## Sheffield <br> Hallam <br> University

# Interactions between canopy cover density and regeneration cores of older saplings in Scots pine (Pinus sylvestris L.) stands 

COBAN, S., COLAK, A.H. and ROTHERHAM, Ian [http://orcid.org/0000-0003-2903-5760](http://orcid.org/0000-0003-2903-5760)

Available from Sheffield Hallam University Research Archive (SHURA) at:
http://shura.shu.ac.uk/13937/

This document is the author deposited version. You are advised to consult the publisher's version if you wish to cite from it.

## Published version

COBAN, S., COLAK, A.H. and ROTHERHAM, Ian (2016). Interactions between canopy cover density and regeneration cores of older saplings in Scots pine (Pinus sylvestris L.) stands. Forest Systems, 25 (3), e073.

## Copyright and re-use policy

See http://shura.shu.ac.uk/information.html

# Interactions between canopy cover density and regeneration cores of older saplings in Scots pine (Pinus sylvestris L.) stands 

S. Coban ${ }^{1}$, A.H. Colak ${ }^{2}$, I.D. Rotherham ${ }^{3}$<br>${ }^{1-2}$ Istanbul University, Faculty of Forestry, Department of Silviculture, Turkey<br>${ }^{3}$ Sheffield Hallam University, Geography, Tourism \& Environment Research Unit, UK


#### Abstract

Aim of study: This paper provides an analysis of growth and survival of twenty-year-old Scots pine saplings in relation to canopy cover density (CCD) gradients, from dense (D-CCD), sparse (S-CCD), and gap (G-CCD) situations. Area of study: Aladag (Bolu) in northern Turkey. Material and methods: Sparse canopy cover density (S-CCD), dense canopy cover density (D-CCD) and gap canopy (G-CCD) were chosen within ten different strip sample plots ( $10 \times$ $50 \mathrm{~m})$ with sapling regeneration cores. Those regeneration cores were divided into two portions (individuals at the edge and middle of the regeneration cores) and from each portion three individuals was were obtained from a sample. The growth relationships of individual saplings were calculated with stem analyses. Honowski Light Factor (HLF) (ratio of Terminal sprout length (T) to Lateral sprout length (L)) was used to present growth potential measure of seedlings.


Main results: The largest sapling regeneration cores were found in the G-CCD followed by SCCD, and finally D-CCD, all tested for significance with Kruskal-Wallis Test. Compared with saplings in the middle of regeneration cores (crop saplings), those at the edge were always reduced in terms of mean height. Significant difference was only found between the 'Main Crop' and the 'Edge 1' of the regeneration cores for G-CCD suggesting that sapling regeneration cores are more typical under G-CCD conditions. HLF ratios were greater than 1 with high growth potentials for both CCD gradients (G-CCD and S-CCD) and there were no significant variations between G-CCD and S-CCD for main crop and edges. The thinning after 12-14 years increased sapling growth. However, under D-CCD, growth had virtually ceased.
Research highlights: Naturally occurring Scots pine saplings are suppressed by a dense canopy. However, they are tolerant of shade to the extent that they can survive over relatively long timeperiods (10-12 years) and can exploit subsequent opportunities should a canopy gap occur.
Keywords: Gap regeneration, sapling growth, light regime, canopy cover density, irregular silviculture

## 1 Received: Accepted:

2 Copyright © 2015 INIA. This is an open access article distributed under the Creative 3 Commons Attribution License (CC by 3.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

5 Funding: This study was supported by the Research Fund of Istanbul University (Project
6 Number: T-60/05122006).
7 Competing interests: The authors have declared that no competing interests exist.
8 Correspondence should be addressed to Suleyman Coban: scoban@ istanbul.edu.tr 9

## Introduction

In forest understoreys, tree seedling survival and growth are determined mainly by light, water and nutrient availability. Interspecific competition and herbivory may result from variation in canopy cover density (CCD) in the overstorey (Löf et al., 2007), and this may also affect growth and survival. Although forest ecologists and silviculturalists have emphasized the importance of creating canopy gaps (G-CCD) to generate spatial variation in order to promote tree regeneration, the effects of G-CCD on seedling recruitment may be offset by the development of dense forest understoreys. Light is a resource that limits tree seedling recruitment in many forest understoreys and G-CCD can raise light levels leading to increased seedling recruitment. However, many forests support dense understoreys that may compete with tree seedlings for resources such as light. This limits seedling recruitment even in gap conditions (G-CCD), and reduces the effectiveness of gaps in promoting seedling recruitment (Beckage et al., 2005; Ruuska et al., 2008). Understanding the behaviour of the seedlings of different tree species in relation to canopy shade is therefore important.

Studies of gap dynamics have contributed significantly to an understanding of the role of small-scale disturbance in forest ecosystems. Yet these have hardly been used by foresters for predicting tree responses to partial cutting (Coates, 2000). It is clear that interactions between heterogeneity in the forest overstorey (e.g. canopy gap or closed canopy) and understorey micro-environments may affect seedling performance. The presence of gapunderstorey interactions may influence both seedling competitive environments and the nature of resource limitation on seedling growth and survival. For example, understorey herbs, ferns, and shrubs may increase in response to high light availability in gaps (G-CCD) and so may compete with tree seedlings. Conversely, micro-environments characterized by high mineral nutrient availability or soil moisture may have disproportionate effects on seedling performance in high light environments (G-CCD), and little effect in light-limited environments (e.g. dense canopy (D-CCD)) (Beckage \& Clark, 2003).

Scots pine (Pinus sylvestris L.) is the most widely distributed pine species and one of the most important timber species in Eurasia. It has high commercial and ecological values (Oleksyn et al., 2002; Stanners \& Bourdeau, 1995; Figure 1). Natural Scots pine forests have a wide distribution in Turkey covering nearly 760,000 ha (Figure 1). There is an abundant literature on the factors affecting natural regeneration in Scots pine forests. Scots pine seed trees have an effect on the structure of pine seedlings (i.e. morphological characteristics), their spatial pattern, and their size distribution. Both height and seedling density decrease close to
the parent trees (Siipiletho, 2006). Competition from the mother trees inhibits development of saplings in close proximity (Montes \& Canellas, 2007). However, the growth of naturally occurring saplings in response to variations in CCD of Scots pine stands are poorly studied in the southern zone of its distribution area (Beckage et al., 2005; Coates, 2000; Löf et al., 2007; Pukkale et al., 1993; Cameron \& Ives, 1997; Andrzejczyk, 2007). Studies on regeneration and advance growth have shown that the effects of the long-term retention of seed trees has a strong negative impact on the development of young Scots pine stands, especially on relatively infertile sites in northern areas of its natural distribution (Ruuska et al., 2008). The research reported in this paper was designed to address four questions:
(1) Are different CCD gradients good predictors of regeneration cores of Scots pine saplings?
(2) How do CCD gradients affect the growth of Scots pine saplings?
(3) How was growth affected by the position of the sapling within the regeneration core in Scots pine stands?
(4) Do these responses vary with the shade tolerance rankings of Scots pine saplings?

## Materials and Methods

## Site description

Much of current knowledge of tree species in relation to canopy development is based on studies of trees occurring in naturally regenerated forest communities (Ellenberg, 1996). This research was therefore undertaken in naturally regenerated Scots pine forest in Aladag (Bolu) in northern Turkey (Figure 1: latitude between $40^{\circ} 30^{\prime}$ and $40^{\circ} 42^{\prime} \mathrm{N}$, longitude between $31^{\circ} 39^{\prime}$ and $31^{\circ} 52^{\prime} \mathrm{E}$ ) which is characterised by a high degree of naturalness (Colak et al., 2003). The research area is typically covered by 120-140 years-old-stands of Scots pine located at $1.380-1.420 \mathrm{~m}$ altitude. Silviculture in the area is based on natural regeneration following a shelterwood system and silvicultural interventions are not frequent at early stages of development (Coban, 2007).


Figure 1. Natural distribution range of Pinus sylvestris L. (EUFORGEN, 2009) and location of sample plots.

The climate shows Euxinian influences, with mean annual precipitation of 883 mm and mean annual temperature of $5.7^{\circ} \mathrm{C}$, cool winters, and sub-humid summers without significant droughts (Serin, 1998). The Euxinian region covers the whole of the EuroSiberian phyto-geographical region and is effectively referred to as the Euxinian province. This is an area that covers much of Georgia and the Caucasus, the Istiranca Mountains of European Turkey, and south-east Bulgaria (Davis, 1965-1988). The soils are mainly brown podzols (Tolunay, 1997), and the site quality class of for the research area is I (I-V: "I" shows the high and "V" the low site quality) (Forest Management Plan, 2005-2014; Tolunay, 1997).

