Quinonoid constituents as contact sensitisers in Australian blackwood (Acacia melanoxylon RBR)*

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ABSTRACT Australian blackwood (Acacia melanoxylon RBR) is a valuable commercial timber that since 1925 has been incriminated as being injurious to health. In addition to toxic effects numerous cases of allergic contact dermatitis and bronchial asthma have been observed in woodworkers. Several constituents have been identified in recent years, but none of them could be considered as aetiological factors. Sensitising experiments performed with blackwood heartwood extracts corroborated the described sensitising properties. Chemical studies showed the occurrence of two or possibly three quinones that produced positive skin reactions in the sensitised guinea pigs. The main contact allergens were isolated and identified by x-ray analysis. The first, a yellow quinone, was identified as 2,6-dimethoxy-1,4-benzoquinone while the second, a red quinone, has the structure of 6-methoxy-2-methyl-3,5-dihydrobenzofurano-4,7-dion and was named acamelin. Whereas 2,6-dimethoxy-p-benzoquinone is already known from natural sources, acamelin is new and belongs to the rare group of naturally occurring furanoquinones. Though the obtained sensitising capacity of A melanoxylon RBR in respect of its quinones is not high, it should be considered as a possible source of allergic contact dermatitis, especially as greater amounts of Australian blackwood may enter European countries in the near future.

Australian blackwood (Acacia melanoxylon RBR) belongs to the family of Leguminosae-Mimosaceae and is one of at least 100 different wood species of commercial value that are capable of inducing allergy of the cell-mediated type (delayed type of allergy, allergic contact dermatitis) in men handling it.1 A melanoxylon RBR is native to Western Australia. Owing to its dark brown colour and fine and decorative appearance it is used in Australia not only for high quality furniture, panelling, joinery, turnery, and shop and bank fittings, but also for bent work in coach- and boat-building, for handles, and even for parts of musical instruments.2

For decades Australian blackwood has also been cultivated in East and South Africa and considerable amounts are exported to Britain. As it is intended to introduce this timber to other European countries in the near future, we have investigated this species for sensitising constituents.

Contact dermatitis

Cases of contact dermatitis as well as outbreaks of bronchial asthma after the handling and inhalation of fine wood dust and shavings have been described since 1925. Mainly workers in joineries and motor-boat factories have been affected, developing itching, weeping dermatitis and circumscribed skin lesions on the forearms, neck, and face.3 4 Nott5 suffered from a papular rash on his hands and forearms after handling the wood at his home. Robertson6 described 10 cases, one of which had bronchial asthma but nine had contact dermatitis. Though his patch tests remained negative, Robertson suggested that sensitisation had played a part. Further cases without detailed descriptions of the observations have been mentioned by Behl et al,7 Cleland,8 Hurst,9 and two anonymous authors.10 11 Recently three new cases of allergic contact derma-
titis were reported by Burry. He stated that Australian blackwood is specially mentioned in the Workmen's Compensation Act, 1965, of South Australia. This Act implies that any dermatitis occurring in a patient in contact with blackwood is considered to be the result of exposure to blackwood until proved otherwise.

In addition to Australian blackwood, some other related species of the genus *Acacia* may cause dermatitis and asthma—for instance, *A. cyperophylla* F MUELLER (Red mulga),* A. shirleyi* MAIDEN (Lancewood), and *A. harpophylla* F MUELLER (Brigalow). Among these wood species, Brigalow is the most notorious, causing what is known as "Brigalow itch."15

**Previous chemical investigations**

In addition to the observed skin irritant and sensitising properties of *Acacia* species there is evidence that several species are toxic to stock and man. At least 20 species, native to Australia or cultivated or growing wildly in New Zealand, have been examined for the occurrence of saponins. Large amounts of alkaloid-like substances such as β-phenyl ethylamine and tryptamine were isolated from most of them, including *A. melanoxylon* RBR. Phytochemical studies on the heartwood constituents of Australian and African species of *Acacia* have been performed by Tindale and Roux and others, yielding evidence for the wide distribution of flavonoids in this genus. From *A. melanoxylon* RBR higher amounts of (-)-melacacidin and (-)-isomelacacidin were isolated.

