in white spruce at tree line to the effects of snow abrasion, four full-crowned and four abraded-crowned white spruce trees were initially chosen as a branch source. The eight trees were 40-60 years old, apparently healthy and appeared representative of local populations. For each tree the (undamaged) vertical growth over the last 12 years was measured and recorded.

The impact of wind and ice crystals on needle abrasion was examined during mid-February with 24 three-year-old tree-line white spruce branchlets. The branchlets were cut from the sample trees at random, weighed to the nearest 0.05 gm with a triple beam balance (Ohaus, Florham Park, N.J. 07932, U.S.A.), and then stapled to wooden poles at 20 cm intervals (Scott and Hansell, 1992). In an exposed area each of the poles was pushed into the snow so that the branchlets ranged from 20 to 160 cm above the snow surface and was readjusted when necessary to maintain the branchlets at these levels. After 10 days each branchlet was removed from the posts and reweighed. The entire process was undertaken outside at ambient temperature.

To determine the susceptibility of the similar-aged needles from branches of variable age to abrasion, 96 two-year-old branchlets were cut from north-, south-, east- and west-facing branches of the third, seventh and eleventh whorls of the 8 white spruce sample trees. (Relative to needle viability, three-year-old branches would be young, seven-year-old mature and eleven-year-old senescent [Clark, 1961].) The branchlets were weighed in the field at ambient temperature with a sheltered triple beam balance. Random pairs were stapled horizontally to wooden stakes and maintained 10 cm above the snow surface for 7 days. The branchlets were examined and reweighed to determine change in weight. The change in water content during the experiment was corrected for by a negative exponential regression (see Scott and Hansell, 1992). The residual change in weight was attributed to needle loss.

**Wind Speed and Snow Distribution**

Patterns of wind stress around trees were examined by measuring wind speed at 1 m above the snow surface and snow depths on grids of 1 m intervals around a sparsely needled white spruce island, a densely needled spruce island and spruce trees within a woodland. Measurements began at the edge of the tree crowns on the four cardinal directions. Wind speeds were recorded using the dual anemometers described above and converted to percentages relative to an undisturbed reference value. Wind speed measurements were integrated over three 20-second intervals and replicated three times. Snow depths were also measured on the 1 m grid by pushing a calibrated metal rod through the snow layer.

The distribution of snow on typical exposed forest tundra was examined by measuring the snow depth on an 86-point grid at 10 m intervals. Snow depths were also measured at selected locations along a transect that passed through a
The wind profiles (Fig. 1) were created during an ambient wind speed of around 20 km·h⁻¹ and 30 km·h⁻¹, typical of conditions during February. Wind speed increased exponentially above the snow surface, reaching 79% of the reference value by 25 cm and exceeding 90% by 75 cm. Snow crystals were most numerous near the surface, where wind speed was least. There was a high frequency of hexagonal crystals (including sub-hedral, dihedral, II [c], I, a; after Nakayana, 1954) near the snow surface, with increasing proportions of hollow tubes and horns (IV [CP] 3) above 10 cm and with winged crystals (III [C], I, a, c) dominating above 80 cm. The cross-sectional area of the crystals within 1 cm of the snow surface was 0.230 mm² (SD ± 0.190). Crystal size declined rapidly with height, until at 10 cm the average crystal was 0.022 mm² (SD ± 0.017). The average cross-sectional area at 24 cm height was 0.009 mm² (SD ± 0.009) and increased slightly above this, where a higher proportion of winged or kite-shaped crystals appeared.

The vertical growth of the abraded-crowned, forest-tundra trees averaged 11.8 cm (SD ± 2.22), as compared to 13.2 cm (SD ± 1.28) for the full-crowned, open forest trees over the past 12 years. In total, the average height growth of the eight sample trees was 12.5 cm (SD ± 1.72), similar to other height increments measured in the area (Scott et al., 1987, 1988).