## Selection of Sample Plots

By the end of 1987, shelterwood cutting areas in the 120-year-old Scots pine stands were at a uniform level of shade (dense: D-CCD) resulting from regular thinning and felling. There were 2- to 5 -year-old Scots pine seedlings here. After 10 years, in the second half of 1997, the CCD in these stands was assessed and placed into three broad gradients: (1) Dense (D-CCD), (2) Sparse (S-CCD), and (3) Gap (G-CCD) overstorey conditions (Figure 2). In
autumn 2007, samples were separated depending on typical sapling regeneration cores under different CCD within these stands. Sampling in the stands was conducted with a simple random sampling method. Measurements were taken for 10 rectangular ( $10 \times 50 \mathrm{~m}$ ) sample plots with different CCD gradients and chosen from natural regeneration cores of Scots pine saplings (Figure 2).


Figure 2. The CCD gradient models of Scots pine stands with three gradients i.e. dense (DCCD: Canopy cover is $55 \%$ ), sparse ( $\mathrm{S}-\mathrm{CCD}$ : Canopy cover is $43 \%$ ) and gap (G-CCD: gap size is $87 \mathrm{~m}^{2}$ ) (Coban, 2007).

## Gradients and measurements of CCD

CCD refers to the proportion of the forest floor covered by the vertical projection of the tree crowns (Figure 2). This is analogous to the use of the term 'cover' by ecologists and silviculturalists to refer to the proportion of the plan ground area occupied by the above ground parts of plants. Measures of CCD assess the presence or absence of canopy vertically above a sample of points across an area of forest. Tree height does not affect CCD, since only the
vertical projection of the crown is assessed. CCD is a measure reflecting the dominance of a site by trees or by particular species of tree (Jennings et al., 1999). The Scots pine stands in 1997 were allocated into 1 of 3 different CCDs (G-CCD, D-CCD and S-CCD) distinguished by CCD gradients in the shelterwood. These three CCD gradients (Ewald, 2007) are:
(1) Dense (D-CCD): CCD over 50\% (50-80\%; Percent canopy cover; Figure 2).
(2) Sparse (S-CCD): CCD up to $50 \%$ (20-50\%; Percent canopy cover; Figure 2).
(3) Canopy gap (G-CCD): no cover; the gap size $25-100 \mathrm{~m}^{2}$ (All sample plots areas in the canopy gaps were between 25.09 and $95.42 \mathrm{~m}^{2}$, Figure 2).

According to the definition by Jennings et al. (1999), if CCD is to be measured correctly, the measurements should be made in exact vertical direction (Korhonen et al., 2006). The following is the equation (Eq. 1) used to calculate the percentage of tree CCD (CCD-D and CCD-S) in the stand projection (Klumpp et al., 2002; Globe, 2005; Jennings et al., 1999; Figure 2):
$\operatorname{CCD}(\%)=\frac{\text { VPTC }}{\text { MA }} 100$
VPTC: Vertical projection of the tree crowns ( $\mathrm{m}^{2}$ )
MA: Forest floor cover $\left(\mathrm{m}^{2}\right)$ of measured area
G-CCD and S-CCD area calculated as a ration of the measured transect part by gap area (Figure 2).

Field procedures and calculations/equations
The standard alternative to CCD for the regeneration cores is by means of 'rectangular sample plots' (10-50 m) and shows longitudinal (profile) and vertical projection of the stand (Aksoy, 1978). For different CCDs, transects are taken from the strip plots. In this study ten transects from each S-CCD, D-CCD and G-CCD were chosen within ten different strip sample plots with sapling regeneration cores (Figure 3).

Figure 3. Longitudinal (profile) and vertical stand projection of all sample plots with different CCD gradients (D-CCD: Dense; S-CCD: Sparse; G-CCD: Gap) (Stand profiles from Coban, 2007).

The ages of individual saplings were assessed with stem sections cut in order to estimate the age by ring counting (González-Martínez \& Bravo, 2001). The saplings were scored considering the position of individuals in regeneration cores: (1) "Main crop sapling" the highest score in both variables (dominant and healthy) and (2) "Edge sapling" with the lowest score in both variables (dominant and healthy). The main crop saplings which were measured were located in the middle of the typical natural sapling regeneration cores (Figure 2). The term "main crop sapling", that is the trees selected to become a component of a future commercial harvest, refers to those saplings with the highest score in both variables (dominant and healthy) (Gonzales-Martinez \& Bravo, 2001). Those individuals at the edge of the natural sapling regeneration cores were selected as "Edge saplings" (Figure 2; G). The individuals were distinguished for stem-analysis as follows: Each core divided into three portions (Edge 1, Edge 2 and Main Crop). From each of the edges three individuals were taken (totalling six for edges) and three individuals were taken from the middle (Figure 2).

## Analysis of saplings

## Sapling-stem analysis

Sample saplings were cut down to ground level and stem cuts were taken at 1 m intervals for stem analysis (Atici, 2003, 1998; Kalipsiz, 1981). For the sapling-stem analysis of the increment and growth data of individual trees the "Computer Supported Statistical Analysis Program (GOVAN)" was used (Atici, 2003). GOVAN is computer software, which provides an opportunity to make computer-based stem analyses to determine the growth relationships of individual trees. Two different forms of graphs of absolute and relative age-height and stem models were drawn with this program. Correlation and regression analyses of the statistical model used for drawing the age-height graph were carried out (Atici, 2003).

## Honowski light factor (HLF ratio)

The ratio of Terminal sprout length (T) to Lateral sprout length (L) was presented by Fabjanowski et al. (1974) as the growth potential measure of seedlings and saplings under the canopy cover in coniferous species. The factor is referred to as the HLF ratio (Eq. 2). According to this value, the growth condition can be defined as 'well' or 'weak' (after Fabjanowski et al. 1974 from Schütz 2001). The individuals for the HLF ratio were selected from Edge 1, Edge 2 and Main Crop portions and from each part a mean set of data was obtained from a sample of three individuals.

1

HLF $=\frac{\mathrm{T}}{\mathrm{L}} \quad$ (Eq. 2; Schütz 2001)
Where: T: Terminal sprout length (cm); L: Lateral sprout length (cm)
HLF ratio: $1.0>$ growth well; 1.0-0.5 growth under the good; $0.5-0.25$ growth not good; 0.25 < growth very low.

## Data analysis

The following equation (Eq. 3) was used to calculate $95 \%$ confidence intervals of populations of all measured data (Atici et al., 2008; Kalipsiz, 1981; Sachs, 1972):
$\mu=\bar{x} \pm t S E_{\overline{\mathrm{x}}}$ (Eq. 3)
Where $\bar{x}$ is arithmetic mean; $S E_{\bar{x}}$ is std. error; $t$ is Student's t coefficient $\left(\mathrm{t}_{1-\alpha / 2 ;}\right.$ n-1 $)$; for 9 degrees of freedom $=2.262, \mathrm{n}$ is 10 number of samples.

Statistical evaluation including nonparametric test (Kruskal-Wallis H Test), t -tests, one-way variance analyses (ANOVA), and Student-Newman-Keuls (SNK) test were applied to the data collected using SPSS 5.01 software for Windows.

## Results

## The effect of CCD gradients on formation of regeneration cores

The results show that different CCD gradients result in major differences for sapling regeneration cores. The maximal sapling regeneration cores were found in the G-CCD, followed by S-CCD and finally D-CCD (Table 1). These differences were shown to be statistically significant by the Kruskal-Wallis H test. This test was applied to the difference in the CCD gradients of regeneration cores, and as a result two typical separate groups (1: DCCD; 2: S-CCD and G-CCD) were determined ( $\mathrm{P}<0.001$, Table 1). This situation was consistent in all sample plots with longitudinal (profile) and vertical projection of stands (Figure 3). Accordingly sapling regeneration cores do not occur in D-CCD.

