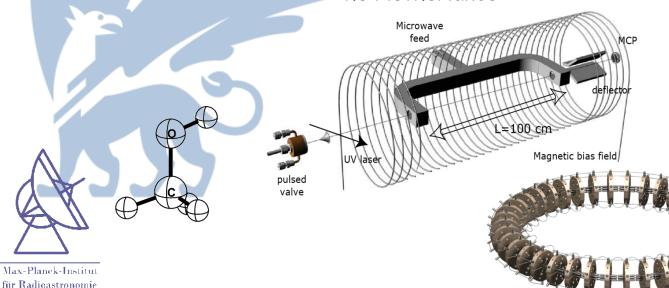
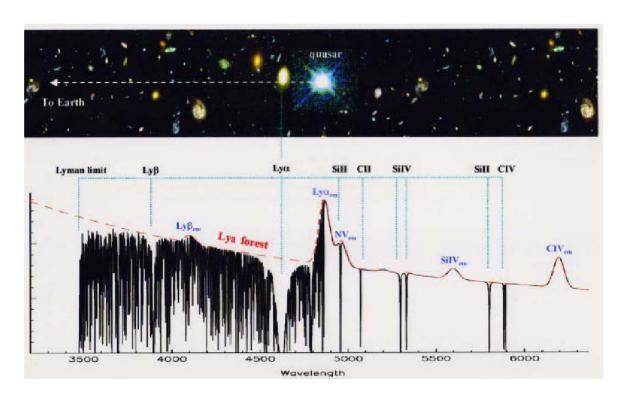
Probing time-variation of fundamental constants in cold and not so cold molecules.

Rick Bethlem

LaserLaB, Department of Physics and Astronomy
VU University
de Boelelaan 1081, 1081 HV Amsterdam
The Netherlands



varying a and μ from astronomical data



 $\Delta a/a = (-0.72 +/- 0.18) \times 10^{-5}$ over the last 12 Gyr [1]

If variation is linear: $0.6 \times 10^{-15}/\text{yr}$

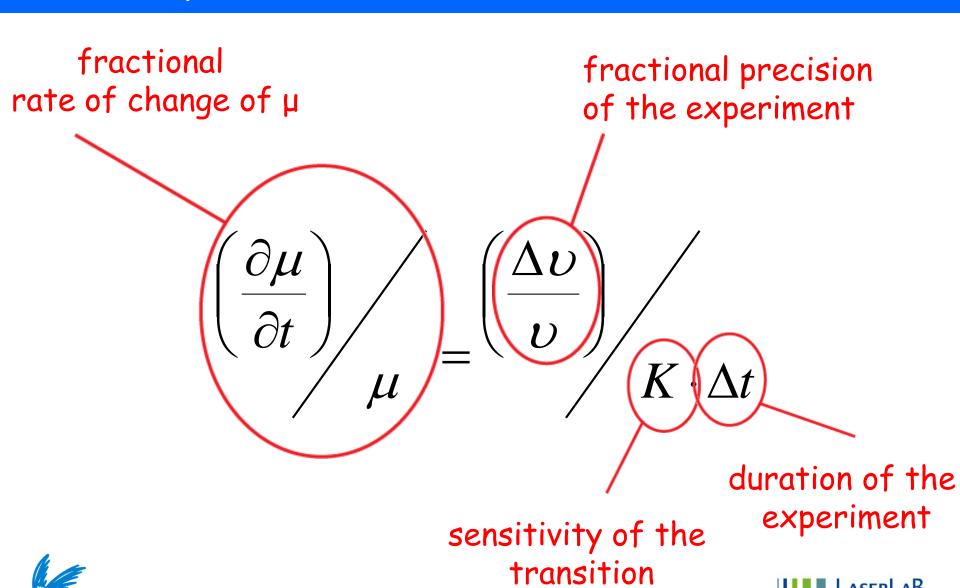
 $\Delta\mu/\mu = (2.4 + /- 0.6) \times 10^{-5}$ over the last 12 Gyr [2]

