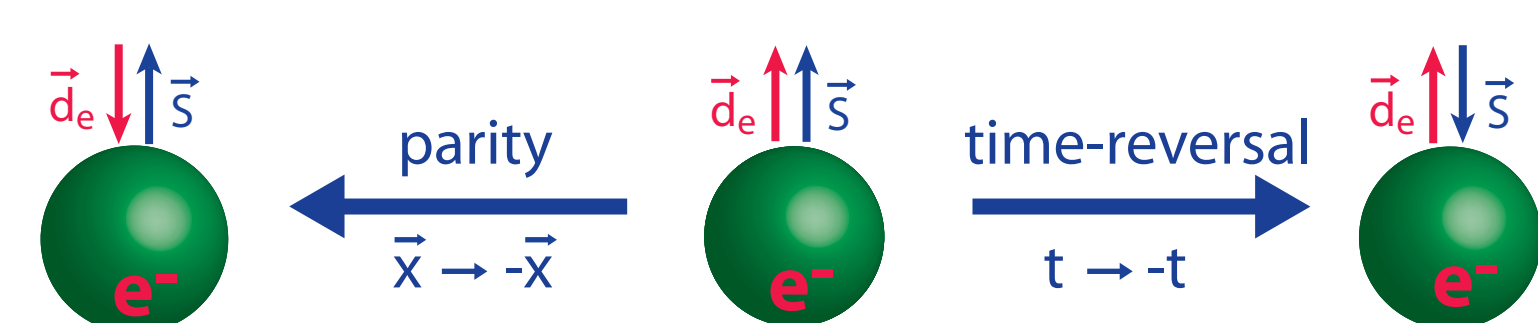


ACME collaboration

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## Why measure the electron EDM?

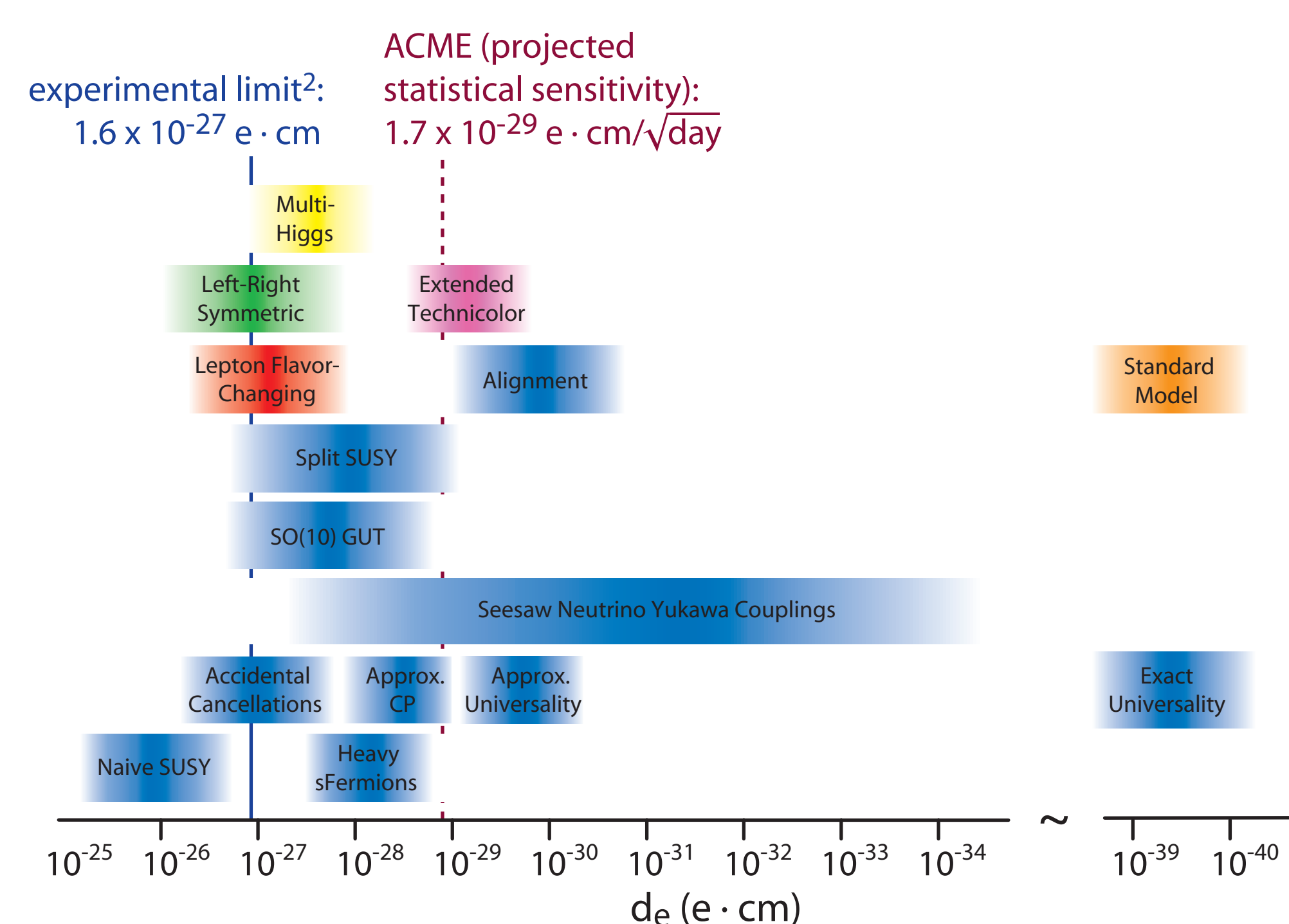
EDM of a fundamental particle violates both parity (P) and time-reversal (T) symmetries:



T-violation in the Standard Model (SM) is not sufficient to explain the observed dominance of matter over antimatter in the universe  $\Rightarrow$  additional sources of T-violation (beyond the SM) must exist

Additional T-violation in the lepton sector arises in most proposed extensions to the SM, which generically predict  $d_e \gtrsim 10^{-29} e \cdot \text{cm}$ . An experimental limit on  $d_e$  below  $\sim 10^{-29} e \cdot \text{cm}$  would also put strong constraints on possible mechanisms of electroweak baryogenesis<sup>1</sup>.

