



Interactions Between Pairs Of Cs Rydberg Atoms



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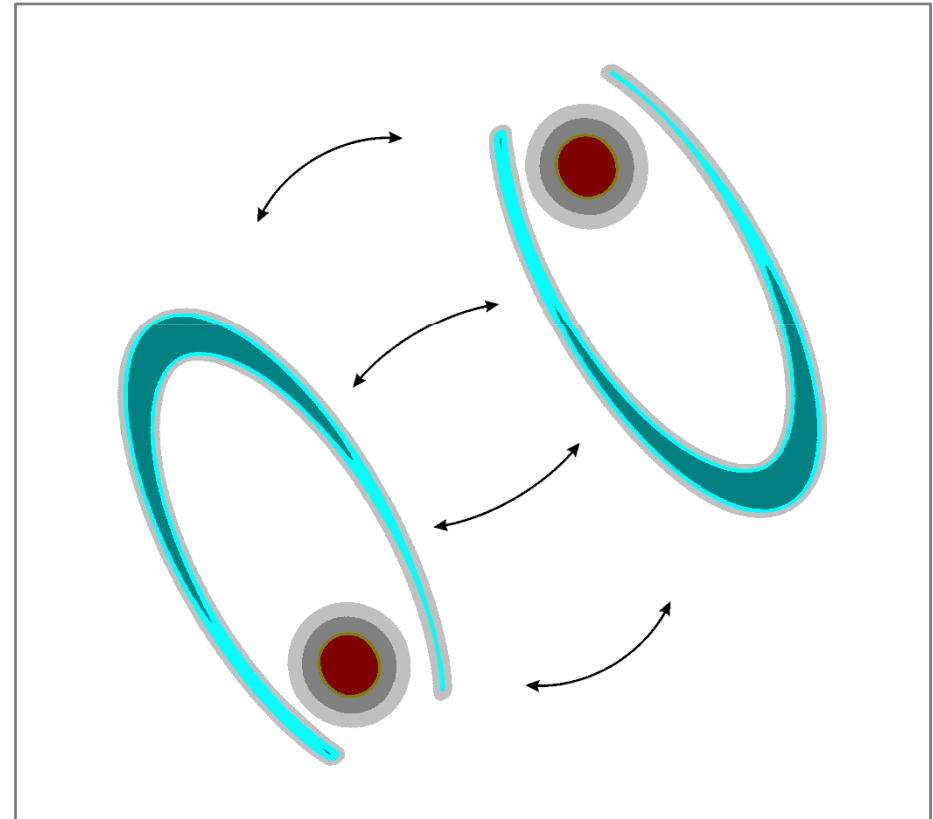
Outline

- Pair Interactions
- Experimental Apparatus
- Time-of-Flight Velocity Distributions
- Photo-Initiated Collision Measurement
- Macrodimers Measurements



Rydberg Atom Pair Interactions

- Interesting for a variety of reasons
 - Resonant energy transfer
 - Dipole Blockade
 - Exotic states of matter
 - *Macrodimers*
- Requires detailed knowledge of Pair interaction potentials





Calculations by Matrix Diagonalization

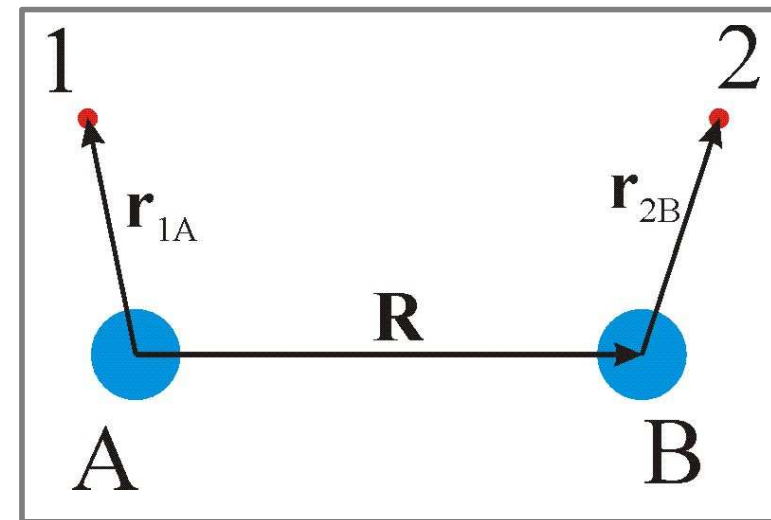
- Includes dipole and quadrupole contributions
- Diagonalized in the Stark shifted basis with $\mathbf{E} \parallel \mathbf{R}$
– *Thanks Arne!*

$$V(\mathbf{R}, \mathbf{r}_{1A}, \mathbf{r}_{2B}) = \sum_{L_1, L_2=1}^N \sum_{M=-L}^L \frac{(-1)^{L_2} f_{L_1 L_2 M}}{R^{L_1+L_2+1}} Q_{L_1 M}(\mathbf{r}_{1A}) Q_{L_2 - M}(\mathbf{r}_{2B})$$

where the multipole operator is

$$Q_{LM}(\mathbf{r}) = \left\{ \frac{4\pi}{2L+1} \right\}^{1/2} r^L Y_{LM}(\hat{r})$$

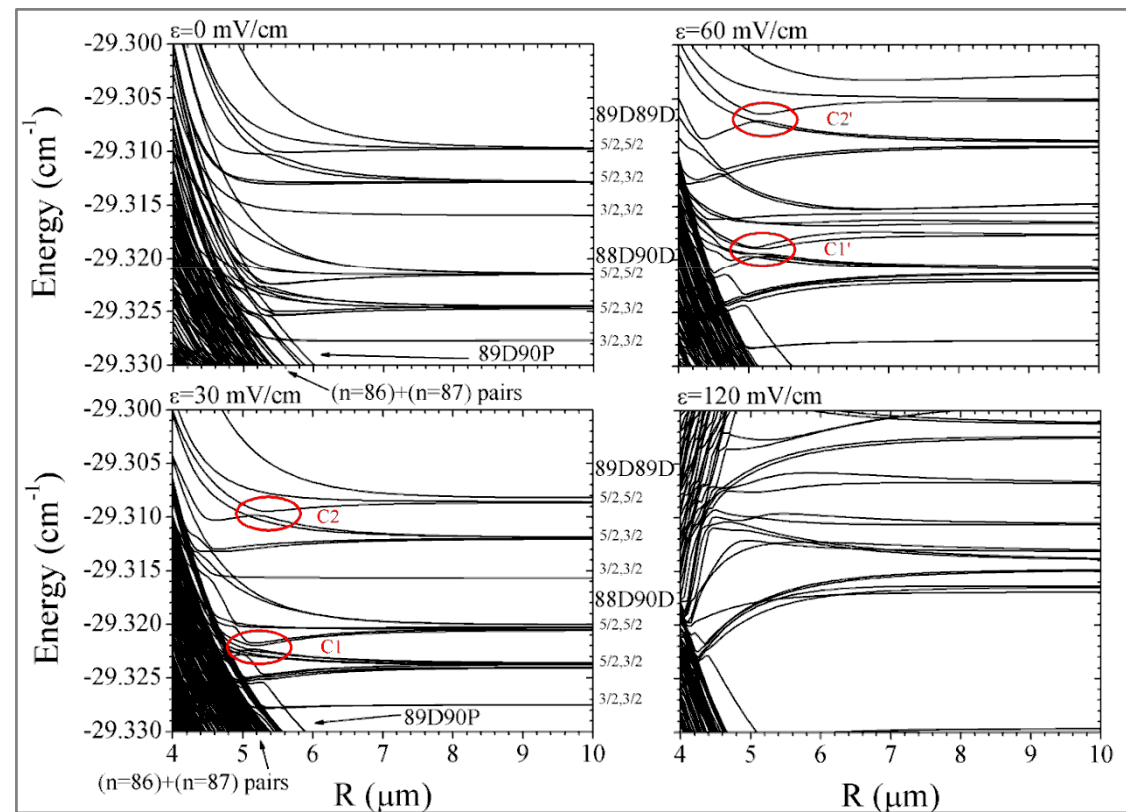
and $f_{L_1 L_2 M} = \frac{(L_1 + L_2)!}{[(L_1 + M)!(L_1 - M)!(L_2 + M)!(L_2 - M)!]^2}$



