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# **Sensitive detection of aerosols and gases using Raman scattering**

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**2nd International Conference & Exhibition on Lasers,  
Optics and Photonics**

**September 08-10, 2014**



This work was sponsored by the Edgewood Chemical Biological Center portion of this work was sponsored by the Defense Threat Reduction Agency under Air Force Contract FA8721-05-C-0002. Opinions, interpretations, conclusions, and recommendations are those of the authors and are not necessarily endorsed by the United States Government.

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# Outline

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- **Introduction to Raman scattering**
- **Schematic of the Raman setup**
- **Detection of aerosols**
- **Detection of gases**
- **Summary**
- **Acknowledgments**



# Introduction to Raman Scattering

- Raman scattering (inelastic light scattering due to molecular vibrations and/or rotations) was first observed by Prof. C. V. Raman on February 28, 1928 using the light from a Hg arc lamp. The results of this experiment were published on March 31, 1928.
- Lasers are now used for Raman scattering experiments.
- Raman spectrum of a material is its molecular *fingerprint*.
- Raman cross sections are very very small ( $1 \times 10^{-30}$  -  $1 \times 10^{-25}$  cm<sup>2</sup>).

Sir C. V. Raman (1888 – 1970)  
1930 Nobel Prize for Physics





# Raman Signal

Raman signal  $S_R$  (CCD counts) is given by

$$S_R = \eta_c \eta_q \sigma_R N_m L (P_L \tau / h \nu_L)$$

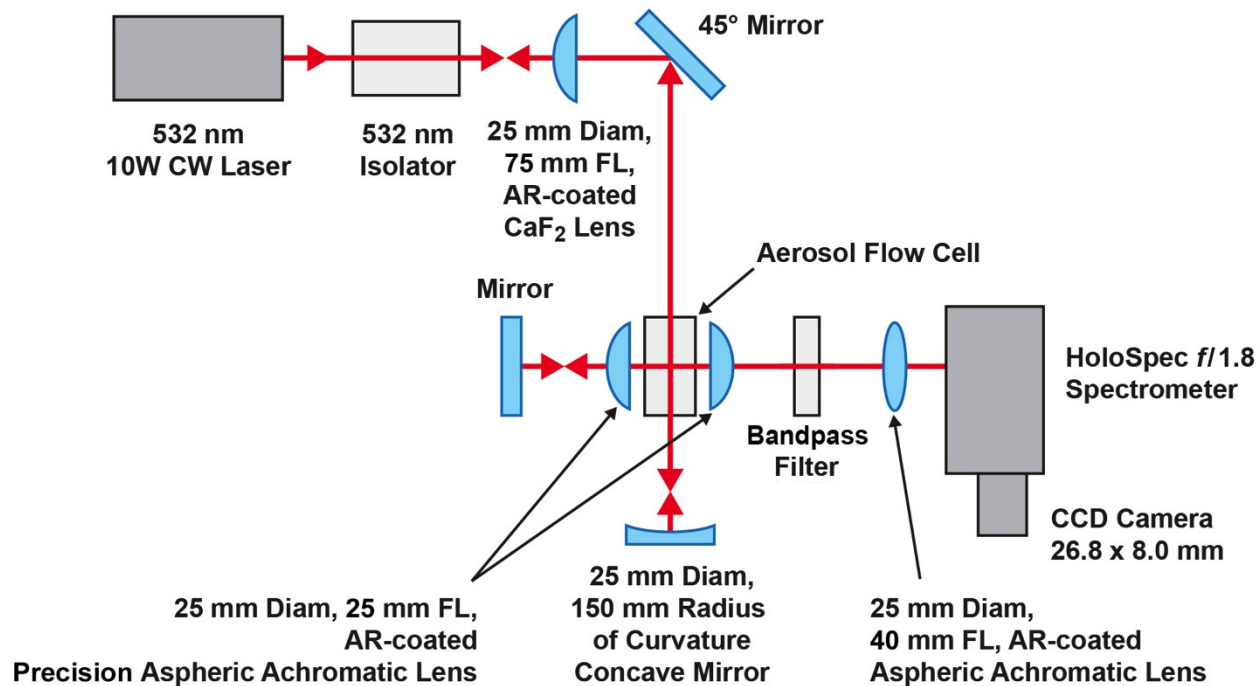
where

- $\eta_c$  = Collection efficiency
- $\eta_q$  = CCD quantum efficiency
- $\sigma_R$  = Raman cross section (cm<sup>2</sup>)
- $N_m$  = Molecular concentration (cm<sup>-3</sup>)
- $L$  = Laser path length (cm)
- $P_L$  = Laser power (W)
- $\tau$  = Integration time (s)
- $h \nu_L$  = Laser photon energy (J)

Relative Raman signals for different materials are determined by the value of the product of  $\sigma_R$  and  $N_m$ .



