

The structure and sites of biochemical interaction of cotton defensive proteins and secondary metabolites

CRDC number: ULA5C

Principal investigator:

Dr Roland Chung

Present address: Victorian College of Pharmacy, Department of Medicinal Chemistry, Monash University, 381 Royal Parade, Parkville, Victoria 3052

Telephone: (03) 9903 9000

Fax: (03) 9903 9582

Sponsor/team leader/co-investigator:

Dr Gideon Polya

School of Biochemistry, La Trobe University, Bundoora, Victoria, 3083

Telephone: (03) 9479 2157

Fax: (03) 9479 2467

e-mail: g.polya@latrobe.edu.au

Plain English summary:

A variety of proteins were purified from cotton seeds and were analysed by amino acid sequence determination (automated Edman degradation) and by state-of-the-art electrospray ionization mass spectrometry (ESMS). The amino acid sequences of about 40 vicilin-related cotton proteins were precisely defined. The molecular masses of the deduced sequences were in agreement with the masses observed by ESMS to within about 1-2 Da (i.e. to within the mass of 1-2 hydrogen atoms). A further major cotton seed protein was purified and shown to be a γ -conglutin-related protein with 2 component subunits and a molecular mass of 46250.3 ± 1.3 Da. The vicilin fractions and the γ -conglutin variously have anti-fungal activity against a wide range of fungi tested including the cotton pathogens *Fusarium oxysporum* and *Verticilium dahliae*. This precise work is potentially useful for classical and transgenic approaches to cotton plant breeding for pest resistance. Cotton leaves also contain anti-fungal components and a defensive component with larvicidal activity against the blowfly *Lucilia cuprina* and the mosquito *Aedes camptorhynchus*. Large-scale purification procedures were developed to enable further biological testing. Protein and non-protein fractions from cotton are variously active as inhibitors of animal cyclic AMP-dependent protein kinase and of the proteases trypsin and chymotrypsin.