



Influence of storage on the physical and chemical properties of Scots pine bark

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Abstract

Bark is currently used mainly to produce energy, but the extraction of valuable compounds before combustion offers an interesting cascading use for debarking biomass. Buffer storage is an inevitable part of bark biomass logistics, but substantial dry matter and extractive losses can degrade the properties and reduce the economic value of the raw material during storage. In this study, moisture and ash content, calorific value, and extractives content and composition of Scots pine (*Pinus sylvestris*) sawmill bark were determined over 2 months of buffer storage, and the change in energy content during storage was calculated. The results showed that the energy content (MWh m⁻³) of the bark increased 3% during storage, while at the same time the moisture content decreased 16%. The content of acetone-soluble extractives decreased markedly, with only 56% of the original amount remaining after 8 weeks of storage. In particular, hydrophilic, phenolic extractive compounds were rapidly lost after debarking and piling of the bark. About 60% of condensed tannins (CT) and about 26% of the quantified lipophilic compounds were lost after 2 weeks of storage. The fastest rate of decrease and the most significant changes in extractives content and composition occurred within the first 2 weeks of storage. Utilization of these valuable compounds necessitates fast supply of material for further processing after debarking. The comprehensive utilization of bark requires efficiency at all levels of the supply chain to ensure that tree delivery times are kept short and loss of bark is avoided during harvest and transport.

Keywords Bark · *Pinus sylvestris* · Moisture · Energy content · Extractives · Debarking

Introduction

EU-wide environmental targets and policy objectives for the period from 2021 to 2030 are presented in the 2030 Climate and Energy Framework. One of the key targets is achieving at least a 32% renewable share of energy in the EU [1]. The share of renewable fuels in the Finnish national energy supply was set to be increased to 38% of total energy consumption by

2020 [2], and this goal was actually achieved in 2014 [3]. The success was achieved mainly by increasing the use of various types of biomass, especially forest chips and forest industry by-products, in energy generation. According to the latest statistics [4], the main solid wood fuel used in heat and power production was bark, with 7.7 million m³ (47.5 TJ) in 2018. Bark is a by-product from the sawmill and pulp industry. As the Finnish milling and pulp industry processes mainly coniferous wood, debarking residues from coniferous trees are the most common bark used as fuel.

The debarking of logs is usually the first processing stage in the mechanical and chemical forest industry. The aim of debarking is to avoid many harmful effects in subsequent process stages [5].

The pulp industry and sawmills use different debarking technologies. In the debarking drum, pulpwood logs are tumbled against each other to remove the bark, and the process is driven by a low amount of water or steam. Cutting season, temperature, and the quality of the raw material, i.e. species, log freshness, and log dimensions, affect the wood loss in the debarking drum. Koskinen [6] and Agin and Svensson [7] reported that the wood loss in the debarking drum varied

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between 1 and 6%. The end product, especially in chemical and mechanical pulping, mainly determines the requirements for bark removal, which is measured by the log cleanliness measurement system. Log cleanliness is defined as the percentage of the log surface that is free of bark after debarking [8]. The debarking result is a compromise between wood loss and bark removal.

Rotary debarking is the preferred method in the sawmill and veneer industries. The logs of the primary wood product industries are debarked, because processing residues (chips) are commonly used as a raw material in chemical and mechanical pulping.

The value chain of forest biomass for energy always includes biomass storage [9]. As supply volumes and commercial values have increased, the economic losses associated with the poor storage management of biomass have become obvious. The energy yields per unit of delivered biomass can be maximized, and emissions can be minimized, through the careful establishment and location of the storage, prediction and measurement of the changing moisture content, and the ability to match supply with demand [9]. Storage of bark has been overlooked in the scientific literature to date [10]. Research has focused on studying stem wood, wood chips, and forest residues.

The content of extractives in Scots pine (*Pinus sylvestris* L.) bark is generally between 16 and 25.9 as a percentage of dry matter (d.m.), whereas the extractives content in *P. sylvestris* stem wood is usually between 1 and 6.8% (d.m.) [11–14]. This makes pine bark an attractive raw material for biorefining. There are large volumes of potentially available Scots pine (*P. sylvestris*) bark, as it is one of the main softwood species in Fennoscandia [15]. Currently, pine bark is commonly used to produce heat at mills or power plants.

Scots pine bark has an attractive chemical composition for the production of high-value biochemicals. Miranda et al. [16] reported hydrophilic extractives content of 14.4% in pine bark (d.m.). This group of extractives includes valuable phenolic compounds. For example, Scots pine bark contains relatively high content of condensed tannins, between 3.2 and 8.5% (d.m.), which have applications in adhesives, rigid insulating foams, as flocculants in wastewater treatment, and in pharmaceuticals, food supplements, animal feeds, and cosmetics [17–21]. In addition, the content of lipophilic compounds in Scots pine bark is high, between 4.4 and 7.3% (d.m.) [16, 22], and these lipophilic compounds also have potential high-value applications as epoxy resins, surfactants, defoaming agents, emulsifiers, soaps, paper sizing agents, printing ink resins, cosmetics, and many others [23–25].

Major changes to woody biomass during storage are caused by three key mechanisms: (1) living cell respiration, (2) biological degradation, and (3) thermo-chemical oxidative reactions [10]. The microbial activity in storage piles, along

with other factors including biological and chemical processes and moisture levels within the stack, causes problems such as heat buildup, biomass loss, and degraded fuel quality. Dry matter loss (DML) of between 0.4 and 10.2% per month has been observed in stored bark [26–29]. The biological processes can lead to various chemical processes that occur at higher temperatures. In large biomass piles, self-heating can be a hazard, when the internal temperature of the pile increases over time and self-ignition occurs [30].

Many of the potentially valuable extractive compounds are either rather volatile or chemically unstable. The extractives content starts to decrease immediately after tree felling, and this degradation continues throughout the supply chain [31–34]. As a result, the chemical composition of the extractive-based fraction also changes over time.

The major reactions of lipophilic extractive compounds during wood storage can be divided into three types: (1) hydrolysis of triglycerides (rapid reaction) and steryl esters and waxes (proceeds slower); (2) oxidation/degradation/polymerization of resin acids, unsaturated fatty acids, and—to some extent—other unsaturated compounds; and (3) evaporation of volatile terpenoids, mainly monoterpenes [33]. Phenolic extractives are hydrophilic by nature. Thus, in addition to microbial and chemical degradation reactions, they are also lost due to leaching. Phenolic compounds may also experience photodegradation during storage, through the formation of phenoxy radicals facilitated by UV light exposure and enzymatic activity [35, 36]. Phenoxy radicals are highly unstable and may lead to polymerization or cyclization reactions [37]. High temperatures in piles can also lead to the thermal decomposition of some of the chemical components of bark [30]. Some compounds are released in the form of gaseous organic pollutants (volatile organic compounds, VOC), and some compounds are released via leachate.

Increasing the value of bark for bioenergy and as a raw material for biorefineries is the key issue in this study. The aim of the study was to determine the changes in quality of Scots pine sawmill bark during storage. In addition, the amount of extractives was measured after 2, 4, and 8 weeks of storage to monitor the changes in the chemical composition of the bark.