Predictions for the electron EDM from the SM and some proposed extensions:



<sup>1</sup>Y. Li, S. Profumo, and M. Ramsey-Musolf, arXiv:0811.1987v1 [hep-ph].  
<sup>2</sup>B. C. Regan et al., Phys. Rev. Lett. **88**, 071805 (2002).

## Advantages of ThO

### Advantages for statistical sensitivity

The effect of an electron EDM is amplified in a polar molecule like ThO due to the large internal electric field  $E_{eff}$  of the molecule. For ThO in the H state,  $E_{eff} = 104 \text{ GV/cm}$ .<sup>3</sup>

ThO can be made into a high-flux, cryogenic beam.

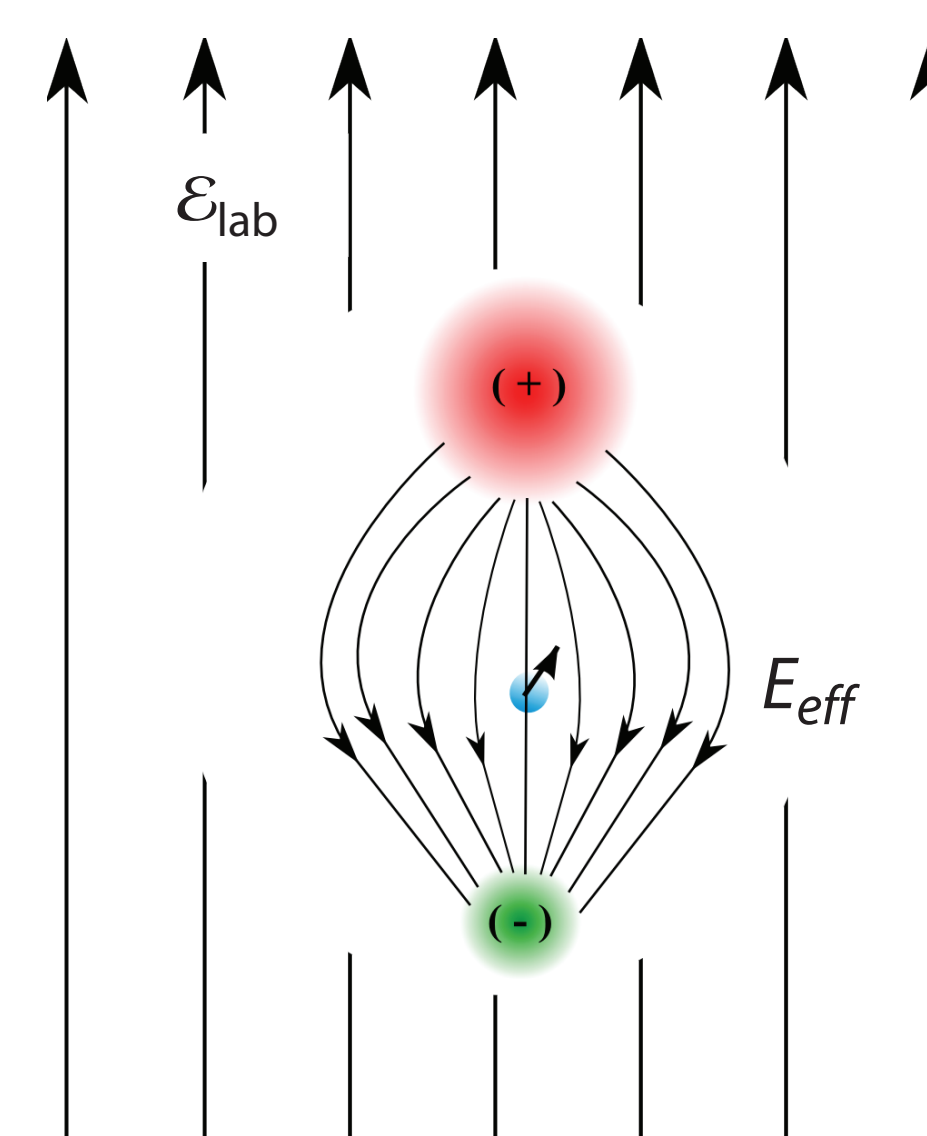
### Properties that reduce systematic effects

The molecule can be almost completely polarized by applying a small electric field  $\mathcal{E}_{lab} \sim 10 \text{ V/cm}$ . Smaller applied field  $\Rightarrow$  smaller leakage currents.

$\Omega$ -doublet in H state of ThO  $\Rightarrow$  two closely spaced, opposite-parity states with equal and opposite values of  $E_{eff}$ . Use of both  $\Omega$ -doublet states enables rejection of systematic effects associated with reversal of the applied electric field.

Suppressed magnetic moment in the  $H^3\Delta_1$  state  $\Rightarrow$  lower sensitivity to magnetic noise and systematics.

<sup>3</sup>E. R. Meyer and J. L. Bohn, Phys. Rev. A **78**, 010502 (2008).

