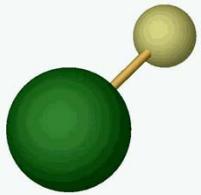


The electric dipole moment of the electron

- Electric dipole moments (EDMs) and new particle physics
- How to detect an EDM
- Upper bound from the ACME experiment
- What it means, and where we go next

DeMille



Group

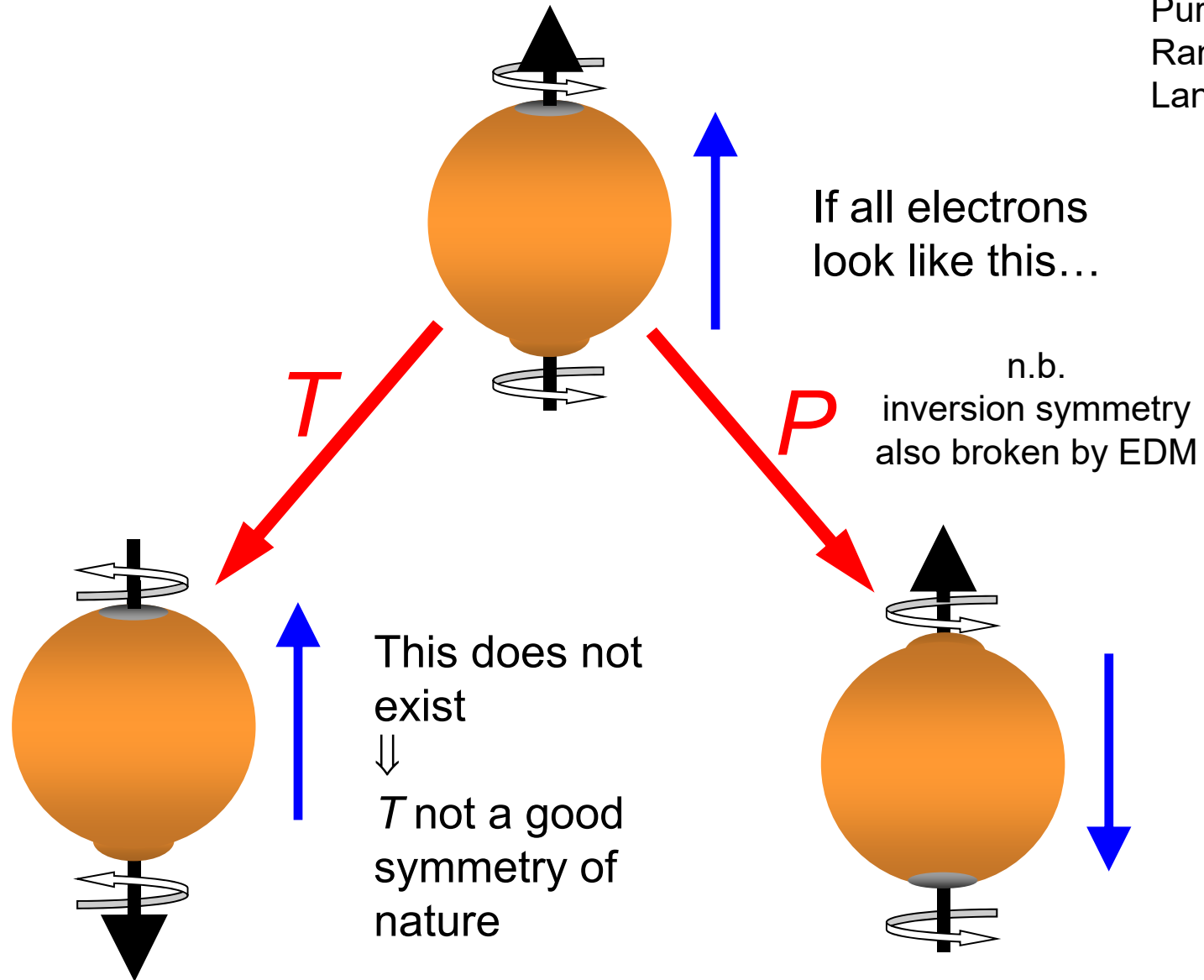
Dave DeMille
Physics Department
Yale University

Funding
NSF



An EDM violates time-reversal symmetry

Purcell
Ramsey
Landau



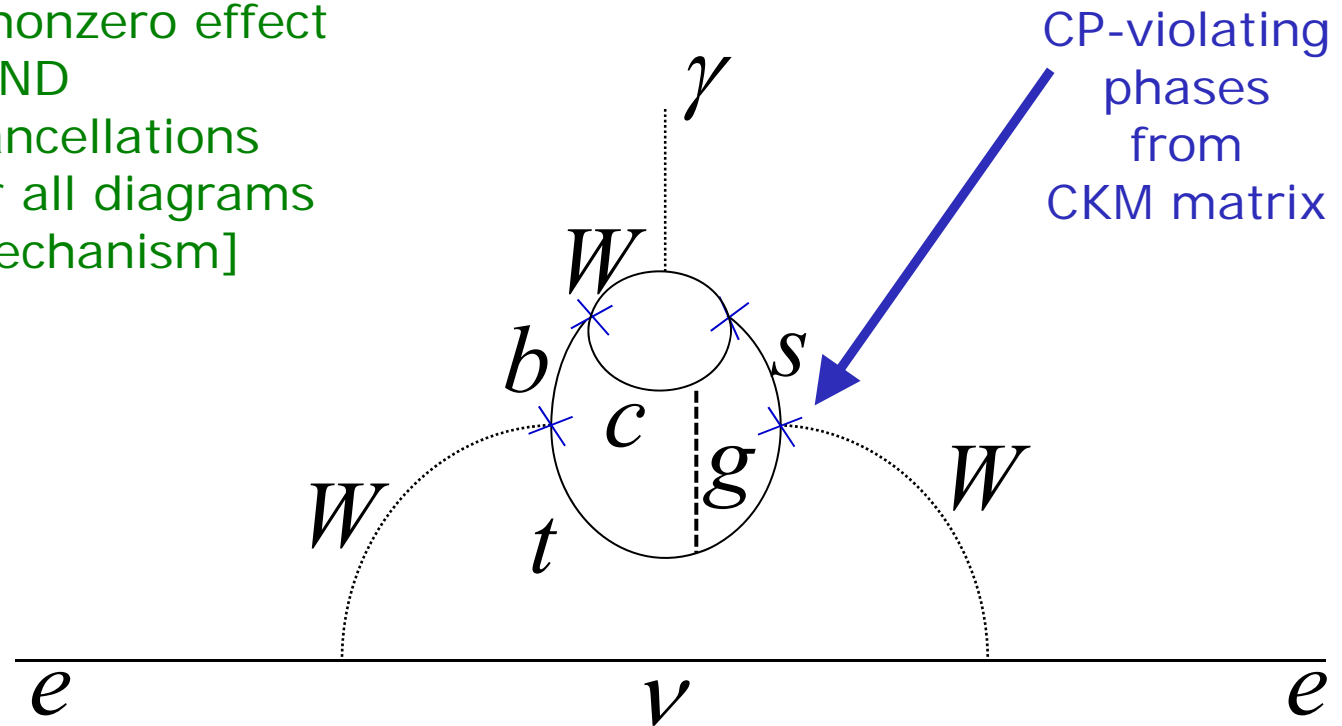
CPT theorem \Rightarrow T-violation = CP-violation

Q. How can a T -violating electron EDM arise?

A. From radiative corrections, with CP-violating interactions

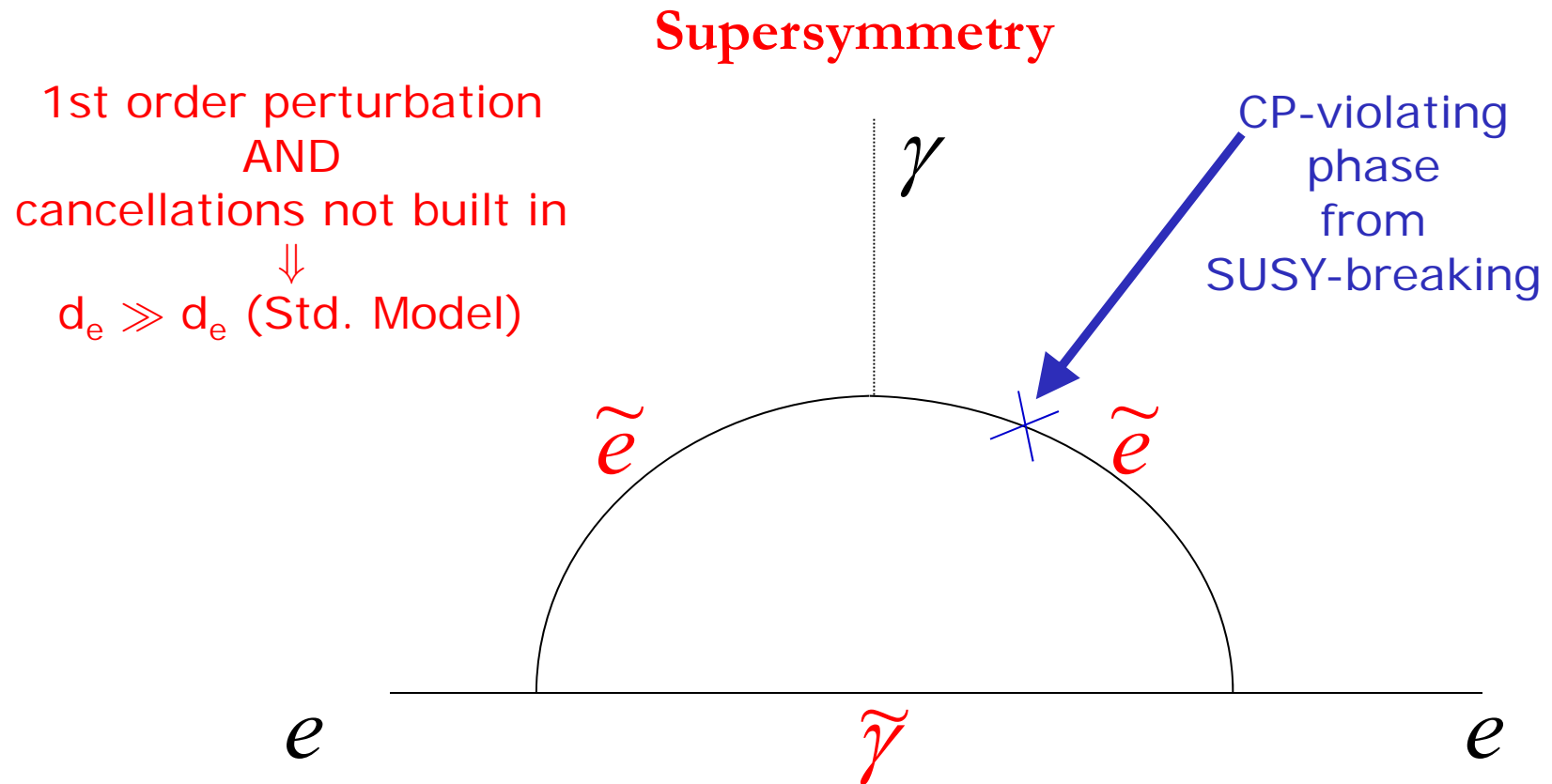
Standard Model

4-loops for nonzero effect
AND
severe cancellations
in sum over all diagrams
[GIM mechanism]

