An Attempt to Analyze the Influence of Properties of Five African Wood Species on Cemented Carbide Tool Wearing

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The dulling of cemented carbide cutting tools after milling five African wood species was studied. The wood specimens investigated varied in high temperature corrosivity (HTC), hard mineral contamination (HMC), and density (D). Attempts to replace the combustion method of evaluating the content of natural HMC in wood with spectral mass analysis methods, such as energy dissipative spectrum (EDAX) or atomic absorption spectrometry (AAS), failed. Experiments showed that scanning electron microscope (SEM) analysis of the ash yielded information on the size and shape of small-dimension, natural HMC particles present in the wood specimens. Employing theoretical multi-variable simulation, the combined effects of the HTC, the HMC, and the D on the cemented carbide tool wear was modeled and appeared to be a good explanation of the results observed.

Keywords: Cemented carbide tool; Wearing; Milling; African Wood Species; Hard mineral contamination; High temperature corrosivity; Wood density

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INTRODUCTION

Previous studies have reported that the machining of some tropical wood species in a dry state excessively dulls high speed steel (HSS) cutting tools (Porankiewicz et al. 2003, 2006; Porankiewicz et al. 2008). This phenomenon was explained to be a result of synergistic effects of mechanical wear (abrasion) caused by the hard mineral contamination (HMC) and high temperature corrosivity (HTC). Chemical and mechanical wearing of the HSS and the cemented carbide tools by tropical wood machining was also the subject of work by Darmawan et al. (2006), but no HTC was considered in that work. The synergistic effect also explained the high tool wear observed in the case of wood with low HMC and high HTC. The effect of wood density was of lesser importance. In recent works by Cristóvão et al. (2009; 2010), studies of the wear of cemented carbide cutting tools when milling low- and high-density, dry, tropical African wood species was examined. Several issues discussed in these works are controversial. In the opinion of the authors of this study, cutting tool wearing when machining wood in the dry state versus in the green state is markedly different. Examining the role of extracts in the case of dry wood cutting experiments has limited value. Similarly, in conducting wearing experiments with dry, solid wood, it is inappropriate to quote studies of cutting tool dulling where green wood or fiber and particle board is cut and machined (in a dry state). The authors’ previous findings,
from Porankiewicz (2003a,b) and Porankiewicz et al. (2006, 2008), state that the basic wear mechanism on dry wood and secondary wood products machining is the combined effects of the HMC and the HTC, whereas when cutting wood in the green state, wet corrosion is of the greatest importance for some wood species and tool materials.

Another controversial issue that should have been mentioned in recent works (Cristóvão et al. 2009, 2010) relates to the cutting tool dullness parameters chosen for wearing experiments. Several parameters for the determination of tool bluntness have been used by researchers because of the irregularity of the real edge round up. In the case of machining solid wood by low HMC and with low HTC, a radius of edge round up, \( \rho \), measured on a plane perpendicular to the edge, seems sufficient. In this case, the edge recession is regular and is the same along the rake, \( A_\gamma \), and the clearance surface, \( A_\alpha \). In the case of presence of HMC and HTC, edge recession along the clearance face \( V_B F \) is usually larger, so this parameter should be chosen. In this case, an average value of the \( \rho \) must be evaluated. In the case of edge chipping, the parameter describing the largest edge wear should be measured. Parameters based on the area of edge wear measured in any plane provide averaged information useless from the point of view of acceptable machining quality and the corresponding edge dullness (Staniszewski and Porankiewicz 1978; Porankiewicz 2014).

The next controversial issue presented in recent works (Cristóvão et al. 2009, 2010) relates to an acceptable content of the HMC. In the case of solid wood from a hot, tropical climate, the term “acceptable HMC” does not seem useful; however, the term “acceptable HMC” is otherwise important in the case of particle and fiber boards because the HTC can be reduced by the modernization of technological processes.

A very important issue related to the influence of the HMC on tool wearing is the size of contaminant particles. This issue has been omitted from many works (Cristóvão et al. 2009, 2010), although the effect of the size of HMC particles on tool wear during solid wood milling has already been reported (Porankiewicz et al. 2003, 2006; Porankiewicz 2008).

