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Laser spectroscopy of iridium monoboride

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High resolution laser induced fluorescence spectrum of IrB in the spectral region between 545 and 610 nm has been recorded and analyzed. Reacting laser-ablated iridium atoms with 1% B₂H₆ seeded in argon produced the IrB molecule. This is the first experimental observation of the IrB molecule. Four vibronic transition bands, (*v*, 0) with *v*=0–3 of an electronic transition system, have been observed. Spectra of all four isotopic molecules, ¹⁹¹Ir¹⁰B, ¹⁹³Ir¹⁰B, ¹⁹¹Ir¹¹B, and ¹⁹³Ir¹¹B, were recorded. Isotopic relationships confirmed the carrier of the spectra and the vibrational quantum number assignment. Preliminary analysis of rotational lines showed that these vibronic bands are with $\Omega' = 2$ and $\Omega'' = 3$. The electronic transition identified is assigned as the [16.5]³Π₂–X³Δ₃ system. Partially resolved hyperfine structure which conforms to the Hund's case *a*_β coupling scheme has been observed and analyzed. The bond length *r*₀ of the lower X³Δ₃ state of IrB was determined to be 1.7675 Å. © 2008 American Institute of Physics. [DOI: 10.1063/1.2901964]

I. INTRODUCTION

Metal borides are refractory compounds of considerable interest in thin film coating because of their attractive surface properties and exceptional corrosion resistance.¹ Many transition metal borides are known to be good catalysts for hydrogenation of alkenes and alkynes, reduction of nitrogenous functional groups, and deoxygenation reactions.² In addition, it is recently known that solid magnesium diboride (MgB₂) is a superconductor at very low temperature.³ Despite these important characteristics, experimental study of metal boride compounds is still very limited. In the past few years, some efforts have been made to determine the spectroscopic properties of diatomic metal borides in gas-phase experimentally^{4,5} and also to understand their bonding characteristics theoretically.^{6–10} Iridium is also an important metal catalyst in the formation of carbon-hydrogen and carbon-oxygen bonds.^{11,12} Iridium compounds' catalytic properties in chemical reactions are well known, but their spectroscopic properties are not readily available. Gas-phase diatomic iridium compounds with one main group element such as IrO (Refs. 13 and 14), IrN (Refs. 15–17), and IrC (Refs. 17–19) are the only ones that were characterized spectroscopically.

A very effective means of studying the bonding nature of small molecules is from an analysis of the optical spectra recorded at sufficiently high resolution to resolve not only the rotational structure but also the spin-fine and hyperfine structures. Rotational structure gives details on the bond length and, if available, the first lines of the rotational branches help to confirm the assignment of electronic state.²⁰ The spin-fine and hyperfine structures provide information concerning electron occupation in different molecular orbitals (MOs). The magnetic hyperfine parameters measured ex-

perimentally can be related to averages of various spin densities, of the unpaired valence electrons at the nuclei.^{21,22}

In this paper, we report the analysis of an electronic transition system of the IrB molecule recorded using the technique of laser vaporization/reaction free jet expansion and laser induced fluorescence (LIF) spectroscopy. The electronic transition identified is assigned as the [16.5]³Π₂–X³Δ₃ system. The four IrB isotopic molecules are ¹⁹¹Ir¹⁰B, ¹⁹³Ir¹⁰B, ¹⁹¹Ir¹¹B, and ¹⁹³Ir¹¹B and their relative natural abundance are, respectively, 1.0, 1.68, 4.02, and 6.77. In our study, the spectra of ¹⁹¹Ir¹¹B and ¹⁹³Ir¹¹B were recorded with good intensity; however, we could not resolve Ir¹⁰B isotope transition lines except in the (0, 0) band. The (2, 0) band was found to be perturbed. Furthermore, hyperfine interaction arising from unpaired electrons and the magnetic moment of the boron atom with nuclear spin of *I*=3/2 have also been observed and analyzed.