M. R. Flannery, D. Vrinceanu, and V. N. Ostrovsky, J. Phys. B **38**, S279 (2005)

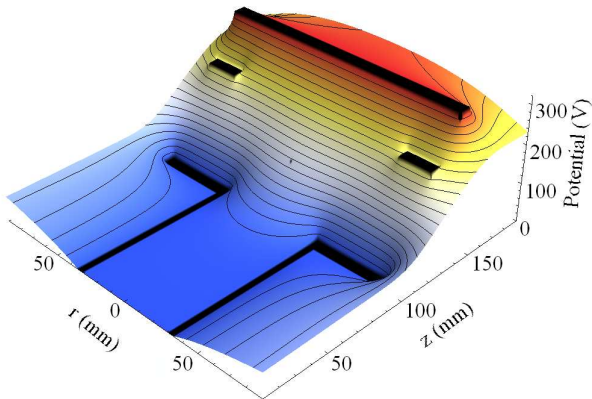
E-Fields and Avoided Crossings

- Electric fields have a strong influence on avoided crossings
 - Existence of wells depends on E
 - Pairs may be bound or dissociative
 - *Photoinitiated (PI) Collisions*
 - *Macrodimers*

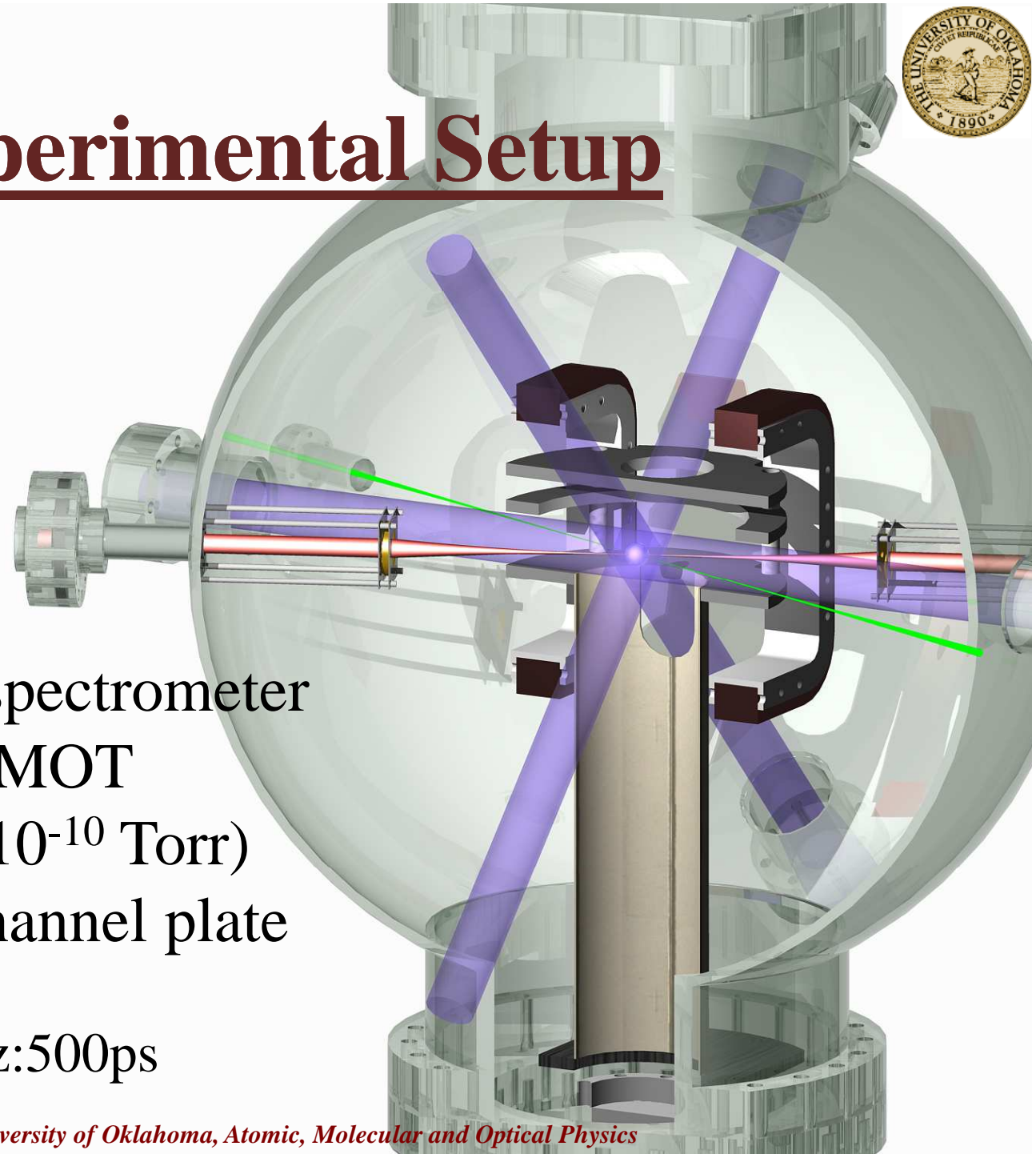




Experimental Setup

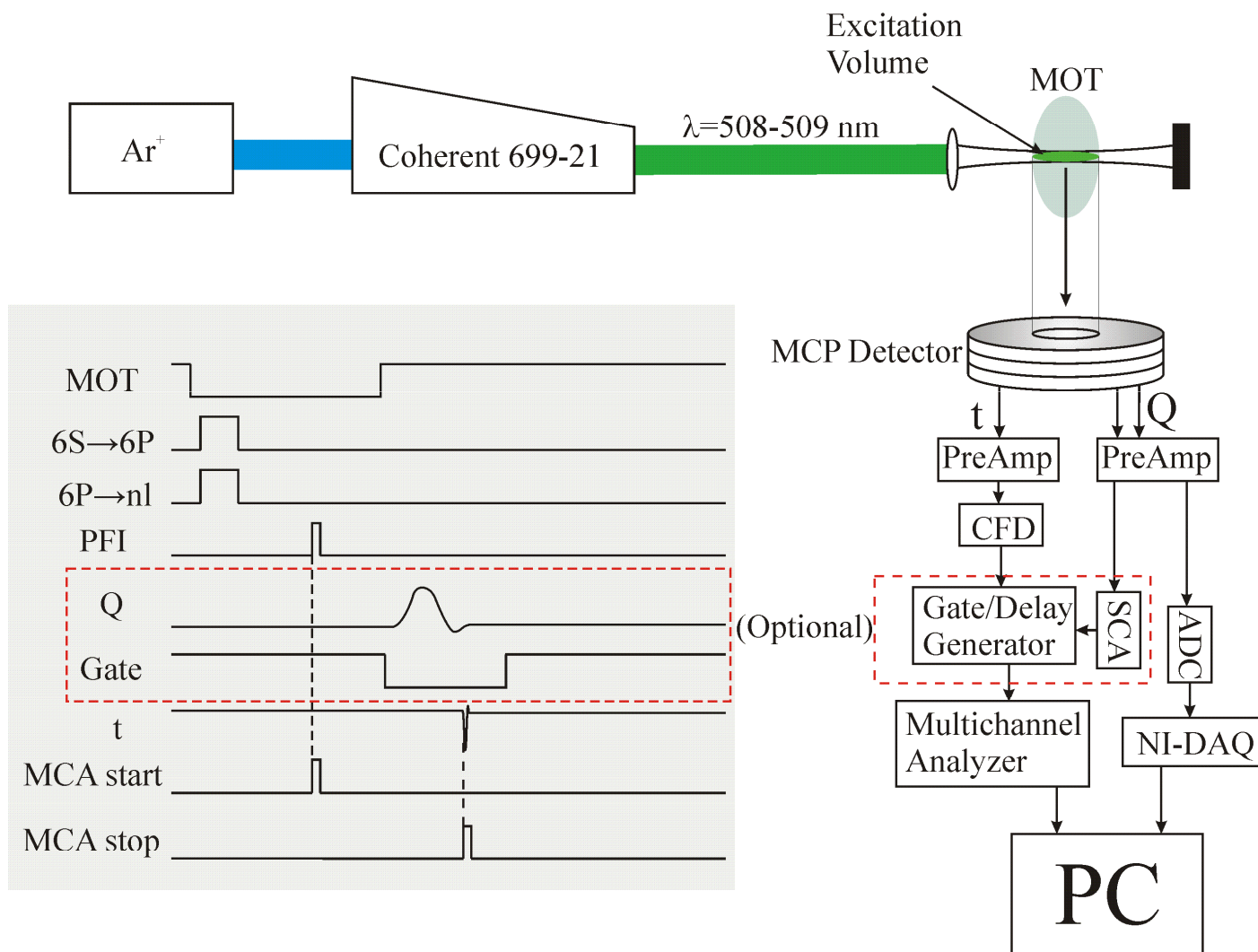


