

Towards measuring the electron's electric dipole moment using trapped molecules

Bart Schellenberg

NNV Soesterberg

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NL-eEDM collaboration

Particle Physics

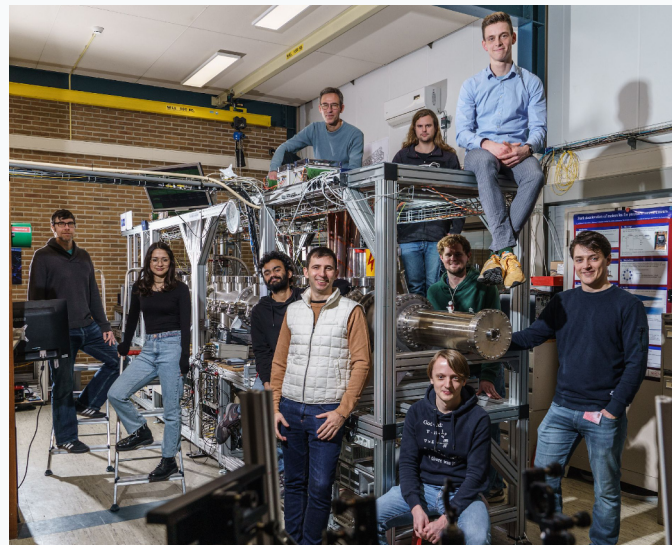
Jordy de Vries
Heleen Mulder
Rob Timmermans

Quantum Chemistry

Anastasia Borschevsky
Lukáš Pašteka
Agustín Aucar
Eifion Prins

Experiments

Steven Hoekstra
Hendrick Bethlem
Lorenz Willman
Steven Jones
Wim Ubachs
Joost van Hofslot
Jelmer Levenga
Bastiaan Nijman
Ties Fikkers
Lucas van Sloten
Bart Schellenberg
Nithesh Balasubramanian
Izabella Thompson
Jason Arzinos
Marianne Westerhof




eEDM as a probe for BSM theories


The EDM of a fundamental particle describes a CP violating interaction.

At $\mathcal{O}^{d=6}$

$$H_{\text{eEDM}} = 2d_e \vec{S} \cdot \vec{E}$$

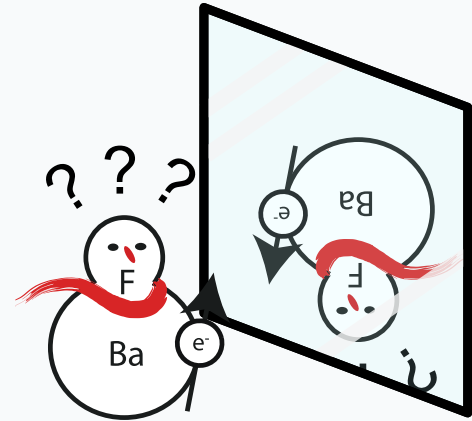


pseudo-vector



vector

CP violation in SM insufficient to explain observed matter-anti-matter asymmetry.



eEDM as a probe for BSM theories

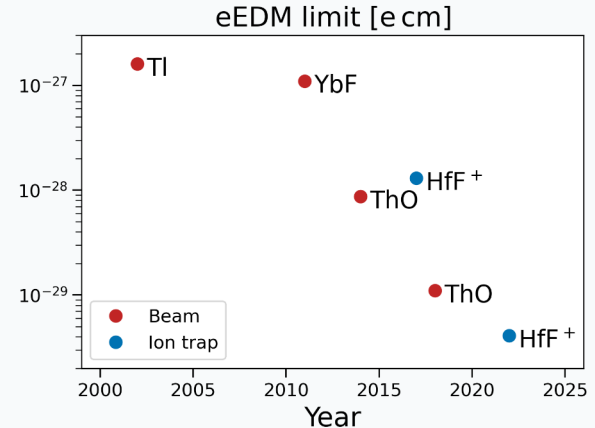
Measure shift in molecular energy that correlates with electric field reversal.