Although these constituents throw some light on the chemotaxonomic and phylogenetic aspects of *Acacia* species, none of the compounds detected so far can seriously be referred to as contact sensitisers.

**Material and methods**

To prove the sensitising capacity of heartwood extracts of *A. melanoxylon* RBR and to isolate their aetiologically active constituents, sensitising experiments were carried out in guinea pigs, and chemical studies were performed by column and thin-layer chromatography (TLC).

Different samples of *A. melanoxylon* RBR were kindly supplied by:

1. Pieces of heartwood and sapwood and branches from (a) 8 km E of Sandy Hill on the Bruxner highway, New South Wales; (b) 82 km ENE of Armidale on the Dorrigo road, NSW, 30°22' S, 152°30' E; and (c) Swamp Creek, 20 km NE of Tenterfield, NSW (all by Dr Tindale).
2. Pieces of heartwood from a timber merchant, Geelong, Victoria (Mr Lindemann).
3. Pieces of heartwood from a timber merchant in Sidney (Dr Taylor).
4. Pieces of heartwood grown in South Africa (Mr Gottwald).

Altogether 1600 g of wood samples were available and have been used for sensitising experiments, patch tests, and chemical investigations.

**OPEN EPICUTANEOUS SENSITISATION PROCEDURE**

Ten female albino guinea pigs of the Pirbright white-strain were sensitised with a blackwood extract, made by extracting the heartwood shavings for 24 hours with pure ethanol at room temperature. The residue left after evaporation of the solvent at reduced pressure (water pump vacuum) and

<table>
<thead>
<tr>
<th>No of animals</th>
<th>Sensitised with</th>
<th>Challenged with</th>
<th>Concentration (%)</th>
<th>Challenge after 24 hours</th>
<th>Challenge after 48 hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>10</td>
<td>Ethanol extract</td>
<td>Ethanol extract</td>
<td>10</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>10</td>
<td>Quinone 1</td>
<td>Quinone 1</td>
<td>1</td>
<td>--</td>
<td>2</td>
</tr>
<tr>
<td>5</td>
<td>Quinone 2</td>
<td>Quinone 2</td>
<td>1</td>
<td>--</td>
<td>2</td>
</tr>
<tr>
<td>5</td>
<td>Quinone 3</td>
<td>Quinone 3</td>
<td>1</td>
<td>--</td>
<td>2</td>
</tr>
<tr>
<td>5</td>
<td>Quinone 4</td>
<td>Quinone 4</td>
<td>1</td>
<td>--</td>
<td>2</td>
</tr>
</tbody>
</table>

**Table 1 Results of sensitising experiments and challenges with chromatographically separated fractions and purified constituents of blackwood ethanol extracts**

Evaluation of the intensity of skin reactions:

<table>
<thead>
<tr>
<th>No</th>
<th>Reaction</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>No reaction.</td>
</tr>
<tr>
<td>1</td>
<td>Slight erythema (visible and distinguishable from a negative reaction).</td>
</tr>
<tr>
<td>2</td>
<td>Distinct erythema restricted to the test area.</td>
</tr>
<tr>
<td>3</td>
<td>Intense confluent erythema and infiltration, slight swelling, but restricted to the test area.</td>
</tr>
<tr>
<td>4</td>
<td>Intense erythema, infiltration and exudation with swelling and spreading over the test area.</td>
</tr>
</tbody>
</table>
low temperature (39°C) was dissolved in acetone to a 10% solution. Each day 0.05 ml of this crude extract was applied to a 2 cm² test area on the clipped and shaved flank of the animals. Nine days after beginning this treatment a slight erythema developed that increased steadily until it reached a maximum of a ++ + + reaction on day 17. The treatment was then stopped.