II. EXPERIMENT

The electronic transition spectrum of IrB was obtained using a laser vaporization/reaction free jet expansion LIF spectrometer. Descriptions of the spectrometer and the laser ablation system were given in earlier publication.²³ Only a brief description of the relevant experimental conditions for obtaining the IrB spectrum is given here. Pulses of laser radiation at 532 nm, 5 mJ, and 10 ns from a Nd:YAG (yttrium aluminum garnet) laser were focused onto the surface of a iridium rod to generate iridium atoms. A pulsed valve, synchronized with appropriate delay time, released a gas mixture of 1% B₂H₆ in argon to react with the iridium atom to produce IrB. The Nd:YAG-pulsed valve system was operated at 10 Hz. A diode-pumped solid-state laser with 6 W output power and at 532 nm was used to pump the ring dye laser operating with R560 and R6G dyes in the visible region, which was used to excite the jet cooled IrB molecules. LIF signal was collected by a lens system and detected by a photomultiplier tube. Linewidths of about 250 MHz were

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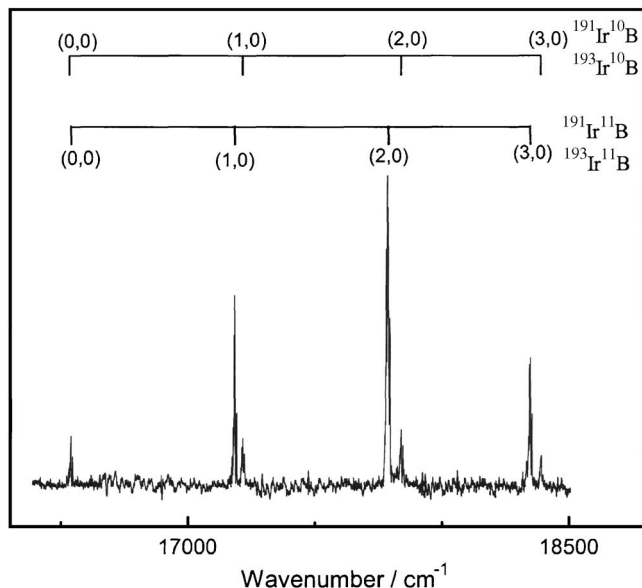


FIG. 1. Low-resolution LIF spectrum of the $[16.5]^3\Pi_2-X^3\Delta_3$ band system of IrB.

obtained. The wavelength of the dye laser was measured using a wavemeter with an accuracy of one part in 10^7 . The absolute accuracy of the measured line position is about $\pm 0.002 \text{ cm}^{-1}$.

III. RESULTS AND DISCUSSION

A. Low-resolution broadband spectrum

Low-resolution LIF spectrum of IrB in the visible region between $16\,000$ and $18\,500 \text{ cm}^{-1}$ has been recorded. Figure 1 shows a broadband low-resolution scan of the IrB spectrum. There are two vibronic sequences belonging to the two isotopes: Ir¹⁰B and Ir¹¹B. With the exception of the (0, 0) band, we were able to resolve the transition bands due to the ¹⁹¹Ir and ¹⁹³Ir isotopes. It is easily noticed that the (2, 0) band is a stronger transition. We have recorded high-resolution spectrum of the (ν , 0) bands with $\nu=0-3$.

B. Assignment of the observed system

Each of the recorded bands at high resolution shows resolved P , Q , and R branches. Line assignment was straightforward. The (0, 0) band head was observed at $16\,547 \text{ cm}^{-1}$,

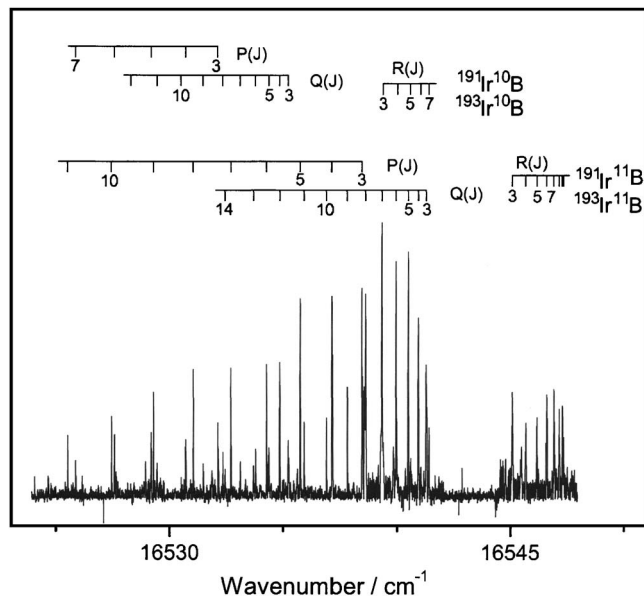


FIG. 2. The (0, 0) band of the $[16.5]^3\Pi_2-X^3\Delta_3$ transition of IrB.

