

POTENTIAL OF *UPUNA BORNEENSIS* AND *SHOREA LONGISPERMA* SEED EXTRACTS AGAINST *COPTOTERMES GESTROI*

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Toxicity and repellency of *Upuna borneensis* and *Shorea longisperma* seed extracts were evaluated against the subterranean termite, *Coptotermes gestroi*. Chemical composition of the seed extracts was examined by GC–MS. A total of 17 compounds were extracted from the cotyledon of *S. longisperma* and another 17 were obtained from the seed coat whereas 12 and 14 compounds were extracted from the cotyledon and seed coat of *U. borneensis* respectively. The major compound in *S. longisperma* cotyledon extract was Germacra-4(15),5,10(14)-trien-1-alpha-ol, while the major compounds in seed coat extract were Dihydro-eugenol acetate, β -Thujaplicinol and Torulosol. *Upuna borneensis* cotyledon extract contained four major compounds: 2,4,6-Trimethoxy-toluene, Syringaldehyde, β -Cyclocitral and Allo-cedrol, while three compounds were identified from the seed coat extract: 2'-Hydroxyacetophenone, Methyl p-tert-butylphenylacetate and Iso-Jasmone. Cotyledon and seed coat extracts of *U. borneensis* showed strong repellency against *C. gestroi*. Conversely, cotyledon and seed coat extracts of *S. longisperma*, attracted more termites at higher concentrations. The results suggested that extract of wasted seeds might be used as candidates in termite control (repellents, anti-repellent or antifeedant).

Keywords: Termiticidal, seed extract, subterranean termite, seed coat, cotyledon

INTRODUCTION

Wood can give many years of excellent service if it is properly treated/preserved with biocidal compounds. However, the public is increasingly concerned about the harmful environmental effects of a number of excellent wood preservatives such as chromated copper arsenate. Malaysia, due to its humid weather, loses valuable wood in service as a result of insect attacks and other biodegrading agents. Each year, massive economic losses resulting from the maintenance and repair of wood structures can be attributed to termite attack.

In recent years, much research has focused on the development of safer wood preservatives which are toxic against wood damaging agents but have minimal impact on the environment. This research includes many plant-derived compounds that are considered to be eco-friendly (Antwi-Boasiako & Damoah 2010). Plant-derived materials are known to be effective against insect pests (George et al. 2014), inciting investigators to screen for their toxic effects for use as insecticides (Singh & Singh 2012).

The use of seed extracts for wood preservation has been reported before as many of these compounds have anti-termite and repellent properties (Yang 2009). For example, using treated filter papers, *Jatropha curcas* seed extract at different concentration (20, 30 and 35 g mL⁻¹) and *Azadirachta indica* (35 g mL⁻¹) killed 100% of *Macrotermes* spp. workers after 72 hours of exposure in laboratory conditions (25 ± 3 °C and 60–70% relative humidity) (Addisu et al. 2014). Other studies found varying degrees of termiticidal activity of *Parkia biglobosa* seed extract against *Coptotermes intermedius* and antitermitic activities of *Mentha arvensis*, *Ocimum bacilicum*, *Plantago ovate* and *Cichorium intybus* against *Coptotermes heimi* (Olugbemi 2012, Aihetasham et al. 2016). Even at a low concentration of 1%, extract of *Jatropha curcas* seeds was proven to be effective against *Microcerotermes beasoni* (Singh & Sushilkumar 2008).