The present work is an attempt to analyze cemented carbide tool wearing after the longitudinal milling of five tropical wood species originating from Mozambique.
EXPERIMENTAL

Experiments were performed using an SCM milling machine at Luleå University of Technology, Wood Technology Faculty Skellefteå, according to the works of Cristóvão et al. (2009, 2010) using the following machining parameters: rotational speed, $n = 2900 \text{ min}^{-1}$; diameter of cutter head, number of cutting edges $z = 1$, $D_{CH} = 154 \text{ mm}$; feed speed, $v_F = 3 \text{ m/min}$; depth of cut, $c_d = 1 \text{ mm}$; cutting length per cut, $l_C = 12.41 \text{ mm}$; cutting width, $w_C = 3.9 \text{ mm}$; rake angle, $\gamma_F = 30^\circ$; and clearance angle, $\alpha_F = 15^\circ$. Up-milling of the longitudinal cutting case was used and had an edge orientation angle of $\phi_E = 90^\circ$ and average vector of the cutting speed and cutting plane orientation angles $\phi_V = \phi_S = 9.24^\circ$. The average moisture content was $m_C = 8$ to $9\%$.

The cutting blade was made of grade 701 cemented carbide provided by Sandvik Hard Materials Company (Sweden). This grade contains $89.5\%$ tungsten carbide ($WC$), $10\%$ cobalt ($Co$), and $0.5\%$ other compounds. The Vickers hardness of the cutting blade material was $1600 \text{ HV}_30$, and the Rockwell hardness was $92.1 \text{ HRA}$. The wood specimens examined in the present study are listed in Table 1 and shown in Fig. 2.

Table 1. Wood Specimens Used in the Experiment

<table>
<thead>
<tr>
<th>no.</th>
<th>Common name</th>
<th>Scientific name</th>
<th>Wood density ($\text{kg/m}^3$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Ntholo</td>
<td><em>Pseudolachnostylis maprounefolia</em></td>
<td>726</td>
</tr>
<tr>
<td>2</td>
<td>Namuno</td>
<td><em>Acacia nigrescens</em></td>
<td>1100</td>
</tr>
<tr>
<td>3</td>
<td>Muanga</td>
<td><em>Pericopsis angolensis</em></td>
<td>920</td>
</tr>
<tr>
<td>4</td>
<td>Metil</td>
<td><em>Sterculia appendiculata</em></td>
<td>550</td>
</tr>
<tr>
<td>5</td>
<td>Iron wood</td>
<td><em>Swartzia madagascariensis</em></td>
<td>1100</td>
</tr>
</tbody>
</table>

Fig. 2. Sections in direction towards wood grains of specimens examined: a - perpendicular; b - parallel

The content of the HMC, $C_{MC}$, (mainly silica based compounds), and content of the ash, in the wood specimens were evaluated using a combustion method. The oven-dry wood specimens of weight of about $1 \text{ g}$ were dried to absolute dry state in a chamber dryer in temperature $105 \ ^\circ\text{C}$, then cooled down in an exsiccator and weighted before burning with accuracy of $0.0001 \text{ g}$. For evaluation of the HMC, the ash was etched $30 \text{ min}$ in hot hydrochloric acid and filtered with use of the Whatman glass filters. After filtering, the
glass filters were washed in distilled water and dried out in a chamber dryer. The difference of the mass of dry glass filters after and before filtering was the amount of the HMC. The content of the HMC and content of the ash were related to absolute dry wood specimens before burning. In order to characterize the size, $S_{MC}$, of the HMC particles, 4 fractions (used in previous experiments) were assumed: $f_1 <0; 50>$, average 25 $\mu$m; $f_2 <50; 75>$, average 63 $\mu$m; $f_3 <75; 100>$, average 88 $\mu$m; $f_4 <100; 200>$, average 150 $\mu$m. The size was an average value from maximum and minimum dimensions of particles measured on a scanning electron microscope (SEM) images of Whatman glass filters. The ash was examined, with regards to the size and shape of the HMC particles, using SEM, type FEI Quanta 250 FEG, in a low pressure mode (110 Pa). The ash was also analyzed, with regard to its chemical composition, using an energy dissipative spectrometer EDAX type Genesis XM 2i equipped with a nitrogen-free EDAX Apollo 10 detector.

