Optimizing and Modeling Brightness Development in Peroxide Bleaching of Thermomechanical Pulp

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ABSTRACT

Alkaline peroxide bleaching of (chemi)mechanical pulp is a very complicated system where various process factors affect the bleaching performance and pulp properties. Traditional on-factor-at a time method is ineffective and costly in finding the optimal bleaching conditions. In this study, statistical experimental methods which include three steps, i. e, screening, response surface modeling and optimization, were used to find the conditions for maximal brightness development during one stage alkaline peroxide bleaching of TMP which had an initial brightness of 54.5% Elerpho. The TMP was pretreated with EDTA (0.5% on O. D. pulp, 3% pulp consistency, 30°C for 60 minutes) and bleached in a 2L Mark V Quantum Reactor at 750rpm, 7.5% of bleaching consistency and with 0.05% magnesium sulfate addition. The ranges of other factors studied were 1~5% hydrogen peroxide on O. D pulp, 1~4% sodium hydroxide on O. D pulp and 1~4% sodium silicate on O. D pulp, reaction temperature 50~90°C and reaction time 40~180minutes. A models with good predictability was established and the maximal brightness after one stage bleaching was found to be 70% Elerpho at 50°C, 50 minutes, 5% hydrogen peroxide on O. D. pulp, 3.2~3.4% sodium hydroxide on O. D. pulp and 4% silicate on O. D. pulp. However, further studies on other pulp properties such as strength and brightness stability shall be carried out in order to find out the optimal bleaching conditions.

INTRODUCTION

In an increasingly competitive economic climate, pulp and paper producers are looking for

new means of reducing costs and improving product quality. Unfortunately most pulping and paper making processes are quite complex and identifying optimal operation conditions is easier said than done¹⁾. Alkali peroxide bleaching of (chemi)mechanical pulp is one of such cases. Factors influence alkaline peroxide bleaching performance include furnish type, preceding pulping process, pretreatment, bleach liquor stability, bleaching consistency, bleaching agent charge, total alkalinity level, retention time, temperature, etc. Many of these factors are interrelated.

Many experimental studies are still carried out by the so-called one-factor-at-a-time method, i. e., changing the setting of one process variable at a time and recording the results ^{2, 3, 4)}. Despite that it is easy to be used and understood in getting basic information, this method is not only extremely inefficient in its use of resources, but also very difficult for the user to study synergistic effects or interactions both in this peroxide bleaching case and many other cases commonly found in pulping and papermaking processes.

The object of this study is to develop models that can describe the impact of one-stage alkaline peroxide bleaching on the brightness development of thermomechanical pulp. The study demonstrates how statistical multiple regression analysis and response surface methodology can be an efficient tool in identify statistically significant factors and provide predicative models necessary to simulate and optimize the bleaching process.

EXPERIMENTAL

TMP used in this study was made from 10% Korean red pine and 90% Korean spruce. The original brightness of TMP was 55.4% Elrepho. The pulp was first pre-treated with DTPA (0.5% O. D. Pulp) at 3% pulp consistency, 30°C for 60 minutes. Then the pre-treated pulp was dewatered to 30% pulp consistency by centrifugation. The bleaching reactions were carried out in a Quantum Mark V Reactor. The pulp bleaching consistency was kept to 7.5% and the treatment of magnesium sulfate at 0.05% on O. D. pulp. Other factors, including chemical charge of hydrogen peroxide, sodium hydroxide and sodium silicate, reaction time and reaction temperature were varied according to Table 1 for screening experiments

and Table 2 for RSM (Response Surface Modeling) experiments. The residual part of the pulp was neutralized to pH 5.5 by 15% sodium bisulfate. Then handsheets were made according to Tappi standard and brightness was measured with an Elrepho 3000 spectrophotometer.

