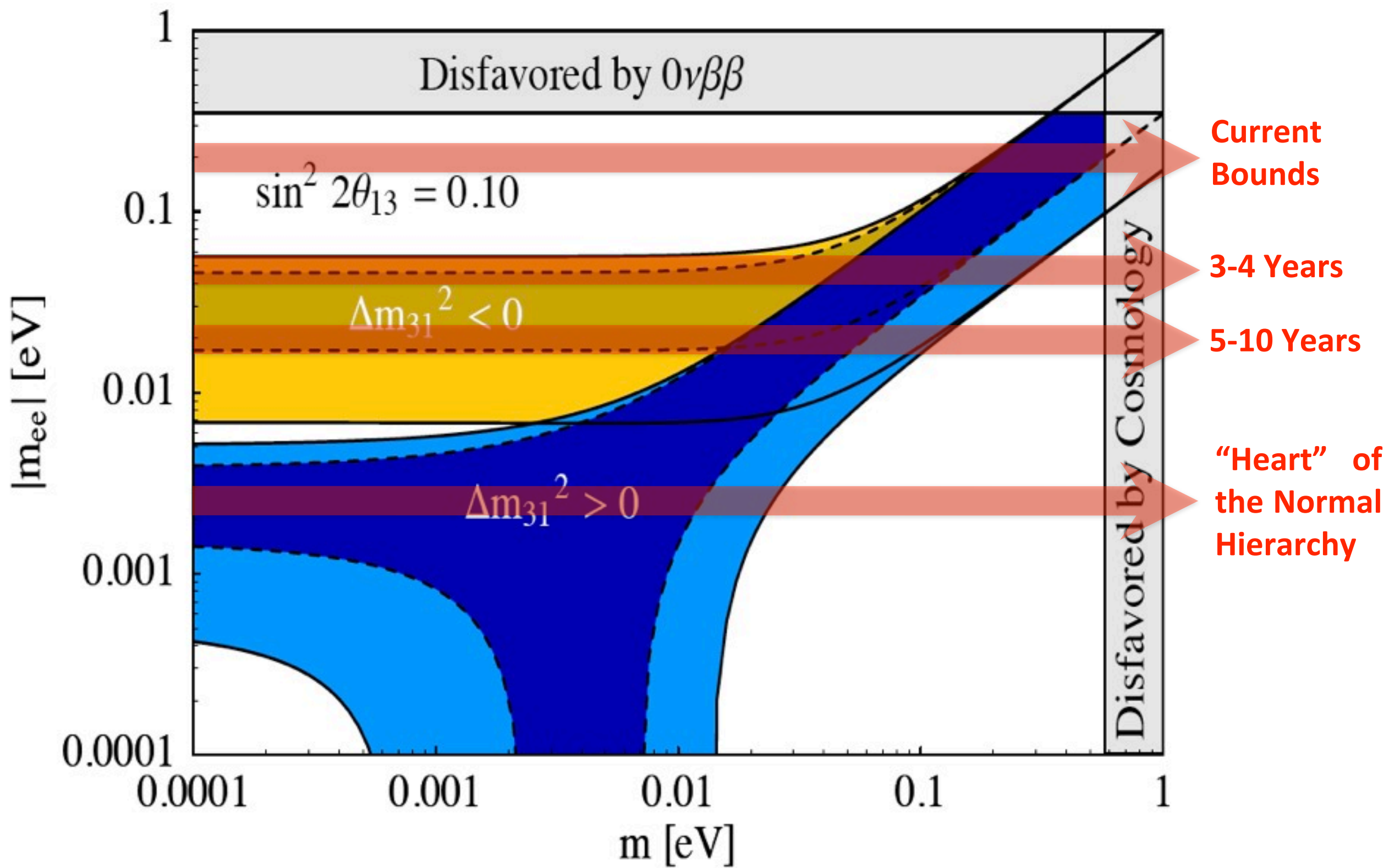


Speculations on Accessing
the Non-Degenerate Normal
Neutrino Mass Hierarchy
with $0\nu\beta\beta$