Thermogravimetric analysis (TGA) was used to evaluate the HTC of the products of the thermal degradation of wood towards Co and iron ($Fe$), the binders of the tool materials. TGA was repeated twice for each wood specimen. The conditions of the analysis, which was performed with a Shimadzu TGA-50 apparatus, were as follows: free flow of ambient air; up to 650 °C; heating rate of 50 °C/min; platinum crucible; proportion of $Co$ and $Fe$ and wood powders was 17 to 20 mg and 1 to 2 mg, respectively; and smoothing filter in TASYS Software 40. The most important part of the method of evaluating the HTC using the Shimadzu TGA-50 apparatus was the unbalanced magnetic field (as high as $2.5 \cdot 10^{-4}$ N/(A·m), at 50 Hz) inside the oven. The HTC peak on the TGA plot is shown in Fig. 3.

![Fig. 3. Parameters of the corrosion peak used for calculations of the quantifier, $Rx$; $dm/dt$ – first derivative of mass, $m$, against time, $t$; $rx$ – height of corrosion peak; $Tx$ – temperature of maximum corrosion peak; $A$ – area of corrosion peak.](image)

The HTC quantifier was calculated using Eq. 1 as the average from two replicate TGA analyses. According to a previous work (Porankiewicz 2008), small peaks were ignored in the evaluation of the HTC quantifier, $Rx$,

$$Rx = rx / m \quad \text{(min}^{-1})$$

where $rx$ is the maximum height of the corrosion peak after maximum degradation of mass (point X in Fig. 3) of the wood specimen (mg/min) and $m$ is the mass of the metal specimen (mg).
According to the functional spectral signature method, carbonyl powder particles of Co and Fe were approximately 2.5 µm in diameter. More details regarding the methodology of the HTC evaluation were described in an earlier work (Porankiewicz 2003b).

The edge recession along the clearance surface, $VB_e$ (Fig. 1) and the edge round up, $\rho$, were evaluated in the works of Cristóvão et al. (2009, 2010) using a light microscope (LM) and are collected in Table 2. A way of evaluating (minimum or average value) of the $\rho$ was not explained in the works quoted. According to Fig. 4, the flank wear, $VB_a^{\phi}$, which is an approximation of edge abrasion on the clearance surface (Fig. 1), because the clearance surface of the blades was not precisely oriented perpendicularly to the optical axis of the microscope, was evaluated.

It must be noted that the choice to use a LM to measure the tool wear parameters, in the case of presence of HMC in the machined wood, is improper because the blade surfaces can be polished at some distance without measurable abrasion (on the order of a hundredth of a micron). The use of contact methods would be more suitable (with the application of a profilograph meter or replica method).

![Fig. 4. LM images of the clearance surface of edges after cutting wood specimens: a – Ntholo, b – Namumo, c – Muanga, and d – Metil, according to Cristóvão et al. (2010)](image)

Numerical analysis of the experimental data collected in Table 2 ($\rho^{\phi}$ and $VB_a^{\phi}$) was performed at the Poznań Supercomputing and Networking Center (PCSS) on a heterogeneous, hybrid computing cluster GPGPU (67x Supermicro H8DGG and 42x Supermicro X9DRT) using an algorithm based on the modified theoretical simulation method developed by Porankiewicz (2008). The quality of the approximation provided by the theoretical simulation was characterized by the summation of the square residuals, $S_R$, as well as the square of the correlation coefficient, $R^2$. 
Table 2. Radius of Edge Round Up, $\rho$, and Edge Recession, $VB$, of Cemented Carbide 701 Edge, Before Machining ($\rho_B$, $VB_{FB}$) and Observed ($\rho^O$, $VB_{\alpha O}$) after Cutting Path, $CP$