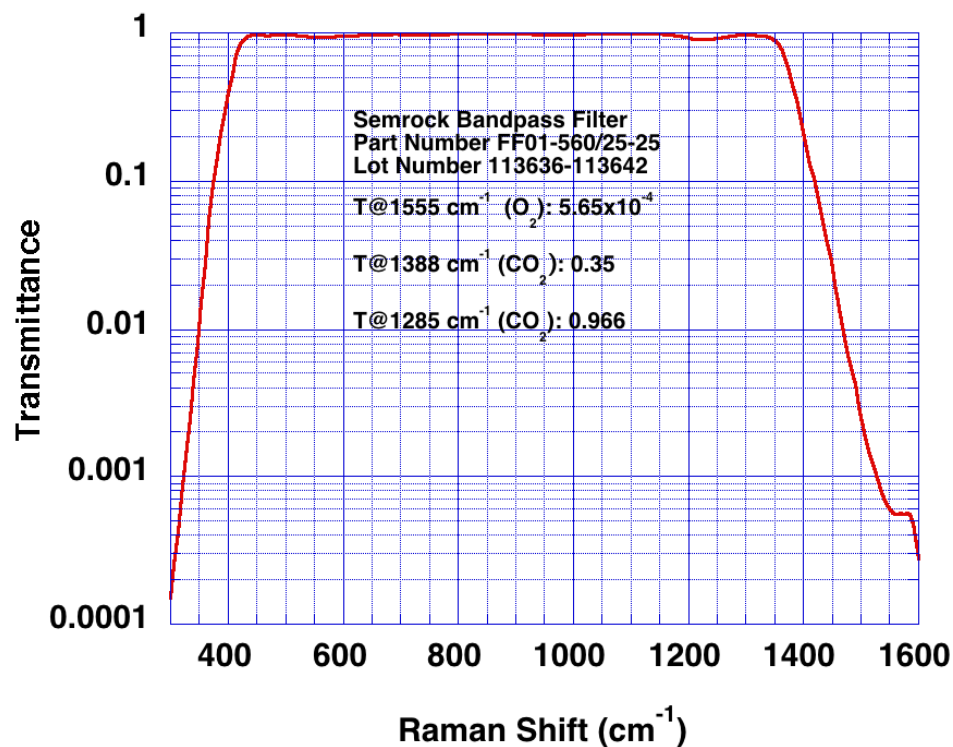
# Schematic of the Raman Setup



**The spectral resolution is  $10 \text{ cm}^{-1}$  using the standard grating and  $4 \text{ cm}^{-1}$  using the high-dispersion grating.**



