

**Spectroscopy of Cs Atoms**

\$\$ National Science Council, Taiwan \$\$  
\$\$ 國科會 \$\$

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**What is spectroscopy?**

Fig. 3.—The HCl band at  $3.46 \mu$ , mapped with 7500-line grating. HCl at atmospheric pressure.

What are the important issues on the spectrum?  
**Line position, Intensity and Shape**

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**Outline**

Two-photon transitions of Cs atoms

Cs 6s-8s TPTs

Cs 6s-7d TPTs

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**Two-Photon Spectroscopy**

**6s-8s Two-photon transition in Cesium  
(Cell at 50.0 °C)**

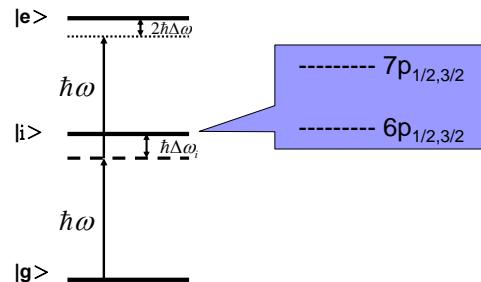
Measurement of the hyperfine structure of the  
6s - 7d( $J=3/2, 5/2$ )  
transitions by two-photon spectroscopy

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## Two-Photon Transition probability

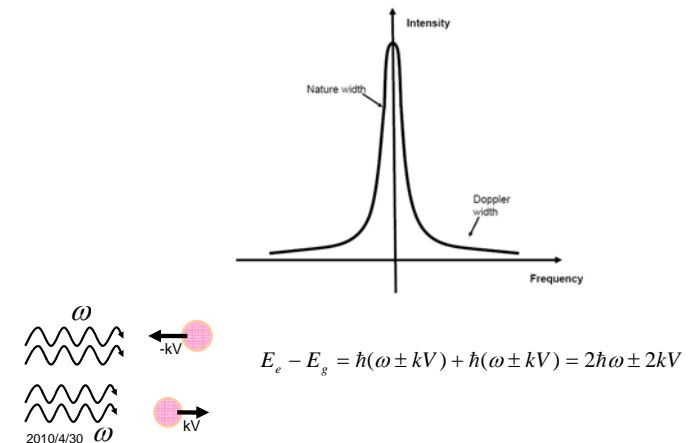
$$P_{ge}^{(2)}(\Delta\omega) = \left| \sum_i \frac{\langle e | H_1 | i \rangle \langle i | H_2 | g \rangle + \langle e | H_2 | i \rangle \langle i | H_1 | g \rangle}{\Delta\omega_i} \right|^2 \frac{\Gamma_e}{4(\Delta\omega)^2 + \Gamma_e^2/4}$$



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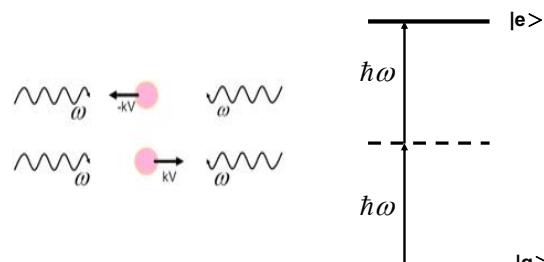
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## Two-Photon Transition/ Doppler Wings



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## Two-Photon Transition/ Doppler Free

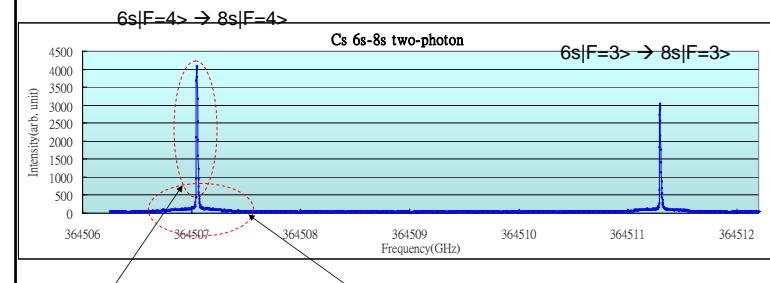


$$E_e - E_g = \hbar(\omega + kV) + \hbar(\omega - kV) = 2\hbar\omega$$

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## Two-Photon Transition/ Expt spectrum

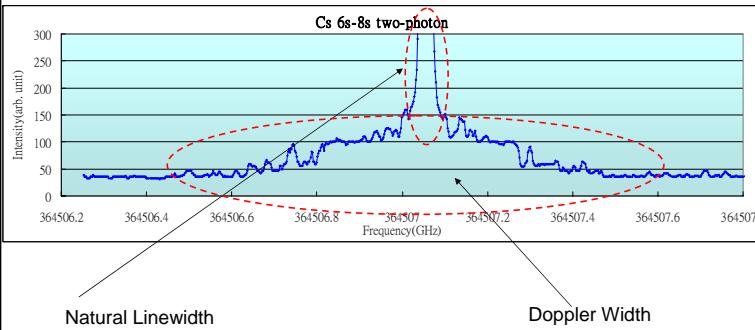


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## Two-Photon Transition/ Expt spectrum

$6s|F=4\rangle \rightarrow 8s|F=4\rangle$

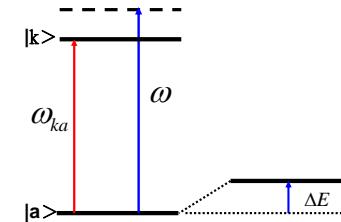


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## Light Shift or AC Stark Effect