If variation is linear: $-2.0 \times 10^{-15}/yr$

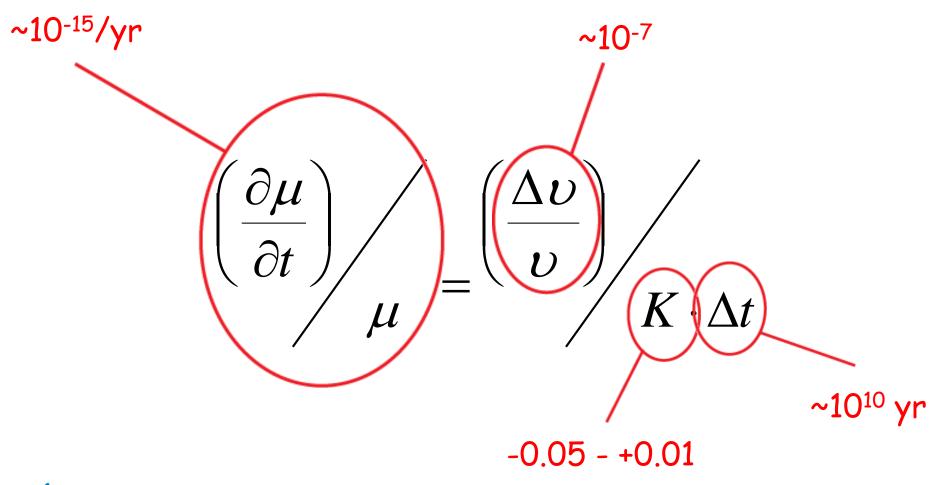




Ability to detect a time-variation of μ



Constraint from H₂ observations







Probing time-variation of μ using:

- Cold ammonia molecules in a molecular fountain. very high precision (10^{-12} ?), high sensitivity factor ($0 \leftrightarrow -4.2$), short time interval (2 yr).
- Not so cold CO molecules in a beam. High precision (10^{-11} ?), huge sensitivity factor ($-300 \leftarrow +200$), short time interval (2 yr).
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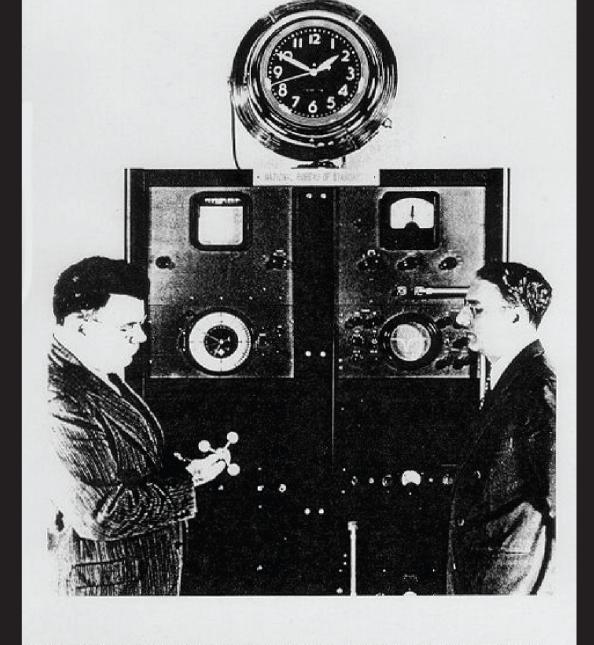
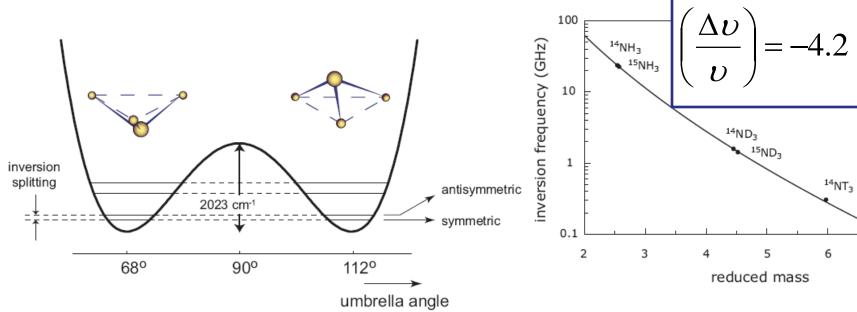


Figure 1: The World's first atomic clock; the ammonia absorption cell atomic clock at the National Bureau of Standards (now the National Institute of Standards and Technology) first operated in August 1948. Dr. Harold Lyons, inventor, is at the right; Dr. Edward U. Condon, Director of NBS, is at the left. The ammonia absorption cell is the coil of waveguide surrounding the clock face.

