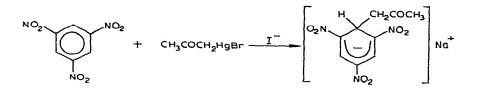
## **Preliminary communication**

## Jackson-Meisenheimer complexes in reactions of organomercury compounds with trinitrobenzene

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We have found that organomercury compounds,  $R_2$  Hg and RHgX, with electronwithdrawing groups react with trinitrobenzene (TNB) under conditions of iodide ion catalysis and that Jackson-Meisenheimer complexes are formed in this reaction which is a novel reaction in the organomercury series. We have also shown that  $(CH_3COCH_2)_2$  Hg and  $CH_3COCH_2$  HgBr form Janowsky complexes<sup>1</sup> with TNB in the presence of I<sup>-</sup> in deuteroacetone or dimethylsulfoxide ( $\tau$  8.37, singlet; 5.10, triplet; 2.60, doublet)



Organomercury compounds,  $R_2$  Hg and RHgX, which have electron—withdrawing substituents  $[R = C_6F_5, PhC \equiv C, (CF_3)_2$  CH,  $C_6H_5$  CHCOOC<sub>2</sub>H<sub>5</sub>, o-carboranyl] react, while  $(C_6H_5)_2$  Hg and Alk<sub>2</sub>Hg do not react with TNB under these conditions. We think that the first step of the reaction is ionisation of the organomercury compound accompanied by C-Hg bond rupture<sup>2</sup> [ $S_E 1(I^-)$ ], with subsequent interaction of the kryptocarbanion formed with TNB, the latter step probably involves a single electron transfer. Studies of the reaction mechanism are in progress.

## REFERENCES

- 1 M.J. Strauss, Chem. Rev., 70 (1970) 667.
- 2 LP. Beletskaya, K.P. Butin and O.A. Reutov, Organometal. Chem. Rev., 7 (1971) 51.