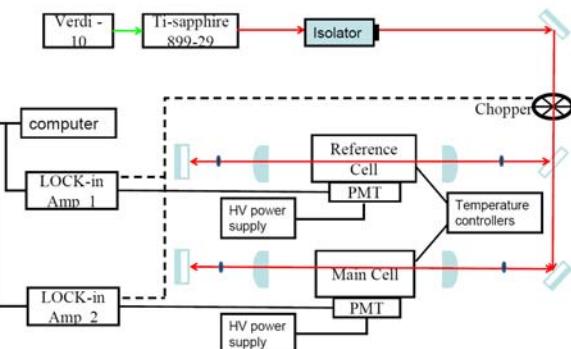
$$\overline{\Delta E_a} = -\frac{E_0^2}{2\hbar} \sum_{k \neq a} \left| \langle k | D_z | a \rangle \right|^2 \frac{\omega_{ka}}{\omega_{ka}^2 - \omega^2}$$



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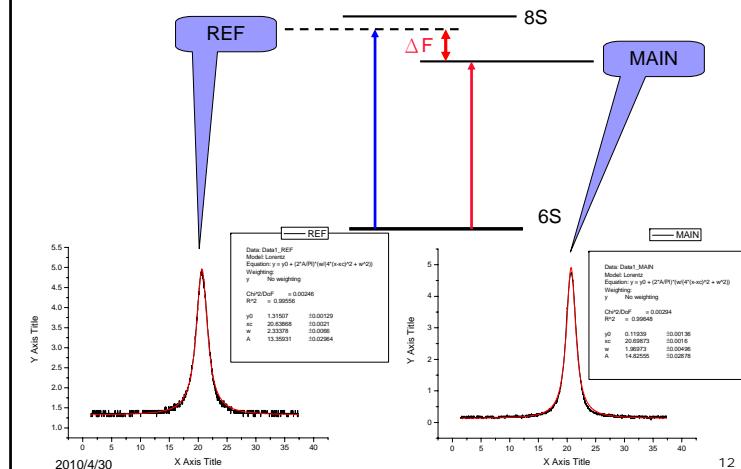
## Experimental Setup



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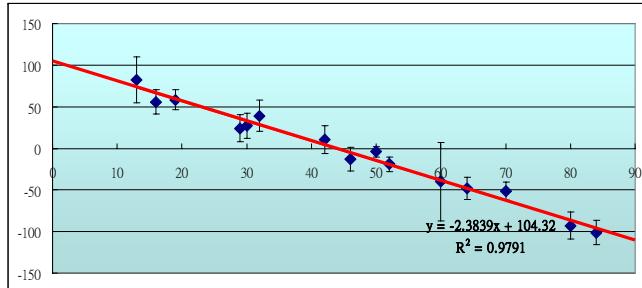
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## Light Shift Measurement



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## Light Shift Measurement



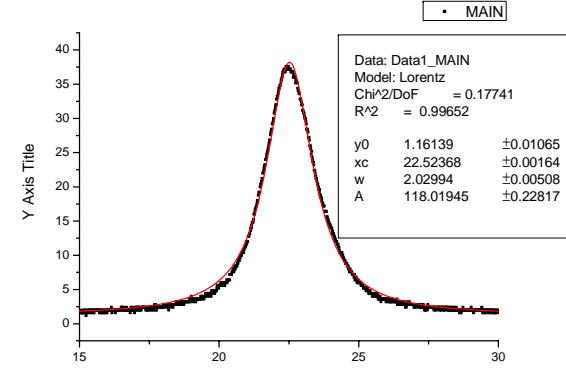
5 measurements on each data point for giving laser power.

$$\text{Light shift for Cs } 8S \text{ level is } -8.6 \frac{\text{Hz}}{\text{mW/mm}^2}$$

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## Natural Linewidth Determination

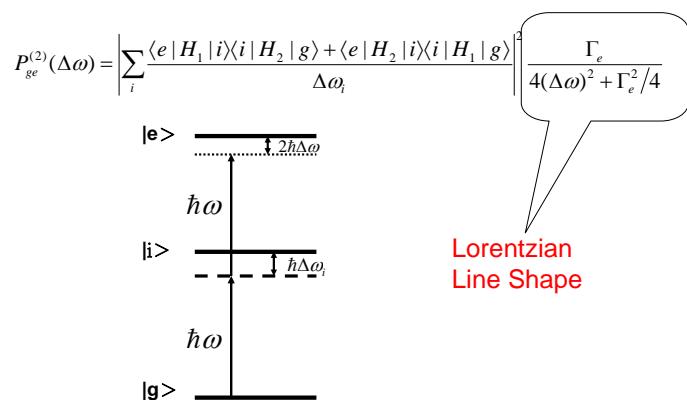


Fit with Lorentzian

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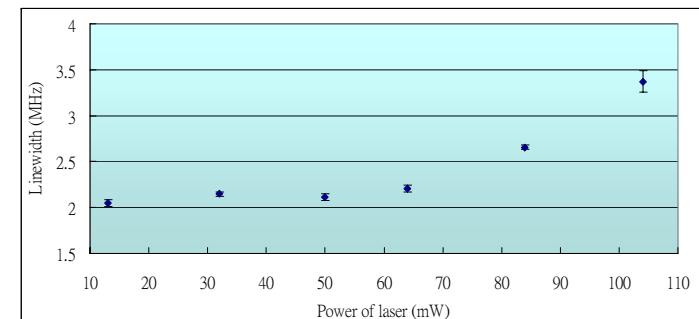
## Two-Photon Transition in a cell



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## Natural Linewidth Determination

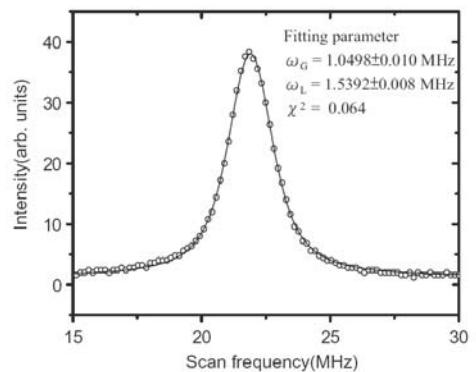


Fit with Lorentzian (5 measurements per data point).  
Power Broadening, Laser linewidth, Collision Broadening...?