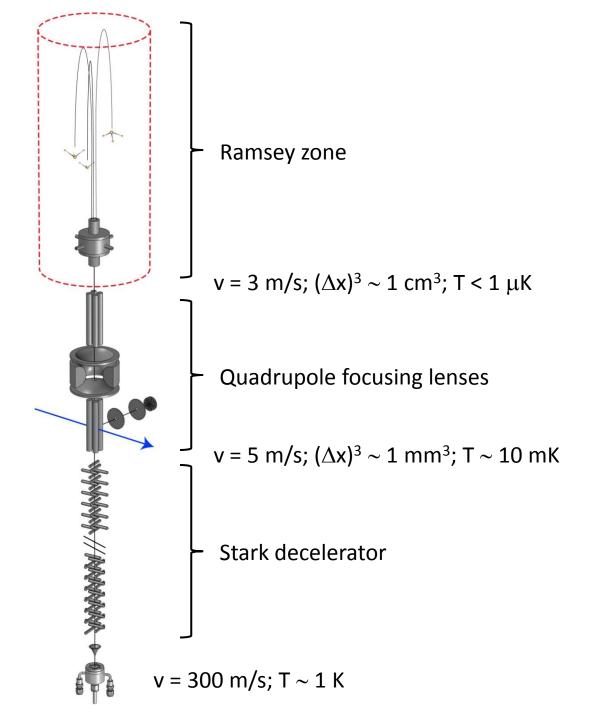
Inversion splitting in Ammonia

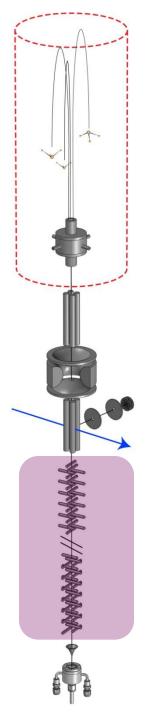
- Ammonia has a double-well potential, giving rise to inversion doubling
- Inversion frequency strongly depends on proton mass







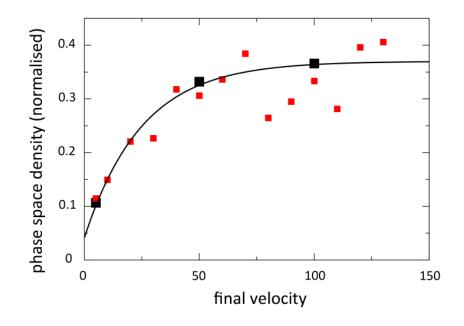


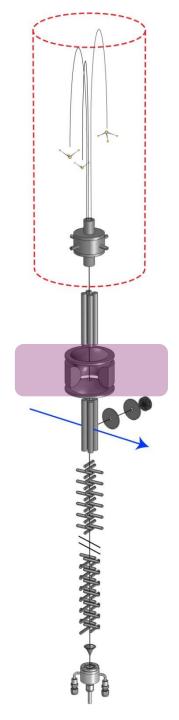


PROBLEMS:

Decelerator: Poor efficiency at low velocities

- Coupling between longitudinal and transverse motion.
- · Over-focusing in last stages.





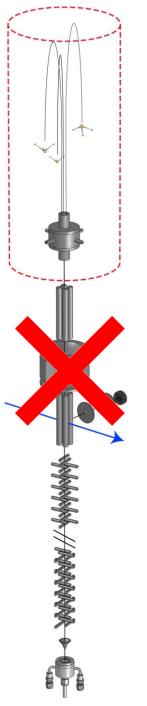
PROBLEMS:

Decelerator: Poor efficiency at low velocities

- Coupling between longitudinal and transverse motion.
- Over-focusing in last stages.

Buncher: No longitudinal focusing observed

- The buncher is most effective for v ≤ 8 m/s; while the lowest velocity observed after the buncher is 12 m/s.
- Bunching depends very critically on the timing of the applied voltages.



Solution (?):

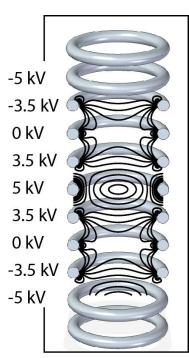
Replace focusing section by a ringtype decelerator



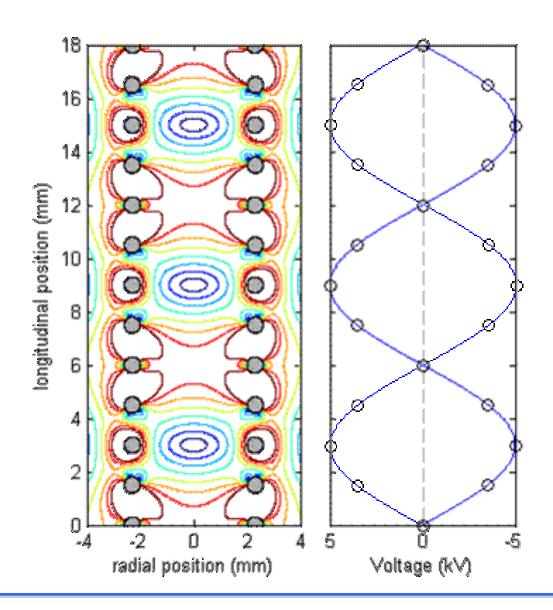
decelerate from ~100m/s to 3m/s.