Upuna borneensis is an endangered monotypic plant from the family of Dipterocarpaceae. It is endemic to Borneo and commonly known as

upun, *upun batu*, *penyau*, *balau penyau*, *cangal tanduk* or *penyau tanduk*. It is a large tree with low stout buttress. The bark is dark brown, finely flaky with non-laminated inner bark. The wood is commonly used for construction, heavy-duty furniture, railway sleepers and boat making due to its high density (945–1040 kg m⁻³ at 15% moisture content) (Wong 1982). *Upuna borneensis* has more than 14 resveratrol derivatives, including upunaphenols B (1), C (4), D (5) (resveratrol tetramer) and E (6, resveratrol dimer with a C6–C1 unit), which were isolated from acetone-soluble part of its stem (Ito et al. 2005). Stilbenes (resveratrol) which are present in many naturally durable woods have been associated with the durability of heartwood (Schultz & Nicholas 2000, Hassan et al. 2018). Resveratrol causes weight reduction of the groundnut pests, *Spodoptera litura* and *Amsacta albistriga*, and attracts the egg parasitoid *Trichogramma chilonis* under laboratory conditions (Sambangi & Rani 2016).

Shorea longisperma, commonly known as *meranti damar hitam*, is a threatened timber tree also in the family Dipterocarpaceae. *Shorea longisperma* has large, diffuse, cauliflower-shaped crown that can grow up to 70 m tall. The tall, straight, cylindrical bole can be up to 290 cm in diameter with stout buttresses up to 4 m high and extending outwards (Soepadmo et al. 2002). The wood is lightweight, soft, moderately durable but susceptible to biological attack and the density is variable (370–860 kg m⁻³ at 15% moisture content). This timber species can be found in Brunei, Indonesia (Sumatra) and Malaysia (Sarawak) (Yong et al. 2011).

Previous studies found that this genus contains lignostilbenes such as resveratrol (Ito 2011) which exhibited a variety of bioactivity including anti-termite and anti-fungal properties

(Roszaini & Hale 2019). In this study, we analysed the chemical constituents of extracts from waste cotyledons and seed coats of *U. borneensis* and *S. longisperma*. These species were chosen because of the inability of the seeds to germinate in storage. Potential uses for these seeds that otherwise would have been wasted, were being examined. We also investigated the toxicity and repellent effects of both seed extracts against the subterranean termite, *Coptotermes gestroi*.

MATERIALS AND METHODS

Termites

Coptotermes gestroi (soldiers and workers) were collected from active field colonies around the Forest Research Institute Malaysia (FRIM) campus using previously described method by Kirton et al. (1998). Cardboard pieces were moistened with distilled water, stacked together in a large basin, covered with aluminium foil and placed in dark plastic bags that were attached to trees containing termite colonies for two weeks. Termites moved from the tree into the cardboard where they could be easily collected into the basin.

Plant materials

Seeds of *U. borneensis* and *S. longisperma* were procured from trees at the FRIM campus and stored in airtight containers. Seed coat (wing) and cotyledon were separated (Figure 1) using a knife. The cotyledon was split to get its contents. Seed coat and cotyledon were air-dried and stored in an airtight container before being separately ground to pass through 250-µm sieves.

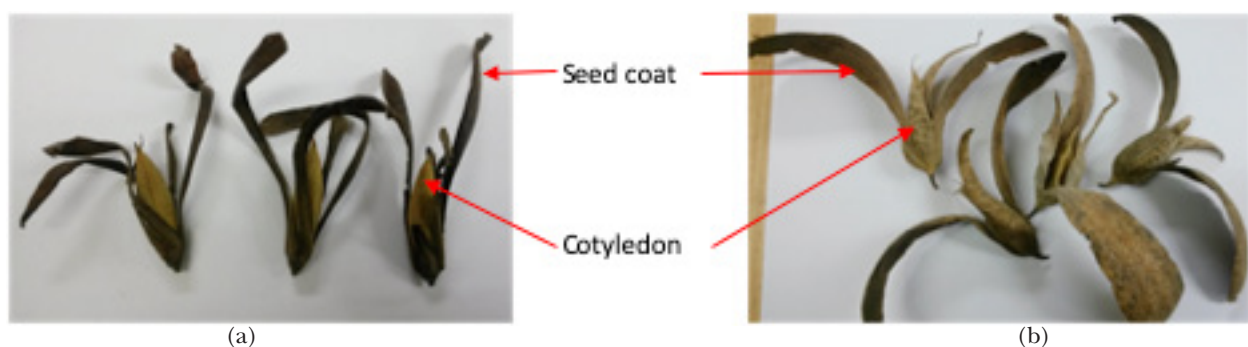


Figure 1 Seeds of *Upuna borneensis* (a) and *Shorea longisperma* (b)