Table 1．The effect of CCD gradients on formation of sapling regeneration cores．The data and statistical analysis from 30 sapling regeneration cores with different CCD gradients．This was confirmed by Kruskal－Wallis H test（Level：0：saplings without regeneration cores；1： saplings with sapling regeneration cores）

| Number of sample plots | Frequency distribution of natural sapling under different CCD gradients |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{gathered} \text { Dense } \\ (\mathrm{D}-\mathrm{CCD}) \end{gathered}$ |  | $\begin{gathered} \text { Sparse } \\ \text { (S-CCD) } \end{gathered}$ |  | Canopy gap(G-CCD) |  |
|  |  |  |  | Jِ |  | ס্ভ |
| 1 | D1 | 0 | S1 | 1 | G1 | 1 |
| 2 |  |  | S2 | 1 | G2 | 1 |
| 3 | D2 | 0 |  |  | G3 | 1 |
| 4 | D3 | 0 | S3 | 1 | G4 | 1 |
|  |  |  | S4 | 1 |  |  |
| 5 | D4 | 0 |  |  | G5 | 1 |
|  |  |  |  |  | G6 | 1 |
| 6 | D5 | 0 | S5 | 0 | G7 | 1 |
|  |  |  | S6 | 1 |  |  |
|  |  |  | S7 | 1 |  |  |
| 7 | D6 | 0 | S8 | 1 |  |  |
|  | D7 | 1 | S9 | 1 |  |  |
|  | D8 | 0 |  |  |  |  |
| 8 | D9 | 0 |  |  | G8 | 1 |
| 9 | D10 | 0 | S10 | 1 | G9 | 1 |
| 10 |  |  |  |  | G10 | 1 |
|  | Frequency distribution |  |  |  |  |  |
|  | D－CCD |  | S－CCD |  | $\mathrm{G}-\mathrm{CCD}$ |  |
| Level | 0 | 1 | 0 | 1 | 0 | 1 |
| Total | 9 | 1 | 1 | 9 | 0 | 10 |
|  | Descriptive Statistics |  |  |  |  |  |
|  | N |  | Mean |  | Std．Deviation |  |
|  | 30 |  | ． 6667 |  | ． 47946 |  |
|  | Test Statistics |  |  |  |  |  |
|  | Ch－Square |  | 21.170 |  |  |  |
|  | df |  | 2 |  |  |  |
|  | Asymp．Sig． |  | p＜0．001 |  |  |  |

## The properties of individuals in regeneration cores

Compared with saplings in the middle of a regeneration core or cluster，those on the edge were always shorter with $\mu$ value（Table 2 ）．These $\mu$ value differences were found for the S－CCD（Edge 1： $1.86 \pm 0.57 \mathrm{~m}$ ；Main Crop： $2.27 \pm 0.51 \mathrm{~m}$ ；Edge 2： $1.92 \pm 0.37 \mathrm{~m}$ ）and for the G－ CCD（Edge 1： $1.79 \pm 0.49 \mathrm{~m}$ ；Main Crop： $2.83 \pm 0.89 \mathrm{~m}$ ；Edge 2： $2.07 \pm 0.43 \mathrm{~m}$ ）．Because of height differentiation between edges and main crop the regeneration core form was determined（Table 2；Figure 3）．The distribution of saplings in different height classes in the sapling regeneration cores revealed they were shorter beneath the canopy than beyond the canopy（Figure 3）．These were statistically significant between the Main Crop and Edge 1 of the regeneration core for $\mathrm{G}-\mathrm{CCD}(\mathrm{t}=-2.317 ; \alpha=0.036)$ ，but not significant for $\mathrm{S}-\mathrm{CCD}(\mathrm{t}=-$

1 1.213; $\mathrm{P}=0.24$ ) (Table 2). This suggests that sapling regeneration cores were more typical 2 under G-CCD conditions than under S-CCD.

3 Table 2. The effect of CCD gradients on height of saplings in the sapling regeneration core. 4 Data and statistical analysis from twenty regeneration cores ( $n=10, v=9, t=2.262$ ), $\mu$ (Eq.3). 5 This was confirmed by Student's $t$-test ( $\alpha=0.05$ ): 95\% confidence interval for arithmetic mean.

| Number of sample plots | Height (m) |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Sparse (S-CCD) |  |  |  |  | Canopy gap (G-CCD) |  |  |  |  |
|  | Place in sample plots (Figure 3) |  | Edge | Main crop | Edge | Place sample p (Figure |  | Edge | Main crop | Edge |
| 1 | S1 |  | 3.1 | 2.8 | 1.9 | G1 |  | 2.0 | 3.8 | 2.1 |
| 2 | S2 |  | 2.4 | 3.0 | 1,8 | G2 |  | 2.0 | 4.6 | 1.8 |
| 3 | S3 |  | 2.2 | 2.8 | 2.1 | G3 |  | 3.6 | 5.2 | 3.7 |
| 4 | S4 |  | 1.6 | 2.5 | 1.7 | G4 |  | 1.6 | 2.5 | 1.7 |
| 5 | S5 |  | 1.4 | 2.4 | 2.0 | G5 |  | 1.5 | 2.1 | 1.8 |
| 6 | S6 |  | 0.9 | 0.9 | 2.2 | G6 |  | 1.7 | 1.9 | 2.0 |
| 7 | S7 |  | 0.8 | 1.4 | 0.7 | G7 |  | 1.4 | 2.4 | 2.2 |
| 8 | S8 |  | 2.2 | 2.1 | 1.9 | G8 |  | 1.1 | 2.1 | 1.6 |
| 9 | S9 |  | 2.8 | 3.0 | 2.7 | G9 |  | 1.5 | 1.9 | 1.9 |
| 10 | S10 |  | 1.2 | 1.8 | 2.2 | G10 |  | 1.5 | 1.8 | 1.9 |
| $\bar{x}$ | - |  | 1.86 | 2.27 | 1.92 | - |  | 1.79 | 2.83 | 2.07 |
| $s^{2}$ | - |  | 0.63 | 0.51 | 0.26 | - |  | 0.48 | 1.54 | 0.36 |
| $s$ | - |  | 0.80 | 0.71 | 0.51 | - |  | 0.69 | 1.24 | 0.60 |
| $S E_{\overline{\mathrm{x}}}$ | - |  | 0.25 | 0.23 | 0.16 | - |  | 0.22 | 0.39 | 0.19 |
| $n$ | - |  | 10 | 10 | 10 | - |  | 10 | 10 | 10 |
| $\mu$ | - |  | $\begin{gathered} 1.86 \\ \pm \\ 0.57 \\ \hline \end{gathered}$ | $\begin{gathered} 2.27 \\ \pm \\ 0.51 \\ \hline \end{gathered}$ | $\begin{gathered} 1.92 \\ \pm \\ 0.37 \\ \hline \end{gathered}$ | - |  | $\begin{gathered} 1.79 \\ \pm \\ 0.49 \\ \hline \end{gathered}$ | $\begin{gathered} 2.83 \\ \pm \\ 0.89 \\ \hline \end{gathered}$ | $\begin{gathered} 2.07 \\ \pm \\ 0.43 \\ \hline \end{gathered}$ |
| Independent Samples Test |  |  |  |  |  |  |  |  |  |  |
|  | Levene's Test for Equality of Variances |  |  |  | t-test for Equality of Means |  |  |  |  |  |
|  | F | Sig. | t | df | $\begin{gathered} \text { Sig. } \\ (2- \\ \text { tailed) } \end{gathered}$ | Mean Dif. | Std. <br> Error <br> Dif. | 95\% Confidence Interval of the Difference |  |  |
|  |  |  |  |  |  |  |  |  | er | Upper |
| $\begin{array}{ll} \hline & \text { Edge } 1 \\ \text { and } \\ \text { main crop } \end{array}$ | . 415 | . 527 | -1.213 | 18 | . $241{ }^{\text {NS }}$ | -. 41000 | . 33805 | -1.12022 |  | . 30022 |
| Edge 2 and main crop | 2.273 | . 149 | -1.261 | 18 | . $224{ }^{\text {NS }}$ | -. 35000 | . 27763 | -. 93328 |  | . 23328 |
| $\begin{array}{ll} \hline & \text { Edge } 1 \\ \text { and } \\ \text { main crop } \end{array}$ | 5.242 | . 034 | -2.317 | 14.089 | . 036 * | -1.04000 | . 44883 | -2.00 | 206 | -. 07794 |
| Edge 2 and main crop | 7.409 | . 014 | -1.744 | 12.996 | $.105^{\text {NS }}$ | -. 76000 | . 43566 | -1.7 | 122 | . 18122 |

$$
y
$$

$$
12
$$

One-way Variance Analyses (ANOVA) were carried out to test the differences in sapling age under the G-CCD and S-CCD between Main Crop saplings and both Edge saplings in the regeneration cores. There was no significant difference (G-CCD: F=1.891; p= $0.17, S-C C D ; F=1.122 ; p=0.340$ ) for sapling age (Table 3).