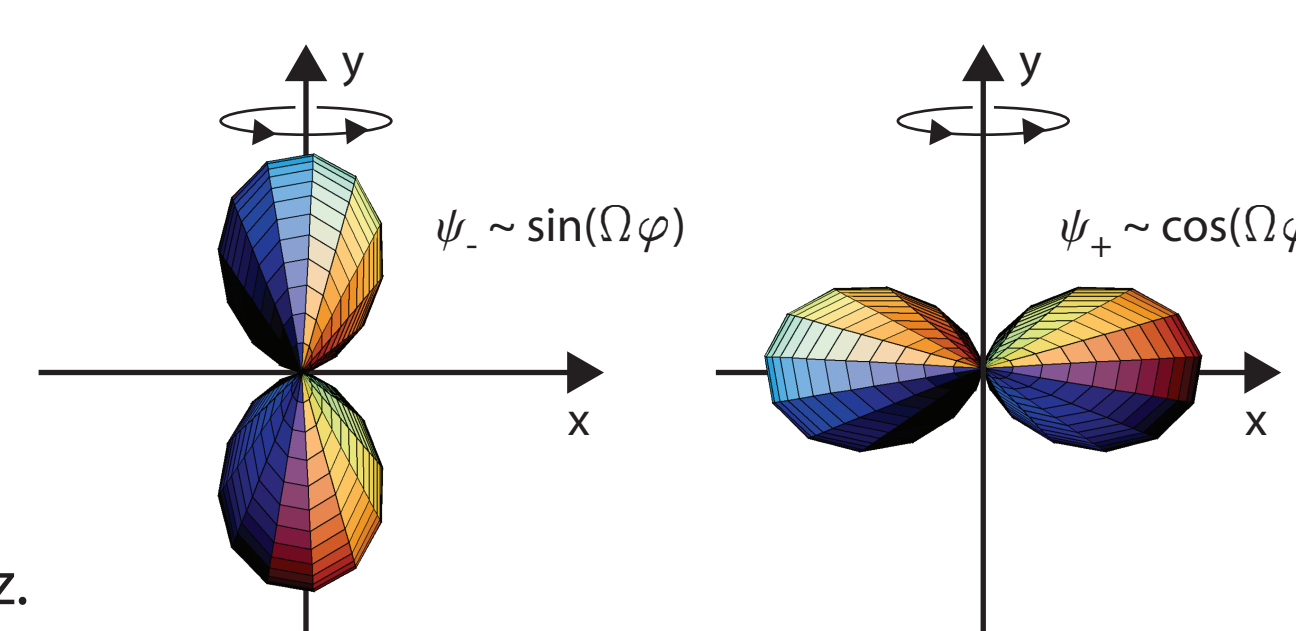


### Parity doublets in molecules

Electronic states with non-zero angular momentum ( $\Omega \neq 0$ ) have parity doublets ( $\Omega$ -doublets).

Parity doublets arise due to the coupling of the electronic and rotational motion.

Energy separation between  $\Omega$ -doublet components  $\Delta E \sim \frac{m_e}{M_n} B_e \approx 10 \text{ MHz}$ .

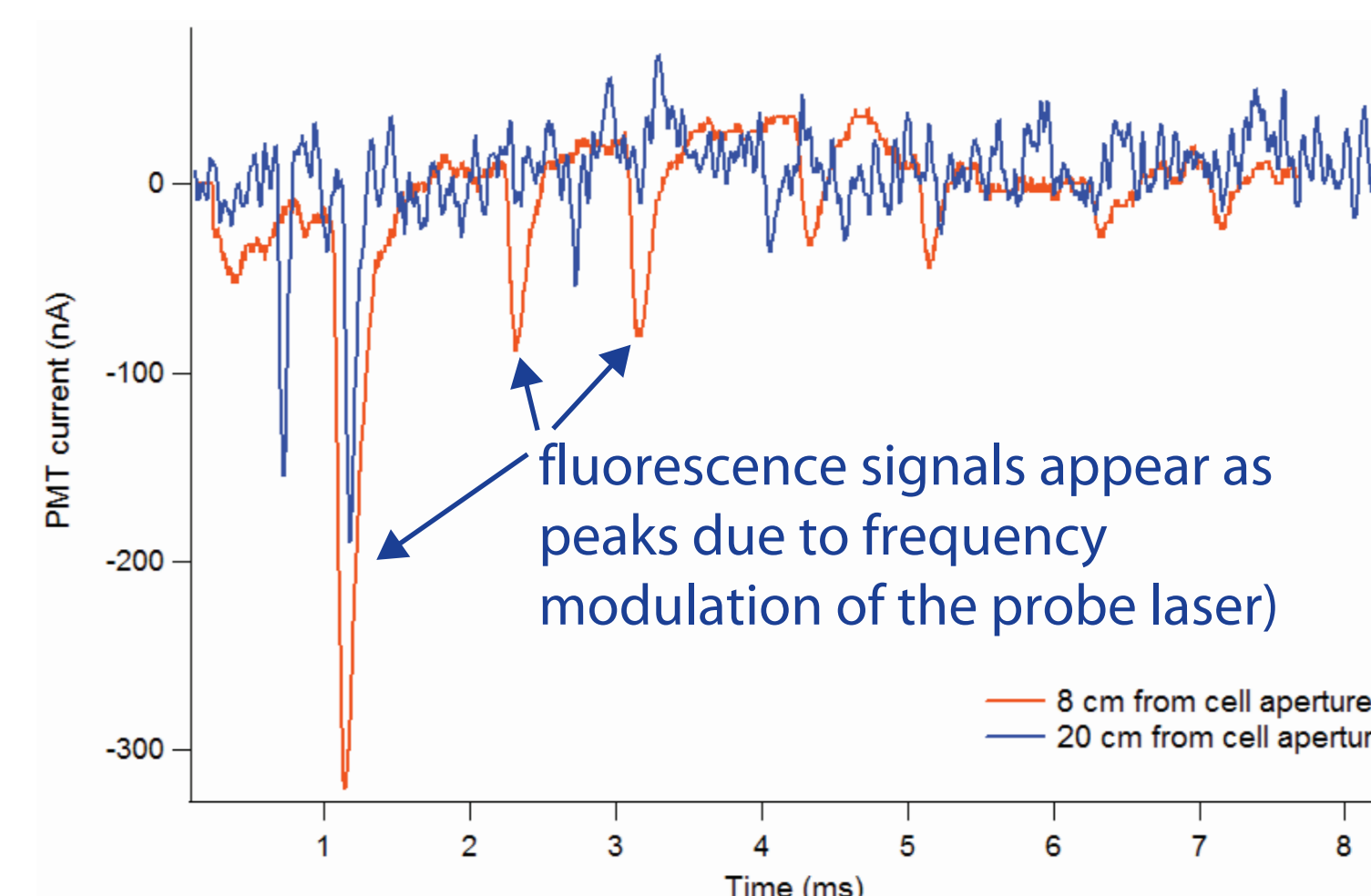
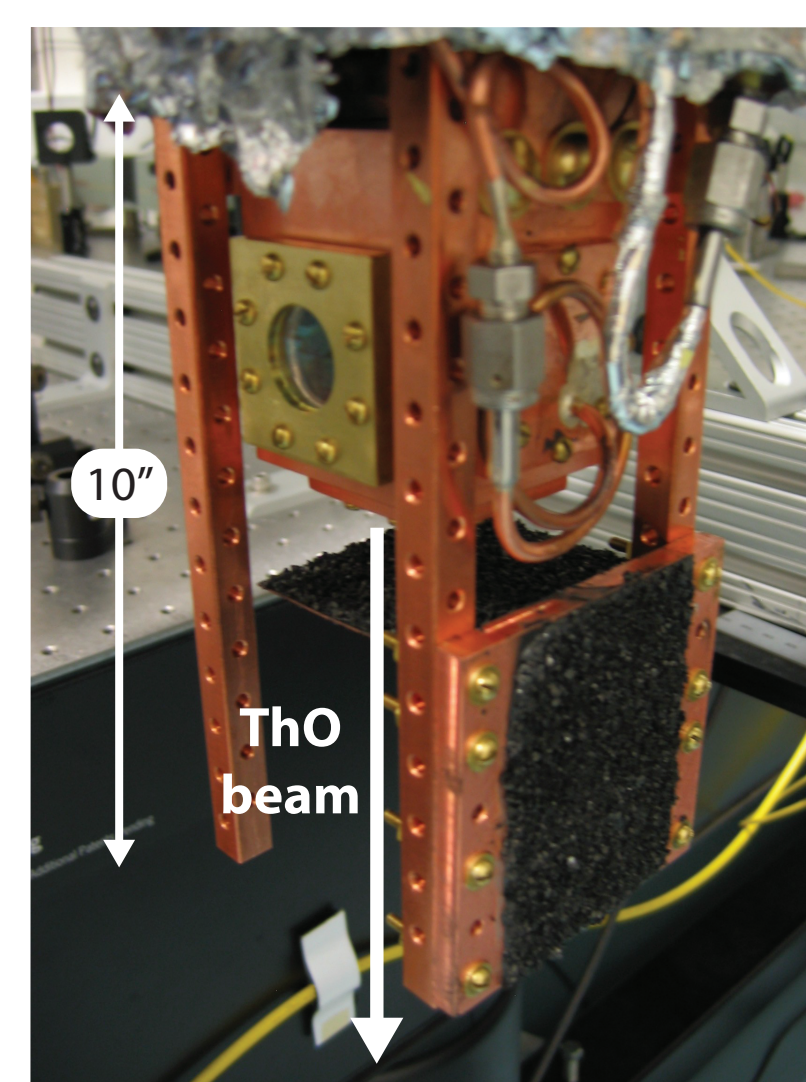


