練習実験報告

肖宇笑 May 16, 2024

Glavano Sepctrum

REMPI scan

Selected peaks

Radius and angular distributions

Peak assignments

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Glavano Sepctrum

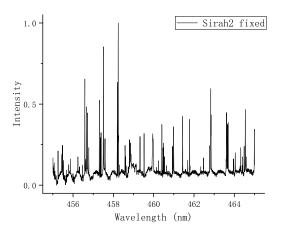


Fig. 1: Wavelen. correction

Glavano Sepctrum



Fig. 1: Wavelen. correction

Glavano Sepctrum

Correction



Fig. 2: Correction function

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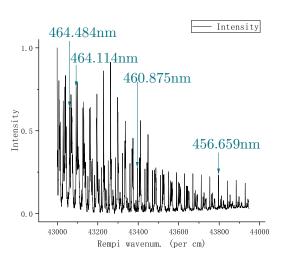
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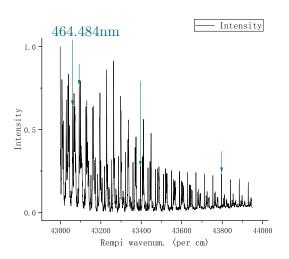
Glavano Sepctrum

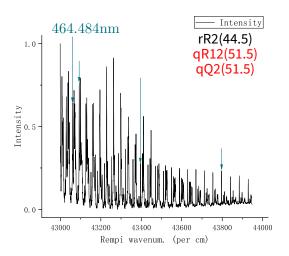
REMPI scan

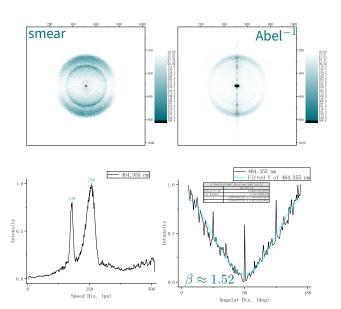
Selected peaks

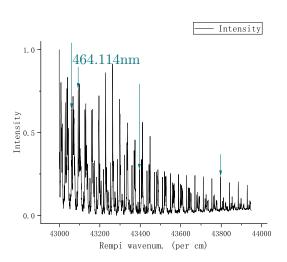
Radius and angular distributions

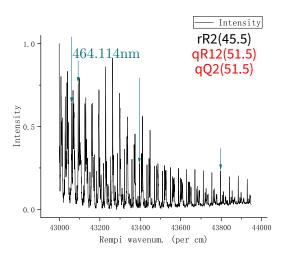
Peak assignments

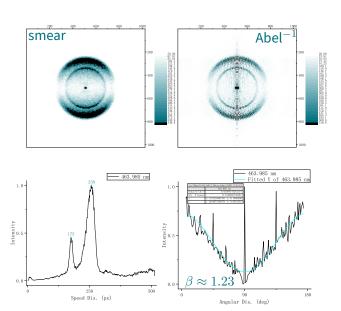


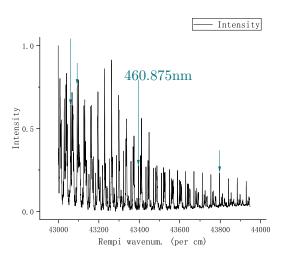


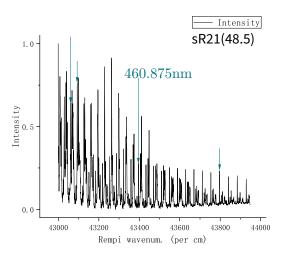


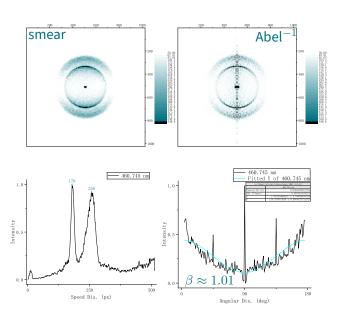


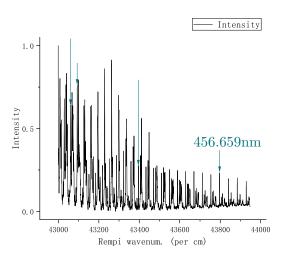


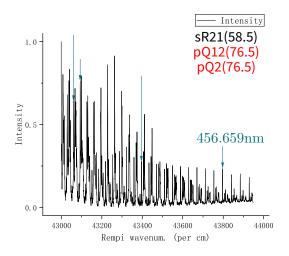


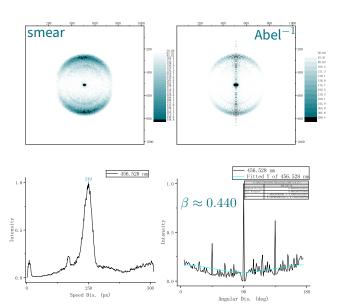












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464.484 nm ≈ 43058.49 cm ⁻¹	464.114 nm ≈ 43092.81 cm ⁻¹	460.875 nm $\approx 43395.69 \text{cm}^{-1}$	456.659 nm $\approx 43796.34 \text{cm}^{-1}$
px = 258	px = 258	px = 258	px = 249
$rR2\ 44.5$ $qR12\ 51.5$ $qQ2\ 51.5$	$rR2\ 45.5$ $qR12\ 51.5$ $qQ2\ 51.5$	sR21~48.5	$sR21\ 58.5$ $pQ12\ 76.5$ $pP2\ 76.5$