Steve Biller, Oxford University



Note on Sensitivity Scaling

$$T_{1/2} \sim m^2 \quad m_{\text{limit}} \sim \sqrt{\frac{1}{T_{1/2}}} \sim \sqrt{\sqrt{B/S}} \quad \text{Gaussian limit}$$

$$S \sim M_{\text{isotope}} T_{\text{live}} \quad B \sim T_{\text{live}}$$

In the non-Gaussian limit ($B \ll 1$): $m_{\text{limit}} \sim T^{-1/2}$

but otherwise:

$$m_{\text{limit}} \sim T^{-1/4}$$

If B is ~independent of M_{isotope} :

$$m_{\text{limit}} \sim M$$

but otherwise:

$$m_{\text{limit}} \sim M^{-1/4}$$

Killers!

Standard Approach:

On a small scale, all isotopes have roughly the same intrinsic rate predictions. So, just try different isotopes and various approaches to see how far each technique can be pushed and worry about scaling things up later.

Alternative Approach:

On a large scale, all isotopes are NOT the same owing to backgrounds, cost, logistics of implementation etc. So, define the practical requirements for a normal hierarchy scale first and use that to identify which isotopes and techniques to pursue.

My Rules for the Game:

- Sensitivity should be at least that needed to achieve a 90% CL at 2.5 meV with 5 years of data using average matrix element values.
- Must observe candidate events at a reasonable rate, greater than 1 per year.
- Must be “practical”: logistically feasible and having an estimated cost of less than one billion US dollars.
- Must be based on modest extrapolations from current techniques and costs (minimal dependence on miracles).

Isotope

Isotope	Q (MeV)	percent natural abund.	element cost [5] (\$/kg)	$G^{0\nu}$ ($10^{-14}/\text{yr}$) [6]	$M^{0\nu}$ (avg) [7]	$T_{1/2}^{0\nu}$ for 2.5meV (10^{29}yrs)	tons of isotope for 1 ev/yr	equivalent natural tons	annual world production [5] (tons/yr)	natural elem. cost (\$M)	enriched at \$20/g (\$M)	$0\nu/2\nu$ rate [2][8] (10^{-8})
^{48}Ca	4.27	0.19	0.16	6.06	1.6	2.70	31.1	16380	2.4×10^8	2.6	622	0.016
^{76}Ge	2.04	7.8	1650	0.57	4.8	3.18	58.2	746	118	1221	1164	0.55
^{82}Se	3.00	9.2	174	2.48	4.0	1.05	20.8	225	2000	39	416	0.092
^{96}Zr	3.35	2.8	36	5.02	3.0	0.93	21.4	763	1.4×10^6	27	427	0.025
^{100}Mo	3.04	9.6	35	3.89	4.6	0.51	12.2	127	2.5×10^5	4.4	244	0.014
^{110}Pd	2.00	11.8	23000	1.18	6.0	0.98	26.0	221	207	5078	521	0.16
^{116}Cd	2.81	7.6	2.8	4.08	3.6	0.79	22.1	290	2.2×10^4	0.81	441	0.035
^{124}Sn	2.29	5.6	30	2.21	3.7	1.38	41.2	736	2.5×10^5	22	825	0.072
^{130}Te	2.53	34.5	360	3.47	4.0	0.75	23.6	68	~150	24	471	0.92
^{136}Xe	2.46	8.9	1000	3.56	2.9	1.40	45.7	513	50	513	914	1.51
^{150}Nd	3.37	5.6	42	15.4	2.7	0.37	13.4	240	$\sim 10^4$	11	269	0.024

Detector Technology

Solid State (CUORE, GERDA, Majorana, COBRA...)

Current background target ~ 0.01 counts/kg/keV/yr and ~ 5 keV resolution (CUORE), corresponding to $\sim 10^4$ counts/yr in ROI for 100 tonnes. The means by which to reduce this by 3-4 orders of magnitude is unclear. Cost of technology dominates and more or less scales with isotope mass. The means by which to tackle practical and logistical issues of scaling the technology by ~ 3 orders of magnitude is also unclear.

