Electronic Spectra of Antimony Monobromide Part II*

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(Z. Naturforsch. **30 a**, 695–696 [1975];
received March 8, 1975)

A new group of bands, designated as system A, of antimony monobromide, lying in the region 5340–4905 Å, has been photographed and completely analysed. Two characteristic continua with maxima at 5320 Å and 6500 Å have been observed. The electronic states with the recorded band systems have been tentatively assigned.

In an earlier paper (1971) the senior author had analysed the band structures of the two band systems lying in the regions 3340–3050 Å and 3020–2950 Å. These were designated as systems B₁ and B₂ respectively.

Another system, extending from 5340–4905 Å, was also mentioned. Heretofore, it will be designated as system A. The general characteristics of this system are the same as those of systems B₁ and B₂. The vibrational analysis of this new system and a general consideration of the electronic structure of the molecule are the subject matter of this paper.

System A (5340–4905 Å)

Using the same experimental set up and conditions as described earlier, the band system under discussion was photographed with a Fuess spectrograph. In this region a large number of atomic lines have been recorded, indicating a good deal of dissociation of the molecule into its constituent elements. Two continua, one situated just at the long wavelength end of the observed discrete band system with a maximum at about 5320 Å, and the other near 6500 Å, have been observed. An enlargement of the spectrum is given in Figure 1.

The wavenumbers in vacuum of the band heads, their visually estimated relative intensities, vibrational analysis along with ν obs. and O–C values are given in Table 1. All the bands are degraded towards the longer wavelength side.

Table 1. Band head data and vibrational analysis of the system A (5340–4905 Å).

Wave number in vacuum (cm ⁻¹)	Relative intensity	ν' , ν''	O–C values (cm ⁻¹)
18732	8	2,6	10
18966	5	2,5	5
18984	8	1,4	–2
19200	8	2,4	–1
19230	7	1,3	4
19253	3	0,2	1
19375	4	4,5	–12
19445	3	2,3	4
19470	8	1,2	2
19494	6	0,1	0
19612	3	4,4	–15
19706	8	1,1	–3
19736	7	0,0	0
19771	1	—	—
19831	2	5,4	–8
19900	3	3,2	5
19951	2	1,0	0
20140	2	3,1	3
20166	2	2,0	0
20380	9	3,0	1

A faint line lying on a continuum sometimes gives the impression of a band. Whenever such a suspicion arose, the authors have preferred not to mention it in the above table. The total number of such bands is four.

The Franck-Condon parabola representing the relative intensity distribution in the band system is similar to that obtained in the system B₁. The values

Fig. 1. Emission spectrum of SbBr molecule.

* This work was done at Banaras Hindu University, Varanasi (India) and subsequently repeated at the University of Jodhpur, Jodhpur (India).

of ω_e' and ω_e'' for the states involved conform with the nature of the F-C parabola.

The $\Delta G''(v+1/2)$ values clearly show that the lower state of this system can be identified with the lower state of the systems B_1 and B_2 . This fact was taken into account while assigning v'' values to different bands of the system. The value of ω_e'' obtained tallies very well with that obtained for the systems B_1 and B_2 .

All the bands of the system A can fairly well be represented by the following equation:

$$\nu = 19736 + (215.45 v' - 0.35 v'^2) - (242.6 v'' - 0.33 v''^2).$$

The ω_e values are correct to $\pm 3 \text{ cm}^{-1}$, while the $\omega_e \omega_e$ values are correct to the first decimal place only.

On the basis of a comprehensive discussion (Ph.D. Thesis (1964) of the senior author, submitted at Banares Hindu University, Varanasi, India) the ground state of the molecule was suggested to be $^3\Sigma^{-1}$ and the two excited states to be $^3\Sigma^{-1}$ and

$^3\Pi$. The bands system was attributed to the following transitions.

It is well known that a correct species of electronic states involved in the different transitions can only be decided by rotational analysis. Since the molecule is heavy, the bands are slightly diffuse, and SbBr_3 is deliquescent, the rotational analysis is not practically possible. Hence the assignments may only be treated as tentative.

Acknowledgement

The authors are thankful to Prof. N. L. Singh and Prof. D. K. Rai of Physics Department, Banares Hindu University, Varanasi (India) for valuable discussions and suggestions and to Prof. A. N. Nigam of Jodhpur University for giving facilities and continuous encouragement. One of us, Miss A. R. Sud is grateful to U.G.C. (India) for the award of Junior Research Fellowship.

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