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## Natural Linewidth Determination/Fit with Voigt

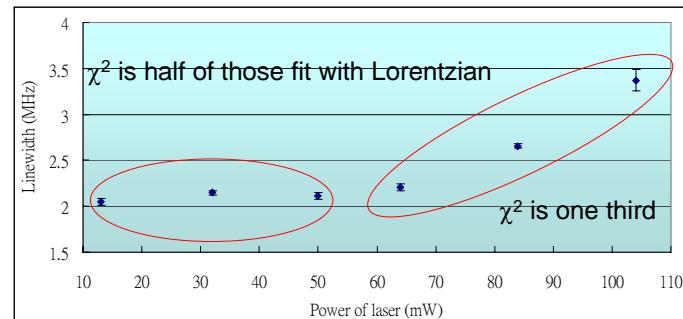


The observed spectrum is fitted with a Voigt Distribution to separate the Lorentzian width (Natural linewidth) and Gaussian width (other broadenings).

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## Natural Linewidth Determination



Fit with Lorentzian (5 measurements per data point).  
 Power Broadening, Laser linewidth, Collision Broadening...?

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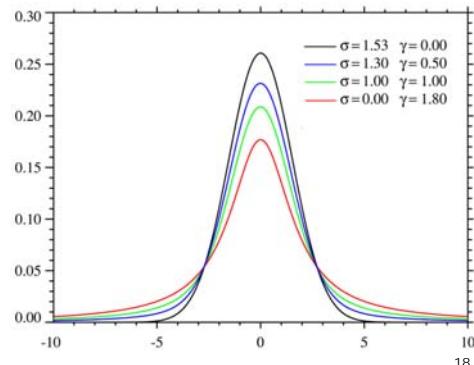
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## Voigt profile

$$V(x; \sigma, \gamma) = \int_{-\infty}^{\infty} G(x'; \sigma) L(x - x'; \gamma) dx'$$

$$L(x; \gamma) \equiv \frac{\gamma}{\pi(x^2 + \gamma^2)}.$$

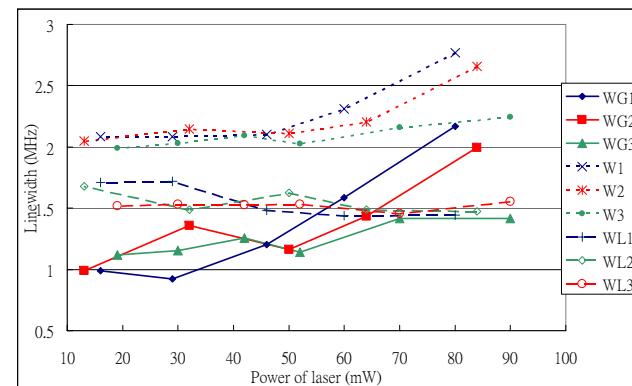
$$G(x; \sigma) \equiv \frac{e^{-x^2/(2\sigma^2)}}{\sigma\sqrt{2\pi}}$$



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## Natural Linewidth Determination

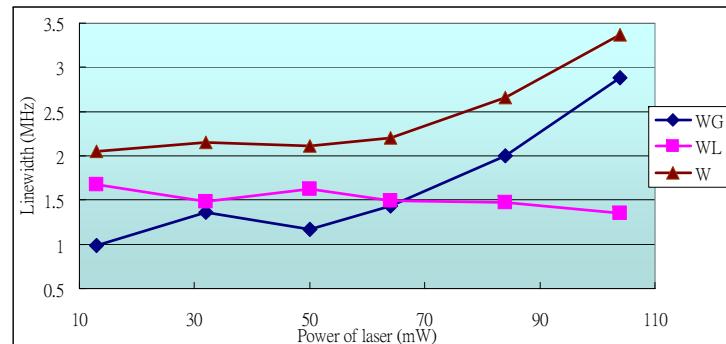


Natural Linewidth (Lorentzian) is 1.51 MHz

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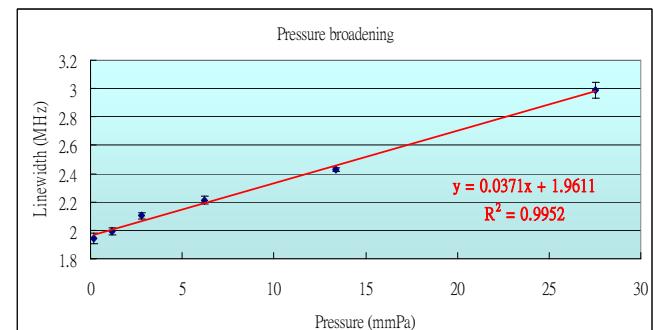
## Natural Linewidth Determination



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## Pressure broadening

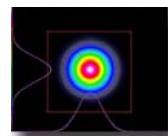


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## Power Broadening

A steady state of two-level system:

$$S_{\text{steady}} \propto \langle P_2 \rangle = \frac{1}{2} \frac{(I/I_{\text{sat}})}{1 + (\Delta/\beta)^2 + (I/I_{\text{sat}})}$$

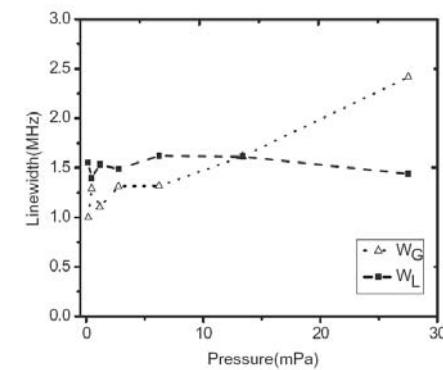


Special mode of laser is Gaussian

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## Pressure broadening/Voigt fit



Collision broadening should be Lorentzian???

