

**UNIVERSITI TEKNOLOGI MARA**

**PHYTOCHEMICAL STUDIES AND  
BIOACTIVITIES OF *XYLOPIA FERRUGINEA*  
HOOK. (ANNONACEAE) AND *SHOREA*  
*MAXWELLIANA* KING.  
(DIPTEROCARPACEAE)**

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Thesis submitted in fulfilment of the requirements  
for the degree of  
**Master of Science**


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## AUTHOR'S DECLARATION

I declare that the work in this thesis was carried out in accordance with the regulations of Universiti Teknologi MARA. It is original and is the result of my own work, unless otherwise indicated or acknowledged as referenced work. This thesis has not been submitted to any other academic institution or non-academic institution for any degree or qualification.

I, hereby, acknowledge that I have been supplied with the Academic Rules and Regulations for Post Graduate, Universiti Teknologi MARA, regulating the conduct of my study and research.

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## ABSTRACT

In the present work, phytochemical studies were conducted on two species from different family. The studied plant samples were the stem bark and leaves of *Xylopia ferruginea*; and the stem bark of *Shorea maxwelliana*. Phytochemical investigation on the crude alkaloidal extracts of the stem bark and leaves of this plant yielded five oxoaporphinoid alkaloids and three flavonols, characterised as *O*-methylmoschatoline, lysicamine, liriodenine, atherospermidine, oxostephanine, quercetin, kaempferol and afzelin. Phytochemical investigation carried out on the stem bark of *Shorea maxwelliana* King has yielded five oligostilbenoids identified as  $\alpha$ -viniferin, maximol A, vaticanol A, suffruticosol A and vaticanol G. Structure elucidation and characterization of isolated compounds were done based on 1D, 2D NMR, IR, UV, MS, optical rotation, melting point and in good agreement with those previously reported for these compounds. In addition, the antibacterial activity of major isolated compounds was tested against seven bacteria. Our findings showed that *O*-methylmoschatoline is the most active compound as it inhibited the growth of all tested bacteria and the highest inhibition were shown in *S. aureus* (20 mm) and *S. epidermis* (23 mm). Vaticanol G showed moderate and slight activity against *S. aureus* and *E. coli*, while maximol A exhibited slight activity against *E. coli* and *P. aeruginosa*. Maximol A, suffruticosol A, vaticanol A and vaticanol G possessed active cytotoxic activity against HL60 cell line with IC<sub>50</sub> values range of 10.1 - 220  $\mu$ M. Only maximol A was active against A549 and MCF-7 cell lines, with both IC<sub>50</sub> value of 220 and 99.2  $\mu$ M.

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