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## MATHEMATIGAL BIOMASS MODELS

FOR YOUNG INDIVIDUALS OF FOREST TREE SPECGES
IN THE REGION OF THE WESTERN GARPATHIANS

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## MATHEMATICAL BIOMASS MODELS <br> FOR YOUNG INDIVIDUALS OF FOREST TREE SPECIES IN THE REGION OF THE WESTERN CARPATHIANS

## MATHEMATICAL BIOMASS MODELS FOR YOUNG INDIVIDUALS

## OF FOREST TREE SPECIES IN THE REGION OF THE WESTERN CARPATHIANS

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## 1. Introduction

We present the target subject, i.e. the development and presentation of mathematical models for tree species biomass estimation in the introductory chapter in a broader context in order to understand its socio-economic importance. We outline the importance of forest biomass in the process of climate change mitigation and the related requirements for the exact quantification of biomass, or the amount of carbon sequestered in the tree layer. The term "climate change" is commonly used to refer to the changes in the Earth's atmosphere that have both regional and global implications. The inherent phenomena of the climate change are gradually becoming the main risks for further development of human civilisation, or for the existence of some species of flora and fauna. Thus, this issue is still gaining its importance and from the long-term perspective it will be the subject of research (including forestry) or implementation of scientific knowledge into practice.

The United Nations Framework Convention on Climate Change (UNFCCC; known as the "Kyoto Protocol") from the year 1997 set the target of stabilising greenhouse gas concentrations in the atmosphere. At the same time, it prescribed a gradual reduction of dangerous anthropogenic interference in the Earth's climate system. Approximately 20 years later, specifically in December 2015, the United Nations Paris Conference on Climate Change took place. The Paris Agreement will replace the Kyoto Protocol after its ratification by national parliaments in 2020. The Paris Conference has committed the countries to keep global warming well below the two degrees Celsius above the state in the pre-industrial era.

The climate change and its inherent phenomena have various negative impacts on environment including its destructive effects on forest ecosystems. However, here we have to note that forests are not only passive objects affected by climate change, but thanks to their ability to absorb and accumulate carbon they can significantly influence this process. Carbon sequestration of forests as one of the complex factors can play an important role in climate change mitigation. Forests are estimated to store as much as $80 \%$ of above-ground and $40 \%$ of below-ground (i.e. roots, plant litter, and soil) terrestrial carbon in their biomass (Dixon et al. 1994). It is also widely known that European forests represent a globally important carbon sink. Although their stocks are increasing, carbon sequestration forest function needs to be further strengthened.

In forest ecosystems, carbon is stored in soil, as well as in plant biomass, particularly in trees. It is the carbon in forest biomass that a man can efficiently influence by its intentional activities. In practice it means that the stand growing stock is gradually increasing thanks to the improvement of production characteristics of forest trees and stands. At the same time, the occurrence of natural disturbances in forests should be avoided or at least reduced. Apart from promoting the sequestration function of forests we need to provide reliable information on the amount of carbon stored in forest biomass, or on the trends of its previous development and future estimates. Thus, the importance of precise determination of biomass in forest trees has been increasing.

In the past, merchantable biomass (the stem part of the tree, or the timber over a certain diameter threshold) was primarily evaluated. A number of practical methods have been developed for the (volume or mass) quantification of merchantable biomass. However, with regard to the climate change, the researchers have started to evaluate all tree components from the point of their energy utilisation, as well as their carbon stocks. This created the need to develop and improve technologies for rapid and statistically representative evaluation of other than stem biomass. One of the most efficient ways to achieve this goal is to use allometric equations based on easily measured (traditionally used) tree characteristics.

According to our literature review, the research in this field has predominantly concentrated on older developmental phases (e.g. Eckmüllner 2006; Seidl et al. 2010; Vejpustková et al. 2015; Krejza et al. 2017). Young trees or forest stands have usually not been included in this research.

Some models developed for young individuals of selected tree species often do not include be-low-ground parts of the biomass (e.g. Annighöfer et al. 2016).

De facto, mathematical models for the biomass estimation of tree components in young (small) trees are still missing. They are necessary because our observations (e.g. Konôpka et al. 2011) suggested that biomass allocation to individual tree components in mature individuals substantially differs from the allocation in young trees. Due to this, the already existing equations for biomass estimation of mature trees and stands are not generally applicable to young trees. Hence, the need to develop specific models for the biomass estimation of all tree components (roots, stem, branches, and foliage) of young trees of different tree species has been identified. They will be useful for quantifying biomass stocks of young regenerated stands, the area of which has substantially increased over the last years.

For these reasons, the presented work focuses on the presentation of mathematical models for the estimation of biomass of young individuals in different tree species. The publication summarises our results, which have been achieved during the last ten years. The prevailing part of the work has been funded by the Slovak Research and Development Agency. The particular projects were: Quantification of biomass in forest stands of $1^{\text {st }}$ age class (2005-2007), Comparative studies of structure of net primary production in beech and spruce stands (2011-2014), Mathematical models of biomass allocation in young stands of selected broadleaved tree species (2013-2017) and Production and ecological studies of tree and ground vegetation after largescale disturbances (2015 onwards).

The research material (whole tree samples) originated from the vast majority of the territory of Slovakia. Hence, the results can be generalised to the conditions of the Western Carpathians. The samples represented eleven tree species. The publication does not deal only with the main commercial tree species, but it also contains the data on some other tree species, which have various functions for the ecological balance of landscape, including their importance for carbon sequestration. Apart from the summary of the existing knowledge on the biomass structure of tree species, this work has an ambition to compare inter-species differences in biomass allocation to individual tree components, and to interpret the differences from production and ecological aspects. In addition, we also wanted to outline the possibilities of further scientific implementation of biomass models of young trees in different tree species.

## 2. Analysis of the subject field and goals of work

As we have already mentioned in the introduction, the volume of merchantable timber was the main subject of interest in the calculation of biomass in forest stands. In Slovakia, this category is characterised by the volume of timber with minimum diameter of 7 cm under bark. Its main proportion is in a stem, the volume of which can be determined using a great number of methods (Hakkila 1989; Petráš \& Pajtík 1991; Husch et al. 2003). Since the time when it was found that biomass allocation significantly determines carbon sequestration and its cycle in a forest ecosystem (Litton et al. 2007), the interest in developing methods for the assessment of non-stem forest biomass has been growing (e.g. Pregitzer \& Euskirchen 2004; Lehtonen 2005).

Models created for older (big) trees are generally not applicable to young individuals (Wirth et al. 2004). Hence, formulas for the calculation of biomass of individuals in initial growth stages need to be derived. The unsuitability of models developed for older trees results from different patterns of biomass allocation in young and old stands. For example Lehtonen et al. (2004) showed that the ratios of individual components of tree biomass depend on the age (or size). In addition, it also reflects different growth strategies of individual tree species and the impact of different forest management or the genesis of previous development. Another reason why the models of older trees cannot be applied to young individuals is the fact that stem diameter at 1.3 m height above ground level is the most common independent variable used to derive the biomass of older trees. This characteristic is not available or cannot be recorded in the case of the youngest trees.

Under the Slovak conditions, Pajtík et al. (2008) emphasised the growing importance of biomass models specifically constructed for young trees. This is related to the increasing area of young stands over the last years (Konôpka et al. 2014). The observed increase is caused by natural disturbances (especially by wind) and secondary pests (in spruce forests mainly bark-beetles; Kunca et al. 2015). Another argument is the concept of uneven-aged forest stands, i.e. often with some proportions of young trees, which is nowadays preferred in many European countries.

The calculation of biomass stock of individual tree components is usually performed using one of the basic methods:

1) regression equations,
2) biomass expansion factors.

The advantage of using regression equations for biomass calculation is that they are frequently based on a larger data set than biomass expansion factors. Another advantage is that they use easily measurable tree characteristics (stem diameter and/or tree height). The regression equations cover the differences in stand structures and can be easily applied to national inventories of carbon stocks. On the other hand, the advantage of biomass expansion factors is their simplicity and more general usage. However, in the case of young stands this advantage is lost because they are derived from stem volume (which is the most readily available information in the case of mature trees, but not for young trees). Another disadvantage is that the values of biomass expansion factors significantly vary in young age classes. On the contrary, the values of mature individuals are more or less stabilised in relation to the tree size, and can be used as a single default (i.e. constant, stable) value (Lehtonen 2004). Our research confirmed significant disadvantages of applying biomass expansion factors to young tree individuals (Pajtík et al. 2008). Due to this, the publication presents only the results related to regression equations.

Biomass regression equations for individual tree species occurred in ecological and forestry literature in the 50s of the last century as a response to the requirement on biomass assessment. Biomass estimates are necessary pre-conditions for the studies of forest production, biochemical or nutrient cycles, biomass energy use, carbon stocks, and carbon sequestration in forests.

The first studies that arose from the need to determine biomass production of different tree species were the works of Burger $(1945,1953)$ dealing with larch and spruce in Switzerland. Subsequently, the scientists started to focus on dry mass determination of individual tree components (usually differentiated to: roots, stem over or under bark, stem bark, branches, foliage), but most frequently of those that were more important to forestry companies.

Ecological and physiological works of that period showed the interest of scientists to contribute to the development of simple methods for biomass determination, especially for the quantification of foliage (Kittredge 1944; Ovington 1957). A number of forestry works have developed regression equations for specific geographic areas and tree species. Biomass equations of a tree as well as of its components are usually based on the relationship to stem diameter (most commonly measured at 1.3 m height above the ground). Some authors used a tree height as an input variable, or a combination of both independent variables (e.g. Satoo \& Madgwick 1982; Ter-Mikaelian \& Korzukhin 1997; Khan \& Faruque 2010; Vahedi et al. 2014). Less frequently, other independent variables are used, e.g. crown length, crown width, ratio of crown length to tree height, ratio of crown width to crown length or $\mathrm{h} / \mathrm{d}$ ratio, i.e. ratio of tree height to stem diameter (e.g. Eckmüllner 2006; Hochbichler et al. 2006; Ledermann \& Neuman 2006; Cienciala et al. 2008).

In the cases when the equations were derived for the biomass calculation at a stand level, site or stand variables were used. Out of them the most frequently applied variables were: number of trees per hectare, basal area, stand top height at a specific stand age, elevation. The number of studies dealing with the assessment of forest biomass has increased over the last decades. They often account for the importance to include a wide number of tree species and different site conditions (Zeide 1987). At the same time, an attempt to create generalised biomass models universally applicable to large regions has also occurred. For example Zianis et al. (2005) made a review of biomass and volume equations for tree species of Europe. In the work they included more than 600 equations, out of which a substantial part originated from Central and Northern Europe. The majority of the studies in this category deals not only with biomass quantification but also with more global aspects. From them we can name e.g. timber utilisation (for the production of pulp, fuel timber, etc.) and application of acquired knowledge in related research areas (e.g. in the studies of carbon cycle, or nutrient balance of forest ecosystems).

The issue of biomass estimation of young stands gained attention only several years ago. Dutca et al. (2010) derived biomass conversion expansion factors (BCEF) for young spruce stands grown on non-forest sites of the Eastern Carpathians. Blujdea et al. (2012) derived allometric equations for the calculation of biomass of young broadleaved trees growing on plantations of Romania. Under the Slovak conditions, Pajtík et al. $(2008,2011)$ derived regression equations and BCEF for spruce, pine, beech, and oak stands in the 1st age class, i.e. younger than 10 years old. Over the last years, the equations for young stands of the following tree species were developed: European ash (Fraxinus excelsior L.), Sycamore (Acer pseudoplatanus L.; e.g. Konôpka et al. 2012, 2015), European larch (Larix decidua Mill.; Pajtík et al. 2015), Goat willow (Salix caprea L.) and Rowan (Sorbus aucuparia L.; Pajtík et al. 2015). The newest summary work focusing on biomass estimation of 19 European tree species in a juvenile stage based on stem base diameter $d_{0}$ (also called root-collar diameter abbreviated as RCD) and tree height was written by Annighöfer et al. (2016). However, the authors of this work did not include the below-ground part of tree biomass.

Several authors (Kozak 1970; Cunia \& Briggs 1984; Parresol 1999; Bi 2004) pointed out at the shortage of many published equations because they did not include the additivity between the equations of individual components and hence were not efficiently determined. It means that the equations were derived for every component separately without accounting for:

1) the correlation between the biomass components measured at the same sample trees,
2) the logical restriction between the sum of the predicted biomass of tree components and the prediction for the whole tree.
The missing additivity in the models causes inconsistency in the predicted values calculated by summing up the equations of individual tree components and the values predicted from the equation for the whole tree biomass. To eliminate this incompatibility, several models and calculation methods were proposed (Chyienda \& Kozak 1984; Cunia \& Briggs 1985).

Allometric equations have an important role among the regression functions. Allometry is the study of varying proportions of organisms dimensions associated with the changes in their size either in the context of the individual growth (ontogenetic allometry), or in comparison to related organisms of different sizes (phylogenetic allometry). This term is also often used to indicate imbalanced growth (development) as an opposite to isometry - balanced growth. Growth allometry is expressed using the allometric equation with the following basic form:

$$
\begin{equation*}
Y=a \cdot X^{b} \tag{1}
\end{equation*}
$$

where $Y=$ a dependent variable, $X=$ an independent explanatory variable, $a, b$ are model coefficients.

Kittredge (1944) was among the first who applied this equation in forestry. Over time, this method has become the most common approach in the studies dealing with biomass quantification (e.g. Marklund 1987; Neumann \& Jandl 2005; Gschwantner \& Schadauer 2006; Ledermann \& Neumann 2006). The reason for its popularity is its flexibility, because it can be easily expanded to a multiple power function in the form:

$$
\begin{equation*}
Y=a_{0} \cdot X_{1}^{b_{1}} \cdot X_{2}^{b_{2}} \cdot X_{3}^{b_{3}} \ldots \ldots . \cdot X_{n}^{b_{n}} \cdot \theta \tag{2}
\end{equation*}
$$

where $Y=$ a dependent variable, $X_{1}-X_{n}=$ independent explanatory variables, $a_{0}-b_{n}=$ model coefficients, and $\theta$ is the error (multiplicative error term).

Frequently, the logarithmic form of the equation is used, because parameters can be estimated using a linear regression. Apart from this advantage, the logarithmic transformation compensates for the tendency to the accelerating increase of the dependent variable with the tree size (the heteroscedasticity of residuals, which is always present in the case of this type of data). Thanks to this approach, the model satisfies the assumption of constant variance. Based on this we can write the equation as follows:

$$
\begin{equation*}
\ln Y=b_{0}+b_{1} \cdot \ln X_{1}+b_{2} \cdot \ln X_{2}+b_{3} \cdot \ln X_{3}+\ldots+\ln X_{n}+\varepsilon \tag{3}
\end{equation*}
$$

where $b_{0}=\ln a_{0}$, and $\varepsilon=\ln \theta$ is the error (additive error term). The logarithmic transformation of the dependent variable causes bias. The bias occurs after the inverse transformation of the logarithmic form to the original one (Baskerville 1972; Ledermann \& Neumann 2006). Hence, when the equations are transformed back, they need to be corrected for the logarithmic bias. For this purpose, a correction factor referred to as $\lambda$ is used.

Finney (1941) and Baskerville (1972) were the first authors who dealt with the calculation of the correction factor for logarithmically transformed allometric equations. However, in spite of the right intention, the formulation of the correction factor was often incorrect at that time. The bias is eliminated by multiplying the result with the correction factor, which is calculated from the standard error of estimates SEE of the regression calculated using the formula:

$$
\begin{equation*}
S E E=\sqrt{\frac{\sum\left(\operatorname{lny}_{i}-\widehat{\ln }_{i}\right)^{2}}{D F}} \tag{4}
\end{equation*}
$$

where $\ln y_{i}$ is the value of the dependent variable, $\widehat{\ln }_{i}$ is the respective predicted value calculated from the equation, and DF is the degrees of freedom, which is calculated as $N-p$, where $N$ is the number of observations and $p$ is the number of equation parameters.

Sprugel (1983) pointed out at the incorrect derivation of SEE by some authors (Snedecor \& Cochran 1967; Whittaker \& Woodwell 1968), who used the values of $N-1$ in the denominator.

The correction factor is then expressed using SEE as follows:

$$
\begin{equation*}
C F=\exp \left(S E E^{2} / 2\right) \tag{5}
\end{equation*}
$$

However, the application of this correction factor requires normal logarithmic distribution of the dependent variable Y, otherwise it causes its overestimation (Marklund 1987). Therefore, instead of this correction factor the method presented by Marklund (1987) is used, who calculates the correction factor using the formula:
$\lambda=\frac{\sum_{i=1}^{n} Y_{i}}{\sum_{i=1}^{n} e^{\widehat{\operatorname{mn}} Y_{i}}}$, where $n$ is the number of trees.
In the case of biomass calculation at a tree level, Ledermann \& Neumann (2006) recommend to use the formula:

$$
\begin{equation*}
\bar{\lambda}=\frac{1}{n} \sum_{i=1}^{n} \frac{Y_{i}}{e^{\widehat{\Pi Y}_{i}}} \tag{7}
\end{equation*}
$$

The calculation of different types of correction factors used for the logarithmic transformation of power functions and their mutual comparison were thoroughly studied by Clifford et al. (2013).

Using a linearised model requires that a user obtains non-transformed biomass values back. For this, the following retransformation is used:

$$
\begin{equation*}
Y=e^{\left(b_{0}+b_{1} \cdot \ln X_{1}+b_{2} \cdot \ln X_{2}+b_{3} \cdot \ln X_{3}+\ldots+b_{n} \cdot \ln X_{n}\right)} \cdot \lambda \tag{8}
\end{equation*}
$$

Recently, the development of non-linear regression methods has raised the question whether it is not more convenient to use allometric equations in their power forms. With this approach we can avoid their logarithmic transformation. Linearisation allows us to use common regression analysis methods, and the calculation process is simpler, especially if several independent variables are included. The disadvantages of this approach are that the logarithmic transformation deforms original data, and the correction factor needs to be used for its retransformation. Cienciala et al. (2006) analysed the effect of linearisation on the calculation of biomass in pine components while deriving regression equations. The authors found that the average biomass predicted using the non-linear regression coincides with the measured values better than when the linearised regression is applied. Other regression statistics, namely the standard error of estimates (SEE), coefficient of determination ( $\mathrm{R}^{2}$ ), and the mean square of residuals (MSR), were slightly better for the linearised two-parameter equations used for calculating the above-ground biomass. On the other hand, the non-linear approach gave better values of the statistical indicators in the case of more complex equations with at least four parameters. On the contrary, Lai et al. (2013) found that in the case of the models for calculating root biomass from tree diameter a linear regression of logarithmically transformed data is more accurate than a non-linear regression. In addition, they revealed that inappropriately used non-linear regressions lead to great inaccuracies in determined biomass at a stand level. This was especially true in the case of stands dominated by small trees. Mascaro et al. (2014) answered the question if logarithmic
transformation is necessary in allometry as follows: "Ten, hundred, thousand times yes". Since the opinions about these two methodical approaches still differ, scientific attention should be paid to this issue also in the future.

The main goal of this work was to summarise and clearly present the results of the long-term scientific work in the field of tree species allometry in juvenile stages of their development, which mainly focused on the development of regression equations for the calculation of biomass of individual tree components, volume, and density of stem and bark. Next, the attention was paid to the inter-species differences in the amount of the total biomass and its allocation to tree components.

The goals set out for this work are related to important commercial tree species, i.e. Common beech, Norway spruce, Sessile oak, and Scots pine. In addition, the research also covered some other tree species, namely European hornbeam, Sycamore, European ash, Goat willow, European larch, Rowan, and Common aspen. These tree species were chosen for our research purposes due to their relatively significant proportion in the tree species composition of our forests or for some other ecological (e.g. pioneer tree species on disturbed plots, amelioration effects on soil) or biological (e.g. their trophic importance for wildlife, or biological protection of target tree species) reasons.

The order of the analysed tree species was selected on the base of the assessment of their importance on the overall (i.e. regardless of age) tree species composition of Slovakia derived from the results of the second round of the National Forest Inventory and Monitoring (NFIM2) of the Slovak Republic (SR). We preferred their area-based occurrence on the whole territory to the traditional comparison based on the contribution to volume or basal area, which are more important from the production view of wood biomass utilisation. The eleven selected tree species are ordered on the base of the relative frequency values from the most to the least common species, while the first six tree species dominate also in the absolute values of all tree species in the Slovak forests, the total number of which is approximately 70 according to the NFIM2 SR.

Based on the area coverage (as well as volume), Common beech ( $1^{\text {st }}$ ) is the most common tree species in Slovakia followed by Norway spruce ( $\left.2^{\text {nd }}\right)$. The third most common tree species is European hornbeam ( $3^{\text {rd }}$ ) that is commercially less important, but currently covers a larger area than the Sessile oak $\left(4^{\text {th }}\right)$ or Scots pine $\left(5^{\text {th }}\right)$. Sycamore $\left(6^{\text {th }}\right)$ occurs in the Slovak forests much more frequently than the European ash $\left(7^{\text {th }}\right.$ ). Relatively equal proportions were found for the pioneer tree species Goat willow ( $8^{\text {th }}$ ), commercially important European larch ( $9^{\text {th }}$ ), and Rowan $\left(10^{\text {th }}\right)$ that is an important admixed amelioration species. The lowest share among the assessed tree species was found for the Common aspen ( $\left.11^{\text {th }}\right)$.

Here we have to note that our allometric equations for young individuals of tree species should complement already existing models for older (bigger) trees. The existing models usually represent the biomass of individual tree components of trees with stem diameter (measured at 1.3 m height above ground) exceeding 7 cm . The persistent problem is that older models usually do not include the amount of stump and roots.

In the following text we present short characterisations of the individual tree species included in this study to better understand their importance. Some information is derived from the most recent data of NFIM2 SR performed in the years 2015-2016.

## 3. Material and methods

We selected several naturally-regenerated stands for each assessed tree species that were in the initial developmental stages from the regeneration stage up to the thicket stage (mostly at the age of 2 to 10 years) from the current national database of young forest stands created from the Forest Management Plans. The proportion of the particular tree species in the area of the selected stands was from 90 to $100 \%$. Basic characteristics of the selected sites are presented in Table 1. Their position inside the area of Slovakia is in Fig. 1.


Fig. 1. Map of the Slovak Republic showing the forest area and the position of sample sites for the following tree species: Common beech (BE), Norway spruce (NS), European hornbeam (HO), Sessile oak (SOK), Scots pine (SP), Sycamore (SY), European ash (AH), Goat willow (GWL), European larch (EL), Rowan (ROW), and Common aspen (ASP).

During the project aims solution, i.e. from the year 2005 to 2016 the methodology was modified and the number of assessed variables increased. At the beginning, when we focused on commercial tree species, i.e. spruce, beech, oak, pine, and larch, the sample trees were not debarked (the methodology was gradually clarified). Hence, the data on dry mass weight of stem bark, volume of stem under bark, bark density, mass and volume proportions of bark are missing. We did not determine the relationship of stem diameter at breast height (hereafter referred to as $d_{1.3}$ diameter) to stem base diameter (hereafter referred to as $d_{0}$ diameter) for spruce, beech, oak and pine. The empirical material is mostly of a national character, only in the case of several tree species it is only of a local character (rowan, goat willow, and larch). The list of sites, from which the sample trees of individual tree species were taken, is presented in Table 1.

In each stand we selected three circular plots, which should represent the whole stand. Their radii varied depending on the stand density to ensure that at least 30 trees occurred within the plot area, which is the number appropriate for statistical evaluation with sufficient confidence. At each plot, we determined the number of individuals and we measured stem base diameter $d_{0}$ (two perpendicular measurements) and height of all trees. From these data we calculated the number of trees per hectare and volume per hectare (the results of these measurements are not included in this monograph). At the same time, these measurements were used to derive diameter and height structures of the stands needed for the subsequent selection of the sample trees.

The stands, from which the sample trees were taken, were selected to ensure that they covered the entire age range from the youngest ( $1-2$ years old) up to the oldest ones ( $10-12$ years). At each site we dug out $20-25$ sample trees that represented the diameter and height range of the individuals at the entire plot. To ensure good coverage of size distribution, we divided the trees to 10 stand-specific height classes of equal widths. Afterwards, we randomly selected and dug out 2 or 3 sample trees from each height class. The trees that were deliberately selected grew in typical stand conditions. For example, we avoided solitary trees, individuals at the vicinity of paths, or those at the forest edge. We also excluded the individuals that were deformed, damaged, or with reduced foliage.

The trees were sampled at the end of the growing season, when the growth of all components was finished. The sample trees were divided to roots, stem, branches, foliage, and stem bark. The samples were packed into the marked paper bags and transported to laboratories for further processing. Every sample was stored in a dry, ventilated room for one month. Afterwards, it was dried in an electric oven at a temperature of $105^{\circ} \mathrm{C}$ until it reached constant weight.

The described method of sample tree selection ensured that at each plot we chose trees from all sociological positions (dominant, co-dominant, sub-dominant, suppressed). On the other hand, this approach led to substantially left-skewed distributions of diameters and heights of the whole set of measured trees, where the greatest diameters and heights were represented only by the trees from the main canopy. The sub-dominant and suppressed trees of such dimensions would have had to be searched for in older stands. However, that was not the goal of this work, which aimed at evaluating only the stands of juvenile growth stages. In the case of the left-skewed distribution of the values, the linearisation of the allometric equation and its reverse re-transformation can in some cases cause that the predicted values substantially differ from the measured values, which is true mainly for the high values.

Model development was primarily focused on the calculation of dry mass weight of tree components using regression functions as well as biomass conversion expansion factors (not shown in this work). Only gradually, in the course of the project solution, we began to develop also the models for the volume of stem, and density and proportion of bark.

When calculating dry mass of tree components, the biomass of individual tree components given in mass units was the dependent variable. Due to the small tree dimensions, $d_{1.3}$ diameter could not be used as an independent variable. Instead, stem base diameter $d_{0}$ was applied. Although the models with height as the only independent variable are used only rarely, we applied also this model. The reason for this was that in the youngest developmental stages height is more easily measured than $d_{0}$ diameter. At the same time, height can be used to couple the models of mature stands with the models of the stands in the initial growth stages. We tested three functions, in which the independent variables were $d_{0}$ diameter, tree height, and their combination.

$$
\begin{gather*}
W_{i}=e^{\left(b_{0}+b_{1} \cdot \ln d_{0}\right)} \cdot \lambda  \tag{9}\\
W_{i}=e^{\left(b_{0}+b_{1} \cdot \ln h\right)} \cdot \lambda  \tag{10}\\
W_{i}=e^{\left(b_{0}+b_{1} \cdot \ln d_{0}+b_{2} \cdot \ln h\right)} \cdot \lambda \tag{11}
\end{gather*}
$$

where:
$W_{i} \quad=$ biomass weight of $i^{\text {th }}$ tree component ( g of dry matter expressed at a tree level),
$d_{0} \quad=$ stem base diameter $(\mathrm{mm})$,
$h \quad=$ tree height $(\mathrm{m})$,
$b_{0}, b_{1}, b_{2}=$ equation coefficients,
$\lambda \quad=$ correction factor.

