How may ybrid a Hybride Filler of Precipitated
Calcium Carbonate and Wood Flour be
Employed to Improve Containing Wood Flour
for Paper Quality? Applications—A Comparative
Handsheet Study

ABSTRACT

The main objective of this paper handsheet study is to investigate if a hybrid filler material containing wood flour and precipitated calcium carbonate can replace and/or supplement commercial available ground calcium carbonate and precipitated calcium carbonate mineral filler material for papermaking. The handsheet study contains 25 different furnish mixtures. Four different types of wood flour were used to manufacture the hybrid filler material, including two blends with a strengthening agent. All handsheet were manufactured with an 80% harwood and 20% softwood mix. The filler content varied between 10, 15 and 20%. The study showed that the hybrid filler material achieved a retention of up to 92.68% as well as higher caliper of up to 208 μm compared to commercial ground and precipitated calcium carbonate of 120.4 μm and 145.6 μm respectively. Tensile and tear strength did not show an improvement. Elongation and TEA did improve by up to 30% with the additive containing hybrid filler material. Opacity was improved by up to 10% with the hybrid filler material. Brightness and colour values were lower due to the natural brown colour of the wood flower material.

Comment [1]:

Keywords: Wood. flour, filler, PCC, GCC, additive, strength, retention, papermaking, handsheets, paper properties

1. INTRODUCTION

In the last decade, worldwide efforts in technology and society have been made to find ways to replace ephemeral plastic products with environmentally friendly materials. A promising product with a wide range of uses is paper. In order to further increase its ability to compete with established plastic products, the paper-making process must also be viewed critically with regard to the use of energy and of raw material usage [1]. The following work focuses on the replacement of mineral fillers with Wood Flour (WF). The filler to be replaced in this study is calcium carbonate in the form of Ground Calcium Carbonate (GCC) and Precipitated Calcium Carbonate (PCC), the most commonly used filler material in papermaking [2]. The share of mineral filler in the total raw material input of the paper industry is currently 8%. However, the trend of recent years shows a clear skew_trend towards the production of papers containing higher filler [3].

The application of GCC and PCC mineral filler material in the paper manufacturing process lowers production cost due to the lower cost of filler to raw **fiberfibre** material [2, 3]. The application of filler materials increase optical properties such as whiteness, brightness, and opacity and can have a positive impact on formation [5].

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GCC occurs by the degradation of calcium carbonate. This is a naturally occurring material whose raw stone is extracted from chalk, lime and marble deposits. A great difference to other fillers are its rhombohedral calcite crystals. After several comminution steps and removal of the chemical impurities by flotation, the final particle size distribution is achieved by micronization. Solid contents of the commercial slurries of GCC are about 78%[3]. PCC, on the other hand, belongs to the group of synthetic carbonates, which are obtained through a three-stage process. First, quicklime is produced by thermal removal of CO_2 . Subsequently, the quicklime is converted with H_2O to calcium hydroxide. The reintroduction of CO_2 causes the precipitation of PCC at a solids content of 20-35%, a level much lower than comparable GCC slurries [3].

Since the beginning of the twentieth century, wood flour has been used as an extender for glue and absorbents of explosives. As a filler in the production of plastic parts, wood flour was first processed in 1916_. [6]_ The term "wood flour," for which no clear-cut definition has been adopted, is applied somewhat loosely to wood reduced to finely divided particles approximating those of cereal flours in size, appearance, and texture. A specific method of production is not a criterion often used involved in characterising the name "wood flour." [7] In practical ityly speaking, wood flour usually refers to wood particles that are small enough to pass through a screen with 850-micron openings (20 US standard mesh)

[7]. Earlier studies have shown that the use of wood flour can be an alternative cellulosic based wood additive [8]. A deterioration in brightness and smoothness could be compensated for by the combined use of wood flour with calcium carbonate. [9]

2. MATERIAL AND METHODS

This section describes the materials, **standardized**standardised TAPPI test methods, and procedures, used for this study. Repeatability of the results stayed in between the allowable margins of the TAPPI testing standards.

2.1 TAPPI Methods

Pulp refining was done according to T 200 sp-06 "Laboratory beating of pulp (Valley beater method) [10], Handsheets for physical testing were prepared <u>in</u> accordance with T 205 sp-06 [11], As was tested with T 211 0m-02, "Ash in wood pulp, paper and paperboard <u>were</u>: combustedion at 525°C" [12].

