

B n)

B (north, Kermadec,
Hawaiian & Tonga. p. 72)

1980

93: 211164q A critical evaluation of the thermodynamic data for boron ions, ion pairs, complexes, and polyanions in aqueous solution at 298.15 K and 1 bar. Bassett, R. L. (US Geol. Surv., Denver, CO 80225 USA). *Geochim. Cosmochim. Acta* 1980, 44(8), 1151-60 (Eng). A polemic. Thermodyn. data

m. g. u. h.
c. b. - b. c.

for B ions, ions pairs, complexes, and polyanions in aq. soln. are critically evaluated.

C. A. 1980, 93 v 22

B⁺

1984

12 Л98. Возобновленное изучение однократно ионизованного бора. A renewed study of singly ionized boron. Bashkin S., McIntyre L. C., Von Buttlar H., Ekberg J. O., Martinson I. «Nucl. Instrum. and Meth. Phys. Res.», 1985, B9, № 4: Phys. Highl Ionised Atoms. Proc. Int. Conf., Oxford, 2—5 July, 1984, 593—597 (англ.)

С помощью классической эмиссионной спектроскопии, а также при возбуждении пучково-пленочным методом выполнено новое исследование спектра В II. Предложены новые идентификации спектра, приведшие к пересмотру энергий уровней $2s3s^1S$ и $2s5p^1P^o$. Установлены также значения уровней $2p3s^1P^o$ и $2s6p^1P^o$. Измерены времена жизни для уровней $2s2p^1P^o$, $2p^2^1S$, $2s3d^1D$ и $2p^2^3P$. Найдены силы осцилляторов переходов $2s^2^1S—2s2p^1P^o$, $2s2p^1S—2p^2^1S$ и $2s2p^3P^o—2p^2^3P$. Найденные значения находятся в хорошем согласии с теоретич. величинами.

А. Н. Рябцев

ε, спектр

ср. 1985, 18, N 12

$B_2(P)$

(DM. 24028)

1986

Lilov S.K.,

K_p , $\Delta_f H$;

J. Phys. and Chem.
Solids, 1986, 47,

N 3, 245-250.

B₁₂

1991

Kleinman Leonard.

Boron-Rich Solids: Pap. 10th

Int. Symp. Boron. Borides,
and Relat. Compounds, Al-
buquerque, N. M. 1990. New
York (N. Y.), 1991. C. 13-20.

ΔH_f ,
pccrēm

(see BC_x; I)

B50

1992

116: 266135c Computational search for the real tetragonal boron (B_{50}). Lee, Seungbok; Bylander, D. M.; Kim, Suck Whan; Kleinman, Leonard (Dep. Phys., Univ. Texas, Austin, TX 78712 USA). *Phys. Rev. B: Condens. Matter* 1992, 45(7), 3248-51 (Eng). Using an expansion of $\sim 12,700$ plane waves, the lattice consts. and heat of formation of tetragonal B_{50} , $B_{48}C_2$, $(B_{50}C_2)_n$, $(B_{50}C_2)_2$, $(B_{52}C_2)_n$, and $(B_{52}C_2)_2$ were calcd. Of these only $(B_{50}C_2)_2$ has both a and c within 1% of their x-ray values as well as having, by far, the least neg. heat of formation.

(fraction $\Delta_f H$)

(+) $B_{48}C_2$ u gp

C.A. 1992, 116, N26

B₂³⁺

1992

116: 224106r Molecular ion stability and populations in tandem accelerator mass spectrometry. Matteson, S.; Weathers, D. L.; Kim, Y. D.; Arrale, A. M.; McDaniel, F. D.; Duggan, J. L.; Anthony, J. M.; Douglas, M. A. (Cent. Mater. Charact., Univ. North Texas, Denton, TX 76203 USA). *Nucl. Instrum. Methods Phys. Res., Sect. B* 1992, B64(1-4), 330-5 (Eng). The success of tandem accelerator mass spectrometry (AMS) rests, in part, on the dissociation of interfering mol. species in high charge states. Previous work in this lab. has detected persistent mols. of B₂³⁺. MO calcns. suggest that the mol. ion is bound in an excited state, but unbound in the ground state. Other binary homonuclear and heteronuclear ions have been examd. theor., and AlO³⁺ appears to be bound in the ground state. In the present work, exptl. detn. of the relative abundance, dissociation cross section in the stripper gas and the charge fraction branching ratios for the at. fragments of B₂ are presented. These results permit an assessment of the expected background levels to be encountered in the application of AMS to stable isobar mass spectrometry.

(масс-спектр.)

⊗ AlO³⁺

C. A. 1992, 116, N 22