Prior to the calculation of stem volume, a stem was divided to at least $3-4$ sections. The sections were measured before drying, i.e. the volume represents the fresh state. The section diameters were measured in two perpendicular directions with a vernier calliper with a precision of one tenth of millimetre. Tree height was determined with a precision of one centimetre. The stem section volume was calculated using the Newton formula:

$$
\begin{equation*}
V=\frac{L\left(A_{b}+4 A_{m}+A_{s}\right)}{6} \tag{12}
\end{equation*}
$$

where:
$V=$ stem volume $\left(\mathrm{cm}^{3}\right)$,
$L \quad=$ section length (cm),
$A_{b}=$ cross-sectional area at the bottom end of the section $\left(\mathrm{cm}^{2}\right)$,
$A_{m}=$ cross-sectional area in the middle of the section $\left(\mathrm{cm}^{2}\right)$,
$A_{s}=$ cross-sectional area at the top end of the section ( $\mathrm{cm}^{2}$ ).
The total stem volume was calculated as a sum of volumes of all sections.
The Newton formula is considered to be the most accurate and flexible formula for the calculation of volume of stem parts (logs, sections), because it is suitable for the volume calculation of cylindrical and conical objects, but also of paraboloids and neloids (Wiant et al. 1992; Harmon \& Sexton 1996; Woldendorp et al. 2002). The calculated values were validated using a pycnometer (a cylinder filled with a liquid used for measuring an object volume). The differences between the calculated and measured volumes were from -2 to $+5 \%$. Since such a volume determination is appropriate only in laboratory conditions, in forestry practice stem volume is usually calculated using one or two easily measurable characteristics. Most stem volume equations use $d_{1.3}$ diameter and tree height as independent variables. Because equations with $d_{1.3}$ as an independent variable are not applicable to young stands, we derived three allometric equations to determine stem volume. In the first case, stem base diameter $d_{0}$ was used as an independent variable; in the second case it was the tree height $(h)$, and in the third case we used both variables $\left(d_{0}, h\right)$. Due to the above-mentioned shortages of logarithmically transformed equations we used non-linear formulas:

$$
\begin{align*}
& V=b_{0} d_{0}^{b_{1}}  \tag{13}\\
& V=b_{0} h^{b_{1}}  \tag{14}\\
& V=b_{0} d_{0}^{b_{1}} h^{b_{2}} \tag{15}
\end{align*}
$$

where:
$V \quad=$ stem volume $\left(\mathrm{cm}^{3}\right)$,
$d_{0} \quad=$ stem base diameter (mm),
$h \quad=$ tree height ( m ), and
$b_{0}, b_{1}$ and $b_{2}=$ equation coefficients.
Stem volume was determined in the fresh (moist) state after the sample trees were dug out as the volume of stem over bark (hereafter as SOB), and after debarking we calculated also the volume of stem under bark (hereafter as SUB). Bark volume was calculated as the difference between these two values. Stem volume was calculated as a sum of volumes of individual sections [equation 12]. Wood density was determined as a reduced wood density $\rho_{r f}$ in the fresh state (reduced wood density) defined as a ratio of wood mass in the absolutely dry state $m_{0}$ to the fresh wood volume $V_{\max }$ :

$$
\begin{equation*}
\rho_{r f}=\frac{m_{0}}{V_{\max }} \cdot 1000 \tag{16}
\end{equation*}
$$

where:

```
\(\rho_{r f}=\) reduced wood density in fresh state \(\left(\mathrm{kg} . \mathrm{m}^{-3}\right)\),
\(m_{0}=\) wood mass in absolutely dry state (g),
\(\mathrm{V}_{\text {max }}=\) volume of wood with moisture above the hygroscopic threshold \(\left(\mathrm{cm}^{3}\right)\).
```

The fresh wood moisture is above the hygroscopic threshold. Reduced wood density above this threshold reaches its minimum value. This is caused by the fact that at these values it does not depend on wood moisture, because wood does not swell any longer and its volume is at maximum $V_{\max }$.

For the calculation of volume, basic density and proportion of bark we used allometric equations [13] - [15] with one independent variable $d_{0}$, or $h$, and two independent variables $d_{0}$ and $h$. The relationship of $d_{1.3}$ diameter to $d_{0}$ diameter was described using the following linear equation:
$d_{1,3}=b_{0} d_{0}+b_{1}$
where:
$d_{1,3}=$ stem diameter at a height of 1.3 m above ground ( mm ),
$d_{0}=$ stem base diameter (mm),
$b_{0}$ and $b_{1}$ are equation coefficients,
and the relationship of tree height $h$ to stem base diameter $d_{0}$ was described using the following equation:
where:
$h=\frac{d_{0}^{2}}{b_{0}+b_{1} d_{0}+b_{2} d_{0}^{2}}$
$h \quad=$ tree height (m),
$d_{o}=$ stem base diameter ( mm ),
$b_{0}, b_{1}$ and $b_{2}$ are equation coefficients.
Stem bark mass proportion was calculated using the formula:
where:

$$
\begin{equation*}
R_{w b}=\frac{100 w_{b}}{m_{s O B}} \tag{19}
\end{equation*}
$$

$R_{w b}=$ bark mass proportion (\%),
$w_{b}=$ dry mass of stem bark (g),
$m_{\text {SOB }}=$ dry mass of stem over bark (g).
Stem bark volume proportion was calculated using the formula:
where:
$R_{V b}=\frac{100 V_{b}}{V_{S O B}}$
$R_{V b}=$ volume proportion of bark (\%),
$V_{b}=$ fresh bark volume after debarking $\left(\mathrm{cm}^{3}\right)$,
$V_{S O B}=$ volume of fresh stem over bark $\left(\mathrm{cm}^{3}\right)$.

Root/shoot ratio represents the ratio of the dry root mass (below-ground part) to the aboveground dry mass, i.e:

$$
\begin{equation*}
\frac{R}{S}=\frac{w_{r}}{w_{\text {abvg }}} \tag{21}
\end{equation*}
$$

$\frac{R}{S}=$ the ratio of the dry root mass (below-ground part) to the above-ground dry mass,
where:
$w_{r}=$ dry root mass (g),
$w_{\text {abvg }}=$ aboveground dry mass (g).
Mass proportion of a component $i$ (foliage, branches, stem over bark, roots) was calculated using the formula:

$$
\begin{equation*}
R_{w i}=\frac{100 w_{i}}{w} \tag{22}
\end{equation*}
$$

where:
$R_{w i}=$ mass proportion of $i^{\text {th }}$ component (\%),
$w_{i}=$ dry mass of $i^{\text {th }}$ component (g),
$w \quad$ dry mass of the whole tree ( g ).
When visualising the data in graphs, we followed the principle of applying the same range of the values of dry matter of a particular component to $y$ axis for all assessed tree species. This allows simple comparison of the component quantities between tree species.

Table 1. List of sites, from which the sample trees of individual tree species were taken.

| Tree species | No. | Name of site | Elevation (m) | N latitude $\left({ }^{\circ}\right)$ | $\begin{gathered} \text { E longitude } \\ \left({ }^{\circ}\right) \\ \hline \end{gathered}$ | Aspect | Slope (\%) | Soil | Bedrock | Site order | Group of forest types |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | ŠLPI | 710 | 48.6318 | 19.0048 | W | 41 | Mesotrophic cambisols | Andesites | Fertile | Fagetum typicum |
|  | 2 | ŠLP II | 675 | 48.6454 | 19.0531 | SW | 19 | Mesotrophic cambisols | Andesites | Fertile | Fagetum typicum |
|  | 3 | Zvolen | 460 | 48.5523 | 19.1251 | N | 30 | Mesotrophic cambisols | Andesites | Fertile | Fagetum pauper |
|  | 4 | Králová | 550 | 48.5343 | 19.1584 | NW | 30 | Mesotrophic cambisols | Andesites | Fertile | Fagetum pauper |
|  | 5 | Sekier I | 670 | 48.4987 | 19.2223 | N | 31 | Mesotrophic cambisols | Andesites | Fertile | Fagetum pauper |
|  | 6 | Sekier II | 660 | 48.4990 | 19.2200 | N | 28 | Mesotrophic cambisols | Andesites | Fertile | Fagetum pauper |
|  | 7 | Hrochot | 620 | 48.6579 | 19.2820 | N | 13 | Oligotrophic cambisols | Quartzites | Fertile | Fagetum pauper |
|  | 1 | Polana I | 985 | 48.6415 | 19.5164 | SW | 34 | Mesotrophic cambisols | Andesites | Fertile | Fagetum typicum |
|  | 2 | Polana II | 790 | 48.6418 | 19.5358 | NE | 28 | Mesotrophic cambisols | Andesites | Fertile | Abieto-Fagetum |
|  | 3 | Drakšiar I | 625 | 48.8490 | 19.7897 | SE | 11 | Mesotrophic cambisols | Sandstones | Fertile | Fageto-Abietum |
|  | 4 | Drakšiar II | 635 | 48.8493 | 19.7919 | S | 16 | Mesotrophic cambisols | Sandstones | Fertile | Abieto-Fagetum |
|  | 5 | Bacúch | 840 | 48.9009 | 19.8149 | S | 52 | Rendzinic cambisols | Limestones | Fertile | Abieto-Fagetum |
|  | 6 | Čierny Váh I | 820 | 48.9840 | 19.9627 | N | 41 | Mesotrophic cambisols | Melaphyres | Fertile | Fageto-Abietum |
|  | 7 | Čierny Váh II | 830 | 48.9837 | 19.9632 | N | 38 | Mesotrophic cambisols | Melaphyres | Fertile | Fageto-Abietum |
| $\begin{aligned} & \text { HO - European hornbeam } \\ & \text { (Carpinus betulus) } \end{aligned}$ | 1 | Píla | 313 | 48.3941 | 17.2944 | S | 1 | Typical paternia | Alluvium | Water- <br> logged | Fraxineto-Alnetum |
|  | 2 | Rudica | 475 | 48.5935 | 18.5497 | W | 23 | Mesotrophic cambisols | Andesites | Fertile | Fageto-Quercetum |
|  | 3 | Antol | 516 | 48.3955 | 18.9564 | W | 13 | Ilimerised soil | Clay loess | Fertile | Fageto-Quercetum |
|  | 4 | Breziny | 432 | 48.5271 | 19.0816 | NE | 20 | Mesotrophic cambisols | Andesites | Fertile | Fageto-Quercetum |
|  | 5 | Cerovo | 560 | 48.2475 | 19.2295 | SW | 8 | Mesotrophic cambisols | Andesitic tuff | Fertile | Querceto-Fagetum |
|  | 6 | Soroška | 567 | 48.6109 | 20.6057 | NW | 9 | Moderrendzinas | Limestones | Fertile | Fagetum pauper |
|  | 7 | Budimír | 295 | 48.7938 | 21.2916 | SW | 3 | Illimerised soils | No data | Fertile | Fageto-Quercetum |
|  | 8 | Zubné | 350 | 49.0459 | 22.0874 | N | 34 | Illimerised soils | Sandstones | Fertile | Fagetum pauper |
|  | 1 | Ladzany I | 480 | 48.2904 | 18.8576 | S | 9 | Illimerised soils | Clay loess | Fertile | Fageto-Quercetum |
|  | 2 | Ladzany II | 500 | 48.2921 | 18.8473 | SE | 6 | Illimerised soils | Clay loess | Fertile | Fageto-Quercetum |
|  | 3 | Antol | 560 | 48.3696 | 18.9534 | E | 30 | Illimerised soils | Andesites | Fertile | Fageto-Quercetum |
|  | 4 | Žibritov I | 480 | 48.3790 | 19.0312 | SW | 16 | Illimerised soils | Clay loess | Fertile | Fageto-Quercetum |
|  | 5 | Krupina I | 460 | 48.3382 | 19.0209 | E | 49 | Mesotrophic cambisols | Andesites | Fertile | Fageto-Quercetum |
|  | 6 | Žibritov II | 480 | 48.3810 | 19.0267 | S | 20 | Illimerised soils | Clay loess | Fertile | Fageto-Quercetum |
|  | 7 | Krupina II | 380 | 48.3785 | 19.0831 | N | 22 | $\begin{gathered} \text { Illimerised } \\ \text { soils } \\ \hline \end{gathered}$ | Clay loess | Fertile | Fageto-Quercetum |


| Tree species | No. | Name of site | Elevation (m) | N latitude $\left({ }^{\circ}\right)$ | E longitude $\left({ }^{\circ}\right)$ | Aspect | Slope (\%) | Soil | Bedrock | Site order | Group of forest types |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | Kopčany | 165 | 48.7539 | 17.0806 | NW | 1 | Ranker cambisols | Drifted sand | Fertile | Carpineto-Quercetum |
|  | 2 | Kopčany II | 165 | 48.7547 | 17.0792 | NW | 0 | Ranker cambisols | Drifted sand | Fertile | Carpineto-Quercetum |
|  | 3 | Žiar n/Hr. | 380 | 48.6026 | 18.8742 | SE | 4 | Pseudogley | Andesites | Fertile | Fageto-Quercetum |
|  | 4 | Kovácová | 380 | 48.5934 | 19.0717 | S | 15 | Illimerised soils | Andesites | Fertile | Fageto-Quercetum |
|  | 5 | Zolná | 430 | 48.6193 | 19.2209 | S | 1 | Pseudogley | Andesites | Fertile | Fageto-Quercetum |
|  | 6 | Kišovce | 640 | 49.0282 | 20.3715 | SW | 20 | Typical pararendzina | Calcareous sandstone | Limestone | Pinetum dealpinum |
|  | 7 | Levoča | 620 | 49.0182 | 20.5126 | N | 24 | Mesotrophic cambisols | Sandstones | Fertile | Piceeto-Pinetum |
|  | 1 | Devínska Kobyla | 456 | 48.1876 | 16.9991 | SE | 10 | Illimerised soils | Clay loess | Nitrophillic | Querceto-Fagetum tiliosum |
|  | 2 | Devínska Kobyla | 415 | 48.1853 | 17.0022 | SE | 12 | Illimerised soils | Clay loess | Fertile | Querceto-Fagetum |
|  | 3 | Tužina | 644 | 48.9151 | 18.5992 | SW | 38 | Mesotrophic cambisols | Granites | Fertile | Fagetum typicum |
|  | 4 | Oščadnica | 684 | 49.4640 | 18.8723 | NE | 22 | Mesotrophic cambisols | Sandstones | Fertile | Abieto-Fagetum |
|  | 5 | Lohyňa | 740 | 48.4780 | 19.2957 | W | 51 | Mesotrophic cambisols | Andesites | Fertile | Fagetum pauper |
|  | 6 | Chvojno | 406 | 48.5360 | 19.3378 | N | 23 | Mesotrophic cambisols | Andesites | Fertile | Querceto-Fagetum |
|  | 7 | Snohy | 790 | 48.6276 | 19.5384 | SE | 14 | Mesotrophic cambisols | Granodiorites | Fertile | Fagetum typicum |
|  | 8 | Nad nádržou | 639 | 48.6106 | 19.5674 | SW | 27 | Mesotrophic cambisols | Granodiorites | Fertile | Fagetum typicum |
|  | 9 | Vrchslatina | 970 | 48.6472 | 19.6037 | W | 2 | Mesotrophic cambisols | Granodiorites | Nitrophillic | Fageto-Aceretum |
|  | 10 | Lom | 950 | 48.6407 | 19.6287 | SW | 28 | Mesotrophic cambisols | Granodiorites | Fertile | Abieto-Fagetum |
|  | 11 | Kravany | 850 | 49.0088 | 20.2131 | NE | 52 | Mesotrophic cambisols | Sandstones | Fertile | Fageto-Abietum |
|  | 12 | Jahodná | 550 | 48.7599 | 21.1380 | SW | 37 | Mesotrophic cambisols | Rhyolites | Fertile | Fagetum typicum |
|  | 13 | Čermel' | 480 | 48.7564 | 21.1704 | NE | 33 | Mesotrophic cambisols | Rhyolites | Fertile | Fagetum typicum |
|  | 1 | Lohyňa | 736 | 48.4780 | 19.2963 | W | 48 | Mesotrophic cambisols | Rhyolites | Fertile | Fagetum pauper |
|  | 2 | Sliačska <br> Polana | 731 | 48.4880 | 19.3914 | SW | 4 | Mesotrophic cambisols | Andesites | Fertile | Fagetum tiliosum |
|  | 3 | Sliačska <br> Polana | 792 | 48.4998 | 19.3842 | E | 13 | Mesotrophic cambisols | Andesites | Nitrophillic | Fagetum typicum |
|  | 4 | Snohy | 781 | 48.6277 | 19.5379 | SE | 13 | Mesotrophic cambisols | Andesites | Fertile | Fagetum typicum |
|  | 5 | Vrchslatina | 848 | 48.6368 | 19.5914 | SW | 27 | Mesotrophic cambisols | Granodiorites | Fertile | Fagetum typicum |
|  | 6 | Vrchslatina | 860 | 48.6374 | 19.5912 | SW | 25 | Mesotrophic cambisols | Granodiorites | Fertile | Fagetum typicum |
|  | 1 | Husárik | 800 | 49.4124 | 18.7693 | SE | 26 | Mesotrophic cambisols | Sandstones | Fertile | Fageto-Abietum |
|  | 2 | Tatry | 1030 | 49.1323 | 20.2032 | S | 13 | Podsolic cambisols | Granites | Acidic | Piceeto-Abietum |


| Tree species | No. | Name of site | Elevation (m) | N latitude $\left({ }^{\circ}\right)$ | E longitude <br> $\left({ }^{\circ}\right)$ | Aspect | Slope <br> (\%) | Soil | Bedrock | Site order | Group of forest types |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | SmokovceFIR | 1060 | 49.1351 | 20.1976 | SE | 24 | Podsolic cambisols | Granites | Acidic | Piceetum abietinum |
|  | 2 | Stará LesnáKolbach | 840 | 49.1522 | 20.2691 | S | 11 | Podsolic cambisols | Granites | Acidic | Piceetum abietinum |
|  | 3 | Stará Lesnánad campom | 834 | 49.1520 | 20.2794 | E | 4 | Podsolic cambisols | Granites | Acidic | Pineto-Piceetum |
|  | 4 | Matliare- <br> Biela Voda | 910 | 49.1855 | 20.2923 | E | 7 | Podsolic cambisols | Fluvioglacial | Acidic | Piceetum abietinum |
|  | 5 | Matliare- <br> Zubry | 810 | 49.1656 | 20.3129 | S | 5 | Oligotrophic cambisols | Granites | Acidic | Pineto-Piceetum |
|  | 6 | Matliare- <br> Rozengart | 790 | 49.1604 | 20.3222 | S | 3 | Oligotrophic cambisols | Granites | Acidic | Pineto-Piceetum |
|  | 1 | Zruby | 988 | 49.1284 | 20.1987 | SE | 10 | Ranker cambisols | Granites | Acidic | Piceeto-Abietum |
|  | 2 | Smokovec | 950 | 49.1285 | 20.2161 | S | 11 | Oligotrophic cambisols | Fluvioglacial | Fertile | Piceeto-Abietum |
|  | 3 | Šart | 1122 | 49.1839 | 20.2620 | E | 18 | Podsolic cambisols | Fluvioglacial | Acidic | Lariceto-Piceetum |
|  | 4 | Jamy | 941 | 49.1628 | 20.2645 | NE | 10 | Oligotrophic cambisols | Fluvioglacial | Acidic | Piceetum abietinum |
|  | 5 | Nad nádržou | 960 | 49.1692 | 20.2643 | SE | 15 | Podsolic cambisols | Granodiorites | Acidic | Piceetum abietinum |
|  | 1 | Kašova Lehôtka | 610 | 48.6204 | 19.0288 | SW | 12 | Mesotrophic cambisols | Andesites | Fertile | Fagetum typicum |
|  | 2 | Stráže | 335 | 48.5864 | 19.0897 | SW | 13 | Illimerised soils | Andesites | Fertile | Fageto-Quercetum |
|  | 3 | Dobrá Niva | 365 | 48.4522 | 19.1003 | NE | 7 | Mesotrophic cambisols | Andesites | Fertile | Fageto-Quercetum |
|  | 4 | Suchán̆ | 540 | 48.2896 | 19.1023 | N | 14 | Mesotrophic cambisols | Andesitic tuff | Fertile | Fageto-Quercetum |
|  | 5 | Opava | 525 | 48.1998 | 19.2235 | SW | 22 | Mesotrophic cambisols | Andesitic tuff | Fertile | Fageto-Quercetum |
|  | 6 | Podkonice | 550 | 48.7930 | 19.2672 | SW | 15 | Moder-rendzinas | Limestones | Limestone | Querceto-Fagetum dealpinum |
|  | 7 | Telgárt | 870 | 48.8359 | 20.1711 | NE | 9 | Mesotrophic cambisols | Gneiss | Acidic | Fageto-Abietum |

## 4. Results

The results about the biomass characteristics are presented in separate chapters for every tree species (i.e. chapters 4.1. to 4.11.). The order of the chapters corresponds to the occurrence frequency of tree species in Slovakia, i.e. from the most frequent (i.e. Common beech) to the least frequent tree species (Common aspen). The last chapter (4.12.) presents the summary results for all tree species together, and aims at comparing the inter-species differences.

### 4.1. Common beech

Common beech (Fagus sylvatica L.) is the most common tree species in Slovakia from the point of species composition and occurrence frequency. It is also commercially the most important tree species with approximately $30 \%$ proportion in total stock (or area). Naturally it grows at a variety of sites in a wide range of all forest elevation zones. Fertile sites suit beech best. It creates either homogeneous stands, or stands composed of two main tree species (mainly with hornbeam or oak). Beech is an important element of the so-called Carpathian mixture together with spruce and fir, though it also creates combinations with other tree species. It is an important component of biotopes protected at national or European levels. The most common forest biotope in Slovakia is Ls5.1 Beech and fir-beech flowery forests (NATURA 2000 defines it as 9130 AsperuloFagetum beech forests), which covers more than half a million of hectares. Other biotopes with a dominant proportion of beech are Ls5.2 Acidophilous beech forests ( 9110 Luzulo-Fagetum beech forests), Ls5.3 Maple beech mountainous forests ( 9140 Medio-European subalpine beech woods with Acer and Rumex arifolius), Ls5.4 Calcareous beech forests (9150 Medio- European limestone beech forests of the Cephalanthero - Fagion).

Beech occurs in all forest vegetation zones (fvz) from the $1^{\text {st }}$ oak zone up to $7^{\text {th }}$ spruce zone with the dominant proportion in $3^{\text {rd }}$ and $4^{\text {th }}$ zones. Its ecological optimum is in $4^{\text {th }}$ beech forest vegetation zone, and its production optimum is in $5^{\text {th }}$ fir-beech zone. On the base of the processed NFIM2 SR data from the years 2015-2016, the minimum and maximum elevations at which beech occurred were 130 m and $1,466 \mathrm{~m}$ a.s.l., respectively, and on average it grew at elevations 600-700 m a.s.l. According to the NFIM2 SR results, it grew at a reduced area of 666 $\pm 39$ thousand ha (the value following $\pm$ sign represents $95 \%$ confidence interval), and occurred at $62 \%$ of forested inventory plots.


Fig. 2. Map of sample sites of Common beech and its distribution in the forests of Slovakia.

The set of beech trees used to derive the biomass regression models consisted of 170 whole tree samples. They were taken from seven sites (see Fig. 2) located in the orographic units of the Kremnické vrchy (sites 1 and 2), Štiavnické vrchy (3), Javorie (4, 5, 6) and Polana (7). The sampled individuals had $d_{0}$ diameters from 4.20 mm to 68.50 mm , and heights from 0.24 m to 5.40 m (Table 2, Fig. 3a). The whole tree dry mass ranged from 2.61 g to $6,148.10 \mathrm{~g}$, and the stem volume ranged from $1.23 \mathrm{~cm}^{3}$ to $5,059.20 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 3. The models use two independent variables, namely $d_{0}$ diameter [Equation 9], tree height [Equation 10], or the combination of these two variables [Equation 11]. Next, we derived the volume of stem over bark, its density, as well as the root/shoot ratio (Table 4). Also in this case we used two independent variables, i.e. $d_{0}$ diameter [Equation 13], tree height [Equation 14], or the combination of both variables [Equation 15].

The scatter plots showing the biomass of individual components (or of the whole trees) of the whole set of the analysed trees in relation to $d_{0}$ diameter with the fitted regression curves (regression models 9) are presented in Fig. 3b - 3f. Similarly, we graphically presented the proportion of the total tree biomass in individual components in relation to $d_{0}$ diameter (Fig. 4a), the volume of stem over bark (Fig. 4b), and the density of stem over bark (Fig. 4c), as well as the ratio of below-ground to above-ground biomass (i.e. "root-shoot ratio" abbreviated as R/S; see Fig. 4d). Further comments on the biomass in individual components of beech and their proportions of the total tree biomass are presented in Chapter 4.12. (Inter-species comparison of biomass characteristics).