- Time-of-Flight spectrometer
- Centered on Cs MOT
- UHV system ($\sim 10^{-10}$ Torr)
- Z-Stack microchannel plate detector
 - res. x-y: $20\mu\text{m}$, z: 500ps





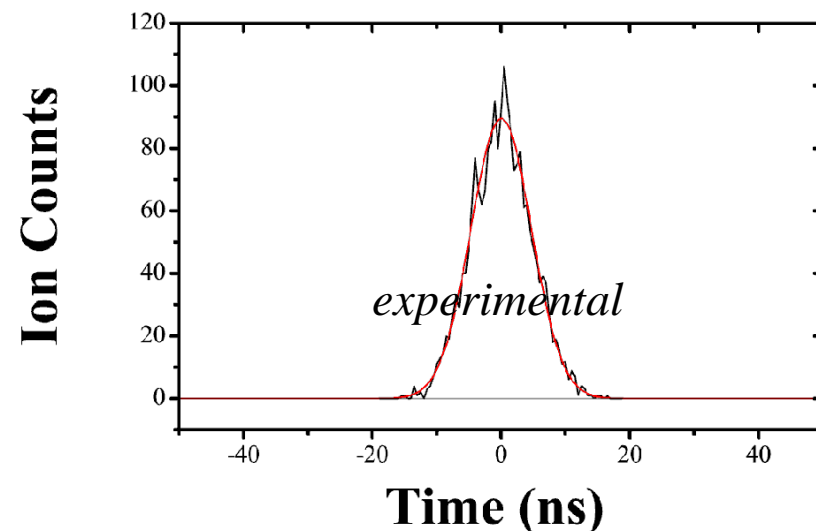
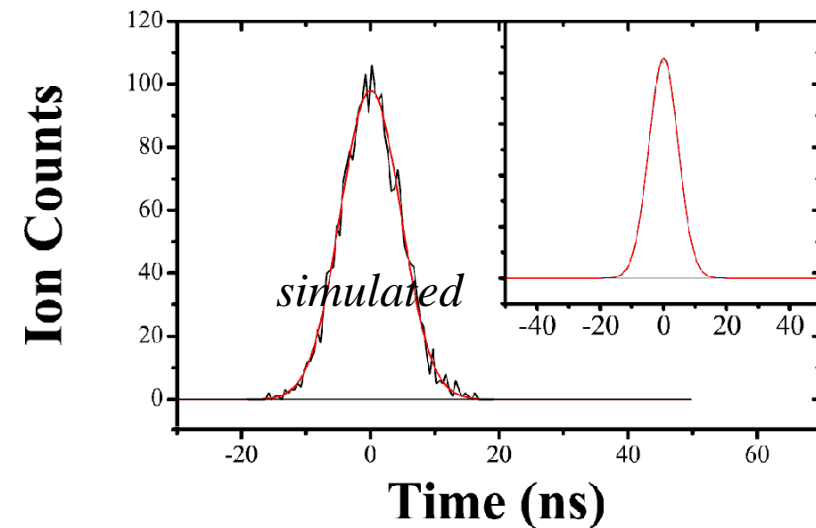
Signals and Timing





Time-of-Flight Distributions

- Expansion of TOF distribution depends on:
 - *Thermal* velocity
 - Recoil velocity determined by *collision* exit channel
 - *Coulomb repulsion*





