Neutrino Physics with Cold Atoms

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Tuesday February 3rd, 2009 Collaborators include Dr. Mark Raizen, Dr. Joshua Klein, and Julia Majors

Neutrino Physics with Cold Atoms

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What can we learn about neutrinos?

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Tritium β-decay:

Has long been a probe of neutrino properties **Atomic Physics:**

Cold SH Source

Recently developed general methods of slowing & cooling

Neutrino Mass

Maki-Nakagawa-Sakata matrix

$$\begin{pmatrix} \mathbf{v}_{e} \\ \mathbf{v}_{\mu} \\ \mathbf{v}_{\tau} \end{pmatrix} = \begin{pmatrix} \mathbf{U}_{e1} & \mathbf{U}_{e2} & \mathbf{U}_{e3} \\ \mathbf{U}_{\mu 1} & \mathbf{U}_{\mu 2} & \mathbf{U}_{\mu 3} \\ \mathbf{U}_{\tau 1} & \mathbf{U}_{\tau 2} & \mathbf{U}_{\tau 3} \end{pmatrix} \times \begin{pmatrix} \mathbf{v}_{1} \\ \mathbf{v}_{2} \\ \mathbf{v}_{3} \end{pmatrix}$$

 $P(v_{\mu} \rightarrow v_{e}) = \sin^{2}(2\theta)\sin^{2}(1.27\Delta m^{2}L/E)$

The probability of flavor change depends on the mass differences between states, not on absolute masses

Oscillations only provide a lower limit on the v-mass scale!

 $m_3 \ge \text{sqrt}(|\Delta m_{\text{atm}}^2|) \sim (0.04 - 0.07 \text{ eV})$

 $\Delta m_{21}^2 = \Delta m_{sol}^2 = 8.0 \text{ x } 10^{-5} \text{ eV}^2$ (KamLAND)

 $\Delta m_{31}^2 \approx \Delta m_{32}^2 = \Delta m_{atm}^2 = 2.4 \text{ x } 10^{-3} \text{ eV}^2$ (Super-K)

Neutrino Mass

- Are neutrino masses hierarchical or degenerate?
- Are neutrinos Dirac or Majorana particles?
- Why are neutrino masses so relatively small?



hierarchical scenarios

degenerate scenarios



Neutrino Mass

Cosmology

- Neutrinos = hot dark matter in the early universe
- Energy density parameter Ω of the universe: Experimental limits: $\Omega_v = 0.003 - 0.25$
- Fits for m_v depend sensitively on other cosmological parameters

Evolution of large scale structures



WMAP data has been fit with cosmological models that estimate $\sum m_v \le 0.6 \text{ eV}$



Neutrinoless Double Beta Decay Experiments

• Neutrino emitted at one beta decay vertex has to be absorbed by the second decay vertex as an antineutrino





- CUORE plans to reach a limit of 20-50 meV on neutrino mass
- Only possible if neutrinos are massive and Majorana, meaning they are their own antiparticles

SuperNEMO GERDA EXO CUORE MOON II COBRA Majorana CANDLES XMASS CARVEL SNO+ many more ...

Tritium beta decay

 $^{3}\text{H} \rightarrow {}^{3}\text{He}^{+} + e^{-} + \bar{\nu}_{e}$

Half life: $t_{1/2} = 12.3$ years Endpoint energy: $E_0=18.6$ keV



Tritium beta decay

Electron energy spectrum of tritium β decay:

$$N(E) = \frac{dN}{dE} = K \times F(E,Z) \times p_e \times E_e \times p_v \times E_v$$
$$= K \times F(E,Z) \times p \times W \times \sqrt{(E_0 - E)^2 - (m_v^2)^2} \times (E_0 - E)$$

- p = electron momentumW = electron total energy
- E = electron kinetic energy
- E_0 = endpoint energy = 18.6 keV
- F(Z,E) = Fermi function, accounting for Coulomb interaction of the outgoing electron in the final state

• K =
$$G_F^2 (m_e^5 / 2\pi^3) \cos^2\theta_C |M|^2$$



Tritium beta decay

What about neutrino mixing?