Table 2. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25.p), 75-percentile (75.p) and skewness of diameter $\left(d_{0}\right)$, tree height ( $h$ ), biomass of stem over bark (SOB), foliage biomass (foliage), branch biomass (branches), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {SOB }}$ ) of individual trees.

|  | Common beech |  |  |  |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | 25.p | 75.p | Skewness |
| $d_{0}(\mathrm{~mm})$ | 170 | 14.50 | 8.67 | 4.20 | 68.50 | 8.65 | 17.35 | 2.47 |
| $h(\mathrm{~m})$ | 170 | 1.25 | 0.81 | 0.24 | 5.40 | 0.70 | 1.56 | 1.78 |
| SOB $(\mathrm{g})$ | 170 | 90.02 | 283.55 | 0.86 | 3197.40 | 8.80 | 62.00 | 8.56 |
| Foliage $(\mathrm{g})$ | 170 | 23.65 | 51.34 | 0.51 | 564.10 | 3.90 | 20.60 | 7.50 |
| Branches $(\mathrm{g})$ | 170 | 32.45 | 127.96 | 0.80 | 1533.60 | 2.20 | 19.00 | 10.04 |
| Roots $(\mathrm{g})$ | 170 | 47.54 | 88.07 | 0.55 | 853.00 | 9.17 | 42.00 | 5.49 |
| Aboveground | 170 | 146.11 | 459.40 | 1.45 | 5295.10 | 14.50 | 100.00 | 8.96 |
| Whole tree $(\mathrm{g})$ | 170 | 193.65 | 542.87 | 2.61 | 6148.10 | 25.36 | 146.18 | 8.47 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 170 | 178.04 | 470.12 | 1.23 | 5059.20 | 13.50 | 135.42 | 7.39 |

Table 3. Common beech, $b_{0}, b_{1}$, $b_{2}$ regression coefficients, their standard errors (S.E.), p-values ( $P$ ), coefficient of determination $\left(R^{2}\right)$, mean square error (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -4.034 | 0.164 | <0.001 | 2.852 | 0.063 | <0.001 |  |  |  | 0.923 | 0.167 | 1.084 | 0.438 |
|  | Branches | -5.982 | 0.218 | <0.001 | 3.117 | 0.084 | <0.001 |  |  |  | 0.891 | 0.295 | 1.142 | 0.592 |
|  | Foliage | -3.750 | 0.183 | $<0.001$ | 2.375 | 0.071 | $<0.001$ |  |  |  | 0.871 | 0.207 | 1.102 | 0.486 |
|  | Roots | -2.960 | 0.179 | $<0.001$ | 2.361 | 0.069 | $<0.001$ |  |  |  | 0.874 | 0.199 | 1.098 | 0.495 |
|  | Aboveground part | -3.288 | 0.144 | $<0.001$ | 2.777 | 0.056 | <0.001 |  |  |  | 0.937 | 0.129 | 1.063 | 0.364 |
|  | Whole tree | -2.521 | 0.132 | $<0.001$ | 2.639 | 0.051 | $<0.001$ |  |  |  | 0.941 | 0.108 | 1.053 | 0.336 |
| [10] | Stem over bark | 3.108 | 0.043 | <0.001 | 2.302 | 0.073 | <0.001 |  |  |  | 0.854 | 0.318 | 1.166 | 0.672 |
|  | Branches | 1.833 | 0.069 | $<0.001$ | 2.324 | 0.117 | $<0.001$ |  |  |  | 0.703 | 0.803 | 1.474 | 1.671 |
|  | Foliage | 2.206 | 0.058 | $<0.001$ | 1.712 | 0.098 | $<0.001$ |  |  |  | 0.643 | 0.573 | 1.308 | 1.068 |
|  | Roots | 2.964 | 0.061 | $<0.001$ | 1.651 | 0.102 | $<0.001$ |  |  |  | 0.607 | 0.620 | 1.298 | 0.949 |
|  | Aboveground part | 3.669 | 0.048 | $<0.001$ | 2.161 | 0.082 | $<0.001$ |  |  |  | 0.806 | 0.396 | 1.216 | 0.855 |
|  | Whole tree | 4.094 | 0.050 | $<0.001$ | 2.000 | 0.085 | $<0.001$ |  |  |  | 0.767 | 0.427 | 1.226 | 0.853 |
| [11] | Stem over bark | -1.530 | 0.163 | $<0.001$ | 1.848 | 0.065 | $<0.001$ | 1.015 | 0.054 | <0.001 | 0.975 | 0.054 | 1.026 | 0.222 |
|  | Branches | -4.768 | 0.364 | $<0.001$ | 2.630 | 0.144 | $<0.001$ | 0.423 | 0.121 | $<0.001$ | 0.901 | 0.270 | 1.130 | 0.560 |
|  | Foliage | -3.286 | 0.317 | $<0.001$ | 2.188 | 0.126 | $<0.001$ | 0.188 | 0.105 | 0.076 | 0.873 | 0.205 | 1.100 | 0.479 |
|  | Roots | -2.898 | 0.314 | $<0.001$ | 2.336 | 0.124 | <0.001 | 0.025 | 0.104 | 0.809 | 0.874 | 0.200 | 1.098 | 0.493 |
|  | Aboveground part | -1.486 | 0.186 | $<0.001$ | 2.054 | 0.074 | $<0.001$ | 0.731 | 0.062 | $<0.001$ | 0.966 | 0.070 | 1.034 | 0.266 |
|  | Whole tree | -1.236 | 0.197 | $<0.001$ | 2.124 | 0.078 | $<0.001$ | 0.521 | 0.065 | $<0.001$ | 0.957 | 0.079 | 1.038 | 0.285 |

Table 3 presents statistical characteristics of the three models derived for the calculation of the dry mass of individual tree components using different independent variables ( $d_{0}, h$, or their combination). Although $d_{0}$ diameter is a variable frequently affected by stem thickening, and it is very difficult to determine its precise position at the stem, our analyses showed that all models using this variable were from the point of their coefficients of determination $R^{2}$ more suitable for the calculation of the dry mass of individual components than the models containing height as an independent variable. The model, which contained both independent variables, had only a slightly higher value of $R^{2}$ if at all, and because the values of $R^{2}$ were high in the case of all the derived models, we did not consider using other independent variables ( $d^{2}, d / h$, crown length, crown width, etc.).

This knowledge is generally valid for all tree species, and hence, we will not repeatedly comment the tables with the statistical characteristics of the models for the calculation of dry mass of individual tree components in the further text.

Table 4. Common beech, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), $p$-values ( $P$ ), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE) for equations [13] - [15].

| Eq. | Dependent variable | $\mathrm{b}_{0}$ | S.E. | P | $\mathrm{b}_{1}$ | S.E. | P | $\mathrm{b}_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [13] | SOB volume | 0.034 | 0.005 | $<0.001$ | 2.821 | 0.035 | $<0.001$ |  |  |  | 0.977 | 4613 |
|  | SOB density | 720.637 | 35.433 | $<0.001$ | -0.038 | 0.019 | 0.050 |  |  |  | 0.022 | 6418 |
|  | R/S ratio | 1.366 | 0.219 | $<0.001$ | -0.374 | 0.066 | <0.001 |  |  |  | 0.185 | 0.046 |
| [14] | SOB volume | 13.984 | 1.205 | $<0.001$ | 3.490 | 0.055 | $<0.001$ |  |  |  | 0.968 | 6518 |
|  | SOB density | 656.288 | 5.906 | $<0.001$ | -0.066 | 0.015 | $<0.001$ |  |  |  | 0.099 | 5912 |
|  | R/S ratio | 0.530 | 0.015 | <0.001 | -0.444 | 0.046 | <0.001 |  |  |  | 0.372 | 0.036 |
| [15] | SOB volume | 0.483 | 0.115 | $<0.001$ | 1.679 | 0.102 | $<0.001$ | 1.282 | 0.121 | $<0.001$ | 0.987 | 2551 |
|  | SOB density | 527.902 | 43.144 | $<0.001$ | 0.087 | 0.032 | 0.008 | -0.124 | 0.027 | $<0.001$ | 0.137 | 5702 |
|  | R/S ratio | 0.371 | 0.091 | $<0.001$ | 0.142 | 0.097 | 0.145 | -0.534 | 0.076 | $<0.001$ | 0.381 | 0.035 |



Fig. 3. Relationship of height a), dry mass of stem over bark b), dry mass of branches c), dry mass of foliage d), dry mass of roots e) and dry mass of the whole tree f) to stem base diameter $d_{0}$ of Common beech.


Fig. 4. Relationship of the component proportion a), volume of stem over bark b), basic density of stem over bark c), $R / S$ ratio d) to stem base diameter $d_{0}$ of Common beech.

### 4.2. Norway spruce

Norway spruce (Picea abies [L.] H. Karst.) is the main and the most frequent coniferous tree species in the forests of Slovakia. Only a single tree species, the most common broadleaved species, i.e. beech, is more frequent. Ecological optimum of spruce is in $7^{\text {th }}$ spruce fvz, where it creates naturally homogeneous stands. It is an important tree species in the forest biotopes of the national and European significance: Ls7.3 Bog spruce forests (*91D0 Bog woodland) Ls9.1 Bilberry spruce forests, Ls9.2 Spruce forests with tall herbs, and Ls9.3 Waterlogged spruce forests, all representing NATURA 2000 biotope 9410 Addophilous Picea forests of the montane to alpine levels (Vaccinio-Piceetea).

On the base of the processed NFIM2 SR data, the minimum and maximum elevations at which spruce occurred were 114 m and $1,676 \mathrm{~m}$ a.s.l., respectively, while on average it most frequently grew at elevations $800-900 \mathrm{~m}$ a.s.l. As a commercially important tree species it occurs in all forest vegetation zones from $1^{\text {st }}$ oak zone up to $8^{\text {th }}$ dwarf pine zone. Naturally, it is distributed at higher elevations from $5^{\text {th }}$ fir-beech forest vegetation zone. It grew at a reduced area of $415 \pm 32$ thousand ha, and occurred at $45 \%$ of the forested inventory plots.

The biomass regression models were derived from the set of 154 spruce individuals. They were taken from seven sites (see Fig. 5) located in the orographic units of Polana (sites 1 and 2), Slovenské rudohorie ( 3 and 4) and Nízke Tatry ( $5,6,7$ ). The samples represented the individuals with $d_{0}$ diameters from 1.55 mm to 97.7 mm , and heights from 0.11 m to 5.3 m (Table 5, Fig. 6a). The dry mass of the whole trees ranged from 0.53 g do $8,757.2 \mathrm{~g}$, and the stem volume ranged from $0.18 \mathrm{~cm}^{3}$ to $8,694.4 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 6. The volume of stem over bark, its density, as well as the root-shoot ratio are presented in Table 7. The regression models, scatter plots, and fitted regression curves were summarised or visualised in a similar way as in the case of Common beech (Chapter 4.1.). More detailed comments on the biomass of the individual components and their proportions of the total tree biomass are presented in Chapter 4.12 (Inter-species comparison of biomass characteristics). The mentioned chapter contains also the interpretations of the volume and density of stem over bark, or of the root-shoot ratio.


Fig. 5. Map of sample sites of Norway spruce and its distribution in the forests of Slovakia.

Table 5. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25.p), 75-percentile (75.p) and skewness of diameter ( $d_{0}$ ), tree height $(h)$, biomass of stem over bark (SOB), foliage biomass (foliage), branch biomass (branches), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {SOB }}$ ) of individual trees.

| Norway spruce |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | 25.p | 75.p | Skewness |
| $d_{0}(\mathrm{~mm})$ | 154 | 23.15 | 22.30 | 1.55 | 97.7 | 7.30 | 33.35 | 1.42 |
| $h(\mathrm{~m})$ | 154 | 1.29 | 1.17 | 0.11 | 5.3 | 0.35 | 2.15 | 1.09 |
| SOB (g) | 152 | 272.96 | 506.38 | 0.20 | 3158.5 | 5.85 | 287.00 | 2.84 |
| Foliage (g) | 151 | 229.70 | 425.70 | 0.03 | 2402.5 | 8.40 | 269.00 | 2.88 |
| Branches (g) | 151 | 175.16 | 353.42 | 0.04 | 2272.0 | 4.30 | 182.00 | 3.49 |
| Roots (g) | 151 | 128.34 | 214.40 | 0.23 | 1090.0 | 5.52 | 135.00 | 2.34 |
| Aboveground (g) | 148 | 675.58 | 1274.25 | 0.27 | 7833.0 | 19.35 | 713.75 | 2.97 |
| Whole tree (g) | 145 | 807.24 | 1490.44 | 0.53 | 8757.2 | 24.86 | 795.69 | 2.83 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 154 | 649.73 | 1300.94 | 0.18 | 8694.4 | 7.85 | 652.90 | 3.14 |

Table 6. Norway spruce, $b_{0}, b_{1}$, $b_{2}$ regression coefficients, their standard errors (S.E.), p-values ( $P$ ), coefficient of determination ( $R^{2}$ ), mean square error (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -3.084 | 0.069 | $<0.001$ | 2.459 | 0.024 | $<0.001$ |  |  |  | 0.986 | 0.090 | 1.046 | 0.324 |
|  | Branches | -3.452 | 0.099 | $<0.001$ | 2.482 | 0.035 | $<0.001$ |  |  |  | 0.971 | 0.185 | 1.093 | 0.488 |
|  | Foliage | -2.589 | 0.099 | $<0.001$ | 2.317 | 0.035 | <0.001 |  |  |  | 0.967 | 0.185 | 1.082 | 0.397 |
|  | Roots | -2.460 | 0.070 | $<0.001$ | 2.114 | 0.025 | <0.001 |  |  |  | 0.980 | 0.095 | 1.046 | 0.309 |
|  | Aboveground part | -1.842 | 0.069 | $<0.001$ | 2.398 | 0.024 | $<0.001$ |  |  |  | 0.985 | 0.088 | 1.043 | 0.296 |
|  | Whole tree | -1.432 | 0.060 | $<0.001$ | 2.333 | 0.021 | $<0.001$ |  |  |  | 0.988 | 0.067 | 1.033 | 0.258 |
| [10] | Stem over bark | 4.010 | 0.035 | $<0.001$ | 2.398 | 0.033 | $<0.001$ |  |  |  | 0.972 | 0.178 | 1.092 | 0.484 |
|  | Branches | 3.704 | 0.056 | $<0.001$ | 2.376 | 0.054 | $<0.001$ |  |  |  | 0.930 | 0.453 | 1.245 | 0.921 |
|  | Foliage | 4.090 | 0.058 | $<0.001$ | 2.200 | 0.055 | <0.001 |  |  |  | 0.914 | 0.486 | 1.232 | 0.816 |
|  | Roots | 3.640 | 0.052 | $<0.001$ | 2.004 | 0.049 | $<0.001$ |  |  |  | 0.919 | 0.386 | 1.198 | 0.748 |
|  | Aboveground part | 5.072 | 0.047 | $<0.001$ | 2.307 | 0.044 | <0.001 |  |  |  | 0.949 | 0.308 | 1.160 | 0.684 |
|  | Whole tree | 5.304 | 0.047 | $<0.001$ | 2.239 | 0.044 | <0.001 |  |  |  | 0.948 | 0.300 | 1.157 | 0.676 |
| [11] | Stem over bark | -0.469 | 0.201 |  | 1.555 | 0.070 | <0.001 | 0.913 | 0.068 | $<0.001$ | 0.994 | 0.041 | 1.020 | 0.205 |
|  | Branches | -2.553 | 0.417 | $<0.001$ | 2.171 | 0.144 | $<0.001$ | 0.313 | 0.141 | 0.028 | 0.972 | 0.180 | 1.089 | 0.460 |
|  | Foliage | -2.487 | 0.426 | $<0.001$ | 2.282 | 0.147 | <0.001 | 0.036 | 0.144 | 0.805 | 0.967 | 0.186 | 1.082 | 0.396 |
|  | Roots | -2.869 | 0.304 | $<0.001$ | 2.254 | 0.105 | <0.001 | -0.142 | 0.103 | 0.169 | 0.980 | 0.094 | 1.045 | 0.306 |
|  | Aboveground part | -0.696 | 0.281 | 0.014 | 2.002 | 0.097 | <0.001 | 0.400 | 0.095 | <0.001 | 0.987 | 0.079 | 1.037 | 0.274 |
|  | Whole tree | -0.579 | 0.251 | 0.023 | 2.039 | 0.087 | <0.001 | 0.297 | 0.085 | $<0.001$ | 0.989 | 0.062 | 1.030 | 0.248 |

Table 7. Norway spruce, $b_{0}, b_{1}$, $b_{2}$ regression coefficients, their standard errors (S.E.), p-values ( $P$ ), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE) for equations [13] - [15].

| Eq. | Dependent variable | $\mathrm{b}_{0}$ | S.E. | P | $\mathrm{b}_{1}$ | S.E. | P | $\mathrm{b}_{2}$ | S.E. | P |
| :--- | :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | SOB volume | 0.500 | 0.171 | 0.004 | 2.068 | 0.079 | $<0.001$ |  |  |  |
| $[13]$ | SOB density | 1205.436 | 37.282 | $<0.001$ | -0.254 | 0.013 | $<0.001$ |  | 0.913 |  |
|  | R/S ratio | 0.630 | 0.038 | $<0.001$ | -0.331 | 0.027 | $<0.001$ | 148956 |  |  |
|  | SOB volume | 108.084 | 9.163 | $<0.001$ | 2.649 | 0.061 | $<0.001$ |  | 0.730 |  |
| $[14]$ | SOB density | 577.291 | 8.493 | $<0.001$ | -0.254 | 0.012 | $<0.001$ | 382 |  |  |
|  | R/S ratio | 0.244 | 0.008 | $<0.001$ | -0.316 | 0.027 | $<0.001$ |  | 0.493 | 0.008 |
|  | SOB volume | 6.194 | 0.919 | $<0.001$ | 0.968 | 0.045 | $<0.001$ | 1.709 | 0.051 | $<0.001$ |
| $[15]$ | SOB density | 613.631 | 89.212 | $<0.001$ | -0.021 | 0.050 | 0.674 | -0.234 | 0.049 | $<0.001$ |
|  | R/S ratio | 0.329 | 0.102 | 0.002 | -0.105 | 0.108 | 0.335 | -0.220 | 0.106 | 0.040 |



Fig. 6. Relationship of height a), dry mass of stem over bark b), dry mass of branches c ), dry mass of needles d), dry mass of roots e) and dry mass of the whole tree f) to stem base diameter $d_{0}$ of Norway spruce.
Note: See also the comment on Fig. 6b-6f, which deals with the fitting of the scatter plot (placed at the end of this chapter, or more detailed information at the end of Chapter 5.).


Fig. 7. Relationship of the dry mass component proportion a), volume of stem over bark b), basic density of stem over bark c), and $R / S$ ratio d) to stem base diameter $d_{0}$ of Norway spruce.

## Note to fitting the scatter plot:

We would like to point out at the deviation of the fitted regression curves from the actual distribution of the observations in the scatter plot in a particular interval of the values. It can be seen in the examples of the relationships of the biomass of some spruce components, namely stem outside bark, branches, foliage, roots, and the whole tree dry mass, to $d_{0}$ diameter for the thickest trees (see Fig. 6b-6f). A similar situation can be observed also for some other tree species presented in the following chapters. A more detailed description of this phenomenon is presented at the end of Chapter 5 (Knowledge synthesis and conclusion).

### 4.3. European hornbeam

European hornbeam (Carpinus betulus L.) is a commercially less important tree species, particularly from the point of assortment quality. It is most frequently used as fuel-wood. Hornbeam occurs as a secondary stand-forming tree species in lower vegetation zones, mainly in $2^{\text {nd }}$ beech-oak and in $3^{\text {rd }}$ oak-beech zones, where it is most abundant, although from all tree species
it reaches only a proportion of $20 \%$. It often forms the under-storey of oak and beech stands. From the silvicultural point of view, it participates in the formation of high-quality assortments of these species. It is an important element of the forest biotopes of European and national significance: Ls 2.2 Pannonian oak-hornbeam forests (NATURA 2000 *91G0 Pannonic woods with Quercus petraea and Carpinus betulus), Ls 2.31 Oak-hornbeam forests with lime (9170 GalioCarpinetum oak-hornbeam forests), Ls 2.33 Oak-hornbeam forests with lime.

Hornbeam occurred at elevations between 100 m (the lowest occurrence) and 809 m a.s.l. (the highest occurrence), most frequently between $350-450 \mathrm{~m}$ a.s.l. (NFIM2 SR data). It is the fourth most common tree species in the forests of Slovakia from the point of its occurrence, while from the point of the spatial proportion it is the third, and when considering the stand stock it is the $6^{\text {th }}$ tree species in the tree species ranking. It grew at a reduced area of $187 \pm 22$ thousand ha, and occurred at $32 \%$ of the forested inventory plots.

The biomass regression models were derived from the set of 200 hornbeam trees. They were taken from eight sites (see Fig. 8), which were located in the orographic units of Malé Karpaty (site 1), Tribeč (2), Štiavnické vrchy (3), Kremnické vrchy (4), Krupinská planina (5), Slovenské rudohorie (6 and 7) and Nízke Beskydy (8). The samples represented the individuals with $d_{0}$ diameters from 0.90 mm to 81.20 mm , and heights from 0.07 m to 7.56 m (Table 8, Fig. 9a). The dry mass of the whole trees ranged from 0.10 g do $5,399.22 \mathrm{~g}$, and the stem volume ranged from $0.02 \mathrm{~cm}^{3}$ to $8,402.30 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 9. In contrast to beech and spruce, for hornbeam we also derived the dry mass of the stem under bark and stem bark.

The regression models, scatter plots, and fitted regression curves were summarised or visualised in a similar way as in the case of Common beech (Chapter 4.1.). Unlike for beech, we graphically presented also bark density. Further comments on the biomass of individual components and their proportions in the total tree biomass are presented in Chapter 4.12. (Inter-species comparison of biomass characteristics).


Fig. 8. Map of sample sites of European hornbeam and its distribution in the forests of Slovakia.

Table 8. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25.p), 75-percentile (75.p) and skewness of diameter $\left(d_{0}\right)$, tree height $(h)$, biomass of stem over bark (SOB), biomass of stem under bark (SUB), foliage biomass (foliage), branch biomass (branches), bark biomass (bark), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {SOB }}$ ) of individual trees.

| European hornbeam |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | 25.p | 75.p | Skewness |
| $d_{0}(\mathrm{~mm})$ | 200 | 17.84 | 13.94 | 0.90 | 81.20 | 7.90 | 23.03 | 1.70 |
| $h(\mathrm{~m})$ | 200 | 2.68 | 1.83 | 0.07 | 7.56 | 1.10 | 3.88 | 0.65 |
| SOB (g) | 199 | 299.34 | 656.22 | 0.03 | 4429.74 | 9.60 | 223.62 | 3.78 |
| SUB (g) | 200 | 263.27 | 596.14 | 0.03 | 4038.19 | 8.01 | 191.30 | 3.84 |
| Foliage (g) | 196 | 31.18 | 59.22 | 0.03 | 346.34 | 1.78 | 35.67 | 3.37 |
| Branches (g) | 199 | 56.10 | 138.84 | 0.00 | 953.15 | 1.65 | 43.15 | 4.28 |
| Bark (g) | 199 | 34.82 | 59.61 | 0.002 | 391.55 | 2.67 | 35.15 | 3.27 |
| Roots (g) | 197 | 78.40 | 185.46 | 0.04 | 1473.50 | 5.83 | 60.50 | 4.61 |
| Aboveground (g) | 194 | 346.45 | 713.01 | 0.06 | 4329.52 | 12.70 | 284.95 | 3.66 |
| Whole tree (g) | 191 | 414.91 | 858.13 | 0.10 | 5399.22 | 19.44 | 341.31 | 3.68 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 200 | 552.55 | 1224.93 | 0.02 | 8402.30 | 17.31 | 448.09 | 3.92 |

Table 9. European hornbeam, $b_{0}, b_{1}$, $b_{2}$ regression coefficients, their standard errors (S.E.), p-values ( $P$ ), coefficients of determination ( $R^{2}$ ), mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -3.402 | 0.081 | $<0.001$ | 2.809 | 0.030 | <0.001 |  |  |  | 0.978 | 0.124 | 1.066 | 0.415 |
|  | Stem inside bark | -3.881 | 0.086 | <0.001 | 2.904 | 0.032 | <0.001 |  |  |  | 0.977 | 0.140 | 1.075 | 0.447 |
|  | Branches | -5.573 | 0.138 | <0.001 | 2.924 | 0.050 | $<0.001$ |  |  |  | 0.946 | 0.295 | 1.143 | 0.596 |
|  | Foliage | -4.127 | 0.165 | <0.001 | 2.354 | 0.061 | <0.001 |  |  |  | 0.884 | 0.497 | 1.225 | 0.700 |
|  | Bark | -4.090 | 0.085 | $<0.001$ | 2.440 | 0.031 | $<0.001$ |  |  |  | 0.971 | 0.110 | 1.056 | 0.356 |
|  | Roots | -3.105 | 0.078 | <0.001 | 2.301 | 0.029 | $<0.001$ |  |  |  | 0.968 | 0.138 | 1.069 | 0.410 |
|  | Aboveground part | -2.919 | 0.068 | $<0.001$ | 2.745 | 0.025 | <0.001 |  |  |  | 0.984 | 0.086 | 1.045 | 0.334 |
|  | Whole tree | -2.389 | 0.062 | <0.001 | 2.640 | 0.023 | $<0.001$ |  |  |  | 0.986 | 0.068 | 1.034 | 0.277 |
| [10] | Stem over bark | 2.281 | 0.048 | <0.001 | 2.407 | 0.041 | $<0.001$ |  |  |  | 0.946 | 0.308 | 1.163 | 0.678 |
|  | Stem inside bark | 1.988 | 0.050 | <0.001 | 2.487 | 0.044 | $<0.001$ |  |  |  | 0.943 | 0.347 | 1.185 | 0.730 |
|  | Branches | 0.419 | 0.089 | <0.001 | 2.392 | 0.079 | $<0.001$ |  |  |  | 0.826 | 0.950 | 1.525 | 1.505 |
|  | Foliage | 0.733 | 0.092 | <0.001 | 1.848 | 0.080 | $<0.001$ |  |  |  | 0.732 | 1.153 | 1.619 | 1.570 |
|  | Bark | 0.831 | 0.038 | <0.001 | 2.111 | 0.033 | <0.001 |  |  |  | 0.846 | 0.576 | 1.345 | 1.224 |
|  | Roots | 1.613 | 0.066 | <0.001 | 1.867 | 0.057 | <0.001 |  |  |  | 0.955 | 0.195 | 1.099 | 0.492 |
|  | Aboveground part | 2.668 | 0.053 | <0.001 | 2.301 | 0.046 | <0.001 |  |  |  | 0.929 | 0.377 | 1.203 | 0.778 |
|  | Whole tree | 2.998 | 0.055 | <0.001 | 2.186 | 0.048 | $<0.001$ |  |  |  | 0.915 | 0.404 | 1.222 | 0.843 |
| [11] | Stem over bark | -1.941 | 0.121 | <0.001 | 1.922 | 0.058 | <0.001 | 0.909 | 0.051 | $<0.001$ | 0.991 | 0.054 | 1.029 | 0.270 |
|  | Stem inside bark | -1.486 | 0.106 | <0.001 | 1.841 | 0.051 | $<0.001$ | 0.896 | 0.044 | $<0.001$ | 0.993 | 0.041 | 1.021 | 0.220 |
|  | Branches | -5.514 | 0.291 | <0.001 | 2.893 | 0.140 | $<0.001$ | 0.029 | 0.123 | 0.816 | 0.946 | 0.296 | 1.143 | 0.594 |
|  | Foliage | -5.423 | 0.366 | <0.001 | 3.008 | 0.177 | <0.001 | -0.599 | 0.152 | $<0.001$ | 0.893 | 0.463 | 1.211 | 0.696 |
|  | Bark | -2.026 | 0.106 | <0.001 | 1.396 | 0.051 | <0.001 | 0.965 | 0.045 | $<0.001$ | 0.971 | 0.109 | 1.055 | 0.353 |
|  | Roots | -3.348 | 0.174 | <0.001 | 2.424 | 0.084 | <0.001 | -0.114 | 0.073 | 0.121 | 0.991 | 0.041 | 1.018 | 0.183 |
|  | Aboveground part | -1.603 | 0.120 | <0.001 | 2.083 | 0.058 | $<0.001$ | 0.605 | 0.050 | $<0.001$ | 0.991 | 0.049 | 1.025 | 0.239 |
|  | Whole tree | -1.445 | 0.121 | <0.001 | 2.165 | 0.058 | <0.001 | 0.433 | 0.050 | $<0.001$ | 0.990 | 0.049 | 1.024 | 0.226 |

Table 10. European hornbeam, $b_{o}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination ( $R^{2}$ ), mean square errors (MSE) for equations [13] - [15].

