

Absorption Spectroscopy of ^{85}Rb Atoms at 420 nm: Pressure Broadening with N_2 Buffer Gas and Saturation Intensity

Xi Zeng and Dmitri L. Boiko

Centre Suisse d'Électronique et de Microtechnique, Jaquet-Droz 1, 2002 Neuchâtel, Switzerland

Email: xi.zeng@csem.ch, dmitri.boiko@csem.ch

Absorption spectroscopy of Rb atoms in the traditional near-infrared D_1 (795 nm wavelength) and D_2 (780 nm) transitions is mature. In contrast, the blue wavelength Rb transitions are not well explored. Here, we present absorption spectroscopy measurements of ^{85}Rb in the $5^2\text{S}_{1/2}$ - $6^2\text{P}_{3/2}$ (420 nm) transition with data on pressure broadening due to N_2 buffer gas and absorption saturation due to high optical intensity. To the best of our knowledge, there is no literature on the effects of N_2 in this atomic transition.

A custom built blue-violet external cavity diode laser was used to perform absorption spectroscopy measurements on ^{85}Rb vapor in the $5^2\text{S}_{1/2}$ - $6^2\text{P}_{3/2}$ transition at 420.18 nm. These vapor cells are antireflection coated and contain N_2 buffer gas at various pressures, ranging from no buffer gas to 200 torr. Figure 1(a) is an example ^{85}Rb absorption spectra, here in 6 torr of N_2 buffer gas at 100 °C. The absorption oscillator strength in this transition line is smaller than that of D lines by an order of magnitude. Absorption spectra are analyzed accounting for all six hyperfine transitions between ground and excited state ($F_g=2 \rightarrow F_e=1,2,3$; $F_g=3 \rightarrow F_e=2,3,4$) using Voigt profiles. The pressure broadening coefficient is then extracted, as shown in Fig. 2(b), and found to be approximately 20 MHz/torr. This is smaller than the coefficients of He (45-50 MHz/torr) and Ar (25-30) buffer gas for the same transition line that were previously reported,¹ and it fits the trend of N_2 coefficient being smaller than He and Ar in D lines.²

Saturation of absorption cross section due to high optical intensity is measured using a cell without buffer gas and at 100 °C as shown in Fig. 1(c). Using a 3 energy level system model, the saturation intensity is found to be 150 mW/cm², which is set by the relaxation time due to collision with the cell walls (30 μs) but not by the excited state ($6^2\text{P}_{3/2}$) lifetime (on the order of 100 ns).

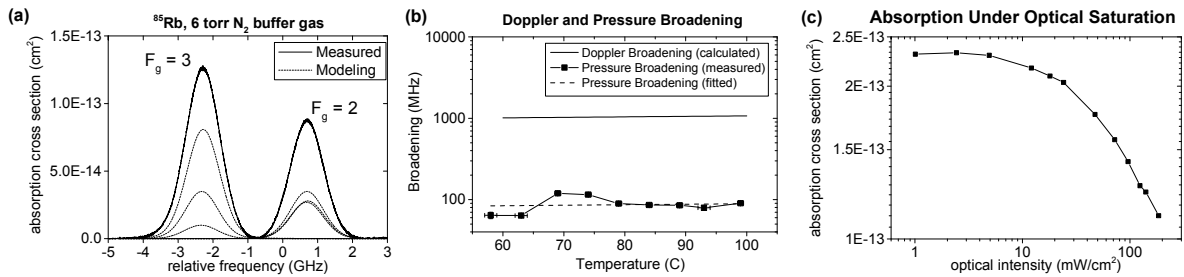


Fig. 1. (a) Absorption spectra of ^{85}Rb in 6 torr of N_2 buffer gas and at 100 °C in the $5^2\text{S}_{1/2}$ - $6^2\text{P}_{3/2}$ transition (420 nm). The dashed curves represent the individual hyperfine transitions. (b) Doppler and pressure broadening in 6 torr of N_2 buffer gas as a function of temperature. (c) Peak absorption cross section of ^{85}Rb , without buffer gas and at 100 °C, as a function of optical intensity.

¹ D. Aumiler, T. Ban, and G. Pichler, "High-resolution measurements of the pressure broadening and shift of the rubidium $5^2\text{S}_{1/2}$ - $6^2\text{P}_{3/2}$ line by argon and helium", Physical Review A, vol. 70, no. 3, p. 032723, 2004.

² M. D. Rotondaro and G. P. Perram, "Collisional broadening and shift of the rubidium D_1 and D_2 lines ($5^2\text{S}_{1/2} \rightarrow 5^2\text{P}_{1/2}$, $5^2\text{P}_{3/2}$) by rare gases, H_2 , D_2 , N_2 , CH_4 , and CF_4 ", J. Quant. Spectrosc. Radiat. Transfer, vol. 57, no. 4, pp. 497-507, 1997.