<table>
<thead>
<tr>
<th>Species no.</th>
<th>$CP$ (m)</th>
<th>$\rho_B$</th>
<th>$\rho^O$</th>
<th>$VB_{FB}$</th>
<th>$VB_{\alpha O}$</th>
<th>$VB_{FC}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Ntholo</td>
<td>4896</td>
<td>2'</td>
<td>43</td>
<td>5'</td>
<td>140</td>
<td>93</td>
</tr>
<tr>
<td>2. Namuno</td>
<td>4896</td>
<td>2'</td>
<td>9</td>
<td>5'</td>
<td>28</td>
<td>55</td>
</tr>
<tr>
<td>3. Muanga</td>
<td>4896</td>
<td>2'</td>
<td>15</td>
<td>5'</td>
<td>70</td>
<td>55</td>
</tr>
<tr>
<td>4. Metil</td>
<td>4896</td>
<td>2'</td>
<td>5</td>
<td>5'</td>
<td>46</td>
<td>46</td>
</tr>
<tr>
<td>5. Iron Wood</td>
<td>2016</td>
<td>2'</td>
<td>8'</td>
<td>5'</td>
<td>20</td>
<td>20</td>
</tr>
</tbody>
</table>

Note: $VB_{\alpha}$ - flank wear; 1 - values taken from work of Cristovao et al. (2009); * - assumed; 2 - calculated; $VB_{FC}$ - values of edge recession measured on the clearance surface, taken from the work of Cristóvao et al. (2011)

The synergistic effect, $SE$, was calculated as,

$$SE = \frac{(W_x - W_i)}{W_x}$$

where $W_x$ is the maximum tool wear ($\rho^O$ or $VB_{\alpha O}$) by the highest value of the $C_{MC}$ and the largest value of the relatively HTC quantifier $R_W$ and $W_i$ is the tool wear ($\rho^O$ or $VB_{\alpha O}$) by the minimum values of the $C_{MC}$ and the minimum values of the relative HTC quantifier, $R_W$.

RESULTS AND DISCUSSION

In Fig. 5 can be seen as rounded, silica based particles of the HMC, recognized with use of EDAX analysis. Measurements performed showed that most of particles were smaller than 50 $\mu$m and can be assigned as “fraction 1” (Table 3).

Fig. 5. Images of the HMC particles found on glass Whatman filters for: a - Ntholo, b – Namuno, c – Muanga, d – Metil, e – Iron wood; SEM
Measurements performed on SEM images and on the glass filters showed that the silica-based particles were smaller than 50 μm in size and can thus be designated as “fraction 1” (Table 3). Small HMC particles on the glass filters were not as easily visible as those in the SEM images. For one Muanga wood specimen, one larger particle was found and measured (on the glass filter) and was assigned to fraction 4. Table 3 contains information on $C_{MC}$ and $S_{MC}$ of the HMC and the HTC for all wood species examined. Wood specimen 1 (Ntholo), having $C_{MC} = 14524$ mg/kg, was the most mineral-contaminated of all specimens examined. The contents of the HMC in wood specimens 2, 3, 4, and 5 were much smaller (Table 3), especially that of specimen 2.

**Table 3.** Content of the HMC, $C_{MC}$, and Size, $S_{MC}$, of Contamination Particles; Ash Content; and HTC Quantifiers $R_X$ and $R_w$ for Co and the Fe Particles for the Wood Specimens Analyzed

<table>
<thead>
<tr>
<th>no.</th>
<th>Common name</th>
<th>$C_{MC}$</th>
<th>$S_{MC}$</th>
<th>Ash</th>
<th>Co $R_X$</th>
<th>Co $R_w$</th>
<th>Fe $R_X$</th>
<th>Fe $R_w$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Ntholo</td>
<td>14524</td>
<td>$f_1$</td>
<td>6.27</td>
<td>0.0221</td>
<td>3.643</td>
<td>0.065</td>
<td>1.0</td>
</tr>
<tr>
<td>2</td>
<td>Namuno</td>
<td>723</td>
<td>$f_1$</td>
<td>1.54</td>
<td>0.029</td>
<td>5.09</td>
<td>0.1052</td>
<td>1.619</td>
</tr>
<tr>
<td>3</td>
<td>Muanga</td>
<td>2393</td>
<td>0.002 $f_1$; 0.998 $f_4$</td>
<td>2.19</td>
<td>0.0158</td>
<td>2.77</td>
<td>0.1024</td>
<td>1.576</td>
</tr>
<tr>
<td>4</td>
<td>Metil</td>
<td>2525</td>
<td>$f_1$</td>
<td>8.34</td>
<td>0.0057</td>
<td>1.0</td>
<td>0.1031</td>
<td>1.787</td>
</tr>
<tr>
<td>5</td>
<td>Iron wood</td>
<td>1865</td>
<td>$f_1$</td>
<td>0.89</td>
<td>0.0153</td>
<td>2.69</td>
<td>0.0904</td>
<td>1.392</td>
</tr>
</tbody>
</table>