Thin Film Tracking (SuperNEMO)

Scale of the technology is large and proportional to the surface area of the film, rendering it impractical for 100 tonne quantities of isotope.

TPC (EXO, NEXT)

Cannot directly use Te as TPC medium and unclear how to do this even indirectly.

Liquid Scintillator (KamLAND-Zen, SNO+)

Practical in terms of construction and instrumentation, even for many kilotonne detector volumes (several percent or more loading levels). Cost of technology does not necessarily scale with isotope mass. Potential for low backgrounds from self-shielding, liquid purification and coincidence-tagging techniques. No inherent Te optical absorption lines. Highly flexible configuration.

Scaling Model:

$0\nu\beta\beta$

Consider only
fundamental
backgrounds

$$S = sM_I T \left(\frac{\langle m \rangle}{2.5\text{meV}} \right)^2$$

signal events
per tonne-year
for $m = 2.5\text{ meV}$

Tonnes of
isotope

Exposure
time

$^8\text{B Solar } \nu$

$$B_{\odot} = b_{\odot} T M_D \left(\frac{L}{1000\text{ pe/MeV}} \right)^{-\frac{1}{2}}$$

^8B events in ROI
per tonne-year for
 $L = 1000\text{ pe/MeV}$

Detector
Mass

Detected
scintillator
light level

$$\sigma \propto 1/\sqrt{L}$$

$2\nu\beta\beta$

$$B_{2\nu} = b_{2\nu} M_I T \left(\frac{L}{1000\text{ pe/MeV}} \right)^{-\frac{5.5}{2}}$$

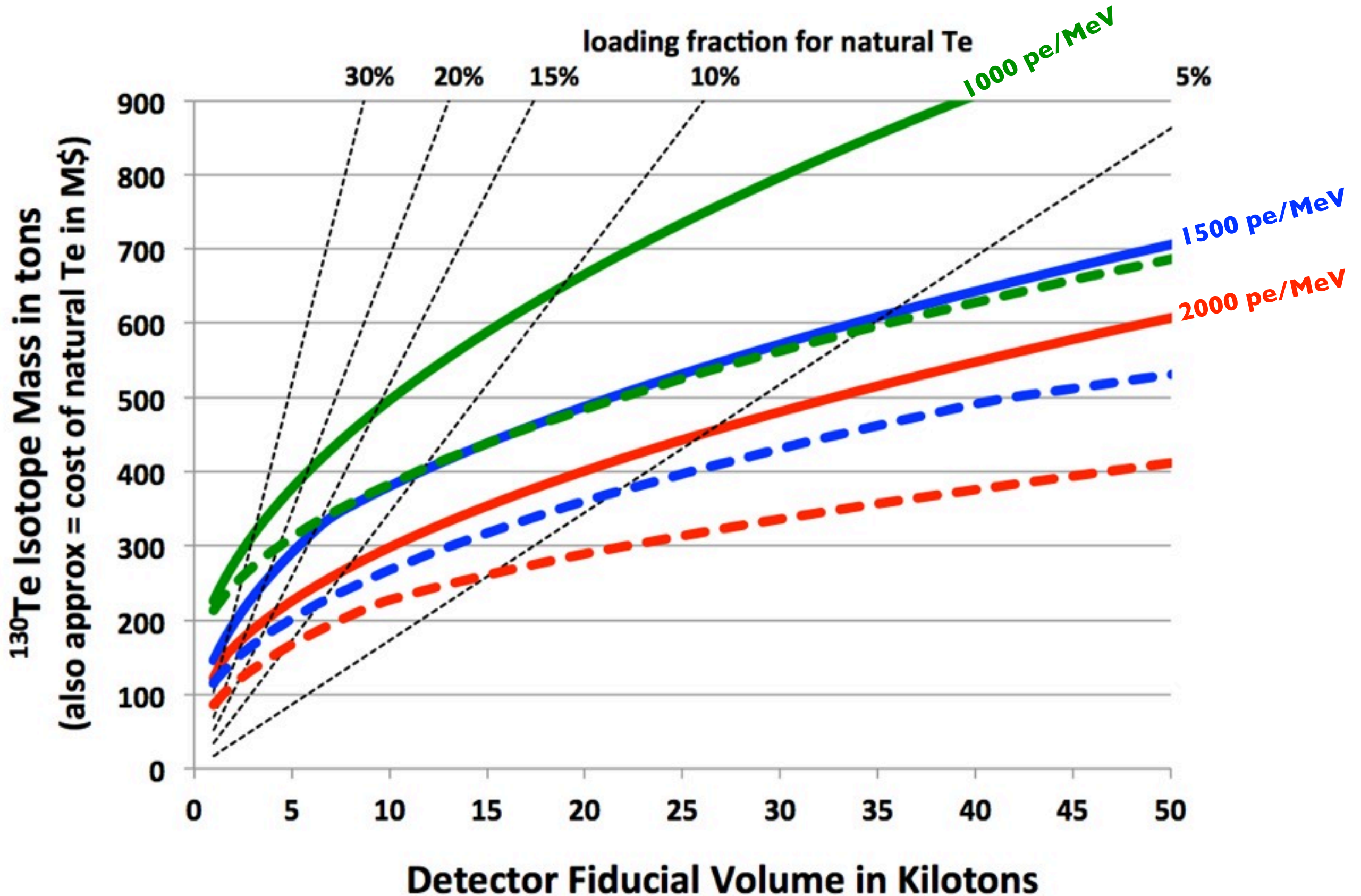
$2\nu\beta\beta$ events in
ROI per tonne-year
for $L = 1000\text{ pe/MeV}$