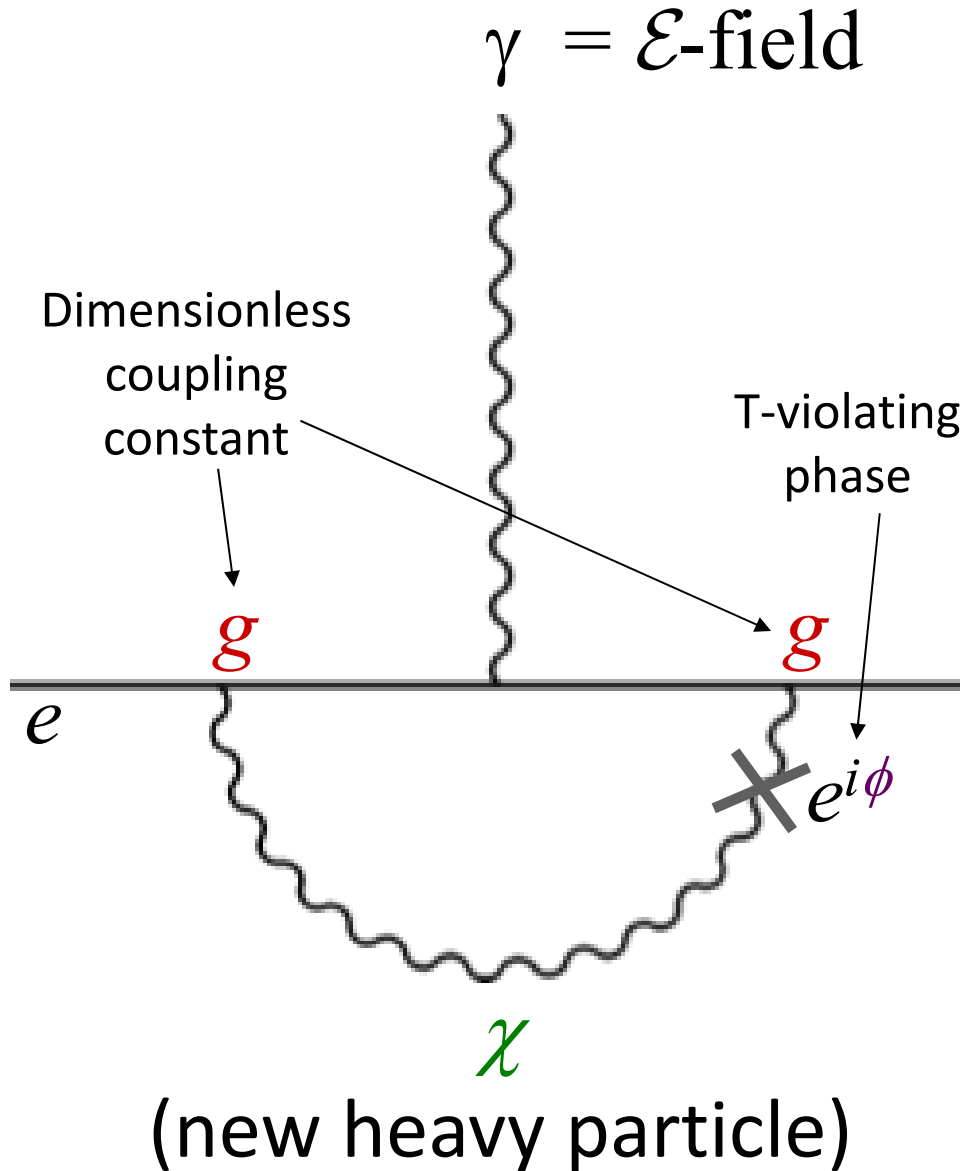


Q. How can a T -violating electron EDM arise?

A. From radiative corrections, with CP-violating interactions



Dimensional estimate for size of EDM



typical e-EDM

$$d_e \sim \mu_B \left(\frac{g^2}{2\pi} \right)^N \left(\frac{m_e}{m_\chi} \right)^2 \sin \phi$$

$N =$ # loops

“natural” assumptions

$$g^2/\hbar c \approx \alpha$$

$$\sin(\phi) \sim 1$$

$$m_\chi \sim 1 \text{ TeV}$$



$$d_e \sim 30 \times \text{current limit}$$

(for $N = 1$ loop)

Artist's impression of an electron EDM...



Note the resemblance to 0...

any EDM must be VERY small

QED radiative correction scale

$$[\alpha/2\pi] \mu_B/e \sim 10^{-13} \text{ cm}$$

:

Size of earth $\sim 10^4$ km

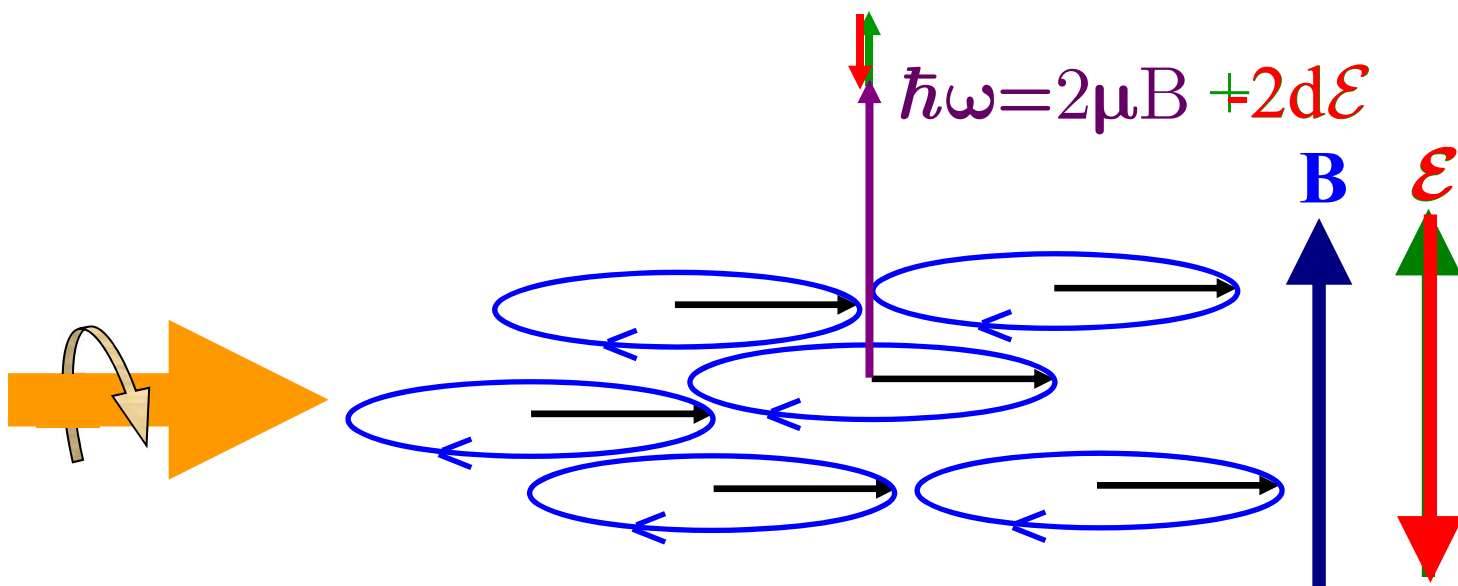
::

eEDM charge displacement

:

<20 nanometers

General method to detect an EDM



Energy level picture:

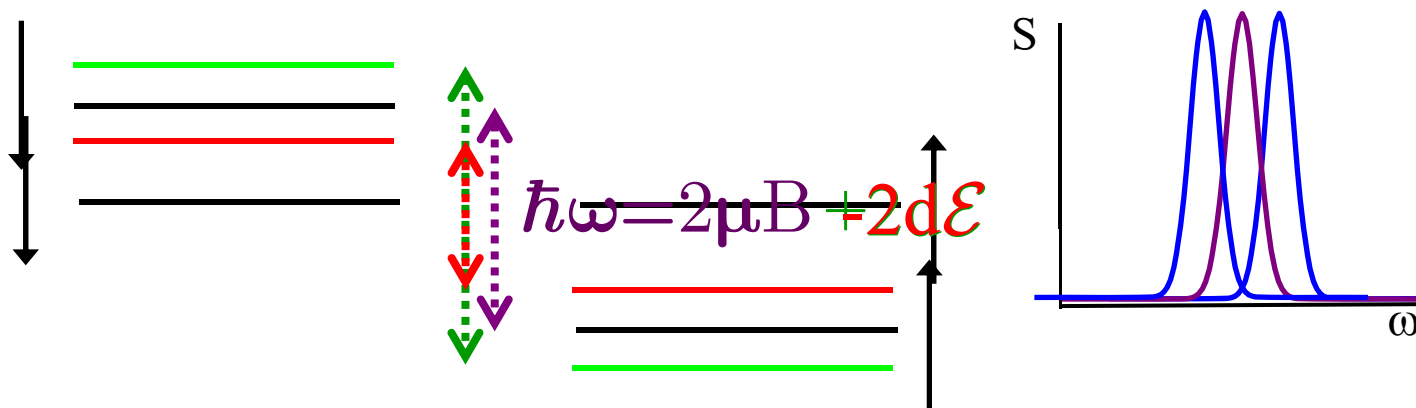
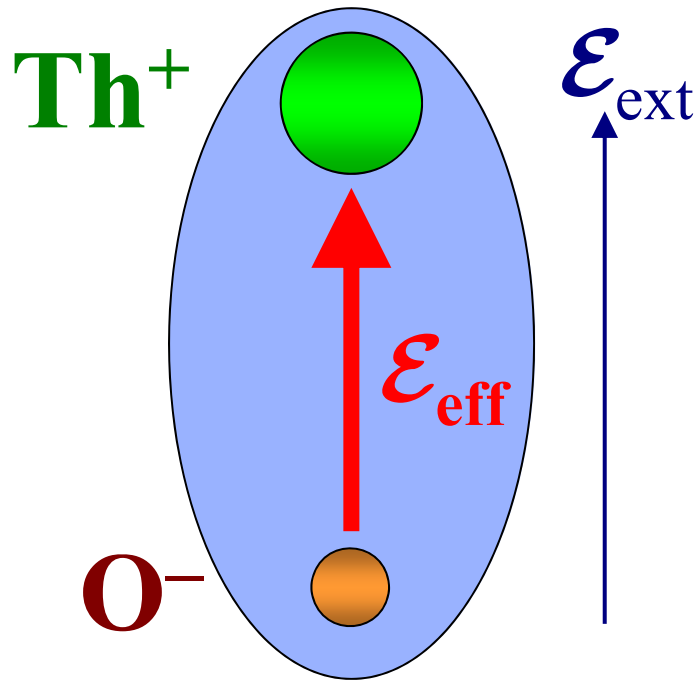


Figure of merit:

$$\frac{\textit{shift}}{\textit{resolution}} = \frac{d\mathcal{E}}{(1/\tau_{coh})(S/N)^{-1}} \propto \mathcal{E} \cdot \tau_{coh} \cdot \sqrt{\dot{N} \cdot T_{int}}$$

Amplifying the electric field \mathcal{E} with a polar molecule



Small energy splittings
(rotational levels)
enables polarization $\mathcal{P} \sim 100\%$
with $\mathcal{E}_{\text{ext}} \sim 10 \text{ V/cm}$

Inside polarized molecule, eEDM acted on by
 $\mathcal{E}_{\text{eff}} \sim \mathcal{P} \alpha^2 Z^3 e/a_0^2$ due to relativistic motion

P. Sandars
1965

$\mathcal{E}_{\text{eff}} \cong 80 \text{ GV/cm}$ for ThO* [near theoretical maximum]

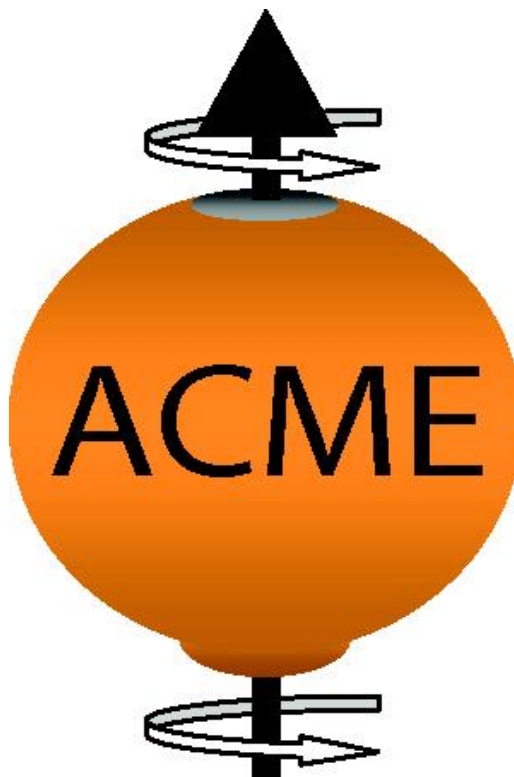
Meyer & Bohn (2008); Skripnikov, Petrov & Titov (2013, 2015); Fleig & Nayak (2014)

Requires unpaired electron spin(s): chemical free radical

Advanced Cold-Molecule Electron EDM



Yale University



Harvard University



The ACME team 2009-2014



Ben
Spaun

Chris Overstreet Nick Hutzler Cris Panda

Adam West

Emil
Kirilov

Max
Parsons

Brendon
O'Leary

Jacob Baron

Paul Hess

Elizabeth Petrik

Wesley Campbell  Amar Vutha 

John Doyle

DPD

Gerald Gabrielse



Yulia
Gurevich

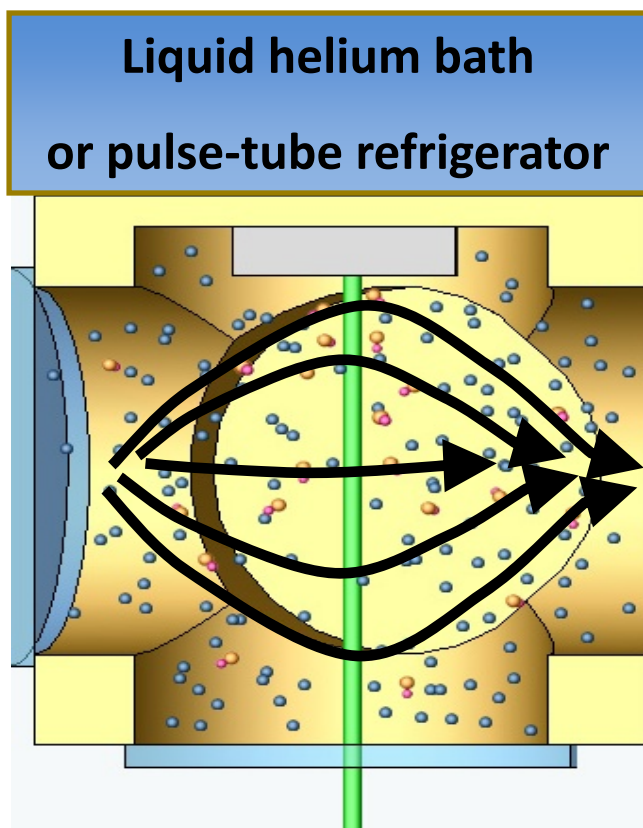


Ivan
Kozyryev



New molecular beam technology: Cryogenic Buffer Gas-cooled Beam (CBGB)