which is shown in Fig. 2. We were able to resolve molecular transition lines with ¹⁰B and ¹¹B atoms; the lines belonging to the Ir¹⁰B isotopes are much weaker in intensity than those of the Ir¹¹B. However, transition lines from the two iridium isotopes, ¹⁹¹Ir and ¹⁹³Ir, were not resolved in the (0, 0) band. The observation of the first lines, $P(3)$, $Q(3)$, and $R(3)$ lines, established that this is a $\Omega'=2$ and $\Omega''=3$ transition. The vibrational quantum number assignment was confirmed by examining the isotopic shift between the Ir¹⁰B and Ir¹¹B. In addition, the R branch is relatively weaker than the P and Q branches, which is consistent with a $\Delta\Lambda=-1$ transition. Since only low J lines ($J < 17$) were observed, the line positions of this band were fitted to the following expression:

$$\nu_o = T_o + B'J'(J'+1) - D'[J'(J'+1)]^2 - \{B''J''(J''+1) - D''[J''(J''+1)]^2\}, \quad (1)$$

where the ' and '' refer to the upper and low states, respectively. Isotopic molecular transition lines of Ir¹⁰B and Ir¹¹B were measured and fitted. The centrifugal constant D was fixed at the calculated values using the Kratzer relation.²⁰ Determined molecular constants of individual bands, equilibrium molecular constants, and bond lengths are given in

TABLE I. Molecular constants for the upper $[16.5]^3\Pi_2$ and the $X^3\Delta_3$ states of IrB (cm^{-1}).

State	Parameter	¹⁹¹ Ir ¹¹ B	¹⁹³ Ir ¹¹ B	¹⁹¹ Ir ¹⁰ B	¹⁹³ Ir ¹⁰ B
$[16.5]^3\Pi_2$	T_3	18 350.286(2)	18 349.943(2)		
	B_3	0.472 35(3)	0.472 04(3)		
	T_2	17 791.129(2)	17 790.787(2)		
	B_2	0.474 55(2)	0.474 24(3)		
	T_1	17 186.970(2)	17 186.679(2)		
	B_1	0.473 21(2)	0.472 94(2)		
	T_0	16 541.845(1)	16 541.845(1)	16 535.824(2)	16 535.824(2)
	B_0	0.473 36(4)	0.473 36(4)	0.518 60(2)	0.518 60(2)
$X^3\Delta_3$	T_0	0.000	0.000	0.000	0.000
	B_0	0.518 09(4)	0.518 09(4)	0.566 84(2)	0.566 84(2)

TABLE II. Equilibrium molecular constants and bond length for the upper $[16.5]^3\Pi_2$ and the $X^3\Delta_3$ states of IrB and the isotopes (values given in parentheses are one standard error in last significant figures quoted).

State	Parameter	$^{193}\text{Ir}^{11}\text{B}$	$^{191}\text{Ir}^{11}\text{B}$		$^{193}\text{Ir}^{10}\text{B}$	
			Obs.	Calc. ^a	Obs.	Calc. ^a
$[16.5]^3\Pi_2$	T_e	16 203.627(2)	16 203.415(2)	...		
	ω_e	686.968(2)	687.436(2)	687.24		
	$\omega_e X_e$	21.067(2)	21.155(2)	21.14		
	B_e	0.473 579(2)	0.473 85	0.473 84		
	α_e	0.000 43	0.000 43	0.000 43		
	r_e (Å)	1.848 7				
	T_0				16 535.824	...
$X^3\Delta_3$	B_0	0.518 09(4)	0.518 09(4)	0.518 38	0.518 60(2)	0.517 92
	r_0 (Å)	1.767 5			0.566 84(2)	0.566 86

^aMolecular constants were calculated using isotopic relationships and the values from $^{193}\text{Ir}^{11}\text{B}$.

