

OPTICAL ABSORPTION SENSITIVITY BETTER THAN 1×10^{-12}

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Improved physical understanding of the REAL sensitivity-limiting processes, along with better technical solutions for cavity-enhanced optical heterodyne spectroscopy recently have allowed remarkable sensitivity improvements, presently at the absorption level of 5.2×10^{-13} for a 1 s integration ^a. Here we review the several problems which led to this NICE-OHMS solution, and report recent progress with active control of the Residual Amplitude Modulation produced by the Electro-Optic Modulator.

Also, saturated absorption spectra near 1064 nm for HCCD, HCCH, and CO₂ are presented. The two additional lines are ¹²C₂H₂ ($2\nu_1 + \nu_2 + \nu_5$) R(12) ^b and ¹²C¹⁶O₂ ($2\nu_1 + 3\nu_3$) R(6) ^c, with their respective transition dipole moments of 50 μ Debye and 6 μ Debye. They are both weaker than our usual C₂HD($\nu_2 + 3\nu_3$) P(5) transition, which has a transition dipole moment of 70 μ Debye, but all are recovered with excellent signal-to-noise ratios. The absolute resonance center frequencies of all three transitions have been measured (\pm 25 kHz) using as reference a Nd:YAG laser locked via frequency doubling on the a₁₀ hyperfine-structure component of the R(56) 32-0 I₂ transition. The C₂H₂ resonance is about 4-fold weaker than that of C₂HD, while the pressure broadening rate of 34(1) MHz/Torr (FWHM) is similar. For the CO₂ transition, however, the saturated absorption signal is much weaker, by more than a factor of 350, and shows an elegant and unexpected lineshape which is believed to result from nearly overlapping one- and two-photon transitions.

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