Physical testing of handsheets was performed in accordance <u>with</u>to T 220 sp-06, "Physical testing of pulp handsheets" [13] <u>and</u>, the freeness of pulp wereas measured as Canadian Standard Freeness (CSF) according to T 227 om-09 "Freeness of pulp (Canadian standard method)" [14].

Conditioning of the paper samples was done according to T 402 sp-08_, ("Standard conditioning and testing atmospheres for paper, board, pulp handsheets, and related products)" [15]. Tensile strength was measured in accordance with T404 cm-92 (, "Tensile breaking strength and elongation of paper and paperboard)" [16]. Basis weight was measured withfollowing T 410 om-08 {. "Grammage of Paper and Paperboard (weight per unit area)}" [17]. The paper thickness was measured by T 411 om-10 (t "Thickness (calipercalliper) of paper, paperboard, and combined board)" [18]. The moisture content of pulp was determined by T412 om-06 ("Moisture in pulp, paper and paperboard)" [19]. The tear strength was done by following the T 414 om-12, ["Internal tearing resistance of paper (Elmendorf-type method)]" [20]. The opacity of paper handsheets was performed according to T 425 om-06, {"Opacity of paper (15/d geometry, illuminant A/2°, 89% reflectance backing and paper backing) [21]. Brightness of pulp was measured according to T 452 om-08, {"Brightness of pulp, paper and paperboard (directional reflectance at 457 nm)}" [22]. Stiffness of the paper was measured according to T 489 om-08_{ }, "Bending resistance (stiffness) of paper and paperboard (Taber-type tester in basic configuration) [23]. The paper colorcolour was measured by T 524 "ColorColour of paper and paperboard" [24].

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2.2 Materials

For the handsheet study, the **fiber**fibre materials used was a blend of 80% Sappi Saiccor Eucalyptus Hardwood and 20% AV Terrace Bay Northern Bleached Softwood Kraft [NBSK]. The furnish was beaten to a 320 ml TAPPI CSF freeness in a valley beater following TAPPI T 200 sp-06 method [9].

For the study, two different commercially available GCC and PCC types were used. The first GCC ([GCC1]] used was a GCC powder with a mean particle size of 0.7 μ m. A 20% slurry using **deionized** water was produced for the application in the handsheets. The second GCC ([GCC2)] applied to the handsheets was a commercially available GCC slurry with 76.72% solids content with a mean particle size of 0.73 μ m.

The first PCC ([PCC1]] used was a PCC powder with a mean particle size of 1.0 μ m to 2.0 μ m. A 20% slurry using **deionized** water was produced for the application in the handsheets. The second PCC ([PCC2]] applied to the handsheets was PCC slurry produced on-site at a paper mill site in the U.S<u>A</u>. with 19.58% solids content and a mean particle size of 1.66 μ m.

For the second trial four different precipitated calcium carbonates (PCC) + Wood Flour (WF) were produced and integrated into the handsheets.

To produce the laboratory hybrid WF-PCC, a calcium hydroxide (Ca(OH)₂) type was used suitable for PCC production in combination with two WF types with and without Strength Additive (SA) that is activated on contact drying. WF1 had a particle size distribution of 20µm to 50µm and WF2 had a particle size distribution of 40µm to 70µm.

The produced hybrid WFPCC1 hadto a solids content of 14.35%; WFPCC1SA of 16.99%; WFPCC2 of 13.28%, and WFPCC2SA of 16/35%.

2.3 PCC + Wood flour material preparation

The preparation of the PCCWF product followed the process sequence laid out in Figure 1 which shows the flow of the production process with all components and important data. A Fischer Scientific Isothemp Lab Stirring Hotplate was used for stirring and heating the suspension in a 2000 ml beaker. Industrial grade CO_2 from a 150 lbs. gas bottle with pressure regulator was used as the precipitation reactant. The CO_2 was dispersed in the beaker with an air stone.

Comment [5]:

Which?

Comment [6]:

To what moisture content?

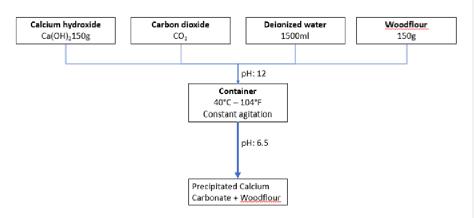


Fig. 1. PCC+ WF process sequence

The preparation of the material started with the weighing of 1500 ml of **deionized**deionised water 150g Calcium di-hydroxide Ca(OH)₂ and 150g of the respective wood flour (Figure 2a). The next process step wasis heating water to 40°C. Subsequently, first GCC and the respective wood flour was added one after the other. In order to avoid clumping and to ensure good mixing, slow addition of the powders and constant agitation was necessary (Figure 2b). After that the addition of CO₂ could be started (Figure 2c). The amount of added CO₂ was determined visually, because the wood flour foamed strongly when too much was added. (Figure 2e). Once the amount was set, it was left **constant**. Throughout the process, the temperature and pH were measured with a pH meter (Figure 2d). The initial pH value was 12.0. The measurement of pH was important because it was the indicator for the precipitation of PCC, the target value of the pH after precipitation was set to a pH of 6.5.