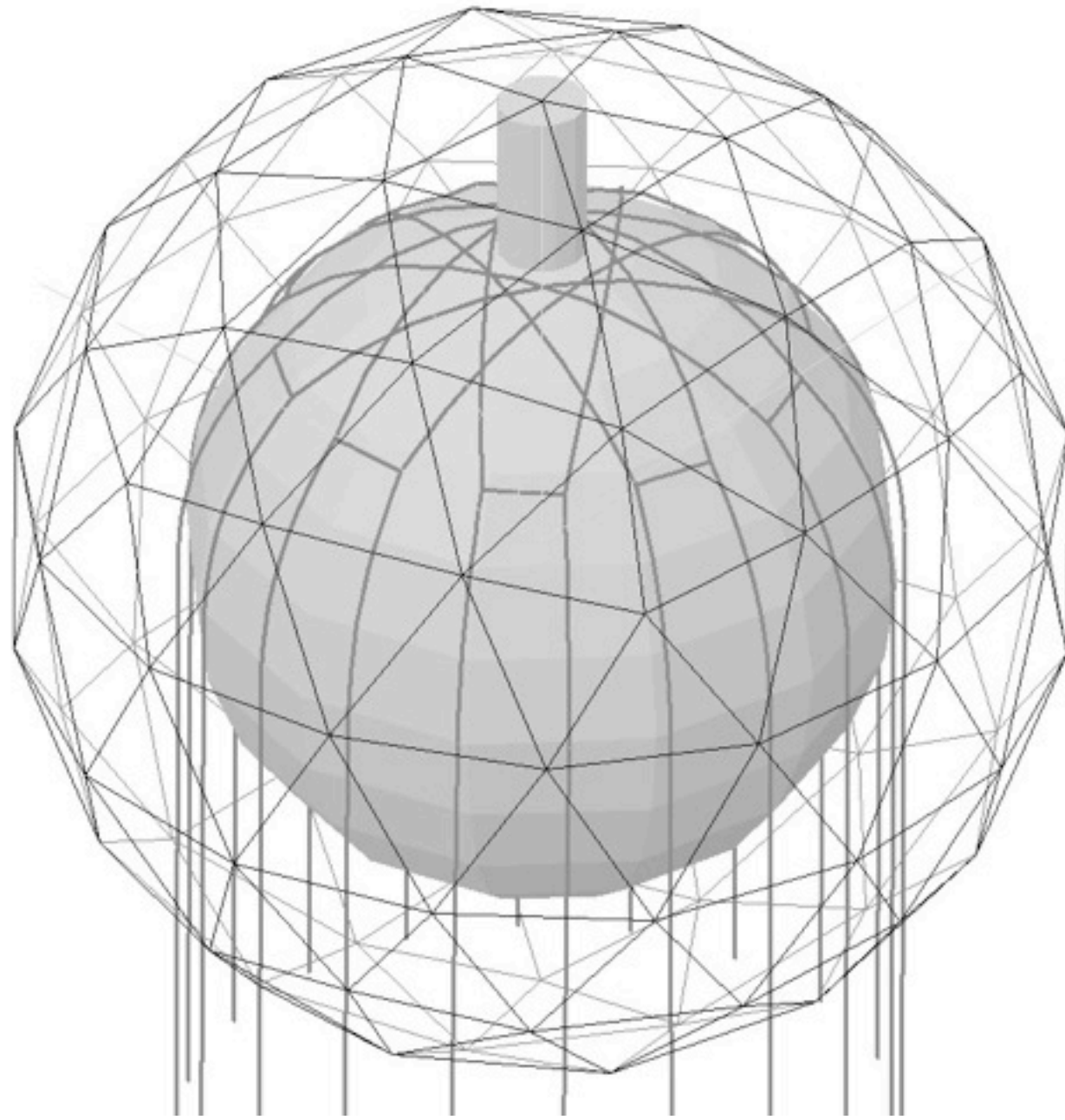
convolved
 $2\nu\beta\beta$ rate
in ROI $\propto \sigma^{5.5}$

