

ought to be measured in biological fluids rather than the major urinary metabolite alone before conclusions can be drawn whether a drug inhibits only prostaglandin biosynthesis or catabolism or both. We are presently exploring this more complete approach towards the effect of drugs on the PG system with several other compounds which we have found to inhibit prostaglandin catabolism^{20, 21}.

²⁰ We thank Drs. J. E. PIKE and U. AXEN, The Upjohn Co., Kalamazoo, U.S.A. for their generous supplies of prostaglandins.

²¹ Supported by a grant No. MA-4181 to C.P.A. from the Medical Research Council of Canada.

Zusammenfassung. Es zeigt sich, dass jedes der 3 Enzyme, die den Abbau von Prostaglandin in den Nieren erwachsener Ratten verursachen, durch schwache Konzentrationen von Indomethacin in steigendem Ausmass inhibiert werden können: 9-PGDH > 13-PGR > 15-PGDH.

C. PACE-ASCIAK and S. COLE

*Research Institute, The Hospital for Sick Children,
555 University Avenue, Toronto M5G 1X8
(Canada), 9 September 1974.*

Identification of Precoccinellin in the Ladybird Beetle, *Coleomegilla maculata*¹⁻³

Coccinellid beetles have long been known to possess defensive compounds associated with 'reflex bleeding'⁴, and the investigation of several European lady bugs has yielded the structures of the specific alkaloids⁵⁻⁷. Likewise, during an investigation of the volatiles of the lady beetle, *Coleomegilla maculata*, we have isolated and identified a defensive alkaloid, precoccinellin.

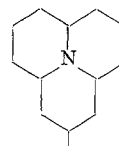
Isolation procedures. The beetles were steam-distilled for 2 h in an all-glass system, and the distillate was extracted with methylene chloride. The extract was concentrated in vacuo and chromatographed by GLC on a 20 ft, 1/8 inch O.D. stainless steel column packed with 10% SE-30. Column temperature was 160°C, and carrier gas pressure at the inlet was 60 ψ . Retention time of the alkaloid was 10 min. $I_b = 1475$. The compound was collected from a stream splitter attachment on the gas chromatograph. *IR-spectrum.* The IR-spectrum in CCl₄ included ν_{max} 1020, 1040, 1120, 1130, 1325, 1385, 1450, 2775, 2860, 2925, 2950 CM⁻¹.

PMR-spectrum. The PMR-spectrum in CCl₄ showed ppm (δ) 0.89, 1.24, 1.38, 1.55, 1.64.

Mass spectrum m/e. 41 (100), 192 (66), 150 (51), 151 (47), 55 (47), 42 (44), 137 (34), 136 (32), 164 (28), 67 (26), 122 (25), 93 (22), 82 (19), 178 (18), 108 (15), 96 (13); M⁺ = 193. Essentially the same as the precoccinellin isolated by TURSCH et al.⁵

Reactions. Hydrogenation (Pd on charcoal; hydrogen under pressure) – no effect. Lithium aluminum hydride reduction – no effect. NaOH treatment – no effect. HCl treatment – hydrochloride formed.

Based on these experiments and on data presented by TURSCH et al.⁵, we propose that the compound isolated from *C. maculata* is precoccinellin:



(Dodecahydro-2-methylpyrido[2,1,6-de]quinolizine)

Extraction of whole beetles with methanol revealed no additional alkaloids. The compound is bitter to the taste, and it is assumed that it performs in a defensive mode as do other similar coccinellid alkaloids.

Zusammenfassung. Isolierung und Strukturzuteilung für ein Alkaloid aus *Coleomegilla maculata* (Coleoptera: Curculionidae).

R. D. HENSON⁸, A. C. THOMPSON⁸, P. A. HEDIN⁸,
P. R. NICHOLS⁹ and W. W. NEEL⁹

*United States Department of Agriculture, Agricultural
Research Service, Boll Weevil Research Laboratory,
P. O. Box 5367, Mississippi State
(Mississippi 39762, USA), 1 August 1974.*

¹ Coleoptera: Curculionidae.

² In cooperation with the Mississippi Agricultural and Forestry Experiment Station, Mississippi State, Mississippi 39762. Received for publication.

³ Mention of a proprietary product does not necessarily imply endorsement of this product by the USDA.

⁴ R. M. HAPP and T. EISNER, *Science* 134, 329 (1961).

⁵ B. TURSCH, D. DALOZE, M. DUPONT, J. M. PASTEELS and MARIE-CLAIRE TRICOT, *Experientia* 27, 1380 (1971).

⁶ B. TURSCH, D. DALOZE, M. DUPONT, C. HOUTELE, M. KAISIN, J. M. PASTEELS, and D. ZIMMERMANN, *Chimia* 25, 307 (1971).

⁷ R. KARLSSON and D. LOSMAN, *J. chem. Soc., Chem. Commun.* 11, 626 (1972).

⁸ Boll Weevil Research Laboratory, USDA, ARS, Mississippi State, Mississippi 39762, USA.

⁹ Department of Entomology, Mississippi State University, Mississippi State, Mississippi 39762, USA.

Microheterogeneity of Staphylococcal Enterotoxin C₂

Staphylococcal enterotoxin C₂ belongs to a closely related group of simple proteins that are causative agents of staphylococcal food poisoning. The toxin has been purified to homogeneity, and several of its physical-chemical properties have been determined¹. We also

isolated enterotoxin C₂ in a homogeneous form as judged by gel filtration, immunodiffusion, ultracentrifugation, and N-terminal amino acid analysis (to be published elsewhere). However, disc-gel electrophoresis showed 2 bands that formed an immunoprecipitate with specific