Extraction

Methanol (analytical grade) was used to extract compounds from the seed parts according to ASTM Standard D1105-96 (ASTM 2001) with slight modifications. Five replicates of 5 g (air-dried at 103 ± 5 °C) of ground seed part were weighed to the nearest 0.1 mg, then extracted using 240 mL methanol in a pre-weighed cellulose thimble (porosity 1, height 95 mm) for 5 hours (4–6 siphons hour⁻¹) in a Soxhlet extractor. The collected solvent extract was then filtered and the filtrate was evaporated to dryness under reduced pressure at 45 °C using vacuum rotary evaporator. The dried extract was used to produce a series of concentrations. The extracts were stored in small vials at room temperature.

Chemical analysis

The chemical composition of the extract was analysed by gas chromatography and gas chromatography mass spectrometry (GC–MS) at similar temperature conditions and parameters as previously described (Roszaini et al. 2013). Quantitative analysis was carried out using GC or GC–MS equipped with fused silica capillary columns CBP5 (25 m × 0.25 mm × 0.25 mm) or HP-5MS column (30 m × 0.25 mm × 0.25 mm) respectively. The GC was equipped with a flame ionisation detector using split-mode injection technique, and the operating parameters were helium gas as the carrier gas at a flow rate of 1 mL min⁻¹, an injector temperature of 250 °C and a detector temperature of 250 °C. With the CBP5 column, the GC was programmed initially at 60 °C for 10 min, followed by 230 °C for 1 min at 3 °C min⁻¹. The temperature programme for GC–MS analysis was set similar to the GC programme. Major chemical constituents were identified by comparison of their retention indices with the literature values (Adam 2001) and their mass spectral data with those of the mass spectral database.

Toxicity assay

The method described by Roszaini et al. (2014) was employed to assess the efficacy of extracts against termites. Dried extracts diluted with methanol were tested at concentrations of 0.5, 1, 2, 3 and 4%. The solutions were applied to filter papers discs (9.0 cm diameter and 1.5 mm thick)

and dried in laminar air flow for 1 hour. The weight of filter paper was measured before and after treatment. Untreated filter paper and filter paper treated with methanol served as controls with each of the tests repeated three times. Fifty active *C. gestoi* (45 workers and 5 soldiers from third instar) were introduced into each Petri dish (9.1 cm diameter and 1.6 cm height). A few drops of water were added periodically to the bottom edge of each Petri dish. The Petri dishes were placed in an incubator in darkness at 26 ± 2 °C and 65 ± 5 % relative humidity. Termite mortality was recorded every 24 hours until 100% mortality was observed. Dose-mortality was developed based on exposure time and the lethal concentration required to kill at least half of the termites exposed (LC₅₀). Probit analysis was used for wood extract concentrations series tested (Finney 1971).

Repellence assay

The ability of the extracts to repel termites was assessed according to McDonald et al. (1970) with some modifications. An amount of 1 mL extracts at different concentrations (diluted with methanol) was applied to half the surface of the 9 cm diameter filter paper (the filter paper was cut into two) while the other half was treated with 1 mL methanol. The Petri dishes were dried in a laminar flow hood for one hour and treated and control halves were attached with adhesive tape. Each dish was placed in a 9.1 cm diameter Petri dish and 50 workers of *C. gestroi* were put in the centre of each dish. The number of termites present in the control (NC) and the treated halves (Nt) was assessed hourly for 4 hours. The percentage of repellent was calculated using the formula below:

$$\text{Repellent percentage} = \frac{[Nc - Nt]}{[Nc + Nt]} \times 100 \quad (1)$$

Statistical analysis

The formula by Abbott (1925) was used for the probit analyses. Changes in properties of treated and untreated specimens were evaluated using a one-way analysis of variance. The effects of concentration of extractives on weight loss, density and feeding inhibition were determined. Duncan's multiple range test was used for ranking of the average values of the measured property.