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## Pressure (Collision )broadening

ening. Although the collision broadening is a homogeneous broadening, however, we increased the temperature to change the vapor pressure but also the velocity of atoms. The increasing of collision rate depends on the atom density and velocity [7]. The velocity dependence will arise the contribution to Gaussian broadening. Under the range of our temperature setting, Fig. 6 shows that the

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## Pressure (Collision )broadening

(e) The line shapes of two-photon transitions are very simple, as they are simple Lorentzian curves, whereas the line shape in the saturation technique is quite complicated (its calculation involves the averaging of a nonlinear effect that depends on the velocity component  $v_z$ ). If collisions are taking place, the two-photon line shape remains Lorentzian and it is easy to measure the broadening and the shift, whereas the velocity-changing collisions complicate still more the saturation line shape.

(f) The finite transit time of the atoms through the laser beam produces analogous broadenings in the two techniques. The large light beams (10 cm or more) used in the saturation technique to reduce this broadening in very narrow molecular transitions seems not to be possible with two-photon transitions because the energy density would be too small. But it is possible to use the Ramsey fringes technique (Baklanov *et al.* 1976; Salour *et al.* 1977).

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## Pressure (Collision )broadening

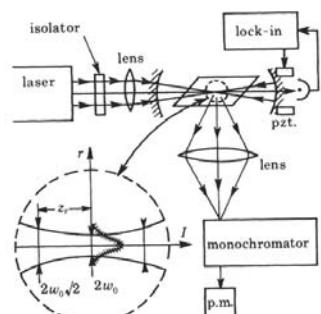


FIGURE 7. Experimental set-up for Doppler-free two-photon spectroscopy. Inside the dotted circle is represented the waist of the laser Gaussian beam which replaces the focus of geometrical optics: the superimposed curve with small bars represents the light intensity  $I$  as a function the radial distance  $r$ .

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## Pressure (Collision )broadening

- (i) natural broadening (which (involves OR affects )the lifetime of the excited level)
- (ii) collisions (pressure broadening)
- (iii) the finite interaction time (transit-time broadening, mainly in beam experiments, and (unimportant OR not important ) here)
- (iv) the second-order Doppler effect
- (v) instrumental width (laser bandwidth)
- (vi) external Fields (Zeeman and Stark effects)
- (vii) residual Doppler broadening (if the beams are not exactly counter-propagating),
- (viii) power broadening (related to the saturation of the transition)
- (ix) the A.C. Stark effect (a shift caused by the electric field of light in the TPT)[20].

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## Cs 8s natural Linewidth

The second-order Doppler effect is approximately 162 Hz

$$\Delta f_{D2} \square \frac{u^2}{c^2} f_0 = \left( \frac{200 \text{m/s}}{3 \times 10^8 \text{m/s}} \right)^2 \frac{3 \times 10^8 \text{m/s}}{822 \times 10^9 \text{m}} = 162 \text{Hz}$$

Ti:sapphire laser (Coherent 899-29) used in this study has a linewidth of  $\Delta v_{RMS} = 500 \text{kHz}$   
 data fitting yields a standard deviation of approximately 80 kHz  
 Error arises from the laser frequency drift during the scan through a 2 MHz spectrum 10 kHz

Lorentzian width (WL, 1.53 MHz)  
 convolution of the 500 kHz laser bandwidth and Errors  
 natural linewidth of 1.03 (+20)MHz.

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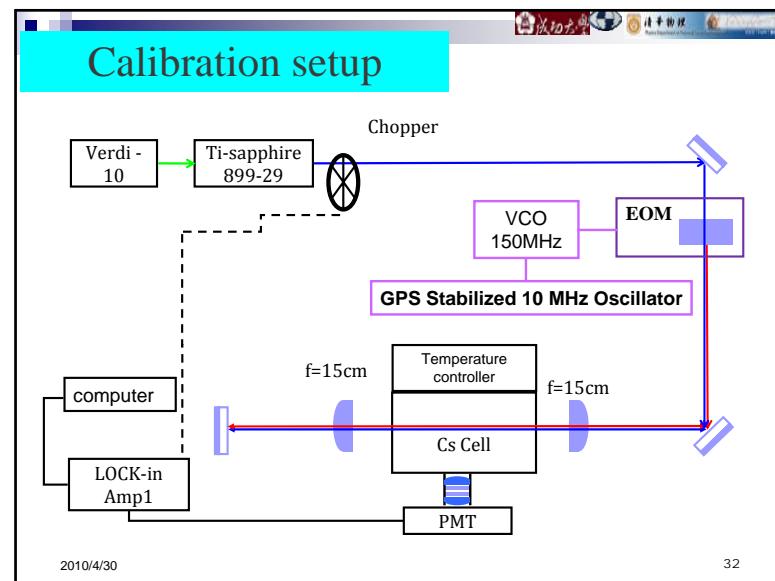
## Energy levels of 6s-7d 2-photon transitions