Shot-noise-limited sensitivity:

$$\sigma = \frac{\hbar}{2 |\Omega| W_d \tau \sqrt{N_p}}$$

Diagram illustrating the components of the shot-noise-limited sensitivity formula:

- $|\Omega|$: state-dependent factor
- W_d : heavy, polar molecule
- τ : interaction time
- $\sqrt{N_p}$: number of molecules



Using polar molecules



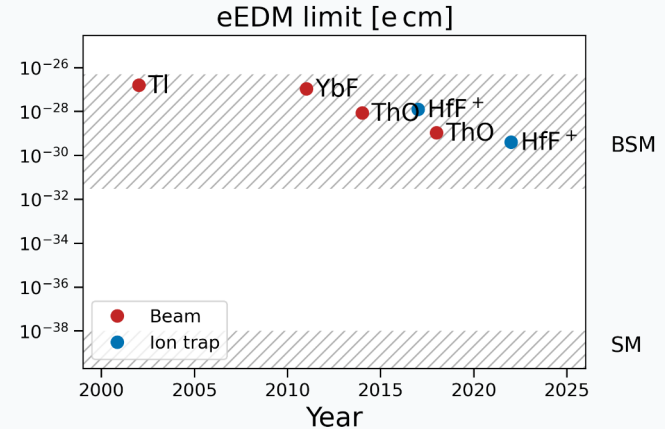
eEDM as a probe for BSM theories

Measure shift in molecular energy that correlates with electric field reversal.

Shot-noise-limited sensitivity:

$$\sigma = \frac{\hbar}{2 |\Omega| W_d \tau \sqrt{N_p}}$$

state-dependent factor $\rightarrow |\Omega|$
 heavy, polar molecule $\rightarrow W_d$
 interaction time $\rightarrow \tau$
 number of molecules $\rightarrow N_p$



Using polar molecules



eEDM as a probe for BSM theories

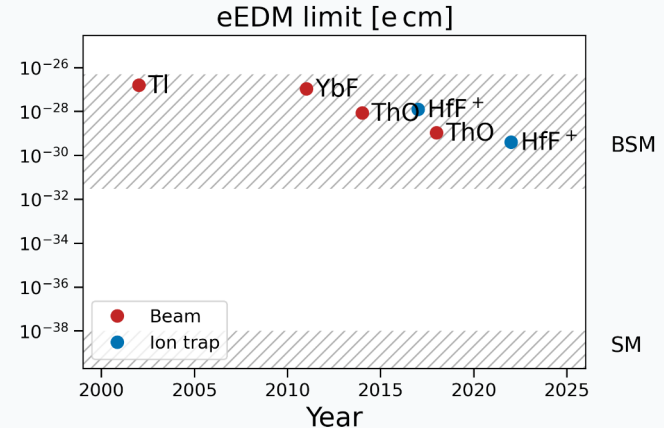
Measure shift in molecular energy that correlates with electric field reversal.

Shot-noise-limited sensitivity:

Can we increase τ , even if that means sacrificing N_p ?

$$\sigma = \frac{\hbar}{2 |\Omega| W_d \tau \sqrt{N_p}}$$

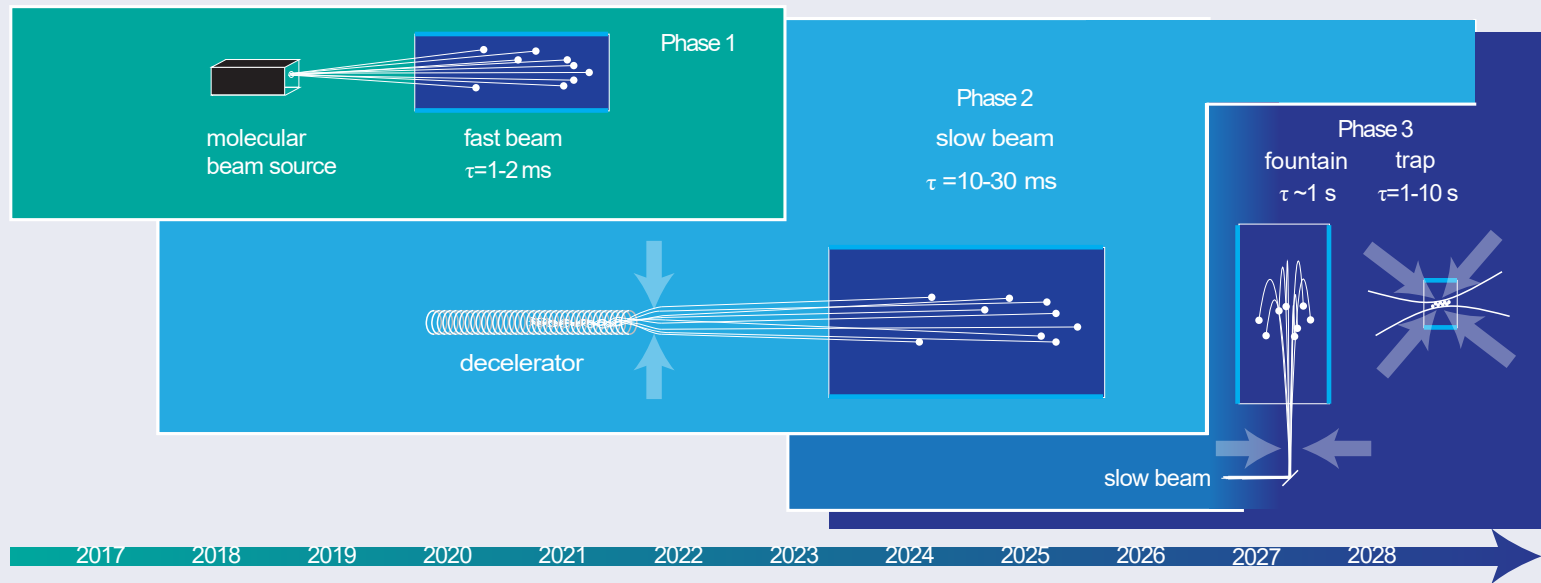
state-dependent factor $|\Omega|$
 heavy, polar molecule W_d
 interaction time τ
 number of molecules N_p



