

Report on the efficacy trials of new pest control products Report for testing Permawood ACQ 1900 for Beat (E.A.) Ltd

Ву

Nellie Oduor Moses Lukibisi Samson Mogire Emmanuel Oduori Peter Sirmah

June 2011

1.0 Introduction

Most untreated wood will be attacked by fungi and insects. Wood in its natural state has got an element of natural durability. This natural durability varies from one species to another and even within the same wood - the heartwood and sapwood. The heartwood, due to its extractives, is more endowed with this durability than the sapwood, which has a lot of sugar/starch deposits that make it susceptible to fungal or insect attack.

Wood is treated by use of chemicals also known as preservatives whose role is to act as a barrier preventing fungi and insects reaching the starch-rich food in the sapwood. Preservatives can be applied by brushing, dipping, sap displacement or pressure impregnation. The Kenya Standard Specification for Preservation of Timber (KS 02-94: 1985 (Confirmed in 1999)) specifies the requirements for preservative treatment of timber; the preservatives, methods of application and suggested average retention levels have all been specified with the object of achieving long service life. Preservatives can be classified into three broad categories: water-borne preservatives, oil-borne preservatives, and light organic solvent preservatives (LOSPs). The efficacy trials will be looking at a water-borne preservative, a copper carbonate and alkyldimethylbenzylammonium also known as Ammoniacal Copper Quat (ACQ).

Objective

- 1. To evaluate the efficacy of **Permawood ACQ 1900** test chemical as a wood preservative against termites and selected fungi
- 2. To test Permawood ACQ 1900 against Tanalith C wood preservative as the standard
- 3. To make recommendation on the suitability of Permawood ACQ 1900 as a wood preservative

2.0 Materials and Methods

2.1 Wood samples

The wood samples measuring 2cm x 2cm x 2cm were sawn from *Eucalyptus grandis* timber. The Eucalyptus species samples were further differentiated by getting samples from the heartwood and sapwood. A total of 168 samples were prepared.

2.2 The treatment of the blocks

One hundred and eight and thirty-six blocks were pressure treated using using Permawood ACQ 1900 and Tanalith C preservatives respectively. The active ingredient(s) for Permawood ACQ 1900 are copper carbonate and alkyldimethylbenzylammonium chloride while that of Tanalith C are copper sulphate, sodium dichromate and arsenic acid. The treatments were carried out at 3 concentrations (2%, 4% and 6%) in ascending order from the lowest to the highest concentrations (to minimize waste of the test preservatives) using vacuum/pressure treatment plant based at the Timber Treatment International (TTI) Eldoret. An initial vacuum of 20mmHg was introduced and held for 5 minutes. Then pressure of 0.7N/mm2 was applied and held for 10 minutes and a final vacuum of 20mmHg was applied and held for 5 minutes. Thus for each concentration, a total of 18 samples were treated while the remaining 54 blocks were untreated (acted as controls of the experiment). Uptake of the preservative was calculated by weighing the wood blocks before and immediately after treatment to the nearest 0.01g. To allow fixation of the preservative in the treated wood samples, the samples were set aside to dry for two weeks.

2.3 Determination of retention

After treatment, the wood blocks were conditioned to allow for distribution of preservative and oven dried to a constant weight. The blocks were then weighed and retention calculated using the following formula;

$$R = \frac{W_2 - W_1}{V} Kg/M^3$$

Where

R = Retention (Kg/ M^3)

- W₁ = Weight of blocks before treatment (Kg)
- W₂ = Weight of blocks after treatment (Kg)
- V = Volume of blocks (M^3)

2.4 Leaching or weathering process

The wood samples were weighed again to the nearest 0.01g. The leaching process used was in accordance with Section 9.3 of ENV 1250-2: 1994. This protocol gives the laboratory method for measuring losses by leaching into water. The weight of the samples was determined. Three conditioned samples per treatment were placed in a beaker with a ballasting device and a magnetic stirrer. 250ml of distilled water (the leaching solution) was added. The contents of the beaker were shaken at a frequency of 60 revolutions per minute. The leaching solution was replaced after 24hr for 4 days. The leachate (the sample of water collected for analysis from the beaker) transferred to another vessel and stored in cold storage facilities for analysis for concentrations of copper, chrome and arsenic using an atomic absorption spectroscopy (AAS). The same procedure would be followed for the second, third and fourth immersions (for each set of treated test specimens) ensuring that in between the immersions the leachate would be transferred and kept aside for analysis

At the end of the leaching process, the wood samples were weighed to the nearest 0.01g and set on racks and exposed to the open laboratory room conditions for 3 days and then set in a conditioning chamber maintained at 25°C and 50% relative humidity for 14 days or until the wood samples reached a constant moisture equilibrium when weighed to the nearest 0.01g.