N(E) =
$$\frac{dN}{dE}$$
 = K × F(E,Z) × p × E × $\sqrt{(E_0 - E)^2 - m_v^2}$ × (E₀ - E)

$$\mathbf{U}_{ei} |^{2} = |\langle \mathbf{v}_{e} | \mathbf{v}_{i} \rangle|^{2}$$

 $m_v^2 =$ "mass" of the electron (anti-)neutrino = $\Sigma | U_{ei} |^2 m_i^2$

The measured neutrino mass from tritium beta decay would **fix the absolute neutrino mass scale** in a degenerate model

Double beta decay experiments actually measure: $m_v = \left| \Sigma |U_{ek}|^2 e^{i\alpha_e k} m_k \right|$ Majorana CP-phases are unknown \Rightarrow cancellations could occur

ITEP	mv	
T ₂ in complex molecule magn. spectrometer (Tret'yakov)	17-40 eV	experimental results
Los Alamos gaseous T ₂ - source magn. spectrometer (Tret'yakov)	< 9.3 eV	
Tokio T - source magn. spectrometer (Tret'yakov)	< 13.1 eV	
Livermore gaseous T ₂ - source magn. spectrometer (Tret'yakov)	< 7.0 eV	-100 -150 - I I I Los Alamos Mainz -150 - Tokio
Zürich T ₂ - source impl. on carrier magn. spectrometer (Tret'yakov)	< 11.7 eV	-200 - Troitsk (step) A Zürich
Troitsk (1994-today) gaseous T ₂ - source electrostat, spectrometer	< 2.5 eV	-250 - electrostatic spectrometers -300 spectrometers
Mainz (1994-today) frozen T ₂ - source electrostat. spectrometer	< 2.2 eV	-350 1986 1988 1990 1992 1994 1996 1998 2000 year

Troitsk & Mainz breakthrough technology: MAC-E-Filter

guiding by magnetic fields (magnetic adiabatic collimation) $\Delta \Omega \sim 2 \pi$

electric (retarding-) field : analysis of electron energies (electrostatic filter) integral transmission : E > U₀

$$\vec{F} = (\vec{\mu} \cdot \vec{\nabla}) \vec{B} + q \vec{E}$$

 $\mu = E_{\perp} / B = const$

adiabatic motion



adiabatic transformation $E_{\perp} \rightarrow E_{\parallel}$

Troitsk and Mainz

• Obtained m_v by fitting the beta spectrum

• Parameters were m_v, endpoint energy, background, and normalization





Troitsk

Mainz

 $m_v^2 = -1.0 \pm 3.0 \pm 2.5 \text{ eV}^2$

 $m_v \le 2.5 \text{ eV} (95\% \text{ CL})$

Source = Windowless gaseous T^2

 $m_v^2 = -1.6 \pm 2.5 \pm 2.1 \text{ eV}^2$

 $m_v \le 2.2 \text{ eV} (95\% \text{ CL})$

Source = Quench condensed T^2 film on graphite

Troitsk and Mainz:

$$m_{v} < 2.2 \text{ eV}$$

Limiting Factors:

- Statistics
- Scattering in source
- Backgrounds
- Energy resolution
- Electronic final state effects
- Tritium source uncertainties



KATRIN

- Scaled-up version of Troitsk experiment
- Plans to reach a neutrino mass sensitivity of 0.2 eV after 5-6 years of data taking



- Windowless Gaseous Tritium Source is a 10m long cylinder
- Main spectrometer is 23m long and 10m in diameter

KATRIN

- Factor of 4 improvement in energy resolution over Troitsk and Mainz
- Increased T₂ source strength (factor 80)
- Low background of 10⁻² counts/s or less is required
- Reduced inelastic scattering events to 2% of signal rate by looking only at the last 25 eV below the endpoint
- Pre-spectrometer rejects all electrons except those close to the endpoint, reducing the count rate to $\sim 1000/s$



Sensitivity (90% CL) $m_v < 0.2 \text{ eV}$

Discovery (95% CL) $m_v < 0.35 \text{ eV}$

KATRIN

Monte Carlo spectra:

- Run time = 3 years
- $\Delta E = 1 \text{ eV}$
- WGTS column density = 5 x 10¹⁷/cm²
- Final state effects included