Using polar molecules



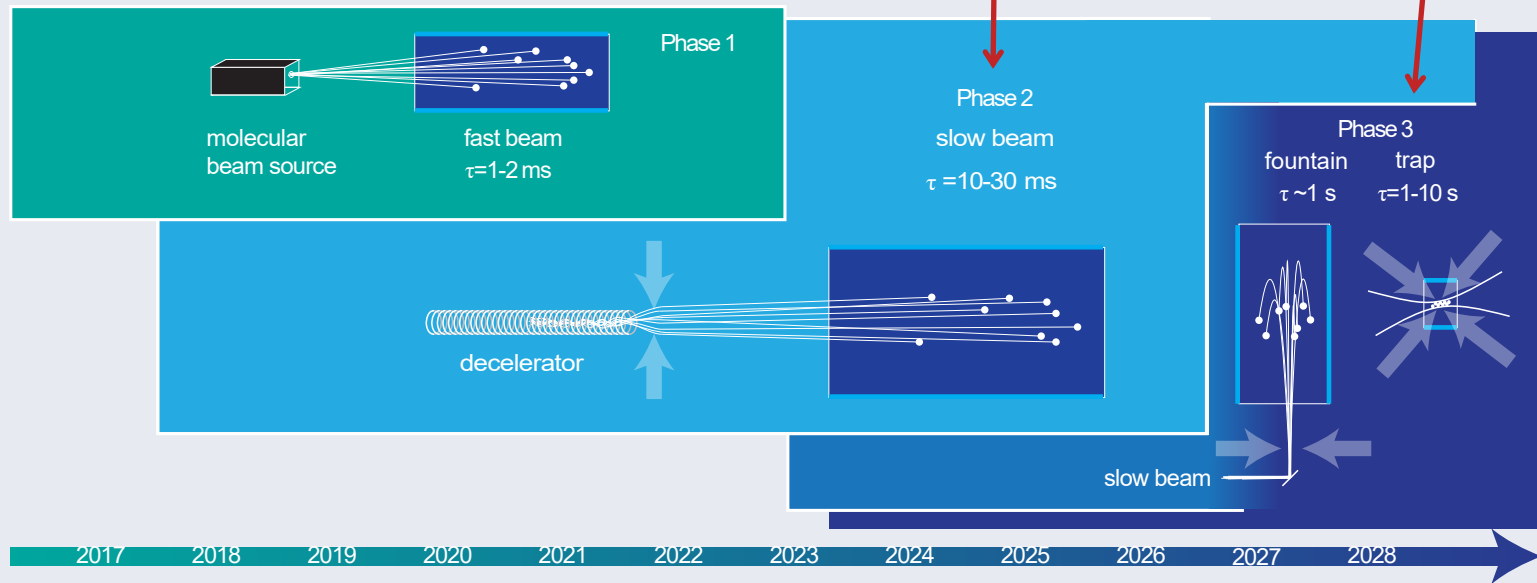
The NL-eEDM experiment



The NL-eEDM experiment

Talk by Bastiaan Nijman:
"Background free detection of BaF for eEDM searches"
(15:20)