Tables I and II. A list of the measured transition line positions of the observed $[16.5]^3\Pi_2-X^3\Delta_3$ system of Ir¹⁰B and Ir¹¹B is available from EPAPS.²⁴

C. Perturbation at the (2, 0) band

We noticed extra transition lines appearing in the spectrum of the (2, 0) band. Figure 3 shows a portion of the spectrum with the P , Q , and R branches labeled and extra lines from the perturbing state are also marked. The transition lines of the (2, 0) band are broad and with unresolved hyperfine structure. The usual P , Q , and R branches were easily identified and assigned. Extra lines that perturbed the $v=2$ level were also assigned. It is easily noted that the line-width of the perturbing line is unusually large, which is due to the unresolved hyperfine structure and the overlapping of isotopic $^{191}\text{Ir}^{11}\text{B}$ and $^{193}\text{Ir}^{11}\text{B}$ molecular lines. Judging from the unresolved isotopic lines, we infer that the perturbing state has vibrational quantum number equal to 0. The perturbing state was found to have $\Omega'=3$ and $\Omega''=3$. Both the $^{191}\text{Ir}^{11}\text{B}$ and $^{193}\text{Ir}^{11}\text{B}$ lines were affected by the perturbing

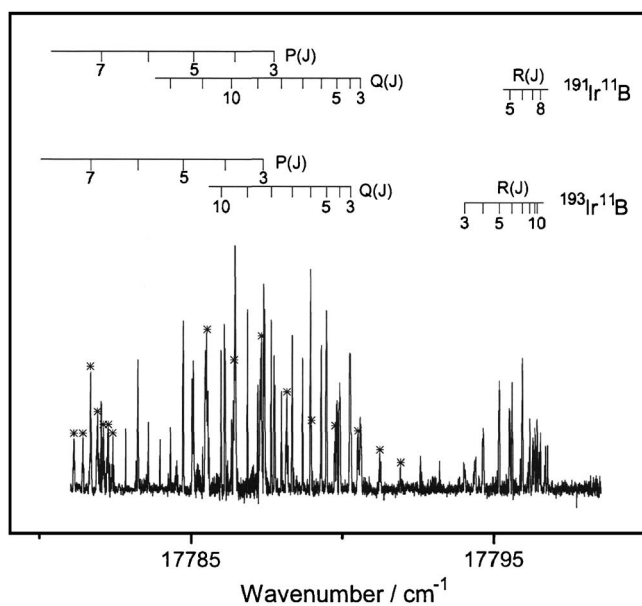


FIG. 3. The (2, 0) band of the $[16.5]^3\Pi_2-X^3\Delta_3$ transition of IrB.

state. Term values of the upper states were obtained by adding the corresponding lower state term values to the transition frequencies. Figure 4 depicts a reduced term value plot of the $v=2$ level of the $[16.5]^3\Pi_2$ state and the $v=0$ of perturbing $\Omega'=3$ level. If the data were fitted to expression (1) without taking into account of the perturbation, the B_2 value obtained was $0.474\ 24\ \text{cm}^{-1}$. However, assuming that no perturbation exists, the calculated B_2 value using known B values from other bands was $0.472\ 50\ \text{cm}^{-1}$. The increase of $0.001\ 74\ \text{cm}^{-1}$ in the rotational constant indicates that the $v=2$ level has been “pushed” by the perturbing state. In addition, the band origin, T_2 , was also pushed up by $0.13\ \text{cm}^{-1}$ from the expected position.

D. Electronic configuration and the electronic states

Marr *et al.*¹⁷ discussed the MO energy level diagram for IrC and IrN. Figure 5 shows a MO energy level diagram formed from the $6s$ and $5d$ atomic orbitals (AOs) of the iridium atom and the $2p$ orbital of a main group element. The lower energy 1σ and 1π MOs and the higher energy 3σ and 2π MOs are formed from the main group $2p$ AO and iridium $5d\sigma$, $5d\pi$, and $6p\sigma$ and $6p\pi$ AOs. The 2σ MO is essentially the iridium $6s$ AO. The 1δ MO is the iridium $5d\delta$ AO, be-

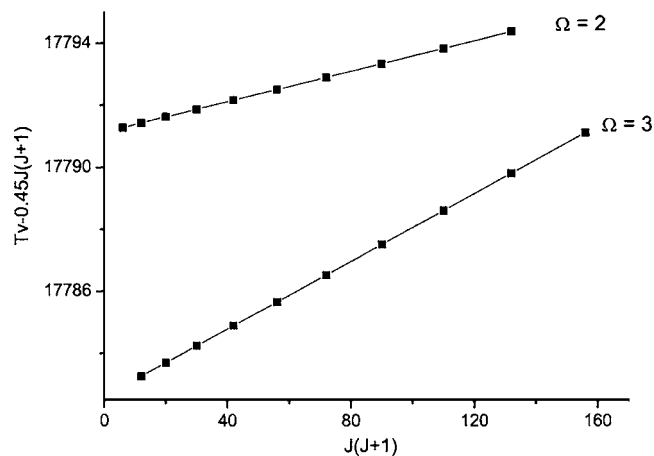


FIG. 4. Reduced term value plot of $v=2$ level of $\Omega=2$ and $\Omega=3$ states of IrB.