| Eq. | Dependent variable | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [13] | SOB volume | 0.837 | 0.156 | $<0.001$ | 2.091 | 0.044 | $<0.001$ |  |  |  | 0.944 | 122079 |
|  | SUB volume | 0.644 | 0.123 | <0.001 | 2.133 | 0.045 | <0.001 |  |  |  | 0.944 | 102312 |
|  | Bark volume | 0.333 | 0.064 | <0.001 | 1.719 | 0.047 | $<0.001$ |  |  |  | 0.899 | 1526 |
|  | SOB density | 880.710 | 26.524 | <0.001 | -0.155 | 0.012 | <0.001 |  |  |  | 0.444 | 6885 |
|  | SUB density | 835.631 | 26.790 | <0.001 | -0.148 | 0.012 | <0.001 |  |  |  | 0.391 | 6913 |
|  | Bark density | 910.292 | 33.268 | $<0.001$ | -0.109 | 0.014 | <0.001 |  |  |  | 0.230 | 15234 |
|  | Bark mass proportion | 54.421 | 0.818 | $<0.001$ | -0.382 | 0.006 | $<0.001$ |  |  |  | 0.631 | 21.17 |
|  | Bark volume proportion | 50.321 | 1.460 | $<0.001$ | -0.406 | 0.014 | $<0.001$ |  |  |  | 0.793 | 14.84 |
|  | R/S ratio | 0.898 | 0.055 | $<0.001$ | -0.437 | 0.029 | $<0.001$ |  |  |  | 0.501 | 0.019 |
| [14] | SOB volume | 0.432 | 0.155 | 0.006 | 4.844 | 0.184 | $<0.001$ |  |  |  | 0.902 | 211954 |
|  | SUB volume | 0.297 | 0.112 | 0.009 | 4.991 | 0.193 | $<0.001$ |  |  |  | 0.901 | 181452 |
|  | Bark volume | 0.412 | 0.111 | $<0.001$ | 3.594 | 0.141 | <0.001 |  |  |  | 0.877 | 1857 |
|  | SOB density | 645.813 | 6.400 | <0.001 | -0.139 | 0.009 | <0.001 |  |  |  | 0.522 | 5915 |
|  | SUB density | 623.866 | 6.327 | $<0.001$ | -0.140 | 0.009 | <0.001 |  |  |  | 0.508 | 5587 |
|  | Bark density | 726.468 | 10.301 | $<0.001$ | -0.087 | 0.012 | <0.001 |  |  |  | 0.196 | 15911 |
|  | Bark mass proportion | 23.889 | 0.345 | <0.001 | -0.247 | 0.012 | <0.001 |  |  |  | 0.662 | 19.52 |
|  | Bark volume proportion | 21.969 | 0.307 | $<0.001$ | -0.331 | 0.011 | <0.001 |  |  |  | 0.783 | 15.49 |
|  | $\mathrm{R} /$ S ratio | 0.371 | 0.010 | $<0.001$ | -0.379 | 0.021 | $<0.001$ |  |  |  | 0.595 | 0.015 |
| [15] | SOB volume | 0.651 | 0.070 | $<0.001$ | 1.313 | 0.037 | $<0.001$ | 1.845 | 0.082 | $<0.001$ | 0.986 | 30603 |
|  | SUB volume | 0.474 | 0.051 | $<0.001$ | 1.345 | 0.036 | <0.001 | 1.895 | 0.082 | $<0.001$ | 0.987 | 24272 |
|  | Bark volume | 0.390 | 0.063 | <0.001 | 1.015 | 0.068 | <0.001 | 1.469 | 0.136 | $<0.001$ | 0.939 | 923 |
|  | SOB density | 644.316 | 42.885 | <0.001 | 0.001 | 0.032 | 0.978 | -0.136 | 0.026 | <0.001 | 0.511 | 6078 |
|  | SUB density | 519.312 | 36.561 | <0.001 | 0.088 | 0.033 | 0.099 | -0.211 | 0.028 | <0.001 | 0.525 | 5423 |
|  | Bark density | 964.983 | 91.240 | <0.001 | -0.138 | 0.046 | 0.003 | 0.026 | 0.039 | 0.506 | 0.232 | 15276 |
|  | Bark mass proportion | 45.331 | 4.604 | <0.001 | -0.313 | 0.050 | <0.001 | 0.006 | 0.041 | 0.876 | 0.716 | 16.45 |
|  | Bark volume proportion | 35.572 | 3.644 | <0.001 | -0.234 | 0.050 | <0.001 | -0.145 | 0.041 | <0.001 | 0.805 | 14.01 |
|  | R/S ratio | 0.217 | 0.046 | <0.001 | 0.256 | 0.099 | 0.011 | -0.581 | 0.081 | $<0.001$ | 0.609 | 0.015 |



Fig. 9. Relationship of tree height a) and $d_{1.3}$ diameter b) to stem base diameter $d_{0}$ of European hornbeam.


Fig. 10. Relationship of dry mass of stem over bark a), dry mass of branches b), dry mass of foliage c), dry mass of roots d) and dry mass of the whole tree e) and proportion of individual tree components f) to stem base diameter $d_{0}$ of European hornbeam.


Fig. 11. Relationship of the basic density of stem over bark a), basic bark density b), mass proportion of bark in SOB mass c), volume proportion of bark in $S O B$ volume d) volume of stem over bark e), and $R / S$ ratio f) to stem base diameter $d_{0}$ of European hornbeam.

### 4.4. Sessile oak

Sessile oak (Quercus petraea [Matt.] Liebl.) is a commercially important tree species of the forests at lower elevations. It predominantly occurs at lower forest vegetation zones from $1^{\text {st }}$ oak to $3^{\text {rd }}$ oak-beech zones, and sporadically also in $4^{\text {th }}$ beech zone, but it does not grow at higher elevations. It is an important tree species in several forest biotopes of European and national significance: Ls2.2 Pannonian oak-hornbeam forests (NATURA 2000*91G0 Pannonic woods with Quercus petraea and Carpinus betulus), Ls2.31 Oak-hornbeam forests with lime (9170 Galio-Carpinetum oak-hornbeam forests), Ls2.33 Oak-hornbeam forests with lime, Ls3.2 Thermophillic Pontic-Pannonian oak forests on loess and sand, Ls3.3 Oak cinquefoil forests, Ls3.52 Xerophilous and Acidophilous oak forests (*9110 Euro-Siberian steppic woods with Quercus spp.), Ls3.4 Sessile oak-Turkey oak forests (91M0 Pannonian-Balkanic turkey oak-sessile oak forests).

On the base of the processed NFIM2 SR data, the minimum and maximum elevations at which oak occurred were 109 m and 833 m a.s.l., respectively, while most frequently it grew at elevations $300-400 \mathrm{~m}$ a.s.l. With regard to the stand stock, it is the third most common tree species in the forests of Slovakia, from the point of the occupied area it is $4^{\text {th }}$, and from the point of occurrence it is $5^{\text {th }}$ in tree species ranking. It grew at a reduced area of $160 \pm 21$ thousand ha, and occurred at $28 \%$ of the forested inventory plots.

The biomass regression models of the Sessile oak were derived from 162 individuals. They were taken from eight sites (see Fig. 12), which were located in the orographic units of the Štiavnické vrchy (sites 1 and 2 ) and Kremnické vrchy (3, 4, 5, 6, 7). The samples represented the individuals with $d_{0}$ diameters from 4.45 mm to 88.75 mm , and heights from 0.28 m to 6.64 m (Table 11, Fig. 13a). The dry mass of the whole trees ranged from 5.17 g do $8,790.0 \mathrm{~g}$, and the stem volume ranged from $1.46 \mathrm{~cm}^{3}$ to $8,300.0 \mathrm{~cm}^{3}$.


Fig. 12. Map of sample sites of Sessile oak and its distribution in the forests of Slovakia.
The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 12. Similarly, the volume of stem outside bark, its density, as well as the root-shoot ratio were derived (Table 13).

The regression models, scatter plots, and fitted regression curves were summarised or visualised in a similar way as in the case of Common beech (Chapter 4.1.). Further comments on the biomass of individual components and their proportions of the total tree biomass are presented in Chapter 4.12. (Inter-species comparison of biomass characteristics).

Table 11. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25. p), 75-percentile (75. p) and skewness of diameter $\left(d_{0}\right)$, tree height ( $h$ ), biomass of stem over bark (SOB), foliage biomass (foliage), branch biomass (branches), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {SOB }}$ ) of individual trees.

| Sessile oak |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | 25.p | 75.p | Skewness |
| $\overline{d_{0}(\mathrm{~mm})}$ | 162 | 26.85 | 20.77 | 4.45 | 88.75 | 11.00 | 40.25 | 1.22 |
| $h(m)$ | 162 | 2.21 | 1.80 | 0.28 | 6.64 | 0.69 | 3.54 | 0.76 |
| SOB (g) | 140 | 541.07 | 1146.41 | 1.30 | 5784.0 | 11.65 | 443.50 | 2.86 |
| Foliage (g) | 140 | 54.36 | 115.40 | 0.20 | 546.0 | 1.60 | 38.00 | 2.76 |
| Branches (g) | 140 | 102.43 | 227.41 | 0.09 | 1132.0 | 2.15 | 65.75 | 3.03 |
| Roots (g) | 162 | 217.20 | 328.98 | 3.03 | 1435.0 | 22.25 | 252.20 | 2.22 |
| Aboveground (g) | 140 | 697.86 | 1485.43 | 2.14 | 7355.0 | 14.66 | 541.25 | 2.87 |
| Whole tree (g) | 140 | 876.25 | 1794.34 | 5.17 | 8790.0 | 33.75 | 705.21 | 2.83 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 162 | 934.07 | 1710.14 | 1.46 | 8300.0 | 20.28 | 900.52 | 2.48 |

Table 12. Sessile oak, $b_{0}, b_{1}, b_{2}$, regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -4.311 | 0.224 | <0.001 | 2.959 | 0.071 | $<0.001$ |  |  |  | 0.942 | 0.240 | 1.121 | 0.544 |
|  | Branches | -6.025 | 0.215 | $<0.001$ | 2.963 | 0.068 | <0.001 |  |  |  | 0.947 | 0.221 | 1.107 | 0.496 |
|  | Foliage | -5.954 | 0.221 | <0.001 | 2.768 | 0.070 | <0.001 |  |  |  | 0.936 | 0.234 | 1.106 | 0.459 |
|  | Roots | -1.954 | 0.150 | <0.001 | 2.066 | 0.046 | $<0.001$ |  |  |  | 0.941 | 0.124 | 1.065 | 0.401 |
|  | Aboveground part | -3.952 | 0.206 | <0.001 | 2.931 | 0.065 | <0.001 |  |  |  | 0.950 | 0.203 | 1.099 | 0.469 |
|  | Whole tree | -2.612 | 0.175 | $<0.001$ | 2.646 | 0.005 | $<0.001$ |  |  |  | 0.956 | 0.146 | 1.070 | 0.382 |
| [10] | Stem over bark | 3.471 | 0.045 | $<0.001$ | 2.516 | 0.046 | $<0.001$ |  |  |  | 0.965 | 0.147 | 1.074 | 0.419 |
|  | Branches | 1.839 | 0.086 | <0.001 | 2.388 | 0.089 | <0.001 |  |  |  | 0.871 | 0.535 | 1.271 | 0.896 |
|  | Foliage | 1.370 | 0.074 | <0.001 | 2.269 | 0.077 | <0.001 |  |  |  | 0.891 | 0.398 | 1.190 | 0.700 |
|  | Roots | 3.511 | 0.053 | <0.001 | 1.719 | 0.051 | <0.001 |  |  |  | 0.898 | 0.213 | 1.112 | 0.553 |
|  | Aboveground part | 3.769 | 0.050 | $<0.001$ | 2.468 | 0.052 | $<0.001$ |  |  |  | 0.954 | 0.186 | 1.095 | 0.483 |
|  | Whole tree | 4.369 | 0.051 | <0.001 | 2.207 | 0.053 | $<0.001$ |  |  |  | 0.942 | 0.189 | 1.099 | 0.503 |
| [11] | Stem over bark | -0.066 | 0.170 | 0.698 | 1.324 | 0.063 | $<0.001$ | 1.491 | 0.053 | $<0.001$ | 0.993 | 0.029 | 1.014 | 0.171 |
|  | Branches | -4.263 | 0.432 | <0.001 | 2.284 | 0.161 | $<0.001$ | 0.619 | 0.135 | $<0.001$ | 0.956 | 0.186 | 1.092 | 0.473 |
|  | Foliage | -3.577 | 0.412 | <0.001 | 1.852 | 0.153 | <0.001 | 0.835 | 0.129 | <0.001 | 0.954 | 0.169 | 1.077 | 0.399 |
|  | Roots | -0.226 | 0.289 | 0.436 | 1.398 | 0.107 | <0.001 | 0.611 | 0.091 | <0.001 | 0.956 | 0.092 | 1.047 | 0.333 |
|  | Aboveground part | -0.201 | 0.194 | 0.303 | 1.486 | 0.072 | <0.001 | 1.317 | 0.061 | <0.001 | 0.991 | 0.038 | 1.019 | 0.199 |
|  | Whole tree | 0.389 | 0.201 | 0.056 | 1.490 | 0.075 | $<0.001$ | 1.054 | 0.063 | $<0.001$ | 0.989 | 0.040 | 1.020 | 0.209 |

Table 13. Sessile oak, $b_{0}, b_{1}, b_{2}$, regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE) for equations [13] - [15].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :--- | :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | SOB volume | 0.235 | 0.062 | $<0.001$ | 2.325 | 0.062 | $<0.001$ |  |  | 0.955 | 133632 |  |
| $[13]$ | SOB density | 896.210 | 37.848 | $<0.001$ | -0.065 | 0.015 | $<0.001$ |  |  | 0.128 | 8334 |  |
|  | R/S ratio | 9.135 | 1.228 | $<0.001$ | -0.888 | 0.059 | $<0.001$ |  |  | 0.698 | 0.113 |  |
|  | SOB volume | 20.139 | 5.424 | $<0.001$ | 3.144 | 0.156 | $<0.001$ |  |  | 0.882 | 348588 |  |
| $[14]$ | SOB density | 751.731 | 8.245 | $<0.001$ | -0.038 | 0.012 | 0.002 |  |  | 0.070 | 8890 |  |
|  | R/S ratio | 0.828 | 0.030 | $<0.001$ | -0.772 | 0.044 | $<0.001$ |  |  | 0.780 | 0.082 |  |
|  | SOB volume | 0.493 | 0.082 | $<0.001$ | 1.676 | 0.052 | $<0.001$ | 1.205 | 0.074 | $<0.001$ | 0.984 | 46900 |
| $[15]$ | SOB density | 1208.970 | 136.502 | $<0.001$ | -0.177 | 0.042 | $<0.001$ | 0.095 | 0.034 | 0.005 | 0.177 | 7929 |
|  | R/S ratio | 1.296 | 0.377 | 0.001 | -0.167 | 0.108 | 0.124 | -0.651 | 0.089 | $<0.001$ | 0.783 | 0.082 |



Fig. 13. Relationship of height a), dry mass of stem outside bark b), dry mass of branches c), dry mass of foliage d) to stem base diameter $d_{0}$ of Sessile oak.


Fig. 14. Relationship of root dry mass a) total tree dry mass b), proportion of individual components c), volume of stem over bark d), reduced stem density over bark e), and $R / S$ ratiof) to stem base diameter $d_{0}$ of Sessile oak.

### 4.5. Scots pine

Scots pine (Pinus sylvestris L.) is a pioneer and a target tree species growing at a wide range of ecological conditions. It tolerates dry and nutrient-poor rocky soils, but it also grows on wet, waterlogged or even peat sites. It sporadically occurs in all forest vegetation zones, and dominates in the area of the Záhorská nížina (plain). It is particularly important in two biotopes: Ls7.2 Bog pine forests (NATURA 2000 *91D0 Bog woodland), and Ls6.2 Relict calcareous pine and larch forests (91Q0 Western Carpathian calcicolous Pinus sylvestris forests).

Pine occurred at elevations between 144 m (the lowest occurrence) and $1,285 \mathrm{~m}$ a.s.l. (the highest occurrence), most frequently between $200-300 \mathrm{~m}$ a.s.l. (NFIM2 SR data). From the point of its stand stock, it is the $4^{\text {th }}$ most common tree species in the forests of Slovakia, according to the area it is $5^{\text {th }}$, and according to its occurrence at the forested inventory plots it was $7^{\text {th }}$ most common tree species. It grew at a reduced area of $106 \pm 17$ thousand ha, and occurred at $17 \%$ of the forested inventory plots.

The biomass regression models of the Scots pine were derived from 175 individuals. The trees were taken from seven sites (see Fig. 15). The sites were located in the orographic units of the Biele Karpaty (sites 1 and 2), Vtáčnik (3), Štiavnické vrchy (4 and 5), and Levočské vrchy (6 and 7). The sample trees represented the individuals with $d_{0}$ diameters from 3.05 mm to 78.10 mm , and heights from 0.13 m to 4.5 m (Table 14, Fig. 16a). The dry mass of the whole trees ranged from 3.05 g do $4,977.0 \mathrm{~g}$, and the stem volume ranged from $0.72 \mathrm{~cm}^{3}$ to $5,619.7 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole pine tree are presented in Table 15. The volume of stem over bark, its density, as well as the root-shoot ratio were derived in a similar way (Table 16). The regression models, scatter plots, and fitted regression curves were visualised in a similar way as in the case of previous tree species. The text describing the biomass of individual components and their proportions in the total tree biomass are presented in Chapter 4.12. (Inter-species comparison of biomass characteristics).


Fig. 15. Map of sample sites of Scots pine and its distribution in the forests of Slovakia.

Table 14. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25. p), 75-percentile (75. p) and skewness of diameter ( $d_{\ell}$ ), tree height ( $h$ ), biomass of stem over bark (SOB), foliage biomass (foliage), branch biomass (branches), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {sob }}$ ) of individual trees.

|  | Scots pine |  |  |  |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | $25 . \mathrm{p}$ | 75.p | Skewness |
| $d_{0}(\mathrm{~mm})$ | 175 | 28.29 | 18.35 | 3.05 | 78.1 | 13.45 | 39.50 | 0.84 |
| $h(\mathrm{~m})$ | 175 | 1.63 | 1.02 | 0.13 | 4.5 | 0.75 | 2.27 | 0.70 |
| SOB $(\mathrm{g})$ | 175 | 315.47 | 438.97 | 0.50 | 2174.0 | 25.85 | 414.00 | 1.89 |
| Foliage $(\mathrm{g})$ | 175 | 175.43 | 219.16 | 1.45 | 1283.0 | 25.60 | 243.00 | 2.17 |
| Branches $(\mathrm{g})$ | 175 | 152.42 | 216.69 | 0.20 | 1077.0 | 13.00 | 200.00 | 2.12 |
| Roots $(\mathrm{g})$ | 175 | 76.55 | 101.98 | 0.50 | 535.0 | 8.00 | 100.00 | 2.15 |
| Aboveground $(\mathrm{g})$ | 175 | 643.31 | 858.13 | 2.25 | 4442.0 | 65.35 | 801.00 | 1.92 |
| Whole tree $(\mathrm{g})$ | 175 | 719.86 | 955.25 | 3.05 | 4977.0 | 72.40 | 900.50 | 1.93 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 175 | 798.17 | 1196.04 | 0.72 | 5619.7 | 50.76 | 918.51 | 2.08 |

Table 15. Scots pine, $b_{o}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | $\mathrm{S.E}$. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :--- | :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Stem over bark | -3.143 | 0.122 | $<0.001$ | 2.480 | 0.038 | $<0.001$ |  |  |  | 0.960 | 0.131 | 1.067 | 0.395 |  |
| [9] | Branches | -3.985 | 0.125 | $<0.001$ | 2.515 | 0.039 | $<0.001$ |  |  |  | 0.960 | 0.138 | 1.072 | 0.435 |
| Foliage | -1.961 | 0.144 | $<0.001$ | 2.014 | 0.045 | $<0.001$ |  |  |  | 0.920 | 0.183 | 1.089 | 0.456 |  |
| Roots | -3.628 | 0.130 | $<0.001$ | 2.236 | 0.041 | $<0.001$ |  |  |  | 0.946 | 0.149 | 1.079 | 0.479 |  |
| Aboveground part | -1.698 | 0.097 | $<0.001$ | 2.290 | 0.030 | $<0.001$ |  |  |  | 0.970 | 0.083 | 1.041 | 0.306 |  |
| Whole tree | -1.531 | 0.089 | $<0.001$ | 2.278 | 0.028 | $<0.001$ |  |  |  | 0.975 | 0.069 | 1.035 | 0.282 |  |
| Stem over bark | 3.958 | 0.043 | $<0.001$ | 2.389 | 0.057 | $<0.001$ |  |  |  | 0.912 | 0.293 | 1.156 | 0.648 |  |
| Branches | 3.246 | 0.062 | $<0.001$ | 2.304 | 0.081 | $<0.001$ |  |  |  | 0.824 | 0.601 | 1.323 | 1.003 |  |
| Foliage | 3.843 | 0.060 | $<0.001$ | 1.805 | 0.078 | $<0.001$ |  |  |  | 0.755 | 0.560 | 1.273 | 0.852 |  |
| Roots | 2.824 | 0.067 | $<0.001$ | 1.965 | 0.087 | $<0.001$ |  |  |  | 0.745 | 0.694 | 1.387 | 1.173 |  |
| [10] | Aboveground part | 4.880 | 0.050 | $<0.001$ | 2.134 | 0.065 | $<0.001$ |  |  |  | 0.862 | 0.388 | 1.200 | 0.725 |
| Whole tree | 5.013 | 0.051 | $<0.001$ | 2.108 | 0.066 | $<0.001$ |  |  |  | 0.854 | 0.402 | 1.209 | 0.752 |  |
| Stem over bark | -0.702 | 0.152 | $<0.001$ | 1.618 | 0.052 | $<0.001$ | 0.944 | 0.052 | $<0.001$ | 0.987 | 0.045 | 1.023 | 0.231 |  |
| Branches | -3.184 | 0.257 | $<0.001$ | 2.232 | 0.089 | $<0.001$ | 0.310 | 0.088 | $<0.001$ | 0.962 | 0.129 | 1.066 | 0.403 |  |
| Foliage | -1.901 | 0.307 | $<0.001$ | 1.993 | 0.106 | $<0.001$ | 0.024 | 0.105 | 0.822 | 0.920 | 0.184 | 1.089 | 0.457 |  |
| Roots | -4.105 | 0.273 | $<0.001$ | 2.405 | 0.094 | $<0.001$ | -0.184 | 0.093 | 0.050 | 0.947 | 0.146 | 1.078 | 0.487 |  |
| Aboveground part | -0.466 | 0.177 | 0.009 | 1.856 | 0.061 | $<0.001$ | 0.476 | 0.060 | $<0.001$ | 0.978 | 0.061 | 1.031 | 0.269 |  |
| Whole tree | -0.502 | 0.167 | 0.003 | 1.914 | 0.057 | $<0.001$ | 0.398 | 0.057 | $<0.001$ | 0.980 | 0.054 | 1.028 | 0.253 |  |

Table 16. Scots pine, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), $p$-values ( $P$ ), coefficients of determination ( $R^{2}$ ), mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [13] - [15].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :--- | :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | SOB volume | 0.235 | 0.062 | $<0.001$ | 2.325 | 0.062 | $<0.001$ |  |  | 0.955 | 133632 |  |
| $[13]$ | SOB density | 896.210 | 37.848 | $<0.001$ | -0.065 | 0.015 | $<0.001$ |  |  | 0.128 | 8334 |  |
|  | R/S ratio | 9.135 | 1.228 | $<0.001$ | -0.888 | 0.059 | $<0.001$ |  |  | 0.698 | 0.113 |  |
|  | SOB volume | 20.139 | 5.424 | $<0.001$ | 3.144 | 0.156 | $<0.001$ |  |  | 0.882 | 348588 |  |
| $[14]$ | SOB density | 751.731 | 8.245 | $<0.001$ | -0.038 | 0.012 | 0.002 |  |  | 0.070 | 8890 |  |
|  | R/S ratio | 0.828 | 0.030 | $<0.001$ | -0.772 | 0.044 | $<0.001$ |  |  | 0.780 | 0.082 |  |
|  | SOB volume | 0.421 | 0.063 | $<0.001$ | 1.961 | 0.040 | $<0.001$ | 0.735 | 0.034 | $<0.001$ | 0.983 | 24279 |
| $[15]$ | SOB density | 1045.979 | 86.230 | $<0.001$ | -0.251 | 0.029 | $<0.001$ | -0.012 | 0.028 | 0.656 | 0.740 | 3196 |
|  | R/S ratio | 0.029 | 0.010 | 0.005 | 0.550 | 0.117 | $<0.001$ | -0.715 | 0.115 | $<0.001$ | 0.201 | 0.004 |



Fig. 16. Relationship of height a), dry mass of stem over bark b), dry mass of branches $c$ ), dry mass of foliage $d$ ), dry mass of roots e) and dry mass of the whole tree f) to stem base diameter $d_{0}$ of Scots pine.


Fig. 17. Relationship of the component proportion a), volume of stem over bark b), basic density of stem over bark c), $R / S$ ratio d) to stem base diameter $d_{0}$ of Scots pine.

### 4.6. Sycamore

Sycamore (Acer pseudoplatanus L.) is the most common species of Acer genus. Unlike other more frequent tree species, it usually does not create homogeneous stands. The exceptions are the groups of sycamore stands at suitable and favourable sites, where sycamore is most competitive. Sycamore is a demanding tree species from the point of soil moisture and air humidity, as well as from the point of nutrient content in the soil. It usually occurs in small clusters, or as an admixture in stands, often on ridges and in screes, in the ends of valleys, and in ravines. Together with other valuable broadleaved species it increases biodiversity of forest ecosystems and has a positive impact on soil conditions. It is an important tree species in forest biotopes of European significance: Ls5.3 Mountainous sycamore-beech forests ( 9140 Medio-European subalpine beech woods with Acer and Rumex arifolius), and together with the lime tree, ash, and other valuable broadleaved species in Ls4 Lime-maple ravine forests ( 9180 Tilio-Acerion forests of slopes, scree and ravines). It occurs practically in all forest vegetation zones from $1^{\text {st }}$ oak up to $7^{\text {th }}$ spruce zones, and is dominant at scree sites.

According to the processed NFIM2 SR data, its lowest and highest occurrences were recorded at elevations of 112 m and $1,365 \mathrm{~m}$ a.s.l., respectively, and its most frequent occurrence between $700-800 \mathrm{~m}$ a.s.l. From the point of occurrence, it is the third most common tree species in the forests of Slovakia (after beech and spruce), considering its proportion in the area it is the $6^{\text {th }}$, and the proportion in stand stock it is $8^{\text {th }}$ in the species ranking. It grew at a reduced area of $79 \pm 15$ thousand ha, and occurred at $32 \%$ of the forested inventory plots.