This talk



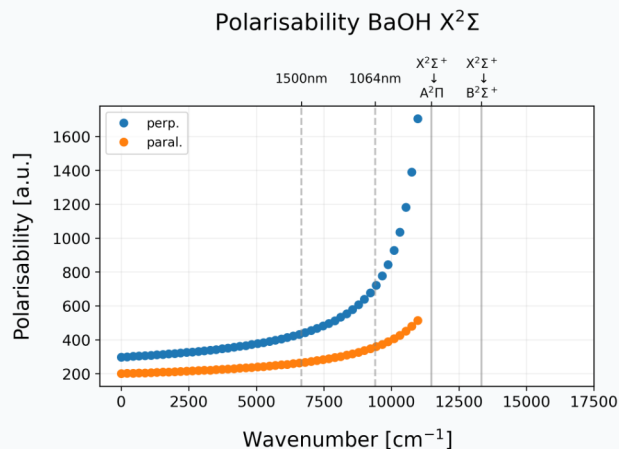
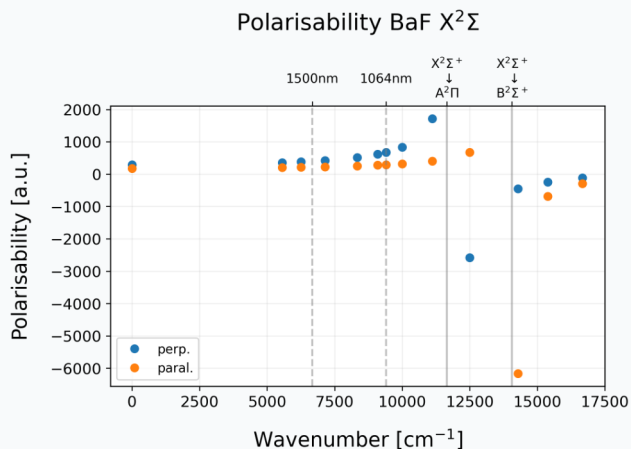
How to make an optical trap

Understanding how molecules interact with light.

$$\Delta E \approx -\frac{1}{4} \alpha \mathcal{E}^2$$

optical field

polarisability



Prinsen et al. arXiv:2510.19564
Schellenberg et al. arXiv:2507.17521

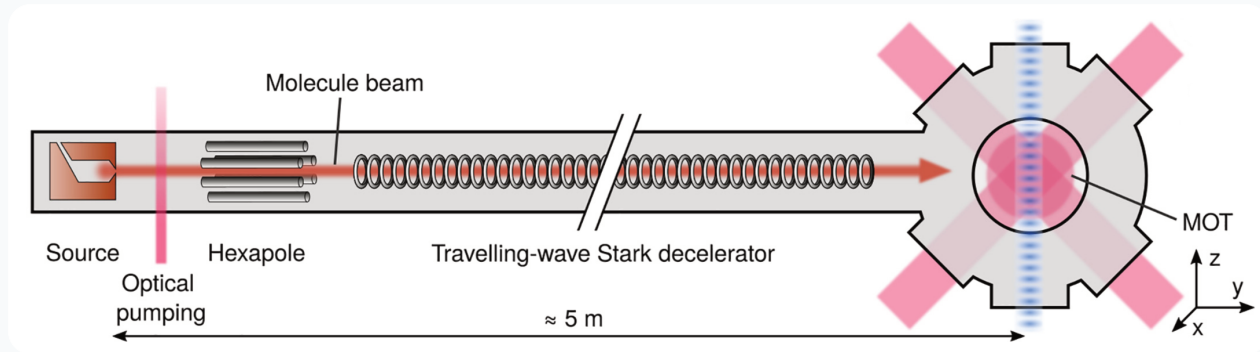


From source:

- $\sim 10^8$ molecules
- Velocity: $\sim 150 - 200 \text{ m/s}$
- Temperature: $\sim 5 - 20 \text{ K}$
- Volume: $\sim \text{cm}^3$

→ **Cooling required**

Sketch of potential loading setup



Production

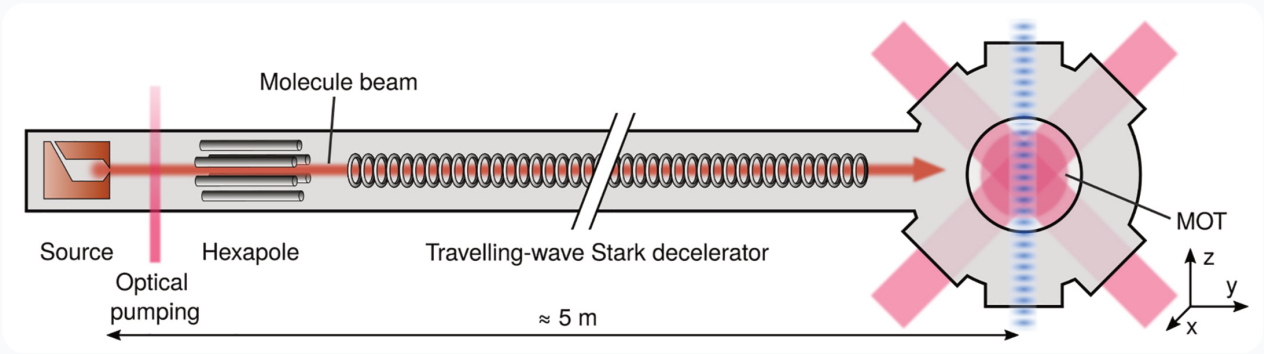
Cooling

Trapping

Bause et al. Phys. Rev. A 111, 062815 (2025)