$$\text{Significance} \sim S / \sqrt{B_{\odot} + B_{2\nu}}$$

Requirements to achieve 90% CL sensitivity at 2.5meV after 5 live years assuming dominant backgrounds are from solar neutrinos and $2\nu\beta\beta$

(S. Biller, PRD 87, 071301(R), 2013)





SNO+ with Tellurium

Potential for ^{130}Te as an ideal isotope for a LS-loaded $0\nu\beta\beta$ experiment

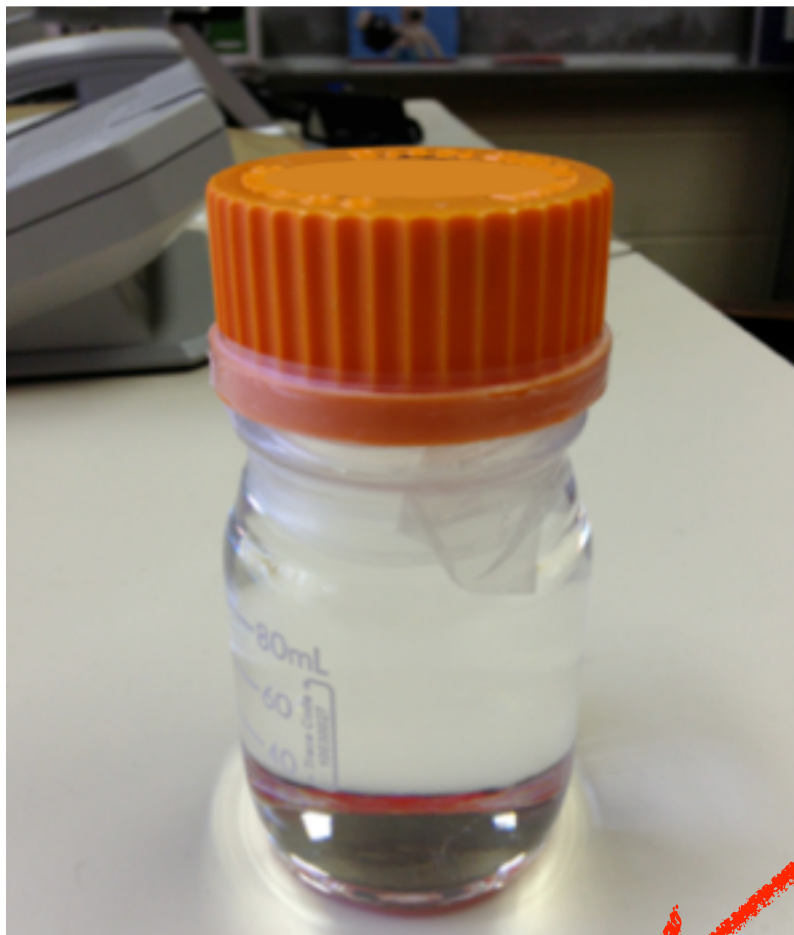
Biller and Chen (Autumn 2011) emphasized potential advantages of Te-loading and initiated development.