Materials and methods

Experimental design

The storage experiment was conducted in Uimaharju (62° 54' N, 30° 14' E), Eastern Finland, at a sawmill storage site and was carried out from August 22, 2018, to October 18, 2018. One pile of fresh pine (*Pinus sylvestris*) sawlog bark was established at the Uimaharju sawmill. The accumulated

rainfall for the 2-month follow-up period was 158 mm, and the mean air temperature was 10.1 °C.

Sawlogs were harvested by a harvester around 2 weeks before storage. Fresh wood was debarked with a Valon Kone VK-820 rotary debarking machine on the same day the pile was established. The bark was piled with a wheel loader. The size of the pile was 16.4 x 6.2 x 3 m (length, width, height). The pine bark contained a large amount of stem wood; the average share of wood pieces in samples in this study was 20%.

Sampling was carried out with balance bags. The samples were individually mixed to homogeneity and divided into two subsamples of about equal weight (1000 g). The first subsample was packed in a plastic bag and set aside for measurement of pre-storage wet-basis moisture content and basic density. The other subsample was packed in a polyester bag (40 x 50 cm) with a 1-mm oval mesh. The bark for the balance bags was thoroughly homogenized to ensure the same starting conditions for all samples within one pile and to achieve minimal variance in the initial data. In constructing the experiment, the balance bags were arranged grid-wise within the piles. A total of 30 balance bags were placed in the pile at three different levels. The first level (bottom), with 11 bags, was at a height of 0.5 m from the ground, the second (middle) was 1.3 m (11 bags), and the third (top) was 2 m (8 bags).

Samples for chemical analyses were collected from different truckloads, and combined and mixed. A sample of approximately 5 L was placed in the sampling bag. Three larger samples were also collected, which were further divided into subsamples at the laboratory.

During the storage period, the bags were extracted after 2, 4, and 8 weeks, using a drag rope and wheel loader/excavator. After 2 weeks, eight bags were removed from the pile, all from the same cross-section, and after 4 weeks, eight bags were removed from another cross-section. At the end of the experiment, the other 14 bags were removed. Samples for determining the extractives amount were taken at the same time and from the same places as the balance bags. The sample for extraction was packed in a 5-L plastic bag.

The in-pile temperature was recorded at 1-h time intervals using miniature temperature data loggers (a-Nap, a-Lab Ltd, Keuruu, Finland) and 3-h time intervals using thermo cables. Five a-Nap data loggers and five thermo cables were placed on the pile at different levels. After a 4-week storage period, one thermo cable broke when the samples were taken.

Sampling methods

The sampling procedure and sampling preparation followed the standard methods valid at the time of the experiments. Moisture content was analyzed according to the Finnish Standards Association SFS-EN 14774-2 standard [38]. Ash content was determined according to SFS-EN 14775 [39]. A

method modified from the Scandinavian Pulp, Paper and Board Testing Committee standard SCAN-CM 43:95 [40] was applied to determine the basic density of the bark samples. Calorimetric heating measurements and calculation of the gross calorific values were performed according to the European Committee for Standardization CEN/TS 14918:2005 methods [41], using an IKA® C 5000 bomb calorimeter (IKA®-Werke GmbH & Co. KG, Germany).

Calculation of energy content and dry matter losses

The energy content of the study piles was calculated at the beginning and end of the experiment, based on the measured moisture content. The net calorific value and energy content of the as-received bark were calculated according to the following equations [42]:

$$q_{p,net,ar} = q_{p,net,d} * \left(\frac{100 - M_{ar}}{100} \right) - 0.02443 * M_{ar}, \tag{1}$$

where

- $q_{p,net,ar}$ net calorific value as received (MJ kg⁻¹),
- $q_{p,net,d}$ net calorific value on a dry basis (MJ kg⁻¹),
- M_{ar} moisture content as received,
- 0.02443 the correction factor of the enthalpy of vaporization at 25°C, and

$$E_{ar} = \frac{1}{3600} * q_{p,net,ar} * BD_{ar} \tag{2}$$

where

- E_{ar} energy content as received (MWh m⁻³),
- $q_{p,net,ar}$ net calorific value as received (MJ kg⁻¹), and
- BD_{ar} density as received (kg m⁻³).

The moisture content, basic density, and net calorific values of dry matter measured in the samples from the balance bags at the beginning and end of the experiment were used in the calculation.

Extractives analysis

Samples were stored in a freezer (< -20 °C). Prior extraction samples were freeze-dried and ground with a Retsch SM 100 laboratory cutting mill (Retsch GmbH, Haan, Germany), equipped with a bottom sieve with trapezoidal holes (perforation size < 1.0 mm). The moisture content of the ground samples was determined according to the SFS-EN 14774-3 standard method by drying 1 g of bark powder at 105 °C in an oven. The extraction of bark samples was done with a Foss Soxtec™ 8000 unit. The extractions were performed in duplicate. Two grams of bark powder was used for each extraction.

Samples were extracted with 80 ml acetone in boiling solvent for 15 min, after which the thimble was raised to the rinse position for 60 min. The acetone extracts were first concentrated by extractor, and drying was finalized before weighing using a nitrogen stream. These duplicate samples were further analyzed with gas chromatography (GC).

The methods used for the analysis of extractive compounds were modified from the original methods proposed by Örså and Holmbom [43]. To analyze individual extractive compounds, about 13 mg of bark extract and 200 µg of internal standards (heneicosanoic acid and betulin) were derivatized with a mixture of pyridine and 1-(trimethylsilyl)imidazole after drying with nitrogen stream. The derivatization of samples was performed by keeping them at room temperature overnight. GC analysis of the individual compounds was carried out on an Agilent 6890 Series GC system equipped with a 7683 injector and an Agilent 5973 mass selective detector (MSD). The capillary column used was the Zebron ZB-SemiVolatiles (30 m × 0.25 mm, film thickness 0.25 µm). The injector temperature was 280 °C, with 1 µl injection volume and split injection mode (split ratio 21.6:1). Helium was used as the carrier gas. The initial GC oven temperature was 180 °C, followed by an increase of 5 °C/min to 320 °C, and 12 min at 320 °C.

A quantitative analysis of triglycerides and steryl esters was performed with the same GC system as the analysis of individual compounds. About 3.5 mg of extract and 200 µg of internal standards cholesteryl heptadecanoate, heneicosanoic acid, and 1.3-dipalmitoyl-2-oleoylglycerol and 75 µg betulin were added to the sample, which was analyzed without derivatization. A short DB-1ht capillary column (15 m × 0.25 mm, film thickness 0.1 µm) was used. The injector temperature was 280 °C, with injection volume of 3 µl and pulsed splitless injection mode. The initial GC oven temperature was 150 °C, followed by an increase of 20 °C/min to 350 °C, and 10 min at 350 °C.