## Cryogenic ThO beam

ThO is produced by laser ablation of solid  $\text{ThO}_2$  with a pulsed Nd:YAG laser ( $\sim 10 \text{ mJ/pulse}$ ,  $5 \text{ ns pulse}$ ).

ThO molecules are cooled to  $4 \text{ K}$  by He buffer gas and entrained in the flow of He out of the cell ( $6 \text{ mm} \times 1 \text{ mm}$  aperture,  $10 \text{ sccm}$  He flow [ $4 \times 10^{18}$  atoms/sec]).

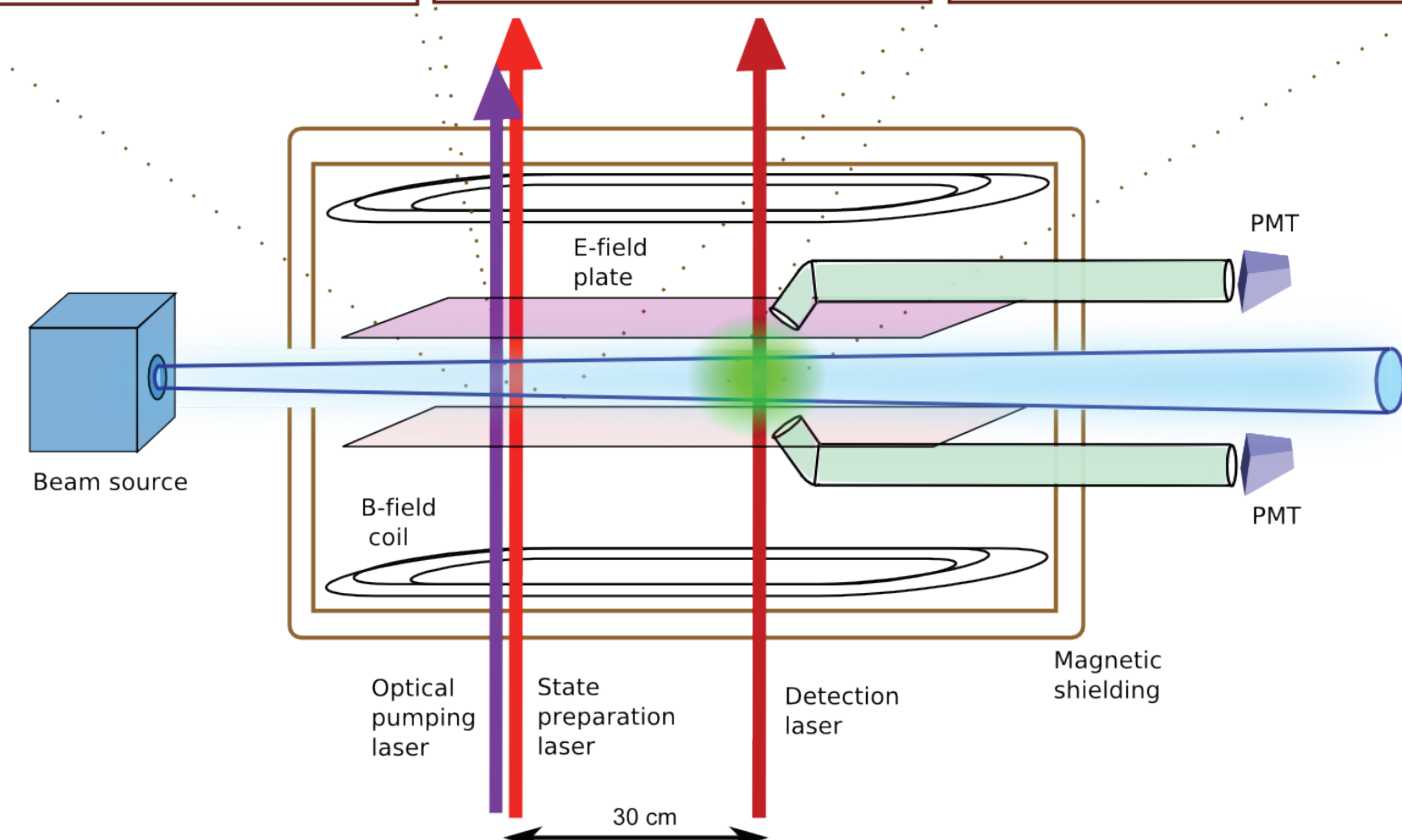
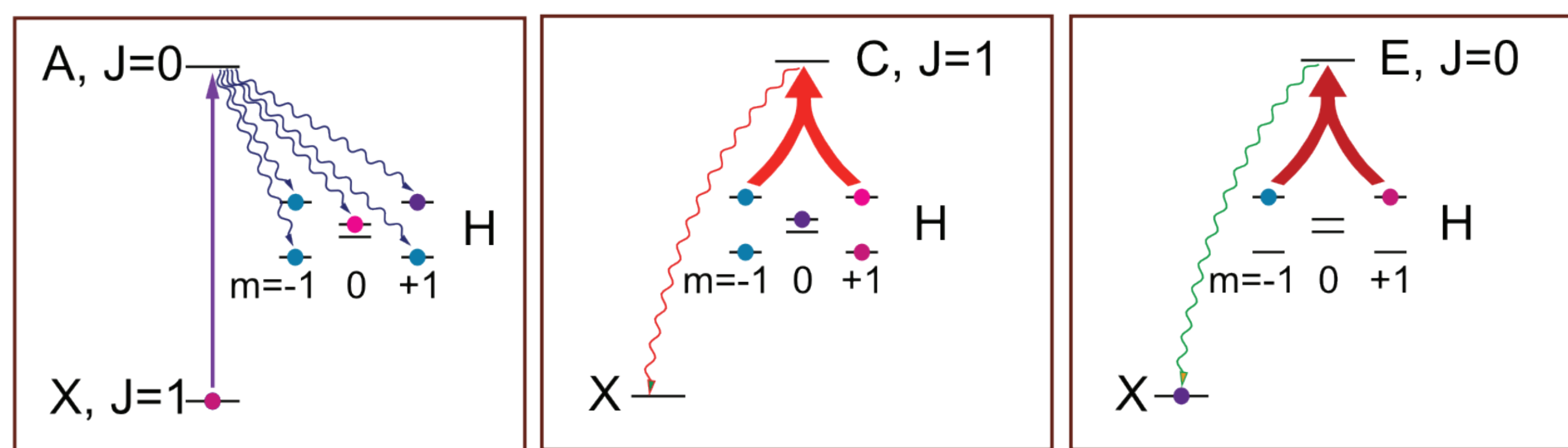
The ThO beam is detected via fluorescence on the  $X \rightarrow C$  transition (typical signals are shown below). Measured beam flux is  $\sim 1.5 \times 10^{10}$  molecules/shot in a single quantum state. Measured beam divergence is  $\Omega_b < 0.1 \text{ sr}$ .