2.5 Exposure of blocks to termites

Sand was collected, washed and then sterilized in a hot plate for 24 hours. This was put in 108 clear plastic test bottles of 300ml, each ½ full. 30 ml of distilled water was sprinkled till the sand was wet and kept for two hours. Two blocks of the treated and untreated blocks were put into the sand in each of the bottle and subterranean termites (*Macrotermes natalensis* widely distributed in Kenya) from a single colony were introduced according to a procedure adapted from AWPA E1-97 standard (Standard method laboratory for evaluation to determine resistance to subterranean termites, 1997). The test bottles were then kept in an incubator at temperatures between 25-28°C. Out of the treated wood blocks, the samples that were exposed to termites were 9 at each concentration.

2.5.1 Assessment of termite attack

The blocks were inspected weekly for visual rating and after four weeks for weight loss techniques in the laboratory test. During each inspection, the blocks were removed, cleaned by scrapping soil or sand off the blocks surfaces and intensity of termite attack assessed. The attack was rated visually and weight loss basis as indicated below:

I. Visual rating:

Rating of the damage was as follows;

Table 1: Visual rating for termite attack

Description of attack	Rating	Percentage
Sound	0	(0% attack)
Trace	1	(1-10% attack)
Slight	2	(11-30% attack)
Moderate	3	(31-50% attack)
Severe	4	(51-80% attack)
Fail	5	(81-100% attack)



Exposure of the samples to termites

2.6 Setting up the accelerated decay test (Exposure of blocks to fungi)

The ASTM Standards D 1413-99 was used to set the accelerated decay tests.

The strains used in this study were the brown-rot fungi *Wolfiporia cocos* and white rot fungi *Trametes versicolor*. They were all provided by the pathology laboratory of KEFRI. The fungus *Wolfiporia cocos* is known to be tolerant to copper compounds, whereas the *Trametes versicolor* is prevalent on hardwoods products. These fungi are recommended in the ASTM standards (D1413-99).

2.6.1 Preparation of soil substrate

The soil substrate used had a water holding capacity of 30% (see Section 2.6.2 on how to determine the water holding capacity). All the soil clumps were broken, mixed and sieved through a sieve of 2mm square to get fine moistened soil. Ensure the soil is not so wet when it is sifted that the particles again stick together. The soil was steam sterilised. The culture containers (of capacity 250ml) were also steam sterilised. Once the sterilised containers were cooled, they were half filled with the sterilised soil.

2.6.2 To determine the water holding capacity of soil

The water holding capacity was determined by first filling a small Bucher funnel of approximately 50 mm in diameter and 25 mm in depth and fitted with rapid-filtering paper with sieved soil. The soil was compacted in the funnel then the soil surface was levelled by cutting off excess soil with a spatula at the top of the funnel without further compaction. The filled funnel was placed in a 400cm³ beaker and retained in an upright position by wedges at the sides of the funnel. Water was added to the beaker to a depth slightly beyond the level of the filter paper. The soil was allowed to wet by capillarity so as to reduce the danger of entrapping air within the column. When the upper soil surface showed signs of wetting more water was added until the water level was approximately the upper surface of the funnel. The beaker was covered and the soil allowed to soak overnight.

The funnel was placed in a suction flask which was connected to the vacuum pump. Full suction was applied for 15 minutes. During suctioning the funnel was covered with a moist cloth on which an inverted cup was placed to prevent evaporation of water from the exposed soil surface. After 15 minutes the funnel was removed from the suction flask, the soil was scraped into a weighed receptacle and weighed to obtain the wet weight W_1 . The soil was oven dried for 24 hours at 105 ± 2 °C and reweighed, W_2 . The water holding capacity (soil moisture) was determined based on the oven dry weight of soil:

Water holding capacity (WHC), $\% = [(W_1 - W_2)/W_2] \times 100$

2.6.3 Preparation of soil culture containers

Plastic containers with holding capacity of 250 g were prepared and surface sterilized using 95% alcohol. The containers were $\frac{1}{2}$ filled with sterile soil. To determine the amount of additional water needed, the volume of soil that will be used to half-fill a plastic container was weighed W₃. This soil was dried at 105±2°C over night and reweighed W₄. The amount of water to be added to each culture container with that particular soil was calculated as follows:

Water required $g = (WHC \times 0.013 \times W_4) + W_4 - W_3$

2.6.4 Sterilization of treated wood samples and placement in the culture containers

The treated wood samples were sterilized by putting them by retention groups into containers and steamed at 100±2°C for 20 minutes. After cooling, the wood samples were aseptically placed on the soil in the culture containers with soil.

2.6.5 Selection and preparation of test fungi strains

Three strains of fungi were identified; these were *Trametes versicolor* and *Wolfiporia cocos*. The *Wolfiporia cocos* fungus was selected because it is known to be tolerant to copper compounds. The preservative Permawood ACQ 1900 is a copper based preservative. The *Trametes versicolor* fungus is a white rot fungus that attacks hardwood products. The cultures containing these fungi were removed from the herbarium culture and left out overnight to attain normal room temperature. After 48 hrs the bottles containing the fungi were taken in the inoculation cabinet for sub-culturing in order to get new and active strains of pathogens. After sub-culturing, the petri dishes containing cultures were then incubated for 2 weeks to allow the fungus to grow in the growth cabinet of $25^{\circ}C - 30^{\circ}C$.