The three-year-old white spruce branchlets placed at 20 cm intervals above the snow surface for 10 days showed clear evidence of needle loss. Under average conditions the rate of needle loss approached 1% of the total branch weight per day in the first 40 cm above the snowpack (Fig. 1). The rate of weight loss declined to less than 10% of this value above 140 cm. Significant and ongoing needle loss was restricted to the first 80 cm above the snowpack. Under severe blizzard conditions even for a few hours, complete loss of needles from branches near the snow surface was observed.

The two-year-old white spruce branchlets placed 10 cm above the snow surface for 7 days lost from 4 to 7% of their initial weight. Although the weight loss due to needle damage was highly variable, some trends are evident (Fig. 2). Branchlets from the forest-tundra trees consistently lost more needles than branchlets from full-crowned trees. This difference is significant for branchlets from the seven-year-old branches (t = 3.75, p < .01), which were most resistant to needle loss. Greatest needle loss occurred in branchlets from three-year-old and eleven-year-old branches. This confirms the field observation that lower branches (older than ten years) often lose all of their needles in a single blizzard, while younger branches are more resistant (Fig. 3).

The wind speed around both islands of white spruce declined to approximately 90% of the reference value on the upwind side and increased by similar proportions on the
cross-wind sides (Fig. 4). When compared to the reference value, the wind speed in the woodland declined to 80% on the upwind and cross-wind sides. The wind velocities immediately downwind of the islands were approximately 40% of the ambient wind speed and only 12% downwind of the woodland tree. In all three studies the minimal wind
speed was located immediately behind the stem(s) and gradually increased with distance downwind. Typical of most trees in the area, the densely needled island under study and some of the woodland trees had a distinct jet of relatively high-speed air moving through channels within or around the tree structure.

At the sparsely needled island, the horizontal depth profile of accumulated snow began at approximately 80 cm depth, 2-3 m upwind of the structure, gradually increasing to 110 cm, 2-3 m downwind (Fig. 4). The resulting winter appearance of the island is that of an aggregation of dead, branchless and severely abraded stems protruding from a snowbank. Snow accumulation near the upwind side of the densely needled island was 75 cm, and it increased to 150 cm within the island. A wind-scoured trough approximately 40 cm wide and 25 cm deep occurs just downwind of the island, but beyond this, in the lee, the snow surface rises to a peak at 200 cm depth approximately 3 m downwind. The woodland tree was situated in approximately 85 cm of snow, with very little upwind or cross-wind redistribution evident. The area underneath the foliage and immediately downwind had less than half the snow depth. Approximately 3 m downwind was a snow drift above the ambient level shaped as a plume. The snow deposition in the open forest was more or less uniform, with subtle contours associated with ground features such as stumps.

Over the five years of study, the mid-February average snow depth in the open forest ranged from 49 cm in 1988 to 75 cm in 1989 (Table 1). During mid-February, 1992, when data were collected from all tree-line environments, approximately 20 cm of snow had accumulated on ice surfaces, while 40 cm had accumulated on the exposed forest tundra. A maximum depth of 420 cm was measured in the forest-tundra woodland and up to 58 cm had accumulated in the open forest. Brown Beckel (1957) reports similar snow depths at Churchill during the early 1950s.

To compare with the measured snowfall at the Churchill Weather Office, the open forest data were converted to snow-water equivalents. The average water equivalent of the exposed forest-tundra snow column was 29% and the powder snow in the open forest was 25%. These values are within the expected range based on measurement in other similar studies (Marchand, 1991; Timoney et al., 1992; Churchill Weather Office, Churchill, Manitoba) and are used here as an average of changing snow densities that will vary with temperature and season.

During early winter, the rate of snow accumulation in the open forest was approximately one and a half times that of the recorded snowfall (Fig. 5). As winter progressed, the rate of accumulation in the open forest increased, reaching three times the recorded snowfall during two of the three years studied.

