## Derivatives of Tetramethylplatinum

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IN a recent paper it is stated that tetramethylplatinum(IV) is nonexistent (private communications by H. C. Brown and by L. F. Dahl to E. L. Amma).<sup>1</sup> This has prompted us to report that we have made some stable derivatives of tetramethylplatinum(IV) of type [PtMe<sub>4</sub>L<sub>2</sub>] with  $L = AsMe_2Ph$ , PMe<sub>2</sub>Ph, or PEt<sub>3</sub>. platinum(II) and platinum(IV) that two equivalent methyl groups in *trans*-position to the phosphines give a complex resonance pattern of the type recently reported for cis-[PtMe<sub>2</sub>(PPh<sub>3</sub>)<sub>2</sub>]<sup>2</sup> except that our compounds, being much more soluble, give better spectra than that published.

Pyrolysis of [PtMe<sub>4</sub>(PMe<sub>2</sub>Ph)<sub>2</sub>] at 160° gives

TABLE

Methyl resonance data for some tetramethylplatinum(IV) complexes measured at 60 Mc./sec. and 34°

		${ m Me}^{f a}$			Meb			Methyls on L		
[PtMe4(AsMe2Ph)2]							~ <b>^</b>			
	Solvent C <sub>6</sub> H <sub>6</sub>	au9·66	Jp	Јр <b>і</b> 44	au 8·83	Jр	J р <b>t</b> 66	au 8·95	Jр	Jр <b>і</b> 6-5
$[PtMe_4(PMe_2Ph)_2]$	C <sub>6</sub> H <sub>6</sub> CHCl <sub>3</sub>	$9.80 \\ 10.23$	$6{\cdot}5 \\ 6{\cdot}4$	43∙5 44	$9.08 \\ 9.61$	*	58 57	8.83 8.60	8·5 8·4	$11.5 \\ 11.5$
$[PtMe_4(PEt_3)_2]$	$C_{6}H_{6}$	9.72	5.8	44	t	t	t	$\operatorname{complex}$		

\* Complex resonance pattern, see discussion; † Obscured by overlap with other resonances.

Treatment of cis-[PtCl<sub>2</sub>(AsMe<sub>2</sub>Ph)<sub>2</sub>] with gives methyl-lithium cis-[PtMe<sub>2</sub>(AsMe<sub>2</sub>Ph)<sub>2</sub>]. This reacts with methyl bromide to give [PtBrMe<sub>3</sub>(AsMe<sub>2</sub>Ph)<sub>2</sub>], configuration (I), which reacts with methyl-lithium to give cis-[PtMe4-(AsMe<sub>2</sub>Ph<sub>2</sub>)<sub>2</sub>], configuration (II). The configuration (II) follows from the dipole moment of 5.4 D and also from the n.m.r. (methyl) resonance pattern (Table). We have similarly made the complex cis-[PtMe4(PMe2Ph)2] and also cis- $[PtMe_4(PEt_3)_2]$  by treating  $[PtCl_2Me_2(PEt_3)_2]$ with methyl-lithium. In these complexes containing phosphine the two mutually trans-methyl groups (platinum bonded) give a simple n.m.r. pattern but we have generally found for both

ca. quantitative yields of ethane and cis-[PtMe<sub>2</sub>(PMe<sub>2</sub>Ph)<sub>2</sub>].

We have also made many complexes of the types [PtMe<sub>x</sub>X<sub>4-x</sub>L<sub>2</sub>] with x = 1-3, X = Cl, Br, I, and L = tertiary phosphine or tertiary arsine.



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<sup>1</sup> R. G. Vranka and E. L. Amma, J. Amer. Chem. Soc., 1967, 89, 3121.

<sup>2</sup> E. O. Greaves, R. Bruce, and P. M. Maitlis, Chem. Comm., 1967, 860.