The regression models of the sycamore were derived from the data gathered from 150 individuals. Sycamore trees were taken from thirteen sites (see Fig. 18). They were located in the orographic units of Malé Karpaty (sites 1 and 2), Strážovské Vrchy (3), Kysucké Beskydy (4), Javorie (5 and 6), Polana (7, 8, 9, 10), Nízke Tatry (11) and Slovenské rudohorie (12 and 13). The samples represented the individuals with $d_{0}$ diameters from 1.75 mm to 104.50 mm , and heights from 0.11 m to 9.87 m (Table 17, Fig. 19a). The dry mass of the whole trees ranged from 0.23 g do $16,932.93 \mathrm{~g}$, and the stem volume ranged from $1.13 \mathrm{~cm}^{3}$ to $23,114.93 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 18. The volume of stem over or under bark, its density, as well as the root-shoot ratio were derived in a similar way (Table 19). The regression models, scatter plots, and fitted regression curves were visualised in the graphs as in the case of the previous tree species. The text describing the biomass of individual components and their proportions of the total tree biomass is presented in Chapter 4.12. (Inter-species comparison of biomass characteristics).


Fig. 18. Map of sample sites of Sycamore and its distribution in the forests of Slovakia.

Table 17. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25. p), 75-percentile (75. p) and skewness of diameter ( $d_{0}$ ), tree height ( $h$ ), biomass of stem over bark (SOB), biomass of stem under bark (SUB), foliage biomass (foliage), branch biomass (branches), bark biomass (bark), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem outside bark ( $V_{\text {SoB }}$ ) of individual trees.

|  | Sycamore |  |  |  |  |  | Max | $25 . \mathrm{p}$ |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | 75.p | Skewness |  |  |
| $d_{0}(\mathrm{~mm})$ | 150 | 27.73 | 22.18 | 1.75 | 104.50 | 11.00 | 37.45 | 1.27 |
| $h(\mathrm{~m})$ | 150 | 3.00 | 2.50 | 0.11 | 9.87 | 1.01 | 4.79 | 0.87 |
| SOB $(\mathrm{g})$ | 150 | 711.72 | 1645.16 | 0.11 | 12190.73 | 19.40 | 690.49 | 3.91 |
| SUB $(\mathrm{g})$ | 150 | 666.92 | 1458.71 | 0.10 | 10948.73 | 14.40 | 562.59 | 4.01 |
| Foliage $(\mathrm{g})$ | 150 | 70.22 | 105.39 | 0.18 | 607.75 | 8.05 | 77.24 | 2.95 |
| Branches $(\mathrm{g})$ | 125 | 92.15 | 196.18 | 0.05 | 1351.80 | 6.40 | 72.65 | 4.03 |
| Bark $(\mathrm{g})$ | 150 | 104.80 | 188.64 | 0.01 | 1242.00 | 110.20 | 188.64 | 3.21 |
| Roots $(\mathrm{g})$ | 150 | 243.57 | 429.48 | 0.03 | 2834.80 | 16.90 | 228.00 | 3.13 |
| Aboveground $(\mathrm{g})$ | 150 | 918.73 | 1908.70 | 0.19 | 14098.13 | 32.25 | 812.84 | 3.88 |
| Whole tree $(\mathrm{g})$ | 150 | 1162.30 | 2325.49 | 0.23 | 16932.93 | 52.35 | 1043.11 | 3.73 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 150 | 1569.80 | 3317.40 | 0.13 | 23114.93 | 31.36 | 1516.02 | 3.71 |

Table 18. Sycamore, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination ( $R^{2}$ ), mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -3.891 | 0.090 | <0.001 | 2.842 | 0.029 | <0.001 |  |  |  | 0.985 | 0.110 | 1.047 | 0.365 |
|  | Stem under bark | -4.157 | 0.095 | <0.001 | 2.857 | 0.030 | <0.001 |  |  |  | 0.983 | 0.122 | 1.051 | 0.382 |
|  | Branches | -6.242 | 0.315 | <0.001 | 2.844 | 0.095 | <0.001 |  |  |  | 0.880 | 0.499 | 1.223 | 0.724 |
|  | Foliage | -2.468 | 0.135 | <0.001 | 1.894 | 0.043 | $<0.001$ |  |  |  | 0.928 | 0.247 | 1.118 | 0.509 |
|  | Bark | -6.119 | 0.176 | <0.001 | 2.996 | 0.057 | <0.001 |  |  |  | 0.950 | 0.420 | 1.201 | 0.731 |
|  | Roots | -3.998 | 0.132 | <0.001 | 2.645 | 0.042 | $<0.001$ |  |  |  | 0.963 | 0.236 | 1.124 | 0.590 |
|  | Aboveground part | -2.931 | 0.077 | <0.001 | 2.645 | 0.025 | <0.001 |  |  |  | 0.987 | 0.080 | 1.041 | 0.301 |
|  | Whole tree | -2.558 | 0.073 | $<0.001$ | 2.627 | 0.023 | $<0.001$ |  |  |  | 0.988 | 0.072 | 1.037 | 0.293 |
| [10] | Stem over bark | 3.032 | 0.043 | $<0.001$ | 2.377 | 0.034 | $<0.001$ |  |  |  | 0.971 | 0.213 | 1.107 | 0.553 |
|  | Stem under bark | 2.803 | 0.043 | <0.001 | 2.394 | 0.034 | $<0.001$ |  |  |  | 0.971 | 0.208 | 1.102 | 0.544 |
|  | Branches | 0.817 | 0.138 | <0.001 | 2.240 | 0.109 | <0.001 |  |  |  | 0.775 | 0.937 | 1.478 | 1.341 |
|  | Foliage | 2.183 | 0.068 | <0.001 | 1.524 | 0.053 | $<0.001$ |  |  |  | 0.847 | 0.526 | 1.254 | 0.818 |
|  | Bark | 1.187 | 0.073 | <0.001 | 2.496 | 0.057 | <0.001 |  |  |  | 0.928 | 0.602 | 1.331 | 1.263 |
|  | Roots | 2.481 | 0.074 | <0.001 | 2.159 | 0.058 | <0.001 |  |  |  | 0.904 | 0.619 | 1.348 | 1.144 |
|  | Aboveground part | 3.521 | 0.046 | <0.001 | 2.201 | 0.036 | $<0.001$ |  |  |  | 0.962 | 0.236 | 1.123 | 0.562 |
|  | Whole tree | 3.857 | 0.050 | <0.001 | 2.174 | 0.040 | <0.001 |  |  |  | 0.953 | 0.289 | 1.155 | 0.652 |
| [11] | Stem over bark | -1.223 | 0.143 | $<0.001$ | 1.736 | 0.058 | $<0.001$ | 0.966 | 0.049 | <0.001 | 0.996 | 0.030 | 1.006 | 0.186 |
|  | Stem under bark | -1.350 | 0.151 | <0.001 | 1.694 | 0.061 | $<0.001$ | 1.016 | 0.052 | $<0.001$ | 0.995 | 0.034 | 1.006 | 0.193 |
|  | Branches | -5.755 | 0.640 | <0.001 | 2.638 | 0.254 | <0.001 | 0.186 | 0.213 | 0.383 | 0.881 | 0.500 | 1.222 | 0.719 |
|  | Foliage | -3.107 | 0.406 | <0.001 | 2.158 | 0.165 | $<0.001$ | -0.231 | 0.139 | 0.098 | 0.929 | 0.244 | 1.118 | 0.516 |
|  | Bark | -3.738 | 0.493 | <0.001 | 2.009 | 0.200 | $<0.001$ | 0.861 | 0.168 | $<0.001$ | 0.957 | 0.359 | 1.176 | 0.768 |
|  | Roots | -3.680 | 0.400 | $<0.001$ | 2.514 | 0.162 | $<0.001$ | 0.115 | 0.136 | 0.401 | 0.964 | 0.236 | 1.123 | 0.589 |
|  | Aboveground part | -0.930 | 0.156 | <0.001 | 1.816 | 0.063 | $<0.001$ | 0.724 | 0.053 | $<0.001$ | 0.994 | 0.036 | 1.018 | 0.193 |
|  | Whole tree | -1.040 | 0.177 | <0.001 | 1.998 | 0.072 | $<0.001$ | 0.549 | 0.060 | <0.001 | 0.993 | 0.047 | 1.024 | 0.232 |

Table 19. Sycamore, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE) for equations [13] - [15].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | SOB volume | 0.028 | 0.011 | 0.010 | 2.917 | 0.087 | $<0.001$ |  |  | $\mathrm{R}^{2}$ |
| SUB volume | 0.016 | 0.007 | 0.018 | 3.016 | 0.095 | $<0.001$ |  |  | 0.945 | 604772 |
|  | Bark volume | 0.047 | 0.015 | 0.002 | 2.294 | 0.073 | $<0.001$ |  |  | 0.943 |
| SOB density | 790.296 | 22.815 | $<0.001$ | -0.128 | 0.010 | $<0.001$ |  | 0.935 | 8287 |  |
| [13] | SUB density | 615.677 | 24.485 | $<0.001$ | -0.064 | 0.013 | $<0.001$ |  | 0.502 | 4457 |
| Bark density | 1485.74 | 96.33 | $<0.001$ | -0.262 | 0.022 | $<0.001$ |  | 0.154 | 3007 |  |
| Bark mass proportion | 58.843 | 1.902 | $<0.001$ | -0.347 | 0.011 | $<0.001$ |  | 0.510 | 14410 |  |
| Bark volume proportion | 34.096 | 2.118 | $<0.001$ | -0.233 | 0.021 | $<0.001$ |  | 0.883 | 3.51 |  |
| R/S ratio | 1.051 | 0.143 | $<0.001$ | -0.310 | 0.047 | $<0.001$ |  | 0.480 | 9.37 |  |


| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [14] | SOB volume | 4.270 | 1.755 | 0.016 | 3.645 | 0.191 | $<0.001$ |  |  |  | 0.867 | 1468819 |
|  | SUB volume | 2.961 | 1.339 | 0.029 | 3.763 | 0.210 | $<0.001$ |  |  |  | 0.861 | 1312206 |
|  | Bark volume | 2.295 | 0.685 | 0.001 | 2.894 | 0.142 | $<0.001$ |  |  |  | 0.877 | 15784 |
|  | SOB density | 580.761 | 5.685 | <0.001 | -0.112 | 0.008 | $<0.001$ |  |  |  | 0.556 | 3974 |
|  | SUB density | 533.178 | 5.731 | <0.001 | -0.069 | 0.009 | <0.001 |  |  |  | 0.281 | 2556 |
|  | Bark density | 774.138 | 13.626 | $<0.001$ | -0.204 | 0.017 | $<0.001$ |  |  |  | 0.506 | 14552 |
|  | Bark mass proportion | 24.707 | 0.254 | <0.001 | -0.262 | 0.010 | <0.001 |  |  |  | 0.826 | 5.22 |
|  | Bark volume proportion | 19.182 | 0.321 | $<0.001$ | -0.189 | 0.016 | $<0.001$ |  |  |  | 0.510 | 8.84 |
|  | R/S ratio | 0.497 | 0.017 | $<0.001$ | -0.277 | 0.034 | $<0.001$ |  |  |  | 0.343 | 0.025 |
| [15] | SOB volume | 0.172 | 0.030 | $<0.001$ | 1.815 | 0.058 | $<0.001$ | 1.452 | 0.072 | $<0.001$ | 0.987 | 143855 |
|  | SUB volume | 0.107 | 0.021 | $<0.001$ | 1.882 | 0.065 | $<0.001$ | 1.487 | 0.080 | $<0.001$ | 0.986 | 137455 |
|  | Bark volume | 0.188 | 0.038 | $<0.001$ | 1.404 | 0.070 | $<0.001$ | 1.214 | 0.087 | $<0.001$ | 0.974 | 3384 |
|  | SOB density | 532.286 | 51.026 | <0.001 | 0.036 | 0.039 | 0.362 | -0.141 | 0.033 | <0.001 | 0.558 | 3979 |
|  | SUB density | 358.061 | 30.039 | $<0.001$ | 0.158 | 0.033 | $<0.001$ | -0.189 | 0.027 | $<0.001$ | 0.386 | 2198 |
|  | Bark density | 1116.778 | 183.790 | $<0.001$ | -0.145 | 0.065 | 0.027 | -0.096 | 0.051 | 0.062 | 0.523 | 14139 |
|  | Bark mass proportion | 49.967 | 4.122 | $<0.001$ | -0.280 | 0.033 | $<0.001$ | -0.055 | 0.025 | 0.033 | 0.887 | 3.42 |
|  | Bark volume proportion | 21.791 | 3.574 | <0.001 | -0.051 | 0.065 | 0.436 | -0.151 | 0.051 | 0.004 | 0.512 | 8.86 |
|  | R/S ratio | 0.184 | 0.065 | 0.006 | 0.394 | 0.139 | 0.005 | -0.575 | 0.111 | $<0.001$ | 0.378 | 0.024 |



Fig. 19. Relationship of height a), $d_{1.3}$ diameter $b$ ), dry mass of stem over bark $c$ ), dry mass of branches $d$ ) to stem base diameter $d_{0}$ of Sycamore.


Fig. 20. Relationship offoliage dry mass a), root dry mass b), dry mass of the whole tree c), proportion of individual tree components d), basic density of stem over bark e), and basic bark density f) to stem base diameter $d_{0}$ of Sycamore.


Fig. 21. Relationship of bark mass proportion of SOB mass a) bark proportion of SOB volume b), volume of stem over bark c) and $R / S$ ratio d) to stem base diameter $d_{0}$ of Sycamore.

### 4.7. European ash

European ash (Fraxinus excelsior L.) belongs together with maple species to the so-called valuable broadleaved species. They represent specific components of stands, which are important for biodiversity, and partially also from the economic point of view. They usually form admixtures to main commercial tree species, and only rarely form small-scale homogeneous stands. Ash is an important admixture practically in all forest vegetation zones from $1^{\text {st }}$ oak up to $7^{\text {th }}$ spruce zones, i.e. it has a wide ecological valency. At specific sites, e.g. at rocky, scree or waterlogged sites, its proportion is greater. In scree and riparian forests it is the main stand-forming tree species. In these forests it grows in the biotopes protected at national and European levels: Ls1.4 Ash-alder sub-montane alluvial forests (NATURA 2000 91E0 Alluvial forests with Alnus glutinosa and Fraxinus excelsior), Ls1.2 Oak-elm-ash plain-land alluvial forests (91F0 Riparian mixed forests of Quercus robur, Ulmus laevis and Ulmus minor, Fraxinus excelsior or Fraxinus angustifolia, along the great rivers).

The lowest and the highest occurrences of ash (NFIM2 SR) were recorded at elevations of 112 m a.s.l. and $1,151 \mathrm{~m}$ a.s.l., respectively, and the most frequent occurrence was recorded between $500-600 \mathrm{~m}$ a.s.l. From the point of its proportion of the area, stock, and occurrence, it is
the $11^{\text {th }}$ most frequent tree species in the forests of Slovakia. It grew at a reduced area of $44 \pm 11$ thousand ha, and occurred at $18 \%$ of the forested inventory plots.

The biomass regression models of the European ash were derived using 81 trees. They were taken from six sites (see Fig. 22). The sites were located in two orographic units, namely Javorie $(1,2,3)$ and Polana $(4,5,6)$. The diameters $d_{0}$ of the trees ranged from 5.35 mm to 51.25 mm , and heights from 0.19 m to 4.24 m (Table 20, Fig. 23a). The dry mass of the whole trees ranged from 5.25 g do $3,064.50 \mathrm{~g}$, and the stem volume ranged from $2.10 \mathrm{~cm}^{3}$ to $2,239.89 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 21. The regression models for the calculation of the volume of stem over or under bark, its density, as well as the root-shoot ratio are presented in Table 22.

The regression models, scatter plots, and fitted regression curves were visualised in the graphs in a similar way as for the previous tree species. Further comments on the biomass of individual components and their proportions in the total tree biomass are presented in Chapter 4.12. (Inter-species comparison of biomass characteristics).


Fig. 22. Map of sample sites of European ash and its distribution in the forests of Slovakia.

Table 20. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25. p), 75-percentile (75. p) and skewness of diameter $\left(d_{0}\right)$, tree height (h), biomass of stem over bark (SOB), biomass of stem under bark (SUB), foliage biomass (foliage), branch biomass (branches), bark biomass (bark), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {SOB }}$ ) of individual trees.

|  |  | European ash |  |  |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | $25 . \mathrm{p}$ | $75 . \mathrm{p}$ | Skewness |
| $d_{0}(\mathrm{~mm})$ | 81 | 23.44 | 11.61 | 5.35 | 51.25 | 13.35 | 31.75 | 0.42 |
| $h(\mathrm{~m})$ | 81 | 2.09 | 1.18 | 0.19 | 4.24 | 1.02 | 2.96 | 0.12 |
| SOB $(\mathrm{g})$ | 80 | 275.90 | 311.12 | 1.70 | 1235.65 | 27.70 | 415.08 | 1.30 |
| SUB $(\mathrm{g})$ | 80 | 218.27 | 253.08 | 1.00 | 1009.20 | 19.65 | 327.48 | 1.36 |
| Foliage $(\mathrm{g})$ | 81 | 52.17 | 57.07 | 0.75 | 307.20 | 8.55 | 74.65 | 1.90 |
| Branches $(\mathrm{g})$ | 81 | 40.56 | 66.07 | 0.00 | 322.55 | 1.95 | 44.80 | 2.49 |
| Bark $(\mathrm{g})$ | 81 | 57.40 | 58.64 | 0.70 | 226.45 | 7.75 | 87.00 | 1.10 |
| Roots $(\mathrm{g})$ | 80 | 133.69 | 179.74 | 1.60 | 1258.00 | 23.08 | 169.90 | 3.63 |
| Aboveground $(\mathrm{g})$ | 80 | 369.10 | 421.27 | 2.45 | 1806.50 | 41.43 | 540.15 | 1.42 |
| Whole tree $(\mathrm{g})$ | 79 | 508.10 | 586.19 | 5.25 | 3064.50 | 64.95 | 700.00 | 1.81 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 81 | 502.06 | 571.05 | 2.10 | 2239.89 | 52.31 | 771.22 | 1.33 |

Table 21. European ash, $b_{o}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination ( $R^{2}$ ), mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -4.374 | 0.229 | $<0.001$ | 2.997 | 0.075 | $<0.001$ |  |  |  | 0.954 | 0.140 | 1.072 | 0.430 |
|  | Stem under bark | -5.129 | 0.247 | <0.001 | 3.147 | 0.081 | $<0.001$ |  |  |  | 0.951 | 0.163 | 1.088 | 0.516 |
|  | Branches | -9.108 | 0.624 | $<0.001$ | 3.738 | 0.197 | $<0.001$ |  |  |  | 0.835 | 0.562 | 1.256 | 0.792 |
|  | Foliage | -3.969 | 0.255 | $<0.001$ | 2.388 | 0.083 | $<0.001$ |  |  |  | 0.912 | 0.174 | 1.085 | 0.435 |
|  | Bark | -4.658 | 0.204 | $<0.001$ | 2.630 | 0.066 | $<0.001$ |  |  |  | 0.952 | 0.111 | 1.053 | 0.331 |
|  | Roots | -3.301 | 0.243 | <0.001 | 2.454 | 0.079 | $<0.001$ |  |  |  | 0.925 | 0.146 | 1.077 | 0.452 |
|  | Aboveground part | -3.839 | 0.209 | $<0.001$ | 2.925 | 0.068 | $<0.001$ |  |  |  | 0.959 | 0.117 | 1.057 | 0.360 |
|  | Whole tree | -2.999 | 0.200 | $<0.001$ | 2.769 | 0.065 | $<0.001$ |  |  |  | 0.959 | 0.099 | 1.049 | 0.332 |
| [10] | Stem over bark | 3.523 | 0.056 | <0.001 | 2.206 | 0.061 | $<0.001$ |  |  |  | 0.944 | 0.171 | 1.096 | 0.601 |
|  | Stem under bark | 3.160 | 0.059 | $<0.001$ | 2.317 | 0.065 | $<0.001$ |  |  |  | 0.942 | 0.193 | 1.119 | 0.773 |
|  | Branches | 0.698 | 0.168 | $<0.001$ | 2.785 | 0.189 | $<0.001$ |  |  |  | 0.753 | 0.841 | 1.433 | 1.174 |
|  | Foliage | 2.345 | 0.070 | $<0.001$ | 1.721 | 0.077 | $<0.001$ |  |  |  | 0.864 | 0.270 | 1.129 | 0.554 |
|  | Bark | 2.271 | 0.047 | $<0.001$ | 1.942 | 0.052 | $<0.001$ |  |  |  | 0.946 | 0.124 | 1.060 | 0.358 |
|  | Roots | 3.171 | 0.075 | $<0.001$ | 1.780 | 0.084 | $<0.001$ |  |  |  | 0.853 | 0.288 | 1.156 | 0.719 |
|  | Aboveground part | 3.874 | 0.057 | $<0.001$ | 2.140 | 0.062 | $<0.001$ |  |  |  | 0.938 | 0.178 | 1.093 | 0.522 |
|  | Whole tree | 4.279 | 0.062 | $<0.001$ | 2.048 | 0.069 | $<0.001$ |  |  |  | 0.920 | 0.193 | 1.100 | 0.525 |
| [11] | Stem over bark | -0.906 | 0.372 | 0.017 | 1.667 | 0.139 | $<0.001$ | 1.052 | 0.103 | $<0.001$ | 0.980 | 0.060 | 1.038 | 0.399 |
|  | Stem under bark | -1.460 | 0.409 | $<0.001$ | 1.739 | 0.153 | $<0.001$ | 1.113 | 0.114 | $<0.001$ | 0.978 | 0.073 | 1.052 | 0.535 |
|  | Branches | -6.807 | 1.152 | $<0.001$ | 2.830 | 0.431 | $<0.001$ | 0.795 | 0.339 | 0.022 | 0.847 | 0.529 | 1.244 | 0.782 |
|  | Foliage | -2.219 | 0.596 | <0.001 | 1.717 | 0.223 | $<0.001$ | 0.531 | 0.165 | 0.002 | 0.923 | 0.156 | 1.075 | 0.411 |
|  | Bark | -1.502 | 0.321 | $<0.001$ | 1.419 | 0.120 | $<0.001$ | 0.958 | 0.089 | $<0.001$ | 0.981 | 0.045 | 1.022 | 0.222 |
|  | Roots | -2.043 | 0.562 | $<0.001$ | 1.968 | 0.211 | $<0.001$ | 0.394 | 0.160 | 0.016 | 0.931 | 0.137 | 1.073 | 0.450 |
|  | Aboveground part | -0.795 | 0.355 | 0.028 | 1.757 | 0.133 | $<0.001$ | 0.923 | 0.099 | $<0.001$ | 0.981 | 0.055 | 1.031 | 0.315 |
|  | Whole tree | -0.589 | 0.372 | 0.117 | 1.838 | 0.140 | $<0.001$ | 0.754 | 0.106 | $<0.001$ | 0.976 | 0.060 | 1.032 | 0.301 |

Table 22. European ash, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE) for equations [13] - [15].

| Eq. | Tree component | $\mathrm{b}_{0}$ | S.E. | P | $\mathrm{b}_{1}$ | S.E. | P | $\mathrm{b}_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [13] | SOB volume | 0.257 | 0.119 | 0.034 | 2.311 | 0.126 | $<0.001$ |  |  |  | 0.894 | 34847 |
|  | SUB volume | 0.187 | 0.091 | 0.042 | 2.330 | 0.131 | $<0.001$ |  |  |  | 0.889 | 22687 |
|  | Barkvolume | 0.072 | 0.033 | 0.032 | 2.245 | 0.125 | $<0.001$ |  |  |  | 0.890 | 1728 |
|  | SOB density | 731.954 | 47.687 | $<0.001$ | -0.090 | 0.022 | $<0.001$ |  |  |  | 0.173 | 3830 |
|  | SUB density | 563.952 | 36.847 | $<0.001$ | -0.010 | 0.021 | 0.659 |  |  |  | 0.002 | 3432 |
|  | Bark density | 1316.481 | 127.782 | $<0.001$ | -0.267 | 0.034 | $<0.001$ |  |  |  | 0.432 | 11372 |
|  | Bark mass proportion | 80.316 | 4.586 | $<0.001$ | -0.382 | 0.020 | <0.001 |  |  |  | 0.808 | 8.30 |
|  | Bark volume proportion | 46.271 | 4.048 | $<0.001$ | -0.210 | 0.030 | $<0.001$ |  |  |  | 0.365 | 15.19 |
|  | R/S ratio | 1.647 | 0.294 | $<0.001$ | -0.441 | 0.064 | $<0.001$ |  |  |  | 0.366 | 0.023 |
| [14] | SOB volume | 51.973 | 12.498 | $<0.001$ | 2.469 | 0.187 | $<0.001$ |  |  |  | 0.854 | 48182 |
|  | SUB volume | 37.127 | 8.820 | $<0.001$ | 2.542 | 0.185 | $<0.001$ |  |  |  | 0.865 | 27397 |
|  | Bark volume | 15.394 | 4.041 | $<0.001$ | 2.221 | 0.207 | $<0.001$ |  |  |  | 0.792 | 3258 |
|  | SOB density | 581.051 | 7.840 | $<0.001$ | -0.077 | 0.015 | $<0.001$ |  |  |  | 0.243 | 3508 |
|  | SUB density | 554.752 | 7.817 | $<0.001$ | -0.024 | 0.015 | 0.132 |  |  |  | 0.028 | 3345 |
|  | Bark density | 649.378 | 13.624 | <0.001 | -0.190 | 0.024 | <0.001 |  |  |  | 0.426 | 11479 |
|  | Bark mass proportion | 29.185 | 0.385 | $<0.001$ | -0.263 | 0.015 | $<0.001$ |  |  |  | 0.782 | 9.41 |
|  | Bark volume proportion | 26.681 | 0.468 | $<0.001$ | -0.162 | 0.020 | $<0.001$ |  |  |  | 0.433 | 13.56 |
|  | R/S ratio | 0.525 | 0.017 | $<0.001$ | -0.368 | 0.039 | $<0.001$ |  |  |  | 0.500 | 0.018 |
| [15] | SOB volume | 1.501 | 0.554 | 0.008 | 1.397 | 0.129 | $<0.001$ | 1.286 | 0.151 | $<0.001$ | 0.948 | 17360 |
|  | SUB volume | 1.250 | 0.451 | 0.007 | 1.338 | 0.125 | $<0.001$ | 1.407 | 0.150 | $<0.001$ | 0.951 | 10165 |
|  | Bark volume | 0.261 | 0.122 | 0.035 | 1.600 | 0.164 | $<0.001$ | 0.880 | 0.183 | $<0.001$ | 0.917 | 1311.8 |
|  | SOB density | 476.008 | 76.962 | $<0.001$ | 0.075 | 0.060 | 0.219 | -0.129 | 0.044 | 0.005 | 0.258 | 3482 |
|  | SUB density | 366.551 | 57.336 | $<0.001$ | 0.156 | 0.058 | 0.009 | -0.131 | 0.043 | 0.003 | 0.114 | 3088 |
|  | Bark density | 972.677 | 259.621 | $<0.001$ | -0.152 | 0.100 | 0.134 | -0.088 | 0.072 | 0.225 | 0.442 | 11305 |
|  | Bark mass proportion | 56.377 | 8.954 | $<0.001$ | -0.247 | 0.060 | $<0.001$ | -0.099 | 0.042 | 0.021 | 0.822 | 7.82 |
|  | Bark volume proportion | 24.708 | 5.621 | $<0.001$ | 0.029 | 0.085 | 0.736 | -0.181 | 0.061 | 0.004 | 0.434 | 13.71 |
|  | $\mathrm{R} /$ S ratio | 0.205 | 0.095 | 0.033 | 0.352 | 0.171 | 0.043 | -0.610 | 0.123 | $<0.001$ | 0.527 | 0.017 |



Fig. 23. Relationship of tree height a), $d_{1.3}$ diameter b), dry mass of stem over bark c), dry mass of branches d) to stem base diameter $d_{0}$ of European ash.