The ash content in the wood specimens, evaluated using the combustion method, differed very much from that used in the work of Cristóvão et al. (2010). For the Ntholo and Metil wood specimens, the relative ash contents differed by about 2 times. Results of the EDAX analysis, as shown in Table 4, show that the largest silica content ($Si$) was found in specimen 1, in agreement with the results of the combustion analysis. A small presence of $Si$ was confirmed in specimen 5. The presence of $Si$ was not confirmed by the EDAX analysis in specimens 2, 3, or 4, which contradicts the results of the combustion analysis. The HMC content, which used to be associated with the silica content in wood specimens evaluated in the present study, with the use of combustion analysis, was much higher than one evaluated in the work of Cristóvão at al. (2010) with the application of the AAS method. The relative difference for the Ntholo wood specimen was a factor of 6.

**Table 4.** Results of EDAX Analysis of Ashes Remaining after Burning the Wood Specimens

<table>
<thead>
<tr>
<th>Species no. and Name</th>
<th>Ca</th>
<th>K</th>
<th>Si</th>
<th>Mg</th>
<th>Al</th>
<th>Na</th>
<th>P</th>
<th>S</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Ntholo</td>
<td>37.3</td>
<td>1.7</td>
<td>1.0</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>2. Namuno</td>
<td>34.9</td>
<td>12.8</td>
<td>0</td>
<td>4.1</td>
<td>2.2</td>
<td>4.1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>3. Muanga</td>
<td>25.1</td>
<td>29.8</td>
<td>0</td>
<td>5.3</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>4. Metil</td>
<td>16.8</td>
<td>29.1</td>
<td>0</td>
<td>7.4</td>
<td>0</td>
<td>0</td>
<td>3.1</td>
<td>0.7</td>
</tr>
<tr>
<td>5. Iron wood</td>
<td>27.8</td>
<td>2.2</td>
<td>1.1</td>
<td>11.6</td>
<td>1.1</td>
<td>0</td>
<td>0</td>
<td>1.7</td>
</tr>
</tbody>
</table>

For Muanga and Metil specimens, the relative values differed by factors of three and four, respectively. Spectroscopy methods such as the EDAX and AAS seemed unsuitable for evaluating the HMC content of wood specimens. The EDAX analysis (Table 4) also showed that calcium (Ca) was the main element present in the ash of wood specimens 1, 2, and 5. In wood specimens 3 and 4, more potassium (K) was found than Ca.
In the ash of all wood specimens, the presence of magnesium (Mg) was found, with the highest amount of 11.6% found in specimen 5. No relationship between the amount of elements shown in Table 4, especially K, Ca, and S, and tool edge wear was found.

Table 3 shows values of the relatively corrosivity, $R_W$, evaluated from TGA plots, as shown in figures Figs. 6 to 10.

![Fig. 6. Plots of TG (m) and dTG (dm/dt) for Co and Fe together for wood specimen Ntholo](image6.png)

It can be seen from these figures that in all wood specimens, the corrosion peaks were large. For Co, the largest corrosion peak was seen for Namuno wood (Fig. 7), 0.579 mg/min at a maximum temperature of 373 °C. With the exception of specimens of Metil wood, all of the HTC peaks for Co were seen at temperatures just after the maximum intensity of the thermal degradation of wood. For the Fe HTC peak, the largest corrosion peak was seen for Muanga wood (Fig. 8), at 2.33 mg/min at a maximum temperature of 288 °C. A very high corrosion peak was also observed for Namuno wood, at 2.294 mg/min at a maximum temperature of 279 °C.