## EDM measurement with ThO

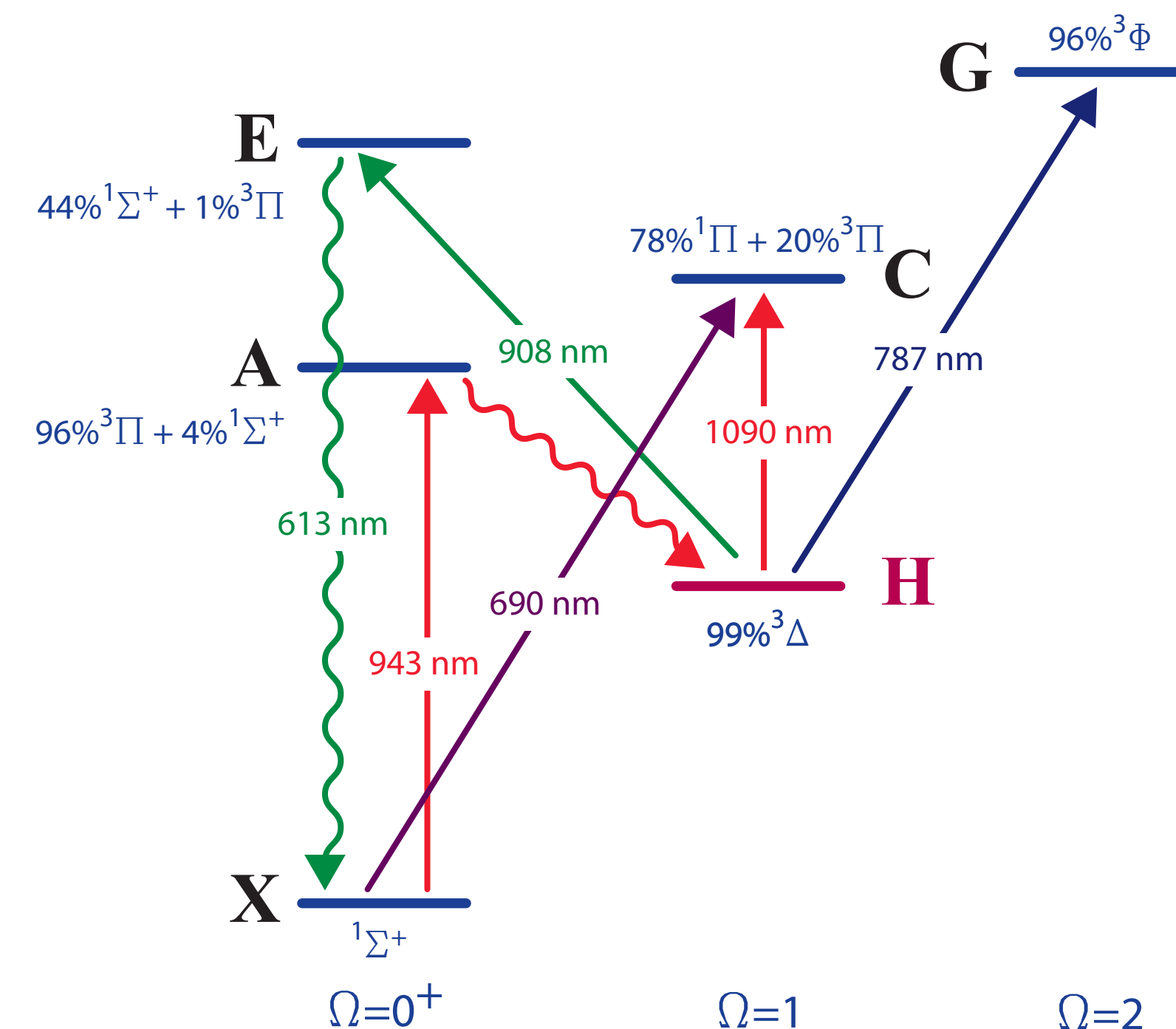
Optical pumping to A, which spontaneously decays to H, followed by depletion using x-polarized light on the  $H \rightarrow C$  transition produces a coherent superposition of molecules in the  $J=1$  rotational level of the H state:  $|\psi(0)\rangle = |m=+1\rangle + |m=-1\rangle$ .