[Maxwell *et al.* PRL 2005; Patterson & Doyle J Chem Phys 2007;
Barry *et al.* PCCP 2011; Hutzler *et al.* PCCP 2011]



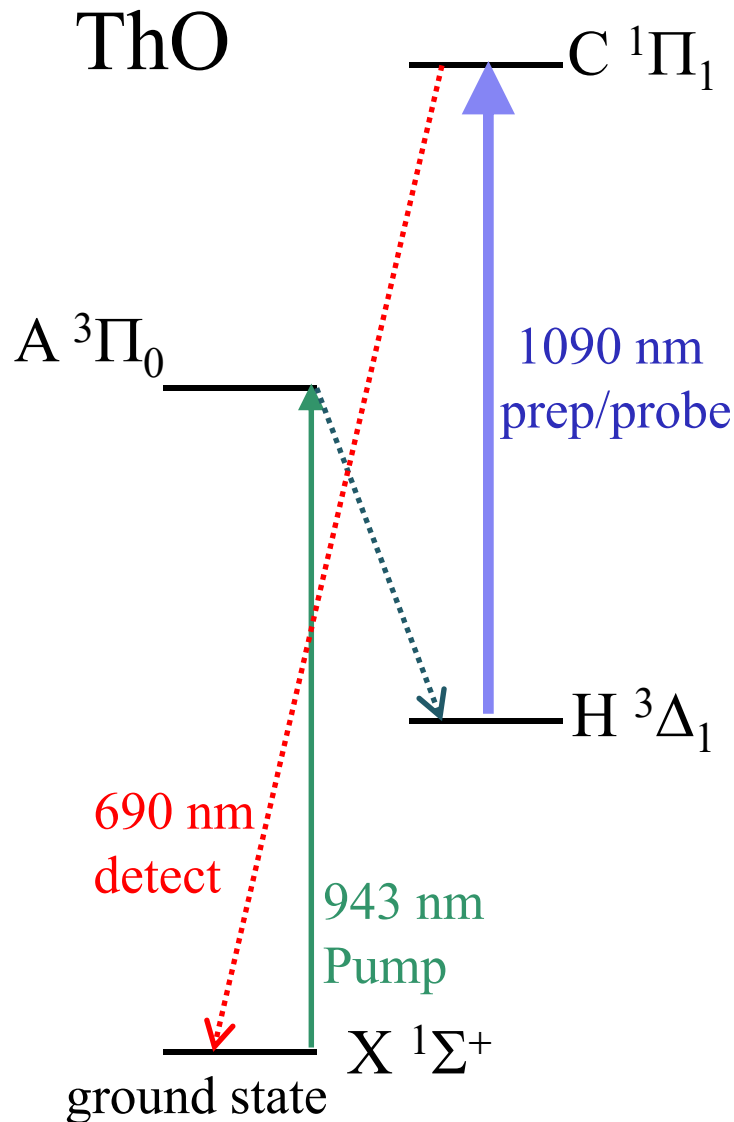
- Inject hot molecules (e.g. via laser ablation)
- Cool w/cryogenic buffer gas @high density
- **Efficient** extraction to beam via “wind” in cell: $10^{-4} \rightarrow 10\%-40\%$
- “Self-collimated” by extraction dynamics
- **Rotational cooling** in expansion: $T \sim 1 - 4\text{K}$
- **Moderately slow:** $v \sim 130-180 \text{ m/s}$

Beam brightness [=flux/divergence] $\sim 10^3 - 10^4 \times$ **larger**
vs. other sources for refractory/free radical species
typically $\sim 2 \times 10^{11} \text{ mol/sr/state/pulse}$ @ $\gtrsim 10 \text{ Hz}$ rep. rate

Used for **SrF**, **ThO**, BaF, YO, CaF, O₂, NH₃, Yb, ...

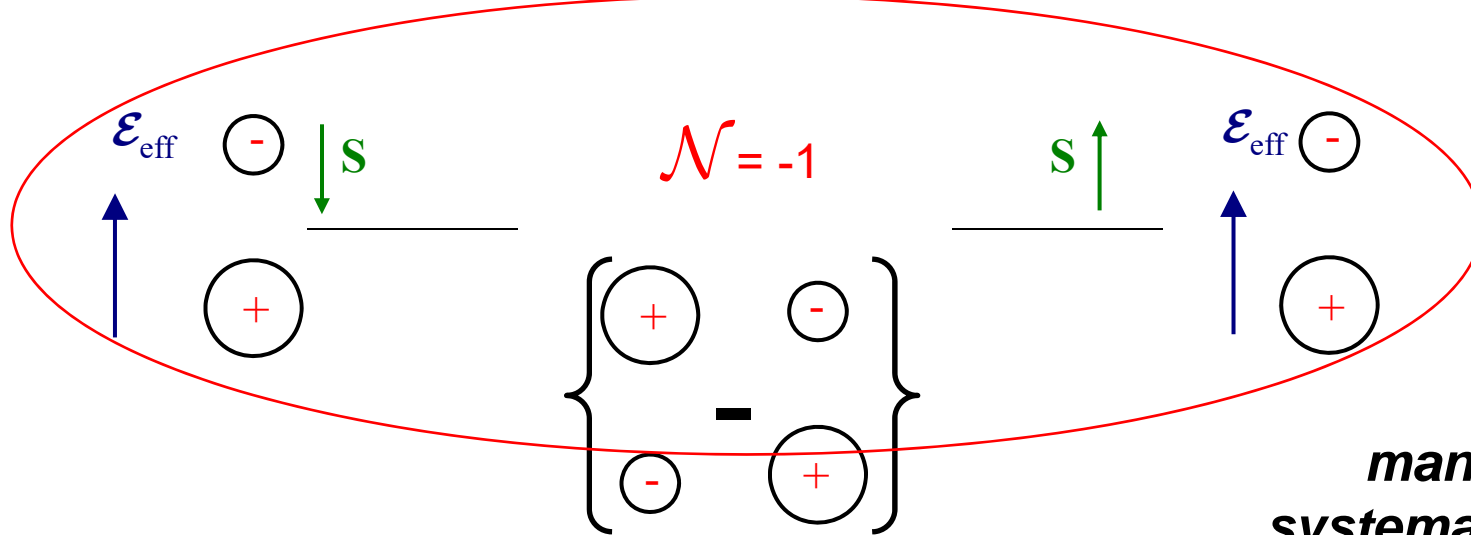
"New" molecular species: ThO^*

[A.C. Vutha *et al.* J. Phys B 2010]

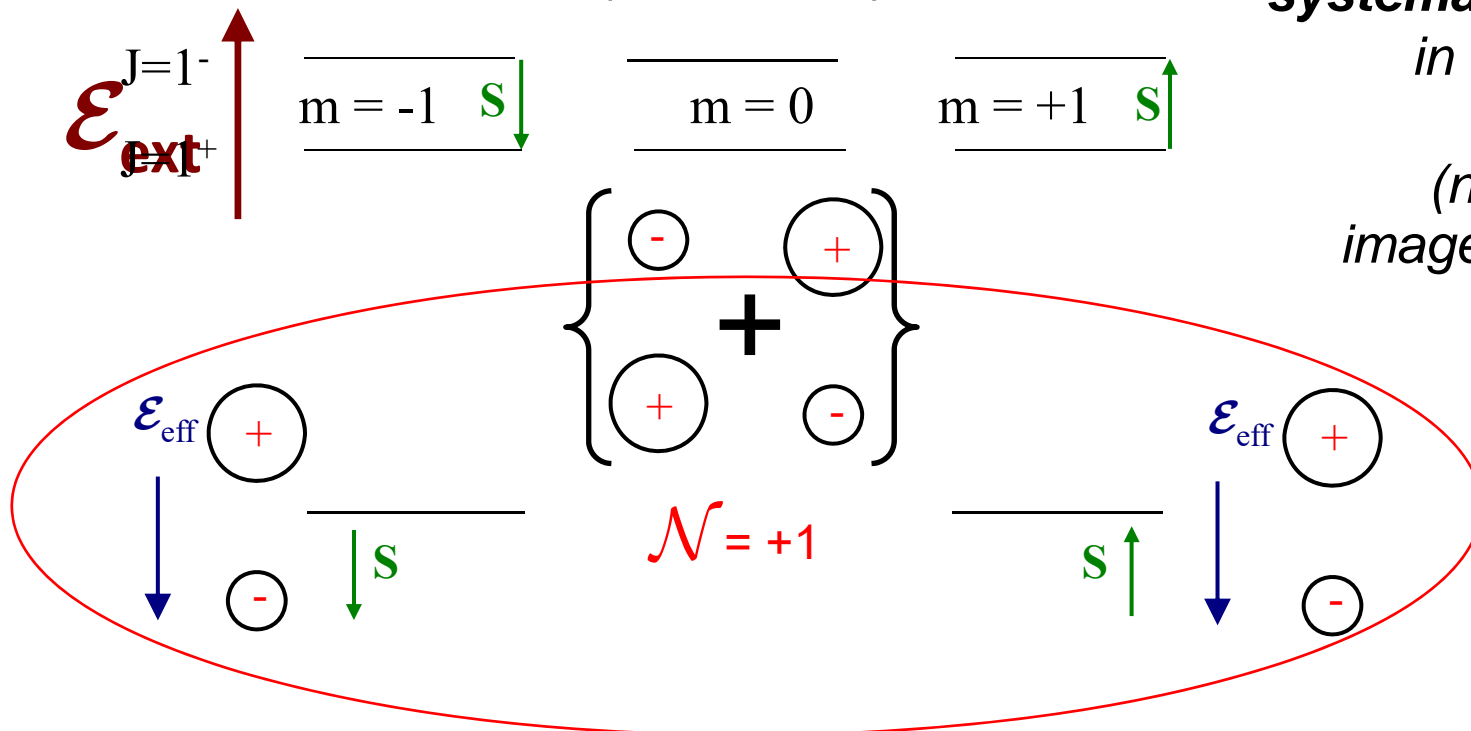


- **Largest effective internal \mathcal{E} : ~ 80 GV/cm**
[Titov *et al.* 2013/2015, Fleig *et al.* 2014]
- **Sufficient coherence time (~ 1.9 ms max)**
lifetime of metastable state $H^3\Delta_1$
- **Suppressed magnetic moment**
 $< 0.01 \mu_B$ in $H^3\Delta_1$ reduces B -field systematics
[Idea: Meyer, Bohn, Cornell *et al.* (JILA);
Measured: A.C. Vutha *et al.*, PRA 2011]
- **Omega-doublet co-magnetometer**
suppresses many possible systematics
- All spectroscopic data previously known
- State preparation and readout
w/standard, robust diode & fiber lasers
- Blue-shifted fluorescence from probe laser
 \Rightarrow no problem with backgrounds
- **High beam source yield**

EDM measurement with Ω -doublet states



many dangerous systematic errors cancel in comparison between (near-)mirror image $\mathcal{N} = \pm 1$ states



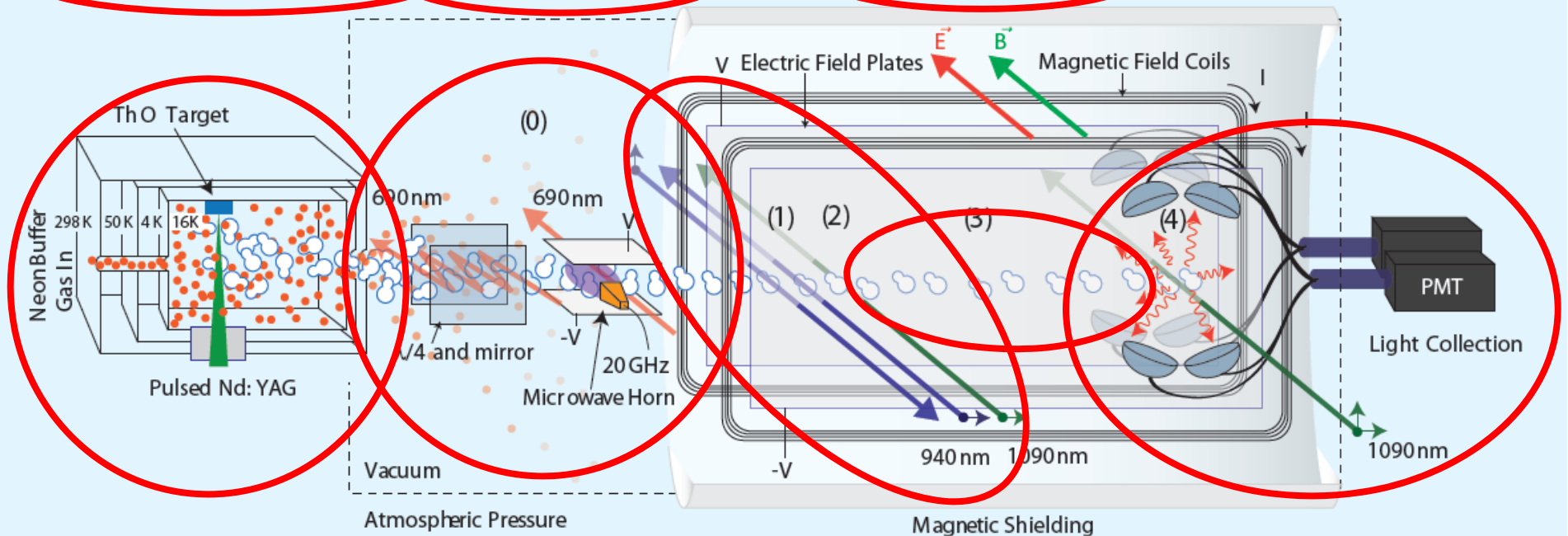
[DD et al., AIP Conf. Proc. 596, 2001]