**DISCUSSION**

Moving away from the snow surface, the increasing wind speed initially corresponds to a decrease in size and number of ice crystals. Frequent collisions with structures resistant to the airflow, such as spruce needles, creates the zone of abrasion, which is largely limited to the first 80 cm above the snow. Hollow and aerodynamically complex crystals are relatively sparse and dominate the upper portion of the abrasion zone and higher. The damage to leaf tissues on

![Graph](image)

**TABLE 1. Comparative snow depths in cm for Churchill, Manitoba, during mid-February**

<table>
<thead>
<tr>
<th>Year</th>
<th>Sea ice Avg</th>
<th>Sea ice SD</th>
<th>Lake ice Avg</th>
<th>Lake ice SD</th>
<th>Forest tundra Avg</th>
<th>Forest tundra SD</th>
<th>Cospe peak</th>
<th>Open forest Avg</th>
<th>Open forest SD</th>
</tr>
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<tr>
<td>1992</td>
<td>20.5</td>
<td>8.13</td>
<td>17.2</td>
<td>6.23</td>
<td>42.4</td>
<td>29.26</td>
<td>420.0</td>
<td>57.8</td>
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<td>-</td>
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<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>68.7</td>
<td>7.71</td>
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<td>1990</td>
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<td>-</td>
<td>40.7</td>
<td>24.24</td>
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<td>75.2</td>
<td>8.31</td>
</tr>
<tr>
<td>1988</td>
<td>-</td>
<td>-</td>
<td>12.6</td>
<td>6.16</td>
<td>18.0</td>
<td>18.96</td>
<td>226.0</td>
<td>49.9</td>
<td>6.71</td>
</tr>
</tbody>
</table>

**FIG. 5.** The snow accumulation in the open forest as a percentage of the reported snowfall at the Churchill Weather Office during monthly periods in the winters of 1988-89 (top), 1989-90 (middle) and 1990-91 (bottom). The snowfall and snow depth figures were initially converted to water equivalents.
impact would be greatest when the collision is most localized to the sharp edge of a complex crystal. Above the 80 cm level some needle removal occurs, but most commonly, cuticular abrasion that leads to desiccation during the growing season (Frey, 1983; Hadley and Smith, 1989) is prevalent. Cuticular damage was not quantified during the needle abrasion studies, although all of the trees were evidently already in a state of desiccation associated with the long, very cold winter period just before the study period (Scott and Hansell, 1992).

Needles from different locations throughout the tree have differences in susceptibility to damage by wind abrasion (Hadley and Smith, 1983). It was apparent during our study of white spruce branches in the zone of abrasion that needles of different age were of different size and susceptibility to being completely abraded away. Further, needles of the same age had differing resistance to removal, reflecting the history of the tree (crown form) and the age of the branch. Needles produced on new branches are initially susceptible to being abraded away. As the branches age over an 8-year period, each subsequent crop of needles becomes more resistant to being removed by abrasion. After 8 years, branches again become more susceptible to needle loss, and increasingly the seasonal crop of needles will be lost during the first winter. Consequently, to successfully grow through the abrasion zone, shoot growth should exceed 80 cm in 7 or 8 years.

In this study the trees that were successful at Churchill grew 80 cm in 6.4 years on average. This amount of growth is typical of measurements in other local studies as well (Scott et al., 1987, 1988). Once through the abrasion zone, new branches develop with needles largely remaining intact but are prone to cuticular damage from a lesser intensity of abrasion (Hadley and Smith, 1989). As lower branches become older, needle loss within 80 cm of the snowpack surface accelerates, leading to localized branch loss and stem damage. In fact, most abraded stems of the forest-tundra white spruce are reddish, as they have lost all dead bark on the upwind side, and resin blisters are common, indicating damage to the living phloem within the bark. The removal of needles and branches and bark loss apparently do not significantly impede wood production or height growth in the stem (Fig. 6).

In the extreme condition, low ground temperatures produce poor growth (Delucia, 1986; Scott et al., 1987; Day et al., 1991), resulting in a prolonged period in the most severe zone of abrasion and frequent loss of the terminal buds. The subsequent disorganization leads to even poorer growth, which often results in failure to erect a shoot through the abrasion zone. However, the krummholz tree islands on the forest tundra, mostly stemless masts and cushions (see Payette, 1974; Marr, 1977; Scott et al., 1987; Marchand, 1991; Lavoie and Payette, 1992), are least exposed to abrasion. These islands accumulate snow early in the season and remain buried throughout the winter. Snow infilling also characterizes the islands that are sparsely needled, as depicted in Figure 4 (top). The snow cover may reduce winter desiccation (Tranquillini, 1979; Hadley and Smith, 1987), but delayed snowmelt often results in low ground temperatures during the growing season (Brown Beckel, 1957; Rouse, 1984a).