- 34% natural abundance;
- Internal U/Th can be actively suppressed (Bi-Po α s);
- External gammas can be attenuated (“fiducialisation”);
- $2\nu\beta\beta$ rate is low (~ 100 times smaller than for ^{150}Nd);
- No inherent optical absorption lines;
- Relatively inexpensive ($<$ a tenth the cost of ^{136}Xe).

Initial loading/purification studies by Yeh *et al.* during 2012.

Subsequently underwent thorough, independent internal review from Aug 2012 - Feb 2013. This resulted in the decision to pursue Te as a first priority, which has since been the focus of a full collaboration development effort.

Conventional Loading Method



New loading technique (BNL):

Dissolve telluric acid in **water** (highly soluble), then combine a small fraction (few percent) of this mixture with LAB using a **surfactant**. Clear and stable (> 1 yr explicitly demonstrated)

(M. Yeh et al., paper in preparation)

Spike tests show metal scavengers reduce U/Th by $\sim 4 \times 10^5$ with double pass (target is a reduction of $\sim 10^4$)

Higher U/Th than pure organic, but still low enough with SNO purification levels (10^{-14} - 10^{-15} g/g)

Isotope ($Q > 2 \text{ MeV}$, $T_{1/2} > 20 \text{ days}$)	$T_{1/2}$ [5] [d]	Q-value [5] [MeV]	R (ϕ from [6][7]) [$\mu\text{Bq/kg}$]	Events/yr in ROI after 1 yr surface exposure
^{44}Sc (daughter of ^{44}Ti)	0.17 (2.16E+4)	3.65	1.19 (0.052)	5.41
^{46}Sc	83.79	2.37	1.97	20.3
^{60}Co (direct and daughter of ^{60}Fe)	1925.27 (5.48E+8)	2.82	0.81 (0.367)	834
^{68}Ga (daughter of ^{68}Ge)	4.70E-2(271)	2.92	3.14 (1.28)	344
^{26}Al	2.62E+8	4.00	0.67	2.20E-4
^{82}Rb (daughter of ^{82}Sr)	8.75E-4(25.35)	4.40	(2.44)	440
^{88}Y (direct and daughter of ^{88}Zr)	106.63 (83.4)	3.62	3.14 (8.11)	3.61E4
^{42}K (daughter of ^{42}Ar)	0.51 (12016.73)	3.53	1.33 (0.24)	10.0
^{56}Co	77.2	4.57	0.13	0.350
^{58}Co	70.9	2.31	1.29	0.252
^{110m}Ag ^a	249.83	3.01	2.34	3.61E3
^{110}Ag (daughter of ^{110m}Ag) ^b	2.85E-4	2.89	(0.03)	48.6
^{106}Rh (daughter of ^{106}Ru)	3.47E-4 (371.8)	3.54	(0.06)	21.8
^{126m}Sb (direct and daughter of ^{126}Sn) ^c	0.01 (8.40E7)	3.69	71.42 (7.87)	8.63
^{126}Sb (direct and daughter of ^{126m}Sb) ^d	12.35 (0.01)	3.67	89.65 (^{126m}Sb)	1.29E4
^{22}Na	950.6	2.84	1.01	1.01E3
^{84}Rb ^e	32.8	2.69	1.29	24.2
^{90}Y (daughter of ^{90}Sr)	2.67 (10519.2)	2.28	2.69 (0.165)	7.90E-3
^{102}Rh (direct and daughter of ^{102m}Rh) ^f	207.3	2.32	11.77 (0.03)	35.9
^{102m}Rh ^g	1366.77	2.46	11.77	69.9
^{124}Sb	60.2	2.90	182.0	1.62E5