After incubation, the cultures were taken out (Plate 1 and 2); a sterile cock borer (Plate 3) of about 5mm was used to get an inoculum from the petri dishes of a given species of fungi (Plate4).



Plate 1: The sub-cultured *Trametes versicolor* fungus having been incubated for 2 weeks



Plate 2: The sub-cultured *Wolfiporia cocos* (Syn. *Poria cocos*) fungus having been incubated for 2 weeks



Plate 3: Sterilizing the cock borer over a naked flame Plate 4: Using a cock borer get an inoculum of cultured fungi from the petri dishes

2.6.6 Incubation of test wood blocks

The test wood blocks were weighed before placing them into the culture containers (T_3). During the inoculation process, an inoculum of 5mm from culture plates of a selected fungus was put on the sterilised soil in the culture containers and then a wood block was placed on top of it. (note this was done for all the three fungi). The same procedure of inoculation was done for all concentrations of the test preservative, the standard preservative and controls (one wood block in its own culture container). The culture containers were placed in an incubator set at a temperature of 25°C and kept there for 12 weeks.

2.6.7 Handling of the test wood blocks after exposure to the test fungi

At the end of the 12 weeks the wood blocks were removed from the culture containers. The mycelium was carefully brushed off. Each individual wood block was then weighed to the nearest 0.01 g. The wood blocks were then placed in a conditioning chamber set at 30° C to enable them to attain equilibrium weight. The wood blocks were individually weighed to the nearest 0.01 g (T₄).

The weight loss was calculated from the conditioned weights of the wood block immediately before and after testing as follows:

Weight loss, $\% = (100 (T_3 - T_4)/T_3)$

2.7 Statistical analysis

To evaluate the effect of preservative concentration on the weight loss of the wood blocks for each fungal species, a completely random design was used and then analyzed with the SPSS Statistical Program (Version 17). The variance homogeneity of the variables was verified. The statistical differences were identified using the Tukey test for multiple mean comparisons (P < 0.05).

3.0 Results

3.1 Retention of the chemicals

The amount of chemical/preservative absorbed by the wood blocks is the retention which is given as kilograms per cubic metre (Kg/m^3) of wood.

Chemical	Concentration of chemical	Test material tested	Retention (Kg/m³)
Permawood ACQ 1900	2%	Eucalypt heartwood	1.9
		Eucalypt sapwood	3.9
	4%	Eucalypt heartwood	3.4
		Eucalypt sapwood	6.6
	6%	Eucalypt heartwood	6.8
		Eucalypt sapwood	10.5
Tanalith C	2%	Eucalypt heartwood	2.0
		Eucalypt sapwood	2.9
	4%	Eucalypt heartwood	5.7
		Eucalypt sapwood	9.4
	6%	Eucalypt heartwood	6.5
		Eucalypt sapwood	10.6

Generally the sapwood absorbed more preservative than the heartwood. The heartwood is moderately resistant to preservative treatment, and the sapwood is more permeable.

Collection and identification of termites

Termites that were found on the wood blocks were collected for identification. The specimens collected were mainly workers and soldiers.

3.2 Data Analysis

3.2.1 Analysis of the termite tests

The results of the weight losses of wood blocks and termite survival during the termite tests are as follows. [Table 1 (pg 4) was used to rank termite attack]

Chemical / Preservative	Concentration (%)	sample code	Initial wt (g)	Oven dry wt. (g)	Final wt. (g)	Wt. Loss (g)	% loss	Ranking	no. of termite s after 28 days
Permawood ACQ 1900	2	Eucalypt sapwood	6.238	3.676	3.658	0.018	0.450	1 (trace)	0
	2	Eucalypt heartwood	6.130	3.991	4.020	(0.029)	0.844)	1 (trace)	0
	4	Eucalypt sapwood	6.093	3.732	3.727	0.005	0.122	1 (trace)	0
	4	Eucalypt heartwood	5.514	3.515	3.509	0.005	0.152	1 (trace)	0
	6	Eucalypt sapwood	6.253	4.071	4.045	0.026	0.706	1 (trace)	0
	6	Eucalypt heartwood	6.016	3.911	3.871	0.041	1.057	1 (trace)	0
Tanalith C	2	Eucalypt sapwood	5.35	3.87	3.79	0.08	2.0	1 (trace)	0
	2	Eucalypt heartwood	6.10	5.38	5.26	0.12	2.3	1 (trace)	0
	4	Eucalypt sapwood	5.69	3.88	3.75	0.14	3.5	1 (trace)	0
	4	Eucalypt heartwood	6.01	5.03	4.95	0.07	1.4	1 (trace)	0
	6	Eucalypt sapwood	5.84	4.29	4.22	0.06	1.4	1 (trace)	0
	6	Eucalypt heartwood	6.71	5.70	5.65	0.05	0.9	1 (trace)	0
Control samples	0	Eucalypt sapwood	4.32	3.324	2.726	1.375	34.178	3 (moderate)	0.667
	0	Eucalypt heartwood	4.756	3.748	2.380	1.367	37.713	3 (moderate)	3

