

THF- $d_8$  solution over a 30–60 min period). However, a  $^{13}\text{C}$  NMR spectrum of **2** in the same solvent at room temperature could not be obtained because of rapid decomposition of the sample. Therefore, a low-temperature study ( $-30\text{ }^\circ\text{C}$ ) of a precooled THF- $d_8$  solution was undertaken in the case of **2**.

A strong low-field shift for the *ipso* carbon atom signal (a narrow signal at 195.6 ppm) was observed in the  $^{13}\text{C}$  NMR spectrum of a THF- $d_8$  solution of **1**, while the low-temperature  $^{13}\text{C}$  NMR spectrum of **2** showed a broad and poorly resolved signal at 201 ppm for the *ipso* carbon atom. It was previously shown<sup>5</sup> that  $^{13}\text{C}$  chemical shifts of *ipso* carbon atoms of both monomeric and dimeric solvated lithium aryls are in the range 180–200 ppm. We note that the solution structures of **1** and **2** are not clear. However, our  $^{13}\text{C}$  solution NMR spectroscopic investigations indicate that the asymmetric arrangement found in the solid-state structure of complex **1**, exhibiting different coordination modes for the two lithium atoms, is not retained in THF solution.

### Conclusion

Our results provide more insight into the solid-state structures of two differently aggregated terphenyl-

lithium complexes and demonstrate that the arrangement found in the solid-state structure of THF adducts of terphenyllithium compounds is mostly determined by the nature of the sterically demanding terphenyl ligand. Lower aggregation numbers can be induced by increased crowding at the aryl group, resulting in formation of monomeric aryllithium species: e.g., complex **2**. Our work also allows a detailed comparison of the structural data of **1** and **2** with previously reported solvated and unsolvated terphenyllithium salts.

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**Supporting Information Available:** X-ray crystallographic files in CIF format for the structures of  $[\text{DppLi}]_2(\text{THF})$  (**1**) and  $\text{DnpLi}(\text{THF})_2$  (**2**). This material is available free of charge via the Internet at <http://pubs.acs.org>.

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## Additions and Corrections

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**Sung-Joon Kim, Namkeun Yang, Dae-Hyun Kim, Sang Ook Kang, and Jaejung Ko\*:** New Types of Base-Stabilized Alkyl Aluminum, Gallium, and Indium Complexes.

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