• Analysis window = 5 eV below endpoint



KATRIN:	MARE:				
Karlsruhe Tritium Neutrino Experiment	Microcalorimeter Arrays for a Rhenium Experiment				
• External β-source (³ H)	• β -source = detector (¹⁸⁷ Re)				
• 3 H endpoint = 18.6 keV	• 187 Re endpoint = 2.6 keV				
• 3 H half-life = 12.3 years	• 187 Re half-life = 5×10^{10} years				
• Energy: electrostatic spectrometer	• Energy: single crystal bolometer				
 Measures kinetic energy of β 	Measure entire decay energy				
• Narrow interval close to E _o	Measure entire spectrum				
 Integrated β-energy spectrum 	 Differential β-energy spectrum 				
• Integral design, size limits	• Modular size, expandable				
• $\Delta E_{\text{expected}} = 0.93 \text{ eV}$	• $\Delta E_{\text{expected}} \sim 5 \text{ eV} \text{ (FWHM)}$				

Is there another approach to directly measuring m_v?

Slowing and trapping cold atomic tritium would create a new kind of source for tritium β-decay.

- Supersonic nozzle \rightarrow beam of atoms moving at ~400m/s
- Temperature of beam is very cold (~50mK in co-moving frame)
- Tritium can be entrained into the beam and then slowed for trapping



E. Narevicius, A. Libson, C. Parthey, I. Chavez, J. Narevicius, U. Even, and M. G. Raizen, Phys. Rev. Lett. 100, 093003 (2008)

Use pulsed magnetic fields to decelerate tritium atoms



• Zeeman effect: $\Delta E = - \mu \cdot B$

• Low-field seekers are repelled in high field regions and lose kinetic energy



Use pulsed magnetic fields to decelerate tritium atoms



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Can we further cool tritium once we've trapped it?

Laser cooling: Highly effective but limited to a small group of atoms

- 1997 Nobel Prize: Chu, Cohen-Tannoudji, Phillips
- Repeated scattering of photons reduces atomic momentum
- Requires a cycling transition
- Hydrogen cannot be laser cooled

	_																	
hydrogen																		helium
1.11																		1
H																		не
1.0079	borsillium	i i										6	horon	earbon	nitrogen	opidan	fluorino	4.0026
3	4												5	6	7	8	9	10
1.1	Ro												B	C	N	0	E	No
6.041	0.0122												10.911	12.011	14.007	15.000	19.000	20.190
sodium	magnesium												aluminium	silicon	phosphorus	sulfur	chlorine	argon
11	12												13	14	15	16	17	18
Na	Ma												A	Si	Ρ	S	CI	Ar
22.990	24.305												26.982	28.086	30.974	32.065	35.453	39.948
19	calcium 20		scandium 21	1ttanium 22	vanadium 23	chromium 24	manganese 25	26	27	nickel 28	copper 29	2inc 30	gallium 31	germanium 32	arsenic 33	selenium 34	35	Krypton 36
1Z	Co		60	Ti	V	Cr	Mp	Eo	Co	NI	Cu	Zn	Ga	Go	Ac	80	Dr	Kr.
N	Ca		30	11007	V	CI		ге	00		Cu		Ga	Ge	AS	Se	DI	
rubidium	strontium		44.956 yttrium	47.867 zirconium	niobium	51.996 molybdenum	technetium	ruthenium	rhodium	58,693 palladium	silver	cadmium	69.723 indium	72.61 tin	antimony	78.96 tellurium	iodine	xenon
37	38		39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54
Rb	Sr		Y	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Те		Xe
85.468	87.62		88.906	91.224	92.906	95.94	[98]	101.07	102.91	106.42	107.87	112.41	114.82	118.71	121.76	127.60	126.90	131.29
55	56	57-70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86
Ce	Ra	×	L u	Hf	Ta	1/1/	Ro	0c	Ir	Dt	Διι	Ha	TL	Ph	Ri	Po	Δt	Rn
122.01	127.22	A	174.97	179.40	190.05	102.04	196.21	100.22	102.22	105.08	196.97	200.59	204.28	207.2	209.99	12001	1210	12221
francium	radium	100000000000000000000000000000000000000	lawrencium	rutherfordium	dubnium	seaborgium	bohrium	hassium	meitnerium	ununnilium	unununium	ununbium	204.50	ununquadium	200,50	200	210	222
87	88	89-102	103	104	105	106	107	108	109	110	111	112		114				
Fr	Ra	* *	Lr	Rf	Db	Sq	Bh	Hs	Mt	Uun	Uuu	Uub		Uuq				
[223]	[226]		[262]	[261]	[262]	[266]	[264]	[269]	[268]	[271]	[272]	[277]		[289]				
			Ionthonum	oorium	Inraseodemium	noodymium	promothium	comorium	ouropium	andolinium	torbium	duenroeium	bolmium	orbium	thulium	vitorbium	i.	
*Lant	hanida	corios	57	58	59	60	61	62	63	64	65	66	67	68	69	70		
Lanti	namue	301103	la	Ce	Pr	Nd	Pm	Sm	Eu	Gd	Tb	Dv	Ho	Fr	Tm	Yh		
			138,91	140.12	140,91	144.24	[145]	150.36	151.96	157.25	158.93	162.50	164,93	167,26	168,93	173.04		
** ^ - 4	inida -		actinium	thorium	protactinium	uranium	neptunium	plutonium	americium	curium	berkelium	californium	einsteinium	fermium	mendelevium	nobelium		
^ ^ ACT	iniae se	erles	89	90	91	92	93	94	95	96	9/	98	99	100	101	102		
			AC	In	Ра	U	пр	гu	AM	Cm	BK	UT	ES	r m	IVIC	NO		
			[227]	232.04	231.04	238.03	[237]	[244]	[243]	[247]	[247]	[251]	[252]	[257]	[258]	[259]		