ACME experimental schematic

Buffer Gas Beam Source

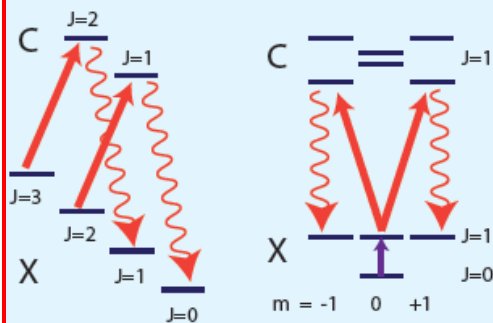
Rotational Cooling

Interaction Region

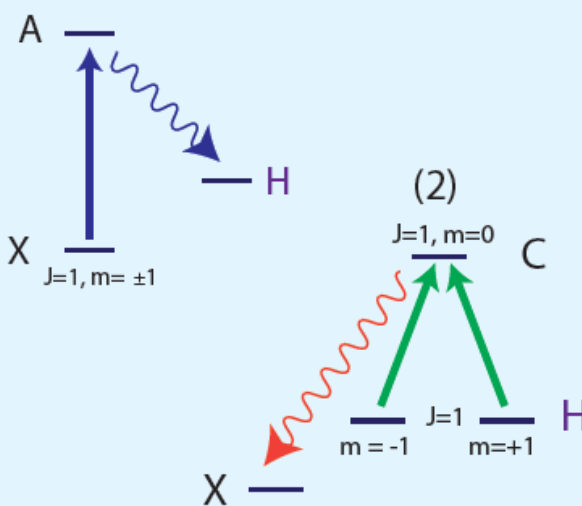


Rotational Cooling

(0)



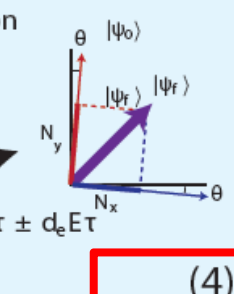
(1) State Preparation



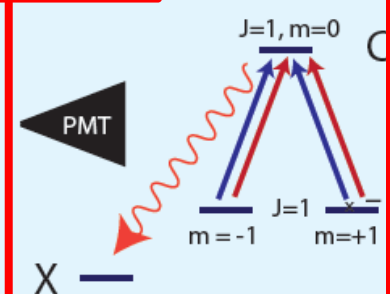
(3) Phase Precession

$$2\phi = \Delta E \tau / \hbar$$

$$\phi \approx g\mu_B \tau \pm d_e E \tau$$



(4) State Detection



ACME apparatus

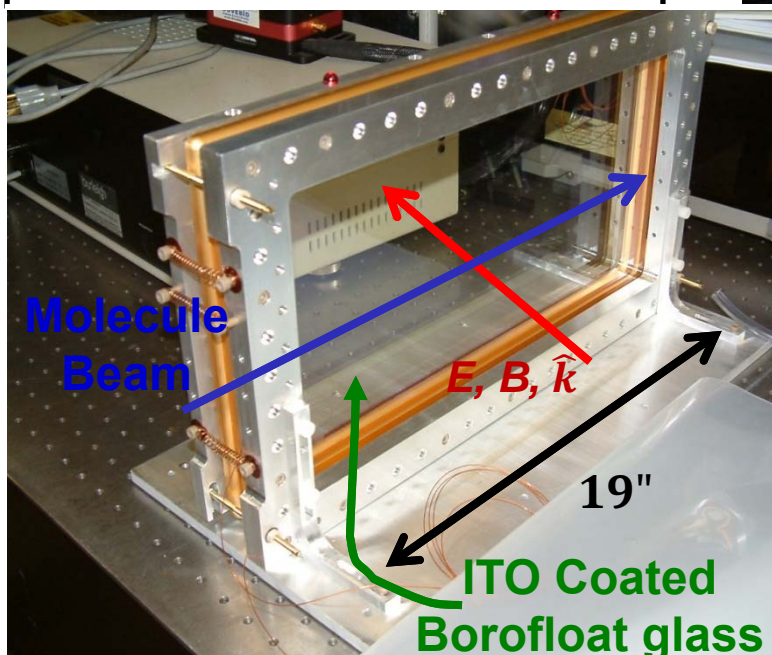
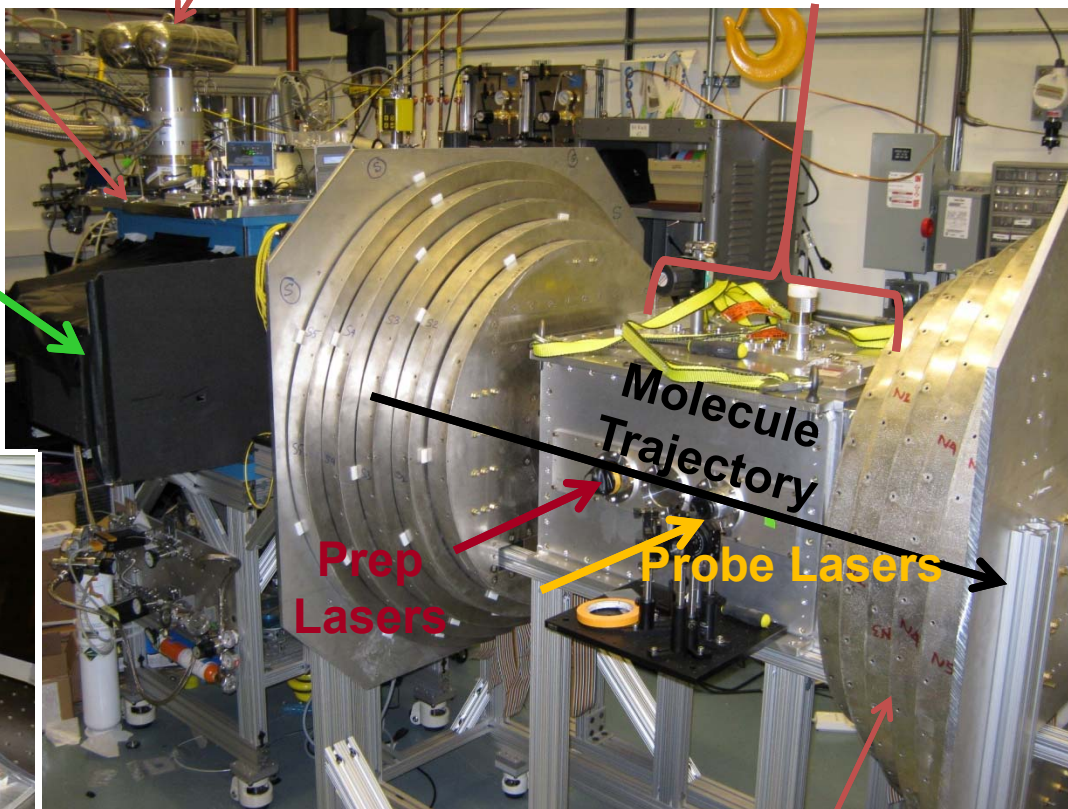
Molecular Beam Source

Pulse Tube Cryo Frig

Interaction Region:
internal \mathcal{E} -field plates & external
 B -field coils
(not shown here)

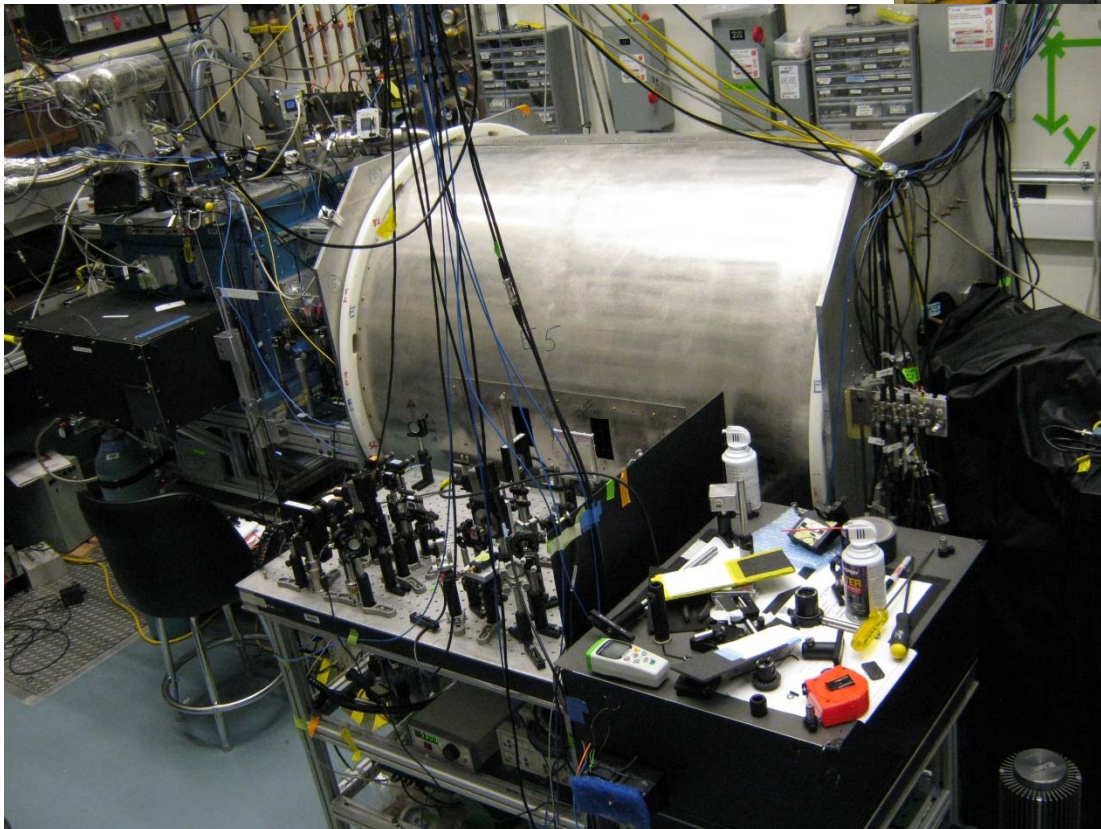
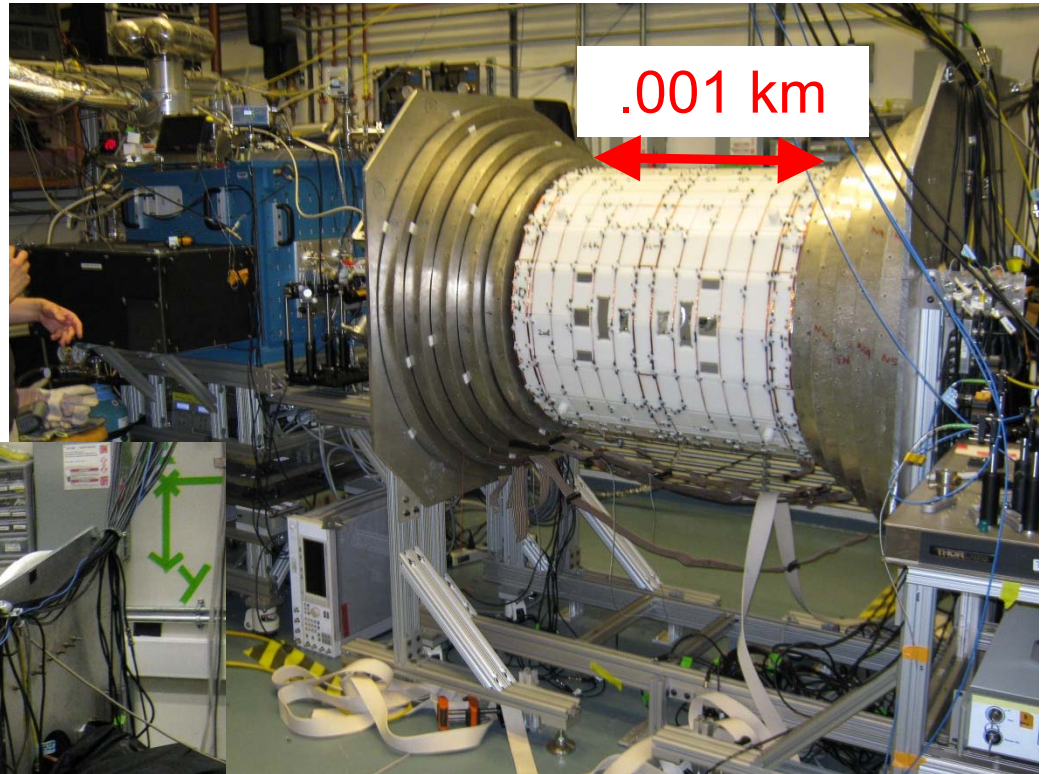
Pulsed ablation laser

