

## Aluminium – Boron – Magnesium

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### Literature Data

Grain refining of boron additions to Al–Mg alloys was the basis for early studies of the Al–B–Mg system [1949Ebo]. No complete phase diagram exists, although several groups investigated the interactions in the ternary system [1959Hof, 1970Mat, 1971Vek, 1981Por, 1983Sho]. Considerable interest was further devoted to the rather hard compound “MgAlB<sub>14</sub>” [1970Mat, 1971Vek, 1983Hig, 1990Hig, 1993Hig]. Single crystals were obtained from aluminum high temperature flux starting from the nominal composition MgAl<sub>31</sub>B<sub>6</sub> which was heated under argon in an alumina crucible to 1500°C, kept at this temperature for 1h and slowly cooled to RT (10K·min<sup>−1</sup>). The excess Al was then dissolved in hot HCl [1983Hig, 1993Hig]. From a batch with ~100 g, crystals were formed up to 5 mm in size and predominantly as plates with the habit {001} [1993Hig]. When starting mixtures with smaller amounts of Mg were used, γAlB<sub>12</sub> type crystals Mg<sub>0.45</sub>Al<sub>0.77</sub>B<sub>12</sub> were obtained, mostly as thin hexagonal plates [1993Hig].

[1959Hof] mixed magnesium, aluminum and boron in order to study the formation of (Al,Mg)B<sub>2</sub> solid solutions. After sintering in argon at temperatures between 725 and 790°C the formation of such a solid solution was identified despite the kinetic difficulty of formation. From 99.9 % pure metals and 99.2 % pure amorphous boron [1971Vek] synthesized 22 ternary alloys. The mixtures were briquetted and heated to temperatures of 850 to 1000°C under an argon atmosphere in sintered alumina crucibles. The reaction products were investigated by X-ray phase analysis. [1970Mat] prepared “MgAlB<sub>14</sub>” by heating a mixture of magnesium, aluminum and boron in atomic proportions of 1:2:14 to 900°C for 6 h. The sample was then cooled and treated with concentrated hydrochloric acid. The crystal structure of this phase was determined by X-ray (probably on single crystal) diffraction.

The present evaluation was published in the MSIT Evaluation Program earlier and reflects today’s state of knowledge.

### Binary Systems

The systems Al–B [1994Dus] and B–Mg [1978Spe] are accepted.

### Solid Phases

According to [1971Vek] the phase “MgAlB<sub>12</sub>” is identical to “MgAlB<sub>14</sub>”. The structure of which was determined in detail from several attempts to obtain single crystal material from high temperature aluminum solutions [1983Hig, 1990Hig, 1993Hig]. In agreement with the chemical analysis (atom emission spectroscopy, Mg<sub>0.79</sub>Al<sub>0.80</sub>B<sub>14</sub>), X-ray single crystal studies revealed significant defects on the metal sites: Mg<sub>0.78</sub>Al<sub>0.75</sub>B<sub>14</sub> [1983Hig, 1990Hig, 1993Hig]. For smaller Mg-concentrations in the Al-melt crystals “Mg<sub>0.5</sub>Al<sub>1.4</sub>B<sub>22</sub>” of the γAlB<sub>12</sub> type were obtained [1990Hig, 1993Hig], suggesting a high temperature solid solution of Mg in γAlB<sub>12</sub> (about 3.5 at.% Mg at ~1500°C).

### Isothermal Sections

A tentative isothermal section at approximately 900°C (Fig. 1) is constructed from the results of [1971Vek]. [1971Vek] reported a region of (Al) solid solution containing more than 65 at.% B, which is quite improbable and might be the result of the difficult formation of the (Al,Mg)B<sub>2</sub> phase as stated by [1959Hof]. For the phase relations at 900°C a small solid solution of Mg in αAlB<sub>12</sub> was assumed as well as for Al in (βB). The ternary compound was taken at the composition Mg<sub>0.78</sub>Al<sub>0.75</sub>B<sub>14</sub> [1983Hig, 1990Hig, 1993Hig].