Condensed tannin (CT, proanthocyanidins) content was determined by high-performance liquid chromatography (HPLC) after thiolytic degradation, following Mattila et al. [44]. Briefly, the samples were weighed (20–30 mg) in 1.5 mL Eppendorf vials, and 1 mL of depolymerization reagent (3 g cysteamine/4 mL 13M HCl/56 mL methanol) was added. The vials were sealed and incubated for 60 min at 65 °C, after which the degradation products, i.e. free flavan-3-ols (terminal units) and their cysteaminyll derivatives (extension units), were separated on a Zorbax Eclipse Plus C18 column (2.1 × 50 mm, 1.8 µm) and determined by HPLC (Agilent 1290 Infinity, Agilent Technologies, Inc., Santa Clara, CA, USA), equipped with diode array (DAD) and fluorescence detection (FLD).

Chemicals

HPLC-grade (≥ 99.8%) acetone (VWR Chemicals BDH®) was used for extraction. The compounds used as internal

standards in the GC analysis of the extractives were heneicosanoic acid (99%), betulin (98%), cholesteryl heptadecanoate, and 1.3-dipalmitoyl-2-oleoylglycerol (>99%), all purchased from Sigma-Aldrich (St. Louis, MO, USA).

The solvents used in the sample preparation of extractives were analytical-grade pyridine (99.5%, Merck KGaA, Darmstadt, Germany) and 1-(trimethylsilyl)imidazole (96%, Sigma-Aldrich, St. Louis, MO, USA).

Standard compounds applied for CT analyses were (+)-catechin (99%, Sigma-Aldrich, St. Louis, MO USA), (–)-epicatechin (98%, Sigma-Aldrich), and procyanidin B2 (>90%, Extrasynthese, Lyon, France). Cysteamine (98%) and formic acid (98%) were obtained from Sigma-Aldrich (St. Louis, MO, USA), and 13M HCl from Fisher Scientific (Waltham, MA, USA). HPLC-grade methanol (≥ 99.8%) and acetonitrile (≥ 99.8%) were obtained from VWR Chemicals BDH®.

Statistical analyses

Statistical analyses were performed using IBM SPSS Statistics version 25 software (IBM Corp., Armonk, NY, USA). The significance level was 5% ($p \leq 0.05$) for all tests. Analysis of variance (ANOVA) was performed for moisture, ash, extractives, and condensed tannin (CT) content, basic density, and energy content data to identify the differences in the pile. A post hoc test, i.e. a Bonferroni-corrected *t* test, was conducted for extractives content measurements and the results of the compound group analysis (steryl esters, triglycerides, phenolic extractives, sterols). For free fatty acid, resin acid, and CT content, ANOVA with a Welch test was carried out due to the violation of homogeneity of variance (significant Levene's test). A Games–Howell post hoc test was conducted for these results.

Variation in moisture content was visualized by interpolating a raster surface from sample points, using an inverse distance weighting method.

Results

Temperature

After the pile was established, the temperature in the middle of the pine bark stockpile rose rapidly, reaching 60 °C within 3 days, but then declined rapidly (Fig. 1). After 2 months, the temperature in the pile had decreased to ca. 25 °C. The temperature was highest in the middle and at the top of the pile for the first 3 weeks in undisturbed parts of the bark pile. At the bottom level, the temperature increased less and declined more slowly. The temperature at the top and in the middle of the pile decreased markedly after 3 weeks, with the temperature fluctuating between 8 and 35 °C at the top of the pile (Fig.

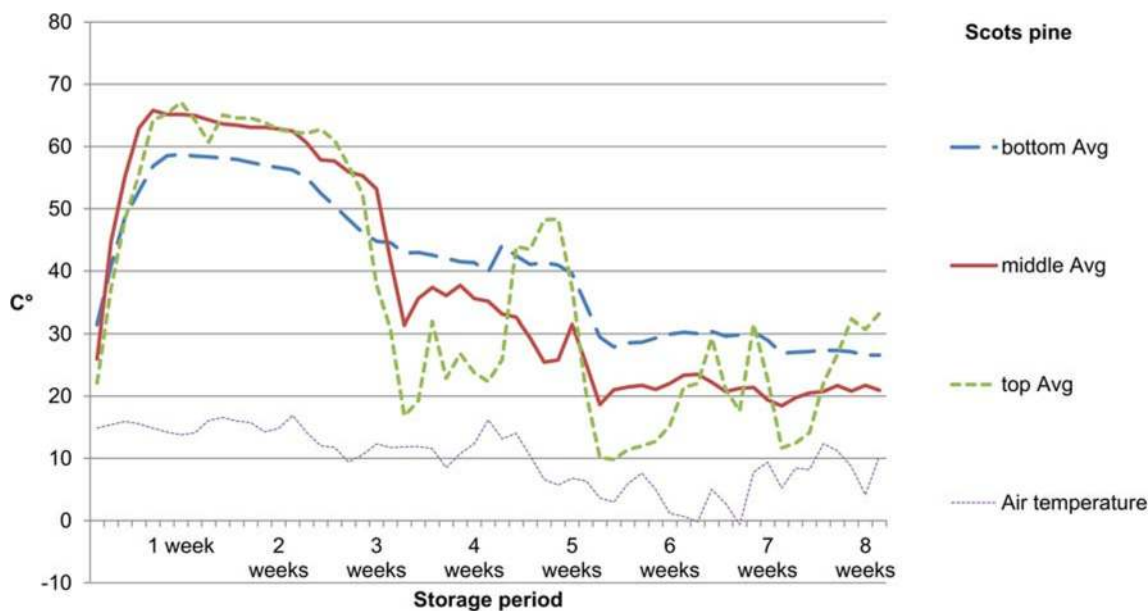


Fig. 1 Temperature development in the pine bark pile at different levels (bottom, middle, and top, undisturbed parts) during the 2-month storage period

1). The disturbance of the pile during sample collection at 2 and 4 weeks caused a brief decrease in the temperature.

Moisture content

The average initial moisture content of the bark pile was $47.5\% \pm 2.05$ (range 43.6–51.7%), and at the end of experiment it had decreased by 16% to $31.4\% \pm 3.92$ (range 25.5–37.7%). The lowest moisture content at the end of the experiment was found in the middle of the pile (Fig. 3), but there were no statistically significant differences ($p = 0.168$) between the bottom, middle, and top parts of the pile. The moisture in the middle of the pile did not differ statistically significantly ($p = 0.575$) from the moisture at the edges of the pile (Figs. 2 and 3).

Ash content

The average ash content of the fresh pine bark was 1.89% (± 0.06) (Table 1). After 8 weeks of storage, the average ash content was 1.97% (± 0.05). The difference was statistically significant ($p = 0.00$). The ash content at the end of the experiment showed no statistically significant differences ($p = 0.612$) at different levels of the pile (bottom, middle, and top).