![Fig. 7. Plots of TG (m) and dTG (dm/dt) for Co and Fe together for wood specimen Namuno](image7.png)

From X-Ray diffraction analyses performed in a previous work (Porankiewicz 2003b), it is known that the rapid mass increase (observed as a corrosion peak on the TGA
plot) was a result of the creation of an oxide phase of the metal (Co, Fe) specimens. The presence of such a metal oxide phase was not found before the corrosion peak appeared.

Fig. 8. Plots of $TG (m)$ and $dTG (dm/dt)$ for Co and Fe together for wood specimen Muanga

Fig. 9. Plots of $TG (m)$ and $dTG (dm/dt)$ for Co and Fe together for wood specimen Metil

Fig. 10. Plots of $TG (m)$ and $dTG (dm/dt)$ for Co and Fe together for wood specimen Iron wood
The largest radius of edge round up, \( \rho^O = 43 \, \mu m \), and the largest flank wear, \( VB_{\alpha}^O = 140 \, \mu m \), were observed for Ntholo wood \( (\text{Table 2}) \) and were associated with the highest HMC content, \( C_{MC} = 14524 \, \text{mg/kg} \). The lowest observed radius of edge round up \( \rho^O = 5 \, \mu m \) and the lowest flank wear \( VB_{\alpha}^O = 20 \, \mu m \) were not observed in the same wood species. The value of \( \rho^O \) followed the value of \( VB_{\alpha}^O \) only for Ntholo and Muanga wood. The reason for this was asymmetrical abrasion of the rake and clearance surfaces, particularly due to the presence of damage on the cutting edge after milling Namuno and Metil wood. The lowest \( VB_{\alpha}^O, 20 \, \mu m \), was observed for \( CP = 2016 \, \text{m} \), at low HMC and HTC \( \text{(it would probably be of value } 24 \, \mu m \text{ for the cutting path of } CP = 4896 \, \text{m, using power law interpolation)} \). In case of the Namuno wood specimen, small flank wear was observed \( \text{at low HMC content, } C_{MC} = 723 \, \text{mg/kg} \) and largest HTC, \( R_W = 5.09 \). On the clearance surface of the tool after cutting the Ntholo wood specimen (Fig. 4a), a wide worn area with a corrosion pattern was seen. A similar but smaller worn area with corrosion patterns was seen on the clearance surface of the tool after cutting the Muanga wood specimen (Fig. 4c). These observations suggest that the combined influence of the HMC and the HTC caused these corrosion patterns. Small width of the visible worn area on the clearance surface of the tool after cutting Namuno wood (Fig. 4b) suggest a lower wearing effect which can be also associated with the lowest HMC. The worn area along the edge after cutting Metil wood (Fig. 4d) was almost invisible, but large scratches were seen. This suggests low abrasion and HTC effects but also suggests the presence of large-sized contaminant particles in the Metil wood specimen, but such large HMC particles were not found earlier (Table 3). It is recommended that a larger amount of Metil wood be examined in subsequent combustion analyses. All of the above remarks imply that the HMC probably plays a dominant role in cemented carbide blade wearing while milling the examined wood specimens.