Notice

Colored assignments are mismatched, and will not be used to calculate.

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	E_{total}	$E_{bond}(\mathrm{O}\!-\!\mathrm{NO})^1$	$E_{int.}(NO)$
Peak 1 464.484nm	$43058.49\mathrm{cm}^{-1}$		E(v = 1) + E(J = 44)
Peak 2 464.114nm	43 092.81cm ⁻¹		E(v = 1) + E(J = 45)
Peak 3 460.875nm	43 395.69cm ⁻¹	25 128.57cm ⁻¹	E(v = 1) + E(J = 48)
Peak 4 456.659nm	43796.34cm ⁻¹		E(v = 1) + E(J = 58)

¹Rémy Jost et al. *The Journal of Chemical Physics* **105**.3 (July 1996).

	E_{total}	$E_{bond}(\mathrm{O}\!-\!\mathrm{NO})^2$	$E_{int.}(NO)$
Peak 1 464.484nm	$43058.49\mathrm{cm}^{-1}$		3525.0743625cm ⁻¹ + $E(J = 44)$
Peak 2 464.114nm	$43092.81\mathrm{cm}^{-1}$		$3525.0743625 \text{cm}^{-1} + E(J = 45)$
Peak 3 460.875nm	43395.69 cm $^{-1}$	25 128.57cm ⁻¹	$3525.0743625 \text{cm}^{-1} + E(J = 48)$
Peak 4 456.659nm	43796.34 cm $^{-1}$		$3525.0743625 \text{cm}^{-1} + E(J = 58)$

²Rémy Jost et al. *The Journal of Chemical Physics* **105**.3 (July 1996).

³J. Danielak et al. Journal of Molecular Spectroscopy **181**.2 (1997), pp. 394–402.

	E_{total}	$E_{bond}(\mathrm{O}\mathrm{-NO})^2$	$E_{int.}(NO)$
Peak 1 464.484nm	$43058.49\mathrm{cm}^{-1}$		3525.0743625cm ⁻¹ + $E(J = 44)$
Peak 2 464.114nm	43092.81 cm $^{-1}$		3525.0743625cm ⁻¹ + $E(J = 45)$
Peak 3 460.875nm	43 395.69cm ⁻¹	25 128.57cm ⁻¹	$3525.0743625 \text{cm}^{-1} + E(J = 48)$
Peak 4 456.659nm	43796.34 cm $^{-1}$		$3525.0743625 \text{cm}^{-1} + E(J = 58)$

Vib. energy level³

$$E(v) = \omega_e \left(v + \frac{1}{2}\right) - \omega_e x_e \left(v + \frac{1}{2}\right)^2 + \omega_e y_e \left(v + \frac{1}{2}\right)^3.$$

²Rémy Jost et al. *The Journal of Chemical Physics* **105**.3 (July 1996).

³J. Danielak et al. Journal of Molecular Spectroscopy 181.2 (1997), pp. 394–402.

	E_{total}	$E_{bond}(O\!-\!NO)^4$	$E_{int.}(NO)$
Peak 1 464.484nm	$43058.49\mathrm{cm}^{-1}$		$6848.563763 \mathrm{cm}^{-1}$
Peak 2 464.114nm	43 092.81cm ⁻¹		6999.631462cm ⁻¹
Peak 3 460.875nm	43 395.69cm ⁻¹	25 128.57cm ⁻¹	7472.976 923cm ⁻¹
Peak 4 456.659nm	43796.34 cm $^{-1}$		$9269.004023\mathrm{cm}^{-1}$

⁴Rémy Jost et al. The Journal of Chemical Physics **105**.3 (July 1996).

⁵J. Danielak et al. Journal of Molecular Spectroscopy **181**.2 (1997), pp. 394–402.