ACTIVIA code, cross sections from Silberberg et al. and TENDL-2009 database, flux parameterisations from Armstrong and Gehrels. Variations from using YIELDX code, TENDL-2012 database, and fluxes from Ziegler change estimated rates by up to a factor of two. Consistency also checked against CUORE beam activation study (Wang et al.) and KamLAND induced backgrounds.

(V. Lozza, paper in preparation)

Requires a reduction factor of $> 10^4$ for these isotopes, which is also comparable to the reduction required for U/Th in “raw” Te material (ICP-MS: $2\text{-}3 \times 10^{-11}$ g/g)

Outline of Te Purification Strategy

(paper in preparation)

(Stage 1)

2 Surface passes:

- Dissolve $\text{Te}(\text{OH})_6$ in water
- Recrystallise using nitric acid
- Rinse with ethanol

> 10^4 reduction

Allow up to 5 hr re-exposure to finish & transport UG

(Stage 2)

2 Underground passes:

- Dissolve in warm water (80°C)
- Cool to Recrystallise thermally

> 10^2 reduction

(~50% Te “loss” recovered by recycling to surface system)

Spike Tests (Ongoing)

Element	Reduction Factor	Assay Technique
Stage 1 Te purification, single-pass spike test		
Co	1555± 326	XRF
Sb	>243	XRF
Sn	> 167	XRF
Fe	> 100	XRF
Na	> 346	XRF
Sc	> 165	XRF
Ge	> 333	XRF
Y	> 278	XRF
Zr	> 278	XRF
Ag	> 278	XRF
Pb-212	299± 22	$\beta - \alpha$ counting
Bi-212	348± 81	$\beta - \alpha$ counting
Ra-224	397± 20	$\beta - \alpha$ counting
Th-228	390±19	$\beta - \alpha$ counting
Stage 1 Te purification, double-pass spike test		
Co	3.7×10^5	XRF
Pb-212	> 10^4	$\beta - \alpha$ counting
Bi-212	> 10^4	$\beta - \alpha$ counting
Ra-224	> 10^4	$\beta - \alpha$ counting
Th-228	> 10^4	$\beta - \alpha$ counting
Stage 2 (UG) Te purification, single-pass spike test		
Co	12	XRF
Ag	> 20	XRF
Zr	17	XRF