Two weeks after the sensitisation procedure, challenge was carried out by epicutaneous application of 0.05 ml of different concentrations of the crude ethanol extract on the opposite flank of the guinea pigs. Readings of the test area after 24 and 48 hours indicated that all animals had been sensitised (table 1).

CHEMICAL EXPERIMENTS

Separation of the ethanol extract in different solvent systems by analytical TLC showed several quinonoid compounds in the heartwood. This was shown by spraying the thin-layer plates with Craven solution.²⁹ The Craven test, a highly specific reagent for benzoquinones and naphthoquinones, requires the presence of a labile hydrogen or halogen atom adjacent to the carbonyl group of the quinoid ring.* Extracts containing such quinones show blue or green spots after separation on thin-layer plates.

Subsequent extraction of the wood shavings with cold ethyl acetate and later with hot ethyl acetate (60°C) yielded further quantities of the detected quinones.

The crude separation of all the heartwood extracts on silica gel columns was followed by further purification steps on preparative TLC-plates using at first a one-dimensional method with chloroform—methanol (100 + 2) (solvent system I). Four fractions were obtained, of which two contained the enriched quinones. In later experiments separation of the quinone-enriched fractions in two dimensions was preferred, using solvent system I in one direction and benzene-acetone (100 + 5) in the other.

The purified quinones and the remaining fractions were used to challenge the sensitised guinea pigs. The positive test responses made clear that the quinones had to be considered the responsible aetiological agents (table 1). Chemical examination of the sapwood and bark were followed up in the same manner. Spraying of the separated bark extract with Craven solution gave no specific colour reaction, whereas the sapwood showed low amounts of the yellow quinone.

*Equal amounts of absolute ethanol and ammonia (25%) are mixed with three to five drops of ethyl cyanoacetate.

Results

The results obtained in the guinea pig experiments with the crude extracts of the heartwood showed a medium sensitising capacity for Australian blackwood. In the material supplied from the Royal Botanic Gardens and National Herbarium of Sydney (Dr Tindale) traces of a third quinone were detectable by the Craven test in addition to the two main quinones. Wood samples supplied later from other sources (Dr Taylor, Mr Gottwald) contained only the two main contact allergens.

After final purification two quinones were isolated:

I  A yellow quinone; 8.6 mg, recrystallised from ether-ethyl acetate, mp: dec 250-252°C.
II A red quinone; 15.8 mg, recrystallised from acetone, mp: 175-176°C (uncorr).

Both quinones were the only constituents to give positive skin responses in the sensitised guinea pigs. During elicitation the red quinone showed a slightly stronger and longer lasting reaction than the yellow quinone (table 1).

Elucidation of the structures was carried out by x-ray analysis. The yellow quinone was identified as 2,6-dimethoxy-1,4-benzoquinone (fig 1). The red quinone could be identified as a new benzofuranquinone that we named acamelin (fig 1). Detailed information on the procedures for elucidation of their structures was given in two preliminary short communications.²³ ²⁴

Discussion

Australian blackwood may affect health in several ways: in addition to the toxic effects allergic dermatitis and bronchial asthma may occur in people handling the wood and inhaling the wood shavings. The toxic properties are probably due to the alkaloid-like constituents occurring in varying concentrations in this species. The constituents responsible for allergic contact dermatitis in wood-workers are two quinones found in the heartwood. The causative agents of the asthmatic reactions remain unidentified.

![Figure 1: Quinonoid contact allergens of Acacia melanoxyylon RBR (Australian blackwood).](http://oem.bmj.com/)

[Fig 1] Quinonoid contact allergens of Acacia melanoxyylon RBR (Australian blackwood).
The sensitising potency of the wood and its quinonoid constituents was shown in guinea pig sensitisation experiments. Though the wood samples were obtained from different sources (Australia, Africa), it is obvious that these two quinones are the main contact sensitisers. It must be remembered, however, that blackwood trees from different localities may yield varying amounts of these constituents and, under certain circumstances, may contain a third quinone. Whether this quinone plays a part in blackwood sensitivity is undetermined.