Transparent Field Plates



ACME apparatus

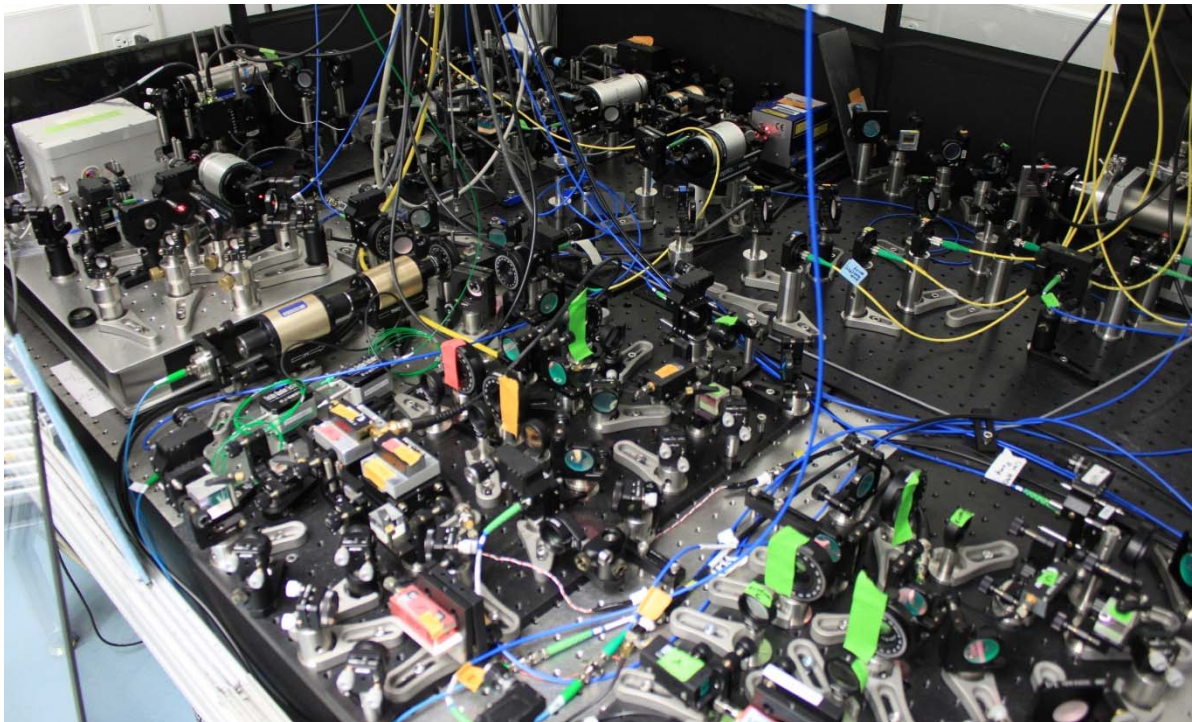
Magnetic field coils
(3 orthogonal components
& all first-order gradients)



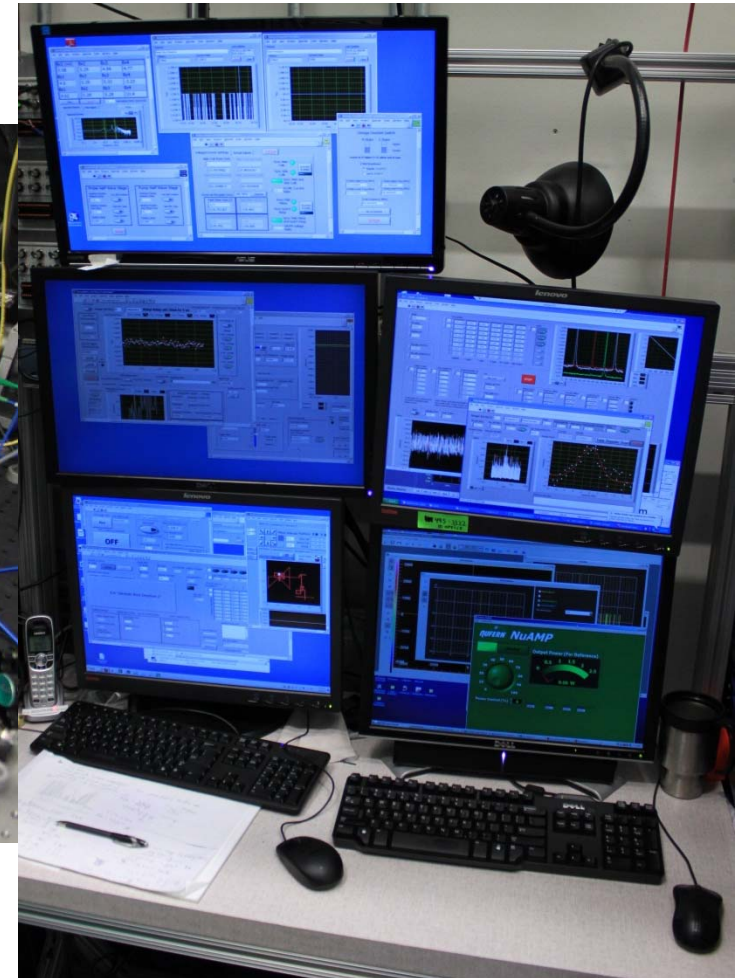
Complete
beam source
& magnetic shields
& last-stage optics

ACME apparatus

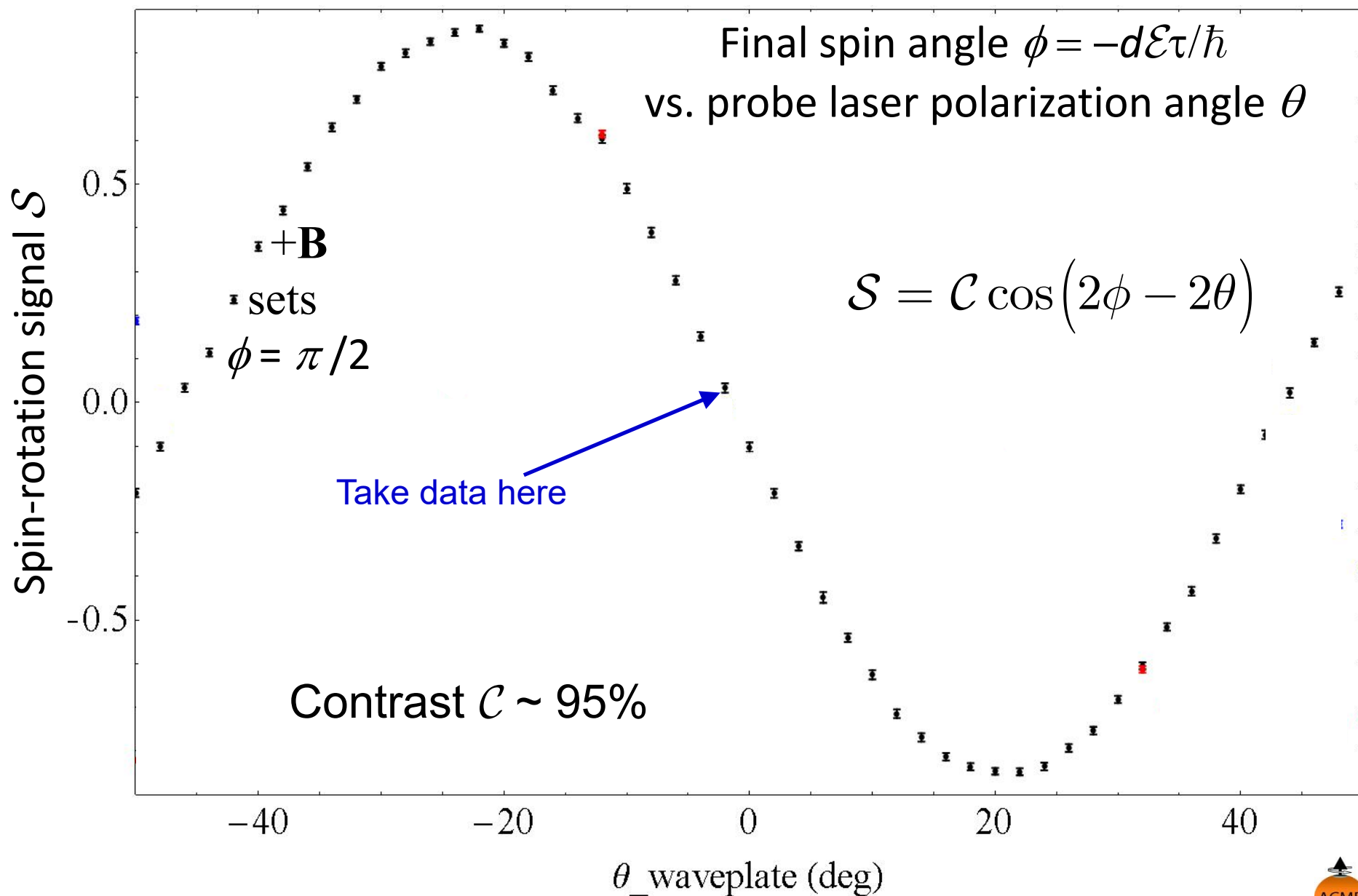
One of several optical tables w/
~ten lasers, dozens of modulators,
hundreds of meters of optical fiber, etc.
spread over two buildings



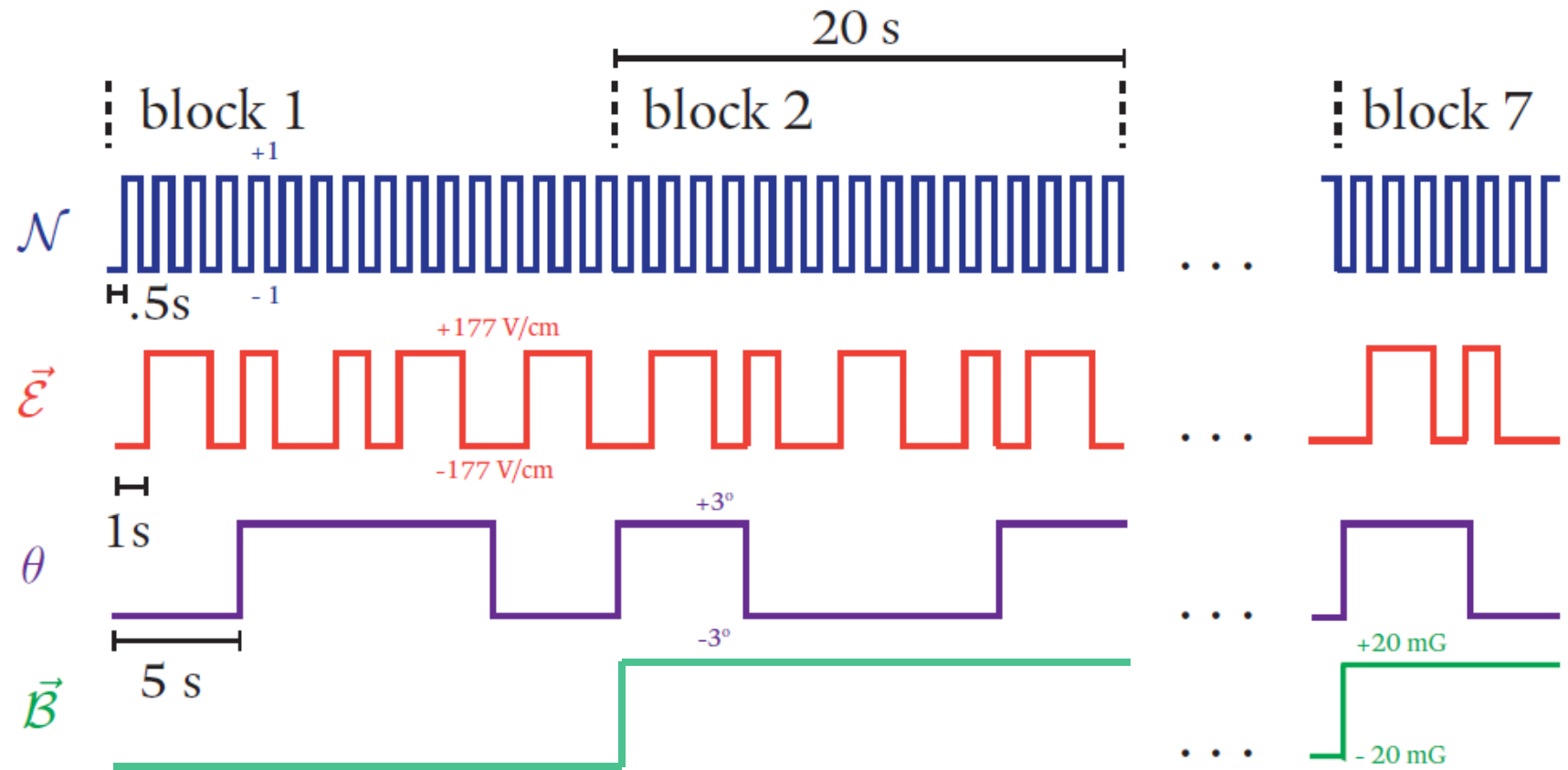
“control room”



Typical spin-rotation fringe signal



Primary machine settings to extract the EDM



--psuedo-random (pair-wise), interleaved reversals

Data sorting & systematic error analysis

Rewrite phase as components correlated w/switches:

$$\phi = \phi^0 + \phi^{\tilde{E}} + \phi^{\tilde{B}} + \phi^{\tilde{N}} + \phi^{\tilde{N}\tilde{E}} + \phi^{\tilde{N}\tilde{B}} + \phi^{\tilde{E}\tilde{B}} + \phi^{\tilde{N}\tilde{E}\tilde{B}}$$

Superscript means
"odd under this reversal"

EDM phase

other phases to
diagnose systematics

Switch-correlated phases measure physical contributions:

$$\phi^{\tilde{N}\tilde{E}} \propto d_e \mathcal{E}_{eff} + \frac{1}{2} \Delta g_N \mu_H \mathcal{B}_{leak} + \frac{1}{2} \Delta g_N \mu_H \mathcal{B}_{nr} \frac{\mathcal{E}_{nr}}{\mathcal{E}_0} + \dots$$

EDM

Spurious terms

Symmetries ensure systematic errors
due ONLY to experimental imperfections e.g.
--leakage current-induced \mathcal{B} -field $\mathcal{B}_{leak} \propto \mathcal{E}$
--non-reversing \mathcal{E} -field \mathcal{E}_{nr}
--etc.

