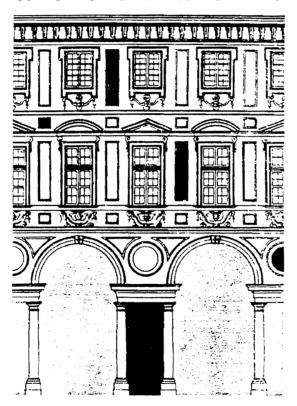
## $\begin{array}{c} \textbf{XIII I} \textbf{International } \textbf{C} \textbf{onference} \\ \textbf{on Organometallic Chemistry} \end{array}$



## abstracts



Torino, September 4-9, 1988

## METAL ALKOXIDE MODIFIED ORGANOMETALLIC REACTIONS. DIRECT SYNTHESIS OF LITHIOOXYALKYL- AND ARYLITHIUM COMPOUNDS

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Greece

Attempts to synthesize  $\mathrm{LiO(CH_2)}_6\mathrm{Li}$  according to the conventional method from  $\mathrm{HO(CH_2)}_6\mathrm{Br}$  led to extensive Wurtz coupling. Similar results were obtained by employing lithium naphthalene radical anion in place of lithium metal.

6-Lithiooxyhexyllithium was prepared in fair yields according to eq. (1).

The organometallic product was characterized by carbonation and by conversion to  $\mathrm{Hg}(\mathrm{CH}_2\mathrm{CH}_2\mathrm{CH}_2\mathrm{CH}_2\mathrm{CH}_2\mathrm{CH}_2\mathrm{OH})_2$ . The function of the alkoxide is to stabilize the organometallic reagent with respect to THF-cleaving reaction, so that solutions of the reagent can be stored at room temperature for 48 hours or longer.

The method has been extended to the preparation of the lithio- oxy- organolithiums of the general types 1 and 2 .

Li(
$$CH_2$$
)<sub>n</sub>-Y-( $CH_2$ )<sub>m</sub>OLi

Y=0,NR

1

METAL ALKONIDE MODIFIED OPGANOMETALLIC REACTIONS. DIRECT SYNTHESIS

OF LITHICONYAUKYD- AND ARYLITHIUM COMPOUNDS

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The method has been extended to the preparation of the lithio- oxy- organolithiums of the general types  ${\bf 1}$  and  ${\bf 2}$  .

Li(
$$CH_2$$
)<sub>n</sub>-Y-( $CH_2$ )<sub>m</sub>OLi  
Y=O,NR

1

O( $CH_2$ )<sub>n</sub>OLi

Li