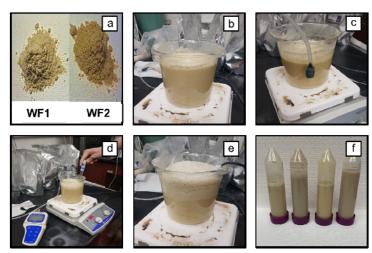


Fig. 2. PCC Production with WF a) WF types, b) WF-CA(OH)₂ slurry, c) CO₂ injection system d) pH and temperature measurement, e) foam built up f) final PCCWF from left: WFPPC1, WFPCC1SA, WFPCC2, WFPCC2SA

Figure 3 shows that all four blends start with the same initial values (pH 12.0, 40° C) and follow the same pattern throughout the process. During the first phase of the reaction the pH stays constant while the temperature increases quickly until a maximum is reached. Striking here is the difference of the length of this phase. The precipitation of WFPCC1 and WFPCC1SA reached a temperature of 56°C is after 33 minutes, and 54°C after 36 minutes respectively. For WFPCC2 and WFPCC2SA precipitation, a temperature a maximum temperature of 55°C after 37 minutes and 52°C after 47 minutes wereas reached respectively. The total precipitation time was between 37 and 47 minutes. As soon as the maximum temperature was reached, the pH level dropped in less than three minutes to the target value of pH 6.5 and the CO₂ influx was turned off. After continuing stirring for 2 minutes without adding more CO₂, the pH levels stabilized around 6.8. The finished slurries were bottled and stored in a cold room at 5°C.

Figure 2f shows the **color**colour differences of the produced WFPCC slurries from left to right:

WFPCC1 with 14.35 % solids content, WFPCC2 with 14.28% solids content, WFPCC1 with 16.99% solids content, and WFPCC2SA with **a** 16.53% solids content. Noticeable is the lighter **color**colour of the WF1PCC after precipitation.

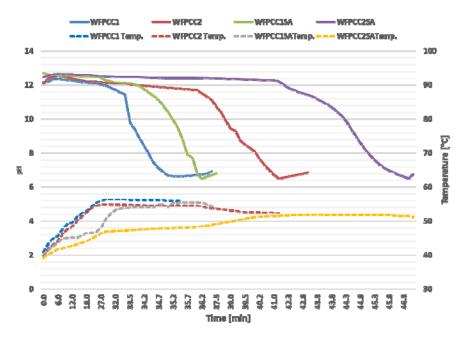


Fig. 3. Process temperature and pH during laboratory PCC production

2.4 Handsheet study

All handsheets contained the same amount of fiberfibre (80% SW and 20% HW) for the base line and the eight filler varieties described above. The target for the basis of the handsheets was 85 g/m². The amount and type of filler was changed in every furnish. The handsheets consisted of a filler variation from 0%, 10%, 15% toand 20% for every filler type that was used. From each variety five handsheets were formed.

The beating of the SW/HW pulp was done in accordance with TAPPI T 200 with a consistency of 1.57±0.04% and a temperature of 23_±_2°C. The pulp was loaded into the Valley Beater, the Valley Beater was operated with no load for 3 min. After that, the initial sample was taken, and the beating was initiated by applying a weight of 5500g to the grinding plate lever. The pulp was refined to CSF value of 320. After the pulp is refined, handsheets are made to the composition mentioned above and tested according to TAPPI standards. When weighing the fillers, it was important to note that the slurries in the bottles were well mixed and all deposits on the bottom were dispelledreleased. Then the weighed fillers were added to the fiberfibre suspension and stirred for about a minute. Subsequently, the <u>mats of</u> handsheets were formed, pressed and <u>conditioned</u> in the laboratory. The sheets which contained strengthener were contact dried after pressing at 120°C for 5 minutes on both sides.

3. RESULTS AND DISCUSSION

All handsheets were tested for basis weight, calipercalliper, stiffness, tear, tensile, elongation, TEA, opacity, brightness and L-/a-/b-values according to the TAPPI testing standards mentioned above and compared to a basis sheet consisting only of ur fiberfibre blend without the addition of any filler.