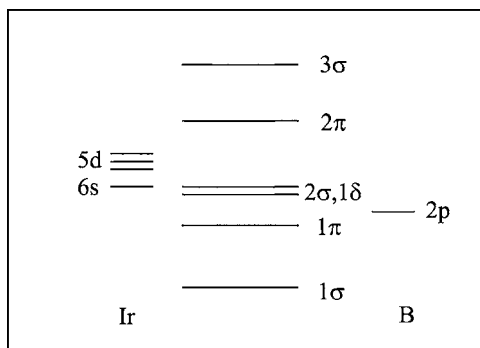
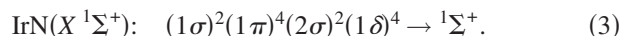
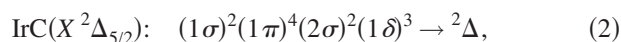
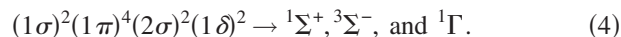


FIG. 5. MO energy level diagram of IrB.

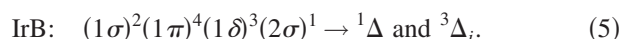
cause there is no other δ symmetry orbital around. The ground state of the diatomic molecules formed from iridium and a main group element are



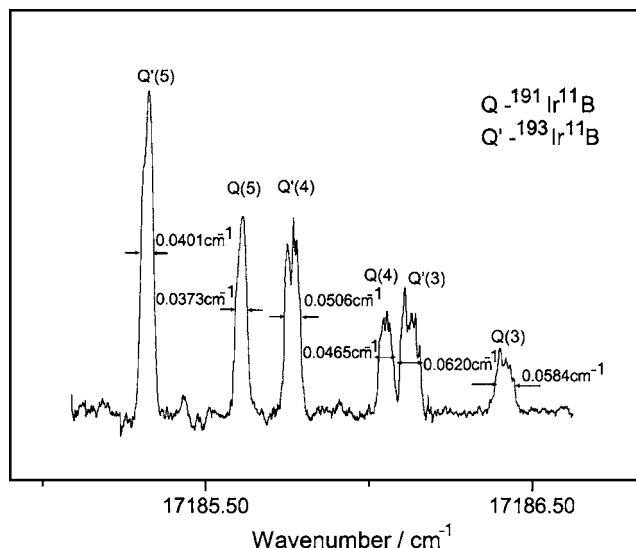
If the trend of putting the electron into the MOs is consistent with the above two iridium compounds, then IrB would have two electrons in the 1δ MO and the electronic configuration obtained is



From the first lines of various rotational branches, we concluded that the ground state has a spin component of $\Omega=3$. Close examination indicates that none of the electronic states in Eq. (4) has a spin component with $\Omega=3$; therefore, the lower state could not possibly arise from such an electronic configuration. Another possible low energy electronic configuration is



The difference between configurations (2) and (5) is that the two MOs, 1δ and 2σ , are so close in energy that in the case of IrB both are partially filled; however, in the IrC, the extra electron goes to the 2σ orbital giving the configuration (2). This energy order of the MOs is similar to that of rhodium carbide, nitride, oxide, and the fluoride molecules in Ref. 25. Due to the fact that the number of electrons in the δ orbital is more than one-half filled, the ${}^3\Delta$ state arising from the above configuration is an inverted ${}^3\Delta$ state. Such a ${}^3\Delta_i$ state has Ω values equal to 1, 2, and 3, and the $\Omega=3$ being the lowest energy. This fits well with the experimental observation and, therefore, is a reasonable assignment for the ground state. This electronic configuration is similar to that of the FeC molecule, which also has $X^3\Delta_3$ as the ground state.²⁶ As for the other spin-orbit components of the ${}^3\Delta_i$ ground state, the 1δ orbital is essentially the iridium $5d\delta$ AO, and it is possible to obtain an estimate of the spin-orbit parameter for the $X^3\Delta$ state from the atomic value of iridium. Lefebvre-Brion and Field²⁷ tabulated a value of $\zeta=3617 \text{ cm}^{-1}$ for iridium atom. Since the first order energy expression for calculating the spin-orbit components is $A\Lambda\Sigma$, the separation between the