Is there a more general cooling method?



2nd law is saved by information carrying entropy

Szilard (1929)

- Demon makes a measurement
- Information entropy

Demon's jobs:

- Measure **r**, **p**
- Operate gate

Single-photon cooling realizes Maxwell's demon:

"Demon" discriminates coldest atoms and releases this info in a single scattered photon

Thought-experiment by Maxwell (1867)

Entropy reduced without expenditure of work



Goal: Transfer atoms from a magnetic to an optical trap via emission of a single photon

- Slowly translate 1-way barrier so that you catch atoms at their classical turning points
- A spontaneous Raman emission could be such a 1-way barrier
- This cooling technique has been demonstrated on ⁸⁷Rb
- (T. Bannerman, G. N. Price, K. Viering, and M. G. Raizen, arXiv:08102239v1)



Allows creation of a tritium source with ~µK temperature



- "Demon" = gravito-optical trap + resonant pump beam
- Approach classical turning points slowly from the left
- If final state has weaker or opposite magnetic coupling, atom is trapped in optical trap

 $U = \mu_B g_F m_F |B| + mgz$





A source of ~10¹¹ trapped atoms allows the ion to escape as well as the β .

$$^{3}\text{H} \rightarrow {}^{3}\text{He}^{+} + e^{-} + \bar{v}_{e}$$

Half life: $t_{1/2} = 12.3$ years Endpoint energy: $E_0 = 18.6$ keV



Direct reconstruction of the neutrino mass!



$$\mathbf{m}_{v}^{2} = (\mathbf{W} - \mathbf{E}_{ion} - \mathbf{E}_{\beta})^{2} - (\mathbf{p}_{x_{ion}} + \mathbf{p}_{x_{\beta}})^{2} - (\mathbf{p}_{y_{ion}} + \mathbf{p}_{y_{\beta}})^{2} - (\mathbf{p}_{z_{ion}} + \mathbf{p}_{z_{\beta}})^{2}$$

- Thin source allows ion detection!
- Don't have to rely only on beta spectrum
- Coincidence measurement \Rightarrow low backgrounds
- Atomic tritium \Rightarrow well-known final state corrections

Direct reconstruction of the neutrino mass!