	Table. 1 Experimental Design in the Screening Stage					
anc	and the Corresponding Brightness Responses.					
Exp.	P	Ak	Si	Ti	Tp	Br
Name	(%)	(%)	(%)	(min)	(°C)_	(%)
NI	1	0.5	0.5	40	95	61.33
N2	5	0.5	0.5	40	50	62.41
N3	1	4	0.5	40	50	61.30
N4	5	4	0.5	40	95	64.20
N5	I	0.5	4	40	50	62.95
N6	5	0.5	4	40	95	62.83
N7	1	4	4	40	95	52.12
N8	5	4	4	40	50	69.16
N9	1	0.5	0.5	180	50	61.77
N10	5	0.5	0.5	180	95	51.94
NII	I	4	0.5	180	95	46.62
N12	5	4	0.5	180	50	69.06
N13	i	0.5	4	180	95	58.46
N14	5	0.5	4	180	50	64.39
N15	1	4	4	180	50	62.88
N16	5	4	4	180	95	59.17
N17	2.25	2.25	2.25	110	72.5	64.38
N18	2.25	2.25	2.5	110	72.5	63.92
N19	2.25	2.25	2.5	110	72.5	65.91

Note: P - peroxide; Ak - alkali, sodium hydroxide; Si - silicate; Ti - time; Temp - Temperature; Br - Brightness

Table. 2 I	Table. 2 Experimental Design in the RSM Stage and				
the Corresponding Brightness Responses.					
Exp.	P	Ak	Si	Br	
Name	(%)	(%)	(%)	(%)	
N1	5	2	2	68.79	
N2	2	4	2	67.36	
N3	5	4	2	69.66	
N4	2	2	4	66.92	
N5	5	2	4	69.41	
N6	2	4	4	67.61	
N7	5	4	4	69.67	
N8	2	2	2.67	67.02	
N9_	5	2.67	2	67.53	
N10_	3	2	2	67.47	
N11	5	3	3	69.50	
N12	3.5	4	3	68.92	
N13	3.5	3	4	69.22	
N14	3.5	3	3	69.25	
N15	3.5	3	3	69.48	
N16	3.5	3	3	69.03	

RESULTS AND DISCUSSION

1. Effects of Bleaching Factors on Brightness Development in Screening Stage

1.1 Modeling of Brightness Development in Screening Stage

Using multiple regression analysis, a model acceptable determination coefficient $(R^2=0.877, R^2_{adj}=0.754)$ and prediction coefficient $(Q^2=0.6645)$ was developed (Table 3). The insignificant effects whose p-values were bigger than the 0.05 were eliminated from the full model in a backward method so that R² does not decrease much while Q2 increases to its maximum. Some factors, like alkali and silicate, though having p-values bigger than 0.05, were not removed from the model according to the hierarchy rule since their interaction with other factors were very significant. The interactions Si*Tp and Ak*Ti, also had p-values over 0.05, were kept in the modeling because eliminating these effects would lead to big decrease in both the R^2 and Q^2 .

Table 3. Model Obtained in the Screening Stage				
Brightness	Scaled & Centered Coefficient	Р		
Constant	61.421	0.000		
Р	1.949	0.021		
Ak	-0.098	0.893		
Si	0.833	0.269		
Тр	-1.376	0.084		
Ti	-3.578	0.001		
P*Ak	2.601	0.005		
Ak*Ti	-1.458	0.070		
Si*Tp	1.106	0.153		
Tp*Ti	-1.661	0.044		
R2	0.877			
R2adj	0.754			
Q2	0.6645			

The model can be summarized as:
Brightness (%) = 61.421 + 1.949P - 0.098Ak +
0.833Si - 1.376Tp - 3.578Ti + 2.601P*Ak 1.458Ak*Ti + 1.106Si*Tp -1.661Tp*Ti, notice
that the coefficients was scaled and centered, so
the levels of factors should also be centered and
scaled when using the model.

From the scaled and centered coefficients listed on table 3 it is seen that the biggest positive effect

on TMP brightness development is the interaction of hydrogen peroxide and alkali, followed by P, Si* Tp and Si. Other factors, ordered by Ti, Ti*Tp, Ak*Ti, Tp, and Ak, have negative effects on brightness development.

1.2 Effect of Hydrogen Peroxide, Alkali and Their Interaction on the Brightness Development of TMP

The effect of peroxide, alkali and their interaction can be seen from Fig. 1 and 2.

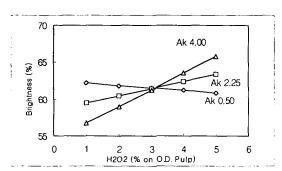


Fig. 1 Effects of H_2O_2 and its interaction with NaOH on brightness illustrated at the center point, i.e., silicate 2.25%, 72.5°C and 110min.