FIG. 6. The (1, 0) band of the $[16.5]^3\Pi_2-X^3\Delta_3$ transition of IrB with partially resolved hyperfine structure.

spin-orbit components is estimated to be very large and probably over 7000 cm^{-1} . With our low temperature jet expansion source, it is expected that the higher energy spin-orbit components would not be appreciably populated. Nevertheless, with the origin of the (0, 0) band at 16541 cm^{-1} , we searched within the energy range of our laser sources between 15000 and 18000 cm^{-1} to look for other electronic transitions or transitions from other spin-orbit components of the ground state, but none was found. As for the assignment of the upper state, our experience is that a relatively strong transition would normally have $\Delta S=0$, when combining with the selection rule of $\Delta\Lambda=-1$ with $\Omega'=2$, it is reasonable that the upper state is assigned to be ${}^3\Pi_2$. Considering the promotion of an electron from the 2σ to the 2π MO, the following electronic configuration is obtained:



The electronic configuration in Eq. (6) gives rise to four excited states of which only the ${}^3\Pi_i$ and ${}^3\Phi_i$ have spin-orbit components with $\Omega=2$, but only the ${}^3\Pi_i$ state would satisfy the selection rule of $\Delta\Lambda=-1$. We have therefore assigned the observed transition as $[16.5]^3\Pi_2-X^3\Delta_3$ system. In view of the fact that the spin-orbit interaction is very large, it might be also appropriate to use Hund's case (c) notation to describe the electronic states, in which case the label would be just $[16.5] 2$.

E. Hyperfine structure in the observed system

Figure 6 shows the band head region of the (1, 0) band of the $[16.5]^3\Pi_2-X^3\Delta_3$ transition of the ${}^{191}\text{Ir}^{11}\text{B}$ and ${}^{193}\text{Ir}^{11}\text{B}$. It is easily noticed that the transition lines have linewidth wider than the expected Doppler width and with partially resolved structure. The unusually large linewidth basically arises from unresolved hyperfine structure, which is due to the interaction between the magnetic moment of unpaired electrons and the magnetic moment of nucleus in the

molecule. Since both ^{191}Ir ($I=3/2$, $\mu/\mu_N=0.146$) and ^{193}Ir ($I=3/2$, $\mu/\mu_N=0.159$) atoms have small nuclear magnetic moment and the ^{11}B ($I=3/2$, $\mu/\mu_N=2.689$) atom has a much larger nuclear magnetic moment,²⁸ it is expected that the hyperfine structure should be dominated by the ^{11}B atom. With only partially resolved hyperfine structure, it is not possible to obtain information concerning the small hyperfine interaction from the Ir atoms. As shown in Fig. 6, the rapid decrease in linewidth as J value increases indicates that the hyperfine coupling case conforms to Hund's case (a_β) coupling scheme.²²

The hyperfine Hamiltonian responsible for the magnetic interactions in a diatomic molecule is

$$\hat{H}_{\text{hfs}} = aI_zL_z + b\hat{I} \cdot \hat{S} + cI_zS_z, \quad (7)$$

where a , b , and c parameters are the same as those defined by Frosch and Foley.²⁹ The three terms in the hyperfine Hamiltonian are, respectively, the nuclear spin-orbit, the Fermi contact, and the dipolar electron spin and nuclear spin interactions. For Π and Δ states with large spin-orbit interaction, the angular momentum coupling case is appropriately described by Hund's coupling case (a_β). In such a coupling case, the grand total quantum number F results from the coupling of the nuclear spin I with angular momentum J , where $F=I+J$. Hyperfine matrix elements of a Π and a Δ state in Hund's coupling case (a_β) can be found in Ref. 30. The matrix elements important for this analysis are

$$\langle J\Omega IF | \hat{H}_{\text{hfs}} | J\Omega IF \rangle = \frac{\Omega \cdot h[F(F+1) - I(I+1) - J(J+1)]}{2J(J+1)}, \quad (8)$$