Table 3. The age distributions in the sapling regeneration cores. Data and statistical analysis from twenty regeneration cores $(\mathrm{n}=10, \mathrm{v}=9, \mathrm{t}=2.262), \mu$ (Eq. 3). This was confirmed by 7 Student's $t$-test $(\alpha=0.05)$ : Arithmetic mean of $95 \%$ confidence interval of total population


## Honowski light factor (HLF) in the regeneration cores

Table 4 shows that for both CCD gradients (G-CCD and S-CCD), as Edge 1, Edge 2 and, Main Crop, the HLF ratios were found to exceed 1 but the different groups varied in their values. One-way variance analyses were carried out to test the differences in the G-CCD and S-CCD between Main Crop saplings and both Edge saplings in the regeneration cores. As a result, the assessment established a significant difference for $\mathrm{G}-\mathrm{CCD}(\mathrm{F}=4.521 ; \mathrm{P}=0.02$ ) but not for S-CCD ( $\mathrm{F}=1.165$; $\mathrm{p}=0.327$ ). Student-Newman-Keuls (SNK) test was applied to the difference in the G-CCD and two typical separate groups were determined 1) Main Crop saplings and 2) Both Edge samples (Table 4). There were no significant variations between the G-CCD and S-CCD for HLF ratios for each zone in the regeneration core (Main Crop saplings: $\mathrm{t}=-1.458 ; \mathrm{P}=0.162$, Edge $1 ; \mathrm{t}=0.243 ; \mathrm{P}=0.811$ and Edge 2: $\mathrm{t}=-0.092 ; \mathrm{p}=0.928$ ) (Tables 4 and 5).

Table 4. Statistical analysis of difference for HLF ratios between Main Crop saplings and both Edge saplings in the regeneration core with different CCD gradients. This was confirmed by one-way variance analyses (ANOVA) and Student-Newman-Keuls (SNK)

| Number of sample plots | HLF ratios |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Sparse (S-CCD) |  |  |  | Canopy gap (G-CCD) |  |  |  |
|  | $\begin{gathered} \text { Place in } \\ \text { sample plots } \\ \text { (Figure 3) } \end{gathered}$ | Edge | Main crop | $\begin{gathered} \text { Edge } \\ 2 \end{gathered}$ | Place in sample plots <br> (Figure 3) | Edge $1$ | Main crop | $\begin{gathered} \text { Edge } \\ 2 \\ \hline \end{gathered}$ |
| 1 | S1 | 1.4 | 1.5 | 1.3 | G1 | 1.4 | 1.5 | 1.5 |
| 2 | S2 | 1.3 | 1.5 | 1.3 | G2 | 1.1 | 1.4 | 1.3 |
| 3 | S3 | 1.2 | 1.3 | 1.4 | G3 | 1.6 | 2.3 | 2.0 |
| 4 | S4 | 1.3 | 1.4 | 1.2 | G4 | 1.3 | 1.4 | 1.2 |
| 5 | S5 | 1.1 | 1.5 | 1.5 | G5 | 1.3 | 2.0 | 1.5 |
| 6 | S6 | 2.2 | 1.6 | 1.4 | G6 | 1.5 | 1.8 | 1.0 |
| 7 | S7 | 0.9 | 1.9 | 1.8 | G7 | 1.1 | 1.5 | 1.5 |
| 8 | S8 | 1.3 | 1.6 | 1.3 | G8 | 1.0 | 1.7 | 1.7 |
| 9 | S9 | 1.4 | 1.3 | 1.4 | G9 | 1.5 | 1.6 | 1.1 |
| 10 | S10 | 1.3 | 1.4 | 1.3 | G10 | 1.3 | 1.4 | 1.2 |
| $\bar{x}$ | - | 1.34 | 1.50 | 1.39 | - | 1.31 | 1.66 | 1.40 |
| s2 | - | 0.11 | 0.03 | 0.03 | - | 0.04 | 0.09 | 0.09 |
| S | - | 0.34 | 0.18 | 0.17 | - | 0.20 | 0.30 | 0.30 |
|  | - | 0.11 | 0.06 | 0.05 | - | 0.06 | 0.09 | 0.10 |
| $S E_{\overline{\mathrm{x}}}$ |  |  |  |  |  |  |  |  |
| n | - | 10 | 11 | 12 | - | 13 | 14 | 15 |
| $\mu$ | - | $\begin{gathered} 1.34 \\ \pm \\ 0.57 \\ \hline \end{gathered}$ | $\begin{gathered} 1.50 \\ \pm \\ 0.58 \end{gathered}$ | $\begin{gathered} 1.39 \\ \pm 0.59 \end{gathered}$ | - | $\begin{gathered} 1.31 \\ \pm \\ 0.60 \\ \hline \end{gathered}$ | $\begin{gathered} 1.66 \\ \pm \\ 0.61 \\ \hline \end{gathered}$ | $\begin{gathered} 1.40 \\ \pm \\ 0.62 \\ \hline \end{gathered}$ |
| ANOVA |  |  |  |  |  |  |  |  |
|  |  | Squares |  | MeanSquare |  | F | Sig. |  |
|  |  |  | 2 |  | . 067 | 1.165 | . 327 NS |  |
|  |  | 1.553 |  | 27 | . 058 |  |  |  |
| Total |  | 1.687 | 29 |  |  |  |  |  |
| Between | roups . 661 |  | 2 |  | . 330 | 4.521 | .020* |  |
|  | oups | 1.973 |  | 27 | . 073 |  |  |  |
|  |  | . 661 |  | 2 | . 330 | 4.521 | . 020 * |  |
|  | Student-Newman-Keuls Test (SNK) |  |  |  |  |  |  |  |


|  | Subset for alpha $=0.05$ |  |  |
| :--- | :---: | :---: | :---: |
|  | N | 1 | 2 |
| Edge 1 | 10 | 1.3100 |  |
| Edge 2 | 10 | 1.4000 | 1.6600 |
| Main crop | 10 |  | 1.000 |
| Sig. |  | .463 |  |

Table 5. Statistical analysis of HLF ratios between G-CCD and S-CCD for each cluster zone

| Independent Samples Test |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{aligned} & \text { HLF } \\ & \text { atios } \end{aligned}$ | Levene's Test for Equality of Variances |  | t-test for Equality of Means |  |  |  |  |  |  |
|  |  |  |  |  |  | Sig. <br> (2- | Mean | Std. <br> Error | 95\% Co Interv Diffe | fidence of the ence |
|  |  | F | Sig. | t | df | tailed) | Dif. | Dif. | Lower | Upper |
|  | Edge 1 | . 230 | . 637 | . 243 | 18 | $.811^{\mathrm{NS}}$ | . 03000 | . 12351 | -. 22949 | . 28949 |
|  | Edge 2 | 4.089 | . 058 | -. 092 | 18 | $.928{ }^{\text {NS }}$ | -. 01000 | . 10899 | -. 23897 | . 21897 |
|  | Main crop | 2.810 | . 111 | -1.458 | 18 | $.162^{\mathrm{NS}}$ | -. 16000 | . 10975 | -. 39057 | . 07057 |

The growth in the regeneration core after the second cutting (thinning and felling) of upper story

The results from examination of the age-height graph of sixty sapling-stems (stem analyses) and derived from correlation and regression analyses (e.g. Figure 4) show major differences for growth in height after the first and second cutting stages in the stand. It was found that high overstorey densities (D-CCD) slightly increased sapling growth (Figures 4 and 5), and lower overstorey densities (G-CCD and S-CCD) substantially increased sapling growth (Figures 4 and 5). D-CCD gradient exhibited reduced growth in height without mortality, but after the second cutting it was found that the saplings (10-14 years in age) grew very well without any obvious slowing or mortality. This indicates that for the time-periods considered (10-12 years), naturally occurring Scots pine saplings are shade-tolerant in that whilst growth is suppressed they do survive.


Figure 4. The examples of the age-height graph of sapling-stems (stem analyses) (Coban, 2007).


Figure 5. Height growth of saplings within regeneration cores under G-CCD and S-CCD (a) and DCCD (b) (only 20 saplings from G-CCD and S-CCD, 8 from D-CCD were shown in the graph).