RESULTS AND DISCUSSION

Extract yields

Table 1 shows the yield of extracts produced using the solvent system. Cotyledon and seed coats from *U. borneensis* produced greater extract yields than *S. longisperma* seeds. *Upuna borneensis* cotyledon produced the highest yield (14.31%) while *S. longisperma* cotyledon, the lowest (6.58%). Variation in extract yield and chemical composition of extracts depends on several factors including growing conditions, wood species (different timber has different characteristics), differences between trees of the same species and within the tree itself (different anatomical structure), the products and reagents used in the extraction, method of extraction, drying temperature, conditions, temperature and time (Xu et al. 2009).

Chemical compounds

Chemical composition of *S. longisperma* and *U. borneensis* cotyledon and seed coat extracts are presented in Tables 2 and 3. The analysis of *S. longisperma* cotyledon extracts revealed that Germacra-4(15),5,10(14)-trien-1- α -ol (16.11%), Spathulenol (9.88%), 10-*Epi*-Cubebol (9.81%), *Allo*-Aromadendrene epoxide (9.15%), 3-Thujopsanone (8.00%), *Allo*-Cedrol (6.72%), 6*Z*, 10*E*-Pseudo phytol (6.46%), Methyl *p*-tert-butylphenylacetate (6.00%) and *Z*-Caryophyllene (5.76%) were the main components. Eugenol acetate (40.71%), β -Thujaplicinol (19.83%), Torulosol (13.11%) and *trans*-Pinane (7.85%) were the main components for *S. longisperma* seed coat.

Table 3 shows the 26 compounds identified from *U. borneensis* seed extracts. 2,4,6-Trimethoxy-Toluene (17.92%), Syringaldehyde (16.49%),

Allo-Cedro (16.36%), β -Cyclocitral (12.18%), Presilphiperfolan-8-ol (9.92%), Carvacrol (8.96%) and *Cis*-Vertocitral C (5.27%) were the major constituents identified in *U. borneensis* cotyledon extracts. Of the 14 compounds identified from seed coat extract of *U. borneensis*, eight compounds were the main components: 2'-Hydroxyacetophenone (19.01%), Methyl *p*-tert-butylphenylacetate (14.84%), *Iso*-Jasmone (13.02%), Methyl linoleate (9.04%), Methyl hexadecanoate (8.51%), Incensole oxide (6.52%), Methyl vanillin (6.08%) and Hillyl acetate (5.43%).

Toxicity

Seed extracts of both species were toxic to the subterranean termite; *C. gestroi*. The amount of filter paper consumed by termites was lower for filter paper treated with different concentrations of seed extracts of both *U. borneensis* and *S. longisperma* than control papers (Table 4). This meant that termites consumed smaller amount of treated paper because the extracts were poisonous to them.

Theoretically, as the concentration of a substance increases, the more constituent content is present in the liquid. If the substance is toxic, the more concentrated the substance, the more toxic it will be. The results of this study were in accordance with this theory. The higher the concentration of the extracts, the lower the percentage of paper consumed. This indicated that the diet of *C. gestroi* was inhibited with the increasing concentration of the tested extract.

Table 4 also shows that even at the highest concentration tested (4%), termites still consumed the filter paper treated with both parts of *U. borneensis* seed extracts. A higher concentration is needed to stop filter paper consumption. Unlike *U. borneensis* (at the same

Table 1 Average extract content from *Shorea longisperma* and *Upuna borneensis* seed coats or cotyledons

Species	Extract yield (%)	
	Cotyledon	Seed coat
<i>S. longisperma</i>	6.58 (1.215) ^{dc}	8.52 (1.400)
<i>U. borneensis</i>	14.31 (0.462) ^a	12.64 (3.190) ^b

Values in brackets are standard deviations; percentage values followed by the same letter are not significantly different in the same group (percentage of extractive yields) at the 0.05 level of probability; n =5