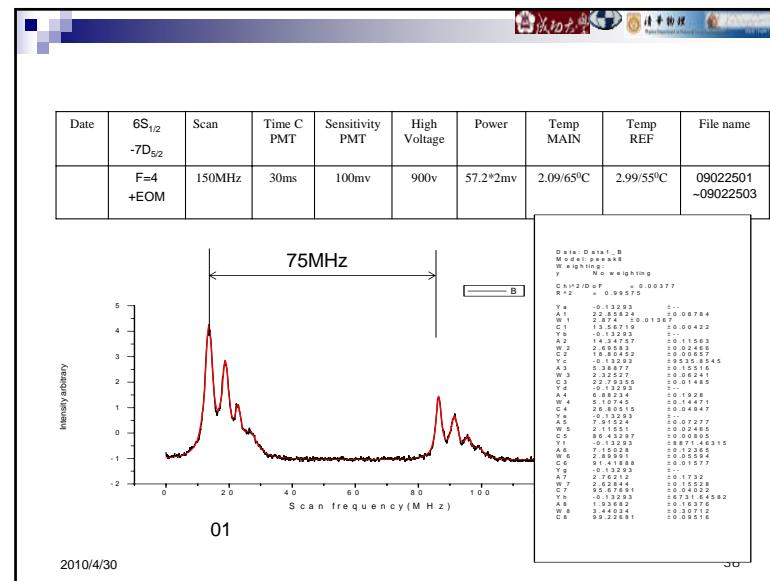
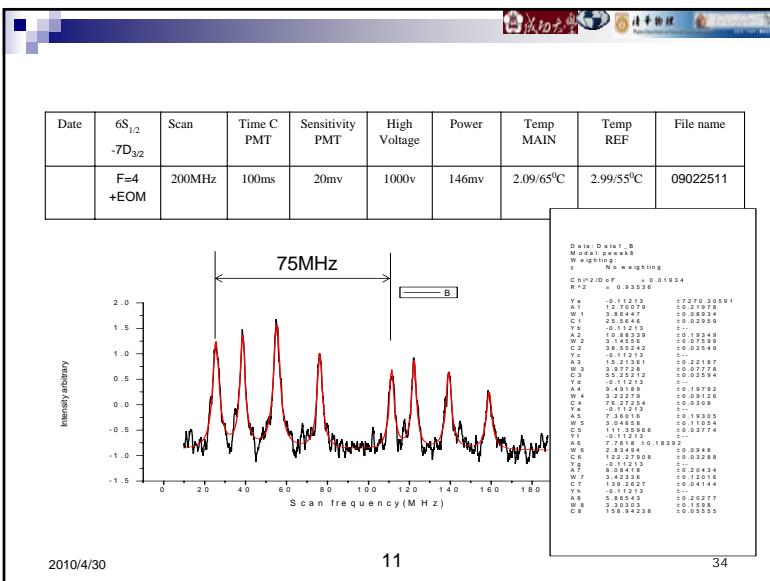
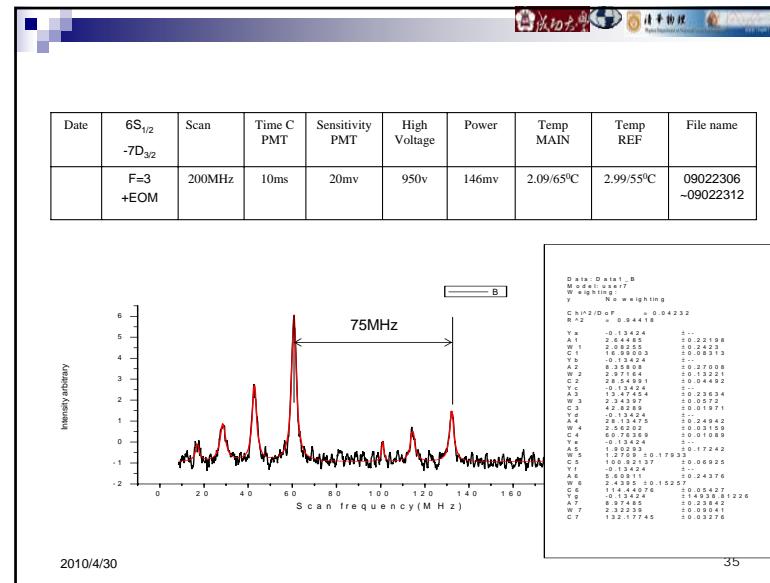
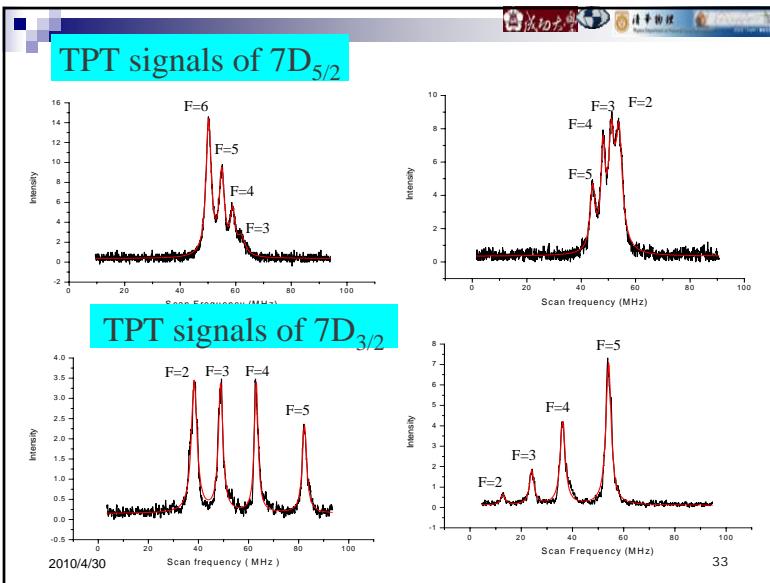
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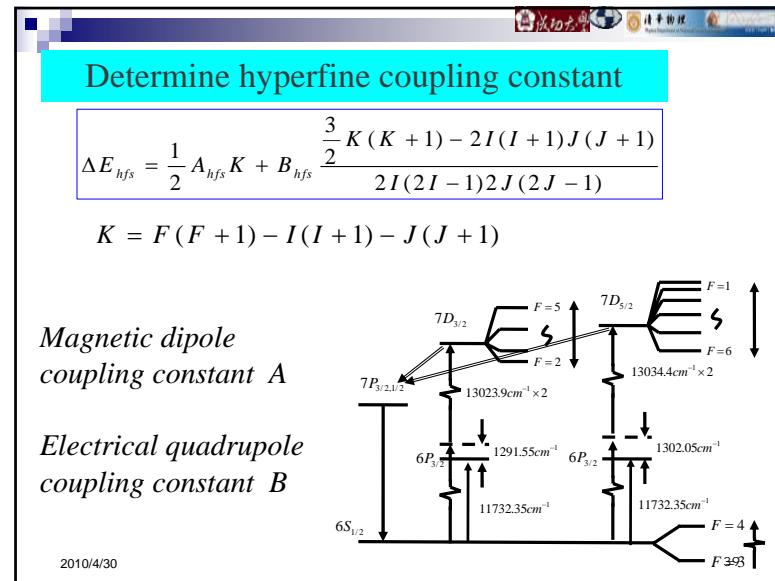
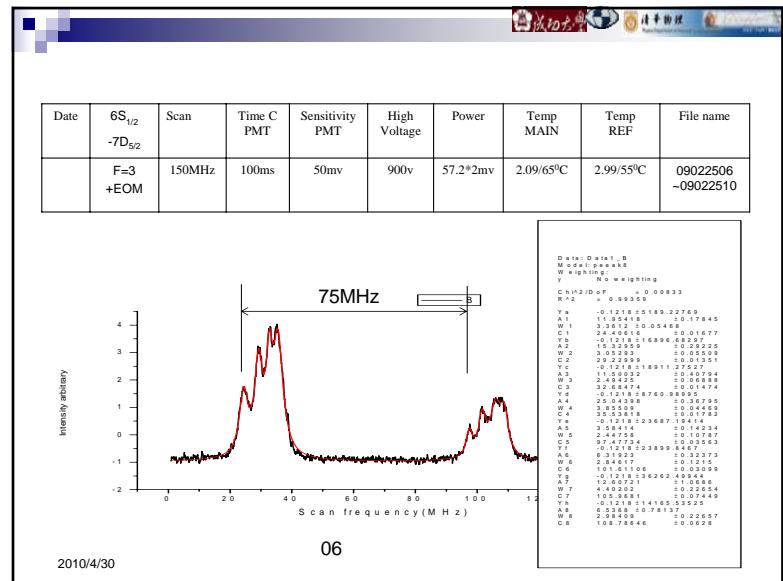
## Two-Photon Spectroscopy