### Notes on Materials Properties and Applications

Microhardness (Vickers hardness under a load of 100g) for  $\text{Mg}_{0.78}\text{Al}_{0.75}\text{B}_{14}$  crystals was found to range from 27.70 to 28.90 GPa [1993Hig].

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**Table 1:** Crystallographic Data of Solid Phases

Phase/ Temperature Range [°C]	Pearson Symbol/ Space Group/ Prototype	Lattice Parameters [pm]	Comments/ References
(Al) < 660.452	<i>cF4</i> <i>Fm<math>\bar{3}m</math></i> Cu	$a = 404.96$	[Mas2]
( $\beta$ B) < 2092	<i>hR333</i> <i>R<math>\bar{3}m</math></i> $\beta$ B	$a = 1093.30$ $c = 2382.52$ $a = 1096.5$ $c = 2386.8$	[1993Wer]  at AlB <sub>31</sub> [V-C2]
Al <sub>2</sub> B <sub>3</sub> ≤ 525	<i>hR*</i> Al <sub>2</sub> B <sub>3</sub> (?)	$a = 1840$ $c = 896$	at 60 at.% B [1992Var]
$\alpha$ AlB <sub>12</sub> ≤ 2050	<i>tP216</i> <i>P4<sub>1</sub>2<sub>1</sub>2</i> $\alpha$ AlB <sub>12</sub>	$a = 1015.8$ $c = 1427.0$ $a = 1018$ $c = 1434.3$	[1994Dus] $\rho_{\text{exp.}} = 2.65 \text{ Mgm}^{-3}$ [1991Pri]
$\gamma$ AlB <sub>12</sub>	<i>oP384</i> <i>P2<sub>1</sub>2<sub>1</sub>2<sub>1</sub></i> $\gamma$ AlB <sub>12</sub>	$a = 1014.4$ $b = 1657.3$ $c = 1751.0$  $a = 1019.5$ $b = 1666$ $c = 1769$	[1994Dus] metastable phase or ternary product stabilized by small amounts of impurity metals present in Al-flux grown material $\rho_{\text{exp.}} = 2.56 \text{ Mgm}^{-3}$ [1991Pri]
Mg <sub>0.25</sub> Al <sub>0.77</sub> B <sub>12</sub>		$a = 1018.7$ $b = 1663.3$ $c = 1754.7$	solid solution of Mg in $\gamma$ AlB <sub>12</sub> [1990Hig, 1993Hig]
(Al <sub>1-x</sub> Mg <sub>x</sub> )B <sub>2</sub> AlB <sub>2</sub> ≤ 975	<i>hP3</i> <i>P6/mmm</i> AlB <sub>2</sub>	$a = 300.6$ $b = 325.2$ $a = 304.7$ $c = 336.6$	$0 < x < 1$ [1959Hof, 1971Vek] at $x = 0$ [1994Dus]  at $x = 0.5$ [1971Vek]
MgB <sub>2</sub> ≤ 1550(BP)		$a = 308.5$ $b = 352.3$	at $x = 1$ [V-C2]
MgB <sub>4</sub> ≤ 1775 (BP)	<i>oP20</i> <i>Pnma</i> MgB <sub>4</sub>	$a = 546.4$ $b = 442.8$ $c = 747.2$	[V-C2]
MgB <sub>7</sub> ≤ 2150 (BP)	<i>oI64</i> <i>Imma</i> MgB <sub>7</sub>	$a = 597.0$ $b = 1048.0$ $c = 812.5$	[V-C2]
* $\tau_1$ , Mg <sub>0.78</sub> Al <sub>0.75</sub> B <sub>14</sub>	<i>oI68</i> <i>Imma</i> MgAlB <sub>14</sub>	$a = 584.8$ $b = 1031.2$ $c = 811.2$	[1970Mat, 1983Hig, 1993Hig]

**Fig. 1: Al-B-Mg.**  
Tentative partial iso-thermal section at about 900°C