Table 2 Major secondary metabolite components found in extracts of *S. longisperma* cotyledon and seed coat

Compound	Class of compounds	Area (%)
Cotyledon		
Germacra-4(15),5,10(14)-trien-1- α -ol	Sesquiterpenes	16.11
Spathulenol	Sesquiterpenes	9.88
10- <i>epi</i> -Cubebol	Sesquiterpenes	9.81
<i>Allo</i> -Aromadendrene epoxide	Sesquiterpenes	9.15
3-Thujopsanone	Sesquiterpenes	8.00
<i>Allo</i> -Cedrol	Sesquiterpenes	6.72
6 <i>Z</i> , 10 <i>E</i> -Pseudo phytol	Sesquiterpenes	6.46
Methyl <i>p</i> -tert-butylphenylacetate	Ester	6.00
<i>Z</i> -Caryophyllene	Sesquiterpenes	5.76
6 <i>E</i> , 10 <i>Z</i> -Pseudo phytol	Sesquiterpenes	4.11
Methyl nidoresedate	Ester	3.72
11,12-Dihydroxy-Valencene	Sesquiterpenes	3.27
<i>Iso</i> -Longifolol	Sesquiterpenes	3.18
Globulol	Sesquiterpenes	3.14
Hillyl acetate	Ester	3.10
β -Acoradienol	Sesquiterpenes	0.97
α -Cedrene epoxide- aroma chemical	Sesquiterpenes	0.62
Total detected (%)		100.00
Seed coat		
Eugenol acetate	Phenyl propanoids	40.71
β -Thujaplicinol	Monoterpenes	19.83
Torulol	Diterpenes	13.11
<i>trans</i> -Pinane	Monoterpenes	7.85
Pogostol	Sesquiterpenes	3.18
Gymnomitrol	Sesquiterpenes	3.05
Thujopsan-2- α -ol	Sesquiterpenes	2.85
Dill ether	Monoterpenes	2.62
3-Isopropyl-2-methoxypyrazine	Sesquiterpenes	1.79
Geranyl anthranilate	Ester	1.37
13- <i>epi</i> -Dolabradiene	Diterpenes	1.21
Zizanal	Sesquiterpenes	1.06
13- <i>epi</i> -Manool	Diterpenes	0.91
<i>Z</i> -Coniferyl alcohol	Lignan	0.44
Total detected (%)		99.98

concentration level), only 0.58 to 0.65% of treated filter paper was eaten for *S. longisperma* cotyledon and seed coat extracts respectively. This may be due to the presence of sesquiterpenes in *S. longisperma*. These compounds and their analogues can act as anti-feedents or insecticidal agents in *S. longisperma* cotyledon and seed coat as reported by Dalia (2011) in her study of *Casimiroa edulis* leaf extract against *Spodoptera littoralis* larvae. An abundance of eugenol acetate (40.71%) and

β -Thujaplicinol (19.83%) was observed in the seed coat of *S. longisperma*. Sesquiterpenes have been found to illicit behavioural responses in subterranean termites (Hassan et al. 2017). For example, caryophyllene is a major compound for effective insecticidal activity against *Aedes aegypti* (Sullamy et al. 2011), *Tribolium confusum* and *Callosobruchus maculatus* (Abbas & Tahere 2012) and stored product pests (*Oryzaephilus surinamensi* and *Trogoderma granarium*) (Janaki

Table 3 Major secondary metabolite components found in the extracts of *U. borneensis* cotyledon and seed coat