The density of a wind barrier is directly proportional to the height of the plume (snow depth) and inversely proportional to its length (Oke, 1987:245). Densely needled islands reduce winter airflow and trap most snow in such a plume on the downwind side. The redistribution of snow appears to enhance establishment of shrubs such as Betula glandulosa, light-tolerant mosses such as Tomentypnum nitens and other snow-bed species. Areas downwind to slightly cross-wind of the islands produced jets where upwardly displaced air moves back down to the surface. On the upwind side, and where jets occur, lichen-heath develops. High-speed air jets develop in front of the cushion (or skirt) of the tree island and on the right-hand side (while viewing from upwind), which partially removes the snow. The cushions with the direct impact from the jets have the most densely needled branches, which often become encrusted with ice and snow.

Over long periods of time there may be a trend towards woodland development on the forest tundra. Marr (1977) suggests that high-speed jets create unfavourable conditions between tree islands, which would become more severe as the gap between them narrowed. This would prevent the islands from growing laterally, coalescing and forming woodlands. At Churchill, woodland formation appears to develop along ridges and in wind-sheltered sites, where spruce islands become grouped. The subsequent height increase from ongoing stem growth and decline in barrier density due to needle loss lengthen the area of snow accumulation downwind, eventually filling in between the islands. Within the woodland complex, high-speed wind jets form an asymmetric horseshoe-shaped vortex (Kind, 1981) around the base of some stems, clearing snow from beneath the raised cushions (Adams, 1981; Kind, 1981). Snow loading on the branches of most trees occurs frequently during quiet periods and in areas with the greatest shelter from the wind. During our study, the snow beneath the woodland study tree was moved out from under the crown and deposited in the branches on the downwind side (Fig. 4, bottom).
In the open forest, the air movement was minimal at the snow surface during periods of measurement and heavy snow load on the trees was common (e.g., Pruitt, 1978; Marchand, 1991). Snow was also absent under open forest trees at the edge of clearings. Such trees develop in association with frost heaving and subsequent palsa development, resulting in "drunken forests" (e.g., Marchand, 1991). Seppala (1986) created similar palsas, without trees, by removing snow from peatlands during the winter.

Snowfall is generally blown southward off Hudson Bay and adjacent lowlands, especially off the lake ice and exposed areas of the forest tundra. The stemless white spruce islands become snow covered early in the season, whereas the islands with stems collect shifting plumes of snow. During mid-winter the forest-tundra woodlands fill to a higher peak and earlier in the season. However during late winter in 1989-90, west winds resulted in snow blowing away from the open forest and the accumulation did not increase. In exposed areas that contribute snow, the distribution of snow is relatively uniform, although the net accumulation is less than the measured rate of snowfall. The distribution of snow is more variable across a gradient of spruce islands and woodlands approaching the tree line. The snow distribution is more uniform again within the open forest.

In spring, snowmelt on the exposed tundra usually occurs in late May (Brown Beckel, 1957; Rouse, 1982). The ground is frozen during snowmelt and most free water will readily run off or evaporate (Lafleur et al., 1992). Consequently, the contribution of snowmelt to ground water recharge and the subsequent availability for biological organisms is limited and can be highly variable (Marsh and Woo, 1985). Snowmelt in the open forest is usually three weeks later than on the exposed forest tundra and tundra. The larger quantities of water result in greater net percolation into the thawing peat (Rouse, 1982). Under near-average conditions, more heat would be required to warm the water immediately after snowmelt and the canopy might further reduce the availability of required latent heat of evaporation, contributing to a delay in warming of the forest (Rouse, 1984b; Scott et al., 1987; Odin and Degermark, 1990). In woodlands, snowmelt is usually completed by mid-July. Woodlands are characterized by snowbed communities commonly found on the downwind side, where the snow accumulation is greatest. Snowfall typically begins during mid-September.

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