Data analysis: diagnosing imperfections

Switch-correlated phases isolate physical contributions:

$$\phi^{\tilde{N}\tilde{\varepsilon}} \propto d_e \mathcal{E}_{eff} + \frac{1}{2} \Delta g_N \mu_H \mathcal{B}_{leak} + \frac{1}{2} \Delta g_N \mu_H \mathcal{B}_{nr} \frac{\varepsilon_{nr}}{\varepsilon_0} + \dots$$

EDM \nearrow
 \downarrow Experimental imperfections
 \downarrow

Most imperfections also appear in *other* correlated phases
BUT GREATLY AMPLIFIED

$$\phi^{\tilde{\varepsilon}} \propto \frac{1}{2} g \mu_H \mathcal{B}_{leak}$$

$g / \Delta g_N \sim 1000$

$$\phi^{\tilde{N}\tilde{\varepsilon}\tilde{\mathcal{B}}} \propto \frac{1}{2} \Delta g_N \mu_H \mathcal{B}_{nr} \frac{\varepsilon_{nr}}{\varepsilon_0}$$

$\mathcal{B} / \mathcal{B}_{nr} \sim 1000$

\Rightarrow "Other" correlated phases diagnose imperfections

Search strategy for systematic errors

Switch-correlated phases isolate physical contributions:

$$\phi^{\tilde{N}\tilde{e}} \propto d_e \mathcal{E}_{eff} + \frac{1}{2} \Delta g_N \mu_H \mathcal{B}_{leak} + \frac{1}{2} \Delta g_N \mu_H \mathcal{B}_{nr} \frac{\mathcal{E}_{nr}}{\mathcal{E}_0} + \dots$$

EDM \nearrow

\mathcal{B}_{leak} \swarrow

$\mathcal{B}_{nr} \frac{\mathcal{E}_{nr}}{\mathcal{E}_0}$ \swarrow

Experimental imperfections \swarrow

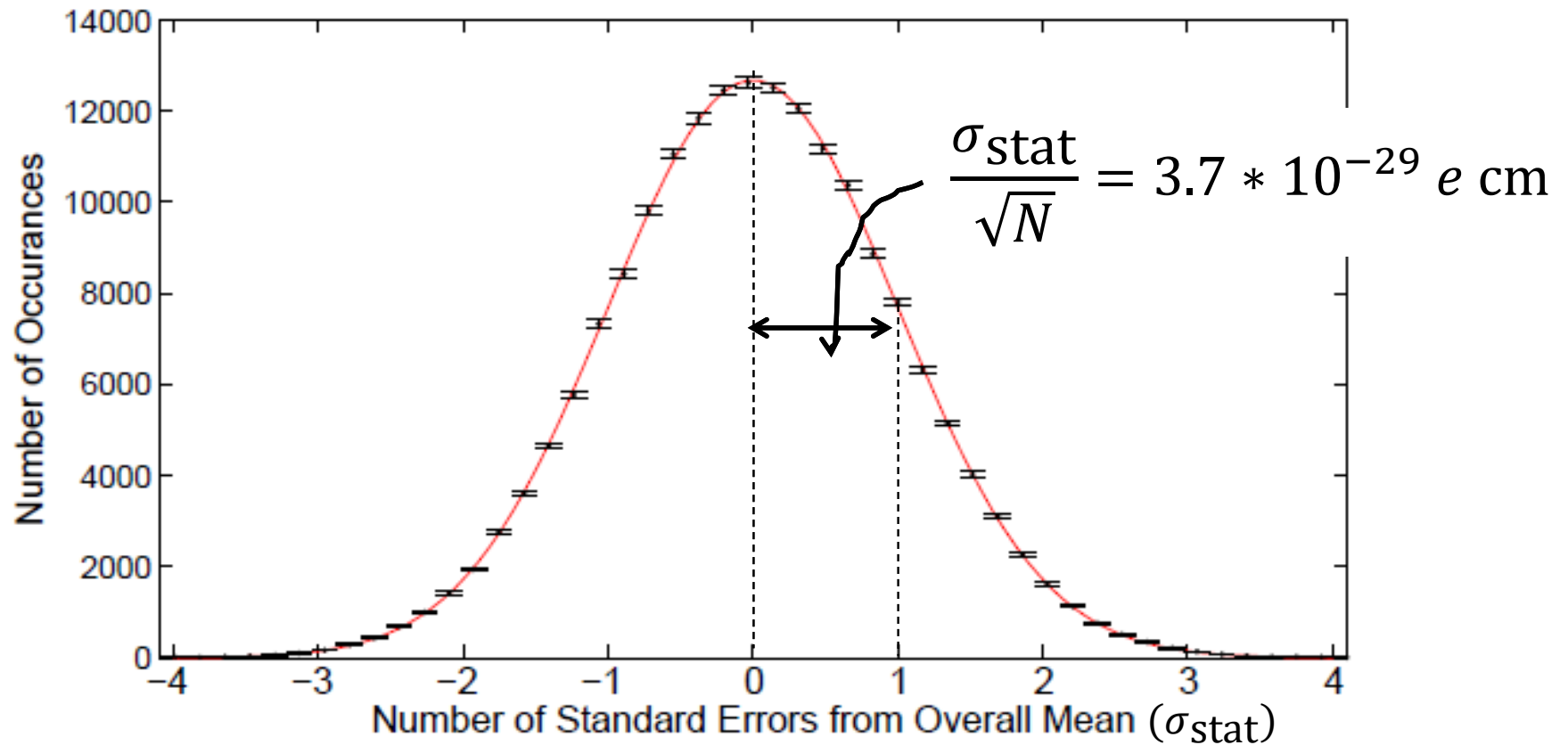
\dots \swarrow

But... what about terms we don't anticipate?

Strategies:

- **Change all possible parameters** that shouldn't affect EDM (\mathcal{E} magnitude, \mathcal{B} magnitude, global polarization, etc. etc.) but *might* couple to unanticipated imperfections
 - **Deliberately amplify imperfections**, understand any changes in correlated phases

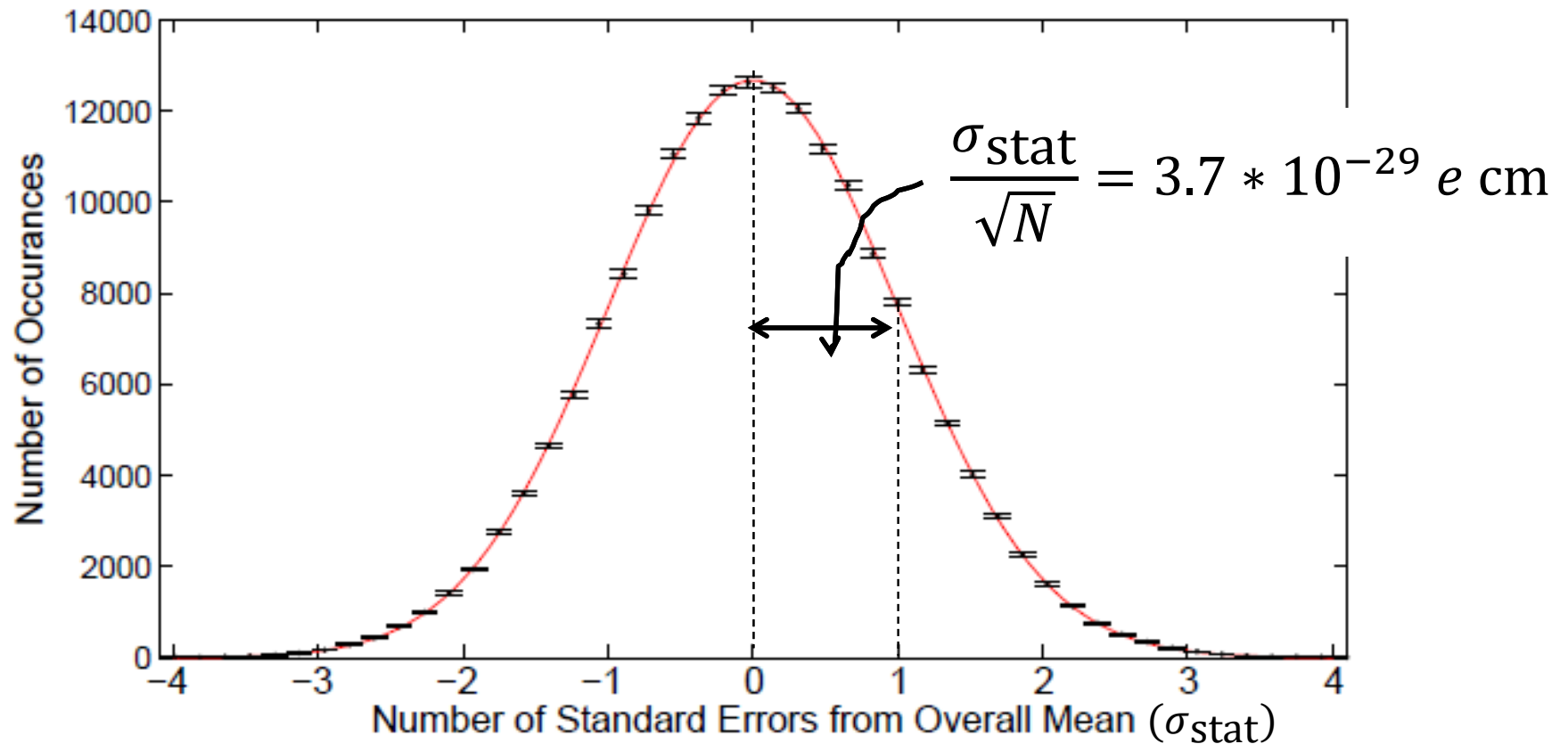
ACME electron EDM data



Blind analysis: randomly chosen offset added to data until analysis complete

$$d_e = (??? \pm 3.7_{\text{stat}} \pm 2.5_{\text{syst}}) \times 10^{-29} e \text{ cm}$$

ACME electron EDM data



Blind analysis: randomly chosen offset added to data until analysis complete

$$d_e = (-2.1 \pm 3.7_{\text{stat}} \pm 2.5_{\text{syst}}) \times 10^{-29} e \text{ cm}$$