Sketch of potential loading setup

Possible, but very challenging



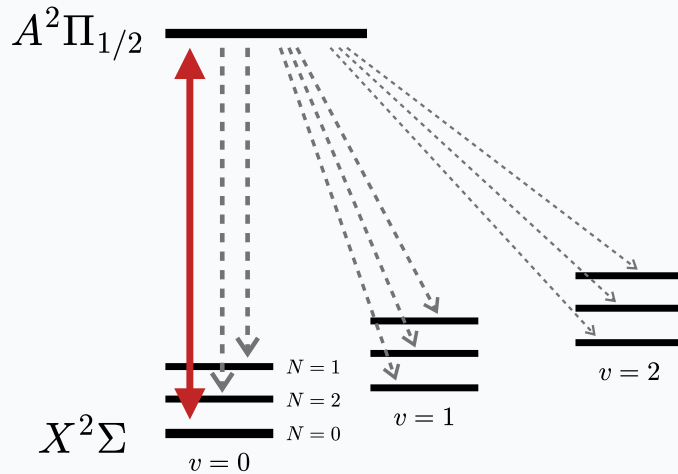
Production

Cooling

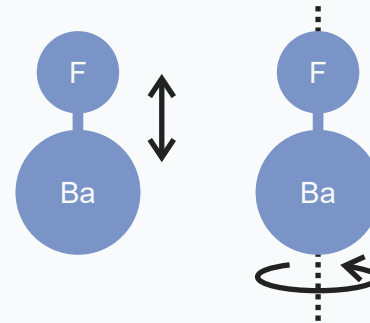
Trapping

Bause et al. Phys. Rev. A 111, 062815 (2025)

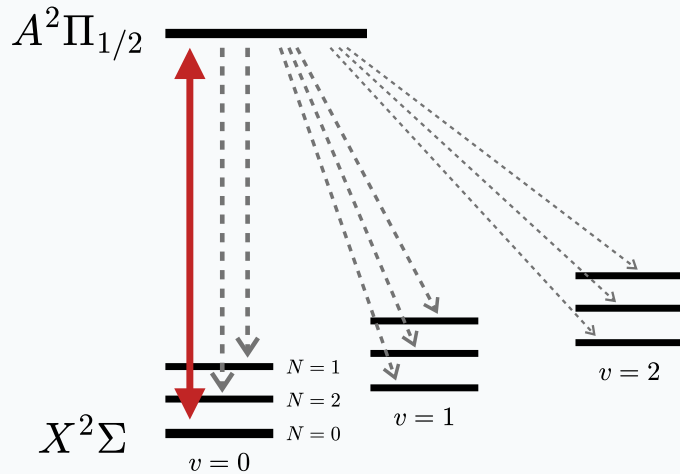
Laser cooling



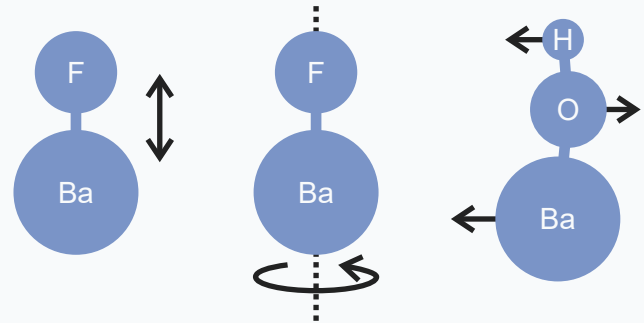
$$H = H_{\text{electronic}} + H_{\text{vibration}} + H_{\text{rotation}} + H_{\text{fine}} + H_{\text{hyperfine}}$$