The data in Tables 2 and 3 show that the HTC of wood toward Co, the binder in cemented carbide tool materials, as expressed relatively by the HTC quantifier \( R_W \), was not well correlated with the observed tool wear rate. The HMC content and the HTC had overlapping effects for specimens 1 and 2. In this case, a synergistic effect of mechanical and HTC wear could take place. Although the densities differed very much, the observed edge wear and density were not correlated. Since no clear correlations between the observed edge wear, the HMC content, and the \( R_W \) and the wood density were seen, an attempt to use multi-parameter theoretical analysis, according to the work of Porankiewicz (2008), was performed. The theoretical model developed is described by Eqs. 3 to 15.
\[ \Delta W_1 = \Delta w \cdot q_{dvb} \cdot q_{1R} \cdot q_{1D} \]  
\[ \Delta w_1 = e_1 \cdot C_{P} \cdot \Delta C_{P - 1}^2 \]  
\[ q_{dvb} = e_3 \cdot e_w^{e4} \cdot \ln(W) + e_5 \]  
\[ q_{1R} = 1 + e_7 \cdot R_W^{e8} \quad \text{for } W^p > e_6 \]  
\[ q_{1D} = 1 + e_{14} \cdot D^{e15} \]  
\[ \Delta w_{2R1} = w_{2R1} \cdot S_{f1} \cdot q_{dvb} \cdot q_{2Rf1} \cdot q_{2D} \]  
\[ \Delta w_{2R2} = w_{2R2} \cdot S_{f2} \cdot q_{dvb} \cdot q_{2Rf2} \cdot q_{2D} \]  
\[ q_{2Rf1} = 1 + e_{10} \cdot R_W^{e9} \quad \text{for } W^p > e_6 \]  
\[ q_{2Rf2} = 1 + e_{11} \cdot R_W^{e9} \quad \text{for } W^p > e_6 \]  
\[ q_{Sf1} = e_{12} \cdot f_1 \]  
\[ q_{Sf2} = e_{13} \cdot f_4 \]  
\[ q_{2D} = 1 + e_{16} \cdot D^{e17} \]  

where for the edge round up, \( W^p = P^p \); for the flank wear, \( W^p = VB_d^p \); \( \Delta w_1 \) is the edge wear increase due to contact with wood; \( w_{2R1} \cdot S_{f1} \) and \( w_{2R2} \cdot S_{f2} \) are the wearing effects of a single HMC particle related from the position against the edge for fractions \( f_1 \) and \( f_2 \), respectively; \( \Delta w_1 \) is the quotient of increase of the edge wear due to contact with wood; \( q_{dvb} \) is the quotient of increase of the tool wear due to HTC expressed by the \( R_W \) quantifier; \( q_{1R} \) is the quotient of increase of the tool wear due to wood density, \( D \); \( \Delta w_{2R1} \) and \( \Delta w_{2R2} \) are the edge wear increases because of contact with particles of mineral contamination for fractions \( f_1 \) and \( f_2 \), respectively; \( q_{2Rf1} \) and \( q_{2Rf2} \) are the quotients of increase of the tool wear due to HTC, as expressed by the \( R_W \) quantifier, together with particles of HMC of fractions \( f_1 \) and \( f_2 \), respectively; \( q_{2D} \) is the quotient of increase of the tool wear due to wood density, \( D \), together with particles of HMC; and \( e_w = 2.7182818 \).

For \( \rho_P \), the quality of approximation reached was as follows: \( R^2 = 0.93 \), and \( S_R = 11.6 \). For \( VB_d^P \), the quality of approximated reached was as follows: \( R^2 = 0.94 \), and \( S_R = 168.5 \). The reason for such discrepancies could be inaccuracies in a LM measurement of an edge dullness parameter (\( \rho \) or \( VB_d \)) or in the evaluation of the HMC content. It must be noted that the values of the observed edge recession along the clearance surface, \( VB_f \), given in the work of Cristóvão et al. (2010) for specimens 1 to 5 did not allow reasonable theoretical simulation results (\( S_R > 1400 \)). Graphical illustrations of the relationships between \( \rho_P \) and \( VB_d^P \) are shown in Fig. 11.

The maximum synergistic effect of the \( C_{MC} \) of the particles of fraction \( f_1 \) and the HTC effect, as expressed by the \( R_W \) quantifier evaluated in Fig. 11, reached as high as 96%. The wearing effect of the HTC itself appeared to be the dominant factor influencing cemented carbide blade dulling in the examined cases. The role of \( C_{MC} \) itself was lower and the effect of the \( D \) was the lowest. For the lowest \( R_W \) and the \( C_{MC} \) and the largest \( D \), after a cutting path of \( C_p = 5082 \) m, the predicted \( \rho_P \) and \( VB_d^P \) values reached 3 and 14 \( \mu m \), respectively.
The theoretical simulation performed for the flank wear, $VB_{\alpha}^p$, with other kinds of quantifiers describing the HTC, was evaluated on basis: all of the corrosion peaks, $R_{mx}$; summation of area, $A$, of all corrosion peaks, $R_A$; weighted average quantifier, $R_{mTA}$, containing maximum temperature, $T_x$, in corrosion peaks and the range of the temperatures of the corrosion peaks, $dT_x$, and corrosion peaks area, $A$; weighted average quantifier with ranks, $R_{mTar}$; were unable to obtain better approximation than one with the $R_x$ quantifier. The quantifiers $R_{mx}$, $R_A$, and $R_{mTA}$ were examined in previous works (Porankiewicz et al. 2003; Porankiewicz et al. 2006; Porankiewicz 2008) in relation to HSS cutting tools.