# Transmittance of the Bandpass Filter



~ 98% transmittance over the 430-1350 cm<sup>-1</sup> spectral range





# The CCD Image Obtained with Standard Grating and 30 s Integration Time

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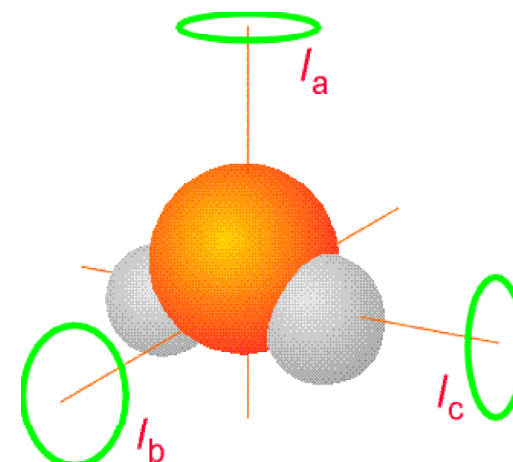
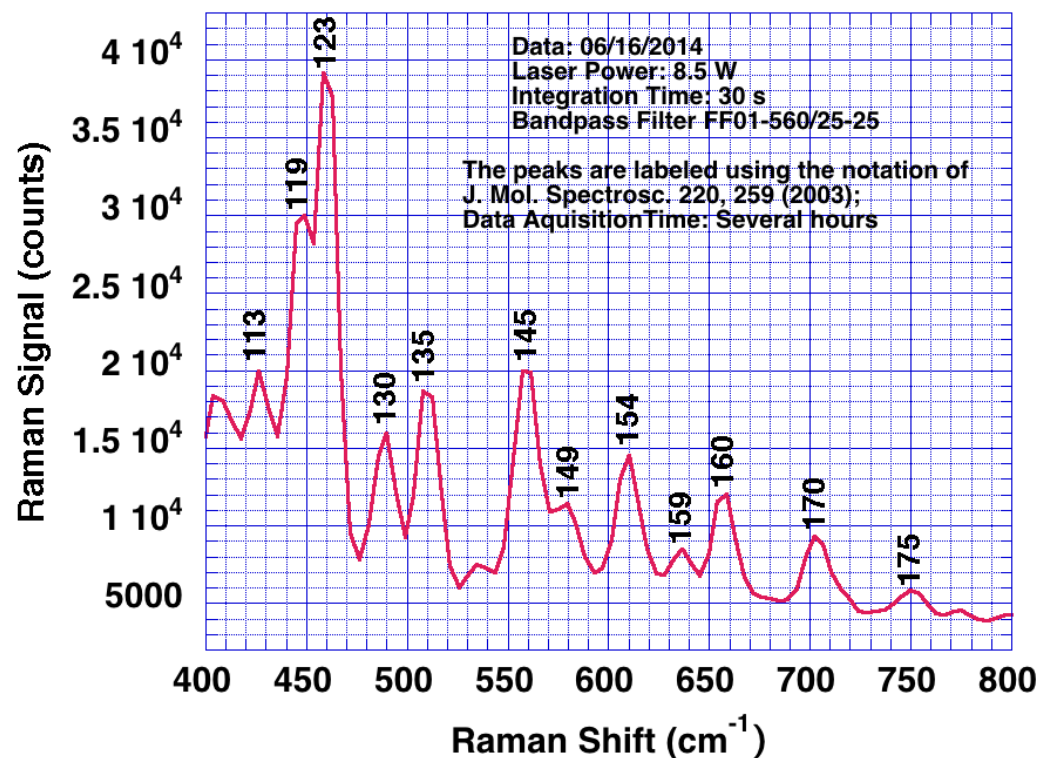
H<sub>2</sub>O