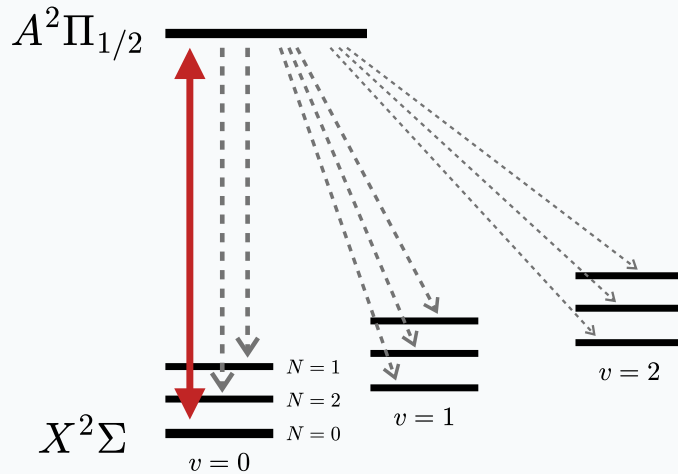
Laser cooling



$$H = H_{\text{electronic}} + H_{\text{vibration}} + H_{\text{bending}} + H_{\text{rotation}} + H_{\text{fine}} + H_{\text{hyperfine}}$$

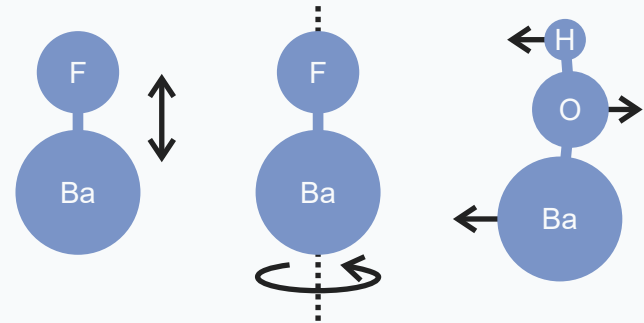


Laser cooling



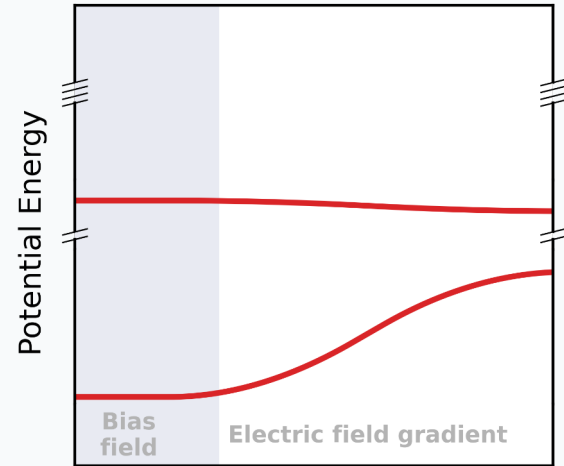
$$H = H_{\text{electronic}} + H_{\text{vibration}} + H_{\text{bending}} + H_{\text{rotation}} + H_{\text{fine}} + H_{\text{hyperfine}}$$

~ 5-10 lasers to keep cycle closed for BaF MOT



Single-photon trap loading

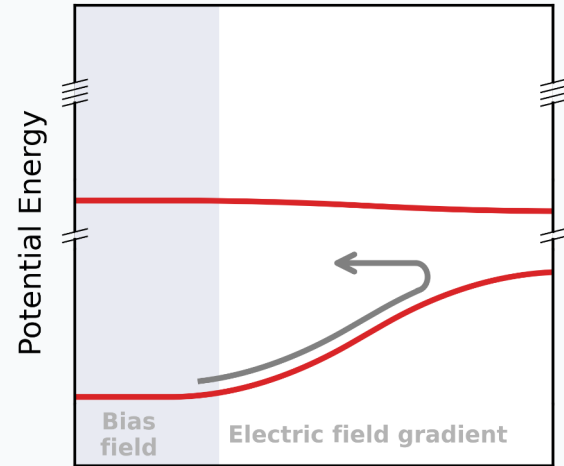
Exploiting state dependent potentials



Schellenberg et al. arXiv:2507.17521

Exploiting state dependent potentials

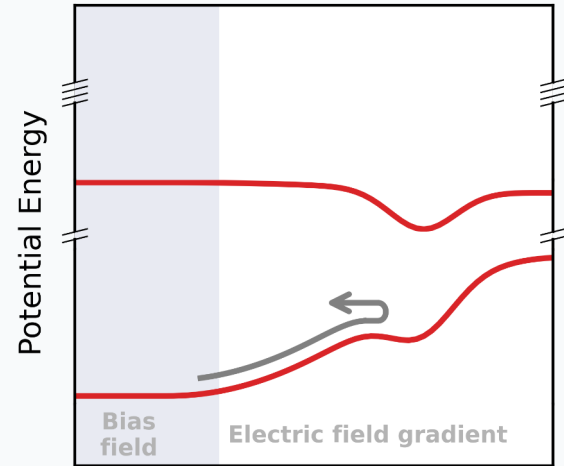
- |1> Molecules fly into potential barrier and are reflected.