More robust cooling:

Performing adiabatic cooling by slowly lowering the voltages applied to the ring-type decelerator.



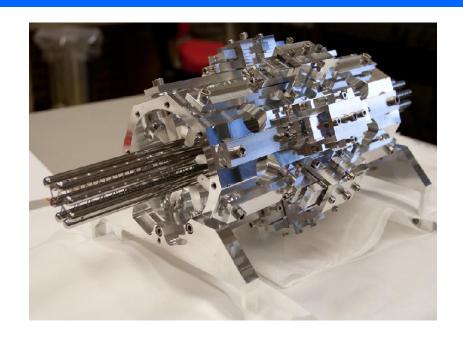
Operation principle of ring-type decelerator







The ring decelerator





Ring decelerator constructed at KVI Groningen similar to the design that was used by Osterwalder et al. [1]

492mm long, consisting of 328 rings with 4 mm inner diameter; Periodicity 12 mm.

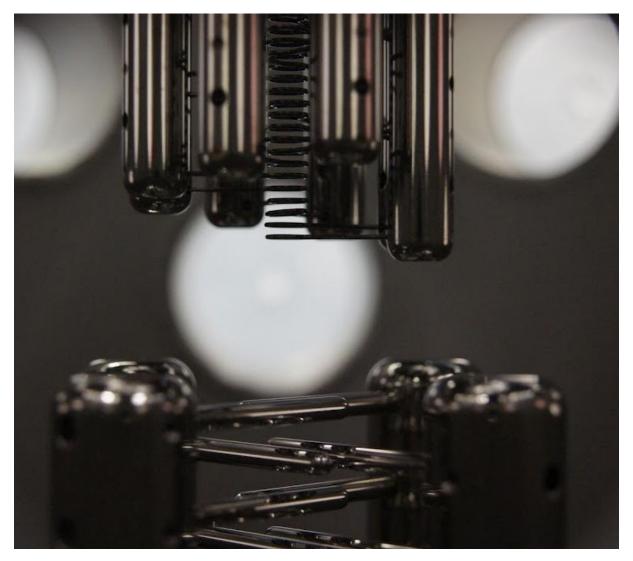
Amplifiers (8 x TREK 5/80-H-CE)

10 kV (peak to peak), frequency 0-15 kHz.





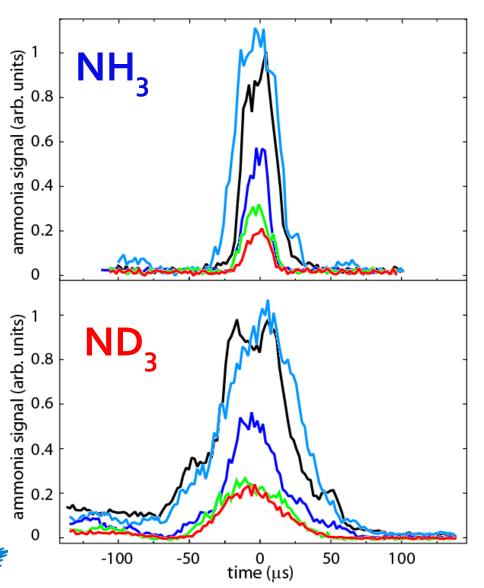
Where two worlds meet

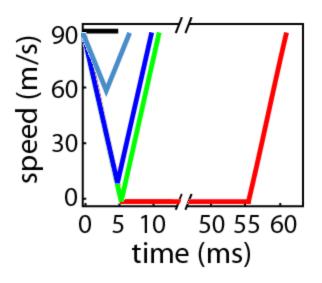






Guiding, Decelerating...