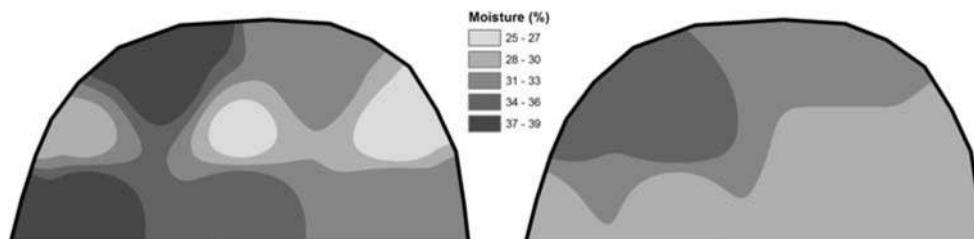
Basic density

At the beginning of the experiment, the basic density of the pine bark was $316 \pm 6.2\text{kg m}^3$ (varying between 301 and 329 kg m^3) (Fig. 4). During the 8 weeks, the storage density declined to $309 \pm 5.8\text{kg m}^3$ (-2.3%). The change was statistically significant ($p = 0.001$). The density at the end of the experiment showed no statistically significant differences ($p = 0.485$) at the different levels of the pile (bottom, middle, and top).

Energy content and calorific value

The average net calorific value ($q_{p,net,d}$), of the pine bark was $19.7 (\pm 0.13)\text{ MJ kg}^{-1}$, and after 8 weeks of storage it was the same, $19.7 (\pm 0.11)\text{ MJ kg}^{-1}$ (Table 1). Correspondingly, the average gross calorific dry basis value (MJ kg^{-1}) (q_{gr}) was $20.99 (\pm 0.13)\text{ MJ kg}^{-1}$, and after 8 weeks of storage it was $21.00 (\pm 0.11)\text{ MJ kg}^{-1}$. The energy content of the fresh pine bark was $1.54 (\pm 0.03)\text{ MWh m}^{-3}$, and during storage it increased slowly to $1.59 (\pm 0.027)\text{ MWh m}^{-3}$ (Table 1). The increase in the energy content was statistically significant ($p = 0.00$).

Fig. 3 Moisture content of the pine bark in various parts of the stockpile at the end of the experiment



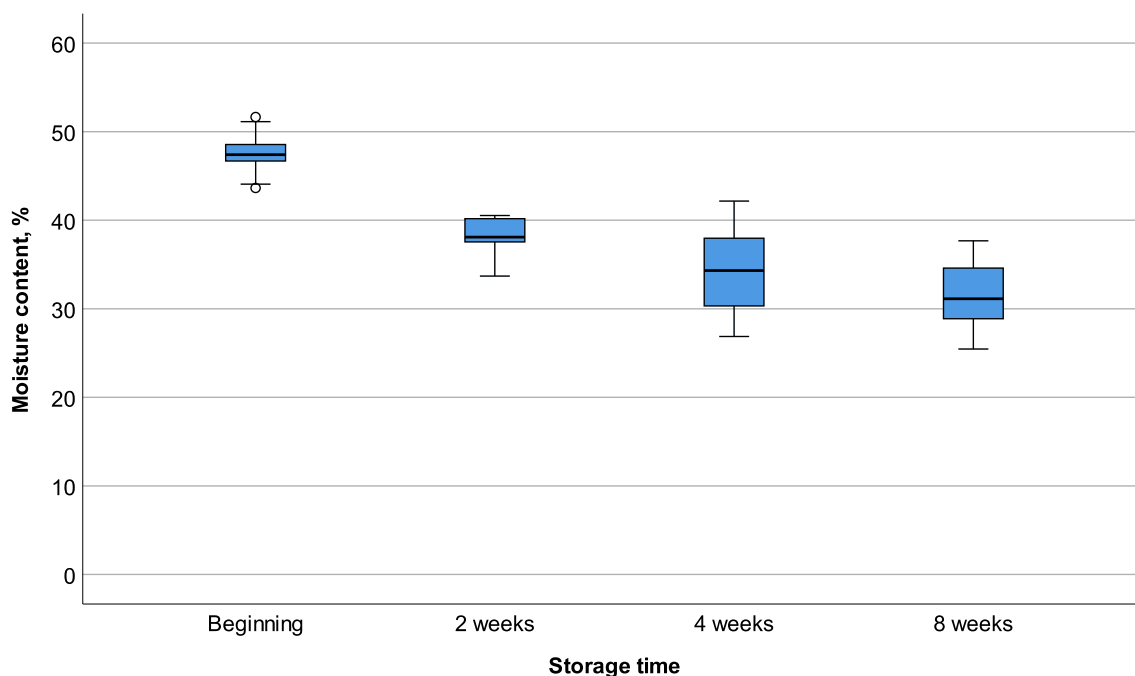


Fig. 2 Average moisture content of bark during 2, 4, and 8 weeks of storage

Extractives

The average extractives content of the fresh pine bark was 9.92% (± 0.43). At the beginning of the storage period, the extractives content decreased rapidly (Fig. 5 and Table 1). The duration of storage significantly affected the extractives content of pine bark ($p = 0.000$). ANOVA for the samples was executed only after storage (2, 4, and 8 weeks). This was due to the small number of samples ($n = 3$) analyzed from fresh bark. A Bonferroni-corrected t test indicated no significant difference in the extract content of the samples between 2 and 4 weeks of storage ($p = 0.993$). However, there was a significant difference in the extractives content of the samples between 4 and 8 weeks of storage ($p = 0.000$). The largest decrease in extractives content occurred

during the first 2 weeks, when approximately 30% of the initial extractives content was lost.

The extractives content at the end of the experiment showed no statistically significant differences ($p = 0.421$ – 1.000) at the different levels of the pile (bottom, middle, and top). ANOVA could not be conducted to compare the extractives content in the middle of the pile with that at the sides of the pile due to the small and uneven number of observations between groups. Instead, the mean values of the extractives content were compared, and the following observations were made: After 4 weeks of storage, the average extractives content (% of d.m.) was 7.61% in the middle of the pile and 7.11% (± 0.54) at the sides of the pile. After 8 weeks of storage, the extractives content was 5.92% and 5.63% (± 0.27) in the middle and at the sides of the pile, respectively.

Table 1 The mean moisture content, basic density, calorific value, energy content, ash content, and extractives content of the pine bark before (initial) and after storage for 2–8 weeks (post-storage)

Time in storage	Moisture content, average, %	Basic density, average, kg m^{-3}	Net calorific value, average, MJ kg^{-1}	Net calorific value as received, average, MJ kg^{-1}	Energy content, MWh m^{-3}	Ash content, average, %	Extractives content, average, %
Fresh bark	47.48 \pm 2.04	316 \pm 6.15	19.70 \pm 0.13	9.19 \pm 0.46	1.54 \pm 0.03	1.90 \pm 0.06	9.92 \pm 0.43
2 weeks in storage	38.24 \pm 2.22	310 \pm 7.80	19.46 \pm 0.07	11.08 \pm 0.48	1.54 \pm 0.03	1.97 \pm 0.05	6.87 \pm 0.37
4 weeks in storage	34.28 \pm 5.35	308 \pm 8.01	19.48 \pm 0.21	11.97 \pm 1.16	1.56 \pm 0.04	2.16 \pm 0.12	7.12 \pm 0.70
8 weeks in storage	31.45 \pm 3.92	309 \pm 5.81	19.71 \pm 0.11	12.74 \pm 0.88	1.59 \pm 0.03	1.97 \pm 0.05	5.55 \pm 0.43