Measurement of the hyperfine structure of the 6s - 7d(J=3/2, 5/2) transitions by two-photon spectroscopy

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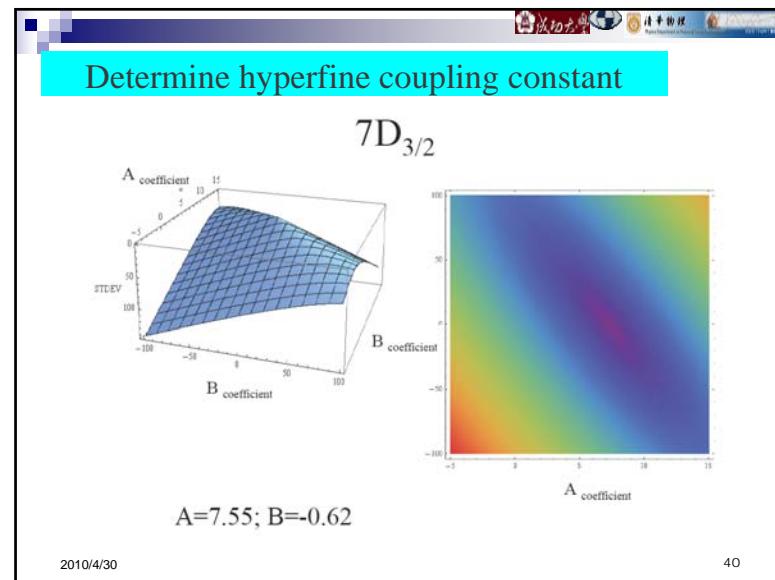