CO<sub>2</sub>

O<sub>2</sub>



# H<sub>2</sub>O Rotational Spectrum Using Standard Grating

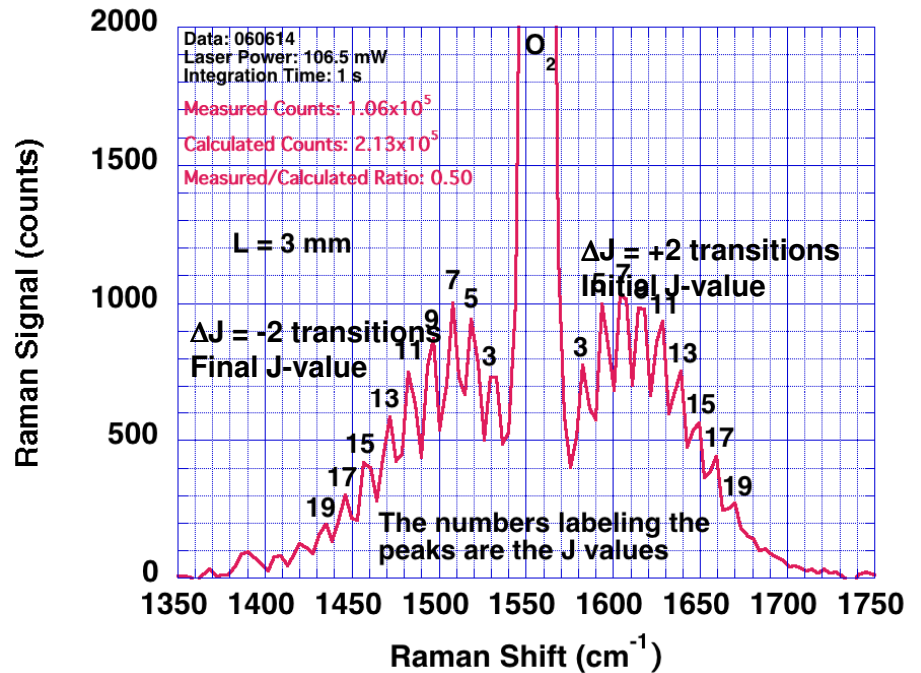
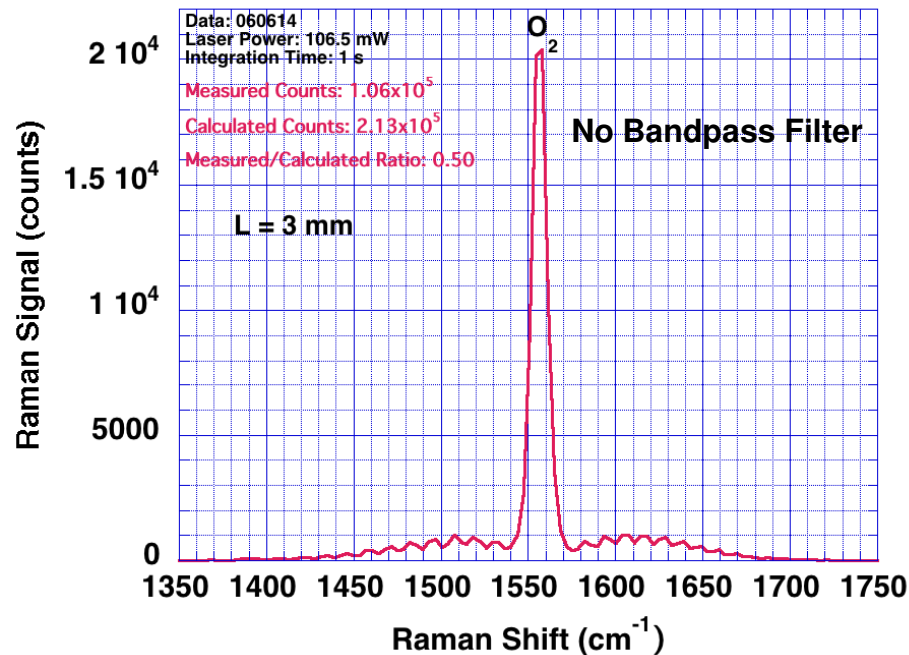


Rotational constants:

$$A = h/8\pi^2 c I_a = 27.9 \text{ cm}^{-1}$$
$$B = h/8\pi^2 c I_b = 14.5 \text{ cm}^{-1}$$
$$C = h/8\pi^2 c I_c = 9.3 \text{ cm}^{-1}$$

The peaks are labeled using the notation of Avila et al., J. Mol. Spectrosc. 220, 259 (2003).

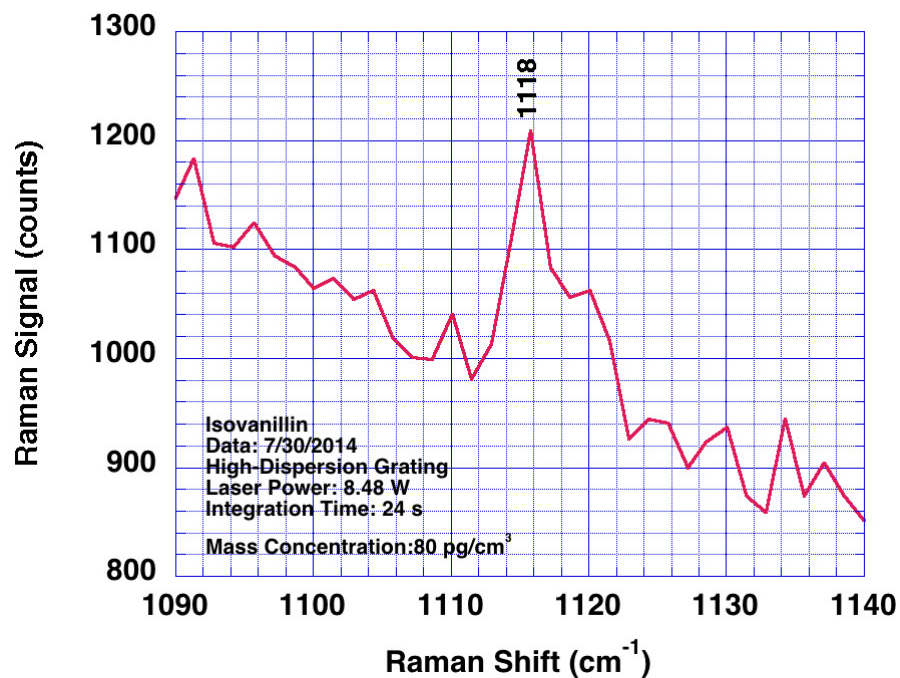
# Performance of Our Raman Setup Using the Standard Grating Without Bandpass Filter