Table 2: Results for testing using termites

*Each value represents the means of six replications

Termite galleries were evident after 28 days on untreated blocks. The galleries increased with increase in blocks exposure time. This was rated as moderate (3). The termites generally did not get in touch with the treated block samples. The termites in these test bottles had slow movement which indicated the effect of the chemicals. Termite survival rate is not related to weight loss at all (Kartel S. N. & F. Green, 2003).

3.2.1.1 Analysis for Permawood ACQ 1900 against termites

The mean weight losses in the wood blocks treated with the different chemical concentrations of Permawood ACQ 1900 shown in Table 3 below indicate that wood blocks that had no chemical treatment (control blocks) had higher weight losses than treated block.

	Ν	Mean	Std. Deviation
control	12	1.4592	.12537
2	12	0052	.09296
4	12	.0053	.00465
6	12	.0330	.04114
Total	48	.3730	.63866

Analysis of Variance (ANOVA) was carried out to find out if the differences in mean weight losses were significant (Table 4)

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	18.884	3	6.295	965.664	.000
Within Groups	.287	44	.007		
Total	19.171	47			

The significance value (0.00) is less than 0.05 implying that the means of the concentration of the chemicals applied were statistically different in their effect on the weight loss of the wood blocks. To establish which concentration levels are different in terms of the weight loss on the wood blocks, post hoc tests were conducted (Table 5).

(I)	(J)				95% Confidence Interval	
concentration of chemical	concentration of chemical	Mean Difference (I-J)	Std. Error	Sig.	Lower Bound	Upper Bound
control	2	1.46442*	.03296	.000	1.3764	1.5524
	4	1.45392 [*]	.03296	.000	1.3659	1.5419
	6	1.42617 [*]	.03296	.000	1.3382	1.5142
2	control	-1.46442 [*]	.03296	.000	-1.5524	-1.3764
	4	01050	.03296	.989	0985	.0775
	6	03825	.03296	.655	1263	.0498
4	control	-1.45392 [*]	.03296	.000	-1.5419	-1.3659
	2	.01050	.03296	.989	0775	.0985
	6	02775	.03296	.834	1158	.0603
6	control	-1.42617 [*]	.03296	.000	-1.5142	-1.3382
	2	.03825	.03296	.655	0498	.1263
	4	.02775	.03296	.834	0603	.1158

Table 5: Multiple comparisons between the various chemical concentrations using the Tukey HSD

*. The mean difference is significant at the 0.05 level.

The multiple comparisons analysis indicates that the control wood blocks (no treatment) were significantly different with all the chemical concentrations. There was no significant difference between the concentrations levels of Permawood ACQ 1900.

3.2.1.2 Analysis for Tanalith C against termites

Analysis for the mean weight losses in the wood blocks treated with the different chemical concentrations of Tanalith C indicated that samples that had no chemical treatment were noted to have higher weight losses than treated blocks (Table 6).

	N	Mean	Std. Deviation
control	12	1.4583	.12443
2	12	.0983	.05237
4	12	.1042	.08028
6	12	.0550	.02714
Total	48	.4290	.60583

Table 6: The mean and standard deviation of the chemical concentration of Tanalith C

To investigate if the difference was significant a one-way Analysis of Variance (ANOVA) was carried out

Table 7: ANOVA of the chemical concentrations of Tanalith C

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	16.971	3	5.657	890.653	.000
Within Groups	.279	44	.006		
Total	17.251	47			

From the ANOVA (Table 7) it was noted that the significance value (0.00) is less than 0.05 implying that the means of the concentration of the chemicals applied were statistically different in their effect on the weight loss of the wood blocks. To establish which concentration levels are different in terms of the weight loss on the wood blocks, post hoc tests were conducted.

(I)	(J)				95% Confidence Interval	
concentration of chemical	concentration of chemical	Mean Difference (I-J)	Std. Error	Sig.	Lower Bound	Upper Bound
control	2	1.36000*	.03254	.000	1.2731	1.4469
	4	1.35417 [*]	.03254	.000	1.2673	1.4410
	6	1.40333 [*]	.03254	.000	1.3165	1.4902
2	control	-1.36000*	.03254	.000	-1.4469	-1.2731
	4	00583	.03254	.998	0927	.0810
	6	.04333	.03254	.548	0435	.1302
4	control	-1.35417 [*]	.03254	.000	-1.4410	-1.2673
	2	.00583	.03254	.998	0810	.0927
	6	.04917	.03254	.440	0377	.1360
6	control	-1.40333 [*]	.03254	.000	-1.4902	-1.3165
	2	04333	.03254	.548	1302	.0435
	4	04917	.03254	.440	1360	.0377

Table 8: Multiple comparisons between the various chemical concentrations using the Tukey HSD

*. The mean difference is significant at the 0.05 level.