After evolution in applied  $\mathcal{E}$ - and  $\mathcal{B}$ -fields for time  $T$ , the state is  $|\psi(T)\rangle = |m=+1\rangle + e^{i\phi}|m=-1\rangle$ , where  $\phi = (2g\mu_B B \pm 2d_e E_{eff})T/\hbar$ . Excitation to the E state with y- or x-polarized light results in a population in E that depends on  $\phi$ :  $P_E \propto 1 \pm \cos\phi$ .

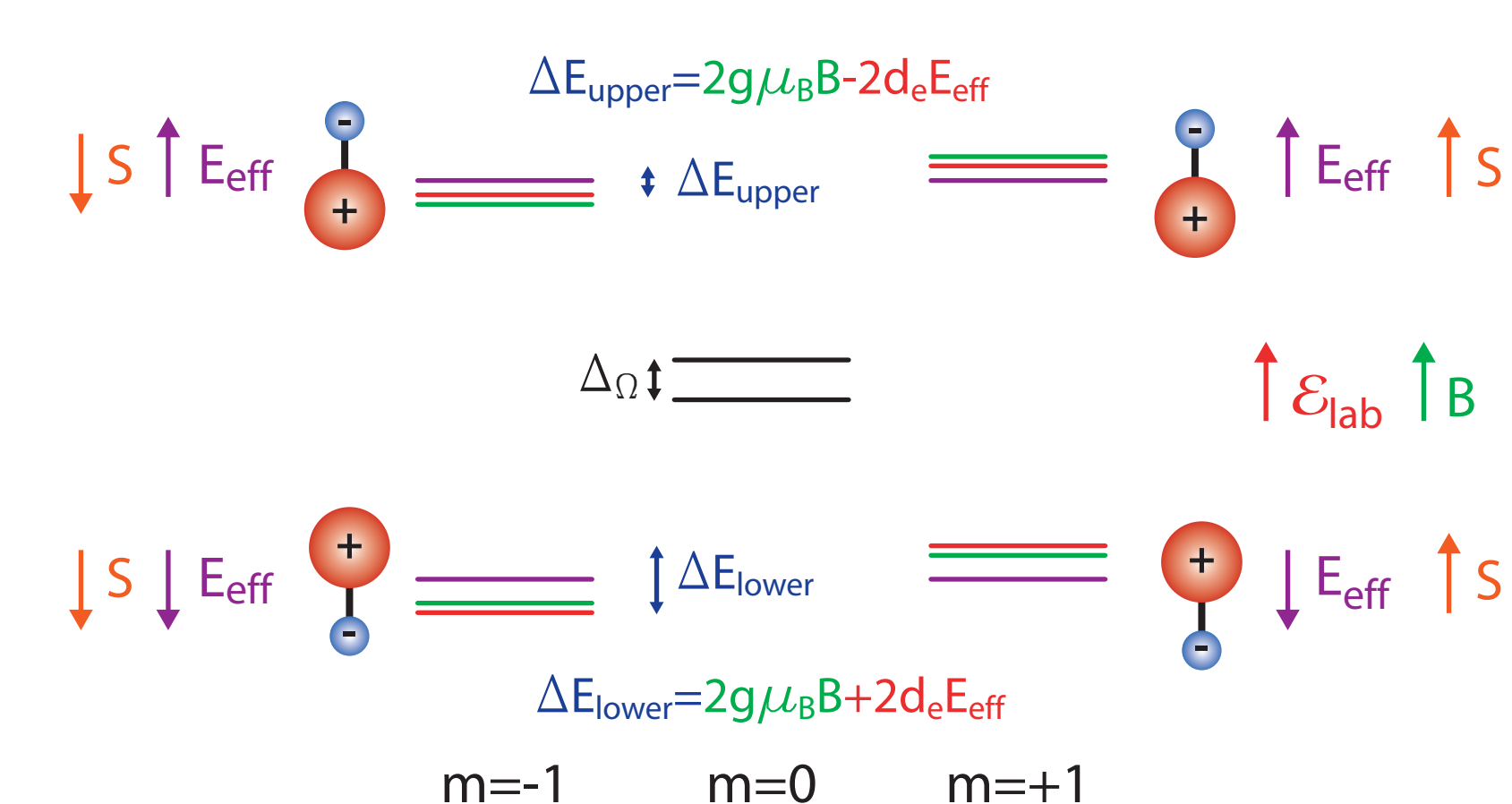


Statistical sensitivity:  $\delta d_e = \frac{1}{2E_{eff}} \frac{\hbar}{\tau \sqrt{N} T}$