Fig. 24. Relationship of foliage dry mass a), root dry mass b), dry mass of the whole tree c), proportion of individual tree components d), basic density of stem over bark e), and basic bark density f) to stem base diameter $d_{0}$ of European ash.


Fig. 25. Relationship of bark mass proportion of SOB mass a), bark volume proportion of SOB volume b), volume of stem outside bark c), and $R / S$ ratio d) to stem base diameter $d_{0}$ of European ash.

### 4.8. Goat willow

Goat willow (Salix caprea L.) is a pioneer tree species, which occurs mainly after disturbances or as an admixture in all forest vegetation zones from $1^{\text {st }}$ oak zone up to $7^{\text {th }}$ spruce zone. Although it is an abundant tree species, it has a negligible economic importance. Therefore, currently it is not assessed as a stand-forming tree species within the forest survey in Slovakia during the renewal of forest management plans except for noting it down in the site description. Goat willow, similarly to rowan or aspen, provides important nutrition to wildlife, and can lure it away from the commercially important tree species and thus reduce economic losses.

According to the NFIM2 SR data, the lowest and the highest occurrences of this tree species were recorded at elevations of 118 m and $1,485 \mathrm{~m}$ a.s.l., respectively, with most frequent occurrence between $600-700 \mathrm{~m}$ a.s.l. Although it is an abundant tree species ranked $12^{\text {th }}$ from all tree species occurring in the forests of Slovakia on the base of its occurrence and its proportion of the
 $\pm 10$ thousand ha, and occurred at $15 \%$ of the forested inventory plots.

The biomass regression models of the Goat willow were derived using 100 individuals. They were taken from two sites (see Fig. 26) located in the orographic units of Kysucké Beskydy (site 1) and Vysoké Tatry (2). The trees covered the interval of $d_{0}$ stem diameters from 3.65 mm to 68.70 mm , and heights from 0.49 m to 4.50 m (Table 23, Fig. 27a). The dry mass of the whole trees ranged from 3.19 g to $3,799.37 \mathrm{~g}$, and the stem volume ranged from $1.83 \mathrm{~cm}^{3}$ to $3,684.05 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 24. The volume of stem outside or inside bark, its density, as well as the root-shoot ratio were derived in a similar way (Table 25).

The regression models, scatter plots, and fitted regression curves were visualised in a similar way as in the case of the previous tree species. Further text description of the biomass of individual components and their proportions in the total tree biomass is presented in Chapter 4.12. (Inter-species comparison of biomass characteristics)


Fig. 26. Map of sample sites of Goat willow.

Table 23. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25. p), 75-percentile (75. $p$ ) and skewness of diameter $\left(d_{0}\right)$, tree height ( $h$ ), biomass of stem over bark (SOB), biomass of stem under bark (SUB), foliage biomass (foliage), branch biomass (branches), bark biomass (bark), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {SOB }}$ ) of individual trees.

| Goat willow |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | 25.p | 75.p | Skewness |
| $d_{0}(\mathrm{~mm})$ | 100 | 25.04 | 13.22 | 3.65 | 68.70 | 14.60 | 33.25 | 1.02 |
| $h(\mathrm{~m})$ | 100 | 2.04 | 0.85 | 0.49 | 4.50 | 1.38 | 2.37 | 0.81 |
| SOB (g) | 100 | 197.64 | 293.03 | 1.15 | 1523.23 | 29.33 | 218.93 | 2.97 |
| SUB (g) | 100 | 152.89 | 239.64 | 0.70 | 1257.93 | 20.10 | 160.30 | 3.10 |
| Foliage (g) | 99 | 78.41 | 87.02 | 0.80 | 467.35 | 20.30 | 110.05 | 2.17 |
| Branches (g) | 99 | 106.06 | 173.06 | 0.44 | 1185.75 | 13.00 | 120.30 | 3.53 |
| Bark (g) | 100 | 44.75 | 54.29 | 0.45 | 265.30 | 9.28 | 54.30 | 2.37 |
| Roots (g) | 100 | 89.08 | 136.30 | 0.80 | 796.50 | 15.33 | 100.43 | 3.19 |
| Aboveground (g) | 100 | 380.26 | 539.34 | 2.39 | 3149.17 | 61.78 | 411.53 | 2.94 |
| Whole tree (g) | 98 | 475.06 | 678.75 | 3.19 | 3799.37 | 79.55 | 513.90 | 2.93 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 100 | 471.69 | 686.23 | 1.83 | 3684.05 | 65.81 | 539.11 | 2.99 |

Table 24. Goat willow, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values ( $P$ ), coefficients of determination ( $R^{2}$ ), mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -3.178 | 0.189 | <0.001 | 2.479 | 0.061 | $<0.001$ |  |  |  | 0.945 | 0.107 | 1.048 | 0.306 |
|  | Stem under bark | -3.886 | 0.202 | <0.001 | 2.601 | 0.065 | <0.001 |  |  |  | 0.943 | 0.122 | 1.055 | 0.331 |
|  | Branches | -5.018 | 0.236 | <0.001 | 2.807 | 0.075 | <0.001 |  |  |  | 0.935 | 0.166 | 1.078 | 0.403 |
|  | Foliage | -2.409 | 0.206 | <0.001 | 2.015 | 0.066 | <0.001 |  |  |  | 0.906 | 0.127 | 1.061 | 0.360 |
|  | Bark | -3.510 | 0.177 | <0.001 | 2.166 | 0.057 | $<0.001$ |  |  |  | 0.937 | 0.094 | 1.043 | 0.290 |
|  | Roots | -3.537 | 0.193 | <0.001 | 2.346 | 0.062 | $<0.001$ |  |  |  | 0.937 | 0.111 | 1.052 | 0.330 |
|  | Aboveground part | -2.305 | 0.149 | $<0.001$ | 2.425 | 0.047 | $<0.001$ |  |  |  | 0.965 | 0.066 | 1.029 | 0.232 |
|  | Whole tree | -2.035 | 0.146 | $<0.001$ | 2.406 | 0.047 | <0.001 |  |  |  | 0.965 | 0.064 | 1.029 | 0.231 |
| [10] | Stem over bark | 2.542 | 0.066 | <0.001 | 3.063 | 0.086 | $<0.001$ |  |  |  | 0.928 | 0.140 | 1.069 | 0.390 |
|  | Stem under bark | 2.106 | 0.065 | <0.001 | 3.230 | 0.086 | <0.001 |  |  |  | 0.935 | 0.138 | 1.068 | 0.386 |
|  | Branches | 1.613 | 0.125 | <0.001 | 3.237 | 0.164 | <0.001 |  |  |  | 0.801 | 0.505 | 1.269 | 0.944 |
|  | Foliage | 2.335 | 0.094 | <0.001 | 2.344 | 0.124 | <0.001 |  |  |  | 0.786 | 0.289 | 1.153 | 0.650 |
|  | Bark | 1.515 | 0.070 | <0.001 | 2.636 | 0.092 | <0.001 |  |  |  | 0.893 | 0.160 | 1.080 | 0.430 |
|  | Roots | 1.953 | 0.091 | <0.001 | 2.780 | 0.120 | <0.001 |  |  |  | 0.846 | 0.271 | 1.135 | 0.573 |
|  | Aboveground part | 3.360 | 0.080 | <0.001 | 2.907 | 0.105 | <0.001 |  |  |  | 0.889 | 0.206 | 1.105 | 0.508 |
|  | Whole tree | 3.589 | 0.080 | <0.001 | 2.879 | 0.105 | $<0.001$ |  |  |  | 0.886 | 0.208 | 1.104 | 0.501 |
| [11] | Stem over bark | -0.809 | 0.212 | $<0.001$ | 1.414 | 0.088 | $<0.001$ | 1.457 | 0.110 | $<0.001$ | 0.980 | 0.039 | 1.019 | 0.195 |
|  | Stem under bark | -1.254 | 0.207 | <0.001 | 1.417 | 0.086 | <0.001 | 1.619 | 0.107 | <0.001 | 0.983 | 0.037 | 1.018 | 0.192 |
|  | Branches | -4.569 | 0.442 | $<0.001$ | 2.605 | 0.184 | $<0.001$ | 0.275 | 0.229 | 0.233 | 0.936 | 0.165 | 1.079 | 0.412 |
|  | Foliage | -1.939 | 0.382 | <0.001 | 1.803 | 0.159 | $<0.001$ | 0.290 | 0.199 | 0.148 | 0.908 | 0.125 | 1.061 | 0.363 |
|  | Bark | -1.834 | 0.263 | <0.001 | 1.413 | 0.109 | $<0.001$ | 1.031 | 0.136 | $<0.001$ | 0.961 | 0.060 | 1.029 | 0.246 |
|  | Roots | -2.441 | 0.336 | $<0.001$ | 1.854 | 0.140 | $<0.001$ | 0.674 | 0.175 | $<0.001$ | 0.945 | 0.097 | 1.048 | 0.322 |
|  | Aboveground part | -0.912 | 0.025 | <0.001 | 1.801 | 0.094 | <0.001 | 0.854 | 0.117 | <0.001 | 0.977 | 0.043 | 1.021 | 0.206 |
|  | Whole tree | -0.705 | 0.225 | 0.002 | 1.810 | 0.094 | <0.001 | 0.815 | 0.117 | $<0.001$ | 0.977 | 0.043 | 1.021 | 0.207 |

Table 25. Goat willow, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE) for equations [13] - [15].

| Eq. | Dependent variable | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [13] | SOB volume | 0.066 | 0.018 | $<0.001$ | 2.598 | 0.068 | $<0.001$ |  |  |  | 0.952 | 22611 |
|  | SUB volume | 0.038 | 0.011 | 0.001 | 2.685 | 0.075 | $<0.001$ |  |  |  | 0.948 | 16280 |
|  | Bark volume | 0.047 | 0.013 | 0.001 | 2.285 | 0.072 | <0.001 |  |  |  | 0.931 | 1273 |
|  | SOB density | 512.437 | 31.057 | $<0.001$ | -0.063 | 0.020 | 0.002 |  |  |  | 0.090 | 2046 |
|  | SUB density | 417.407 | 27.283 | $<0.001$ | -0.009 | 0.021 | 0.666 |  |  |  | 0.002 | 2120 |
|  | Bark density | 783.500 | 64.267 | <0.001 | -0.168 | 0.027 | <0.001 |  |  |  | 0.272 | 5023 |
|  | Bark mass proportion | 68.039 | 4.579 | <0.001 | -0.292 | 0.023 | <0.001 |  |  |  | 0.620 | 13.45 |
|  | Bark volume proportion | 46.300 | 3.622 | $<0.001$ | -0.198 | 0.026 | $<0.001$ |  |  |  | 0.360 | 13.61 |
|  | R/S ratio | 0.293 | 0.038 | <0.001 | -0.074 | 0.042 | 0.079 |  |  |  | 0.029 | 0.003 |
| [14] | SOB volume | 30.423 | 4.321 | <0.001 | 3.128 | 0.105 | $<0.001$ |  |  |  | 0.926 | 34976 |
|  | SUB volume | 20.487 | 2.930 | <0.001 | 3.257 | 0.105 | <0.001 |  |  |  | 0.934 | 20493 |
|  | Bark volume | 10.933 | 1.763 | <0.001 | 2.681 | 0.124 | <0.001 |  |  |  | 0.859 | 2592 |
|  | SOB density | 444.766 | 7.911 | <0.001 | -0.085 | 0.024 | 0.001 |  |  |  | 0.107 | 2008 |
|  | SUB density | 411.380 | 8.079 | <0.001 | -0.021 | 0.026 | 0.413 |  |  |  | 0.006 | 2110 |
|  | Bark density | 529.619 | 12.696 | <0.001 | -0.199 | 0.034 | <0.001 |  |  |  | 0.254 | 5148 |
|  | Bark mass proportion | 35.160 | 0.548 | <0.001 | -0.383 | 0.023 | <0.001 |  |  |  | 0.725 | 9.75 |
|  | Bark volume proportion | 30.006 | 0.573 | $<0.001$ | -0.285 | 0.028 | $<0.001$ |  |  |  | 0.505 | 10.53 |
|  | R/S ratio | 0.250 | 0.010 | $<0.001$ | -0.112 | 0.052 | 0.033 |  |  |  | 0.042 | 0.003 |
| [15] | SOB volume | 1.234 | 0.224 | $<0.001$ | 1.364 | 0.069 | $<0.001$ | 1.513 | 0.085 | $<0.001$ | 0.989 | 5518 |
|  | SUB volume | 0.962 | 0.170 | <0.001 | 1.309 | 0.067 | <0.001 | 1.694 | 0.084 | <0.001 | 0.989 | 3309 |
|  | Bark volume | 0.276 | 0.102 | 0.008 | 1.540 | 0.140 | <0.001 | 0.903 | 0.165 | <0.001 | 0.947 | 975 |
|  | SOB density | 449.175 | 51.316 | $<0.001$ | -0.004 | 0.048 | 0.931 | -0.081 | 0.059 | 0.176 | 0.107 | 2028 |
|  | SUB density | 377.944 | 46.202 | <0.001 | 0.036 | 0.051 | 0.485 | -0.061 | 0.063 | 0.336 | 0.011 | 2121 |
|  | Bark density | 700.079 | 111.695 | <0.001 | -0.117 | 0.067 | 0.082 | -0.068 | 0.083 | 0.413 | 0.277 | 5039 |
|  | Bark mass proportion | 36.932 | 4.273 | <0.001 | -0.021 | 0.048 | 0.669 | -0.360 | 0.059 | $<0.001$ | 0.726 | 9.83 |
|  | Bark volume proportion | 24.309 | 3.255 | <0.001 | 0.088 | 0.056 | 0.115 | -0.383 | 0.068 | <0.001 | 0.518 | 10.37 |
|  | $\mathrm{R} /$ S ratio | 0.233 | 0.058 | <0.001 | 0.028 | 0.103 | 0.783 | -0.145 | 0.128 | 0.261 | 0.042 | 0.003 |



Fig. 27. Relationship of tree height a), $d_{1.3}$ diameter b), dry mass of stem over bark c), dry mass of branches d) to stem base diameter $d_{0}$ of Goat willow.


Fig. 28. Relationship of foliage dry mass a), root dry mass b), dry mass of the whole tree c), proportion of individual tree components d), basic density of stem outside bark e), and basic bark density f) to stem base diameter $d_{0}$ of Goat willow.


Fig. 29. Relationship of bark mass proportion of SOB mass a) bark volume proportion of SOB volume b), volume of stem over bark c) and $R / S$ ratio d) to stem base diameter $d_{0}$ of Goat willow.

### 4.9. European larch

European larch (Larix decidua Mill.) is a light demanding tree species, which is with its ecological demands adjusted to rough continental climate. It is an economically important and in industry well utilisable tree species. It is usually planted, and individually occurs in all forest vegetation zones from $1^{\text {st }}$ oak zone up to $8^{\text {th }}$ dwarf pine zone. Naturally, it dominates at rocky mountainous sites. Larch is typical for the most northern parts of Slovakia, usually occurring at elevations from 400 to $1,600 \mathrm{~m}$ a.s.l. It is normally an admixture in the stands of other main tree species, and outside Lariceto-Piceetum group of forest types (specific for the Vysoké Tatry) larch is not considered to be a native species if its proportion exceeds $5-15 \%$. At lower elevations it is regarded as a completely non-native species, but due to its economical importance it is popular to plant it at lower sites, too. Within the scheme of protected biotopes, larch is more abundant only in one biotope, namely Ls9.4 Larch-Stone pine forests (NATURA 20009420 "Alpine Larix decidua and/ or Pinus cembra forests").

The lowest and the highest occurrences of larch (according to NFIM2 SR) were recorded at elevations of 138 m and $1,465 \mathrm{~m}$ a.s.l., respectively, with most frequent occurrence between 700 and 800 m a.s.l. Its greater economic importance is confirmed by its $9^{\text {th }}$ place in the ranking of tree species according to the stock in the forests of Slovakia, while based on its proportion of the area it was ranked $15^{\text {th }}$, and from the point of the occurrence it was $16^{\text {th }}$ most frequent tree species. It grew at a reduced area of $32 \pm 9$ thousand ha, and occurred at $9 \%$ of the forested inventory plots.

The biomass regression models of European larch were derived from the set of 125 whole tree individuals. The larch trees were taken from six sites alllocated in the post-disturbance area of Vysoké Tatry (see Fig. 30). The trees covered the interval of $d_{0}$ diameters from 3.35 mm to 95.15 mm , and of heights from 0.21 m to 4.89 m (Table 26, Fig. 31a). The dry mass of the whole trees ranged from 3.80 g do $10,133.37 \mathrm{~g}$, and the stem volume ranged from $0.73 \mathrm{~cm}^{3}$ to $8,203.87 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 27. Similarly, the volume of stem over bark, its density, as well as the root-shoot ratio were derived (Table 28).

The regression models, scatter plots, and fitted regression curves were visualised in a similar way as in the case of the previous tree species. The comment on the biomass of individual components and their proportions in the total tree biomass is presented in Chapter 4.12. (Interspecies comparison of biomass characteristics).


Fig. 30. Map of sample sites of European larch and its distribution in the forests of Slovakia.

Table 26. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25. p), 75-percentile (75. p) and skewness of diameter ( $d_{\ell}$ ), tree height ( $h$ ), biomass of stem over bark (SOB), foliage biomass (foliage), branch biomass (branches), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {sob }}$ ) of individual trees.

|  | European larch |  |  |  |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | $25 . \mathrm{p}$ | $75 . \mathrm{p}$ | Skewness |
| $d_{0}(\mathrm{~mm})$ | 125 | 34.93 | 25.74 | 3.35 | 95.15 | 11.95 | 52.55 | 0.63 |
| $h(\mathrm{~m})$ | 125 | 2.05 | 1.32 | 0.21 | 4.89 | 0.82 | 3.23 | 0.31 |
| SOB $(\mathrm{g})$ | 125 | 670.81 | 944.18 | 0.95 | 3931.86 | 15.35 | 986.82 | 1.73 |
| Foliage $(\mathrm{g})$ | 125 | 250.88 | 328.92 | 0.90 | 1228.20 | 14.00 | 326.85 | 1.59 |
| Branches $(\mathrm{g})$ | 125 | 684.71 | 975.55 | 0.95 | 4006.10 | 14.25 | 85.05 | 1.78 |
| Roots $(\mathrm{g})$ | 125 | 263.01 | 395.35 | 0.50 | 1592.13 | 7.05 | 382.31 | 1.92 |
| Aboveground $(\mathrm{g})$ | 125 | 1606.40 | 2225.30 | 3.30 | 8836.06 | 41.50 | 2232.55 | 1.70 |
| Whole tree $(\mathrm{g})$ | 125 | 1869.41 | 2609.92 | 3.80 | 10133.37 | 48.45 | 2630.48 | 1.72 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 125 | 1395.86 | 2004.30 | 0.73 | 8203.87 | 24.37 | 2028.96 | 1.75 |

Table 27. European larch, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination ( $R^{2}$ ), mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -3.511 | 0.126 | $<0.001$ | 2.596 | 0.038 | $<0.001$ |  |  |  | 0.974 | 0.139 | 1.080 | 0.521 |
|  | Branches | -3.576 | 0.132 | <0.001 | 2.614 | 0.040 | <0.001 |  |  |  | 0.973 | 0.152 | 1.104 | 0.770 |
|  | Foliage | -2.825 | 0.133 | $<0.001$ | 2.191 | 0.040 | $<0.001$ |  |  |  | 0.961 | 0.154 | 1.098 | 0.708 |
|  | Roots | -3.962 | 0.131 | $<0.001$ | 2.453 | 0.039 | $<0.001$ |  |  |  | 0.970 | 0.150 | 1.105 | 0.820 |
|  | Aboveground part | -2.271 | 0.115 | $<0.001$ | 2.508 | 0.034 | $<0.001$ |  |  |  | 0.977 | 0.115 | 1.077 | 0.637 |
|  | Whole tree | -2.092 | 0.111 | $<0.001$ | 2.499 | 0.033 | $<0.001$ |  |  |  | 0.979 | 0.109 | 1.075 | 0.648 |
| [10] | Stem over bark | 3.671 | 0.057 | $<0.001$ | 2.705 | 0.061 | $<0.001$ |  |  |  | 0.942 | 0.319 | 1.188 | 1.026 |
|  | Branches | 3.678 | 0.075 | $<0.001$ | 2.667 | 0.079 | $<0.001$ |  |  |  | 0.902 | 0.546 | 1.288 | 1.093 |
|  | Foliage | 3.263 | 0.070 | <0.001 | 2.218 | 0.075 | <0.001 |  |  |  | 0.877 | 0.486 | 1.261 | 1.095 |
|  | Roots | 2.864 | 0.080 | <0.001 | 2.465 | 0.085 | <0.001 |  |  |  | 0.871 | 0.632 | 1.334 | 1.145 |
|  | Aboveground part | 4.683 | 0.065 | $<0.001$ | 2.576 | 0.070 | $<0.001$ |  |  |  | 0.918 | 0.418 | 1.230 | 1.052 |
|  | Whole tree | 4.840 | 0.067 | $<0.001$ | 2.559 | 0.071 | $<0.001$ |  |  |  | 0.913 | 0.437 | 1.237 | 1.043 |
| [11] | Stem over bark | -1.200 | 0.248 | $<0.001$ | 1.751 | 0.088 | $<0.001$ | 0.945 | 0.094 | $<0.001$ | 0.986 | 0.076 | 1.051 | 0.499 |
|  | Branches | -2.609 | 0.338 | $<0.001$ | 2.260 | 0.121 | $<0.001$ | 0.396 | 0.128 | 0.002 | 0.975 | 0.142 | 1.097 | 0.720 |
|  | Foliage | -2.448 | 0.351 | <0.001 | 2.053 | 0.125 | <0.001 | 0.154 | 0.133 | 0.248 | 0.962 | 0.153 | 1.097 | 0.699 |
|  | Roots | -3.993 | 0.348 | <0.001 | 2.464 | 0.124 | <0.001 | -0.013 | 0.132 | 0.924 | 0.970 | 0.151 | 1.106 | 0.822 |
|  | Aboveground part | -0.942 | 0.276 | $<0.001$ | 2.022 | 0.099 | $<0.001$ | 0.543 | 0.104 | $<0.001$ | 0.982 | 0.095 | 1.067 | 0.597 |
|  | Whole tree | -0.953 | 0.275 | $<0.001$ | 2.082 | 0.098 | $<0.001$ | 0.466 | 0.104 | $<0.001$ | 0.982 | 0.094 | 1.067 | 0.606 |

Table 28. European larch, $b_{0}, b_{1}$, $b_{2}$ regression coefficients, their standard errors (S.E.), p-values ( $P$ ), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE) for equations [13] - [15].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :--- | :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [13] | SOB volume | 0.389 | 0.088 | $<0.001$ | 2.169 | 0.049 | $<0.001$ |  |  |  | 0.960 | 349183 |
|  | SOB density | 897.714 | 36.406 | $<0.001$ | -0.153 | 0.013 | $<0.001$ |  |  | 0.530 | 4883 |  |
|  | R/S ratio | 0.195 | 0.019 | $<0.001$ | -0.055 | 0.029 | 0.058 |  |  |  | 0.025 | 0.002 |
|  | SOB volume | 44.476 | 11.589 | $<0.001$ | 3.395 | 0.172 | $<0.001$ |  |  |  | 0.868 | 1140699 |
| $[14]$ | SOB density | 587.870 | 6.987 | $<0.001$ | -0.161 | 0.013 | $<0.001$ |  |  | 0.531 | 4869 |  |
|  | R/S ratio | 0.172 | 0.005 | $<0.001$ | -0.117 | 0.029 | $<0.001$ |  |  | 0.099 | 0.002 |  |
|  | SOB volume | 1.075 | 0.151 | $<0.001$ | 1.520 | 0.047 | $<0.001$ | 1.323 | 0.082 | $<0.001$ | 0.987 | 116292 |
| $[15]$ | SOB density | 731.161 | 76.056 | $<0.001$ | -0.078 | 0.037 | 0.032 | -0.084 | 0.039 | 0.032 | 0.547 | 4745 |
|  | R/S ratio | 0.048 | 0.012 | $<0.001$ | 0.453 | 0.085 | $<0.001$ | -0.564 | 0.087 | $<0.001$ | 0.272 | 0.002 |



Fig. 31. Relationship of height a), dry mass of stem over bark b), dry mass of branches c), dry mass of foliage d), dry mass of roots e) and dry mass of the whole tree f) to stem base diameter $d_{0}$ of European larch.


Fig. 32. Relationship of the proportion of individual tree components a), volume of stem over bark b), basic density of stem over bark c), $R / S$ ratio d) to stem base diameter $d_{0}$ of European larch.

### 4.10. Rowan

Rowan (Sorbus aucuparia L.) is a rare tree species in Slovakia. From the point of its production potential and technical characteristics of the stem it does not belong to the important species for the wood-processing industry. It is a typical pioneer tree species, which increases its abundance after disturbances for a short period. In addition, it is a permanent component of forest communities in mountainous regions of $7^{\text {th }}$ spruce forest vegetation zone in Sorbeto-Piceetum group of forest types (predominantly at elevations from 1,250 to $1,550 \mathrm{~m}$ a.s.l.). Rowan is an important amelioration tree species, which can enhance soil conditions with its leaf litter. It is a fruit-producing tree species and an attractive species for wildlife (mainly red deer) browsing. This fact was also confirmed with the NFIM2 SR results, which showed that from all tree species rowan was attacked by ruminant ungulates most. Game damaged $17 \%$ of rowan individuals (expressed from basal area) by browsing and stripping, while the average damage intensity of all tree species in Slovakia was less than $3 \%$. Young rowan trees in the phase of growth were damaged most frequently, approximately every second individual. It can be assumed that the attractiveness of
rowan for game can be used as a biological protection of other tree species, or to increase the carrying capacity of hunting grounds.

According to the NFIM2 SR data, the minimum and maximum elevations at which rowan occurred were 112 m and $1,603 \mathrm{~m}$ a.s.l., respectively, while most frequently it grew at elevations $900-1,000 \mathrm{~m}$ a.s.l. From the point of its occurrence it is $13^{\text {th }}$ most common tree species in the forests of Slovakia, while from the point of its proportion of the stand stock it is ranked $35^{\text {th }}$. It grew at a reduced area of $32 \pm 9$ thousand ha, and occurred at $15 \%$ of the forested inventory plots.