— 90 m/s

___ 60 m/s

___ 10 m/s

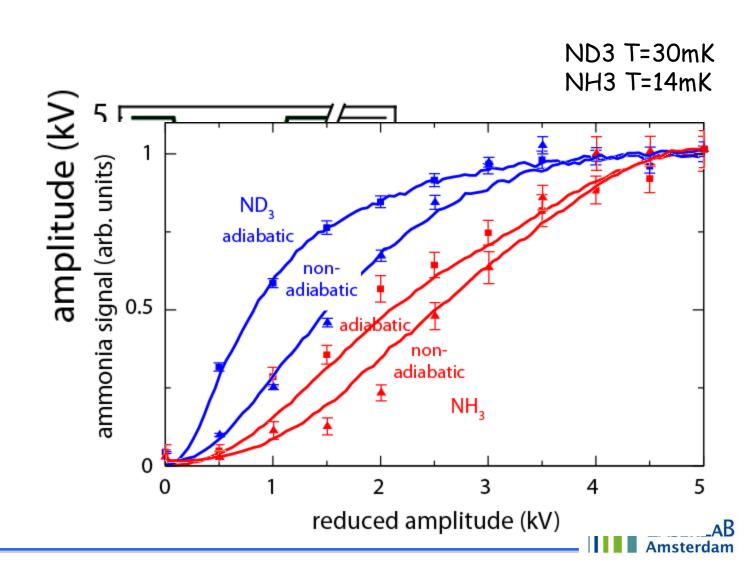
___ 0 m/s

trapped 50 ms



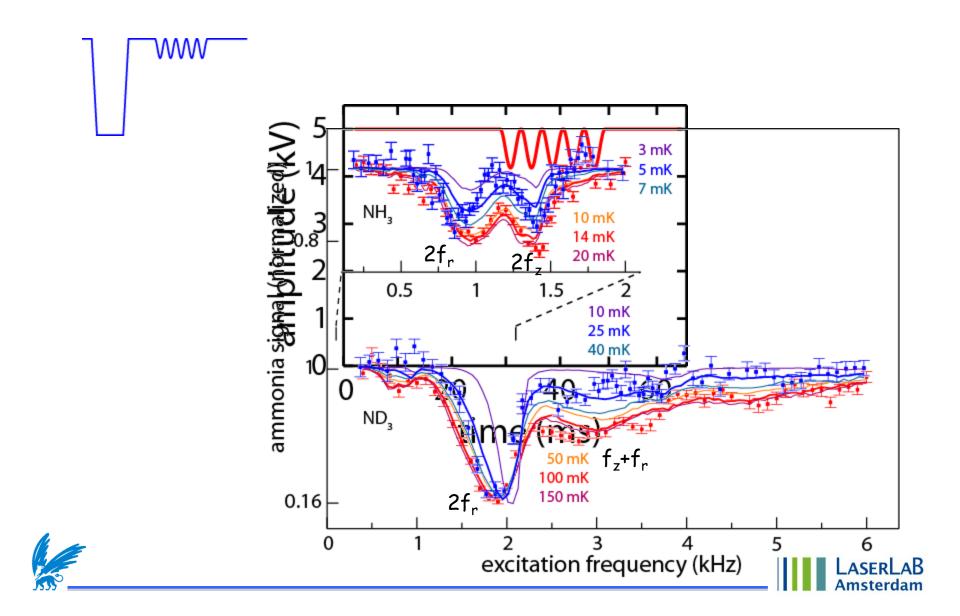


Adiabatic cooling





Exciting the trap motion



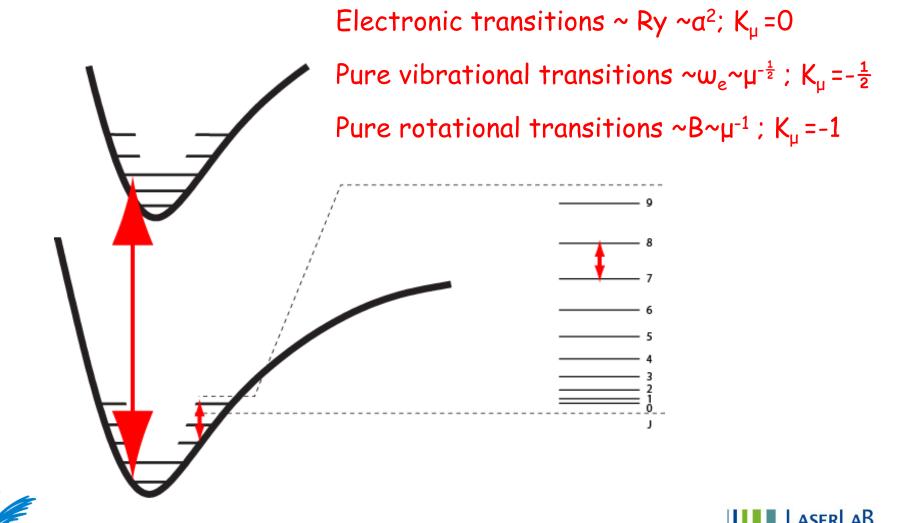
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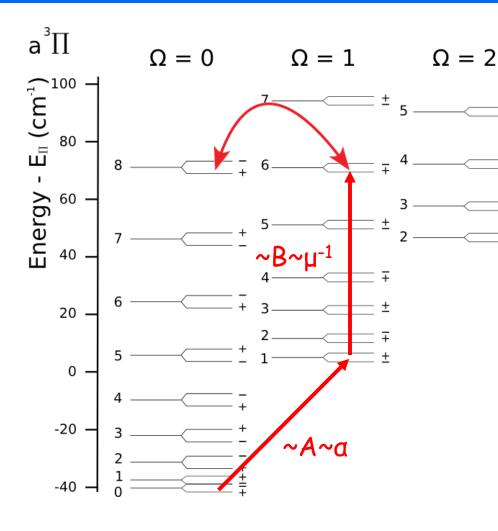