Results are presented with standard deviations

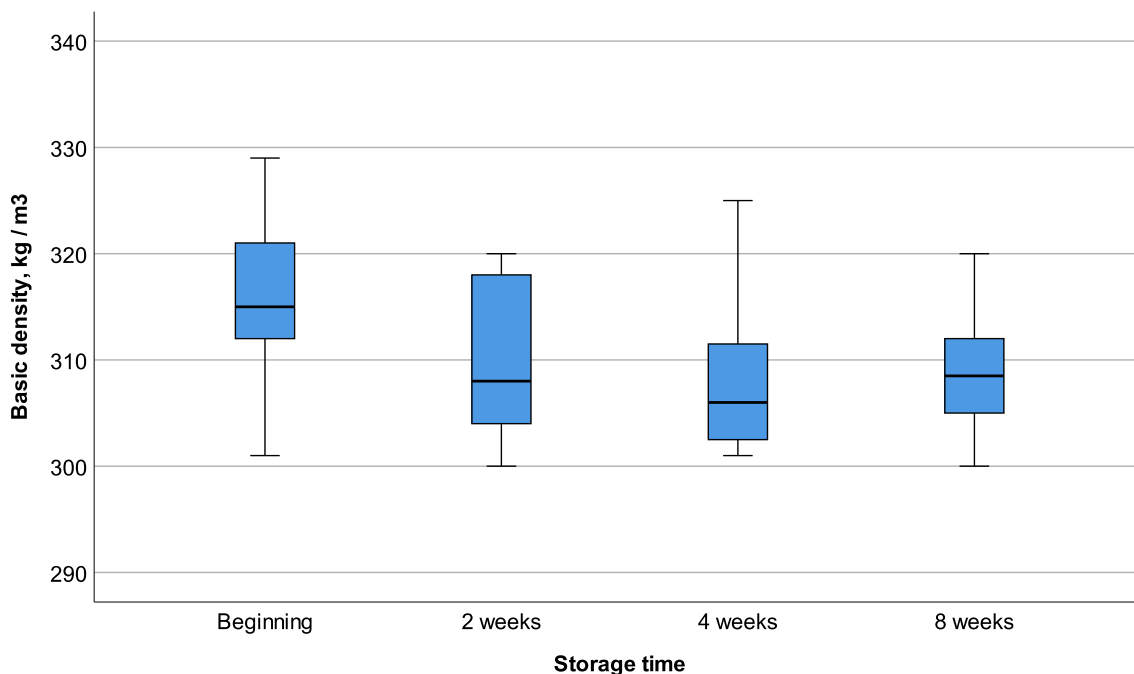


Fig. 4 Basic density change during 2, 4, and 8 weeks of storage

The standard deviation was relatively large, and the values were quite close to each other, so no differences in extractives content regarding its location in the pile were observed.

The qualitative GC/MSD analysis results for acetone extractives are presented in Table 2. The results of quantification are presented in Table 3 and Fig. 6 for different extractive groups.

The content of triglycerides (TG) decreased rapidly at the beginning of the storage period, and was approximately halved during the first 2 weeks of storage (Table 3 and Fig. 6). There was no significant difference in TG content between 2 and 4 weeks of storage ($p = 0.058$), but a significant difference was found between 4 and 8 weeks ($p = 0.005$). Most of the phenolic compounds quantified with GC/MSD (Table 2) were lost during

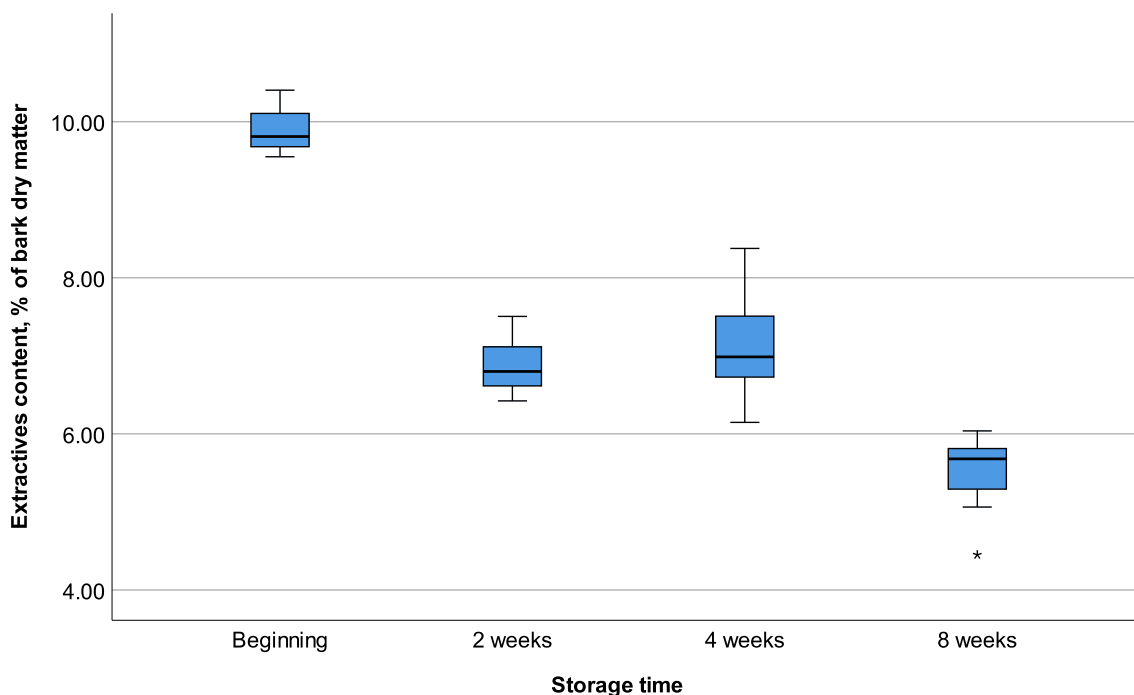


Fig. 5 Changes in gravimetric extractives content during 2, 4, and 8 weeks of storage compared to fresh spruce bark. There was one outlier, a sample taken from the top layer (marked with *) of the pile.

the first 2 weeks of storage, with only a small amount of lignans, dihydroconiferyl alcohol, and taxifolin remaining after the 2-week storage period. No other phenolic extractive compounds could be reliably detected and quantified after this period. The loss of dihydroconiferin was especially noteworthy. Fresh bark

Table 2 Qualitative gas chromatographic analysis results for extractives. Identified compounds are listed in order based on their retention times

