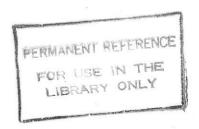
SURA

# CHEMISTRY AND SEARCH FOR ANTIVIRAL /HIV ACTIVITY OF SOME SRI LANKAN CALOPHYLLUM SPECIES

A thesis presented by

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in partial fulfilment of the requirement for the award of



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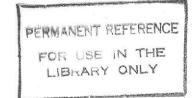
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#### **ABSTRACT**



In this study chemistry and antiviral / HIV activity of *Calophyllum* products were investigated. The root bark of *Calophyllum thwaitesii* has been shown to contain seven xanthones, calozeylanic acid, friedelin and sitosterol. Two of the xanthones have been identified as demethylcalabaxanthone and trapezifolixanthone, two new xanthones as 3,10,11,12-tetrahydro-13-hydroxydipyranoxanthen-14-one [11,12-dihydrothwaitesixanthone] and 6,8-dihydroxy-2,2-dimethyl-7,9-di(3-methylbut-2-enyl)-2H,5H-pyrano(3,2-a)xanthene-5-one [batukinaxanthone]. The other three xanthones are thwaitesi xanthone, calothwaitesixanthone and 6-deoxy-γ-mangostin, which have previously reported from the same species. This is the first report on the isolation of a 1,3,5-trioxygenated xanthone [trapzifolixanthone] from this plant. Stem bark of *Calophyllum cordato-oblongum* gave two coumarins cordatolide B, soulattrolide and cordato-oblongic acid. This is the first report on the isolation of a 4-phenylpyrano coumarin derivative, soulattrolide from this plant.

MeOH extracts of *C. bracteatum, C. cordato-oblongum, C. soulattri* and *C. thwaitesii* inhibited the activity of one of the aspartic proteinase enzymes pepsin-A. Further, bioassay guided fractionation of above extracts showed that water fractions of *C. moonii* and *C. cordato-oblongum* were protective against pepsin A. Above results indicated that further investigation of active fractions is very essential with the view to isolating antiviral / HIV compounds from this genus. Above coumarins are structurally very close to the anti HIV coumarins isolated from the same genus. Therefore antiviral / HIV studies were carried out on them. Preliminary tests showed no activity and this may be due to the solubility problem regarding pure coumarins in aqueous phase.