value and the heat of formation of crystalline polyoxymethylene  $(-40.93 \text{ kcal per CH}_20 \text{ unit})$  obtained by Parks and Mosher<sup>10</sup>. These yield the value  $-23.2 \text{ kcal mole}^{-1}$  for the heat of formation of formaldehyde. It is believed that this figure is more reliable than the presently accepted 'best' value for this quantity.

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## REFERENCES

<sup>1</sup> NORDGREN, G. Acta Path. Bicrobiol. scand. (suppl.), 1939, 40, 21

<sup>2</sup> NIELSEN, H. H. and EBERS, E. S. J. chem. Phys. 1937, 5, 824

<sup>3</sup> DAINTON, F. S., IVIN, K. J. and WALMSLEY, D. A. G. Trans. Faraday Soc. 1959, 55, 61

<sup>4</sup> WALKER, J. F. Formaldehyde, 3rd ed., p 180. Reinhold: New York, 1964

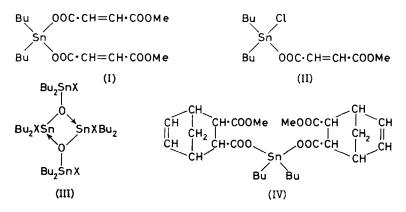
<sup>5</sup> IWASA, Y. and IMOTO, T. J. chem. Soc. Japan, Pure Chem. Sect. 1963, 84, 29

- <sup>6</sup> THOMPSON, J. B. (See WALKER, J. F.) Formaldehyde, 3rd ed., p 180. Reinhold: New York, 1964
- <sup>7</sup> DWORJANYN, L. O. Austral. J. Chem. 1960, 13, 175
- <sup>8</sup> DAINTON, F. S., EVANS, D. M., HOARE, F. E. and MELIA, T. P. Polymer, Lond. 1962, 3, 263
- <sup>9</sup> VON WARTENBERG, H. and LERNER-STEINBERG, B. Angew. Chem. 1925, 38, 591

<sup>10</sup> PARKS, G. S. and MOSHER, H. P. J. Polym. Sci. A, 1963, 1, 1979

## The Stabilization of Poly(vinyl chloride) by Organotin Maleates

ORGANOTIN derivatives of maleic acid are widely used to inhibit degradation of poly(vinyl chloride) by heat and light<sup>1</sup>. It has been suggested<sup>1</sup> that these compounds undergo Diels-Alder addition reactions with diene systems in the partially degraded polymer chains. There is no direct evidence for this and many of the stabilizers have been only vaguely described in the patent literature and are of uncertain composition.



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In recent work<sup>2</sup> on the mechanism of stabilization of poly(vinyl chloride) dibutyltinbis(methyl maleate) (I), prepared<sup>3</sup> from dibutyltin dichloride and potassium methyl maleate, was used but no details of this compound were reported.

An attempt to repeat this preparation gave dibutylchlorotin methyl maleate (II), m.pt 41° to 43°C as the sole product. Compound I was, however, prepared from dibutyltin oxide and methyl hydrogen maleate and isolated as an oil  $n_D^{23}$ 1·4930 which could not be distilled without decomposition. An attempt to convert I to its 2,2'-bipyridyl derivative using the aqueous ethanolic conditions described by Huber *et al.*<sup>4</sup> caused hydrolysis to (III, X=OOC·CH=CH·COOMe) m.pt 91° to 94°C (similar compounds where X=CH<sub>3</sub>·COO are well known<sup>5</sup>). Compound I interacted with dienes such as 2,3-dimethylbuta-1,3-diene but the products were gums and difficult to purify. However, with cyclopentadiene the crystalline Diels-Alder adduct IV m.pt 56° to 58°C was obtained.

Further studies on the mechanism of stabilization of poly(vinyl chloride) by organotin compounds are in progress.

(All the compounds described had satisfactory elemental analyses and infra-red spectra; compound III had molecular weight (found) 1 450, (required) 1 480.)

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## REFERENCES

<sup>1</sup>CHEVASSUS, F. and DE BROUTELLES, R. The Stabilization of Polyvinyl Chloride. Edward Arnold: London, 1963

<sup>2</sup> FRYE, A. H., HORST, R. W. and PALIOBAGIS, M. A. J. Polym. Sci. A, 1964, 2, 1765, 1785, 1801

<sup>3</sup> FRYE, A. H. and HORST, R. W. J. appl. Radiation and Isotopes, 1964, 15, 169

<sup>4</sup> HUBER, F., ENDERS, M. and KAISER, R. Z. Naturf. 1966, 21b, 83

<sup>5</sup> POLLER, R. C. J. Organometal. Chem. 1965, 3, 321