	E_{total}	$E_{bond}(\mathrm{O}\!-\!\mathrm{NO})^4$	$E_{int.}(NO)$
Peak 1 464.484nm	$43058.49\mathrm{cm}^{-1}$		$6848.563763 \mathrm{cm}^{-1}$
Peak 2 464.114nm	43092.81 cm $^{-1}$		6999.631 462cm ⁻¹
Peak 3 460.875nm	43395.69 cm $^{-1}$	25 128.57cm ⁻¹	$7472.976923\mathrm{cm}^{-1}$
Peak 4 456.659nm	43796.34 cm $^{-1}$		9269.004023 cm $^{-1}$

Rot. energy level⁵

$$E\left(J\right) = B_v \left(J^2 + J\right)$$

⁴Rémy Jost et al. *The Journal of Chemical Physics* **105**.3 (July 1996).

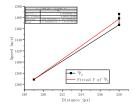
⁵J. Danielak et al. Journal of Molecular Spectroscopy **181**.2 (1997), pp. 394–402.

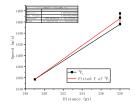
$E_{int.}(O)$	$\begin{split} E_{\text{trans}}(total) &\approx 2.88 E_{\text{trans}}(\text{NO}) \\ &= E_{\text{total}} - E_{\text{bond}}(\text{O-NO}) - E_{\text{int.}}(\text{O}) - E_{\text{int.}}(\text{O}) \end{split}$	$E_{trans}(NO) \\ = \frac{1}{2} m(NO) v^2(NO)$
$^{3}P_{2}$ (0cm^{-1})	$11081.356237 \mathrm{cm}^{-1}$ $10964.608538 \mathrm{cm}^{-1}$ $10794.143077 \mathrm{cm}^{-1}$ $9398.765977 \mathrm{cm}^{-1}$	1923.84656892361cm ⁻¹ 1903.57787118055cm ⁻¹ 1873.98317309028cm ⁻¹ 1631.73020434028cm ⁻¹
$^{3}P_{1}$ (158.625cm ⁻¹)	$10922.731237 \mathrm{cm}^{-1}$ $10805.983538 \mathrm{cm}^{-1}$ $10635.518077 \mathrm{cm}^{-1}$ $9240.140977 \mathrm{cm}^{-1}$	1896.30750642361cm ⁻¹ 1876.03880868055cm ⁻¹ 1846.44411059028cm ⁻¹ 1604.19114184028cm ⁻¹
$^{3}P_{0}$ (226.977cm ⁻¹)	$10854.379237 \mathrm{cm}^{-1}$ $10737.631538 \mathrm{cm}^{-1}$ $10567.166077 \mathrm{cm}^{-1}$ $9171.788977 \mathrm{cm}^{-1}$	1884.44083975695cm ⁻¹ 1864.17214201389cm ⁻¹ 1834.57744392361cm ⁻¹ 1592.32447517361cm ⁻¹

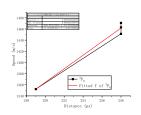
y Trans. energy of NO

$E_{int.}(O)$	$v(NO) = \sqrt{\frac{2E_{trans}(NO)}{m(NO)}}$	Δy
$^{3}P_{2}$ $_{(0\text{cm}^{-1})}$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	258 px 258 px 258 px 258 px 249 px
$^{3}P_{1}$ (158.625cm ⁻¹)	$ \begin{vmatrix} 1475.5645855\mathrm{ms}^{-1} \\ 1467.6576085\mathrm{ms}^{-1} \\ 1456.0353685\mathrm{ms}^{-1} \\ 1357.162647\mathrm{ms}^{-1} \end{vmatrix} $	258 px 258 px 258 px 249 px
$^{3}P_{0}$ (226.977cm ⁻¹)	$\begin{array}{c} 1470.940464\mathrm{ms^{-1}} \\ 1463.0084955\mathrm{ms^{-1}} \\ 1451.3490265\mathrm{ms^{-1}} \\ 1352.133667\mathrm{ms^{-1}} \end{array}$	258 px 258 px 258 px 249 px

y Trans. energy of NO

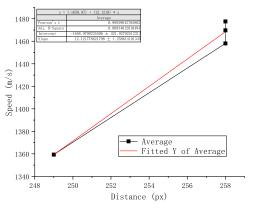






$$\tfrac{dv}{d\,{\rm px}} = 12.04{\rm m\,s^{-1}/px\,@}\,^3P_2 \quad \tfrac{dv}{d\,{\rm px}} = 12.14{\rm m\,s^{-1}/px\,@}\,^3P_1 \quad \tfrac{dv}{d\,{\rm px}} = 12.18{\rm m\,s^{-1}/px\,@}\,^3P_0$$

y Trans. energy of NO



$$rac{dv}{d\,\mathrm{px}} = 12.12\mathrm{m\,s^{-1}/px}$$
 @ Average

Reference

- [1] J. Danielak et al. *Journal of Molecular Spectroscopy* **181**.2 (1997), pp. 394–402.
- [2] Rémy Jost et al. The Journal of Chemical Physics 105.3 (July 1996).

Open source

