LANTHANOID CARBIDE-HALIDES - THE 09.5-12 BORDERLINE FOR METAL-METAL BONDING. BY Hj.Mattausch, <u>C.Schwarz</u>, A.Simon, Max-Planck Institut für Festkörperforschung, D-7000 Stuttgart-80, FRG

The lanthanoids form various metal-rich hal-The lanthanoids form various metal-rich hal-ides. With few exceptions, such as Gd_2X_3 , Tb_2X_3 ; X=Cl,Br, they are most commonly found as ternary compounds with nonmetals like H, B, C, and N. Especially large is the number of car-bide-halides, of which those compounds contain-ing gadolinium were examined in more detail.

The structurally characterized phases are listed in Table 1 according to their "extent of condensation". In this classification scheme lies the fundamental concept that these structures are constructed from edge-sharing $Gd_{6X_{12}}$ clusters, in which their centers are occupied by C atoms or C₂ units. Thus, a strong struc-tural correlation with the metal-rich transition metal compounds, e.g. ZrCl, ZrBr, is obtained. When the number of electrons available for me-tal-metal bonding in clusters of the d-elements is too small, especially for Zr, (e.g. $Zr_6I_{12}X$ with X=B,C; $Zr_6I_{14}C$), one finds nonmetal atoms inserted in these clusters. The number of electrons associated with the interstitial nonmetal atoms increases, while the number of bonding states in the cluster remains constant. The formalism described here is not a usual one for chemical bonds. Extensive charge transfer from metal to nonmetal occurs especially with the electropositive lanthanoid compounds. Therefore, these structures are alternatively described as a defect NaCl variant: in the usual close-packed arrangement of X and C^{4-} or C_2^{-4} ions, the Gd³⁺ ions occupy the octahedral c2. holes around the more highly charged C species. Using the ionic model, the geometrial details of these structures can be simply explained, (e.g. the shifting of the C atoms from the cen-(e.g. the shirting of the C atoms from the cen-ters of the octahedra, the distortion of the octahedra, etc.). Very importantly, a predic-tion of the C_n species is possible from the electron balance with a knowledge of the compo-sition: so in $(Gd^{3+})_{10}(Cl^{-})_{18}(C2^{-})_{24}$ there is a C-C single bond, in $(Gd^{3+})_{24}(Cl^{-})_{24}(C2^{-})$, a C-C double bond. In some compounds up to 3 valence These can partially occupy bonds with metal-tal bonding character, as is well known for $Gd_{10}C_{18}C_4$. There now results a simple picture of their chemical bonds: the strong heteropolar Gd-C and Gd-X interactions are supplemented by the relatively weak metal-metal bonds.

Table 1

Edge sharing d_{Gd-Gd} C-Octahedra [pm] Species

Gd ₁₀ Cl ₁₈ C ₄		two		321-409	c2 ⁶⁻
Gd10Cl17C4		two		312-401	c2 ⁶⁻
Gd ₁₀ I ₁₆ C ₄		two		328-400	c2 ⁶⁻
$Gd_{12}X_{17}C_6(X=Br,I)$	1	-chain		319-427	c2 ⁶⁻
Gd;I5C	1	-chain		333-398	c4-
Gd ₆ X ₇ C ₂ (X=Br,I)	1 &	-double	chain	339-395	c ^{4 –}
Gd3I3C	1	-double	chain	331-391	¢4-
Gd ₂ X ₂ C ₂ (X=C1,Br,I) <u>2</u>	-layers		345-400	c24-
Gd ₂ Br ₂ C	2	-layers		343-382	c4-
Gdécl ⁵ C ³	28	-layers		340-389	c÷-
Gd ₂ IC	2 %	-layers		338-380	c4-
Gd3Cl3C	3	-nets		329-368	c4-

STRUCTURES WITH OLIGOMERIC CLUSTERS 09.5-13 IN THE INDIUMOXOMOLYBDATES: In₁₁Mo₄₀O₆₂, In₃Mo₁₁O₁₇. By <u>Hi.Mattausch</u>, A.Simon, Max-Planck-Institut für Festkörperforschung, 7000 Stuttgart-80, FRG

Infinite chains of Mo_6 -octahedra condensed via opposite edges are present in the structure of NaMo₄O₆ (C.C.Torardi, R.E.McCarley, J.Am.Chem. Soc. 1979, <u>101</u>, 3963). The Mo₄O₆ chain can be interrupted by structural disorder in the cryster tal.

tal. $In_{11}Mo_{40}O_{62}$ is the first compound (Hj.Mat-tausch, A.Simon, E.M.Peters, Inorg. Chem. 1986, <u>25</u>, 3428), where discrete cluster anions with four and five condensed Mo₆-octahedra are ar-ranged in layers, which are stacked in an al-ternating sequence (fig. 1a). A single oligome-ric cluster of five condensed octahedra is rhour in figure 1b. All free edges are bridged shown in figure 1b. All free edges are bridged by O-atoms. The channels between the cluster by O-atoms. The channels between the cluster anions are occupied by linear M-M-bonded poly-cations In_5 ⁺⁻ and In_6 ⁺⁻. The layers seen in fig. la have the general composition $In_{n+1}Mo_{4n+2}O_{5n+4}$, where n indicates the number of MO₆ octahedra in the oligomeric cluster. The number of Mo-Mo-bonding states can be estimated by suitable fragmentation of the oligomeric by suitable fragmentation of the oligomeric cluster into $Mo_6O_{12}+(n-1)Mo_4O_6$ units (A.Simon in A.K.Cheetham and P.Day (Eds.) Inorganic solids, Oxford University Press, in print). Accordingly, one expects, besides "homogeneous structures" (i.e. structures with a single type of layers), also structures with clusters of other length. A first confirmation of their ovistore way provided by bich resolution existence was provided by high resolution transmission electron microscopy (A.Simon, W.Mertin, Hj.Mattausch, R.Gruehn, Angew. Chem. 1986, <u>98</u>, 831). An electron microscopic photo-graph of an $In_{11}Mo_{40}O_{62}$ crystal fragment shows the alternating Sequence of the $(Mo_6)_4$ and $(Mo_6)_5$ -layers. Such alternating layer segments occur in ordered regions of 10° pm extension. In the same sample regions are found in which stacking disorder is recognizable: two consecu-tive In Mo. On layers are followed by two imstacking disorder is recognizable: two consecutive $In_5Mo_{18}O_{28}$ layers are followed by two immediately adjacent $In_6Mo_{22}O_{34}$ layers. Another part of the same sample is exclusively made up of identical $In_6Mo_{22}O_{34}$ layers over distances of 10° pm. In the meantime, the respective new compound $In_3Mo_{21}O_{17}$ has been isolated as a horgeneous phase and have been characterized by X-ray crystallography (a= 3160.8, b= 948.9, c= 983.9 pm).



(ь)

a: Projection of the crystal structure of Fig. $In_{11}Mo_{10}O_{62}$ on [010]. The oligometric clusters $(Mo_6)_4$ and $(Mo_6)_5$ are arranged in layers (small circles Mo-atoms; large circles with crosses 0-atoms)

b: Oligomeric cluster with five edgelinked Mo₆ octahedra in In₁₁Mo₁₀O₆₂.