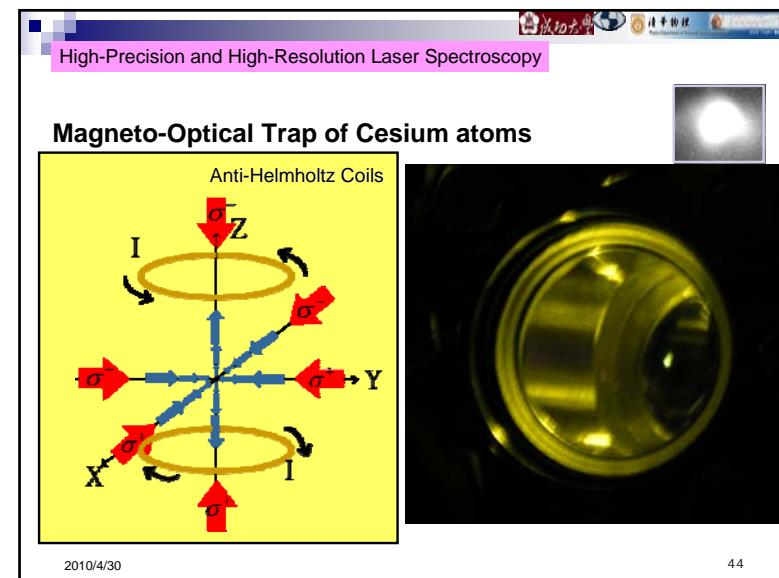
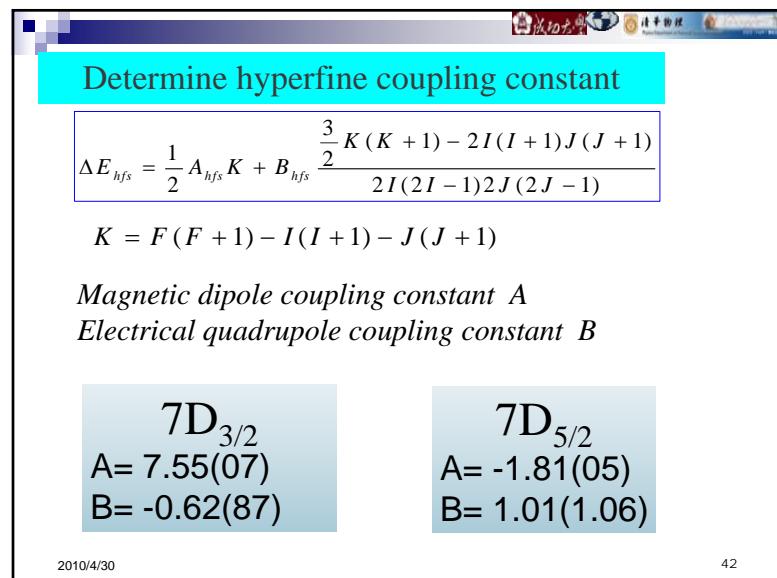
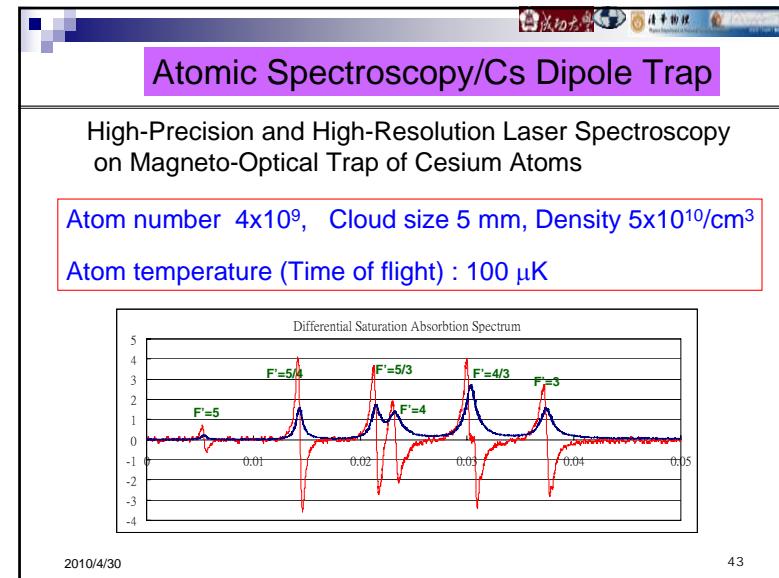
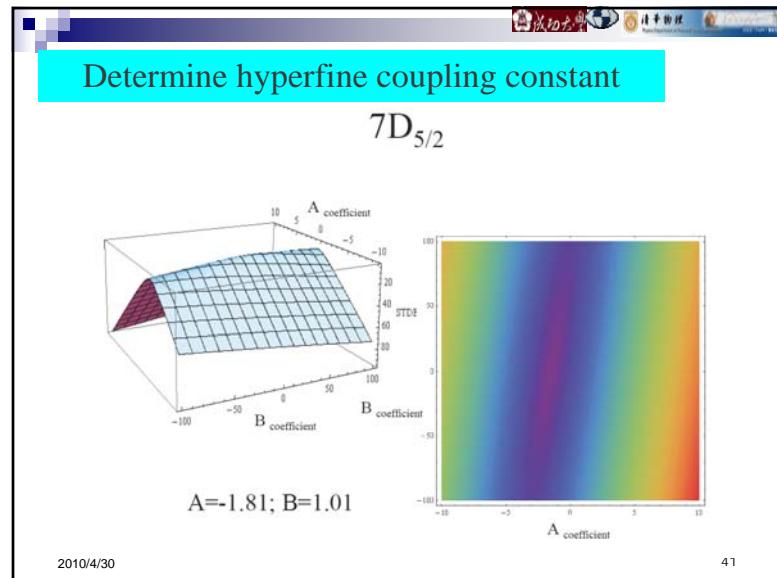