Compound	Class of compounds	Area (%)
Cotyledon		
2,4,6-Trimethoxytoluene	Sesquiterpenes	17.92
Syringaldehyde	Aldehyde	16.49
<i>Allo</i> -Cedrol	Sesquiterpenes	16.36
β -Cyclocitral	Sesquiterpenes	12.18
Presilphiperfolan-8-ol	Sesquiterpenes	9.92
Carvacrol	Monoterpenes	8.96
<i>Cis</i> -Vertocitral C	Monoterpenes	5.27
<i>neo</i> -Intermedeol	Sesquiterpenes	3.40
Sesquithuriferol	Sesquiterpenes	2.86
<i>trans</i> -Pinane	Monoterpenes	2.44
(<i>E,Z</i>)-2,6-Nonadienal diethyl acetal	Aldehyde	2.36
5 <i>E,9Z</i> Farnesyl acetone = ketones	Sesquiterpenes	1.83
Total detected (%)		99.99
Seed coat		
2'-Hydroxyacetophenone	Phenyl propanoids	19.01
Methyl p-tert-butylphenylacetate	Ester	14.84
<i>Is</i> o-Jasmone	Monoterpenes	13.02
Methyl linoleate	Alkaloid	9.04
Methyl hexadecanoate	Ester	8.51
Incensole oxide	Diterpenes	6.52
Methyl vanillin	Aldehyde	6.08
Hillyl acetate	Ester	5.43
Nonenal<dimethyl acetal-(3 <i>Z</i>)->	Aldehyde	4.64
6 <i>Z</i> , 10 <i>Z</i> Pseudo phytol	Sesquiterpenes	4.35
Ethyl hydroquinone	Sesquiterpenes	3.08
Cavacrol, methyl ether	Monoterpenes	2.40
4-Methoxy Stilbene	Monoterpenes	1.64
α -Alaskene	Sesquiterpenes	1.44
Total detected (%)		100.00

et al. 2018). Eugenol and eugenol acetate have toxicity against several insect pests, including termites (*Coptotermes formosanus*) (Cornelius & Lax 2005). The 24-hour LC₅₀ value for *U. borneensis* cotyledons extract was 4.71% and for seed coat, 4.81% (Table 4). The values for *S. longisperma* (3.01 and 3.27% respectively) were lower, and this was attributed to the higher sesquiterpenes content. The higher the total sesquiterpenes content, the higher the resistivity of the wood species.

Repellency test

Extract of *U. borneensis* seed coat showed strong repellent activities against *C. gestroi* at every

concentration tested compared with cotyledon (Table 5). Cotyledon extract was only 5% repellent at 0.5% extract concentration, while seed coat extract was seven times (36%) more active. Activity decreased to 33% for seed coat extract but continued to increase for cotyledon extract (28%) at 1.0% of concentration. As reported by Sbeghen et al. (2002), repellent activity is not dependent on the chemical dosage. Generally, both cotyledon and seed coat extracts of *U. borneensis* showed increased repellency with the concentration of extract used—repellent activity increased with increasing concentration of the extract tested. Hence, *U. borneensis* seed extract, alone or in combination with those obtained from other termite repellent

Table 4 Effect of extracts from *U. borneensis* and *S. longisperma* cotyledons and seed coats on paper consumption and feeding inhibition of *C. gestroi*

Treatment	Concentration (%)	Paper consumption (%)	Feeding inhibition (%) ¹	LC ₅₀ (%)
Control		4.36 (0.44) ^a		
Methanol		1.41 (0.30) ^d		
<i>U. borneensis</i> cotyledons	0.5	2.02 (0.18) ^b	51.43 (2.86) ^b	4.71
	1.0	2.00 (0.11) ^b	52.38 (1.65) ^b	
	2.0	1.55 (0.32) ^c	58.10 (4.36) ^a	
	3.0	1.48 (0.34) ^{cd}	59.05 (4.36) ^a	
	4.0	1.38 (0.23) ^d	60.95 (3.30) ^a	
Control		4.36 (0.44) ^a		
Methanol		1.41 (0.30) ^d		
<i>U. borneensis</i> seed coat	0.5	2.24 (0.15) ^b	45.71 (2.86) ^c	4.81
	1.0	2.23 (0.00) ^b	48.57 (0.00) ^c	
	2.0	1.86 (0.35) ^c	54.29 (4.95) ^b	
	3.0	1.83 (0.36) ^c	54.29 (4.95) ^b	
	4.0	1.37 (0.13) ^d	60.95 (1.65) ^a	
Control		4.36 (0.44) ^a		
Methanol		1.41 (0.30) ^c		
<i>S. longisperma</i> cotyledons	0.5	1.58 (0.17) ^c	54.29 (2.86) ^{cd}	3.01
	1.0	1.41 (0.28) ^c	57.14 (4.95) ^c	
	2.0	1.83 (0.97) ^b	52.38 (2.88) ^d	
	3.0	1.10 (0.55) ^d	62.86 (8.57) ^b	
	4.0	0.58 (0.12) ^e	68.20 (1.11) ^a	
Control		4.36 (0.44) ^a		
Methanol		1.41 (0.30) ^c		
<i>S. longisperma</i> seed coat	0.5	2.18 (0.25) ^b	49.52 (3.30) ^c	3.27
	1.0	1.39 (0.14) ^c	60.95 (1.65) ^b	
	2.0	1.20 (0.18) ^d	62.86 (2.86) ^b	
	3.0	0.75 (0.24) ^e	69.52 (3.30) ^a	
	4.0	0.65 (0.13) ^e	70.48 (1.65) ^a	