Dependence of molecular spectra on μ and a



Energy levels of CO in $a^3\Pi$ state



Enhancement: K₁~A/2v

- $a^3\pi$ is the first excited state of CO at 6eV.
- Decay to the groundstate is spin-forbidden.
- Lifetime 3-200ms.
- Can be directly excited from the groundstate using light around 206nm.





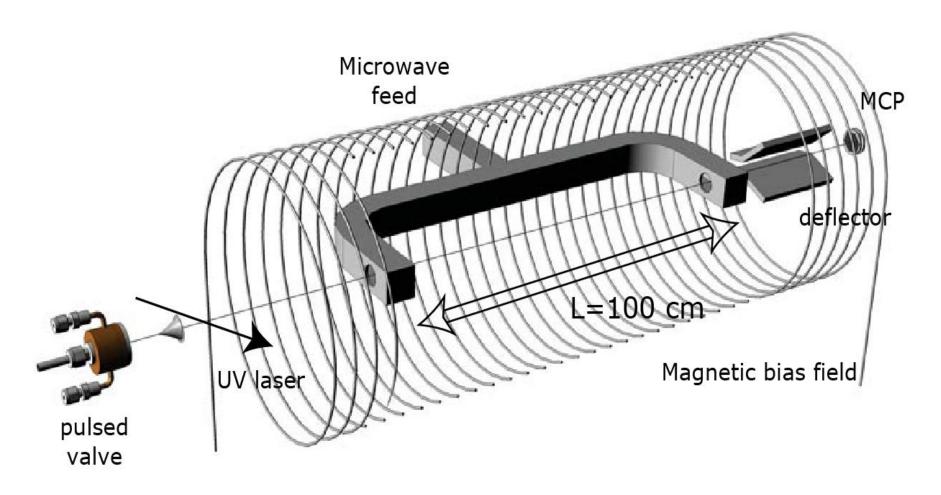
K-coefficients of 2-photon transitions in CO

Isotopologue	Transition	Measured (MHz)	MeasCalc. (MHz)	\mathbf{K}_{μ}
¹² C ¹⁶ O	$J = 6, \Omega = 1, + \rightarrow J = 4, \Omega = 2, +$	19270.1	3.5	27.8
	$J = 6, \Omega = 1, - \rightarrow J = 4, \Omega = 2, -$	16057.7	4.7	33.7
	$J = 6, \Omega = 1, + \rightarrow J = 8, \Omega = 0, +$	-1628.3	-3.3	-334
	$J = 6, \Omega = 1, - \rightarrow J = 8, \Omega = 0, -$	-19406.7	4.5	-27.3
¹³ C ¹⁶ O	$J = 6, \Omega = 1, +, F = 3.5 \rightarrow J = 4, \Omega = 2, +, F = 6.5$	43005.8	0.1	12.9
	$J = 6, \Omega = 1, -, F = 3.5 \rightarrow J = 4, \Omega = 2, -, F = 6.5$	39988.0	0.7	12.2
	$J = 6, \Omega = 1, +, F = 5.5 \rightarrow J = 8, \Omega = 0, +, F = 8.5$	22329.6	6.0	23.5
	$J = 6, \Omega = 1, -, F = 5.5 \rightarrow J = 8, \Omega = 0, -, F = 8.5$	4003.4	5.0	128
13C18O	$J = 6, \Omega = 1, +, F = 3.5 \rightarrow J = 4, \Omega = 2, +, F = 6.5$	69062.0	-7.1	7.22
	$J = 6, \Omega = 1, -, F = 3.5 \rightarrow J = 4, \Omega = 2, -, F = 6.5$	66277.5	-3.6	7.60