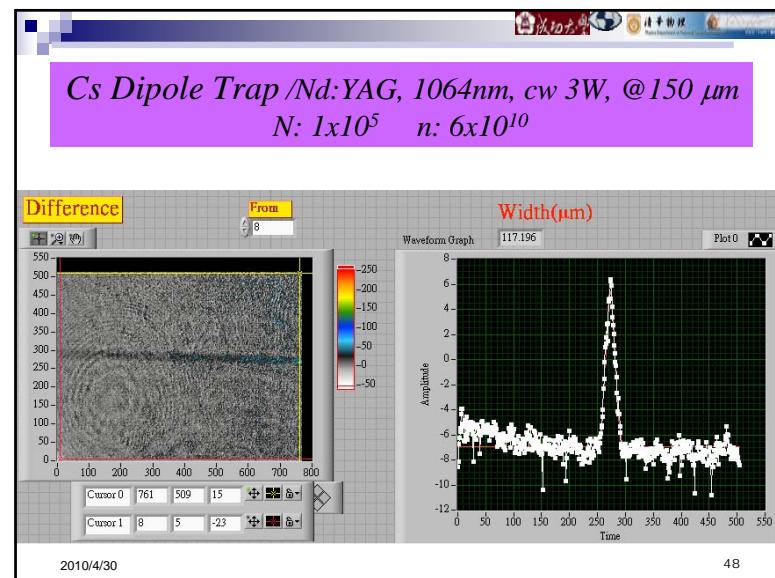
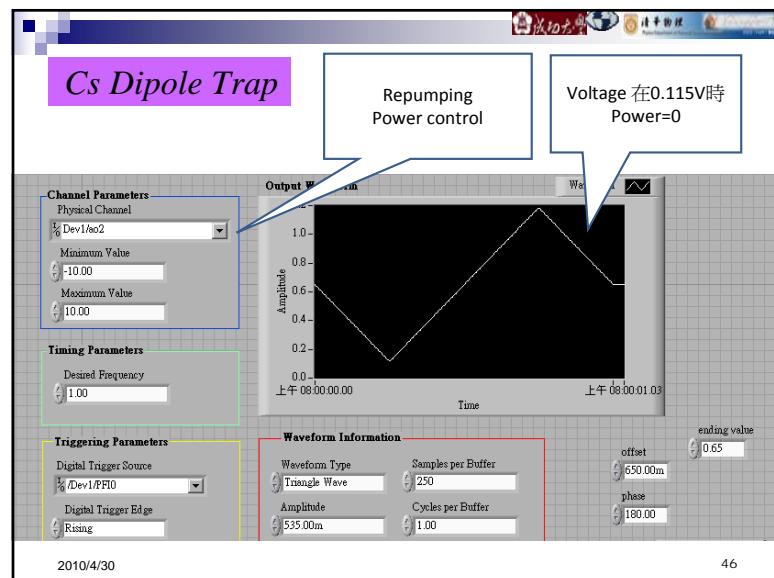
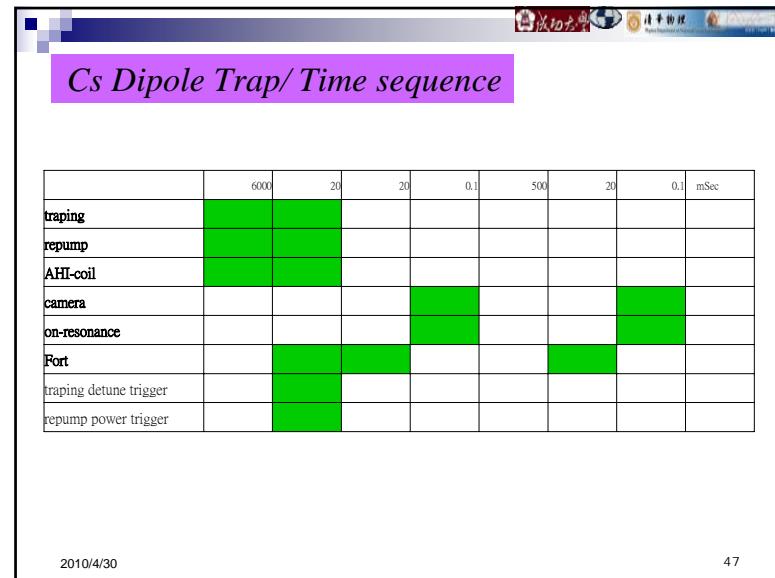
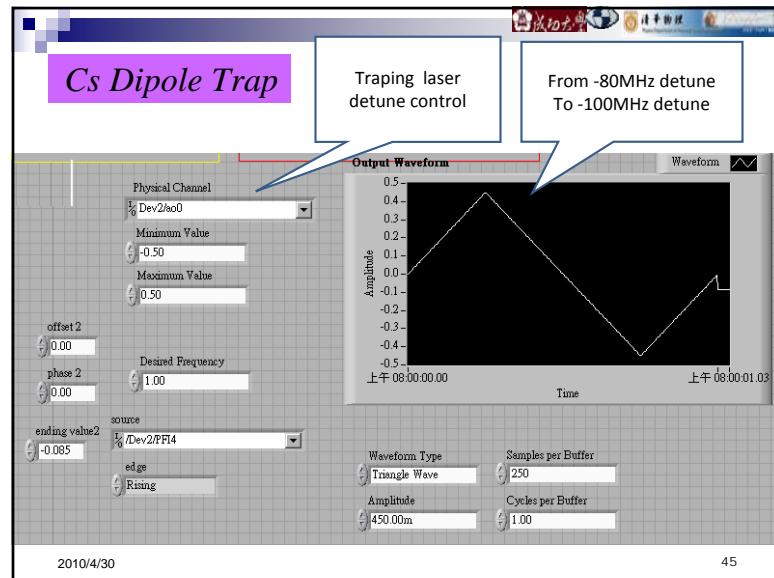
EOM	STDEV(+)	STDEV(-)	Reference	
7D(5/2) F=4	A=-1.77 B=-0.73	A=-1.78 B=0.36	A=-1.78±0.01 B=0.36±0.93	
w/o △F6~6, only F=6 ~ 5 ~ 4	A=-1.78 B=-0.13	A=-1.78 B=0.32	A=-1.78±0.05 B=0.32±0.4	
F=3	A=-1.88 B=0.86	A=-1.73 B=3.47	A=-1.63±0.15 B=3.47±3	
w/o △F5~5, only F=5 ~ 4 ~ 3	A=-1.83 B=1.88	A=-1.74 B=3.01	A=-1.63±0.1 B=3.01±2	
Reference				
7D(3/2) F=4	A=7.6 B=1.85	A=7.35 B=-0.17	A=7.05 B=2.01	A=7.35±0.3 B=0.17±2
w/o △F5~5, only F=3 ~ 4 ~ 5	A=7.5 B=1.41	A=7.31 B=-0.89	A=7.11 B=1.32	A=7.31±0.2 B=0.899±1.5
F=3	A=8.38 B=-16.73	A=8.14 B=-15.07	A=7.89 B=-13.3	A=8.14±0.3 B=-15.07±2
w/o △F3~3, only F=3 ~ 4 ~ 5	A=8.29 B=-15.94	A=8.17 B=-15.11	A=8.04 B=-14.18	A=8.17±0.2 B=-15.11±1

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## *Thanks for your attention!*

Collaboration: 黃守仁老師, Chemistry, NCKU  
Postdoctoral : 李益志博士  
PhD Students: 何宗勳  
And many many master students.....

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我們的實驗室出現在【日本料理】中的帥哥們…  
還有平面模特兒



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