Peak no.	Retention time, min	Compound
1	5.1099	Pinitol
2	5.194	Dihydroconiferyl alcohol ^b
3	7.776	Palmitic acid (16:0)
4	8.728	Margaric acid (17:0)
5	9.015	Thunbergol
6	9.603	Linolenic acid (18:3)
7	10.094	Linoleic acid (18:2)
8	10.158	Oleic acid (18:1)
9	10.524	Stearic acid (18:0)
10	11.597	Monomethyl pinosylvin ^a
11	11.735	Pimaric acid
12	11.9683	Sandaracopimaric acid
13	12.1459	Pinosylvin ^a
14	12.225	Isopimaric acid
15	12.42	Palustric acid
16	12.542	Unidentified resin acid
17	12.613	Eicosatrienoic acid (20:3)
18	12.725	Levopimaric acid
19	12.912	Dehydroabietic acid
20	13.391	Abietic acid
21	14.281	Hydroxydehydroabietic acid
22	14.345	Abietatetraenoic acid
23	14.654	Neoabietic acid
24	14.786	ISTD Heneicosanoic acid (21:0)
25	14.8685	Docosanol
26	14.957	Hydroxydehydroabietic acid
27	15.209	Isodehydroabietic acid
28	15.437	Unidentified hydroxy resin acid
29	15.564	Hydroxydehydroabietic acid
30	15.953	7-Oxo-dehydroabietic acid
31	16.169	Behenic acid (22:0)
32	16.381	Unidentified hydroxy resin acid
33	17.269	Dihydroxydehydroabietic acid
34	17.541	Tricosanoic acid (23:0)
35	17.5876	Tetracosanol
36	17.701	1-Mono-oleoylglycerol
37	18.324	Hydroxy-7-oxodehydroabietic acid
38	18.844	Lignoceric acid (24:0)
39	19.407	Catechin ^a
40	19.788	Dihydroconiferin ^a
41	20.514	Taxifolin ^b
42	20.581	22-Hydroxydocosanoic acid
43	20.925	Secoisolariciresinol
44	21.377	Cerotic acid (26:0)
45	22.935	Behenoyl-glycerol
46	23.287	Matairesinol
47	24.199	Campesterol
48	25.206	Sitosterol
49	26.846	Stigmast-4-en-3-one
50	27.581	ISTD Betulin
51	28.509	Taxifolin glycoside ^a

^a Phenolic extractives lost during the two 2 weeks of storage

^b Phenolic extractives lost between 4 and 8 weeks of storage

contained 2.25 ± 0.09 mg/g dihydroconiferin, which was not detected after 2 weeks of storage. The differences between the content after 2, 4, and 8 weeks were significant ($p = 0.000$), whereas no significant differences in the content of sterols ($p = 1.000$) were observed between 2 and 4 weeks of storage, although differences in content were observed between 4 and 8 weeks ($p = 0.000$). In the case of steryl esters, there was no significant difference in content between 2, 4, or 8 weeks of storage ($p = 1.000$). The content of free fatty acids roughly doubled during the first 2 weeks of storage. The Games–Howell post hoc test indicated that there was a significant difference in the fatty acid content between 2 and 8 weeks of storage ($p = 0.045$). The content of resin acids decreased during the first 2 weeks of storage, and then remained relatively stable, with no significant differences observed between the 2-, 4-, and 8-week storage periods ($p = 0.074–0.745$).

At the beginning of the study, the CT content in the bark was 3.51 g/100 g (± 0.04) (Fig. 7). Thiolytic degradation released subunits (flavan-3-ols) with structures of catechin and epicatechin, indicating that CT in pine bark essentially consisted of procyanidins. CT content was 1.28 g/100 g (± 0.01), 1.18 g/100 g (± 0.01), and 0.91 g/100 g (± 0.15) after storage for 2, 4, and 8 weeks, and the differences in content were significant ($p = 0.001–0.012$). Thus, after 2 weeks of storage, more than 60% of the initial content was lost, after which CT content remained relatively steady, although it still declined slightly. The average degree of polymerization in CT was highest at the beginning of the study and then declined, especially during the first 2 weeks. Generally, no significant differences could be observed in samples taken from different levels of the pile, except after 8 weeks of storage, when CT content at the top (0.735 ± 0.108 g/100 g) of the pile was significantly lower than in the middle (0.976 ± 0.252 g/100 g) or at the bottom (1.020 ± 0.345 g/100 g).

Discussion

The aim of this study was to determine the changes in quality characteristics and extractive content during the storage of pine bark in order to evaluate the effect of storage on bark for bioenergy use and the suitability of bark for biorefining.

The temperature development in the pine bark pile was rapid, reaching 60 °C within 3 days, but declining after 3 weeks to around 30 °C. In coniferous bark piles, the temperature reached $60–65$ °C within a few days, and then stabilized at 60 °C for several weeks, similar to the results in studies by Lehtikangas and Jirjis [45], Krigstin et al. [46], and Routa et al. [29]. A complicating factor is that the sampling of bark during storage causes a disturbance in the piles and can lead to biased results. The moisture content of the pine bark pile at the beginning of the experiment was quite low, and significant drying occurred (by 16%) during the 2-month storage period. The net bag analyses showed that the driest areas at the end of

Table 3 Changes in the content (mg/g dry bark) of different extractive compound groups during storage

Time in storage	Fatty acids	Resin acids	Sterols	Phenolic extractives	Other lipophil.	Steryl esters	Triglycerides
Fresh bark	1.81 ± 0.12	9.93 ± 0.59	1.23 ± 0.04	3.88 ± 0.15	0.40 ± 0.01	2.50 ± 0.05	12.37 ± 0.34
2 weeks	3.58 ± 0.07	7.47 ± 0.27	1.27 ± 0.09	0.30 ± 0.03	0.34 ± 0.01	2.22 ± 0.21	6.13 ± 0.66
4 weeks	3.58 ± 0.21	7.59 ± 0.33	1.24 ± 0.12	0.13 ± 0.02	0.35 ± 0.02	2.21 ± 0.18	5.12 ± 0.40
8 weeks	3.22 ± 0.49	7.05 ± 0.73	1.02 ± 0.10	0.04 ± 0.01	0.35 ± 0.04	2.19 ± 0.38	3.85 ± 1.02

Standard deviations for the results are given in the table. Compound groups fatty acids, resin acids, sterols, phenolic extractives, and other lipophilic compounds were quantified with a long capillary column (ZB-SemiVolatiles), and steryl esters and triglycerides with a short capillary column (DB-1HT). The phenolic compound group in this table includes compounds with peak numbers 2, 10, 13, 39–41, 43, 46, and 51 (Table 2).

the experiment were in the middle of the pile, but the values only refer to their specific position in the pile.

The average net calorific value ($q_{p,net,d}$) of the bark did not change during the experiment, as also reported by Lehtikangas and Jirjis [45]. However, the net calorific value as received ($q_{p,net,ar}$) increased markedly, since the bark moisture content decreased during storage. The energy content ($MWh\ m^{-3}$) of the pile increased during the 8-week storage, and at the same time the moisture content decreased by 16%. The ash content of the pine bark was 1.89% at the beginning of the experiment and 1.97% at the end, similar to the levels reported by Lehtikangas and Jirjis [45] and Saarela et al. [47]. During the 8-week storage, the basic density of the pine bark decreased by 2.3%.