3.1 Retention

Figure 4 displays the differences, regarding retention, between the eight filler types. For determining the level of retention, we averaged the amount of ash of all three handsheets (10%, 15%, 20% target filler value) which were produced using the same filler. One of the most important parameters of our handsheets was the exact loading of certain quantities of filler. Only by achieving constant amounts of filler can, the physical properties of the handsheets can be compared with each other. Based on experience inon filler retention in handsheet moldsmoulds and the knowledge about the much poorer retention, handsheets were first produced with eight times as much filler as to determine the actual retention and to adjust the dosage factor.

Comparing the GCC powder and the GCC slurry, the powder had a three times higher retention rate than the slurry (8.66% versus 2.58%). In contrast, the PCC powder and the PCC slurry show a nearly identical behavior behaviour (34.32% and 34.77%). In addition, PCC shows better retention than GCC. This e reason for this is probably due to the larger surface, the larger particle size and more branched structure of the PCC.

The PCC slurries produced together with the WF have a significantly higher retention than the pure PCC. The integration of wood flour into the filler offers the possibility of hydrogen bonds between the filler particles and the fibersfibres of the paper.

This is not seendoes not exist with pure inorganic filler. It is noticeable that the type WFPCC1 has a retention of 82.98%, a value much higher than that of the type WFPCC2 with 51.51%. The reason for this could again be the larger particle sizes of the WF product. The best retention of 92.68% was achieved by using WFPCC1SA. THE WFPCC2SA also showed a an improvedncreased retention of 79.13% compared to the commercially available filler materials of..., but lower retention compared to the WFPCC1 product of....

Comment [7]:

give full meaning of acronym ().

Comment [8]:

At what and what temperature and/or pressure etc?

Comment [9]:

If averaged, then error bars are expected which are missing, which undermines the scientific basis of this work

Comment [10]:

An assumption is made that all three levels of the same filler will produce equal amounts of ash which is not a priori sported by any literature. How is that?

Comment [11]:

Which?

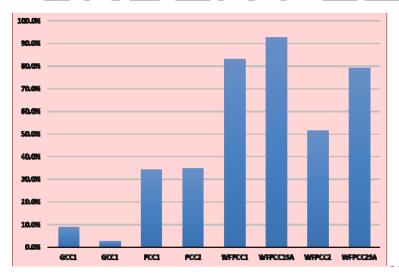
Comment [12]: Which?

Comment [13]:

Impossible to have larger surface area and larger particle size at the same time as the two are variably related. Did you do a detailed particle analysis before coming to this conclusion?

Comment [14]:

Highly untenable, larger particle size does not promote hydrogen bonding but small particle size with more free hydroxyls.



Comment [15]:

No error bars to judge statistical differences between bars which is scientifically unacceptable. GCC2 is sudnely missing, why?

Fig. 4. Filler retention

3.2 Caliper, Basis weight, Stiffness Index

For comparison of the basis weight, the base sheet had 87.54 g/m². The PCC1 handsheet had a basis weight of 87.18 g/m² comparable to the GCC1 basis weight of 85.65 g/m². GCC2 hansheets had a basis weight of 80.94 g/m² while and PCC2 had a basis weight of 92.22 g/m² for PCC2. For handsheets manufactured with wood flour, the basis weight had a range of 107.31 g/m² for (WFPCC1) to 117.03 g/m² for (WFPCC2).

The calipercalliper of the basis sheet and the sheets with GCC filler respectively werehad comparable withproperties of 128.8 µm and 120.4 µm for (GCC1) and 119.2 µm and (GCC2) ...µm for GCC2respectively. The handsheets with the PCC as filler had slightly higher calipercalliper in the tests. The PCC1 had a calipercalliper of 145.6 µm comparable to that ofe PCC2 with 142.8 µm.

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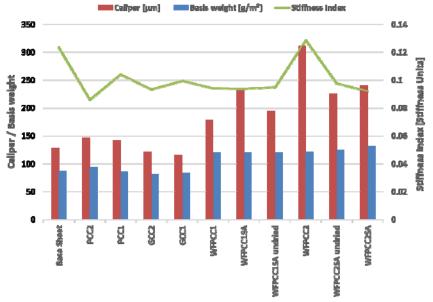
Comparable within which error margins?

Comment [17]:

Comparable within which error margins?