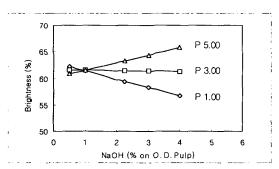


Fig. 2 Effects of NaOH and its interaction with H₂O₂ on brightness illustrated at the center point, i.e., silicate 2.25%, 72.5°C and 110min.

From figure 1 it was seen that increase of peroxide content led to higher brightness when alkalinity is high enough. At very low alkalinity, increase of hydrogen peroxide adversely decreased the pulp brightness. Figure 2 also showed the high dependence of alkalinity on the level of hydrogen peroxide. When hydrogen peroxide content was low (1% on O. D. pulp), increase of alkalinity decreased the brightness development, while when hydrogen peroxide

content was high (5% on O. D. pulp), increase the alkalinity dramatically increased the brightness development of TMP. Interestingly enough, at a medium hydrogen peroxide level of 3%, alkalinity had little influence on brightness of TMP.

The interaction between hydrogen peroxide and alkalinity can be easily explained by the following reaction:

$$H_2O_2 + HO^- \leftrightarrow HOO^- + H_2O$$
 [1]

where the pKa = 11.6 at 25°C.

The hydroperoxide anion or perhydroxyl anion, HOO, is believed to be the principal active species involved in the elimination chromophores in lignin structure. According to the above equilibrium, the concentration of perhydroxyl anion increases as the hydroxide concentration increases, thus peroxide bleaching should be most effective at a high concentration of hydroxide ion, i. e., a high pH. However, this is known not to be the case 5). At least three types of reaction may take place during peroxide bleaching. there are the brightening reactions themselves, which involve reactions between perhydroxyl ions and lignin chromophores. Secondly there are peroxide decomposition reactions as shown by equation [2]. This reaction involves interaction between hydrogen peroxide and its conjugate base, HOO, in the presence of a transition metal catalyst or enzyme 4). Thirdly, there are alkali darkening reactions, which consume caustic, involving formation of colored 6, 7) groups and catalyzed by peroxide decomposition intermediates which include radicals such as the hydroxyl radical (HO•) and the superoxide ion $(O_2 \bullet^{-})^{8}$.

$$H_2O_2 + HOO$$
 $O_2 + OH$ $+ H_2O$ [2]

At too high a pH, decomposition and darkening will likely become dominant while, at too low a pH, the brightening reaction is very slow 9).

1.3 Effect of Time, Temperature and Their Interaction on Brightness Development

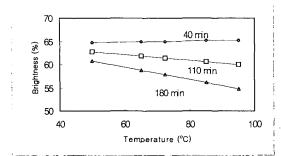


Fig. 3 Effect of temperature and its interaction with reaction time on brightness illustrated at center point, i.e., H_2O_2 3%, NaOH 2.25% and silicate 2.25%

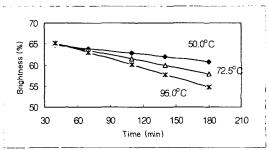


Fig. 4 Effect of reaction time and its interaction with temperature on the brightness illustrated at center point, i.e., H_2O_2 3%, NaOH 2.25% and silicate 2.25%

Figure 3 and 4 showed that increasing either the bleaching temperature or time decreased the brightness development. By higher bleaching temperature and longer bleaching time, detrimental effects on brightness development appeared apparently. This phenomenon was also observed by J. Colodette, et. Al^{-4} , where stonegroundwood pulp was bleached by alkaline peroxide.

The negative effect of long bleaching time at high temperature on pulp brightness is because high temperature and prolonged bleaching time favor decomposition of hydrogen peroxide by a free radical route (equation 3, 4 and 5), releasing oxygen other than singlet oxygen which then initiates the formation of conjugated chromophoric structures and fragmentation of lignin and cellulose ¹⁰⁾.

$$H_2O_2 \rightarrow HOO_{\bullet} + H_{\bullet}$$
 [3]
 $HOO_{\bullet} + OH_{\bullet} \rightarrow O_2 \cdot \bullet + H_2O$ [4]

$$H_2O_2 \rightarrow 2HO_{\bullet}$$
 [5]

It was stated that under routine bleaching conditions at high consistency, time and temperature could be interchanged. An increase in bleaching temperature can compensate for a reduction in retention time up to a point. Conversely, if bleaching temperatures are held below 60°C, extending the reaction time often produces the same brightness response ¹¹⁾. This interchangeable property was not observed in the range investigated in this study, possibly it may be observed under the range of lower temperature and shorter bleaching time. And it is worth mentioning that the exchangeable effect was admitted to have limitations ¹¹⁾.