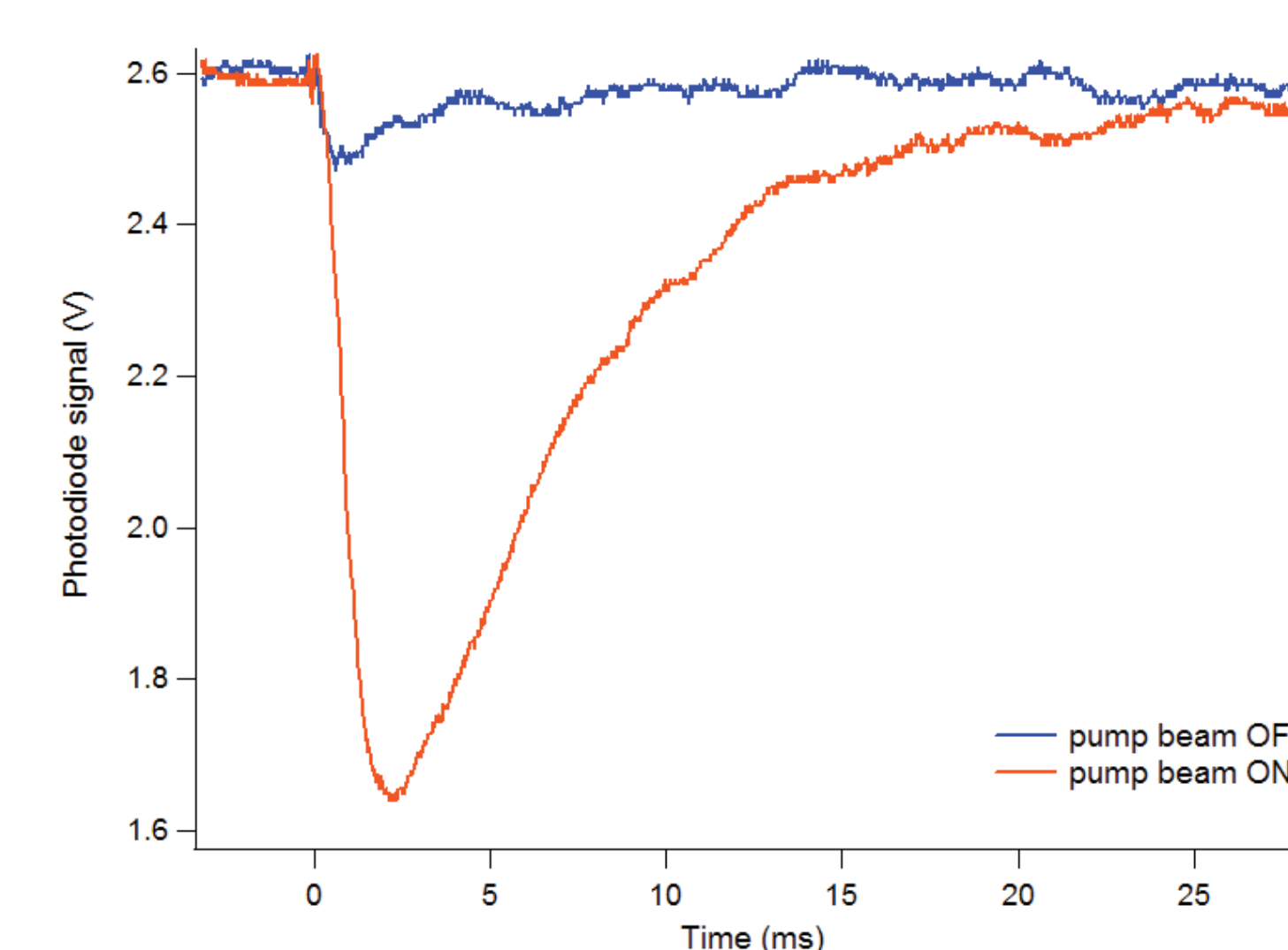
internal electric field  $\rightarrow E_{eff}$ , coherence time  $\rightarrow \tau$ , counting rate  $\rightarrow N$ , integration time  $\rightarrow T$