Temperature Measurement

- Resolution of spectrometer calibrated using thermal expansion

- Gaussian distribution

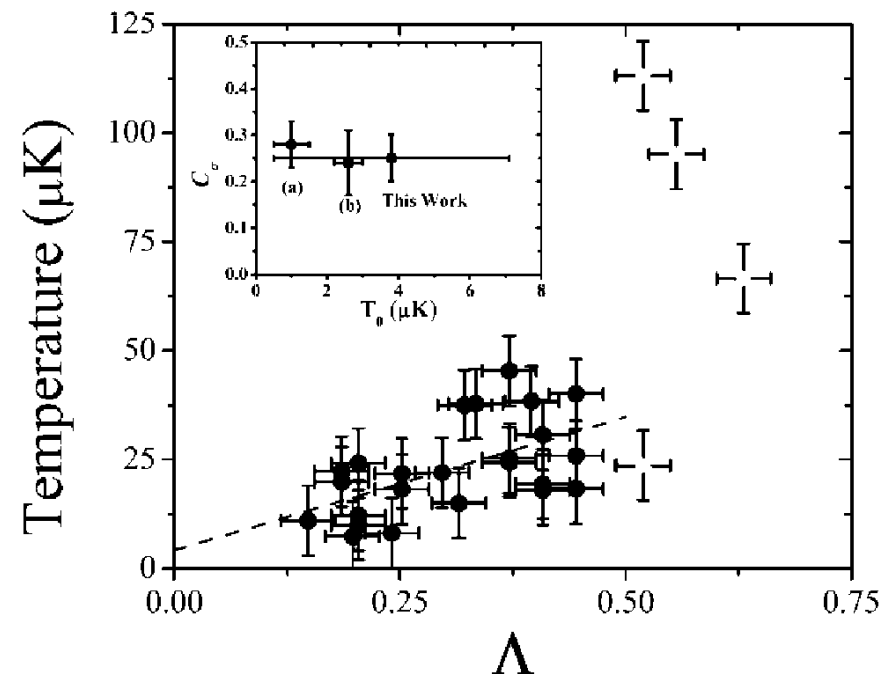
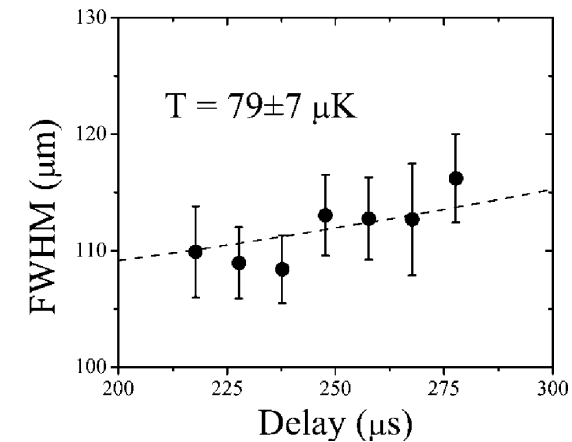
$$f(z, t) \propto e^{-mz^2 4 \ln 2 / \Delta z^2}$$

$$\Delta z = \sqrt{\Delta z_0^2 + \frac{8 \ln 2 k_B T}{m} \tau^2}$$

- Velocity resolution of **2.5 cm/s**

- Light shift parameter
 $\Lambda = \Omega^2 / |\delta| \Gamma$

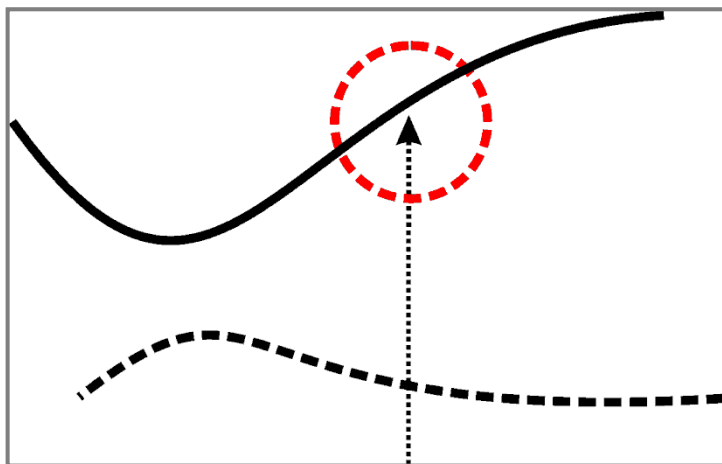
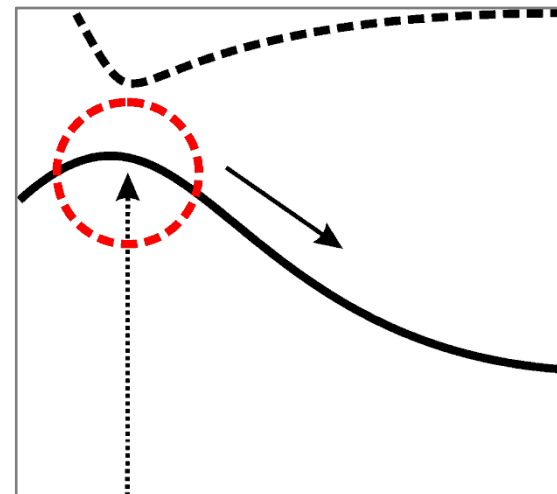
$$T = T_0 + 2 \times C_\sigma T_D \left(\frac{\Omega^2}{|\delta| \Gamma} \right)$$





PI Collisions vs. Macrodimers

- **PI collision** can occur from excitation at a stationary point
 - Collision products gain velocity v_{coll} determined by the energy of the exit channel

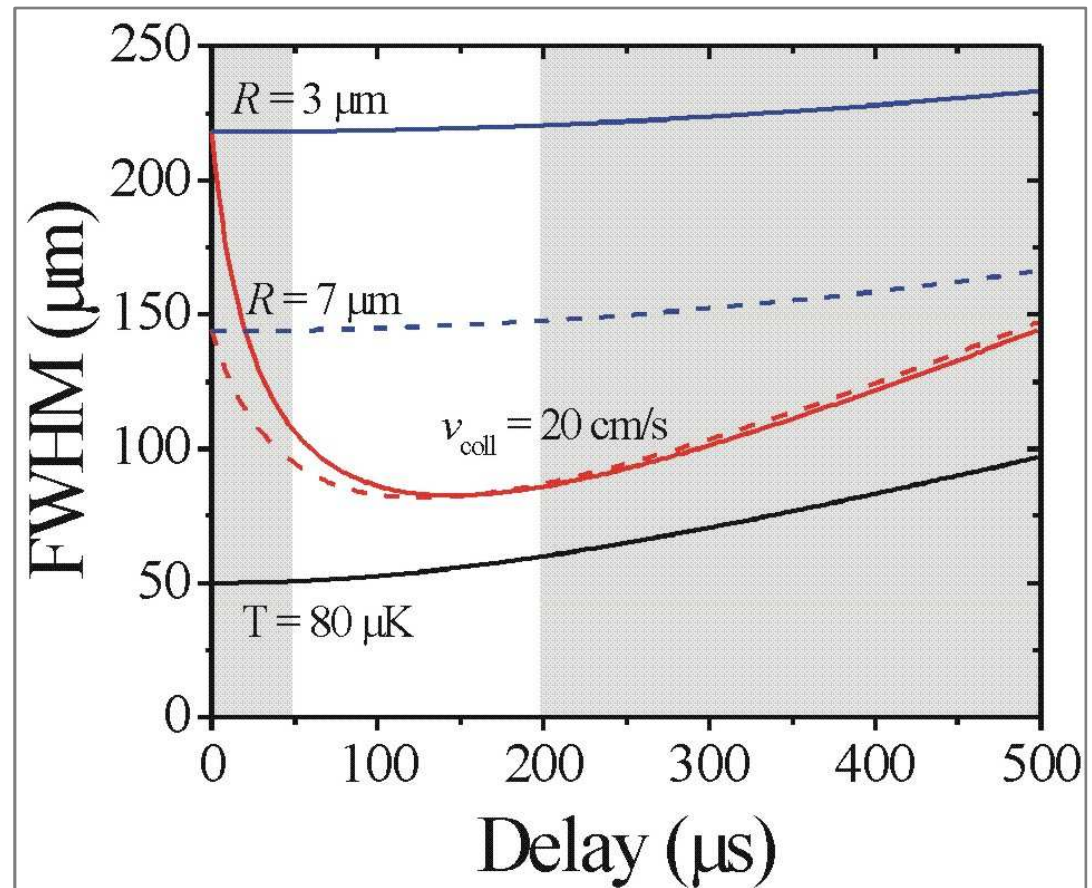


- **Macrodimers** can be excited in a well
 - Excitation with narrow band cw laser light causes R to have *fixed distribution*
 - *Vibrational period* ($\sim 2 \mu s$) < *excitation time* ($\sim 5 \mu s$)