(i) The measured value of the Raman signal is 2x smaller than that of the calculated value, partly due to vignetting of the rays from off-axis points on the laser beam. (ii) The selection rule for rotational transitions is  $\Delta J = \pm 2$ , where  $J$  is the rotational quantum number and even  $J$  transitions are completely missing. The most populated level at 300 K is  $J = (kT/2hcB)^{1/2} - 1/2 = 8$ ;  $B$  is the rotational constant equal to  $1.45 \text{ cm}^{-1}$  for  $\text{O}_2$ .



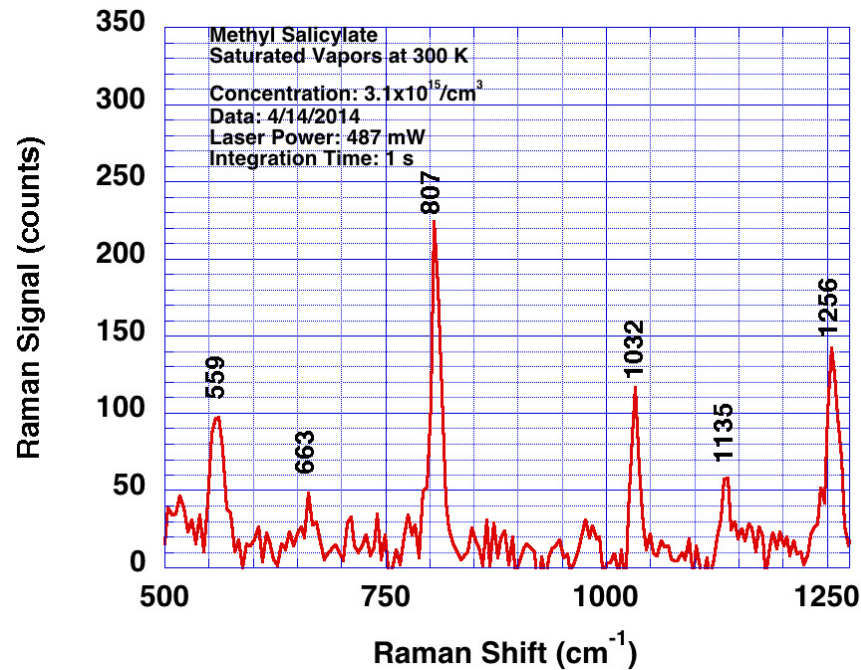
# Detection of Isovanillin Aerosol Using High-Dispersion Grating



**Limit of detection < 80 pg/cm<sup>3</sup>**



# Detection of Methyl Salicylate Vapors Using Standard Grating



- Limit of detection deduced to be  $< 2 \text{ ng/cm}^3$  using 10 W laser power and 30 s integration time.
- $\sigma_R$  for the  $807 \text{ cm}^{-1}$  mode determined to be  $2.80 \pm 0.17 \times 10^{-29} \text{ cm}^2$



# Summary

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- **Limit of detection < 80 pg/cm<sup>3</sup> in 24 s for isovanillin aerosol using the high-dispersion grating.  $\sigma_R$  is equal to  $3.3 \times 10^{-28}$  cm<sup>2</sup> for the 1116 cm<sup>-1</sup> mode.**
- **Limit of detection deduced to be < 2 ng/cm<sup>3</sup> for methyl salicylate vapors for 10 W laser power and 30 s integration time using the standard grating. The value of  $\sigma_R$  has been determined to be  $2.80 \pm 0.17 \times 10^{-29}$  cm<sup>2</sup> for the 807 cm<sup>-1</sup> mode.**





# Acknowledgments

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- **David Sickenberger (ECBC)**
- **Steven Christensen (ECBC)**
- **Max Howe (MIT Lincoln Laboratory)**

**Thanks for listening to me.**



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