The produced handsheets with wood flour showed different caliper_calliper depending on the used WF. It was noticeable that PCCWF2 with distance showed the highest caliper_calliper with 268 μm. WFPCC2SA + WFPCC1SA had comparable values of 198 μm and 208 μm. The lowest caliper_calliper was achieve was had WFPCC1 with 185 μm, which is still higher than that of the purely inorganic fillers and the basis sheet.

The resulting stiffness index of 0.123 was higher for the basis sheet (0.123) and lower for all other fillers except for WFPCC2 of stiffness index of (0.127). In general, the stiffness of each



filler remained about the same. The range was between 0.096 and 0.107 after calculating the index.

Fig. 5. Basis weight/ Caliper/ Stiffness 15 % filler + WF

The high retention of the wood flour + PCC filler resulted in a final ash content of 15% for the 10% handsheet series. Accordingly, all other test series were <u>all</u> well above their targeted ash content. In order to ensure a comparability, only sheets with the same amount of filler were compared with each other. Since the proportion of fibersfibres was not adjusted to the unexpectedly higher retention, higher basis weights resulted accordingly.

A direct result of higher fiber content and thus higher basis weight is a higher calipercalliper. An additional factor for a higher calipercalliper in contrast to the base sheet is the generally larger particle size of the wood flour and the possibility of swelling of the wood particles. In order to compare strength properties despite different basis weights, an index was calculated. Due to the higher volume of wood flour in contrast to the PCC or GCC, the stiffness of WFPCC2 is comparable to the base sheet. Overall, the wood flour has hardly any influence on the stiffness in the finished paper.

All tests were repeated with 20% filler sheets in Figure 6 to reproduce the previous results that are shown in Figure 6. By a higher proportion of wood flour, any influences are to be displayed more clearly. Additionally, two sets of handsheets were made containing both WF with and without SA, which required contact drying. These handsheets were examined by all the above test methods to find the differences that contact drying makes.

Comment [18]:

Who do you mean? interparticulate distance?

Comment [19]:

Comparable within which error margins?

Comment [20]:

No error bars to judge statistical differences between bars which is scientifically unacceptable.

Comment [21]:

How do you mean? Rephrase.

Comment [22]:

Was drying to specific moisture content. If so swelling as a factor should not play.

Comment [23]:

Which?

Comment [24]: Why so?

ny 50.

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Which?

Comment [26]:

Why? was 20% not part of the initial filler levels?

Comment [27]:

Assumption unsupported by literature a priori. Weak as it stands.

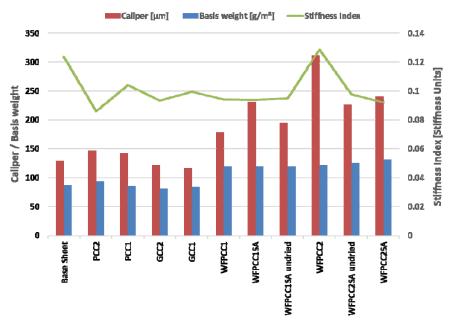
Comment [28]:

To what moisture content?

Comment [29]:

How different or more intense is this contact drying from the one in the previous experiments?

Both basis weight and calipercalliper have increased as expected due to the higher filler load. The stiffness has not changed compared to 15%. The activation process of the SA



shows no influence on **caliper**calliper and stiffness.

Fig. 6. Basis weight/ Caliper/ Stiffness 20 % filler + WF

3.3 Tensile Index and Tear Index

The tensile and tear index in Figure 7 shows a consistently strong correlation. The base sheet has the highest tensile and tear index values (0.69 / 6.07). Adding filler of any kind significantly reduces these strengths. Both PCC2 and PCC2 have comparable values of (0.22/4.76 and 0.22/4.94). When using GCC as the filler, slightly higher values of seen in the tensile and tear index (0.32/5.40) can be seen.

Adding wood flour to the furnish while replacing fibersfibres, lowers the tensile and tear index significantly. Including a SA to the WFPCC shows slightly higher values in comparison with WFPCC without SA. WFPCC1 shows values of tensile and tear index of 0.30/4.31 and values of 0.31/4.64 with SA. WFPCC2 shows values of tensile and tear index of 0.26/4.32 without SA and values of 0.31/4.93 with SA.

Comment [30]:

No error bars to judge statistical differences between bars which is scientifically unacceptable.

Comment [31]:

why so?

Comment [32]:

How did you judge correlation? I do not see an R or R2 value.

Comment [33]:

Why PCC2 and PCC2. No PCC1?

Comment [34]:

How do you mean? Reducing?