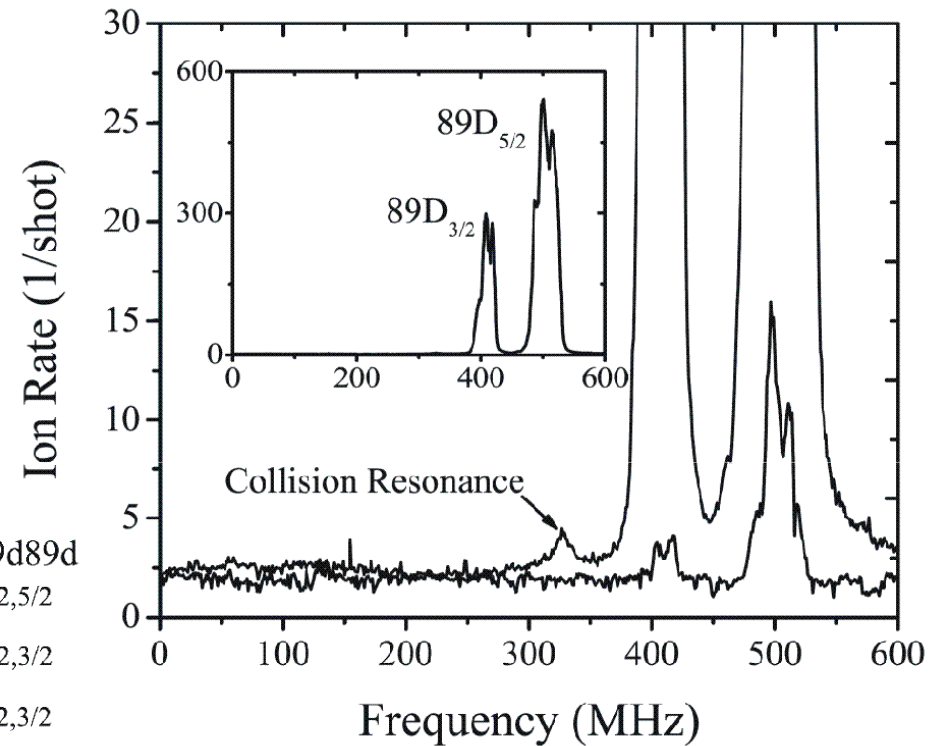
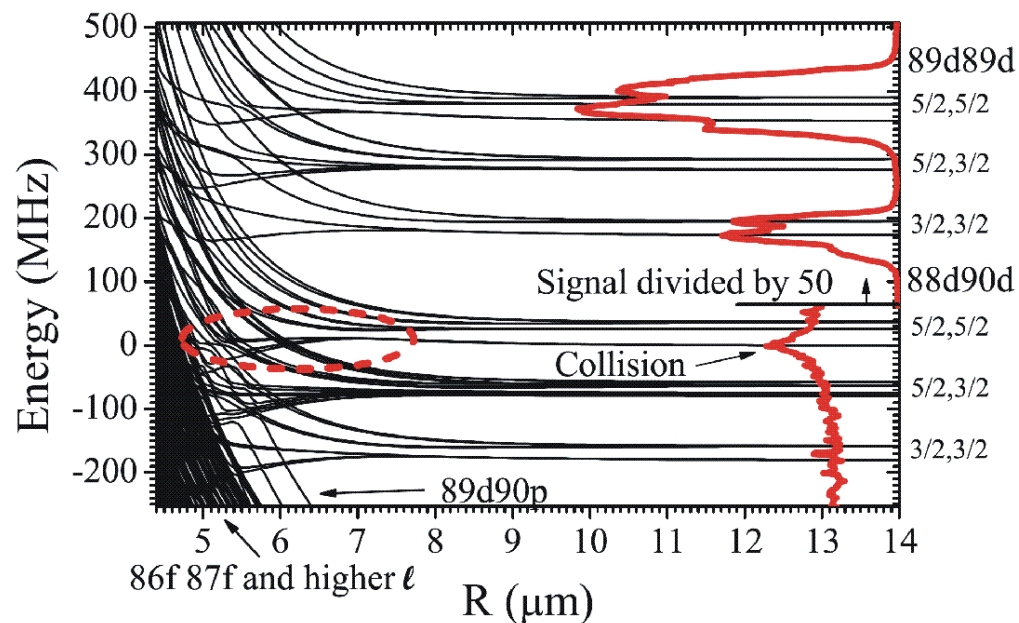
Delay Dependence of TOF Distributions

- Coulomb repulsion in TOF identifies pair interaction
 - Expansion at short delay can identify R as constant
 - Expansion at long delay is a direct measure of v_{coll} .