The biomass regression models of rowan were derived from the data about 93 individuals. These trees were taken from five sites all located in the post-disturbance area of Vysoké Tatry (see Fig. 33). The trees covered the intervals of $d_{0}$ diameters from 4.95 mm to 80.95 mm , and heights from 0.41 m to 4.86 m (Table 29, Fig. 34a). The dry mass of the whole trees ranged from 7.15 g do $9,154.75 \mathrm{~g}$, and the stem volume ranged from $3.68 \mathrm{~cm}^{3}$ to $6,661.10 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 30. The volume of stem outside or inside bark, its density, as well as the root-shoot ratio were derived in a similar way (Table 31).

The regression models, scatter plots, and fitted regression curves were visualised in a similar way as in the case of the previous tree species. Further comments on the biomass of individual components and their proportions in the total tree biomass are presented in Chapter 4.12. (Interspecies comparison of biomass characteristics).


Fig. 33. Map of sample sites of Rowan and its distribution in the forests of Slovakia.

Table 29. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25. p), 75-percentile (75. p) and skewness of diameter ( $d_{0}$ ), tree height ( $h$ ), biomass of stem over bark (SOB), biomass of stem under bark (SUB), foliage biomass (foliage), branch biomass (branches), bark biomass (bark), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {SOB }}$ ) of individual trees.

| Rowan |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | 25.p | 75.p | Skewness |
| $\overline{d_{0}(\mathrm{~mm})}$ | 93 | 36.67 | 21.35 | 4.95 | 80.95 | 17.60 | 53.65 | 0.34 |
| $h(\mathrm{~m})$ | 93 | 2.82 | 1.21 | 0.41 | 4.86 | 1.78 | 3.99 | -0.12 |
| SOB (g) | 93 | 742.94 | 799.21 | 3.20 | 3144.05 | 83.05 | 1199.70 | 1.09 |
| SUB (g) | 93 | 626.35 | 690.01 | 1.90 | 2749.40 | 58.50 | 1003.00 | 1.13 |
| Foliage (g) | 93 | 169.74 | 202.07 | 1.55 | 915.60 | 18.90 | 283.10 | 1.62 |
| Branches (g) | 89 | 283.49 | 472.64 | 0.00 | 2715.30 | 8.00 | 372.60 | 2.86 |
| Bark (g) | 93 | 116.60 | 110.73 | 1.30 | 418.75 | 22.50 | 190.30 | 0.91 |
| Roots (g) | 93 | 432.57 | 525.73 | 2.40 | 2379.80 | 39.40 | 707.80 | 1.49 |
| Aboveground (g) | 89 | 1154.26 | 1396.28 | 4.75 | 6774.95 | 119.75 | 1766.35 | 1.61 |
| Whole tree (g) | 89 | 1564.29 | 1892.83 | 7.15 | 9154.75 | 167.20 | 2407.10 | 1.61 |
| $V_{S O B}\left(\mathrm{~cm}^{3}\right)$ | 93 | 1472.96 | 1577.64 | 3.68 | 6661.10 | 145.16 | 2465.56 | 1.07 |

Table 30. Rowan, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values ( $P$ ), coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -2.515 | 0.088 | $<0.001$ | 2.412 | 0.025 | $<0.001$ |  |  |  | 0.990 | 0.028 | 1.014 | 0.171 |
|  | Stem under bark | -3.218 | 0.094 | <0.001 | 2.546 | 0.027 | <0.001 |  |  |  | 0.990 | 0.033 | 1.016 | 0.183 |
|  | Branches | -7.336 | 0.312 | <0.001 | 3.334 | 0.090 | <0.001 |  |  |  | 0.943 | 0.280 | 1.138 | 0.596 |
|  | Foliage | -3.383 | 0.161 | <0.001 | 2.237 | 0.046 | <0.001 |  |  |  | 0.962 | 0.096 | 1.047 | 0.322 |
|  | Bark | -2.591 | 0.092 | $<0.001$ | 1.966 | 0.027 | <0.001 |  |  |  | 0.984 | 0.032 | 1.016 | 0.179 |
|  | Roots | -3.396 | 0.169 | $<0.001$ | 2.476 | 0.049 | <0.001 |  |  |  | 0.966 | 0.105 | 1.056 | 0.382 |
|  | Aboveground part | -2.423 | 0.099 | $<0.001$ | 2.499 | 0.029 | <0.001 |  |  |  | 0.989 | 0.034 | 1.017 | 0.190 |
|  | Whole tree | -2.067 | 0.104 | <0.001 | 2.485 | 0.030 | $<0.001$ |  |  |  | 0.987 | 0.038 | 1.019 | 0.198 |
| [10] | Stem over bark | 2.861 | 0.078 | <0.001 | 3.066 | 0.073 | $<0.001$ |  |  |  | 0.951 | 0.142 | 1.074 | 0.441 |
|  | Stem under bark | 2.453 | 0.080 | $<0.001$ | 3.242 | 0.075 | <0.001 |  |  |  | 0.953 | 0.149 | 1.078 | 0.450 |
|  | Branches | -0.018 | 0.193 | 0.927 | 4.332 | 0.182 | $<0.001$ |  |  |  | 0.871 | 0.632 | 1.393 | 1.443 |
|  | Foliage | 1.651 | 0.109 | $<0.001$ | 2.792 | 0.103 | <0.001 |  |  |  | 0.890 | 0.278 | 1.154 | 0.704 |
|  | Bark | 1.790 | 0.067 | $<0.001$ | 2.501 | 0.063 | <0.001 |  |  |  | 0.946 | 0.104 | 1.053 | 0.358 |
|  | Roots | 2.188 | 0.122 | $<0.001$ | 3.076 | 0.116 | <0.001 |  |  |  | 0.886 | 0.352 | 1.196 | 0.798 |
|  | Aboveground part | 3.186 | 0.094 | $<0.001$ | 3.135 | 0.090 | <0.001 |  |  |  | 0.934 | 0.202 | 1.112 | 0.584 |
|  | Whole tree | 3.522 | 0.100 | <0.001 | 3.104 | 0.095 | <0.001 |  |  |  | 0.924 | 0.227 | 1.125 | 0.615 |
| [11] | Stem over bark | -1.317 | 0.161 | $<0.001$ | 1.857 | 0.071 | $<0.001$ | 0.749 | 0.092 | $<0.001$ | 0.994 | 0.017 | 1.008 | 0.129 |
|  | Stem under bark | -1.843 | 0.164 | $<0.001$ | 1.909 | 0.072 | $<0.001$ | 0.859 | 0.093 | <0.001 | 0.995 | 0.017 | 1.008 | 0.130 |
|  | Branches | -6.976 | 0.690 | <0.001 | 3.162 | 0.308 | $<0.001$ | 0.244 | 0.416 | 0.560 | 0.943 | 0.282 | 1.137 | 0.592 |
|  | Foliage | -3.389 | 0.390 | <0.001 | 2.240 | 0.171 | <0.001 | -0.004 | 0.222 | 0.987 | 0.962 | 0.097 | 1.047 | 0.322 |
|  | Bark | -1.575 | 0.191 | <0.001 | 1.495 | 0.084 | <0.001 | 0.635 | 0.109 | <0.001 | 0.988 | 0.023 | 1.011 | 0.153 |
|  | Roots | -3.684 | 0.408 | <0.001 | 2.610 | 0.179 | $<0.001$ | -0.181 | 0.232 | 0.438 | 0.966 | 0.106 | 1.056 | 0.379 |
|  | Aboveground part | -1.666 | 0.220 | $<0.001$ | 2.149 | 0.096 | $<0.001$ | 0.470 | 0.124 | $<0.001$ | 0.990 | 0.030 | 1.015 | 0.178 |
|  | Whole tree | -1.586 | 0.243 | <0.001 | 2.262 | 0.106 | <0.001 | 0.299 | 0.137 | 0.032 | 0.988 | 0.036 | 1.018 | 0.196 |

Table 31. Rowan, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values $(P)$, coefficients of determination $\left(R^{2}\right)$, mean square errors (MSE) for equations [13] - [15].

| Eq. | Dependent variable | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [13] | SOB volume | 0,557 | 0,189 | 0,004 | 2,101 | 0,082 | $<0,001$ |  |  |  | 0,951 | 123665 |
|  | SUB volume | 0,356 | 0,127 | 0,006 | 2,163 | 0,085 | <0,001 |  |  |  | 0,950 | 88720 |
|  | Bark volume | 0,299 | 0,102 | 0,004 | 1,827 | 0,082 | <0,001 |  |  |  | 0,932 | 4742 |
|  | SOB density | 821,786 | 33,646 | <0,001 | -0,130 | 0,012 | $<0,001$ |  |  |  | 0,527 | 2028 |
|  | SUB density | 719,209 | 28,331 | <0,001 | -0,087 | 0,012 | $<0,001$ |  |  |  | 0,364 | 1819 |
|  | Bark density | 1271,188 | 78,324 | <0,001 | -0,275 | 0,019 | $<0,001$ |  |  |  | 0,668 | 4913 |
|  | Bark mass proportion | 90,963 | 3,515 | <0,001 | -0,438 | 0,013 | $<0,001$ |  |  |  | 0,925 | 4,06 |
|  | Bark volume proportion | 63,813 | 3,617 | <0,001 | -0,316 | 0,018 | $<0,001$ |  |  |  | 0,762 | 8,39 |
|  | R/S ratio | 0,453 | 0,082 | <0,001 | -0,064 | 0,053 | 0,234 |  |  |  | 0,015 | 0,016 |
| [14] | SOB volume | 32,970 | 11,634 | 0,006 | 3,205 | 0,246 | <0,001 |  |  |  | 0,855 | 363668 |
|  | SUB volume | 23,916 | 8,915 | 0,009 | 3,296 | 0,260 | <0,001 |  |  |  | 0,850 | 263797 |
|  | Bark volume | 10,107 | 2,971 | 0,001 | 2,808 | 0,207 | $<0,001$ |  |  |  | 0,860 | 9840 |
|  | SOB density | 622,663 | 8,229 | <0,001 | -0,178 | 0,013 | <0,001 |  |  |  | 0,629 | 1592 |
|  | SUB density | 599,732 | 8,057 | <0,001 | -0,126 | 0,013 | $<0,001$ |  |  |  | 0,469 | 1518 |
|  | Bark density | 681,152 | 13,846 | <0,001 | -0,336 | 0,022 | $<0,001$ |  |  |  | 0,683 | 4686 |
|  | Bark mass proportion | 32,650 | 0,529 | <0,001 | -0,491 | 0,019 | $<0,001$ |  |  |  | 0,866 | 7,29 |
|  | Bark volume proportion | 30,893 | 0,601 | <0,001 | -0,373 | 0,021 | $<0,001$ |  |  |  | 0,746 | 8,96 |
|  | R/S ratio | 0,408 | 0,027 | $<0,001$ | -0,125 | 0,065 | 0,057 |  |  |  | 0,038 | 0,016 |
| [15] | SOB volume | 0,880 | 0,286 | 0,003 | 1,697 | 0,112 | $<0,001$ | 0,862 | 0,188 | $<0,001$ | 0,960 | 101061 |
|  | SUB volume | 0,555 | 0,189 | 0,004 | 1,759 | 0,116 | <0,001 | 0,870 | 0,196 | <0,001 | 0,959 | 73310 |
|  | Bark volume | 0,512 | 0,176 | 0,005 | 1,413 | 0,125 | $<0,001$ | 0,836 | 0,209 | $<0,001$ | 0,943 | 4049 |
|  | SOB density | 520,806 | 47,552 | <0,001 | 0,078 | 0,039 | 0,050 | -0,272 | 0,049 | <0,001 | 0,644 | 1544 |
|  | SUB density | 477,880 | 41,915 | <0,001 | 0,100 | 0,038 | 0,010 | -0,247 | 0,048 | <0,001 | 0,507 | 1427 |
|  | Bark density | 864,156 | 138,766 | <0,001 | -0,103 | 0,069 | 0,140 | -0,217 | 0,083 | 0,010 | 0,690 | 4627 |
|  | Bark mass proportion | 82,974 | 9,027 | <0,001 | -0,398 | 0,047 | <0,001 | -0,048 | 0,054 | 0,370 | 0,926 | 4,07 |
|  | Bark volume proportion | 49,369 | 7,498 | $<0,001$ | -0,201 | 0,065 | 0,003 | -0,142 | 0,078 | 0,072 | 0,770 | 8,18 |
|  | R/S ratio | 0,172 | 0,075 | 0,024 | 0,375 | 0,186 | 0,004 | -0,572 | 0,231 | 0,015 | 0,084 | 0,016 |



Fig. 34. Relationship of tree height a) and $d_{1.3}$ diameter b) to $d_{0}$ stem base diameter of Rowan.


Fig. 35. Relationship of dry mass of stem over bark a), dry mass of branches b), dry mass of foliage c), dry mass of roots d) and dry mass of the whole tree e) and proportion of individual tree componentsf) to stem base diameter $d_{0}$ of Rowan.


Fig. 36. Relationship of the basic density of stem over bark a), basic bark density b), bark mass proportion of SOB mass c), bark volume proportion of $S O B$ volume d) volume of stem over bark e), and $R / S$ ratio f) to stem base diameter $d_{0}$ of Rowan.

### 4.11. Common aspen

Common aspen (Populus tremula L.) is a pioneer tree species with an important economic significance. Its proportion in the forests of Slovakia is small, based on its occurrence it is ranked $17^{\text {th }}$, from the point of the stock and area it was ranked $18^{\text {th }}$ and $16^{\text {th }}$, respectively. It usually occurs in middle forest vegetation zones from $2^{\text {nd }}$ beech-oak zone up to $6^{\text {th }}$ spruce-beech-fir zone, most frequently at lower elevations. It is not an important tree species in the biotopes of European and national significance, and it is not the main tree species in any biotope. Aspen is an attractive tree species for the nutrition of red deer, hence it increases the carrying capacity of hunting grounds and can ensure biological protection of economically important tree species.

According to the NFIM2 SR data, the minimum and maximum elevations at which aspen occurred were 199 m and $1,466 \mathrm{~m}$ a.s.l., respectively, while most frequently it occurred at elevations $450-550 \mathrm{~m}$ a.s.l. It grew at a reduced area of $17 \pm 7$ thousand ha, and occurred at $8 \%$ of the forested inventory plots.

The biomass regression models for the Common aspen were derived from the data measured at 185 individuals. The trees were taken from seven sites (see Fig. 37) located in orographic units of Kremnické vrchy (1 and 2), Štiavnické vrchy (3 and 4), Krupinská planina (5), Malá Fatra (6) and Nízke Tatry (7). The trees covered the intervals of $d_{0}$ diameters from 3.30 mm to 100.90 mm , and heights from 0.40 m to 10.54 m (Table 32, Fig. 38a). The dry mass of the whole trees ranged from 6.25 g do $15,650.85 \mathrm{~g}$, and the stem volume ranged from $2.76 \mathrm{~cm}^{3}$ to $24,500.28 \mathrm{~cm}^{3}$.

The regression models (i.e. the coefficients and basic statistical characteristics) for the calculation of the biomass in individual tree components, as well as of the whole tree are presented in Table 33. The volume of stem outside or inside bark, its density, as well as the root-shoot ratio were derived using the same approach (Table 34).
The regression models, scatter plots, and fitted regression curves were visualised in a similar way as in the case of the other tree species. The text describing the biomass of individual components and their proportions in the total tree biomass is presented in Chapter 4.12. (Inter-species comparison of biomass characteristics).


Fig. 37. Map of sample sites of Common aspen and its distribution in the forests of Slovakia.

Table 32. Number ( $N$ ), mean, standard deviation (SD), minimum, maximum, 25-percentile (25. p), 75-percentile (75. p) and skewness of diameter ( $d_{0}$ ), tree height ( $h$ ), biomass of stem over bark (SOB), biomass of stem under bark (SUB), foliage biomass (foliage), branch biomass (branches), bark biomass (bark), root biomass (roots), aboveground biomass (aboveground), total tree biomass (tree), and volume of stem over bark ( $V_{\text {SOB }}$ ) of individual trees.

|  | Common aspen |  |  |  |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | N | Mean | SD | Min | Max | $25 . \mathrm{p}$ | 75.p | Skewness |
| $d_{0}(\mathrm{~mm})$ | 185 | 31.39 | 20.00 | 3.30 | 100.90 | 15.50 | 43.10 | 0.89 |
| $h(\mathrm{~m})$ | 185 | 3.81 | 2.42 | 0.40 | 10.54 | 1.87 | 4.96 | 0.75 |
| SOB $(\mathrm{g})$ | 182 | 935.07 | 1654.58 | 1.23 | 10474.65 | 4.29 | 89.89 | 3.10 |
| SUB $(\mathrm{g})$ | 182 | 699.67 | 1293.38 | 0.76 | 8351.95 | 29.95 | 673.93 | 3.26 |
| Foliage $(\mathrm{g})$ | 181 | 93.54 | 130.82 | 0.75 | 736.85 | 10.92 | 128.86 | 2.57 |
| Branches $(\mathrm{g})$ | 179 | 251.49 | 460.01 | 0.00 | 2770.00 | 11.89 | 283.65 | 3.45 |
| Bark $(\mathrm{g})$ | 185 | 235.29 | 364.10 | 0.47 | 2122.70 | 18.34 | 275.70 | 2.57 |
| Roots $(\mathrm{g})$ | 183 | 262.48 | 375.57 | 1.20 | 2190.00 | 23.25 | 310.26 | 2.57 |
| Aboveground $(\mathrm{g})$ | 174 | 1193.43 | 2151.36 | 3.15 | 13460.85 | 69.25 | 1235.58 | 3.38 |
| Whole tree $(\mathrm{g})$ | 172 | 1455.66 | 2530.23 | 6.25 | 15650.85 | 82.88 | 1538.25 | 3.27 |
| $V_{\text {SOB }}\left(\mathrm{cm}^{3}\right)$ | 185 | 2232.89 | 3979.88 | 2.76 | 24500.28 | 106.71 | 2139.86 | 3.22 |

Table 33. Common aspen, $b_{0}, b_{1}, b_{2}$ regression coefficients, their standard errors (S.E.), p-values (P), coefficients of determination ( $R^{2}$ ), mean square errors (MSE), logarithmic transformation bias $\lambda$ and its standard deviation (S.D.) for equations [9] - [11].

| Eq. | Tree component | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE | $\lambda$ | S.D. |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [9] | Stem over bark | -3.612 | 0.107 | $<0.001$ | 2.795 | 0.033 | $<0.001$ |  |  |  | 0.976 | 0.098 | 1.048 | 0.323 |
|  | Stem under bark | -4.454 | 0.120 | $<0.001$ | 2.932 | 0.036 | $<0.001$ |  |  |  | 0.973 | 0.123 | 1.061 | 0.367 |
|  | Branches | -5.683 | 0.196 | $<0.001$ | 3.010 | 0.060 | $<0.001$ |  |  |  | 0.935 | 0.306 | 1.116 | 0.710 |
|  | Foliage | -2.907 | 0.225 | $<0.001$ | 2.020 | 0.069 | $<0.001$ |  |  |  | 0.829 | 0.429 | 1.191 | 0.634 |
|  | Bark | -3.906 | 0.104 | $<0.001$ | 2.528 | 0.031 | $<0.001$ |  |  |  | 0.973 | 0.092 | 1.045 | 0.313 |
|  | Roots | -3.315 | 0.130 | $<0.001$ | 2.410 | 0.039 | $<0.001$ |  |  |  | 0.954 | 0.145 | 1.078 | 0.474 |
|  | Aboveground part | -2.853 | 0.103 | $<0.001$ | 2.687 | 0.032 | $<0.001$ |  |  |  | 0.977 | 0.088 | 1.042 | 0.296 |
|  | Whole tree | -2.379 | 0.092 | $<0.001$ | 2.618 | 0.028 | $<0.001$ |  |  |  | 0.981 | 0.070 | 1.036 | 0.293 |
| [10) | Stem over bark | 2.541 | 0.055 | $<0.001$ | 2.602 | 0.042 | $<0.001$ |  |  |  | 0.956 | 0.182 | 1.090 | 0.450 |
|  | Stem under bark | 1.986 | 0.054 | $<0.001$ | 2.743 | 0.040 | $<0.001$ |  |  |  | 0.962 | 0.172 | 1.085 | 0.439 |
|  | Branches | 1.114 | 0.117 | $<0.001$ | 2.653 | 0.089 | $<0.001$ |  |  |  | 0.834 | 0.786 | 1.402 | 1.154 |
|  | Foliage | 1.676 | 0.111 | $<0.001$ | 1.755 | 0.085 | $<0.001$ |  |  |  | 0.706 | 0.736 | 1.350 | 1.009 |
|  | Bark | 1.680 | 0.060 | $<0.001$ | 2.336 | 0.045 | <0.001 |  |  |  | 0.937 | 0.211 | 1.107 | 0.502 |
|  | Roots | 2.034 | 0.076 | $<0.001$ | 2.198 | 0.057 | $<0.001$ |  |  |  | 0.893 | 0.336 | 1.166 | 0.641 |
|  | Aboveground part | 3.108 | 0.069 | $<0.001$ | 2.451 | 0.053 | $<0.001$ |  |  |  | 0.924 | 0.283 | 1.145 | 0.611 |
|  | Whole tree | 3.425 | 0.069 | $<0.001$ | 2.386 | 0.053 | $<0.001$ |  |  |  | 0.923 | 0.275 | 1.137 | 0.576 |
| [11] | Stem over bark | -1.354 | 0.162 | $<0.001$ | 1.741 | 0.071 | $<0.001$ | 1.040 | 0.067 | <0.001 | 0.990 | 0.042 | 1.021 | 0.210 |
|  | Stem under bark | -1.759 | 0.163 | $<0.001$ | 1.674 | 0.072 | $<0.001$ | 1.240 | 0.067 | <0.001 | 0.991 | 0.043 | 1.021 | 0.211 |
|  | Branches | -6.312 | 0.449 | $<0.001$ | 3.303 | 0.197 | $<0.001$ | -0.286 | 0.184 | 0.122 | 0.936 | 0.304 | 1.162 | 0.689 |
|  | Foliage | -4.306 | 0.505 | $<0.001$ | 2.674 | 0.223 | <0.001 | -0.645 | 0.210 | 0.002 | 0.837 | 0.409 | 1.184 | 0.637 |
|  | Bark | -2.319 | 0.199 | $<0.001$ | 1.787 | 0.088 | <0.001 | 0.732 | 0.082 | $<0.001$ | 0.981 | 0.065 | 1.033 | 0.271 |
|  | Roots | -2.593 | 0.295 | $<0.001$ | 2.072 | 0.130 | $<0.001$ | 0.334 | 0.123 | 0.007 | 0.956 | 0.140 | 1.074 | 0.448 |
|  | Aboveground part | -1.692 | 0.219 | $<0.001$ | 2.145 | 0.096 | $<0.001$ | 0.533 | 0.090 | $<0.001$ | 0.981 | 0.073 | 1.036 | 0.281 |
|  | Whole tree | -1.434 | 0.200 | $<0.001$ | 2.177 | 0.088 | <0.001 | 0.434 | 0.083 | $<0.001$ | 0.983 | 0.060 | 1.031 | 0.269 |

Table 34. Common aspen, $b_{0}, b_{1}$, $b_{2}$ regression coefficients, their standard errors (S.E.), $p$-values ( $P$ ), coefficients of determination ( $R^{2}$ ), mean square errors (MSE) for equations [13] - [15].

| Eq. | Dependent variable | $b_{0}$ | S.E. | P | $b_{1}$ | S.E. | P | $b_{2}$ | S.E. | P | $\mathrm{R}^{2}$ | MSE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| [13] | SOB volume | 0.066 | 0.015 | $<0.001$ | 2.792 | 0.051 | $<0.001$ |  |  |  | 0.962 | 553328 |
|  | SUB volume | 0.038 | 0.009 | $<0.001$ | 2.865 | 0.056 | <0.001 |  |  |  | 0.954 | 405034 |
|  | Bark volume | 0.043 | 0.009 | $<0.001$ | 2.528 | 0.049 | <0.001 |  |  |  | 0.956 | 25170 |
|  | SOB density | 603.039 | 23.540 | $<0.001$ | -0.093 | 0.012 | <0.001 |  |  |  | 0.232 | 2908 |
|  | SUB density | 442.437 | 16.643 | $<0.001$ | -0.022 | 0.012 | 0.058 |  |  |  | 0.019 | 2111 |
|  | Bark density | 953.096 | 49.181 | $<0.001$ | -0.180 | 0.017 | <0.001 |  |  |  | 0.371 | 8153 |
|  | Bark mass proportion | 73.579 | 3.251 | <0.001 | -0.260 | 0.015 | $<0.001$ |  |  |  | 0.628 | 23.73 |
|  | Bark volume proportion | 49.011 | 2.586 | $<0.001$ | -0.186 | 0.017 | $<0.001$ |  |  |  | 0.390 | 21.90 |
|  | R/S ratio | 0.823 | 0.114 | $<0.001$ | -0.338 | 0.048 | $<0.001$ |  |  |  | 0.211 | 0.019 |
| (14] | SOB volume | 10.943 | 2.555 | <0.001 | 3.176 | 0.107 | <0.001 |  |  |  | 0.914 | 1200225 |
|  | SUB volume | 6.331 | 1.540 | $<0.001$ | 3.322 | 0.111 | <0.001 |  |  |  | 0.917 | 735196 |
|  | Bark volume | 6.315 | 1.424 | $<0.001$ | 2.699 | 0.105 | <0.001 |  |  |  | 0.878 | 70190 |
|  | SOB density | 499.005 | 6.590 | <0.001 | -0.103 | 0.011 | <0.001 |  |  |  | 0.323 | 2564 |
|  | SUB density | 426.242 | 5.893 | <0.001 | -0.031 | 0.011 | 0.004 |  |  |  | 0.043 | 2060 |
|  | Bark density | 651.553 | 10.928 | <0.001 | -0.185 | 0.014 | <0.001 |  |  |  | 0.454 | 7085 |
|  | Bark mass proportion | 42.352 | 0.518 | <0.001 | -0.262 | 0.011 | <0.001 |  |  |  | 0.748 | 16.1 |
|  | Bark volume proportion | 33.069 | 0.567 | $<0.001$ | -0.189 | 0.015 | <0.001 |  |  |  | 0.469 | 19.07 |
|  | $\mathrm{R} /$ S ratio | 0.395 | 0.018 | $<0.001$ | -0.323 | 0.043 | $<0.001$ |  |  |  | 0.233 | 0.019 |
| [15] | SOB volume | 0.385 | 0.065 | <0.001 | 1.809 | 0.067 | $<0.001$ | 1.157 | 0.078 | <0.001 | 0.983 | 235109 |
|  | SUB volume | 0.262 | 0.048 | <0.001 | 1.775 | 0.073 | <0.001 | 1.295 | 0.087 | <0.001 | 0.981 | 170898 |
|  | Bark volume | 0.141 | 0.031 | <0.001 | 1.894 | 0.087 | <0.001 | 0.725 | 0.094 | <0.001 | 0.968 | 18440 |
|  | SOB density | 381.359 | 32.960 | <0.001 | 0.119 | 0.038 | 0.002 | -0.208 | 0.035 | <0.001 | 0.359 | 2441 |
|  | SUB density | 339.056 | 29.066 | $<0.001$ | 0.102 | 0.038 | 0.007 | -0.122 | 0.035 | <0.001 | 0.080 | 1990 |
|  | Bark density | 534.357 | 63.217 | <0.001 | 0.088 | 0.052 | 0.092 | -0.262 | 0.048 | <0.001 | 0.462 | 7010 |
|  | Bark mass proportion | 33.678 | 3.068 | $<0.001$ | 0.101 | 0.040 | 0.012 | -0.351 | 0.036 | <0.001 | 0.757 | 15.62 |
|  | Bark volume proportion | 26.944 | 3.272 | $<0.001$ | 0.091 | 0.053 | 0.089 | -0.270 | 0.049 | <0.001 | 0.478 | 18.87 |
|  | R/S ratio | 0.402 | 0.141 | 0.005 | -0.007 | 0.155 | 0.964 | -0.317 | 0.141 | 0.026 | 0.233 | 0.019 |



Fig. 38. Relationship of tree height a) and $d_{1.3}$ diameter $b$ ) to stem base diameter $d_{0}$ of Common aspen.