Schellenberg et al. arXiv:2507.17521

Exploiting state dependent potentials

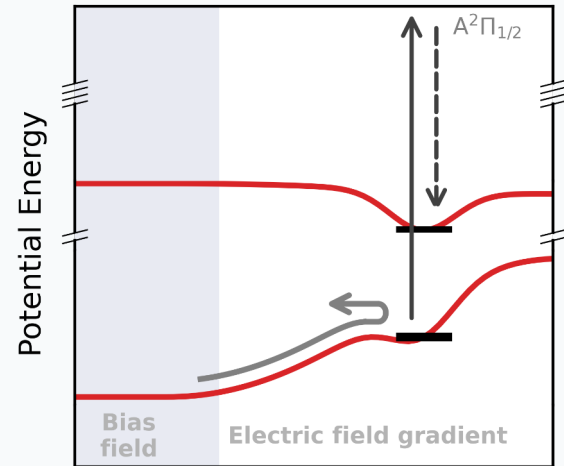
- |1⟩ Molecules fly into potential barrier and are reflected.
Optical trap is placed at the turning point.



Schellenberg et al. arXiv:2507.17521

Exploiting state dependent potentials

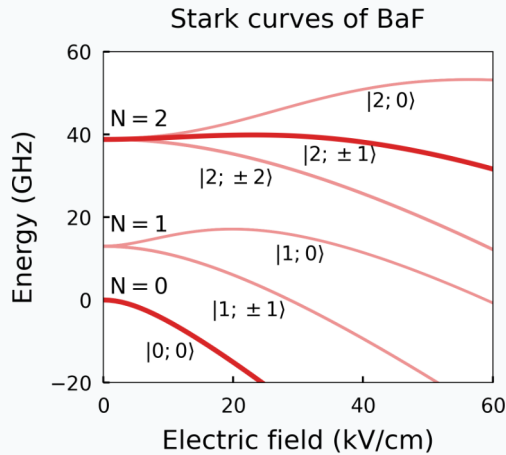
- 1) Molecules fly into potential barrier and are reflected.
Optical trap is placed at the turning point.
- 2) Molecules are excited when their kinetic energy is minimal.
- 3) Spontaneous decay to trapped state, whose coupling to E is small.



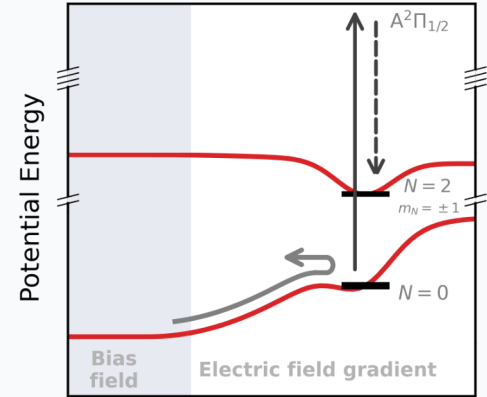
Schellenberg et al. arXiv:2507.17521

Exploiting state dependent potentials

Trap loading without optical cycling.



Rotational structure common in linear molecules.



Estimated Performance (using Stark decelerator)			
Trap depth	7.3 mK	Capture rate	$\sim 0.04\%$
Lifetime	~ 1 second	Number in trap	~ 880

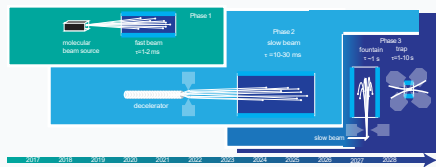
Schellenberg et al. arXiv:2507.17521



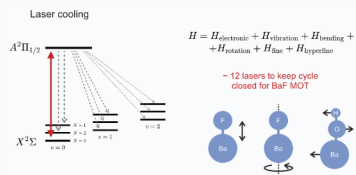
Conclusion

Summary

Traps offer longer interaction time



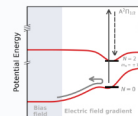
Cooling is challenging



Investigating alternative routes

Exploiting state dependent potentials

- [1] Molecules fly into potential barrier and are reflected. Optical trap is placed at the turning point.
- [2] Molecules are excited when their kinetic energy is minimal.
- [3] Spontaneous decay to trapped state, whose coupling to E is small.