PI Collision Measurement

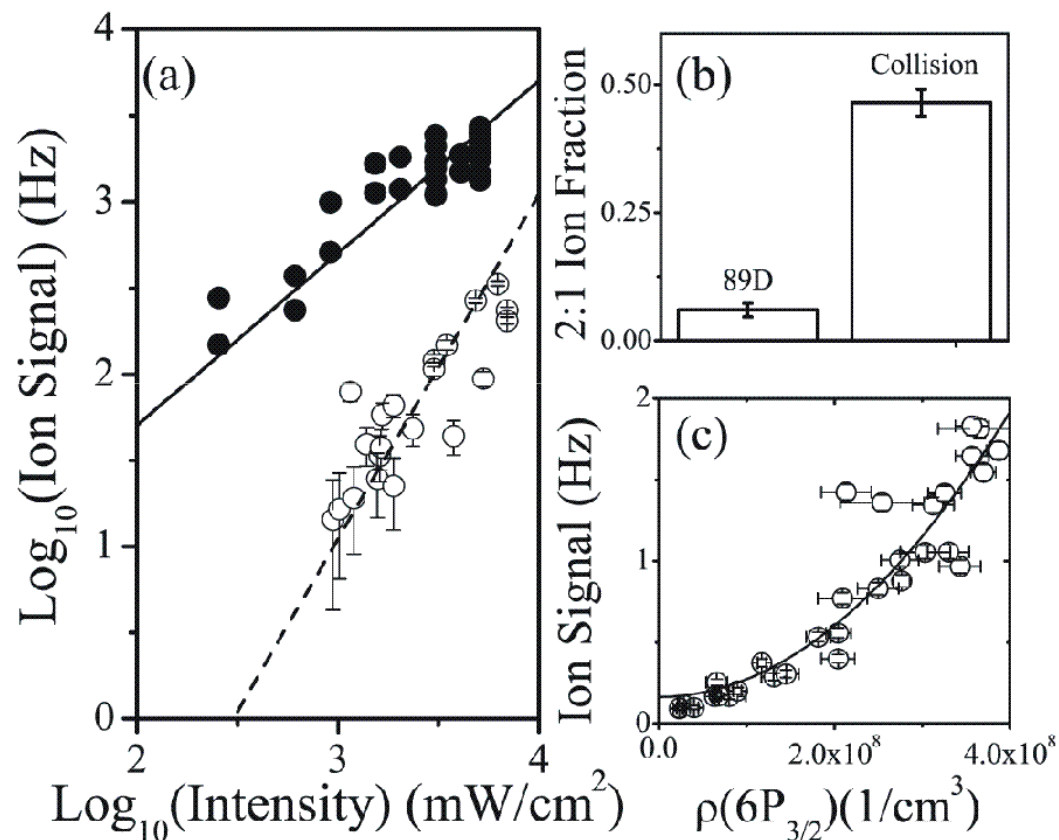
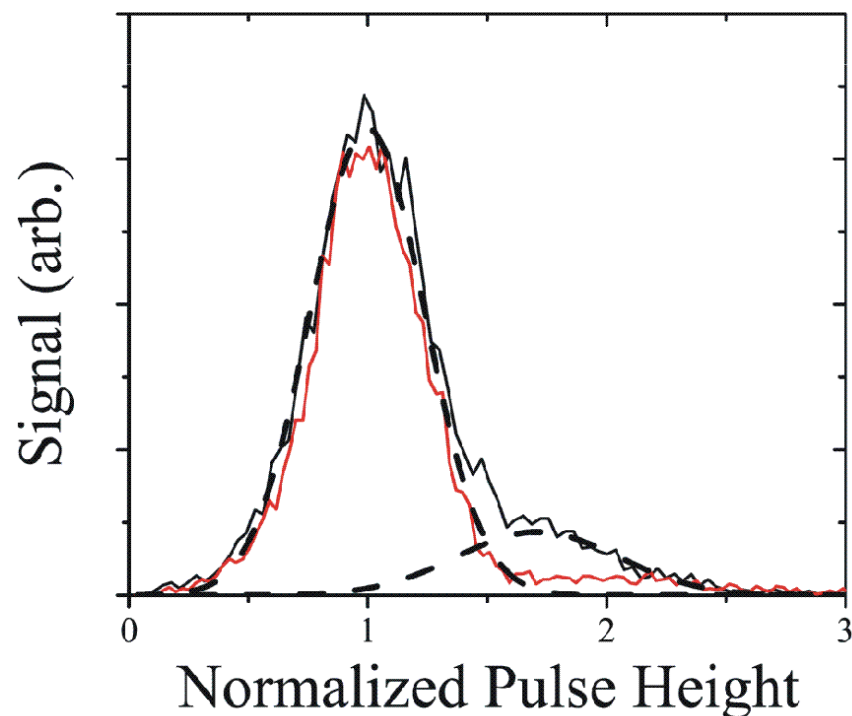
- Pair resonance observed near $89D+89D$
 - No prominent well





Pair Identification

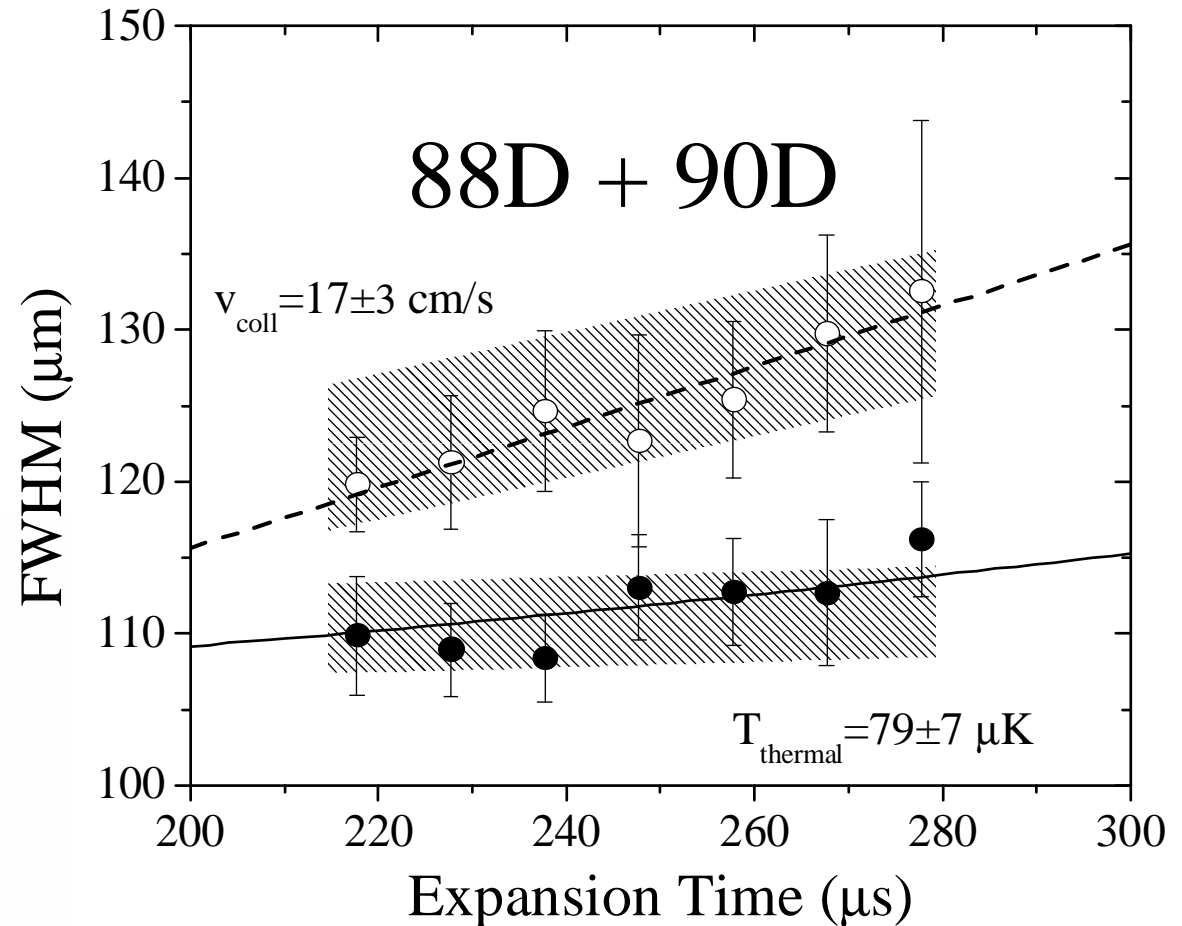
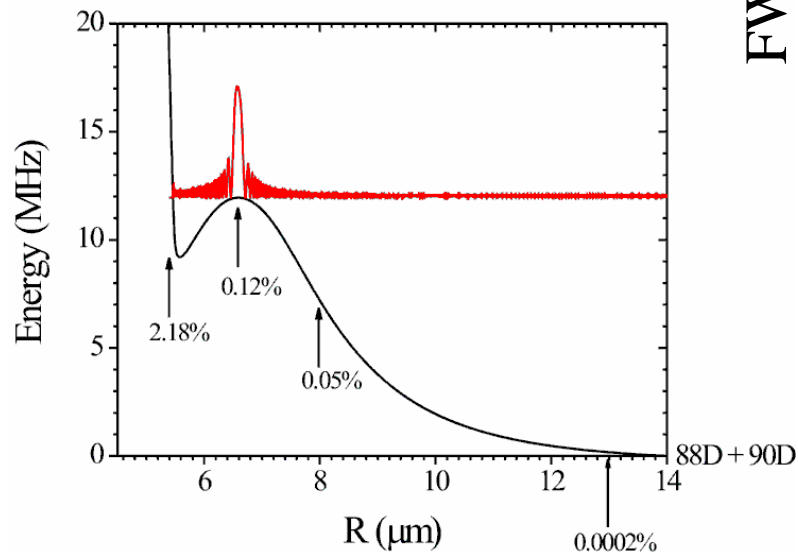
- (a) 2-photon for $6P \rightarrow 89D$
- (b) 2-atom from charge pulse height distribution
- (c) 2-photon $6S \rightarrow 6P$