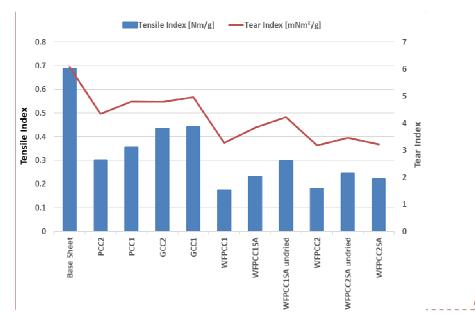
With the use of 20% filler, Figure 8 shows identical results as Figure 7, whereby the individual values have a lower value in comparison to the base sheet. Not to be expected in

<u>handshakes of</u>were the higher strength values of the handsheets with strengthener without contact drying. WFPCC1SA with contact drying had tensile index values of 0.23 and of 0.30 without drying 0.30. The tear index was 3.84 with drying and of 4.23 without. WFPCC2SA with contact drying had tensile index values of 0.22 and of 0.24 without drying 0.24. The tear index was 3.22 with drying and of 3.46 without.

Fig. 7. Tensile / Tear 15% filler + WF

Comment [35]:

No error bars to judge statistical differences between bars which is scientifically unacceptable.



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No error bars to judge statistical differences between bars which is scientifically unacceptable.

Fig. 8. Tensile / Tear 20% filler + WF

3.4 Elongation and TEA

Figure 9 shows elongation and TEA (Tensile Energy Absorption) at 15% filler. The base sheet had a maximum elongation of 1.91% and a TEA of 3.27. In comparison to that, PCC2

values were 1.76% and 1.57 and PCC1 values were 1.94% and 1.75 respectively. GCC2 and GCC1 results were also very close, 1.68% versus 1.72% regarding elongation and 2.10 versus 1.94 regarding TEA respectively.

WFPCC1 without strengthener has similar values similar to those of pure mineral fillers, 1.54% elongation and 1.49 TEA. Adding SA results in an elongation higher than the basis sheet (2.45%) as well as a far less reduced TEA value of 2.32. The positive influence of the strengthener with the bigger wood flour particle size of FLPCC2 is even higher; the TEA value of 3.28 is on the same level as that of thee basis sheet (1.45 WF without SA) of 1.45. The elongation is, with 3.13% higher more than 30% of higher than the maximum elongation of the basic sheet (1.45% WF without SA) of 1.45%.

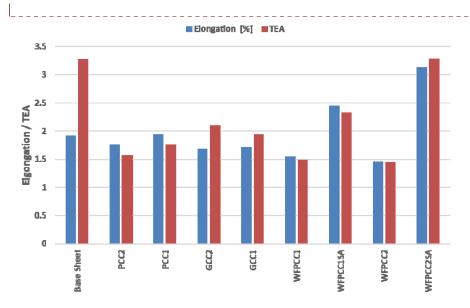


Fig. 9. Elongation / TEA 15% filler + WF

Figure 10 shows that increasing the amount of filler from 15% to 20% changes the influence of the wood flour and the additional strengthener drastically. The pure mineral fillers show a loss of strength in the paper at an increased input. It is noticeable that the use of PCC (1.76%) leads to higher elongation values compared to GCC (1.61%). GCC (1.66), on the other hand, shows a lower decrease in TEA of 1.66 than PCC of (1.57).

The use of wood flour shows <u>an</u> even <u>an</u> greater decrease in <u>relation</u> to elongation and TEA. It can be seen that the wood flour with strengthener exhibit both higher elongation and TEA in the paper. WFPCC1 (1.43% versus 0.88) compared to WFPCC1SA (2.55% versus 2.08). WFPCC2 (1.63% versus 1.37) has also lower values than WFPCC2SA (2.18% / 1.85%). Overall, both values are reduced compared to th<u>ose in whiche</u> <u>use of</u> 15% filler <u>is used</u>.

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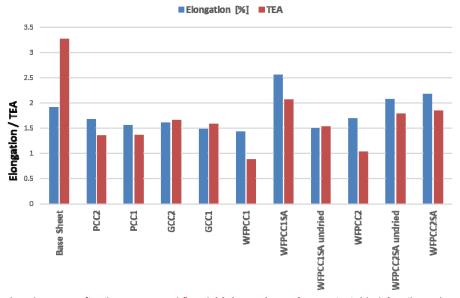
Comment [38]:

Vague. Rephrase to clarity.

Comment [39]:

No error bars to judge statistical differences between bars which is scientifically unacceptable.

