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Takamine Research Laboratory, Sankyo Co., Ltd., Nishi-shinagawa, Shinagawa-ku, Tokyo. Takahiro Nakamura (中村隆洋)

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Decarboxylation of the Permanganate Oxidation Product of Lyoniresinol Dimethyl Ether

In a previous paper¹¹¹ it was reported that lyoniresinol dimethyl ether (I) gave a carboxylic acid (II), $C_{20}H_{22}O_9 \cdot 2\frac{1}{2}H_2O$, m.p. $186.7 \sim 187^\circ$,* by oxidation with potassium permanganate in pyridine and that the molecular formula of (II) agreed with that of galloylgallic acid hexamethyl ether.

A neutral substance (III), m.p. $129.8 \sim 130^{\circ}$ (Anal. Calcd. for $C_{19}H_{22}O_7$: C, 62.97; H, 6.12; O, 30.91. Found: C, 62.71; H, 6.20; O, 31,14. IR ν_{CO}^{KBr} 1660 cm⁻¹), was obtained from (II) by decarboxylation with copper bronze in quinoline and the substance (III) was confirmed as 2,3,4,3',4',5'-hexamethoxybenzophenone by comparison with an authentic sample, m.p. 129.9° (Anal. Calcd. for $C_{19}H_{22}O_7$: C, 62.97; H, 6.12; O, 30.91. Found: C, 62.78; H, 6.25; O, 30.80. IR ν_{CO}^{KBr} 1660 cm⁻¹), by elemental analysis, mixed melting point determination, and from infrared absorption spectra.

The authentic sample of 2,3,4,3',4',5'-hexamethoxybenzophenone²) was prepared by methylation of the condensation product (2-hydroxy-3,4,3',4',5'-pentamethoxybenzophenone) of pyrogallol trimethyl ether and galloyl chloride trimethyl ether.

Thus, it was proved that the two trimethylpyrogallol rings were joined to one another with one carbon atom between them.

Pharmaceutical School, Nagoya City University, Hagiyama-cho, Mizuho-ku, Nagoya. Masaichi Yasue (安江政一) Yoshishige Kato (加藤義成)

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^{*} All m.p.s were determined by micro-method and are uncorrected.

¹⁾ M. Yasue, Y. Kato: Yakugaku Zasshi, 80, 1013(1960).

²⁾ W.H. Perkin, C. Weizmann: J. Chem. Soc., 89, 1665(1906).