Diameter $\mathrm{d}_{0}(\mathrm{~mm})$
Fig. 39. Relationship of dry mass of stem over bark a), dry mass of branches b), dry mass of foliage c), dry mass of roots d) and dry mass of the whole tree e) and proportion of individual tree componentsf) to stem base diameter $d_{0}$ of Common aspen.


Fig. 40. Relationship of the basic density of stem over bark a), basic bark density b), mass proportion of bark in SOB mass c), volume proportion of bark in SOB volume d), volume of stem over bark e), and $R / S$ ratiof) to stem base diameter $d_{0}$ of Common aspen.

### 4.12. Inter-species comparison of hiomass characteristics

Since our results on tree species biomass dealt with eleven species, we had a possibility to compare their inter-species differences. We assumed that the differences in biomass allocation can exist between individual tree species, as well as between groups of tree species. At a tree species group level, we expected differences due to the differences in the leaf-fall cycle (i.e. evergreen versus deciduous) or due to their ecological demands (mainly light demanding versus shade tolerant). Hence, in the following text we will focus on graphical visualisation of potential interspecies differences and their subsequent interpretation.


Fig. 41. Inter-tree species comparison of the relationships of tree height to stem base diameter $d_{0}$.
The height curves based on $d_{0}$ diameter as an independent variable showed substantial interspecies differences. We graphically divided the most important tree species (spruce, pine, beech, oak, larch, hornbeam; Fig. 41a) from the valuable (ash and sycamore) and pioneer tree species (rowan, aspen, and goat willow; Fig. 41b). If we consider equal diameters for all analysed tree species, we see that aspen was the highest tree species followed by sycamore, hornbeam, and oak. Beech was shorter, and the group of shortest trees (very similar in height) comprised ash, rowan, larch, spruce, goat willow, and pine. The relationship of height to $d_{0}$ diameter was described using the regression function [18]. The statistical characteristics of this function for individual tree species are in Table 35.

Table 35. Regression coefficients $b_{0}, b_{1}, b_{2}$, their standard errors (S.E.), p-values ( $P$ ), coefficients of determination $\left(R^{2}\right)$, and mean square errors (MSE) of regression functions describing the relationship of tree height to $d_{0}$ diameter.

| Tree species | $b_{0}$ | S.E. | p | $b_{1}$ | S.E. | p | $b_{2}$ | S.E. | p | $\mathrm{R}^{2}$ | MSE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Beech | 0.493 | 12.795 | 0.969 | 11.204 | 1.279 | <0.001 | 0.019 | 0.747 | 0.456 | 0.779 | 117.311 |
| Spruce | 44.751 | 19.323 | 0.022 | 11.816 | 1.316 | $<0.001$ | 0.103 | 0.17 | $<0.001$ | 0.937 | 150.708 |
| Hornbeam | 16.924 | 6.036 | 0.006 | 3.254 | 0.527 | $<0.001$ | 0.093 | 0.009 | $<0.001$ | 0.864 | 701.942 |
| Oak | 137.581 | 25.176 | <0.001 | 1.971 | 1.385 | 0.157 | 0.127 | 0.017 | $<0.001$ | 0.920 | 424.214 |
| Pine | 52.966 | 38.272 | 0.168 | 9.260 | 2.694 | $<0.001$ | 0.164 | 0.039 | $<0.001$ | 0.741 | 198.814 |
| Sycamore | 84.237 | 25.897 | 0.001 | 3.273 | 1.311 | 0.014 | 0.073 | 0.014 | $<0.001$ | 0.882 | 723.968 |
| Ash | 125.858 | 45.116 | 0.007 | 0.395 | 3.445 | 0.909 | 0.193 | 0.061 | 0.002 | 0.861 | 150.077 |
| Goat willow | 6.921 | 4.691 | 0.143 | 8.127 | 0.542 | $<0.001$ | 0.126 | 0.016 | $<0.001$ | 0.812 | 237.939 |
| Larch | 26.626 | 18.865 | 0.161 | 10.506 | 1.126 | $<0.001$ | 0.113 | 0.013 | $<0.001$ | 0.929 | 281.182 |
| Rowan | 14.640 | 13.260 | 0.273 | 6.486 | 0.969 | $<0.001$ | 0.135 | 0.014 | $<0.001$ | 0.933 | 287.626 |
| Aspen | 8.221 | 11.812 | 0.487 | 7.077 | 0.693 | $<0.001$ | 0.021 | 0.009 | 0.017 | 0.897 | 1218.609 |



Fig. 42. Inter-tree species comparison of the relationships of $d_{1,3}$ diameter to stem base diameter $d_{0}$.
Next, we compared the regression relationships of $d_{0}$ diameter to $d_{1.3}$ diameter between the individual tree species (in the case of the main tree species, i.e. beech, oak, spruce, and pine, which we started our project with, the data on $d_{1.3}$ diameter were missing; see Fig. 42), and the relationships were fitted with linear function [17]. This relationship reflects the stem taper in its bottom part, the greater the values of $d_{1.3}$ diameter in relation to $d_{0}$ diameter are, the less taper the stem is. Based on this relationship, aspen had the least tapered stem, while larch and goat willow had the most tapered stems. The stem tapers of the other tree species (i.e. sycamore, hornbeam, ash, and rowan) were in the middle of the analysed group. The statistical characteristics of the linear functions for individual tree species are in Table 36.

Table 36. Regression coefficients $b_{0}, b_{1}, b_{2}$, their standard errors (S.E.), p-values ( $P$ ), coefficients of determination ( $R^{2}$ ), and mean square errors (MSE) of regression functions describing the relationship of $d_{1.3}$ diameter to $d_{0}$ diameter.

| Tree species | $b_{0}$ | $\mathrm{~S}, \mathrm{E}$, | P | $b_{1}$ | $\mathrm{~S}, \mathrm{E}$, | P | $\mathrm{R}^{2}$ |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Hornbeam | -2.023 | 0.378 | $<0.001$ | 0.664 | 0.013 | $<0.001$ | 0.946 | $\mathrm{MSE}^{2}$ |
| Sycamore | -4.948 | 0.732 | $<0.001$ | 0.718 | 0.017 | $<0.001$ | 0.950 |  |
| Ash | -3.446 | 1.275 | 0.009 | 0.664 | 0.042 | $<0.001$ | 0.827 |  |
| Goat willow | -5.285 | 0.971 | $<0.001$ | 0.620 | 0.031 | $<0.001$ | 0.838 |  |
| Larch | -9.967 | 1.434 | $<0.001$ | 0.698 | 0.023 | $<0.001$ | 0.925 | 11.58 |
| Rowan | -3.629 | 0.711 | $<0.001$ | 0.665 | 0.016 | $<0.001$ | 0.958 |  |
| Aspen | -3.422 | 0.911 | $<0.001$ | 0.763 | 0.022 | $<0.001$ | 0.890 | 7.83 |

The observations of inter-species differences in the total tree biomass in relation to $d_{0}$ diameter revealed interesting information. We divided the tree species to two groups, one group comprising the species with greater quantities (with dry mass exceeding 10 kg at $d_{0}$ diameter equal to 100 mm ; Fig. 43A), namely beech, hornbeam, oak, sycamore, larch, and aspen, and the other group containing the species with lower quantities (below 10 kg ; Fig. 43B), which comprised spruce, pine, ash, goat willow, and rowan. This inter-species comparison indicated that at a specific $d_{0}$ diameter the greatest tree biomass was found for hornbeam and aspen, and the lowest biomass was found for goat willow and pine. However, the reasons for these differences are not clear, and hence, their interpretation is not simple, either. We cannot unambiguously state that at a particular $d_{0}$ diameter a specific group of tree species had a substantially different biomass amount than the other group (e.g. light demanding versus shade tolerant, or coniferous versus broadleaved). We use $d_{0}$ diameter equal to 50 mm as a basis for further interpretation of interspecies differences. Using our models, we estimated the following amounts of tree biomass at this diameter (tree species were ranked bottom-up on the base of their mass): goat willow - 1.65 kg , pine -1.66 kg , rowan -2.15 kg , spruce -2.27 kg , sycamore -2.33 kg , larch -2.34 kg , oak -2.46 kg , beech -2.58 kg , ash $-2,65 \mathrm{~kg}$, aspen -2.69 kg , hornbeam -2.90 kg .


Fig. 43. Inter-tree species comparison of the relationships of tree dry mass to stem base diameter $d_{0}$.


Fig. 44. Inter-tree species comparison of the relationships of volume of stem over bark to stem base diameter $d_{0}$.
Similarly we compared the inter-species differences in the volume of stem over bark (Fig. 44a, 44b). Also in this case we divided the tree species to two groups, one comprising the tree species with the volume of stem over bark exceeding $8,000 \mathrm{~cm}^{3}$ at $d_{0}$ diameter equal to 100 mm (beech, hornbeam, oak, sycamore, ash, and aspen), and the other with the volume below $8,000 \mathrm{~cm}^{3}$ (spruce, pine, goat willow, larch, rowan). Next, we compared the dry mass of stem over bark. At $d_{0}$ diameter equal to 50 mm , we estimated the following quantities of stem over bark for the individual tree species: goat willow -0.71 kg , spruce -0.72 kg , pine -0.74 kg , larch -0.83 kg , rowan -1.03 kg , beech -1.34 kg , sycamore -1.44 kg , aspen -1.59 kg , oak -1.60 kg , ash -1.67 kg , hornbeam -2.10 kg .


Fig. 45. Inter-tree species comparison of the relationships of the basic density of stem over bark to stem base diameter $d_{0}$.


Fig. 46. Inter-tree species comparison of the relationships of bark mass a) and bark volume proportion b) of SOB to stem base diameter $d_{0}$.

Next, we graphically compared the basic density of stem over bark (Fig. 45a, 45b). Basic density of stem over bark is a variable with a great variation of values that depends on a number of factors. The most important factors are site conditions, climatic conditions, and silvicultural treatments. The greatest density was found for the smallest individuals irrespective of tree species. The value of the basic density of stem over bark decreased with the increasing tree size for the majority of the observed tree species, while the reduction was first rapid, then slow. After reaching diameter $d_{0}=40 \mathrm{~mm}$, their values changed only a little, and did not depend on $d_{0}$ diameter. This did not hold for spruce, pine, and larch, for which we observed the dependence on $d_{0}$ diameter in the whole range of diameters. Oak and beech had the greatest density, while aspen and goat willow had the lowest density of stem over bark. The bark mass (Fig. 46a) and volume (Fig. 46b) proportion of stem over bark (SOB) were derived similarly. The highest bark proportion was found for aspen and goat willow, and the lowest for hornbeam and sycamore.


Fig. 47. Inter-species comparison of the relationship of the ratios between bark density and wood density to stem base diameter $d_{0}$.


Fig. 48. Inter-tree species comparison of the relationships between the foliage proportion of the whole tree biomass to stem base diameter $d_{0}$.

We were also interested in the difference between bark density and density of stem wood (Fig. 47). If the density of both components is the same, the ratio between them is 1.0 . The ratio of almost all tree species decreased with the increasing tree size represented by $d_{0}$ diameter. It means that bark density of small trees was greater than the density of their stem wood. A surprisingly opposite tendency was observed for hornbeam, for which bark density increased with the tree size and was always greater than wood density. Here we have to note that in small trees the bark proportion of the stem is very high, and hence, the data on bark density are relevant.

Next, we focused on inter-species differences in biomass allocation, i.e. on the component proportion of the total tree biomass. We found that the proportion of the total tree biomass in foliage biomass (Fig. 48a, 48b) decreased with the increasing $d_{0}$ stem diameter or remained stable. The stable foliage proportion was revealed for oak and spruce. Pine had the highest foliage proportion ( $50 \%$ in the case of small individuals) followed by spruce. On the contrary, oak had the lowest foliage proportion (below 10\%). Hence, the results indicate that ever-green tree species (i.e. pine and spruce) have a higher foliage proportion than deciduous species. Here we
present the foliage proportion of the whole tree biomass at $d_{0}$ diameter equal to 50 mm for individual tree species. The values were as follows: oak $-5.7 \%$, hornbeam $-6.3 \%$, sycamore $-6.4 \%$, aspen $-6.4 \%$, ash $-8.2 \%$, beech $-10.1 \%$, rowan $-10.4 \%$, larch $-14.7 \%$, goat willow $-15.0 \%$, pine $-23.6 \%$, spruce $-30.0 \%$.


Fig. 49. Inter-tree species comparison of the relationships of branch biomass proportion of whole tree biomass to stem base diameter $d_{0}$.


Fig. 50. Inter-tree species comparison of the relationships between the biomass proportion of whole tree biomass in stem over bark to stem base diameter $d_{0}$.

Large inter-species differences existed also in the case of the branch proportion of the whole tree biomass (Fig. 49a, 49b). Here we observed an opposite tendency as in the case of foliage for the majority of tree species, i.e. the increase of branch proportion with the increasing $d_{0}$ stem diameter. High branch proportion (approximately $30-40 \%$ ) was found for larch, followed by goat willow. In contrast, sycamore had the lowest branch proportion (below 10\%). For example, at a diameter $d_{0}$ equal to 50 mm the following proportions of the total tree biomass in branches were estimated: sycamore $-6.6 \%$, ash $-10.9 \%$, oak $-11.5 \%$, hornbeam $-12.8 \%$, rowan $-15.8 \%$, aspen $-17.9 \%$, beech $-20.5 \%$, pine $-21.8 \%$, spruce $-24.4 \%$, goat willow $-24.9 \%$, larch $-36.5 \%$.

Stem over bark was the most important component of mass of all tree species in the greatest part of the diameter range (Fig. 50a, 50b). Nevertheless, also in this case we revealed significant inter-species differences. The proportion increased with the increasing $d_{0}$ stem diameter in the case of all tree species except for rowan (for which we observed a slight decrease). The highest proportion in the stem over bark was found for oak and hornbeam (exceeding 70\% in the case of larger individuals), while spruce and larch had the lowest proportions (around $30 \%$ ). If $d_{0}$ diameter was set to 50 mm , the following stem proportions of the total tree biomass were estimated: spruce $-30.8 \%$, larch $-35.5 \%$, goat willow $-42.6 \%$, pine $-44.0 \%$, rowan $-47.5 \%$, beech $48.4 \%$, aspen $-58.0 \%$, ash $-60.4 \%$, sycamore $-61.0 \%$, oak $-63.5 \%$, hornbeam $-68.5 \%$.


Fig. 51. Inter-tree species comparison of the relationships between the root proportion of the whole tree biomass to stem base diameter $d_{0}$.

Last, we examined the inter-species differences in the root proportion of the total tree biomass (Fig. 51a, 51b). For the majority of tree species, the root proportion decreased with the increasing $d_{0}$ stem diameter, but in the case of pine, larch, sycamore, rowan and goat willow, the proportion was almost stable. The greatest inter-species differences were found for smaller individuals (for diameters $d_{0}$ around 30 mm ), while for larger individuals the differences decreased. In the smallest individuals of oak and ash, the roots accounted for more than $50 \%$ of the whole tree biomass, while in the case of pine and larch it was only around $15 \%$. If we compared the modelled values of the root biomass proportion to the total tree biomass at $d_{0}$ diameter equal to 50 mm , the tree species were ranked as follows: pine $-10.5 \%$, hornbeam $-12.4 \%$, larch $-13.2 \%$, spruce $-14.9 \%$, goat willow $-17.6 \%$, aspen $-17.7 \%$, oak $-19.3 \%$, ash $-20.5 \%$, beech $-21.0 \%$, sycamore $-26.1 \%$, rowan $-26.3 \%$.

## 5. Knowledge synthesis and conclusion

The quantification of the whole tree biomass (or above-ground biomass and biomass of individual components) from easily measurable tree characteristics, i.e. $d_{0}$ diameter and tree height, is the most important outcome of our work. We found great inter-species differences. However, the results did not reveal substantial differences between the groups of coniferous and broadleaved tree species. We mentioned this because some authors tried to develop models of the tree biomass or the biomass of individual components using generalised relationships separately for coniferous and broadleaved tree species (e.g. Teobaldelli et al. 2009; Annighofer et al. 2016). For example, the generalised model of Annighofer et al. (2016) for coniferous tree species estimated the above-ground biomass of a tree with $d_{0}$ diameter to be equal to 1.8 kg . The model for broadleaved species estimated the above-ground biomass of the tree with the same dimension to be 2.3 kg . Our models estimated the amount of the above-ground biomass of a coniferous tree with diameter $d_{0}$ equal to 50 mm in the interval from 1.48 kg (pine) to 2.03 kg (larch). The biomass of spruce, which is the most common coniferous tree species in Slovakia, was 1.96 kg . In the case of broadleaved tree species, the biomass fluctuated from 1.35 kg (goat willow) up to 2.60 kg (hornbeam). The biomass of beech, i.e. our most common tree species in Slovakia, was 2.07 kg .

The results of this work revealed substantial inter-species differences in the biomass allocation of young individuals between the analysed tree species. The tree species had different proportions of the whole tree biomass in foliage, branches, stem over bark, and roots. We confirmed that ever-green tree species (i.e. pine and spruce) had a greater foliage proportion of the whole tree biomass than deciduous tree species (i.e. broadleaved species, and larch).

From all components, the greatest proportion of the tree biomass of the analysed tree species (except for the smallest individuals) was in the stem. The majority of the tree species had surprisingly high root and foliage proportions, which is different from older individuals (outside our interval of observations). This fact was confirmed by comparing our results with the knowledge of other authors (see e.g. Zhou et al. 2006; Kleinn 2007; Skovsgaard et al. et al. 2011; Krejza et al. 2017), who presented models for bigger trees than those included in our set. We also found that the proportions in individual components changed with the tree size, which results from uneven relative increments of individual tree components. For example, we can observe that the biomass proportion in stem, or in branches increases with the tree size at the expense of foliage and roots. This suggests that tree species have a specific growth strategy. More specifically, smaller trees prefer storing carbohydrates in physiologically most active organs. These organs are responsible for photosynthesis (thanks to the active surface of foliage), or for water and nutrient absorption (using root tips). Later, the differences between the relative increments of roots and foliage and the relative increments of stem and branches diminish. As trees develop further, the ratios change in the favour of above-ground "woody" organs. Subsequent preference of stem and branch growth is most probably linked to the need to ensure construction or support tree organs, or to occupy the space under competitive conditions (mainly a struggle for light) in a closed stand canopy.

The knowledge about biomass allocation, its inter-species differences, or its changes related to stand characteristics (e.g. density and vertical structure) and the development of an individual is interesting from several aspects. For example, it can tell us a lot about various growth strategies of individual tree species, i.e. the process of occupying soil by roots and the above-ground area by foliage. This question is particularly up-to-date from the point of management of mixed stands composed of the tree species that differ in their growth characteristics. Next, such knowledge clarifies the periods of carbon storage (or the opposite phenomenon, i.e. the rate of carbon flow) in the biomass of living trees. The duration of carbon storage obviously depends on the life span of a tree organ. While foliage has a life span from one growing season (deciduous tree species) up to a period of several years (ever-green tree species), stem and coarse roots and branches have a life span of several decades up to the maximum equal to a life time of a tree.

Here we have to note that allometric relationships should not be a goal, but only certain means or a tool to secure further scientific or practical intentions (see Čermák et al. 2015). We can demonstrate this with our previous works (see Chapter 7, i.e. List of papers of the authors relevant to the topics), where we used the allometric relationships to derive the component biomass for different scientific aims. They included the assessment of growth or production of tree species and stands (papers No. 5, 10, and 15), tree ontogenesis (papers No. 2 and 8), and physiological indicators of tree species (papers No. 2, 4, and 11), or development of biomass in old disintegrated or young post-disturbance forest stands ( $10,12,19$ ). Other papers dealt with carbon stock and sequestration (papers 12,19, and 20), as well as with the quantification of biomass consumed by red deer and the amount of forage potential of young stands (papers No. 7, 16, and 17).

Our original ambition with our allometric relationships for the estimation of tree biomass and its individual components was to derive models applicable in the western part of the Carpathians, or in the area of the Slovak Republic. The whole tree samples taken from the majority of the Slovak territory occupied by a particular species were thought to be the basis to meet this goal. For objective reasons (mainly problems to find young forest stands originating from natural regeneration and dominated by the particular species, i.e. its proportion had to exceed $90 \%$ in stand species composition), this could not be achieved in the case of goat willow, European larch, and Rowan. The samples of these three tree species were taken from the post-disturbance areas of the northern Slovakia (Vysoké Tatry, or Kysucké Beskydy). Hence, the models of these tree species may not be suitable for the rest of Slovakia, and should serve as framework information.

Another unsolved problem of a wider implementation of our models is the fact that they did not include the trees originating from artificial plantations, where lower numbers of individuals occur per unit area, nor they included the individuals developed under parent stands. In both cases, we can assume different light conditions as in our modelled case (i.e. young forest stands originating from natural regeneration developed without a parent stand), i.e. and probably also different allocation of biomass. The impact of a tree position inside a stand (availability of resources, mainly light) on biomass allocation remains an open question, particularly under the conditions of closed stands. For this purpose, the models for individual bio-sociological positions of trees should be developed. The models for some other tree species, mainly Silver fir and birch, are still missing. The birch samples were collected in the year 2017, but we have not managed to process the data and include it in this publication.

At the end, we need to point out at two most important methodological problems, which occurred while developing these models. One is the selection of the sample trees and the other logarithmisation of the data. When selecting the sample trees, we applied the principle of the equal numbers of sample trees representing individual bio-sociological positions, which were taken from the stands of different ages (usually 1 - 10 years). This sampling design caused that the trees representing subdominant and suppressed individuals had lower ranges of diameters and heights in our dataset. At the same time, it resulted in the considerably left-skewed distributions of diameters and heights. Regarding the logarithmic transformation of allometric equations, our opinion is that it is more suitable to use an allometric equation in its power form. Logarithmic transformation deforms original data, which can cause considerable differences between the models derived with and without the logarithmic transformation, particularly at greater values of an independent variable. We present such an example in Fig. 52 and 53, which shows the power model (b) and the linearised model after its reverse re-transformation (a). The differences in the predicted values of the spruce branch dry mass for the trees with diameter $d_{0}$ equal to 100 mm are substantial ( $3,000 \mathrm{~g}$ versus $2,000 \mathrm{~g}$ ).

Fig. 53 shows the fitting of the scatter plot after the linear transformation, from which 11 thickest trees marked red should be noted first. The same trees are marked red also in Fig. 52a. In both figures we can see that in nine cases the predicted values were greater than the measured ones, while in two cases the values were almost equal. While in Fig. 53 everything seems to be OK and the values of the residuals are not substantially greater, Fig. 52a shows that the values of the residuals after the reverse re-transformation of the linear model are great. This is caused by the fact that the same differences between the predicted and logarithmically transformed measured values, i.e. the same values of residuals, do not represent the same differences in the predicted and measured values of dry mass, but become greater with the increasing $d_{0}$ diameter (see Fig. 53). The deformation of the measured values can also be seen when comparing the widths of the diameter intervals comprising 11 thickest trees. In Fig. 52a the interval is from 60 to 100 mm , which is $40 \%$ of the whole range of diameters, while in Fig. 53 the interval is from 4.1 to 4.6 mm , which is $10 \%$ of the whole range of diameters. It means that if Fig. 53 shows that the linear function overestimates the values at ten per cent of the interval width, in reality it overestimates the values at forty per cent of the interval. The correctness of the logarithmically transformed model expressed by the variability of the predicted values around the observed values (Fabrika \& Pretzsch 2011) mainly depends on how the linear function fits the scatter plot at greater values of the independent variable. In Fig. 54 we can see the residuals of the linear and non-linear models. In the case of the residuals of the linear model, heteroscedasticity is eliminated, but there is a clear trend of overestimation at the smallest and greatest values, which is also visible in Fig. 53. The values of the non-linear model are heteroscedastic, but no signs of underestimation or overestimation in any parts of the interval are visible.


Fig. 52. Scatter plot of measured values of spruce branch dry mass fitted with logarithmic transformation of allometric equation [8] a) and with allometric equation in a non-linear power form [13] b).

We would like to emphasise that in the works of this character it is important to ensure the standardisation of the applied approaches, i.e. to choose one of the available approaches (either logarithmic or other transformation, or non-linear regression), and to apply this approach systematically to all data. In our work we used the method that is most frequently applied in the international scientific literature. This method was appropriate for the majority of the cases. In several (rather rare) cases, other approaches of fitting the scatter plot would be more suitable. However, in order to meet one of our main goals, i.e. to compare inter-species differences in biomass allocation, we did not combine the methods.


Fig. 53. Scatter plot of logarithmically transformed values of spruce branch dry mass fitted with linear function [3].


Fig. 54. Scatter plots of the residuals of the model derived using the logarithmic transformation of data fitted with the linear function a) and of the model derived using the non-linear function.

We intend to address the issue of developing mathematical models for the calculation of biomass of tree species (including the optimal selection of sample trees, and the most appropriate way of fitting the scatter plot) also in the future, and hence, we will fill in the gaps in the knowledge. In spite of some unsolved questions we believe that we have made considerable progress in this area. We are encouraged to continue in the work by a large number of our already published original scientific papers that dealt with this issue (see Chapter 7). We are also pleased with the intense citations of our papers, especially abroad. For example the paper of Pajtík et al. (2008) was cited more than 60 times (according to SCOPUS database).

We hope that the newly developed mathematical biomass models for eleven tree species, or the knowledge on biomass allocation to tree components, will serve as a basis for scientific purposes of many of our colleagues - research workers. The newly acquired knowledge could be applied not only in Slovakia, but also in other European countries, particularly in the regions of the Carpathian Mts., namely in the Czech Republic, Poland, Ukraine, and Romania.

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## 7. List of papers of the authors relevant to the topic

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