## Discussion and Conclusions

There is an abundant literature on population structure and factors affecting natural regeneration in Scots pine in Europe (González-Martínez \& Bravo, 2001; Valkonen et al., 2002; Montes \& Canellas, 2007) and in Turkey (Pamay, 1962). However, the characteristics of naturally-occurring saplings of Scots pine under the different CCD gradients are poorly studied. Estimates of CCD are also becoming increasingly important in forest management
(Ganey \& Block, 1994; Korhonen et al., 2006) and the demand for natural landscapes, the multi-resource use of forests and the high cost of plantations all help to focus European foresters' attention on natural regeneration (González-Martínez \& Bravo, 2001). In this context long-term experiments to determine the interactions between different CCD gradients (G-CCD, D-CCD and S-CCD), the regeneration cores of natural Scots pine saplings (Figure 3 ), and the effects on sapling growth rates (Table 1-5) and more have been reported by Pamay (1962), Genc (2004) and Odabasi et al. (2004).

The practical application of this study requires determination of critical gradients below-CCD (D-CCD, S-CCD and G-CCD) for satisfactory sapling survival and growth of Scots pine. This must then be linked to observations of field light regimes. The studies have revealed a significant relationship between CCD gradients and regeneration core of saplings (P:0.000). Decreased canopy cover had a significant positive effect on sapling growth and this has been found by most studies in the past (Pamay, 1962; Genc, 2004; Odabasi et al., 2004; Valkonen, 2000).

The maximal regeneration core of sapling was found in the $\mathrm{G}-\mathrm{CCD}$, followed by S CCD and finally D-CCD (Table 1). Similar responses were found by Pukkala et al. (1993) with the correlation between the radiation parameters and Scots pine sapling growth significantly positive. The variation in height growth of Scots pine seedlings seems to be caused mostly by the spatial heterogeneity of the stand (i.e. different CCD gradients), which is consistent with some previous studies (Kuuluvainen et al., 1993). As stated by Tegelmark (1998), regeneration core of naturally-occurring Scots pine saplings is potentially important in future stand development and sapling properties change with the evolving stages of the stand. As with Beckage et al. (2005), G-CCD had only a slight positive effect on seedling survival, and the benefit was offset by a large negative effect of understorey shrubs. This study also found, like Pamay (1962) that high overstorey densities (D-CCD) slightly increased sapling growth (Figures 4 and 5). Lower overstorey densities (G-CCD and S-CCD) substantially increased sapling growth (Figures 4 and 5). Other key factors were intraspecific competition (González-Martínez and Bravo, 2001; Kuuluvainen and Juntunen, 1998) between saplings placed differently in the regeneration cores (Pamay, 1962), and root competition with mature trees (Valkonen, 2000; Siipilehto, 2006; Montes \& Canellas, 2007). Compared with saplings in the middle of regeneration cores (crop sapling), those on the edge were shorter with $\mu$ value ( $\alpha: 0.05$; Eq. 3). This also highlights root competition effects and CDD gradients (Valkonen, 2000; Siipilehto, 2006). Examination of the positions of previously removed trees
indicated that root system shape and extent resulted from past competition prior to regeneration works (Valkonen, 2000).

The ideas of shade tolerance suggest that there are species-specific physiological and growth adaptations which influence the ability to survive and grow at different levels of light. For example, in low light, shade-tolerant Abies species exhibit reduced height and diameter growth without mortality, but this is not true for pine species (Kobe \& Coates, 1997; Mason et al., 2004). Scots pine is a typical shade-intolerant pioneer (Coates \& Burton, 1999; Chantal et al., 2003; Ewald, 2007) for which regeneration is practically restricted to open, non-forest vegetation (Ewald, 2007). Its behaviour in native pinewoods in Scotland certainly reflects this. While the broad classification of species as 'shade tolerant', 'intermediate', or 'light demanding' appear to be consistent between regions (Mason et al., 2004). However, the behaviour is not totally fixed and shade tolerance within species may be affected by site quality (Carter \& Klinka, 1992). Consequently, the magnitude of the competition effect may vary between geographical areas along with differences in site productivity. However, there is little published research available to evaluate or quantify this hypothesis (Valkonen, 2000). Sapling establishment and development continues out of the dense groups of the younger cohort, under the protection of the low density groups of remaining mother trees. This semishade tolerant behaviour found in the southern distribution of Scots pine, i.e. the Sistema Central range, the Iberian Mountain Range and other enclaves in Spain, is quite different from the poor shade tolerance shown by the species in the rest of its distributional area (Montes \& Canellas, 2007). Although Scots pine is generally considered a shade intolerant species (Chantal et al., 2003), with increasing site quality it can survive for long periods under a dense forest canopy (Odabasi et al., 2004). Species-specific growth responses show little difference under high available light conditions, but performance at low light levels is generally consistent with shade tolerance rankings in the literature. The exception was that Scots pine shade intolerance was higher [don't you mean lower?? i.e. more shade tolerant less intolerant??] than expected (Claveau et al., 2002). The results of stem sampling and correlation and regression analyses, age-height graph and age-periodical height increment graph evaluations showed naturally-occurring sapling of Scots pine in the study area were shade tolerant (Figures 4, 5). Some previous studies suggest that Scots pine saplings cannot survive long under a dense forest canopy (Ata, 1995; Genc, 2004). However, as found in this study and earlier investigations (e.g. Pamay, 1962), Scots pine saplings can survive 20-25 years under dense forest canopy (Figure 3). According to Pamay (1962), this period may be up to 45-60 years in the case of less dense clustering. Pamay described this situation as the
"semi-shade type" of Scots pine. This is important since a more detailed understanding of species response to different light levels can help develop appropriate silvicultural prescriptions to promote varied forest structures with improved species diversity. Linked to other decision-making tools this can help inform the potential impacts of different stand management regimes (Mason et al., 2004).

Recent studies of shade tolerance have examined the relationships between mortality and growth in varying light conditions (Kobe et al., 1995; Kobe and Coates, 1997; Wyckoff \& Clark, 2002; Kunstler et al., 2005; Löf et al., 2007). In these studies, the interactions between CCD gradients and Scots pine sapling regeneration cores was on the basis of tradeoffs between the ability to survive at $\mathrm{D}-\mathrm{CCD}$ gradients and to achieve a high growth rate at G-CCD and then S-CCD. Edge 1, Edge 2 and, Main Crop HLF ratios ( $\alpha: 0.05$ ) were found to be more than 1 for both CCD gradients ( $\mathrm{G}-\mathrm{CCD}$ and $\mathrm{S}-\mathrm{CCD}$ ). According to this value, the growth potential can be defined as high (Eq. 2; After Fabjanowski, 1974 from Schütz, 2001). The results of HLF ratio assessments and stem analyses drawn by correlation and regression analyses, age-height graphs and age-periodical height increment graphs show the growth of sapling regeneration cores to be affected by CCD gradients. It was found that growth continued rapidly in CCD gradients at G-CCD and S-CCD; a response to thinning after 1214 years suppression by parent trees. In published research it has been suggested that older suppressed saplings were degenerated individuals which under a dense forest canopy lost their vigour (Ata, 1995). However, as this study indicates, these older saplings retain their growth potential during the time of suppression and can recover when the opportunity arises. Vaat and Vildo (2005) concluded that for Scots pine such management intervention with thinning and opening up the canopy needed to be within the first six years and stand densities radically reduced (recommended to be to the to the minimum values allowed by forest legislation or guidance). High-density stands will be unsuitable for shelterwood cutting due to shorter crowns and a higher risk of windfall after repeated overstorey removals. This research found sapling survive for $10-14$ years under a dense overstorey ( $\mathrm{D}-\mathrm{CCD}$ ) without mortality and with growth at a standstill (Figures 4, 5 and 6; Section A). But after the second cutting, lower overstorey densities (G-CCD and $\mathrm{S}-\mathrm{CCD}$ ) released saplings (10-14 years) to growth well and without mortality (Figures 4, 5 and 6).