CONCLUSIONS

1. The Ntholo wood specimen was most hard mineral-contaminated with $C_{MC} = 14524$ mg/kg and particle sizes smaller than 25 µm.

2. A wide worn area with corrosion patterns on the clearance surface of the tool after cutting Ntholo wood was explained by a combined effect of the hard mineral contamination and high temperature corrosivity.

3. The results of the theoretical simulation performed show that the most influencing factor on cemented carbide tool wear was the high temperature corrosivity, together with the hard mineral contaminant content, which had a maximum synergistic effect as high as 95%.

4. The relative quantifier of high temperature corrosivity, $R_W$, based on the average value of the corrosion peaks observed on the TGA plots just the after maximum wood mass degradation rate, provided the best fit between the observed and predicted tool wear values in the theoretical simulation performed.

5. The sizes of rounded, small, hard mineral contamination particles can be measured using SEM images of remainings on Whatman glass filters.
6. Spectroscopy methods such as EDAX and AAS seem unsuitable for the evaluation of the hard mineral contaminant content, $C_{MC}$, in wood specimens with respect to tool wear.

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APPENDIX

The following estimators for Equations 3 to 15 for the edge round up, $\rho^3$, were evaluated: $e_1 = 2.94776 \cdot 10^{-5}$; $e_2 = 5.43699 \cdot 10^{-3}$; $e_3 = 4.50867 \cdot 10^{-5}$; $e_4 = 6.85641 \cdot 10^{-3}$; $e_5 = 6.97489 \cdot 10^{-2}$; $e_6 = 5.5911$; $e_7 = 0.57998$; $e_8 = 0.86285$; $e_9 = 0.71746$; $e_{10} = 2.362887$; $e_{11} = 0.46051$; $e_{12} = 0.55724$; $e_{13} = 0.97752$; $e_{14} = 1.65125 \cdot 10^{-2}$; $e_{15} = 9.91695 \cdot 10^{-3}$; $e_{16} = 8.76272 \cdot 10^{-2}$; and $e_{17} = 0.13401$. Rounding the values of the estimators for each Equation to 6 decimal places yielded acceptable deterioration of fit of less than 1%. Reducing the number decimals to 5, 3, or 2 caused unacceptable deterioration of fit by as much as 96 and 108%, respectively.

The following estimators for Equations 3 to 15 for the flank wear, $VB_a^O$, were evaluated: $e_1 = 9.83182 \cdot 10^{-6}$; $e_2 = 8.9264 \cdot 10^{-4}$; $e_3 = 5.31132 \cdot 10^{-5}$; $e_4 = 3.80234 \cdot 10^{-3}$; $e_5 = 7.05214 \cdot 10^{-3}$; $e_6 = 6.45928$; $e_7 = 0.5719$; $e_8 = 0.76568$; $e_9 = 0.43094$; $e_{10} = 3.01585$; $e_{11} = 0.91149$; $e_{12} = 1.17249$; $e_{13} = 2.14328$; $e_{14} = 1.43351 \cdot 10^{-2}$; $e_{15} = 3.30962 \cdot 10^{-2}$; $e_{16} = 0.22752$; and $e_{17} = 0.17442$. Rounding the values of the estimators for each Equation to 6 decimal places yielded acceptable deterioration of fit of less than 1%. Reducing the number decimals from 5 to 2 caused unacceptable deterioration of the fit by as much as 270%.