However, regarding this strength characteristic, it is interesting to note striking that the activation of the SA by contact drying has a positive effect. Both the elongation and the TEA



show <u>increase after the same wood flour is</u>higher values after contact dr<u>iedying</u> than when using <u>it is</u>the same wood flour _without contact drying. WFPCC1SA without drying shows elongation of 1.50% and a TEA of 1.53. WFPCC2SA without drying shows elongation of 2.07% and a TEA of 1.79.

Fig. 10. Elongation / TEA 20% filler + WF

3.5 Brightness and Opacity

This section describes the results of the optical examinations. Figure 11 shows the comparison of all fillers used in terms of opacity and brightness. Figure 11 only shows the measured values in the use of 20% filler.

As expected, when using mineral fillers, both the opacity and the brightness in the paper increase in comparison to the basis sheet. The base sheet had an opacity of 85.76 and a brightness of 86.91. When PCC was used, the opacity and the brightness increased. For PCC2 a value of 93.41 for opacity and 91.82 for brightness was achieved. When using PCC1 a value of 90.72 for opacity and 90.27 for brightness was achieved. GCC2 showed little differences in the values; of 88.37 opacity and 89.65 brightness. When using GCC1 a value of 89.62 and 88.59 was achieved was achieved for opacity and brightness respectively.

When using PCC + wood flour mixtures, the brightness dropped drastically while the opacity increased significantly. The loss of brightness can clearly be traced back to the naturally darker and unbleached wood flour. The increased opacity can be explained by the increased use of fine material. When comparing the two types of wood flour it was noticeable that the opacity was not dependent on the grade but on the amount of wood flour used. The brightness was higher when using the WF1with smaller particle size than WF2 with the larger particle size.

Comment [40]:

No error bars to judge statistical differences between bars which is scientifically unacceptable.

Comment [41]:

Expatiate. ..as the finer, the more...

Comment [42]:

Highly untenable. Finer particles should make light reflected from the surface of hand-sheets diffuse. That is finer particles will absorb more light and scatter less light, thereby reducing brightness. Present more acceptable explanation.

There were hardly any differences in brightness and opacity within the WF grades. The values ranged between a minimum brightness of 62.91 and a maximum of 64.13. Regarding opacity the minimum value is 98.34 and the maximum 100.46.

WFPCC2 opacity values are very similar, ranging from 98.98 to 100.11. When comparing brightness, the handsheets produced without SA occurs darker (57.35) than the oneshandsheets of the containing WFPCC2SA a composition (62.08 undried and 61.82 dried) which are very similar.

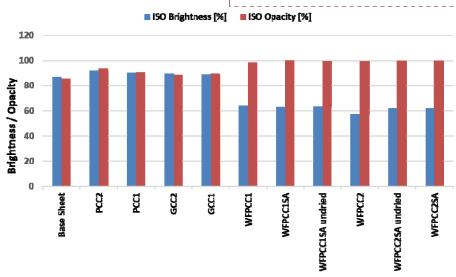


Fig. 11. Brightness / Opacity 20% filler + WF

Figure 12 shows the final produced handsheets with all four wood flour PCC slurries. Picture 12a (from up to down) WFPCC1, WFPCC1SA, WFPCC2SA, and WFPCC2. Picture 12b shows the back sides of the handsheets in opposite direction.

The **color**colour differences were so significant that they were easily visible.

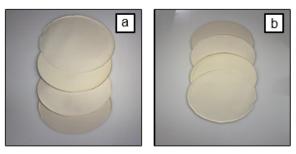


Fig. 12. Handsheets Wood flour and PCC front a) back b)

Figure 13 compares the L-/ a-/ b- values of the handsheets with a 20% filler level to the basis sheet (L: 96.19 / a: $\underline{\ }$ -0.816 / $\underline{\ }$ $\underline{\ }$.2.78).

Comment [43]:

To what? Or you mean they are precise rather in comparison with themselves?

Comment [44]:

To what? Or you mean they are precise rather in comparison with themselves?

Comment [45]:

No error bars to judge statistical differences between bars which is scientifically unacceptable.

The calcium based pure mineral fillers (PCC1, PCC2, GCC1, GCC2) are all in the same range. In general, the L-value of all four papers is higher than the basis sheet, <u>all</u> in a range of 97.7 to 96.47. Looking at the a-values, all four sheets appear less green/ redder than the basis sheet, <u>all</u>. <u>rRanging from -0.426 to -0.542</u>. Comparing b-values, ranging from 1.77 to 2.34 with the handsheets containing filler, the basis sheet appears more yellow/ less blue. Introducing wood flour <u>into</u> the filler blend lowers the L-, <u>then</u> increases the a- and strongly increases the b-value in comparison to the basis sheet.