It is suggested that key elements to the interpretation of this situation are the local differences and distinctiveness of landscapes, together with variations in forest product extraction and management. This finding relates to the idea that application of 'close-tonature' silviculture in Turkey could significantly reduce the problems facing Turkish forests
today. However, it will take time and requires a change from current practices. The application of similar management regimes for all forest zones regardless of stand properties is not sustainable. There is a wealth of good practice and evidence from case studies in Europe that can help inform the future management of this unique resource. In the United Kingdom and Germany, and in mountain regions of Italy and Austria, for example, there are many situations where sustainable forest management is increasingly moving towards 'close-to-nature' silviculture. This is generally incorporated into development plans that help sustain local communities through jobs and economic regeneration; the forest seen as a key to success. In particular, the concept of multi-functional forest management, including timber and wood production, sustainable tourism and leisure, wildlife, heritage and forest culture (with local food and drink), begins to provide a potential framework for long-term remediation (Çolak \& Rotherham, 2006). To conclude, this study supports the point of view that one of the most important rules of close-to-nature silviculture is the protection and generation of irregular stand structures (multi-layer stand, uneven-aged stands etc.). According to the findings of this study, the stands of parent Scots pine and of young-growth stands (old saplings) may occur together under S-CCD and G-CCD gradients. This is particularly the case where site quality is high. This study concluded that stands of young Scots pine can persist under the shelter of the parental canopy. With this information the practice of suitable forest management can be directed to the protection and maintenance of necessary conditions for sustainability. When developing silvicultural systems for Scots pine forests that would produce structural and compositional features as found in natural forests, there must be a better understanding of the role of microhabitats in regeneration dynamics (Kuuluvainen \& Juntunen, 1998).

## Acknowledgements

The authors wish to thank Dr Simay Kirca (Turkey) and Dr Emrah Özdemir for their contributions in preparing the manuscript.

## References

Aksoy H., 1978. Untersuchungen über Waldgesellschaften und Ihre waldbauliche Eigenschaften im Versuchswald Büyükdüz bei Karabük. Istanbul Üniversitesi Orman Fakültesi Yayinlari, 2332 (237), Istanbul. (In: Turkish).

Andrzejczyk T., 2007. Response of Scots pine (Pinus sylvestris L.) young-growth stands to an overstorey canopy in the postoutbreak stands in Pila Forests. Sylwan 151(1): 20-29. Ata C., 1995. Silvikültür Tekniği. Zonguldak Karaelmas Üniversitesi Yayinlari 4(3), Bartin. (In: Turkish).
Atici E., 1998. Volume Table of Oriental Beech (Fagus orientalis Lipsky) and the Comparison of Present Situation. Journal of Poplar and Fast Growing Forest Trees Research Institute, 1(25): 23-48.

Atici E., 2003. Computer supported statistical analysis (GOVAN) of the increment and growth data of individual trees. Review of the Faculty of Forestry, Istanbul University, 53(2): 37-55. (In: Turkish).

Atici E., Colak A.H., Rotherham I.D., 2008. Coarse Dead Wood Volume of Managed Oriental Beech (Fagus orientalis Lipsky) Stands in Turkey. Investigación Agraria: Sistemas y Recursos Forestales 2008 17(3): 216-227.

Beckage B., Clark S.J., 2003. Seedling Survival and Growth of Three Forest Tree Species: The Role of Spatial Heterogeneity. The Ecological Society of America.

Beckage B., Lavine M., Clark J.S., 2005. Survival of tree seedlings across space and time: estimates from long-term count data. Journal of Ecology, 93: 1177-1184.

Cameron A.D., Ives J.D., 1997. Use of hemispherical photography techniques to determine the association between canopy openness and regeneration of Scots pine (Pinus sylvestris L.) and downy birch (Betula pubescens Ehrh.) in Ballochbuie native pinewood, north-east Scotland. Scottish Forestry. 51(3): 144-149.

Carter RE, Klinka K, 1992. Variation in shade tolerance of Douglas fir, western hemlock and western red cedar in coastal British Columbia. Forest Ecology and Management, 55: 87-105.

Chantal M., Leinonen K., Kuuluvainen T., Cescatti A., 2003. Early response of Pinus sylvestris and Picea abies seedlings to an experimental canopy gap in a boreal spruce forest. Forest Ecology and Management, 176 (1-3): 321-336.

Claveau Y., Messier C., Comeau P.G., Coates K.D., 2002. Growth and crown morphological responses of boreal conifer seedlings and saplings with contrasting shade tolerance to a gradient of light and height. Canadian Journal of Forest Research, 32: 458-468.

Coates K.D., 2000. Conifer seedling response to northern temperate forest gaps. Forest Ecology and Management, 127: 249-269.

Coates K.D., Burton P.J., 1999. Growth of planted tree seedlings in response to ambient light levels in northwestern interior cedar-hemlock forests of British Columbia. Canadian Journal of Forest Research 29(9):1374-1382.
Coban S., 2007. Research on the natural generation samples of Scots pine (Pinus sylvestris L.) stands in Bolu-Aladag. Master Thesis, Istanbul University, Graduate School of Natural and Applied Sciences, Istanbul. (In: Turkish).
Colak A.H., Calikoglu M., Rotherham I.D., 2003. Combining Naturalness Concepts with Close-to-Nature Silviculture. Forstwissenschaftliches Centralblatt, 122: 421-431.
Colak AH, Rotherham ID, 2006. A review of the Forest Vegetation of Turkey: its status past and present and its future conservation. Royal Irish Academy, Journal of Biology and the Environment, 106(3): 343-355.
Davis P.H., 1965-1988. Flora of Turkey and the East Aegean Islands. Edinburgh. Edinburgh University Press.
Ellenberg H., 1996. Vegetation Mitteleuropas mit den Alpen in ökologischer, dynamischer und historischer Sicht, 5. Auflage, Stuttgart, Ulmer.
EUFORGEN. 2009. Distribution map of Scots pine (Pinus sylvestris). <www.euforgen.org> Ewald J., 2007. Ein pflanzensoziologisches Modell der Schattentoleranz von Baumarten in den Bayerischen Alpen A Phytosociological Model of Shade Tolerance of Tree Species in the Bavarian Alps. Forum geobotanicum, 3:11-19.
Fabjanowski, J., Jaworski, A., Musiel, W., 1974: The use of certain morphological features of the fir (Abies alba Mill.) and spruce (Picea abies Link.) in the evaluation of the light requirements and quality of their up-growth. Orig. Poln. Acta agraria et silvestris, Series silvestris (Warschau \& Krakau). 14: 3-29.
Ganey J.L., Block, W.M., 1994. A Comparison of Two Techniques for Measuring Canopy Cover. Western Journal of Applied Forestry, 9(1): 21-23.
Genc M., 2004. Silvikültürün Temel Esaslari. Süleyman Demirel Üniversitesi Yayinlari No: 44, SDÜ Basımevi, Isparta. (In: Turkish).
Globe, 2005. Canopy Cover and Ground Cover. Field Guide. Land Cover/Biology. Biometry Protocol, 9(11): 1-3.
González-Martínez S.C., Bravo F., 2001. Density and population structure of the natural regeneration of Scots pine (Pinus sylvestris L.) in the High Ebro Basin (Northern Spain). Ann. For. Sci., 58: 277-288.
Jennings S.B., Brown N.D., Sheil D., 1999. Assessing forest canopies and understorey illumination: canopy cover, canopy cover. Forestry, 72: 59-74.

Kalipsiz A., 1981. Statistical Methods. Istanbul University, publication number: 2837/294, Istanbul. (In: Turkish).
Kalipsiz A., 1999. Dendrometri, Istanbul Üniversitesi Orman Fakültesi Yayinlari, 3194/354, Istanbul. (In: Turkish).
Klumpp R.H., Colak A.H., Pitterle A., 2002. "Waldbauliches Glossar", in Waldbauliche Chancen und Probleme in Österreich, ed. J. Spörk und H. Vacik, eine CD für die forstliche Praxis und universitaere Lehre, Österreichischer Agrarverlag,Wien.

Kobe R.K., Coates K.D., 1997. Models of sapling mortality as a function of growth to characterize interspecific variation in shade tolerance of eight tree species of northwestern British Columbia. Canadian Journal of Forest Research, 27: 227-236.

Kobe R.K., Pacala S.W., Silander J.A., Canham C.D., 1995. Juvenile tree survivorship as a component of shade tolerance. Ecol. Appl., 5: 517-532.
Korhonen L., Korhonen K.T., Rautiainen M., Stenberg P., 2006. Estimation of forest canopy cover: a comparison of field measurement techniques. Silva Fennica 40(4): 577-588.

Kuuluvainen T., Juntunen P., 1998. Seedling establishment in relation to microhabitat variation in a windthrow gap in a boreal Pinus sylvestris forest. Journal Of Vegetation Science, 9(4): 551-562.