1.4 Effect of Alkali and Time Interaction on Brightness Development

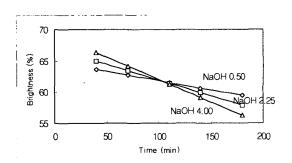


Fig. 5 Interaction of alkali and bleaching time on brightness illustrated at center point, i.e., H_2O_2 3%, silicate 2.25%, temperature 72.5°C.

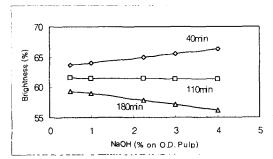


Fig. 6 Interaction of alkali and bleaching time on brightness illustrated at center point i.e., H₂O₂ 3%, silicate 2.25%, temperature 72.5°C.

Table 3 showed that the interaction of alkali and bleaching time was significant at 10% level of significance. Figure 5 and 6 showed when

bleaching time was set at 40 min, increasing the alkali content increased the brightness while at longer reaction time, say over 110min, a reverse result was observed. Besides, Fig 5 showed again that increasing the reaction time decreased brightness over the whole alkali range.

1.5 Interaction of Silicate and Temperature on Brightness Development

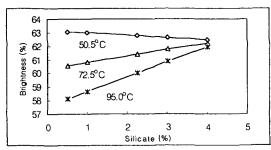


Fig. 7 Interaction of silicate and temperature on brightness illustrated at center point, i.e., H₂O₂ 3%, silicate 2.25%, bleaching time 110 min.

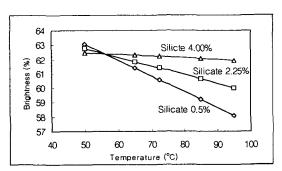


Fig. 8 Interaction of silicate and temperature on brightness illustrated at center point, i.e., H_2O_2 3%, silicate 2.25%, bleaching time 110 min.

Table 3 showed that the interaction of silicate and bleaching temperature was significant at 16% level of significance. As mentioned above, although the 16% level of significance is too big, this effect was not eliminated from the model to keep the large R² and maximum Q² values.

Figure 7 and 8 showed that at high temperature, increasing the silicate content increased the brightness development, but at lower temperatures, increasing the silicate content showed little effect on brightness development. On the other hand, at

low silicate content, negative effect of temperature on brightness appeared, while at high silicate content, brightness is stable over temperature variation.

It was well known that silicate is a costeffective stabilizer for alkaline peroxide bleaching and produces two strong benefits: it significantly reduces peroxide decomposition during bleaching and it improves the internal stability of the bleach liquor itself. Several theories about the role of silicate in the peroxide bleaching of (chemi)mechanical pulp have been suggested. These roles include silicate acting as a peroxide stabilizer, metal ion sequestrant, buffering agent, and metal surface passivator. However, the chemical composition in silicate solution is complicated, it was know that Na2SiO3, Na2Si2O5, Na₂Si₄O₉ and many other polysilicates co-exit in the solution. It is still not very clear how these components contribute to the bleaching process. However, our result indicates that temperature has some effects on the activity of silicate solution.

2. Effects of Bleaching Factors on Brightness Development by RSM (Response Surface Modeling) Methodology

2.1 Model by RSM Methodology

The results obtained from the above screening experiments assume that it will be beneficial to keep both the temperature and reaction time at low levels. Thus in this RSM stage, temperature was set at 50°c and time for 50 minutes. The experiments carried out were shown in table 2. The result of modeling was shown in table 4.