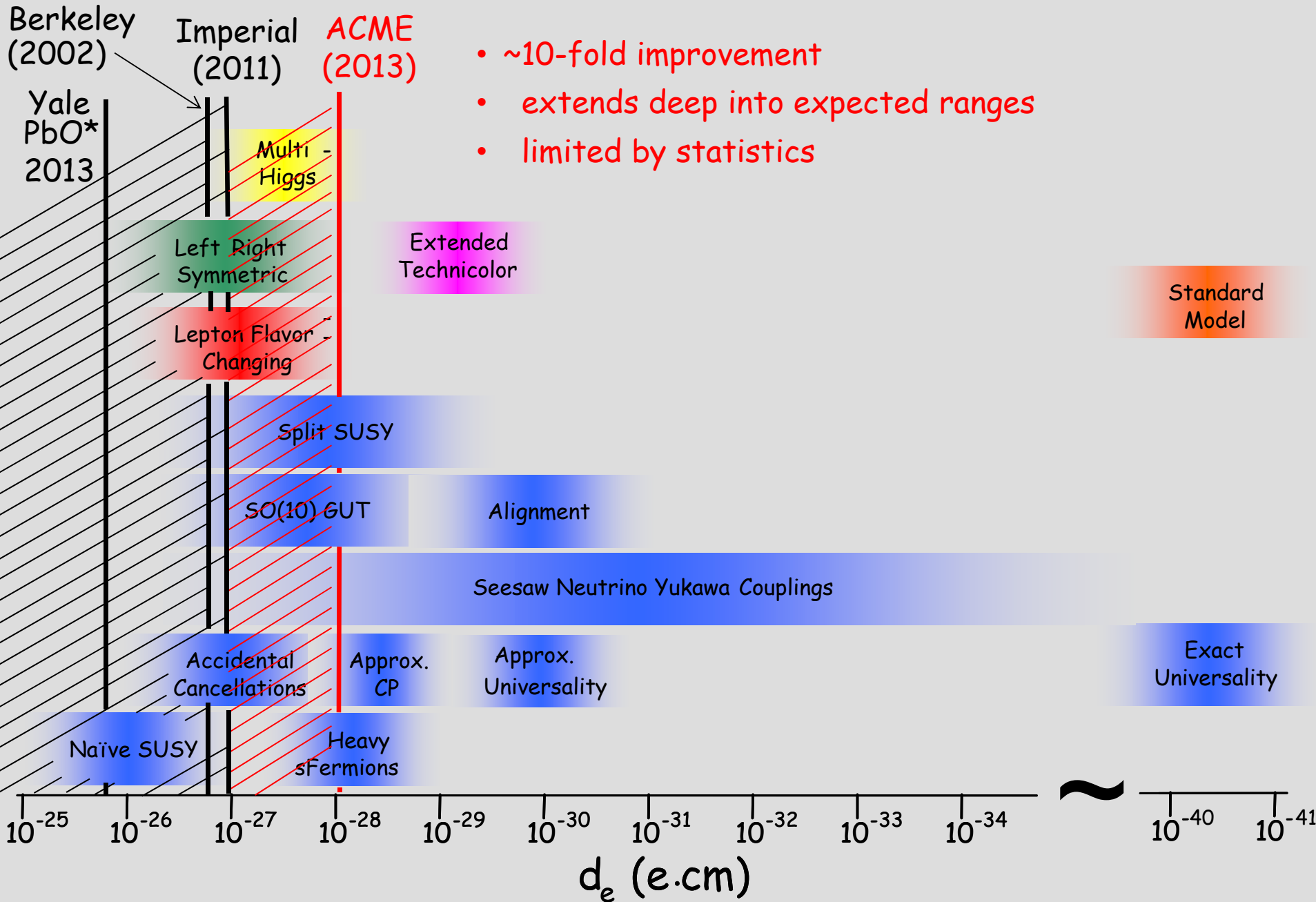
Consistent with zero
 \Rightarrow set upper limit

$$|d_e| < 9 \times 10^{-29} e \text{ cm}$$

J. Baron *et al.*,
 Science (2014)

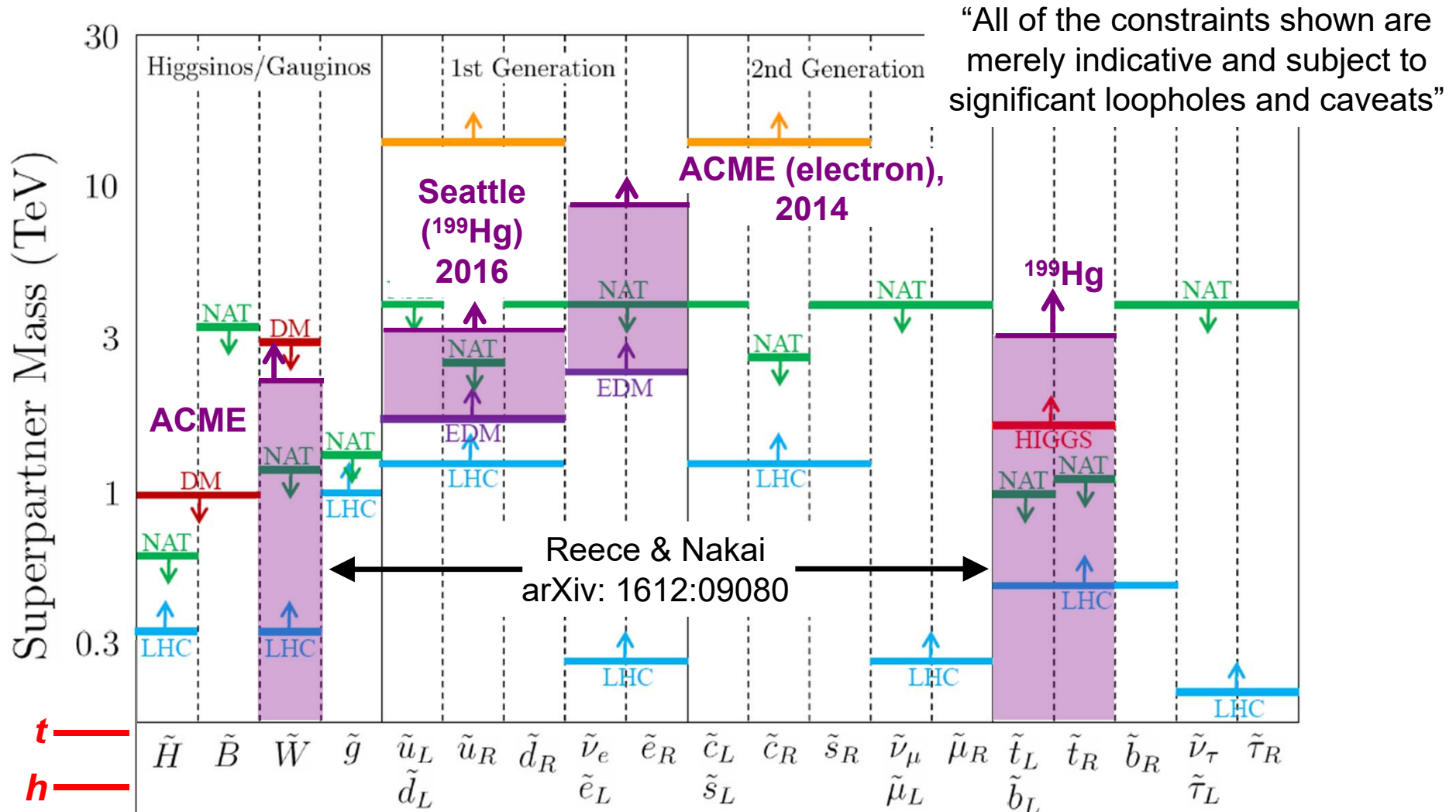


New upper bound on electron EDM from ACME



Impact of EDMs in particle physics

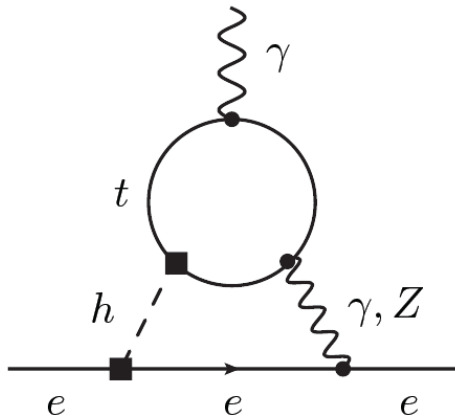
J. Feng: "Naturalness and the status of SUSY", Annu. Rev. Nucl. Part. Sci. (2013)



EDM results push SUSY scale into “unnatural” regions
 --potential for imminent discovery...?

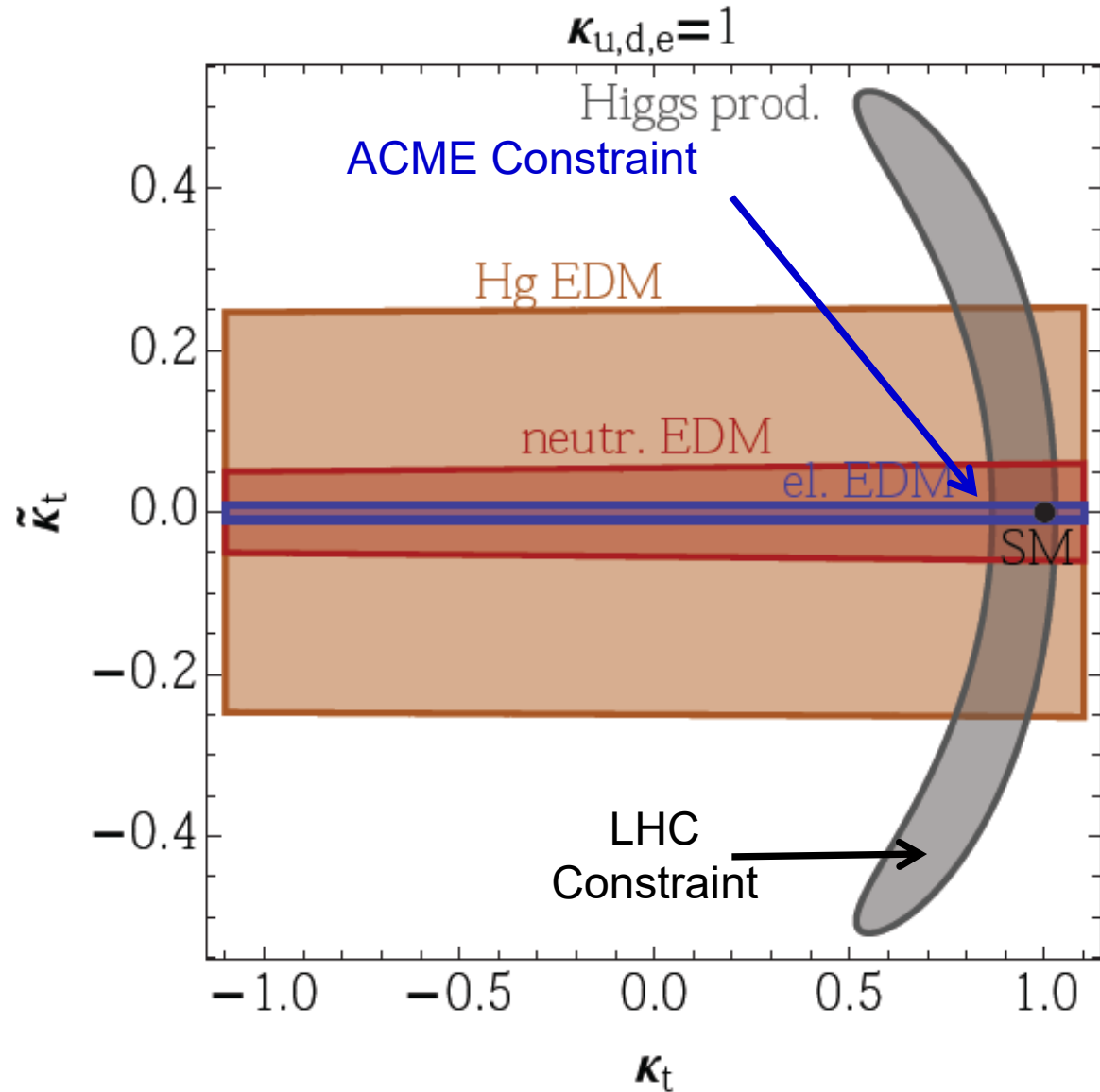
What does the $eEDM$ limit mean for particle physics?

Diagrams with **known** SM particles can rule out non-SM couplings



Example:
CP-violating
Higgs-top coupling

Brod *et al.*,
arXiv: 1310.1385



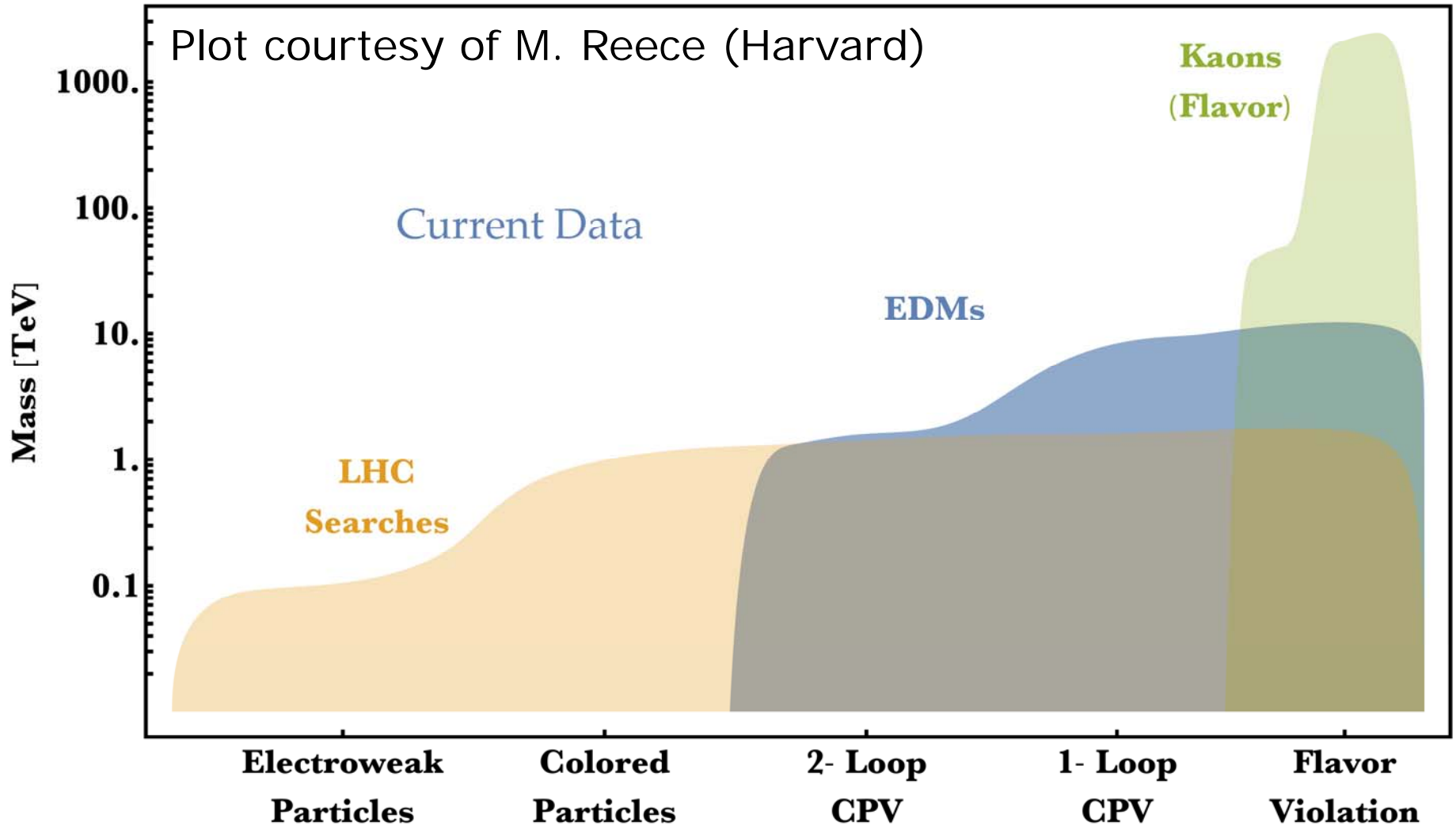
CP-odd/CP-even Higgs-top coupling $< 1\%$ from ACME