The analysis of multiple comparisons indicates that the control wood blocks (no treatment) were significantly different with all the chemical concentrations.

3.2.1.3 Compare the two chemicals (Permawood ACQ 1900 and Tanalith C) against termites

	Ν	Mean	Std. Deviation
ACQ	48	.3730	.63866
Tanalith C	48	.4292	.60624
Total	96	.4011	.62002

 Table 9: Comparison of the mean weight loss after exposure to termites against each chemical

The mean of weight loss for Permawood ACQ 1900 was slightly lower than with Tanalith C (Table 9). To assess if the difference was significant a one-way ANOVA was carried out (Table 10).

Table 10: ANOVA of the chemicals against termite attack

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	.076	1	.076	.195	.660
Within Groups	36.445	94	.388		
Total	36.520	95			

The significance level is 0.660 which is greater than 0.05. This implies that there was no significant difference between Permawood ACQ 1900 and Tanalith C in terms of their performance against termite attack.

Table 11: Performance of the chemicals at various concentrations

	Ν	Mean	Std. Deviation
control	24	1.4592	.12261
2	24	.0465	.09080
4	24	.0547	.07513
6	24	.0440	.03589
Total	96	.4011	.62002

The control samples had the highest mean weight loss while the samples that had 6% chemical applied had least mean weight loss. To assess if the difference was significant a one-way ANOVA was carried out (Table 12).

Table 12: ANOVA of the chemical concentrations

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	35.825	3	11.942	1581.120	.000
Within Groups	.695	92	.008		
Total	36.520	95			

The significance value (0.00) is less than 0.05 implying that the means of the concentration of the chemicals applied were statistically different in their effect on the weight loss of the wood blocks. To establish which concentration levels are different in terms of the weight loss on the sample material, post hoc tests were conducted (Table 13).

(I)	(J)				95% Confide	nce Interval
concentration of chemical	concentration of chemical	Mean Difference (I-J)	Std. Error	Sig.	Lower Bound	Upper Bound
control	2	1.41263*	.02509	.000	1.3470	1.4783
	4	1.40446*	.02509	.000	1.3388	1.4701
	6	1.41517*	.02509	.000	1.3495	1.4808
2	control	-1.41263*	.02509	.000	-1.4783	-1.3470
	4	00817	.02509	.988	0738	.0575
	6	.00254	.02509	1.000	0631	.0682
4	control	-1.40446*	.02509	.000	-1.4701	-1.3388
	2	.00817	.02509	.988	0575	.0738
	6	.01071	.02509	.974	0549	.0764
6	control	-1.41517*	.02509	.000	-1.4808	-1.3495
	2	00254	.02509	1.000	0682	.0631
	4	01071	.02509	.974	0764	.0549

*. The mean difference is significant at the 0.05 level.

The significance values of the comparison between the control and the chemical concentration levels of 2%, 4% and 6% levels were less than 0.05. This means that the chemical concentration levels and the control are significantly different. There was no significant difference between the concentration levels.

3.2.2 Results for the accelerated decay tests

Weight loss resulting from fungal attack is the method most frequently used to determine the effectiveness of a preservative treatment to protect wood from decay. Decay resistance can be expressed as either weight loss or residual weight according to ASTM D 2017-81 shown as follows:

Average weight loss (%)	Average residual weight (%)	Indicated class of resistance to a specified test fungus
0-10	90 - 100	Highly resistant (HR)
11-24	76 – 89	Resistant (R)
25 - 44	56 - 75	Moderately resistant (MR)
45 or above	55 or less	Slightly resistant or non- resistant (NR)

Table 14: Decay resistance expressed	as either weight loss or residual weight
--------------------------------------	--

Table 15: The mean percentage weight loss of the wood blocks and their performance rating

Chemical	Concentration of chemical	Fungus exposed	Test material tested	Mean % weight loss	Rating
Permawood	2%	Trametes versicolor	Eucalypt heartwood	4.0	HR
ACQ 1900			Eucalypt sapwood	8.3	HR
		Wolfiporia cocos	Eucalypt heartwood	5.4	HR
			Eucalypt sapwood	6.4	HR
	4%	Trametes versicolor	Eucalypt heartwood	5.2	HR
			Eucalypt sapwood	5.7	HR
		Wolfiporia cocos	Eucalypt heartwood	4.8	HR
			Eucalypt sapwood	5.0	HR
	6%	Trametes versicolor	Eucalypt heartwood	11.6	R
		Eucalypt sapwood	9.0	HR	
	Wolfiporia cocos	Eucalypt heartwood	5.1	HR	
		Eucalypt sapwood	4.7	HR	
Tanalith C 2%	2%	Trametes versicolor	Eucalypt heartwood	4.0	HR
			Eucalypt sapwood	9.9	HR
		Wolfiporia cocos	Eucalypt heartwood	8.9	HR
			Eucalypt sapwood	9.0	HR
	4%	Trametes versicolor	Eucalypt heartwood	5.0	HR
			Eucalypt sapwood	5.2	HR
		Wolfiporia cocos	Eucalypt heartwood	12.0	R
			Eucalypt sapwood	6.7	HR
	6%	Trametes versicolor	Eucalypt heartwood	5.5	HR
			Eucalypt sapwood	5.5	HR
		Wolfiporia cocos	Eucalypt heartwood	4.4	HR
		_	Eucalypt sapwood	4.7	HR
	Control	Trametes versicolor	Eucalypt heartwood	25.8	MR
			Eucalypt sapwood	11.4	R
		Wolfiporia cocos	Eucalypt heartwood	11.8	R
			Eucalypt sapwood	10.7	R