Exit Velocity Measurements

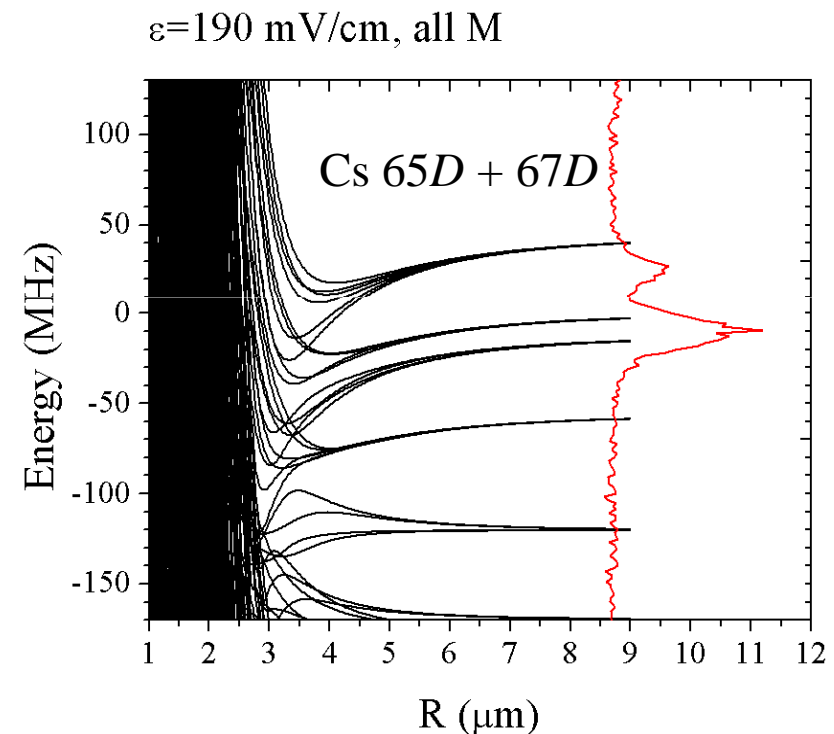
- Expansion the result of thermal and exit velocities
 - 88D + 90D resonance is a *collision* process.
 - Exit velocity of 17 ± 3 cm/s





Choosing n for Molecular States

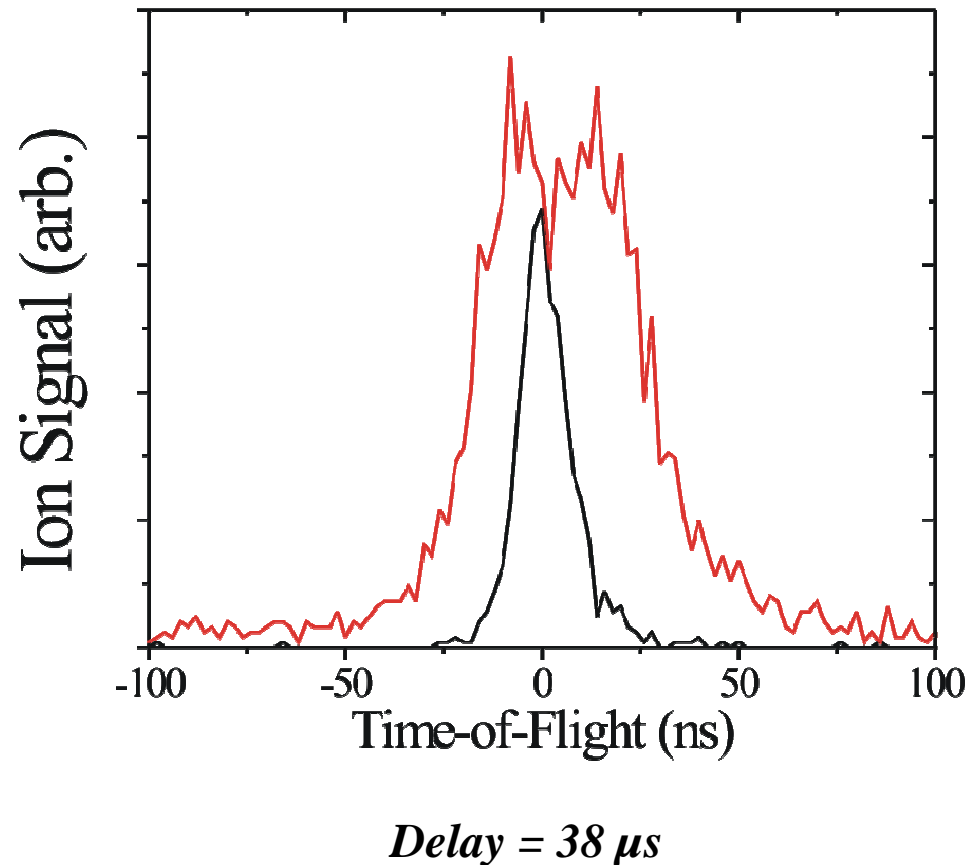
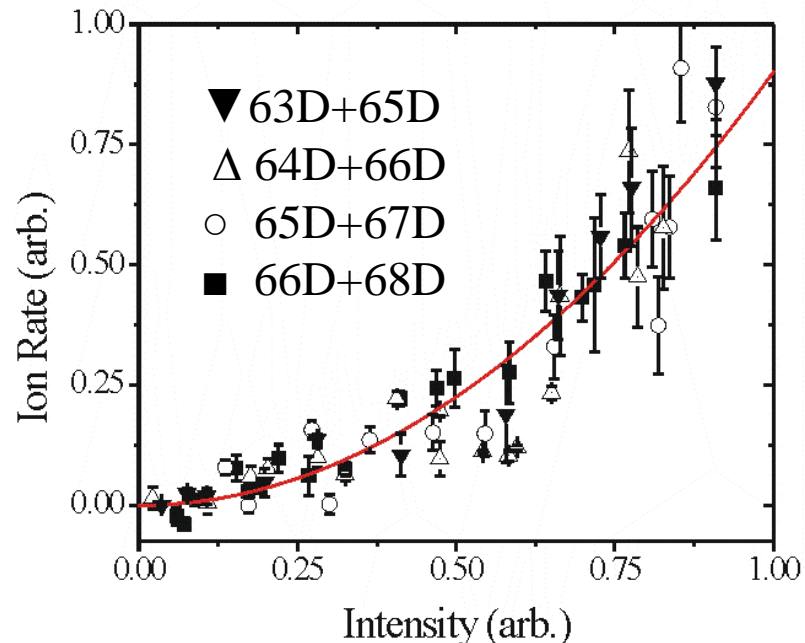
- Lower principal quantum number *advantageous* for measurements of Coulomb repulsion for bound states.
 - Lower $n \rightarrow$ resonances are farther from atomic lines
 - **Pros:**
 - Less atomic background signal
 - Deeper wells
 - **Con:**
 - Less oscillator strength