The simple yellow 2,6-dimethoxy-1,4-benzoquinone has previously been discovered and isolated from 25 different wood and plant species and was recently detected in an additional 21 species. Its sensitising capacity has been proved in guinea pig sensitisation experiments and was compared with that of 10 related naturally occurring quinones (table 2). The results showed a slight sensitising power that indicates in single cases of plant and wood dermatitis 2,6-dimethoxy-p-benzoquinone should be recognised as a possible sensitisier.

Acamelin, the red quinonoid allergen of Australian blackwood is a 6-methoxy-2-methyl-3,5-di-hydrobenzo-(b)furano-4,7-dion. Naturally occurring furanoquinones are rare. Four similar but naphtho-furanoquinones were isolated and identified in 1968 from Peroba wood Paratecoma peroba KUHLM (family: Bignoniaceae) (fig 2). Although these quinones have not been examined with respect to their sensitising properties, it is known that Peroba wood is also responsible for allergic contact dermatitis. The present findings show once more that quinonoid constituents of plants and woods are of considerable and ever increasing dermatological interest. Numerous benzo- and naphtho-quinones have been isolated from natural sources in the past two decades and their sensitising capacity has been confirmed in animal experiments. Furthermore, recent research has shown that cross-reactivities exist between different but chemically related benzo- and naphtho-quinones. This knowledge is important in view of the possible recurrence of allergic contact dermatitis, which may develop in individuals who are sensitive to distinct plant or wood quinones. In those patients who acquired their quinone-sensitivity occupationally or through a hobby, recurrent dermatitis may develop after contact with other quinone-containing plants, wood species, or natural products, such as wooden shoes (clogs), knife-handles, recorders, and wooden jewellery by accident or changing a working place.

Table 2 Composition of sensitising capacity of 2,6-dimethoxy-1,4-benzoquinone with related naturally occurring quinones obtained by guinea pig maximisation sensitisation experiments

<table>
<thead>
<tr>
<th>Quinone</th>
<th>Occurrence</th>
<th>Name</th>
<th>Obtained sensitising capacity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thymoquinone</td>
<td>Calocedrus decurrens (TORR) FLORIN</td>
<td>Incense cedar</td>
<td>Strong</td>
</tr>
<tr>
<td>Obtusquinone</td>
<td>Dalbergia retusa HEMSLEY</td>
<td>Cocobolo</td>
<td></td>
</tr>
<tr>
<td>Macassar quinone*</td>
<td>Diospyros celebica BAKH</td>
<td>Macassar ebony</td>
<td>Medium</td>
</tr>
<tr>
<td>2,6-dimethoxy-1,4-benzoquinone</td>
<td>Acacia melanoxylon RBR</td>
<td>Australian blackwood</td>
<td></td>
</tr>
<tr>
<td>Toluquinone</td>
<td>Brachiaus crepitans L</td>
<td>Anthrodopes</td>
<td></td>
</tr>
<tr>
<td>Lapachol</td>
<td>Tectona grandis L</td>
<td>Teakwood</td>
<td></td>
</tr>
<tr>
<td>Rapanone</td>
<td>Ardista macrocarpa WALLICH</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Quinone Q1</td>
<td>Diospyros melanoxylon GILL &amp; PERR</td>
<td>Coromandel ebony</td>
<td>Negative</td>
</tr>
<tr>
<td>2,5-dimethoxy-1,4-benzoquinone</td>
<td>Diospyros melanoxylon GILL &amp; PERR</td>
<td>Coromandel ebony</td>
<td></td>
</tr>
</tbody>
</table>

*Occurs in the wood as the precursor macassar II.

Fig 2 Naturally occurring furanoquinones from peroba wood (Paratecoma peroba KUHLM).
References