### Energy shifts in $J=1$ level of H state



## Optical pumping into H state

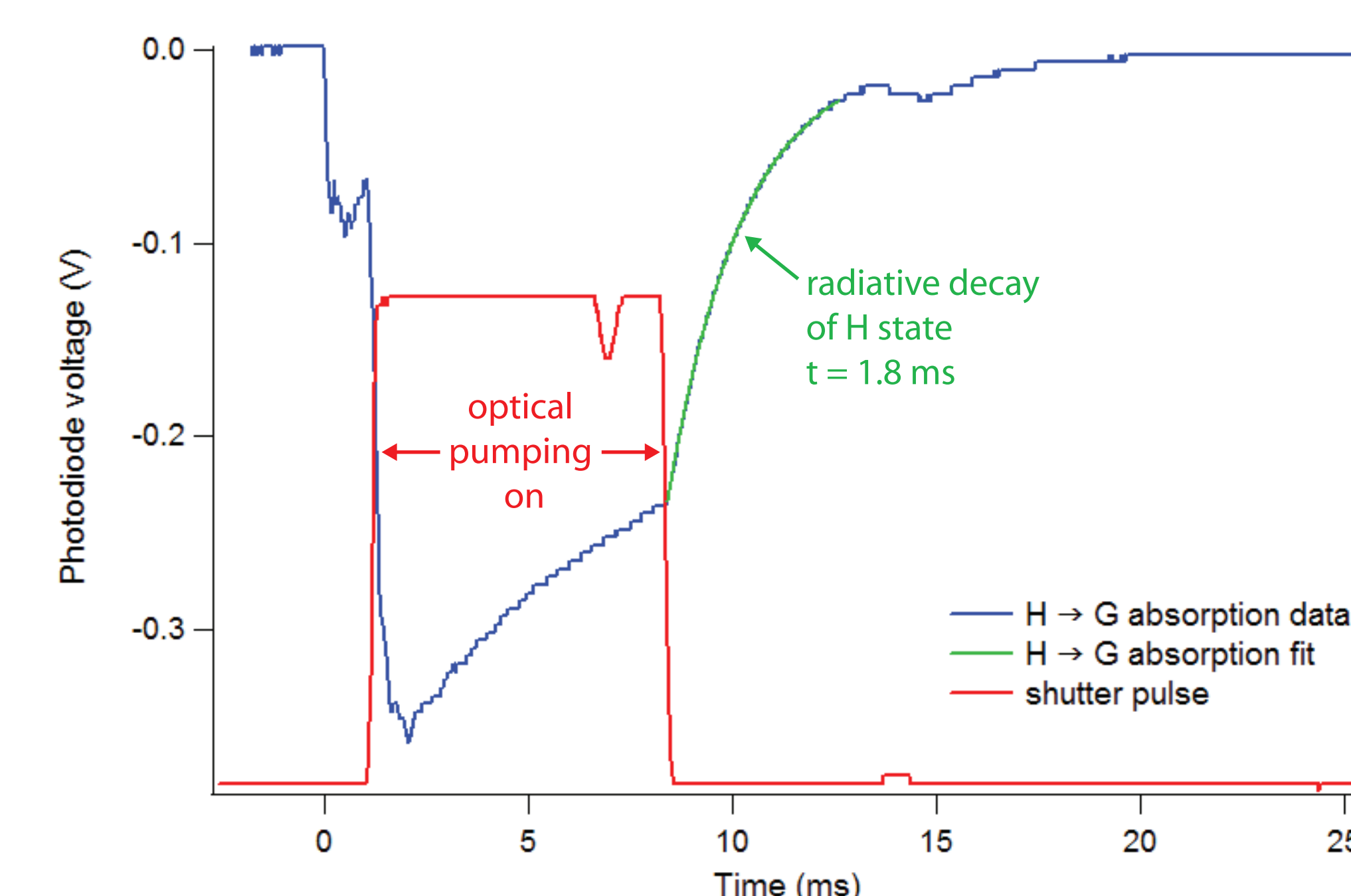


The plot at left shows absorption on the  $H \rightarrow G$  transition with (orange) and without (blue) the presence of a pump beam tuned to the  $X \rightarrow A$  transition.

The data shown here were taken in a buffer gas cell. We have also demonstrated optical pumping in a beam.

## H state lifetime

An exponential fit to the absorption data after the optical pumping pulse is turned off yields a lifetime of  $\tau = 1.8 \text{ ms}$ , which gives a lower bound on the H state radiative lifetime.



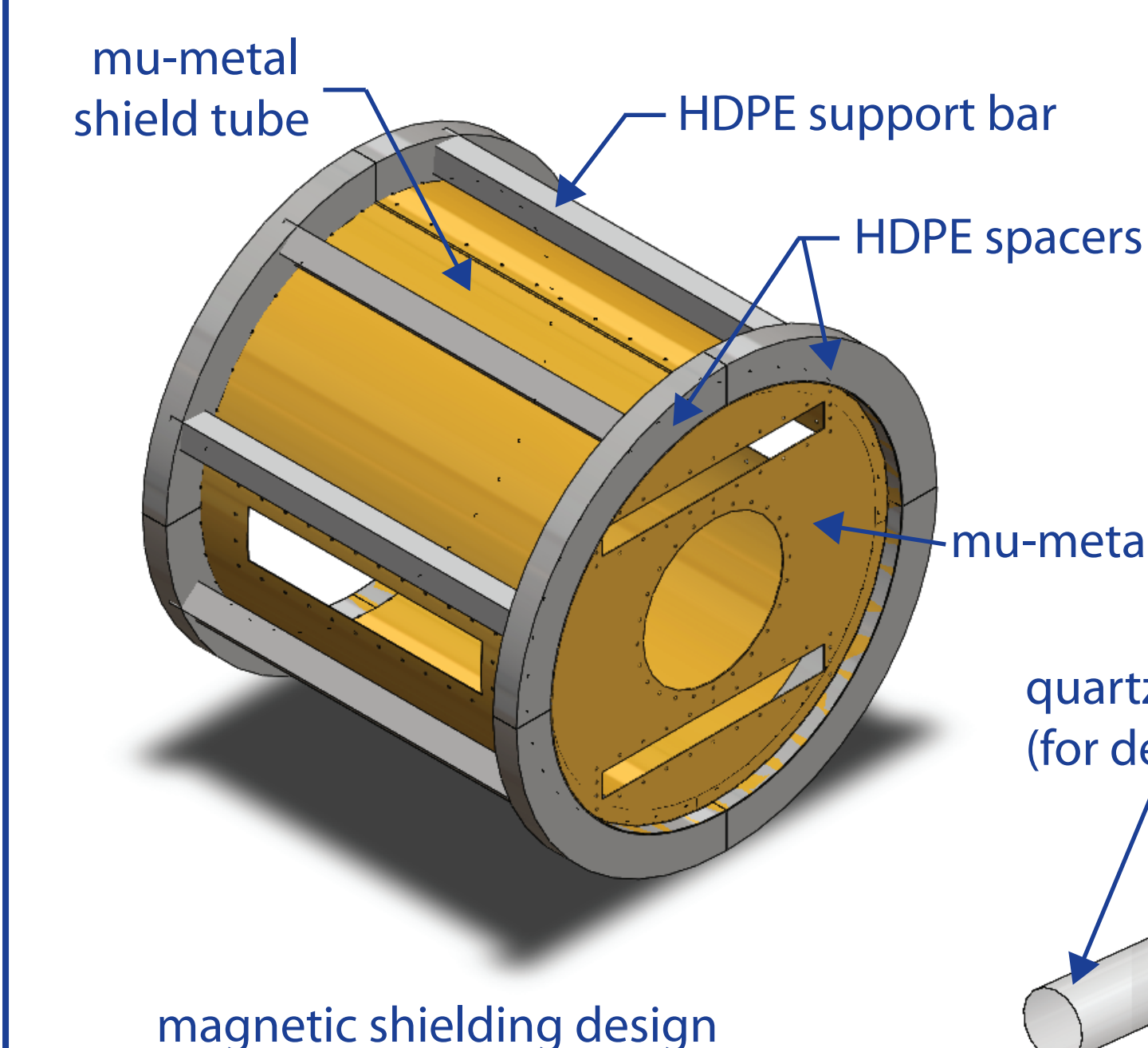
## Current work

- High flux beam source development
  - Demonstrated production of atomic Yb beam with  $4 \text{ K}$  He buffer gas
  - He beam studies started with Yb
  - In progress
    - Detailed beam studies and optimization
    - Investigation of neon buffer gas<sup>4</sup>
    - Demonstration of high flux molecular beam

<sup>4</sup>D. Patterson, J. Rasmussen, and J. M. Doyle, arXiv:0812.2212 [physics.atom-ph].

### Interaction region

- Transparent electric field plates (ITO-coated glass) under fabrication
- Interaction region vacuum chamber under fabrication
- Magnetic shielding design and fabrication in progress



### Laser systems and spectroscopy

- Lasers for state preparation and detection
- Digital locking system that can keep laser frequencies stable for  $> 12 \text{ hrs}$
- Measurements of dipole moment and g-factor for H state
- Demonstration of state preparation and detection of the accumulated phase