The results of the ranking fungal attack (Table 15) shows generally the chemical treatments made the wood blocks resistant to highly resistant to fungal attack. However, the control blocks were rated resistant to moderately resistant with weight loss ranging between 11- 24% and 25 - 44%.

3.2.2.1 Analysis for Permawood ACQ 1900 against fungi

Analysis for the mean weight losses in the wood blocks treated with the different chemical concentrations of Permawood ACQ 1900 indicated that samples that had no chemical treatment were noted to have higher weight losses than treated blocks (Table 16).

Table 16: Performance of the Permawood ACQ 1900 at various concentrations against selected fungi

	Ν	Mean	Std. Deviation
control	12	.6808	.53695
control	12	.0000	.53095
2% concentration of chemical	12	.2267	.11268
4% concentration of chemical	12	.2133	.03869
6% concentration of chemical	12	.2533	.13131
Total	48	.3435	.33731

The mean weight loss from the control wood blocks was higher than that obtained by the treated blocks. To investigate if the difference was significant, a one-way Analysis of Variance (ANOVA) was carried out

Table 17: ANOVA of the chemical concentrations of Permawood ACQ 1900

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	1.830	3	.610	7.632	.000
Within Groups	3.517	44	.080		
Total	5.347	47			

The significance value (0.000) is less than 0.05 implying that the means of the concentration of the chemicals applied were statistically different in their effect on the weight loss of the sample material. To establish which concentration levels are different, post hoc tests were conducted.

(1)	(L)				95% Confide	ence Interval
concentration of the	concentration of the	Mean Difference				
chemical applied	chemical applied	(I-J)	Std. Error	Sig.	Lower Bound	Upper Bound
control	2% concentration of chemical	.45417 [*]	.11543	.002	.1460	.7624
	4% concentration of chemical	.46750 [*]	.11543	.001	.1593	.7757
	6% concentration of chemical	.42750 [*]	.11543	.003	.1193	.7357
2% concentration of	control	45417 [*]	.11543	.002	7624	1460
chemical	4% concentration of chemical	.01333	.11543	.999	2949	.3215
	6% concentration of chemical	02667	.11543	.996	3349	.2815
4% concentration of	control	46750 [*]	.11543	.001	7757	1593
chemical	2% concentration of chemical	01333	.11543	.999	3215	.2949
	6% concentration of chemical	04000	.11543	.986	3482	.2682
6% concentration of	control	42750 [*]	.11543	.003	7357	1193
chemical	2% concentration of chemical	.02667	.11543	.996	2815	.3349
	4% concentration of chemical	.04000	.11543	.986	2682	.3482

Table 18: Multiple comparisons between the various chemical concentrations using the Tukey HSD

*. The mean difference is significant at the 0.05 level.

The significance values of the comparison between the control and the chemical concentration levels of 2%, 4% and 6% levels were less than 0.05. This means that the chemical concentration levels and the control are significantly different. However, the different chemical concentration levels are not statistically significant from each other in their effect on the weight loss of the wood block.

To investigate the effect of Permawood ACQ 1900 on the selected the fungi ANOVA was carried out after computing the mean weight loss for each fungus (Table 19)

	N	Mean	Std. Deviation
trametes	24	.4117	.45003
poria	24	.2754	.14243
Total	48	.3435	.33731

Table 19: Effect of Permawood ACQ 1900 on the test fungus

The weight loss with *Trametes versicolour* was higher than that of *Poria cocos*. An ANOVA was carried out to find out if the difference was significant (Table 20)

Table 20: ANOVA for effect of Permawood ACQ 1900 on the test fungus

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	.223	1	.223	2.000	.164
Within Groups	5.125	46	.111		
Total	5.347	47			

The significance value (0.164) is greater than 0.05 implying that the means of the weight loss against the two fungi is was not significantly different.

To investigate the effect of Permawood ACQ 1900 on the selected wood block sections (heartwood or sapwood) an ANOVA (Table 22) was carried out after computing the mean weight loss for different wood sections (Table 21).