Use Rydberg atoms to measure β momentum non-invasively:



Rydberg atoms have valence electrons with high principle quantum number:

$$r = \frac{n^2 \bar{h}^2}{k Z e^2 m} \quad \Longrightarrow \quad$$

High n implies large orbital radius and easily perturbed or ionized electrons

Use Rydberg atoms to measure β momentum non-invasively:



What about the opening angle uncertainty?

$$\begin{split} \tilde{\mathbf{p}}_{v} \cdot \tilde{\mathbf{p}}_{v} &= \mathbf{m}_{v}^{2} \qquad \tilde{\mathbf{p}}_{v} + \tilde{\mathbf{p}}_{ion} + \tilde{\mathbf{p}}_{\beta} = \tilde{\mathbf{p}}_{3H} & & & \\ \mathbf{m}_{v}^{2} &= \tilde{\mathbf{p}}_{v} \cdot \tilde{\mathbf{p}}_{v} = (\tilde{\mathbf{p}}_{3H} - \tilde{\mathbf{p}}_{ion} - \tilde{\mathbf{p}}_{\beta}) \cdot (\tilde{\mathbf{p}}_{3H} - \tilde{\mathbf{p}}_{ion} - \tilde{\mathbf{p}}_{\beta}) & {}^{3He^{+}} \\ \mathbf{m}_{v}^{2} &= \mathbf{W}^{2} - 2\mathbf{W}\mathbf{E}_{ion} - 2\mathbf{W}\mathbf{E}_{\beta} + \mathbf{m}_{ion}^{2} + \mathbf{m}_{\beta}^{2} + 2|\mathbf{p}_{ion}||\mathbf{p}_{\beta}|\cos\theta \\ & \delta\theta \frac{\partial \mathbf{m}_{v}}{\partial\theta}^{2} = -2|\mathbf{p}_{ion}||\mathbf{p}_{\beta}|\sin\theta & \sim \delta\theta\sin(\theta)10^{10}\,(eV/c)^{2} \end{split}$$

 $\theta \approx \pi$

e

How do we avert disaster?

- Opening angle is almost π , which makes sin θ small
- The uncertainty of the mean goes like $1/N^{1/2}$
- $\delta\theta\sin(\theta)10^{10} (eV/c)^2 = 10^{-5} (\sin(\pi 10^{-4}))10^{10} (eV/c)^2 = 10 (eV/c)^2$

ROOT simulation: based on kinematics (no particle tracking)

What's in the simulation?

- atomic tritium source is 100µm diameter sphere
- Tritium source atoms at 1µK with a Gaussian momentum smear of width mkT
- Electron TOF Gaussian smear of 20ps
- Electron energy resolution of 5 meV
- Final state effects: ground state 70%, 1st excited state 30%
- 1 year assumed runtime

- Electron momentum resolution of 15meV/c to 2.7 eV/c
- Geometrical acceptance for the β limited by 10 beams of Rydberg atoms (width 1cm) placed 5m from source
- MCP: 2 micron binning, 44% geometrical acceptance, 100cm wide and 20cm tall, placed 5m from source
- Ion TOF Gaussian smear of 20ps
- Gravity correction for the ion ~0.5 microns

PHOIBOS hemispherical analyzer 225 HV

 For electron energies up to 15 keV
 Different modes of operation (UPS, XPS and HXPS)
 Ultra high energy resolution in UPS (<1 meV), XPS (<7 meV) and HXPS (<15 meV)
 Angular Mapping (Δθ < 0.1°)
 CCD, DLD and DLD/SPIN detection available

- 15 keV energy
- Small geometrical acceptance
- Potential calibration source: ^{83m}Kr conversion electron with energy of 17.8 keV and width of 2.7 eV

SPECS

Burle 2-micron MCP detector



We need:

- 2-10 µm spacing
- ~20ps timing

• Large area: ~ 1m wide x 20cm tall

- Detects position and time-of-flight (TOF)
- 2 micron holes spaced 3 microns center-to-center
- 350 ps pulse width resolution

Background test:

- Randomize MCP hits
- Randomize ion TOF
- Leave beta unchanged

 10^{-5} background rejection, not including β -coincidence





Magnitude of p_v is increased 2-3 times, while E_v changes only slightly $\rightarrow m_v$ always reconstructs extremely negative for background events

- Fit utilizes data up to 500eV away from the endpoint energy
- Minuit log-likelihood fit using 2D probability density functions (pdf)
- Find m_v by interpolating between pdfs of different neutrino masses



Assumed m _v (eV)	Fit m _v	(+) error	(-) error
0.2	0.190	0.166	0.139
0.4	0.407	0.211	0.160
1.0	0.778	0.268	0.240



Statistics gained by moving far from the endpoint improve precision on m_v even though the spread in reconstructed mass gets broader.