Values in brackets are standard deviations; percentage values followed by the same letter are not significantly different in the same group (percentage of extractive yields) at the 0.05 level of probability; n =5; ¹values represent the degree of difference between mass losses of the controls and the extractive tree. LC₅₀ = lethal concentration which causes a 50 % reduction in feeding as compared to the non-treated control

Table 5 Ability of seed coat and cotyledon extracts from *U. borneensis* and *S. longisperma* to repel *Coptotermis gestroi*

Species	Seed part	Repellence activity (%)					
		Control	0.5	1.0	2.0	3.0	4.0
<i>U. borneensis</i>	Cotyledons	21 (0.56) ^c	5 (0.22) ^d	28 (1.00) ^b	33(0.42) ^b	40(0.66) ^a	41(1.02) ^a
	Seed coat	21(0.82) ^c	36(0.87) ^d	33(1.22) ^d	40(0.82) ^c	53(0.11) ^b	63(0.49) ^a
<i>S. longispema</i>	Cotyledons	21(0.33) ^c	55(0.64) ^a	51(0.56) ^a	36(0.22) ^b	24(1.00) ^c	13(0.33) ^d
	Seed coat	21(0.62) ^c	32(0.11) ^a	24(0.49) ^b	24(0.33) ^b	16(0.56) ^c	15(0.88) ^c

Values in brackets are standard deviations; percentage values followed by the same letter are not significantly different in the same group (percentage of extractive yields) at the 0.05 level of probability; n =3

plant species could potentially be used for the preparation of termite repellent products.

Shorea longisperma seed extracts showed a reverse trend of repellency compared with *U. borneensis*. *Shorea longisperma* seeds showed less activity in repelling *C. gestroi* (Table 5). Extracts of both parts showed higher performances at 0.5% concentration (55% for cotyledon extract and 32% for seed coat extract) but attracted more termites when the concentration was increased. As discussed above, the effectiveness of the *S. longisperma* extract may be attributed to the presence of sesquiterpenes. The presence of monoterpenes other than sesquiterpenes in *S. longisperma* seed coat extracts also makes it more repellent against *C. gestroi*. Monoterpenes are highly efficient as inducers of mortality or repellents against insect species (Reis et al. 2016).

CONCLUSIONS

Upuna borneensis seeds contained higher levels of extractives compared with *S. longisperma* seeds. *Upuna borneensis* seed coat extract effectively repelled termites, but did not cause high termite mortality. Cotyledon extract of *S. longisperma* attracted and killed termites even at the lowest concentration. Hence, extract of *S. longisperma* cotyledon could potentially be used as wood preservative. It was effective against termites even at the lowest concentration compared with the rest of the samples (*S. longisperma* seed coat, *U. borneensis* cotyledon and *U. borneensis* seed coat). Evaluation of efficiency, stability and production costs of seed coat and cotyledon extracts as wood preservatives need to be investigated.

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