$$\langle J\Omega IF | \hat{H}_{\text{hfs}} | J-1, \Omega IF \rangle = \frac{h \cdot \sqrt{J^2 - \Omega^2} \sqrt{(J+I+F+1)(F+J-I)(J+I-F)(F+I-J+1)}}{2J\sqrt{2J+1}(2J-1)}, \quad (9)$$

where $h = \Lambda \cdot a + (b+c) \cdot \Sigma$. For the upper $^3\Pi_2$ ($\Lambda=1$ and $\Sigma=1$) state and the ground $^3\Delta_3$ ($\Lambda=2$ and $\Sigma=1$) state, the h' and h'' are $a+b+c$ and $2a+b+c$, respectively. We used the linewidth (FWHM, full width at half maximum) of the hyperfine lines and the diagonal matrix elements only [Eq. (8)] to obtain an estimate of the hyperfine parameters. As shown in Fig. 6, the FWHM linewidth of $Q(3)$ and $P(3)$ lines were, respectively, measured to be 0.0584 cm^{-1} and 0.0426 cm^{-1} , from which the hyperfine parameters for the upper and the lower states were estimated to be $h'=0.012 \text{ cm}^{-1}$ and $h''=0.029 \text{ cm}^{-1}$. The magnetic hyperfine parameter h is related to the expectation values of the coordinates of electron near the spinning nucleus³¹ and can be expressed as averages over electronic coordinates and is listed as follows (in cm^{-1} unit):

$$a = \left(\frac{\mu_0}{4\pi\hbar c} \right) g g_N \mu_B \mu_N \sum_i \langle r_i^{-3} \rangle, \quad (10)$$

$$b_F = \left(\frac{\mu_0}{4\pi\hbar c} \right) \frac{8\pi}{3} g g_N \mu_B \mu_N \left(\frac{1}{2S} \right) \sum_i \langle \psi_i^2(0) \rangle, \quad (11)$$

$$c = \left(\frac{\mu_0}{4\pi\hbar c} \right) \frac{3}{2} g g_N \mu_B \mu_N \left(\frac{1}{2S} \right) \sum_i \langle 3 \cos^2 \theta_2 - 1 \rangle \langle r_i^{-3} \rangle, \quad (12)$$

$$b = b_F - \frac{1}{3}c. \quad (13)$$

A method of obtaining estimates of the hyperfine parameters is using *ab initio* results for the atom. The a , b , and c parameters can be estimated using the expressions in Eqs. (10)–(13) and the values of $\langle r_i^{-3} \rangle$, $\langle r_i^{-3} \rangle^{24}$, $\langle \psi_i^2(0) \rangle$, and

$\langle 3 \cos^2 \theta_i - 1 \rangle$ from Morton and Preston.³² We obtained $a = 0.005 \text{ cm}^{-1}$, $b = 0.043 \text{ cm}^{-1}$, and $c = -0.002 \text{ cm}^{-1}$. The calculated h' and h'' are, respectively, 0.046 and 0.051 cm^{-1} , which are larger than those values determined experimentally by almost a factor of 2. This discrepancy could be due to the fact that the estimated h' and h'' values from experimental data are only preliminary and have relatively large errors because broad overall linewidths were measured. In addition, there have not been any theoretical calculations to estimate the contributions of various AOs to the MOs responsible for the hyperfine structure. Moreover, the effect of the nuclear quadrupole interaction, e^2Qq , has not been included in both experimental and theoretical treatments. A high resolution and detailed study of the hyperfine structure arising from both nuclei is important to resolve the noted discrepancy.

In summary, we report the first observation of an electronic transition spectrum of the IrB molecule. The observed transition has been assigned as the $[16.5]^3\Pi_2 - X^3\Delta_3$ transition. The bond length of the lower state has been determined to be 1.7675 \AA . The hyperfine parameters obtained also provide reasonable support to the assignment of the electronic states in this transition.

