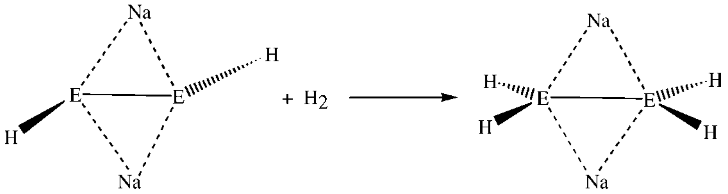
Reaction enthalpies are calculated for the hydrogenation reactions of main group hydrides with the potential for multiple bonding, and thus the *unsaturated* character of these species was determined. The study includes species such as $[HGaGaH]^{2-}$ and $Na_2[HGaGaH]$ (see scheme, E = B, Al, or Ga).



Supporting information on the WWW (see article for access details).

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All the Tables of Contents from 1998 onwards may be found on the WWW under **http://www.wiley-vch.de/home/chemistry/**

CORRIGENDA

In the paper by K. C. Nicolaou et al. published in *Chem. Eur. J.* **2001**, *7*, 5359–5371, the enantiomeric structures of the Diels–Alder components in the transition states $\mathbf{TS_a}$ and $\mathbf{TS_b}$ are required to account for the observed enantiomers **8**, (−)-**5**, **8′**, and **5′** shown in Scheme 9.

In the paper by J. Jiang, N. Kobayashi, and D. K. P. Ng et al. in *Chem. Eur. J.* **2001**, *7*, 5059–5069, the true space group for all the double-deckers $\mathbf{4}\cdot 0.5\,C_6H_{12}$, **6**, $\mathbf{9}\cdot 0.5\,C_6H_{12}$, and **14** is *Pnam* (No. 62; interchange of the *b* and *c* axes as reported in the paper leads to the standard symbol *Pnma*). The $[M^{III}(nc)(oep)]$ molecule lies on a crystallographic mirror plane that contains the lanthanide atom, two *trans*-related isoindole N donor atoms of the nc ring, and a pair of *trans*-related *meso* CH groups on the oep ligand. In the complexes $\mathbf{4}\cdot 0.5\,C_6H_{12}$ and $\mathbf{9}\cdot 0.5\,C_6H_{12}$, the chair-shaped cyclohexane molecule has half-site occupancy with two of CH_2 groups lying in a mirror plane. The authors apologize for these errors.