Table 21: The effect of the weight l	ss on the selected wood	d block sections of either heartwood or
sapwood		

	N	Mean	Std. Deviation
eucalypt heartwood	24	.3792	.45708
eucalypt sapwood	24	.3079	.14467
Total	48	.3435	.33731

Interestingly the weight loss was higher in heartwood sections than with the sapwood. An ANOVA was carried out to find out if the difference was significant (Table 22).

Table 22: ANOVA for effect of Permawood ACQ 1900 on selected wood block sections of either heartwood or sapwood

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	.061	1	.061	.530	.470
Within Groups	5.287	46	.115		
Total	5.347	47			

The significance value (0.470) is greater than 0.05 implying that the means of the weight loss against the two wood sections is not significantly different.

3.2.2.2 Analysis for Tanalith C against fungi

Analysis for the mean weight losses in the wood blocks treated with the different chemical concentrations of **Tanalith C** indicated that samples that had no chemical treatment were noted to have higher weight losses than treated blocks (Table 23).

	Ν	Mean	Std. Deviation	Maximum
control	12	.5042	.12894	.71
2% concentration of chemical	12	.3267	.14896	.58
4% concentration of chemical	12	.2975	.23814	.86
6% concentration of chemical	12	.2125	.08946	.45
Total	48	.3352	.18906	.86

Table 23: Performance of the Tanalith C at various concentrations against selected fungi

To investigate if the difference was significant a one-way Analysis of Variance (ANOVA) was carried out

Table 24: ANOVA of the chemical concentrations of Tanalith C

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	.541	3	.180	6.970	.001
Within Groups	1.139	44	.026		
Total	1.680	47			

The significance value (0.00) is less than 0.05 implying that the means of the concentration of the chemicals applied were statistically different in their effect on the weight loss of the sample material. To establish which concentration levels are different in terms of the weight loss on the sample material, post hoc tests were conducted

Table 25: Multiple comparisons between the various chemical concentrations using the Tukey HSD

(1)	- (J)	Mean			95% Confide	nce Interval
concentration of	concentration of	Difference			Lower	Upper
the chemical applied	the chemical applied	(I-J)	Std. Error	Sig.	Bound	Bound
control	2% concentration of chemical	.17750 [*]	.06568	.046	.0021	.3529
	4% concentration of chemical	.20667*	.06568	.015	.0313	.3820
	6% concentration of chemical	.29167 [*]	.06568	.000	.1163	.4670
2% concentration of chemical	control	17750 [*]	.06568	.046	3529	0021
	4% concentration of chemical	.02917	.06568	.970	1462	.2045
	6% concentration of chemical	.11417	.06568	.317	0612	.2895
4% concentration of chemical	control	20667*	.06568	.015	3820	0313
	2% concentration of chemical	02917	.06568	.970	2045	.1462
	6% concentration of chemical	.08500	.06568	.571	0904	.2604
6% concentration of chemical	control	29167 [*]	.06568	.000	4670	1163
	2% concentration of chemical	11417	.06568	.317	2895	.0612
	4% concentration of chemical	08500	.06568	.571	2604	.0904

*. The mean difference is significant at the 0.05 level.

The significance values of the comparison between the control and the chemical concentration levels of 2%, 4% and 6% levels were less than 0.05. This means that the chemical concentration levels and the control are significantly different from the control on their effect on the weight loss of the wood blocks. However, the different chemical concentration levels are not statistically significant from each other in their effect on the weight loss of the wood blocks.

Table 26: Effect of Tanalith C on the test fungus

	N	Mean	Std. Deviation
trametes	24	.3096	.17758
poria	24	.3608	.20035
Total	48	.3352	.18906

Table 27: ANOVA on the of Tanalith C on the test fungus

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	.032	1	.032	.880	.353
Within Groups	1.648	46	.036		
Total	1.680	47			

The significance value (0.353) is greater than 0.05 implying that the means of the weight loss against the two wood sections is not significantly different.

Table 28: The effect of the weight loss on the selected wood block sections of either heartwood or sapwood

	N	Mean	Std. Deviation
eucalypt heartwood	24	.3454	.21871
eucalypt sapwood	24	.3250	.15809
Total	48	.3352	.18906

Table 29: ANOVA of the weight loss on the selected wood block sections of either heartwood or sapwood

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	.005	1	.005	.137	.713
Within Groups	1.675	46	.036		
Total	1.680	47			

The significance value (0.713) is greater than 0.05 implying that the means of the weight loss against the two wood sections is not significantly different.

3.2.2.3 Compare the two chemicals (Permawood ACQ 1900 and Tanalith C) against fungi

	Ν	Mean	Std. Deviation
ACQ	48	.3435	.33731
Tanalith C	48	.3352	.18906
Total	96	.3394	.27201

Table 30: Comparison of the performance of the chemicals applied against selected fungi

.

The mean of weight loss with Permawood ACQ 1900 was slightly higher than that with Tanalith C. To assess if the difference was significant a one-way ANOVA was carried out.

Table 31: ANOVA of the chemicals against fungal attack

......