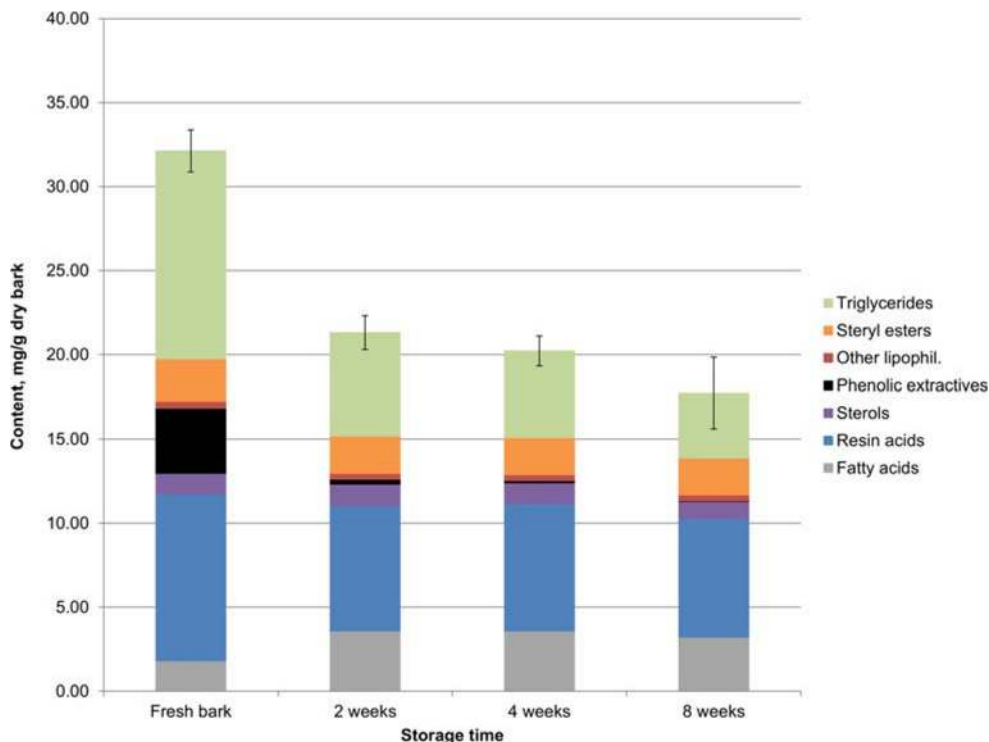
According to Koskinen [6] and Agin and Svensson [7], wood loss of 1–6% is normal with drum-debarked pulp wood. In this study, rotary debarking was used for sawlogs, and a

relatively high amount of wood (19.6%) was observed in saw bark streams. In this experimental pile, such a high percentage of wood may have affected the heat development in the pile compared with other study results.

In favorable weather conditions, the bark pile can dry markedly during storage. The bark was stored from August to October, which is a relatively short storage period for an industry in which production runs during all seasons. However, sawmills are usually shut down in July for maintenance. After the shut-down, the bark starts to accumulate, and the need for energy is low, whereas large amounts of bark need to be stored. Thus, the most problematic time is in the autumn season, and therefore the storage test was timed specifically for that time period.

During the storage period, the ambient temperature and total precipitation were slightly higher than the long-term average (1981–2010) obtained from the Finnish Meteorological

Fig. 6 Changes in the content of different extractive groups during storage. Results are based on GC/MS analysis of acetone extracts. Standard deviation for the quantification of all the compounds is shown in the figure



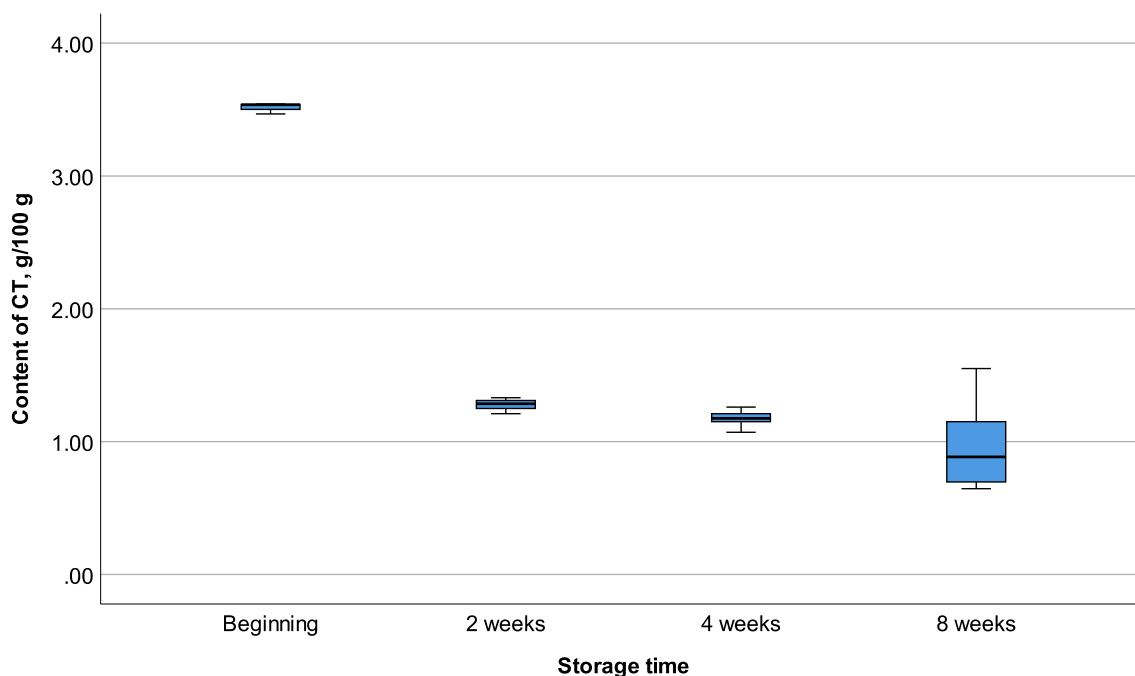


Fig. 7 Decreases in content of condensed tannins during 2, 4, and 8 weeks of storage compared to fresh spruce bark

Institute [48]. The experiment ended in the middle of October, and so the pile was not exposed to heavy rain or snow. If the pile had been stored over the winter, it would have had to have been covered. Covering the biomass to protect it from rain and snow can both increase the drying performance and reduce the risk of dry matter loss [28, 49]. However, the cover material applied must allow ventilation of water vapor and heat.