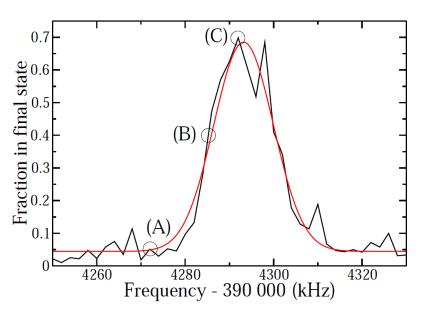
Setup to measure 2-photon transitions



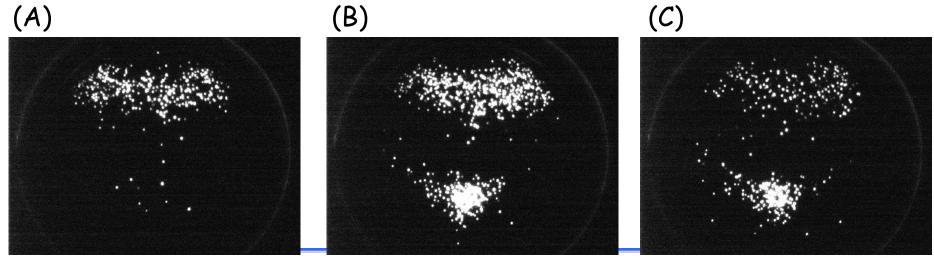




Microwave measurements



- MW-frequency scanned
- Fraction of molecules in final state calculated
- ~250 molecules per shot
- Shot-to-shot noise cancels



Probing time-variation of μ using:

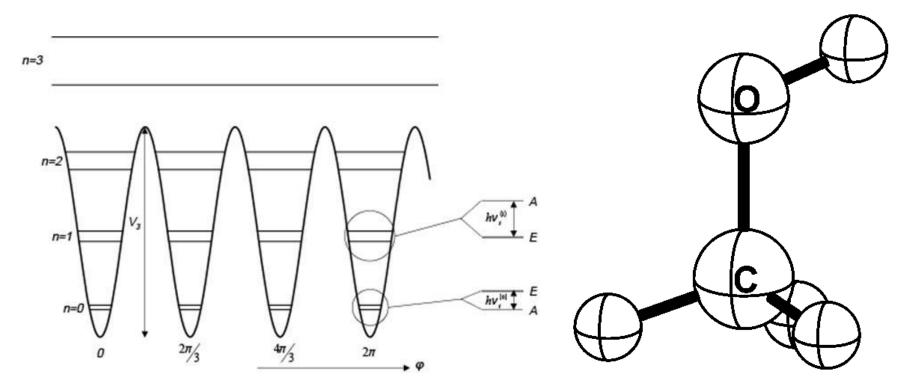
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Hindered rotation in Methanol

- The internal rotation is hindered by a barrier, leading to a splitting between levels of A and E symmetry.
- The torsional splittings strongly depends on proton mass