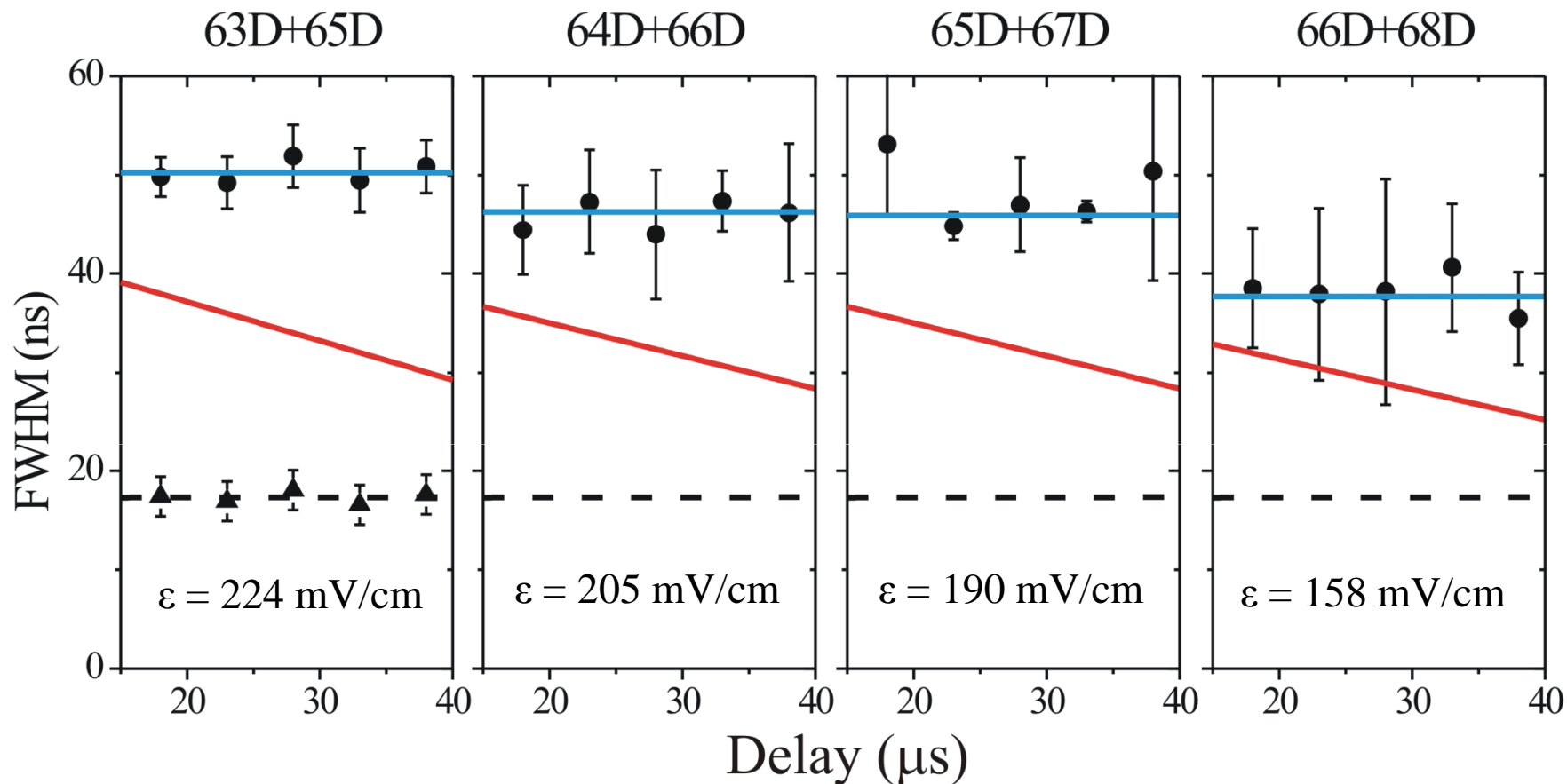
Ion Rates and TOF Velocity Distributions

- Ion rate is *quadratic*
- Coulomb repulsion easily resolved from TOF distributions
 - Black: atomic state
 - Red: molecular resonance





Width vs. Delay Measurements



- The red line is a *Monte Carlo* simulation of collision with thermal velocity recoil.
- Circles are molecule data
- Triangles are atomic data (dashed line is fit to thermal expansion)



Future Directions

- Investigate angular distribution of Macrodimers
 - 3D imaging to study applied E spatial dependence
- Measure macrodimer *lifetimes* by observing state distribution of products
- Perform detailed spectroscopy of wells
 - Electric field dependence of wells



The Group

- James Shaffer
 - Arne Schwettmann: contributed calculations of pair interactions
 - Jonathan Tallant: assisted with experiments
 - Donald Boothe: currently assisting Arne with calculations



Publications

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- Arne Schwettmann, Jack Franklin, K. Richard Overstreet, and James P. Shaffer, “Stark slowing asymmetric rotors: Weak-field-seeking states and nonadiabatic transitions” J. Chem. Phys. **123**, 194305 (2005)
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- J. Tallant, K. R. Overstreet, A. Schwettmann, and J. P. Shaffer, “Sub-Doppler magneto-optical trap temperatures measured using Rydberg tagging” Phys. Rev. A **74**, 023410 (2006)
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- K. Richard Overstreet, Arne Schwettmann, Jonathan Tallant, and James P. Shaffer, “Photoinitiated collisions between cold Cs Rydberg atoms” Phys. Rev. A **76**, 011403 (2007)
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- Arne Schwettmann, K. Richard Overstreet, Jonathan Tallant, and James P. Shaffer, “Analysis of long-range Cs Rydberg potential wells” J. Mod. Opt. **54**, 2551 (2007)
- K. R. Overstreet, A. Schwettmann, J. Tallant, D. Booth, and J. P. Shaffer, “Observation of Cs Rydberg atom macrodimers” (*in submission*)



Pair Excitation

- Near resonant 2-photon transition
 - Excitation rate increased due to proximity to nearest atomic duplicate pair state
 - Pair interact strongly at short R and mixes in nD state character
 - Excitation rate higher for higher n