The most significant changes in extractives content and composition occurred within the first 2 weeks of storage. A similar rate of decrease was reported in Norway spruce bark extractives content during the first 2 weeks of storage, although the extractives content of spruce bark was higher ($11.83 \pm 0.13\%$ d.m.) at the beginning [29]. However, a greater share of the extractives were lost from Scots pine bark during the 8-week storage. The content of pine bark extractives decreased from 9.92 ± 0.43 to 5.55 ± 0.43 (% d.m.) during the 8 weeks, whereas the spruce bark extractives content after 8 weeks of storage was still 7.83 ± 0.29 (% d.m.). It should be noted that these storage studies were conducted simultaneously, and the environmental conditions were therefore the same. The experimental setups were also similar, there were no significant differences in pile size, and storage locations were in close proximity to one another. Thus, the observed differences in the extractives losses may be a result of the differences in the properties of the bark materials, i.e. in their extractive composition or chemical composition in general, and the structural properties of the bark. The structure and chemical composition of woody material significantly influence its degradation by microorganisms [50]. In the case of softwood bark (mixture of Norway spruce and Scots pine pulp mill barks), the most significant changes in extractives content were likewise observed at the beginning of the storage period, i.e. after the first

week of storage [51]. The amount of extractives in the bark material was almost halved during the first 4 weeks of storage [14, 51].

Neither the Norway spruce bark [29] nor the Scots pine bark storage in this experiment showed statistically significant differences in extractives content at the different levels of the pile (bottom, middle, and top). This result differs from the results for Norway spruce bark storage by Halmemies et al. [52], who observed that the different sampling locations in the pile had a clear effect on extractives content. They found that the rate of decrease in extractives content was fastest at the top of the pile, slowest in the middle, and somewhere in between at the sides of the pile. These differences between the studies may be due to the different tree species, pile dimensions, sampling locations, and storage seasons. The differences may also be partly explained by the differences in extraction methods and the solvents used. There may also have been differences in the properties of the stored bark (e.g. age and origin of harvested stems, particle size of bark, and density of the bark pile).

The composition of the extractives fraction changed during the 8-week storage period. The content of triglycerides decreased rapidly at the beginning of the storage period, and was approximately halved during the first 2 weeks of storage. As a result of the hydrolysis of triglycerides and possibly other compounds (e.g. steryl esters), the amount of free fatty acids increased at the beginning of the storage period. The increase in the amount of 18:1, 18:2, and 22:0 fatty acids was most remarkable. Most of the phenolic compounds quantified with GC/MS were lost during the first 2 weeks of storage. Only a small amount of lignans, dihydroconiferyl alcohol, and taxifolin remained after the 2-week storage period. After 8 weeks of

storage, the lignans were the only phenolic compounds that could be detected from acetone extract with GC/MS. The content of resin acids decreased during the first 2 weeks of storage, after which it remained relatively stable. The content of levopimaric, isopimaric, and neoabietic acids decreased most significantly. A decrease in the concentration of abietic acid was also noted. Lappi et al. [51] also observed a decrease in TG content and the content of interesting phenolic compound groups such as lignans during softwood pulp mill bark storage.

CTs in Scots pine bark essentially comprised procyanidins, in line with previous studies [53, 54]. However, both Bianchi et al. [53] and Matthews et al. [54] determined somewhat higher CT content in pine bark than that in the present study. Naturally, some variation is expected, but the relatively low CT content in the present study may be explained in part by the relatively high proportion of non-bark material, i.e. wood, included in the samples. Wood is not expected to contain CT.

CT content in the pine bark decreased during storage, especially in the first 2 weeks. Similar results were found in a study by Mupondi et al. [55], in which storage of pine bark for 90 days led to significant CT loss. While CT is known to be a relatively resistant substance and not easily decomposed in nature, it is also known that CT-rich material such as tree bark is often occupied by CT-utilizing bacteria and fungi [56–58]. Rapid self-heating of the bark piles indicates high microbial activity, and it is very likely that CT loss in bark during storage is largely accounted for by microbial activity. The microbial degradation of CT has long been documented [59]. The generation of heat may also accelerate the oxidation of CT, resulting in further loss. Over time, CT can also form complexes with proteins and other macromolecules, and become more resistant for thiolytic [56, 60], and therefore partially escape detection with the method used in the present study.

The utilization of bark extractives as raw materials for high-value applications requires raw material freshness. Due to the rapid changes occurring in bark extractive fractions during pile storage, it can be concluded that bark should be sent for further processing as soon as possible after debarking. Otherwise, sufficient yields in further processing phases will be compromised as a result of decreased extractives content and changes in the chemical composition of the extractive fractions. In this study the yield of acetone extract decreased about 30% after 2 weeks of storage. The CT content decreased even more, as over 60% of CTs were lost during the same time. This is bound to have an effect on feasible production of CTs and CT-based products, as not only does the amount change, but some less soluble CT-based compounds may also be formed.

The time between harvesting and debarking, the transportation method (by truck or log floating), and the harvesting method (cut-to-length or tree length) affect the characteristics of the bark. The whole supply chain must be efficient to ensure fast delivery; the loss of bark has to be avoided during harvest and transport, and the bark must be kept clean from impurities such as mineral

soil. This is not a problem for harvesting companies using the cut-to-length method and modern enterprise resource planning systems for harvest and transportation, and it also supports the supply of industrial wood, in which the goal is to deliver the trees to the factory in as fresh a state as possible.

The storage behavior of bark extractives is not a well-studied topic. Previous studies of bark storage have been conducted mainly in the context of bioenergy production, and thus the changes in chemical composition have not been investigated extensively. In order to fill the gaps in current knowledge and to better serve the emerging biorefining industry, it would be beneficial to conduct more industrial-scale storage studies for softwood bark and to perform detailed chemical characterization of storage samples.

Systematic studies of the effect of season on the pile storage of bark are also needed, as well as comparative investigations of the effects of different storage methods on bark chemistry. Industrial biorefineries utilizing extractive compounds of wood biomass have more steady year-round demand for feedstock than, for example, the bioenergy sector, which is more dependent on the changing seasons. For this reason, the effect of season on changes in the chemical composition of bark is an important topic that warrants further research.

Conclusions

Biomass storage can have a substantial influence on bioenergy or biofuel economic feasibility, and on the potential environmental benefits. From the energy perspective, this study clearly showed that the energy content (MWh m^{-3}) of the bark pile can increase during storage, while at the same time the moisture content decreases. On the other hand, in poor storage conditions, the energy content and the quality of fuel can decrease markedly.

The results of this study confirm that extractive losses occur rapidly after debarking of wood and piling of the bark, which means that to utilize these valuable compounds, the supply chain from the forest should be accelerated, and material should be sent for further processing as soon as possible after debarking. The results clearly demonstrate that hydrophilic, phenolic extractive compounds in particular are rapidly lost after debarking and piling of bark. About 60% of CTs were lost after 2 weeks of storage, after which the CT content remained rather steady, albeit still declining slightly. About 26% of the quantified lipophilic compounds were lost after 2 weeks of storage, and 63% of the lipophilic compounds still remained at the end of the storage period. The magnitude of the changes in extractives fraction was thus highest in the beginning of the storage period, when most of the changes and losses occurred in both lipophilic and hydrophilic extractives.

Improving bark storage conditions could increase the energy content and decrease the moisture content of the bark, thus improving the raw material for energy use. However, storage

of bark significantly decreases the quality of the raw material for biorefineries. Therefore, the comprehensive utilization of bark requires that efficiency be maintained throughout the supply chain to ensure that tree delivery times are kept short and that loss of bark is avoided during harvest and transport.

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