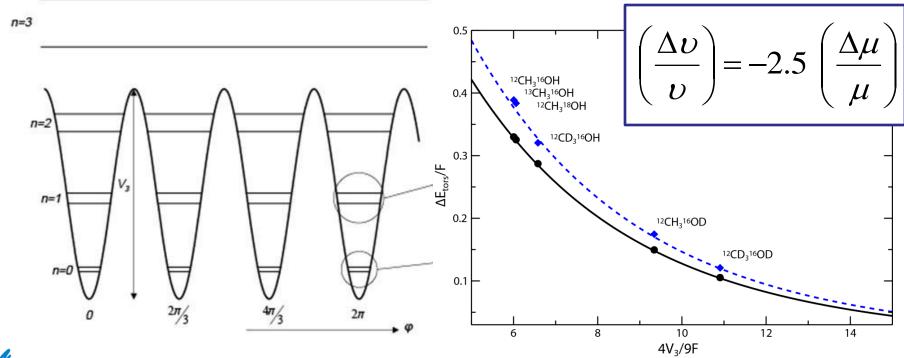






Hindered rotation in Methanol

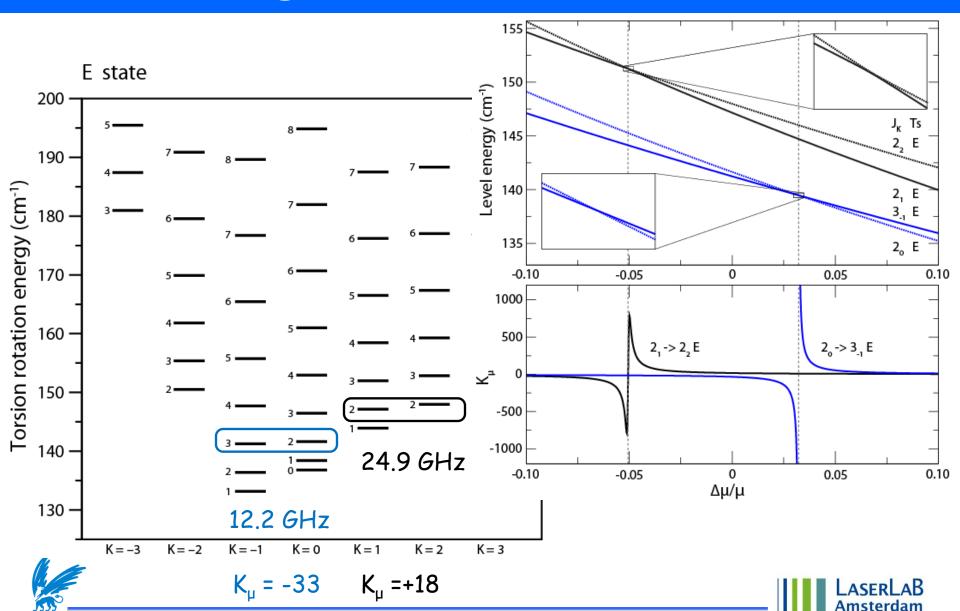
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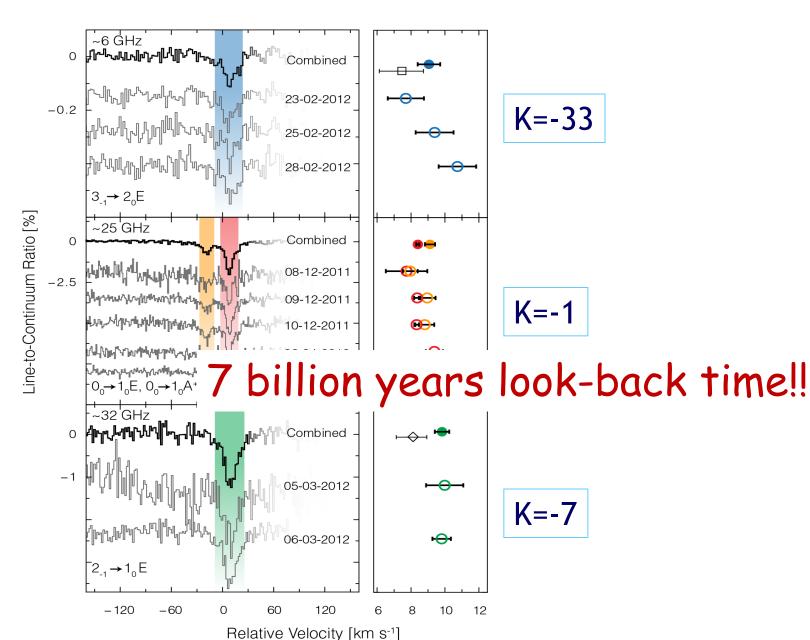


Near degeneracies in level scheme

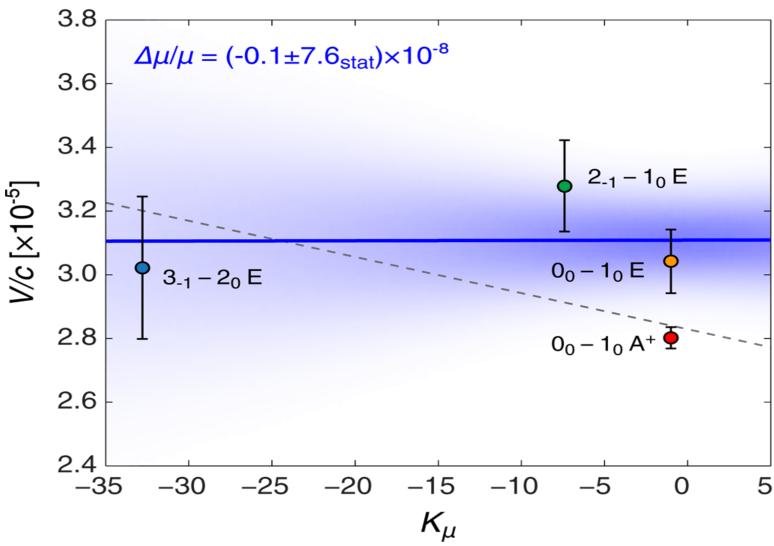




Methanol in PKS-1830-211



Constraint on time-variation of μ









LaserLaB, Amsterdam: Paul Jansen Marina Quintero Pérez Thomas Wall Adrian de Nijs Aernout van der Poel Peter Zieger Julija Bagdonaite Rob Kortekaas Wim Ubachs Rick Bethlem

KVI Groningen: Joost van den Berg Steven Hoekstra

LISA et Université's Paris 7 et Paris Est: Isabelle Kleiner

Max Planck Institut für Radioastronomie, Bonn: Christian Henkel Karl Menten