Table 4. Me	Table 4. Model Obtained in the RSM Stage				
	Scaled & Centered Coefficient	P 1.513e-24			
Constant	69.144				
P	1.048	4.870e-8			
Ak	0.315	1.442e-3 0.079			
Si	0.141				
P*P	-0.350	0.031			
Ak*Ak	-0.537	3.240e-3			
R ²	0.9671				
R ² adj	0.9507				
O^2	0.9098				

The model can be summarized as:

Brightness (%)=69.144+1.048P+0.315Ak+ 0.141Si-0.350P²-0.537Ak², where the levels of factors should be centered and scaled.

2.2 Effect of Peroxide and Alkali on Brightness Development of TMP

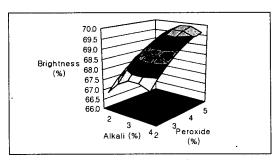


Fig. 9 Effect of Peroxide and Alkali on Brightness.

From figure 9 it can seen that increase the peroxide content increased the brightness development, but at the high level of peroxide addition, the ratio of brightness increase slowed down gradually. On the other hand, alkali showed maximal effects on brightness at a level around 3.5%. No interaction between hydrogen peroxide and alkali was found in the range investigated at this RSM condition.

2.3 Effect of Silicate on Brightness Development of TMP

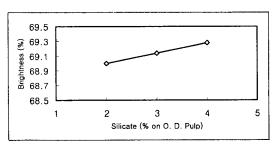


Fig. 10 Effect of silicate on brightness

Figure 10 showed that increasing the silicate content increased the brightness development, but this effect, with the p-value of 0.079, is not significant at 5% level of significance. Thus silicate might not be considered as very important factor.

3. Optimization and Predication

Based on the above RSM model, Modde uses steepest ascent approaches or simplex designs to achieve the optimal conditions which were listed in table 5:

	Table. 5. Optimal bleaching conditions suggested by					
M	Modde and the practical predictability of the model.					
	P (%)	Ak (%)	Si (%)	Brightness Predicated	Brightness Practical	
1	4.9997	3.2419	3.9995	70.028	70.05	
2	5.0000	3.2986	3.9994	70.029	-	
3	4.9984	3.2871	3.9995	70.029	-	
4	5.0000	4.0000	4.0000	69.760	69.70	
5	4.9999	3.4041	3.9905	70.021	70.20	
6	4.9984	3.3128	3.9993	70.028	-	
7	5.0000	4.0000	4.0000	69.761	•	
8	5.0000	3.0000	3.0000	69.842	69.50	

Among the eight experiments predicated by Modde, 1, 4, 5 and 8 was repeated and the result also listed in table 5. It can be seen that the predicated and practical values coincided very well, which means the model developed in the RSM step is very successful.

Thus it can be seen that under the current experimental and the factor ranges investigated, the optimal condition for maximum brightness (70% Elerpho) is: bleaching temperature 50°C, bleaching time 50 min, hydrogen peroxide 5%, sodium hydroxide 3.2~3.4%, and silicate 4%.

However it must be clear that conditions where maximal brightness was achieved, in most cases, are not the optimal condition of bleaching. Other pulp properties, such as strength properties and brightness stability should also be considered in determining the optimal bleaching conditions. This unresolved problem will be studied further in our later research.

CONCLUSIONS

In this study, TMP (10% red pine and 90% spruce) with an initial brightness of 55.4% Elrepho was bleaching by one-stage alkaline peroxide bleaching. The TMP was pretreated with EDTA (0.5% on O. D., 3% pulp consistency, 30°C for 60 minutes) and bleached in a 2L Mark V Quantum Reactor at 750rpm, 7.5% bleaching

consistency and with 0.05% magnesium sulfate addition. The ranges of other factors studied were 1~5% hydrogen peroxide on O. D. pulp, 1~4% sodium hydroxide on O. D. pulp and 1~4% sodium silicate on O. D. pulp, reaction temperature 50~90°C and reaction time 40~180 minutes. Under the above experimental conditions and by statistical experimental methods which include three steps, i. e., screening, response surface modeling and optimization, the maximal brightness development was found to be 70% Elerpho at 50°C, 50 minutes, 5% hydrogen peroxide on O. D. pulp, 3.2~3.4% sodium hydroxide on O. D. pulp and 4% silicate on O. D. pulp. The model developed at the RSM step showed very good predictability on brightness. However, Further study shall be carried out considering other properties such as pulp strength and brightness stability.

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