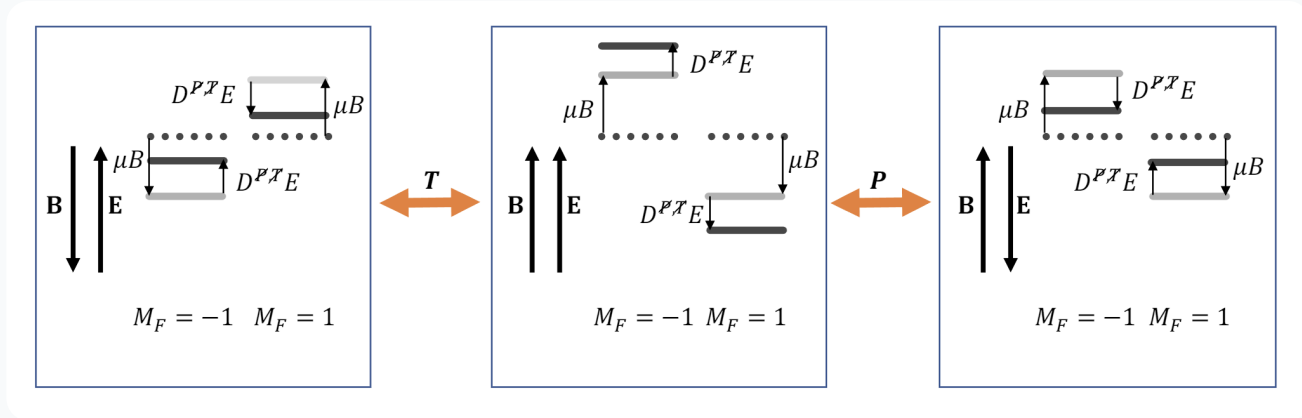


Backup slides

eEDM as a probe for BSM theories

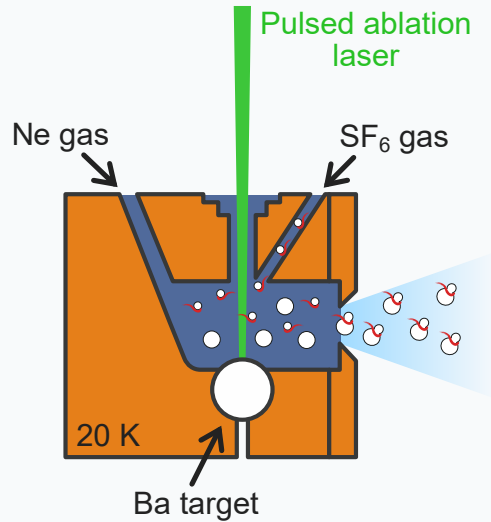
Molecular energy levels: $H = \dots - \vec{\mu} \cdot \vec{B} - \vec{D}^{\mathcal{PT}} \cdot \vec{E}$

Levels shift in
E and B field:



Measuring Ramsey interference between hyperfine M_F levels.

Figure adapted from A. Boeschoten (2023) [PhD thesis]



Output:

150 - 200 m/s

5 - 20 K

$\sim 10^8$ molecules

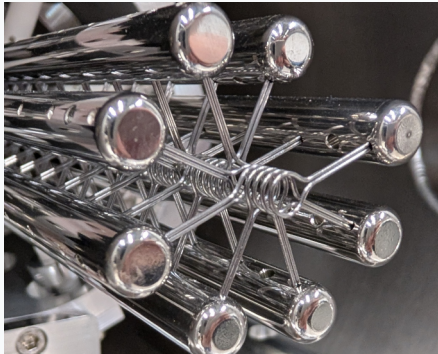
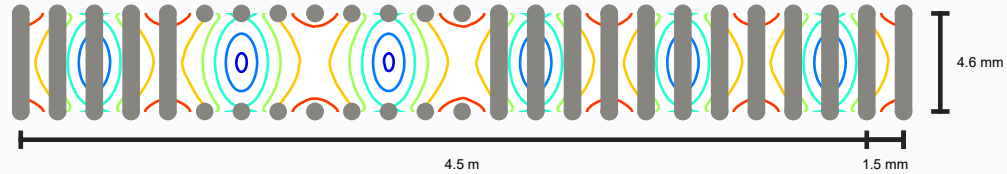
Work by:

* Ties Fikkers

* Nithesh Balasubramanian

(Longitudinal) Stark deceleration

3D moving electrostatic traps to
decelerate molecules



Output:

0 - 30 m/s

500 mK

$\sim 10^6$ molecules

Work by:

- * Lucas van Sloten
- * Marianne Westerhof