Impact of EDMs in particle physics

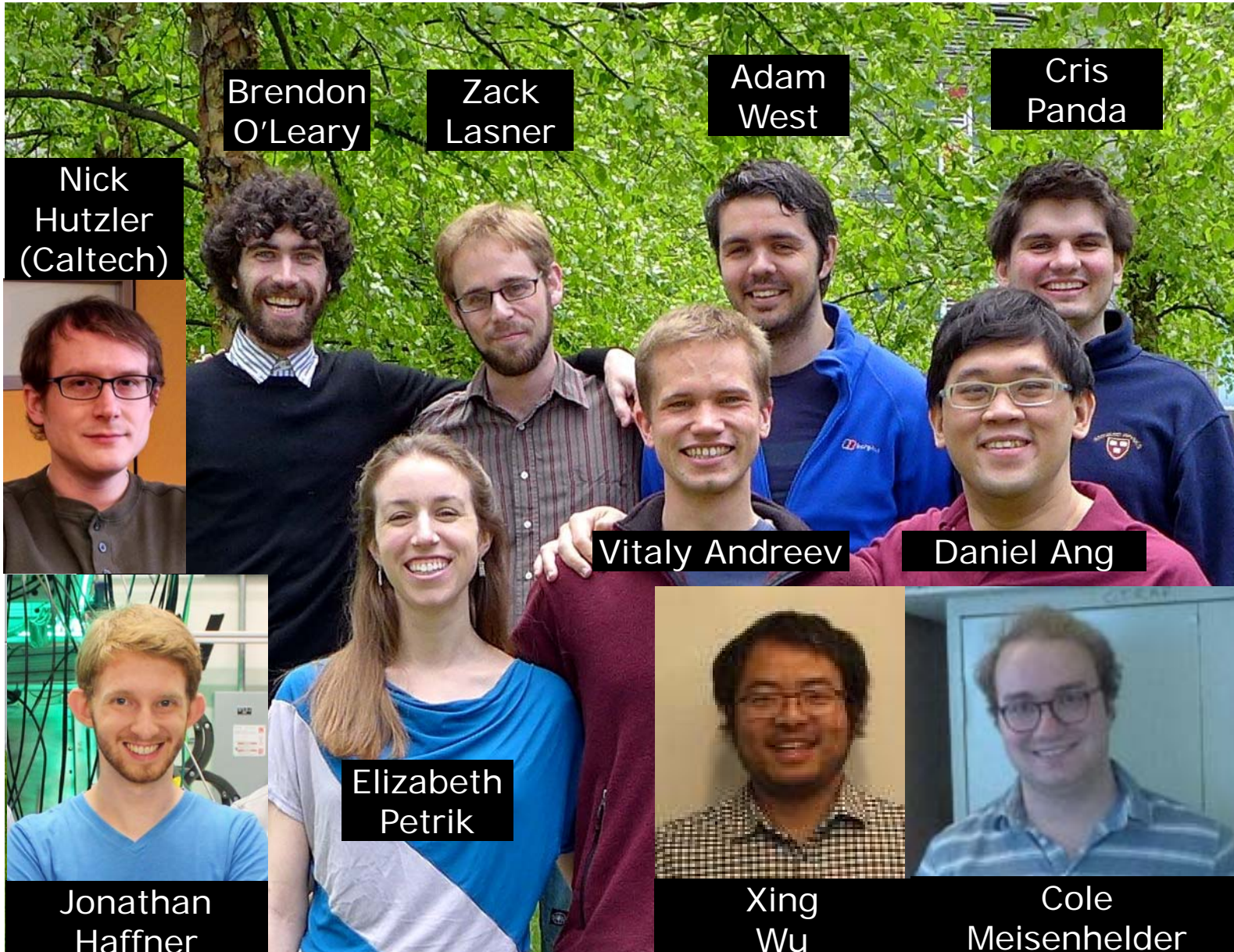
(A very generic view)

Breadth of new physics versus depth of mass reach

← **Genericity**



The ACME II team



Gerald Gabrielse

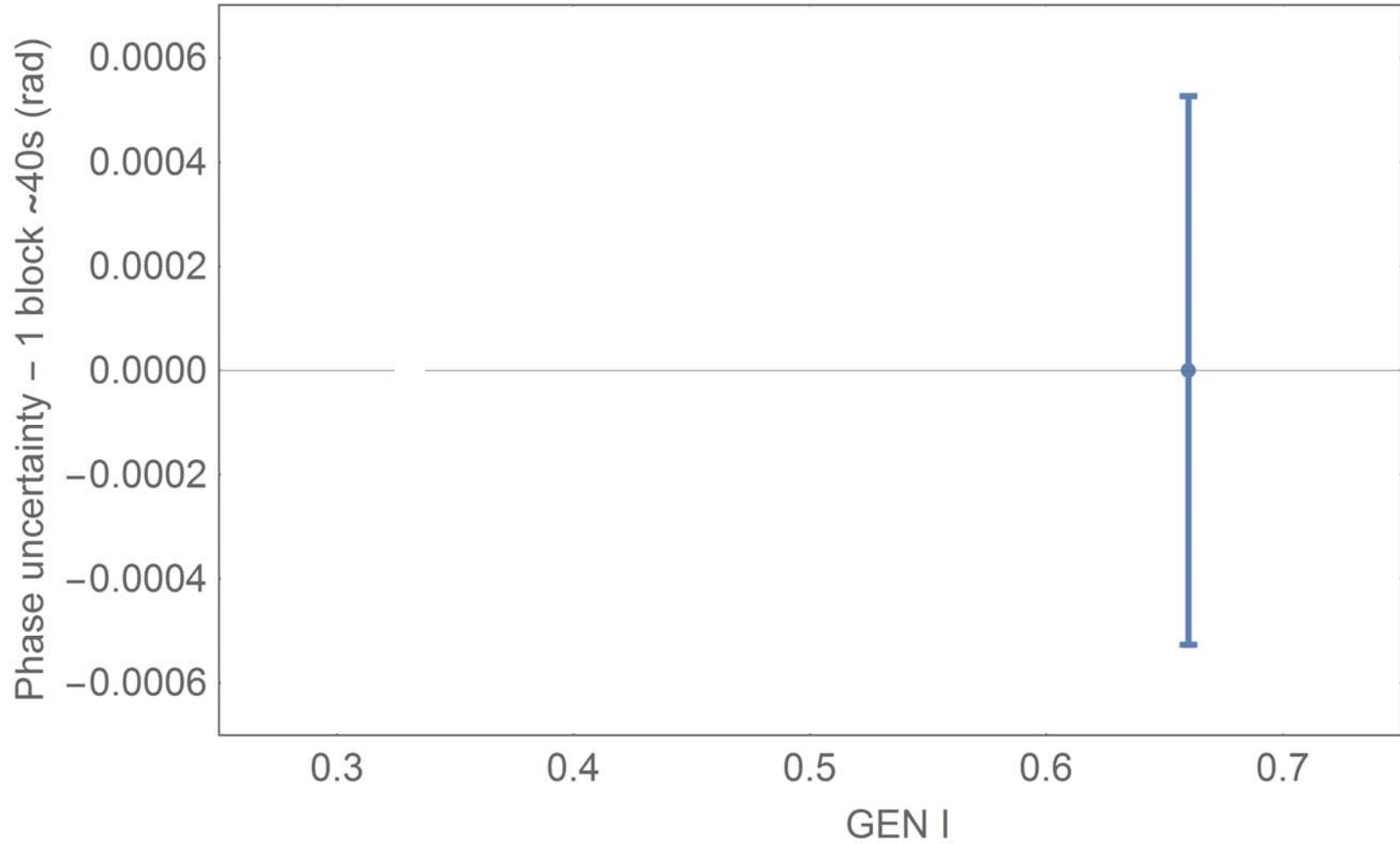


John Doyle

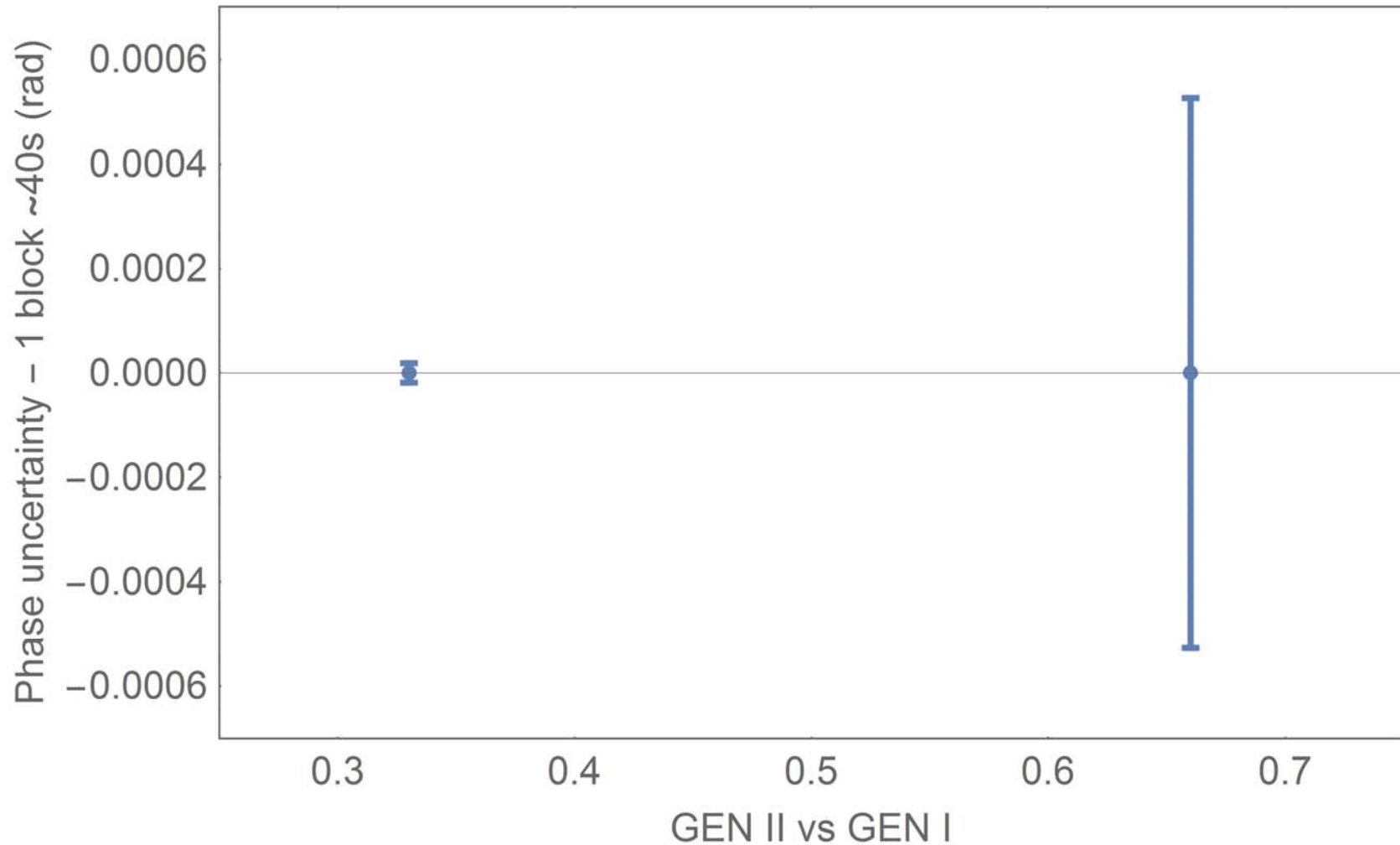


DD

ACME II statistical sensitivity



ACME II statistical sensitivity



~10x improved EDM statistical sensitivity/unit time

Systematic error studies ~complete, final data taking underway

Near-future discovery potential with the electron EDM