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- ¹M. Trenary, in *Materials Science of Carbides, Nitrides and Borides*, NATO Science Series Vol. 68, edited by Y. G. Gogotsi and R. A. Andrievski (Kluwer Academic, Dordrecht, 1998).
- ²B. Ganem and J. O. Osby, *Chem. Rev. (Washington, D.C.)* **86**, 163 (1986).
- ³H. J. Choi, D. Roundy, H. Sun, M. L. Cohen, and S. G. Louie, *Nature (London)* **418**, 758 (2002).
- ⁴P. K. Chowdhury and W. J. Balfour, *J. Chem. Phys.* **124**, 216101 (2006).
- ⁵C. R. Brazier, J. I. Ruiz, and S. V. Parks, *J. Mol. Spectrosc.* **241**, 1 (2007).
- ⁶C. W. Bauschlicher, Jr., and S. R. Langhoff, *J. Chem. Phys.* **101**, 80 (1994).
- ⁷F. R. Ornellas and A. R. S. Valentim, *J. Phys. Chem.* **98**, 12570 (1994).
- ⁸A. Ricca and C. W. Bauschlicher, Jr., *Chem. Phys. Lett.* **241**, 241 (1995).
- ⁹M. Pelegrini, O. Roberto-Neto, and F. B. C. Machado, *Int. J. Quantum Chem.* **95**, 205 (2003).
- ¹⁰I. Černušák, M. Dallos, H. Lischka, T. Müller, and M. Uhlár, *J. Chem. Phys.* **126**, 214311 (2007).
- ¹¹M. A. Ciriano, L. A. Oro, and M. T. Camellini, *Organometallics* **14**, 4764 (1995).
- ¹²R. Jimenez-Cataño and M. B. Hall, *Organometallics* **15**, 1889 (1996).
- ¹³V. Raziunas, G. Macur, and S. Katz, *J. Chem. Phys.* **43**, 1010 (1965).
- ¹⁴K. Jansson and R. Scullman, *J. Mol. Spectrosc.* **43**, 208 (1972).
- ¹⁵T. C. Steimle, A. J. Marr, S. A. Beaton, and J. M. Brown, *J. Chem. Phys.* **106**, 2073 (1997).
- ¹⁶R. S. Ram, J. Lievin, and P. F. Bernath, *J. Mol. Spectrosc.* **197**, 133 (1999).
- ¹⁷A. J. Marr, M. E. Flores, and T. C. Steimle, *J. Chem. Phys.* **104**, 8183 (1996).
- ¹⁸K. Jansson and R. Scullman, *J. Mol. Spectrosc.* **36**, 246 (1970).
- ¹⁹T. Ma, J. W.-H. Leung, and A. S.-C. Cheung, *Chem. Phys. Lett.* **385**, 259 (2004).
- ²⁰G. Herzberg, *Spectra of Diatomic Molecules* (Van Nostrand, New York, 1950).
- ²¹H. Lefebvre-Brion and R. W. Field, *The Spectra and Dynamics of Diatomic Molecules* (Elsevier, New York, 2004).
- ²²T. Dunn, in *Molecular Spectroscopy: Modern Research*, edited by K. N. Rao, (Academic, New York, 1972), Vol. III, Chap. 4.4.
- ²³Q. Ran, W. S. Tam, C. Ma, and A. S.-C. Cheung, *J. Mol. Spectrosc.* **198**, 175 (1999).
- ²⁴See EPAPS Document No. E-JCPSA6-128-011815 for the observed line positions of the $[16.5]^3\Pi_2-X^3\Delta_3$ transition. For more information on EPAPS, see <http://www.aip.org/pubservs/epaps.html>.
- ²⁵R. Li, R. H. Jensen, W. J. Balfour, S. A. Shepard, and A. G. Adam, *J. Chem. Phys.* **121**, 2591 (2004).
- ²⁶W. J. Balfour, J. Cao, C. V. V. Prasad, and C. X. W. Qian, *J. Chem. Phys.* **103**, 4046 (1995).
- ²⁷H. Lefebvre-Brion and R. W. Field, *Perturbations in the Spectra of Diatomic Molecules* (Academic, New York, 1986).
- ²⁸I. Mills, T. Cvitaš, K. Homann, N. Kallay, and K. Kuchitsu, *Quantities, Units and Symbols in Physical Chemistry* (Blackwell Scientific, London, 1993).
- ²⁹R. A. Frosch and H. M. Foley, *Phys. Rev.* **88**, 1337 (1952).
- ³⁰Y. Azuma, J. A. Barry, M. P. J. Lyne, A. J. Merer, J. O. Schroder, and J. L. Femenias, *J. Chem. Phys.* **91**, 1 (1989).
- ³¹C. H. Townes and A. L. Schawlow, *Microwave Spectroscopy* (Dover, New York, 1975).
- ³²J. R. Morton and K. F. Preston, *J. Magn. Reson. (1969-1992)* **30**, 577 (1978).