What are the strengths of this technique?

- Extremely thin source \rightarrow low scattering
- Atomic tritium -> simpler final state effects Utilizing data 500eV from endpoint
- β coincidence \rightarrow low backgrounds



• Direct m, reconstruction & β -spectrum

- Valid for Dirac and Majorana neutrinos

Experimental challenges:

- Trapping 2x10¹³ tritium atoms
- Large MCP with binning 2-10µm
- 5-50meV energy resolution
- Non-invasive momentum measurement for β

Boundstate β-Decay: 2-Body

$$^{3}\text{H} \rightarrow ^{3}\text{He} + v_{e}$$

$$v_{Recoil} = \frac{[(M_{3H} - M_{3He})^2 - (m_v c^2)^2]^{1/2}}{M_{3He}c}$$

- Measure ³He recoil velocity
- 0.69% of all ³H decays are boundstate
- 3% of boundstate He³ atoms are in an excited state and emit a 706.52nm photon



Boundstate β-Decay: 2-Body

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Boundstate β-Decay: 2-Body

Boundstate beta decay does not currently offer a competitive limit on m_v.



Ordinary Mossbauer effect: photons emitted recoillessly by one nucleus can be resonantly absorbed by another nucleus of the same type



Nuclei must be bound in a lattice for significant recoilless emission or absorption.

v's emitted recoillessly from boundstate decay of ³H can be resonantly absorbed by ³He



Boundstate tritium beta decay:

$$^{3}\text{H} \rightarrow ^{3}\text{He} + \bar{\nu}_{e}$$

Reverse tritium beta decay: $\bar{\nu}_e + {}^{3}\text{He} \rightarrow {}^{3}\text{H}$

Debye temperature = temperature of a crystal's highest normal mode of vibration $f_{recoilless} = \exp\{(-E^2/(2Mc^2)*(3/2k_B\theta_D))\}$ where θ_D is the Debye temperature

We can get a very high Debye temperature by going to high pressures



- High pressures raise the Debye temperature, which increases $f_{recoilless}$
- Volume not likely to exceed 0.004cm³



Tuning the pressure allows us to align emission & absorption peaks!

$$\sigma_{\text{resonant}} = 4.18*10^{-41}*g_0^2*\rho(\text{E}^{\text{res}}_{v_e})/ft_{1/2} \approx 10^{-32}\text{cm}^2$$
(assuming linewidth ~10⁻¹²eV)

• Linewidth dominated by inhomogeneous broadening (impurities, lattice defects, ect.)

• Narrow linewidth implies we must be able to tune energy shifts to observe resonance

• Very cold temperatures reduce Doppler shifts

• Isomer shift (from changes in atomic radius) can be canceled by zero-point energy shift:

 $\Delta E/E = (9k_{\rm B}/16Mc^2)^*(\theta_{\rm emitter} - \theta_{\rm absorber})$



We estimate a Debye temperature of ~700K Simulation results: ~31755 events per week



But how do you detect the tritium in the helium-3 absorber?

• Magnetic slowing enables trace element detection so we can actually detect the ³H in the ³He absorber! (\sim 1/1000 detection efficiency)

Physics motivation:

• θ_{13} measurement from rates taken at distances 1cm-10m



$$P(v_{\alpha} \rightarrow v_{\beta}) = \sin^2(2\theta)\sin^2(1.27\Delta m^2 L/E)$$

A large L is unnecessary if E=18.6keV

Conclusions

Slowing and trapping cold ³H atoms \rightarrow exciting potential v experiments:

- First atomic source ever utilized in tritium β -decay
- Three-body β -decay: Fundamentally new way of measuring v mass
- Boundstate β -decay: Unique (if uncompetitive) constraint on the neutrino mass
- Neutrino Mössbauer effect: Trace element detection through magnetic slowing may enable v research at tabletop scales



M. Jerkins, J. R. Klein, J. H. Majors, and M. G. Raizen, arXiv:0901.3111

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