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	.002	1	.002	.022	.882
Within Groups	7.027	94	.075		
Total	7.029	95			

The significance level is 0.882 which is greater than 0.05. This implies that there was no significant difference between the Permawood ACQ 1900 and the Tanalith C in terms of their effectiveness on weight loss of the wood sample. They were both effective in controlling weight loss (decay).

Table 32: Performance of the chemicals at various concentrations against fungal attack

	N	Mean	Std. Deviation	Maximum
control	24	.5925	.39241	2.13
2% concentration of chemical	24	.2767	.13890	.58
4% concentration of chemical	24	.2554	.17230	.86
6% concentration of chemical	24	.2329	.11184	.58
Total	96	.3394	.27201	2.13

The control samples had the highest weight loss mean while the samples that had 6% chemical applied had least weight loss mean. To assess if the difference was significant a one-way ANOVA was carried out.

Table 33: ANOVA of the chemical concentrations against fungal attack

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	2.073	3	.691	12.829	.000
Within Groups	4.956	92	.054		
Total	7.029	95			

The significance value (0.00) is less than 0.05 implying that the means of the concentration of the chemicals applied were statistically different in their effect on the weight loss of the sample material. To establish which concentration levels are different in terms of the weight loss on the wood block, post hoc tests were conducted.

Table 34: Multiple comparisons between the various chemica	al concentrations using the Tukey HSD
--	---------------------------------------

(1)	- (J)	Mean		95% Confidence Interval		
()	concentration of the chemical applied		Std. Error	Sig.	Lower Bound	Upper Bound
control	2% concentration of chemical	.31583	.06700	.000	.1405	.4911
	4% concentration of chemical	.33708 [*]	.06700	.000	.1618	.5124
	6% concentration of chemical	.35958 [*]	.06700	.000	.1843	.5349
2% concentration of	control	31583	.06700	.000	4911	1405
chemical	4% concentration of chemical	.02125	.06700	.989	1541	.1966
	6% concentration of chemical	.04375	.06700	.914	1316	.2191
4% concentration of	control	33708 [*]	.06700	.000	5124	1618
chemical	2% concentration of chemical	02125	.06700	.989	1966	.1541
	6% concentration of chemical	.02250	.06700	.987	1528	.1978
6% concentration of	control	35958 [*]	.06700	.000	5349	1843
chemical	2% concentration of chemical	04375	.06700	.914	2191	.1316
	4% concentration of chemical	02250	.06700	.987	1978	.1528

*. The mean difference is significant at the 0.05 level.

The significance values of the comparison between the control samples and the chemical concentration levels of 2%, 4% and 6% levels were less than 0.05. This means that the chemical concentration levels and the control are significantly different. However, the different chemical concentration levels are not statistically significant from each other in their effect on the weight loss of the sample material.

4.0 Conclusions

The test chemical Permawood ACQ 1900 compared well with the Tanalith C chemical for both tests of resistance against termites and selected fungi. There was no significant difference between the Permawood ACQ 1900 and the Tanalith C.; they are both effective in deterring termites and fungal growth on wood blocks.

The three concentrations for each of the chemicals were also not significantly different with both the termite and fungal tests. The chemicals deterred the termites and any weight loss was not significant.

Permawood ACQ 1900 is suitable to be used as a wood preservative against subterranean termites and fungal attack. It is recommended at least 4% concentration to be used for Permawood ACQ 1900 to have full protection against termites and fungi. The ACQ preservatives have similar concentration specifications as those of CCA.

Bibliography

- ASTM, 1999 D 1413-99. Standard test method for wood preservatives by laboratory soil-block cultures.
- ASTM, 1998 D 2017-81. Standard method of accelerated laboratory test of natural decay resistance of woods.
- American Wood Preservers' Association. 1997. AWPA E1-97 Standard method laboratory for evaluation to determine resistance to subterranean termites.
- ENV 1250-2: 1995. Wood preservatives- methods of measuring losses of active ingredients and other preservative ingredients form treated timber. Part 2: Laboratory method for obtaining samples for analysis to measure losses by leaching into water or synthetic sea water.
- Kartel S. N. & F. Green, 2003. Decay and termite resistance of medium density fibreboard (MDF) made from different wood species. International Bio-deterioration and biodegradation 51 (2003) 29-35.
- KS 02-94: 1985 (Confirmed in 1999). Kenya Standard for specification for preservation of timber. Kenya Bureau of Standards, Nairobi Kenya
- Panshin, A. J and Zeeuw, C. 1980. Text Book of Wood Technology. MCGraw-Hill Book Company, New York, ISBN 0-07-048441-4, 720pp.

Signed by:

Nellie Oduor – KEFRI Senior Research Scientist	
Moses Lukibisi – KEFRI Technologist	
Samson Mogire– KEFRI Technologist	
Emmanuel Oduori – KEFRI Technologist (Pathology Laboratory)	
Peter Sirmah – Moi University Lecturer (for termite tests)	