Kuuluvainen T., Hokkanen T.J., Järvinen E., Pukkala T., 1993. Factors related to seedling growth in a boreal Scots pine stand: a spatial analysis of a vegetation-soil system, Can. J. For. Res. 23: 2101-2109.

Kunstler G., Curt T., Bouchaud M., Lepart J., 2005. Growth, mortality, and morphological response of European beech and downy oak along a light gradient in sub-Mediterranean forest. Canadian Journal of Forest Research, 35: 1657-1668.
Löf M, Karlsson M, Sonesson K, Welander TN, Collet C, 2007. Growth and mortality in underplanted tree seedlings in response to variations in canopy cover of Norway spruce stands. Forestry, 80(4): 371-383.

Mason W.L., Edwards C., Hale S.E., 2004. Survival and early seedling growth of conifers with different shade tolerance in a Sitka spruce spacing trial and relationship to understorey light climate. Silva Fennica, 38(4): 357-370.
Montes F., Canellas I., 2007.The spatial relationships between post-crop remaining trees and the establishment of saplings in Pinus sylvestris stands in Spain. Applied Vegetation Science, 10: 151-160.

Odabasi T., Caliskan A., Bozkus H.F., 2004. Silvikültür Tekniği. Istanbul Üniversitesi Orman Fakültesi, I.Ü. Yayın no: 4459, Istanbul. (In: Turkish).

Oleksyn J., Reich P.B., Zytkowiak R., Karolewski P., Tjoelker M.G., 2002. Needle nutrients in geographically diverse Pinus sylvestris L. populations. Ann. For. Sci. 59: 1-18.
Pamay B., 1962. Türkiye'de Sarıçam (Pinus sylvestris L.) in tabi Gençleşmesi Imkanları Üzerine Araştırmalar, Orman Genel Müdürlüğü Yayınları No: 337/31, Istanbul. (In: Turkish).
Pukkala T., Kuuluvainen T., Stenberg P., 1993. Below-Canopy Distribution Of Photosynthetically Active Radiation and its Relation to Seedling Growth In A Boreal Pinus sylvestris Stand - A Simulation Approach. Scandinavian Journal Of Forest Research, 8(3): 313-325.
Ruuska J., Siipılehto, J., Valkonen, S., 2008. Effect of edge stands on the development of young Pinus sylvestris stands in southern Finland. Scandinavian Journal of Forest Research, 23(3), 214-226.

Sachs L., 1972. Statistical Methods (Statistiche Auswerttungsmethoden). Springer-Verlag, Berlin, Heidelberg, New York.

Schütz J-PH, 2001. Die Technik der Waldverjüngung von Wäldern mit Ablösung der Generationen. Skript zur Vorlesung Waldbau. ETH-Zentrum, Zürich.
Serin, M., 1998. Climatical data for 21 years of Bolu-Serif Yuksel Research Forest Meteorological station. Batı Karadeniz Ormancılık Araştırma Enstitüsü Dergisi. Seri no 1.

Siipilethto J, 2006. Height Distributions of Scots Pine Sapling Stands Affected by Retained Tree and Edge Stand Competition. Silva Fennica, 40(3): 473-486.

Stanners D., Bourdeau P., 1995. Europe's Environment-The Dobris Assessment Report. European Environment Agency, Copenhagen.

Tegelmark D.O., 1998. Site factors as multivariate predictors of the success of natural regeneration in Scots pine forests. Forest Ecology and Management, 109: 231-239.
Tolunay D., 1997. Aladag'da (Bolu) Sıklık Cağındaki Sarıcam (Pinus sylvestris L.) Mescerelerinde Bakimlarin Madde Dolasimina Etkileri. Doktora Tezi, Istanbul. (In: Turkish).

Vaat T, Vildo M, 2005. Shelterwood cutting in Scots pine ( $P$. sylvestris L.) stands. Transactions of the Faculty of Forestry, Estonian Agricultural University Issue, 38: 5364.

Valkonen S., 2000. Effects of retained Scots pine trees on regeneration, growth, form and yield of forest stands. Invest. Agraria fuera ser., 1:121-146.

Valkonen S., Ruuska J., Supılehto J., 2002. Effect of retained trees on the development of young Scots pine stands in Southern Finland. Forest Ecology and Management, 166: 227-243.

Wyckoff P.H., Clark J.S., 2002. The relationship between growth and mortality for seven cooccurring tree species in the southern Appalachian Mountains. J. Ecol., 90: 604-615.


Figure 1. Natural distribution range of Pinus sylvestris L. (EUFORGEN, 2009) and location of sample plots.


Figure 2. The CCD gradient models of Scots pine stands with three gradients i.e. dense (D3 CCD: 55\%), sparse (S-CCD: 43\%) and gap (G-CCD: $87 \mathrm{~m}^{2}$ ) (Coban, 2007).

Figure 3. Longitudinal (profile) and vertical stand projection of all sample plots with different CCD gradients (D-CCD: Dense; S-CCD: Sparse; G-CCD: Gap) (Stand profiles from Coban, 2007).


Figure 4. The examples of the age-height graph of sapling-stems (stem analyses) (Coban, 2007).


Figure 5. Height growth of saplings within regeneration cores under G-CCD and S-CCD (a) and DCCD (b) (only 20 saplings from G-CCD and S-CCD, 8 from D-CCD were shown in the graph).

Table 1. The effect of CCD gradients on formation of sapling regeneration cores. The data and statistical analysis from 30 sapling regeneration cores with different CCD gradients. This was confirmed by Kruskal-Wallis H test (Level: 0: saplings without regeneration cores; 1: saplings with sapling regeneration cores)

|  | Frequency distribution of natural sapling under |  |  |
| :---: | :---: | :---: | :---: |
|  |  |  |  |


|  |  | J |  | J |  | ָ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | D1 | 0 | S1 | 1 | G1 | 1 |
| 2 |  |  | S2 | 1 | G2 | 1 |
| 3 | D2 | 0 |  |  | G3 | 1 |
| 4 | D3 | 0 | S3 | 1 | G4 | 1 |
|  |  |  | S4 | 1 |  |  |
| 5 | D4 | 0 |  |  | G5 | 1 |
|  |  |  |  |  | G6 | 1 |
| 6 | D5 | 0 | S5 | 0 | G7 | 1 |
|  |  |  | S6 | 1 |  |  |
|  |  |  | S7 | 1 |  |  |
| 7 | D6 | 0 | S8 | 1 |  |  |
|  | D7 | 1 | S9 | 1 |  |  |
|  | D8 | 0 |  |  |  |  |
| 8 | D9 | 0 |  |  | G8 | 1 |
| 9 | D10 | 0 | S10 | 1 | G9 | 1 |
| 10 |  |  |  |  | G10 | 1 |
|  | Frequency distribution |  |  |  |  |  |
|  | D-CCD |  | S-CCD |  | G-CCD |  |
| Level | 0 | 1 | 0 | 1 | 0 | 1 |
| Total | 9 | 1 | 1 | 9 | 0 | 10 |
|  | Descriptive Statistics |  |  |  |  |  |
|  | N |  | Mean |  | Std. Deviation |  |
|  | 30 |  | . 6667 |  | . 47946 |  |
|  | Test Statistics |  |  |  |  |  |
|  | Ch-Square |  | 21.170 |  |  |  |
|  | df |  | 2 |  |  |  |
|  | Asymp. Sig. |  | $\mathrm{p}<0.001$ |  |  |  |