The three handsheets containing WFPCC1 all have a very similar L- (89.35 to 89.55) and a- (-0.69 to -0.76). The blend without SA has a b-value of 9.73, both sheets containing the strengthener appear more yellow (11.04 and 10.08). Contact drying to activate the strengthener does not interfere with the **color**colour.

The three handsheets containing WFPCC2 are lower in L-readings, WFPCC2 with the WF C 750 FP type being by far the darkest (84.9), the WFPCC2 types containing the SA have a L-values of 88.23 and 88.2. Comparing a- and b-readings, all three papers lay within a very narrow range (a: -0.21 to -0.03 and b: 9.58 to 9.77). The additional drying did not appear to influence **on** the optical properties.

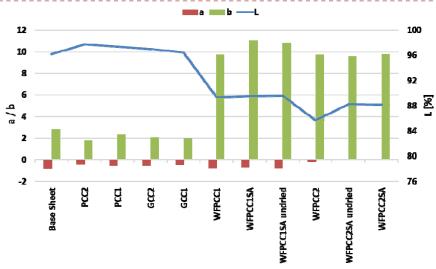


Fig. 13. L - a - b 20% filler + WF

3.6 Suggestions

During the research and the discussion of the results, several proposals are made for future research projects, which could deal more intensively with partial aspects.

To compensate for the significant losses in brightness and whiteness, the wood flour could be bleached.

Another possibility would be the comparison of low filler amounts. Smaller gradings would be advantageous to achieve an ideal ratio. This is mainly because the best results were achieved with the lowest amount of filler in this research work. One possible avenue for further research is the usage of wood flour as a retention aid, specifically, determining how as little wood flour can be added yet still produce an appreciable effect. In this scenario, sheet formation would be an area of interest; conventional retention aids (typically charged polyacrylamides) tend to damage formation by flocculating the fibersfibres. [25, 26] It would be interesting to know if wood flour could aid retention without damaging formation

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No error bars to judge statistical differences between bars which is scientifically unacceptable.

4. CONCLUSION

The main objective of this study <code>iswas</code> to investigate if wood flour can replace commercial mineral fillers for papermaking with a handsheet study. The handsheet study contains 25 different furnish mixtures. For the handsheet study, the base sheet contains a 80%SW/20% HW. The filler content in the use of wood flour and mineral fillers varied between 10, 15 and 20%

The study showed that when_calcium carbonate precipitated in the presence of wood flour, wer the retention of filler increased significantly up to over 92.68% whereas commercial PCC and GCC achieved a maximum of 8.66% and 34.77% respectively. The use of WFPCC resulted in a higher calipercalliper of up to 208 µm compared to commercial GCC and PCC filler which achieved a calipercalliper of 120.4 µm and 145.6 µm respectively.

Tensile and tear strength did not show an improvement for the WFPCC usage in comparison to the commercial GCC and PCC filler material. Usage of WFPCC with SA showed an improvement in comparison to WFPCC without SA.

Elongation and TEA <u>did</u>could did not show an improvement <u>infor</u> the WFPCC usage in comparison to the commercial GCC and PCC filler material. The WFPCC including the SA showed a clear improvement of plus 30% for the 15% filler containing handsheets. For the 20% filler-containing hansheets the improvement is only up to 2%.

The opacity of the paper was increased <u>by</u>by the use of WFPCC <u>from</u>of up to 10 opacity points <u>up</u> to 100.46 <u>points</u> compared to commercial filler material with a maximum opacity of 93.41.

The **coloring**colouring of the <u>wood flour</u> filler has a significantly greater influence on the final **color**colour of the paper than the conventional fillers, even in small amounts. This has resulted in lower brightness of up to 30 brightness points compared to the commercial filler material. In addition and more yellow/cream tones of the paper could be observed.

Promising uses for this special filler type would be, for example, **usage** in corrugationg mediumm **grades**, as properties such as stiffness and **calipers** play a major role in these grades. The optical properties, on the other hand, are rather irrelevant in the middle layer. For these reasons, these fillers could also be used as an internal layer of multi-ply, carton board and coaster board. Wood flour-containing fillers could facilitate the development of cream **colored** grades of paper that use little or no synthetic dye. **Coloring** paper in this way could also be beneficial from an preservative, environmental and marketing perspective.

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Comment [48]:

Wood flour filler did not show any significant improvement in stiffness. So what is this recommendation based on?

Comment [49]:

How do you mean? Middle layers of what?

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References not in conformity with the Harvard system. Please revise.