2 Table 2. The effect of CCD gradients on height of saplings in the sapling regeneration core.
3 Data and statistical analysis from twenty regeneration cores ( $\mathrm{n}=10, \mathrm{v}=9, \mathrm{t}=2.262$ ), $\mu$ (Eq.3).
4 This was confirmed by Student's t-test ( $\alpha=0.05$ ): Arithmetic mean of $95 \%$ confidence 5 interval of total population

| Number of sample plots | Height (m) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Sparse (S-CCD) |  |  |  | Canopy gap (G-CCD) |  |  |  |
|  | Place in sample plots (Figure 3) | $\begin{gathered} \text { Edge } \\ 1 \\ \hline \end{gathered}$ | $\begin{aligned} & \text { Main } \\ & \text { crop } \end{aligned}$ | $\begin{gathered} \text { Edge } \\ 2 \\ \hline \end{gathered}$ | Place in sample plots (Figure 3) | $\begin{gathered} \text { Edge } \\ 1 \\ \hline \end{gathered}$ | $\begin{aligned} & \text { Main } \\ & \text { crop } \end{aligned}$ | $\begin{gathered} \text { Edge } \\ 2 \\ \hline \end{gathered}$ |
| 1 | S1 | 3.1 | 2.8 | 1.9 | G1 | 2.0 | 3.8 | 2.1 |
| 2 | S2 | 2.4 | 3.0 | 1,8 | G2 | 2.0 | 4.6 | 1.8 |
| 3 | S3 | 2.2 | 2.8 | 2.1 | G3 | 3.6 | 5.2 | 3.7 |
| 4 | S4 | 1.6 | 2.5 | 1.7 | G4 | 1.6 | 2.5 | 1.7 |
| 5 | S5 | 1.4 | 2.4 | 2.0 | G5 | 1.5 | 2.1 | 1.8 |
| 6 | S6 | 0.9 | 0.9 | 2.2 | G6 | 1.7 | 1.9 | 2.0 |
| 7 | S7 | 0.8 | 1.4 | 0.7 | G7 | 1.4 | 2.4 | 2.2 |
| 8 | S8 | 2.2 | 2.1 | 1.9 | G8 | 1.1 | 2.1 | 1.6 |
| 9 | S9 | 2.8 | 3.0 | 2.7 | G9 | 1.5 | 1.9 | 1.9 |
| 10 | S10 | 1.2 | 1.8 | 2.2 | G10 | 1.5 | 1.8 | 1.9 |
| $\bar{x}$ | - | 1.86 | 2.27 | 1.92 | - | 1.79 | 2.83 | 2.07 |
| $s^{2}$ | - | 0.63 | 0.51 | 0.26 | - | 0.48 | 1.54 | 0.36 |
| $s$ | - | 0.80 | 0.71 | 0.51 | - | 0.69 | 1.24 | 0.60 |
| $S E_{\overline{\mathrm{x}}}$ | - | 0.25 | 0.23 | 0.16 | - | 0.22 | 0.39 | 0.19 |
| $n$ | - | 10 | 10 | 10 | - | 10 | 10 | 10 |
|  |  | 1.86 | 2.27 | 1.92 |  | 1.79 | 2.83 | 2.07 |
| $\mu$ | - | $\pm$ | $\pm$ | $\pm$ | - | $\pm$ | $\pm$ | $\pm$ |
|  |  | 0.57 | 0.51 | 0.37 |  | 0.49 | 0.89 | 0.43 |
| Independent Samples Test |  |  |  |  |  |  |  |  |

$\left.\begin{array}{lllllllllll}\hline & & \begin{array}{c}\text { Levene's } \\ \text { Test for } \\ \text { Equality of } \\ \text { Variances }\end{array} & & & & & \text { t-test for Equality of Means }\end{array}\right]$

1 Table 3. The age distributions in the sapling regeneration cores. Data and statistical analysis from twenty regeneration cores $(\mathrm{n}=10, \mathrm{v}=9, \mathrm{t}=2.262), \mu(\mathrm{Eq} .3)$. This was confirmed by Student's $t$-test $(\alpha=0.05)$ : Arithmetic mean of $95 \%$ confidence interval of total population

|  | Age (year) |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Sparse (S-CCD) |  |  |  | Canopy gap (G-CCD) |  |  |  |
|  | Place in sample plots (Figure 3) | Edge 1 | Main crop | $\begin{gathered} \text { Edge } \\ 2 \end{gathered}$ | Place in sample plots (Figure 3) | Edge 1 | Main crop | Edge 2 |
| 1 | S1 | 22 | 22 | 18 | G1 | 18 | 21 | 18 |
| 2 | S2 | 16 | 20 | 13 | G2 | 15 | 21 | 16 |
| 3 | S3 | 21 | 22 | 19 | G3 | 23 | 32 | 19 |
| 4 | S4 | 19 | 22 | 19 | G4 | 19 | 22 | 20 |
| 5 | S5 | 15 | 17 | 18 | G5 | 17 | 17 | 17 |
| 6 | S6 | 19 | 16 | 18 | G6 | 16 | 19 | 20 |
| 7 | S7 | 12 | 16 | 16 | G7 | 15 | 19 | 20 |
| 8 | S8 | 19 | 20 | 19 | G8 | 14 | 16 | 16 |
| 9 | S9 | 20 | 22 | 22 | G9 | 20 | 19 | 16 |
| 10 | S10 | 20 | 21 | 20 | G10 | 18 | 16 | 19 |
| $x$ | - | 18.30 | 19.80 | 18.20 | - | 17.50 | 20.20 | 18.10 |
| $s^{2}$ | - | 9.34 | 6.40 | 5.73 | - | 7.39 | 21.51 | 2.99 |
| $s$ | - | 3.06 | 2.53 | 2.39 | - | 2.72 | 4.64 | 1.73 |
| $S E_{\overline{\mathrm{x}}}$ | - | 0.97 | 0.80 | 0.76 | - | 0.86 | 1.47 | 0.55 |
| $n$ | - | 10 | 10 | 10 | - | 10 | 10 | 10 |
| $\mu$ | $\begin{gathered} 18.30 \\ \pm \\ 2.19 \end{gathered}$ |  | $\begin{gathered} 19.80 \\ \pm \\ 1.81 \end{gathered}$ | $\begin{gathered} 18.20 \\ \pm \\ 1.71 \\ \hline \end{gathered}$ | - | $\begin{gathered} 17.50 \pm \\ 1.94 \end{gathered}$ | $\begin{gathered} 20.20 \\ \pm \\ 3.32 \end{gathered}$ | $\begin{gathered} 18.10 \pm \\ 1.24 \end{gathered}$ |
| Independent Samples Test |  |  |  |  |  |  |  |  |
| Levene's  <br> Test for  <br> Equality of t -test for Equality of Means <br> Variances  |  |  |  |  |  |  |  |  |
|  | F | Sig. | t | Sig. (2tailed) | Mean Dif. | Std. <br> Error <br> Dif. | 95\% Confidence Interval of the Difference |  |
|  |  |  |  |  |  |  | Lower | Upper |
|  | \% ${ }^{.194}$ | . $665-1$ | $95 \quad 18$ | . $247{ }^{\text {NS }}$ | S -1.500 | 1.2547 | -4.1361 | 1.1361 |


|  | Edge 2 <br> and <br> main crop | . 514 | . 483 | -1.453 | 18 | . $164{ }^{\text {NS }}$ | -1.600 | 1.101 | -3.914 | . 71420 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Ô } \\ & \text { û } \end{aligned}$ | Edge 1 <br> and main crop | . 643 | . 433 | -1.588 | 18 | . $130{ }^{\text {NS }}$ | -2.700 | 1.700 | -6.271 | 0.871 |
|  | Edge 2 <br> and main crop | 2.022 | . 272 | 1.342 | 18 | . $196{ }^{\text {NS }}$ | 2.100 | 1.565 | -1.188 | 5.388 |

2 Table 4. Statistical analysis of difference for HLF ratios between Main Crop saplings and 3 both Edge saplings in the regeneration core with different CCD gradients. This was confirmed
4 by one-way variance analyses (ANOVA) and Student-Newman-Keuls (SNK)


Table 5. Statistical analysis of HLF ratios between G-CCD and S-CCD for each cluster zone

| Independent Samples Test |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{aligned} & \text { HLF } \\ & \text { atios } \end{aligned}$ | Levene's Test for Equality of Variances |  | t-test for Equality of Means |  |  |  |  |  |  |
|  |  | F | Sig. | t | df | $\begin{gathered} \text { Sig. } \\ (2- \\ \text { tailed) } \end{gathered}$ | Mean Dif. | Std. <br> Error <br> Dif. | 95\% Confidence Interval of the Difference |  |
|  |  |  |  |  |  |  |  |  | Lower | Upper |
|  | Edge 1 | . 230 | . 637 | . 243 | 18 | $.811^{\text {NS }}$ | . 03000 | . 12351 | -. 22949 | . 28949 |
|  | Edge 2 | 4.089 | . 058 | -. 092 | 18 | . $928{ }^{\text {NS }}$ | -. 01000 | . 10899 | -. 23897 | . 21897 |
|  | Main crop | 2.810 | . 111 | -1.458 | 18 | $.162^{\text {NS }}$ | -. 16000 | . 10975 | -. 39057 | . 07057 |