acid-induced
recrystallisation
+ ethanol wash

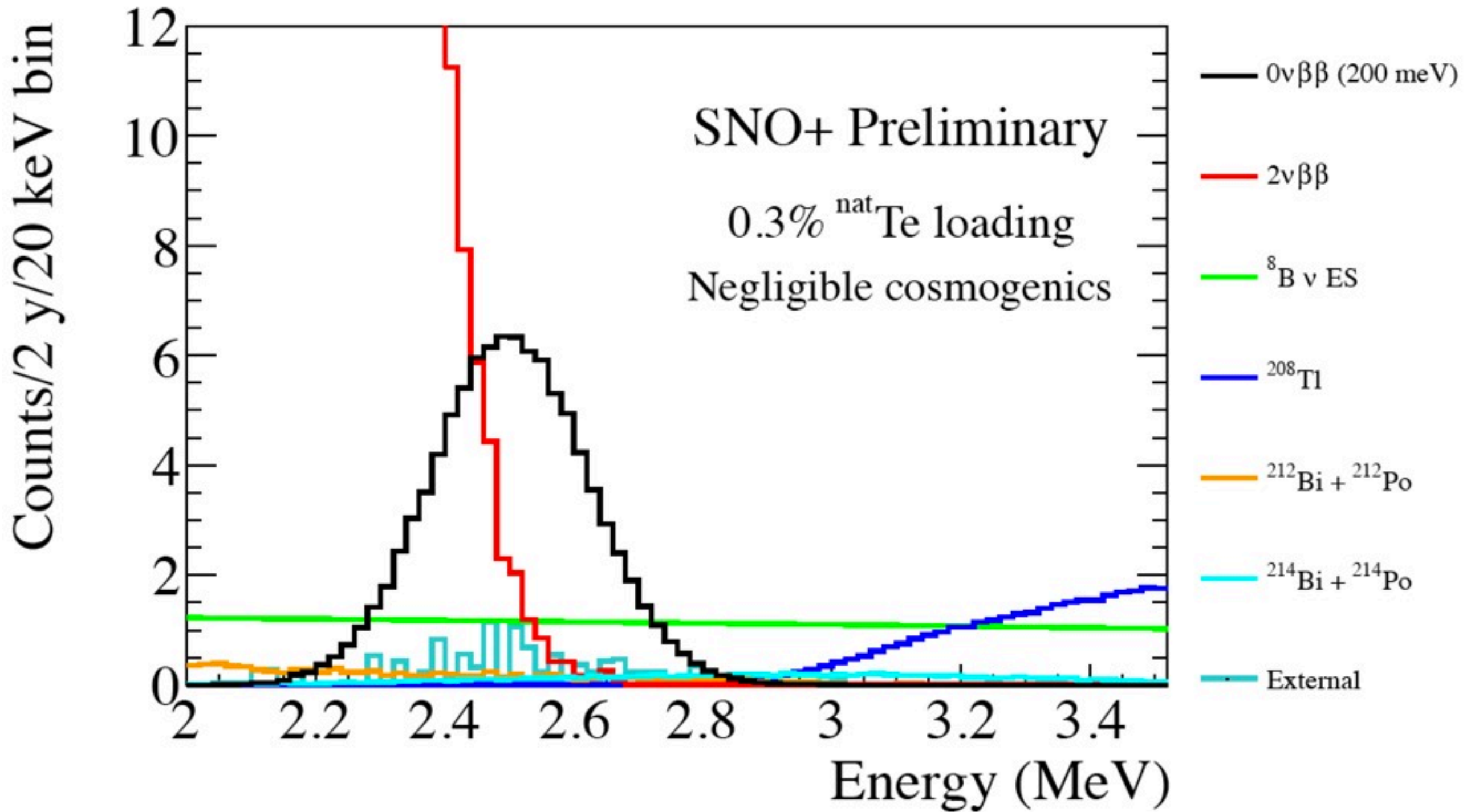
thermal
recrystallisation

Isotope ($Q > 2$ MeV, $T_{1/2} > 20$ days)	Events/yr in ROI after 1 yr surface exposure	After stage 1 purification plus 5h re-exposure	After stage 2 (UG) purification plus 6 months “cool-down”
^{44}Sc (daughter of ^{44}Ti)	5.41	1.80	1.89E-5
^{46}Sc	20.3	0.04	9.30E-5
^{60}Co (direct and daughter of ^{60}Fe)	834	0.511	4.79E-3
^{68}Ga (daughter of ^{68}Ge)	344	0.703	2.03E-3
^{26}Al	2.20E-4	2.63E-7	2.63E-9
^{82}Rb (daughter of ^{82}Sr)	440	2.56	1.74E-4
^{88}Y (direct and daughter of ^{88}Zr)	3.61E4	37.9	0.213
^{42}K (daughter of ^{42}Ar)	10.0	0.90	7.72E-5
^{56}Co	0.350	6.83E-4	1.33E-6
^{58}Co	0.252	5.29E-4	8.88E-7
^{110m}Ag ^a	3.61E3	3.60	0.022
^{110}Ag (daughter of ^{110m}Ag) ^b	48.6	0.05	2.93E-4
^{106}Rh (daughter of ^{106}Ru)	21.8	0.022	1.58E-4
^{126m}Sb (direct and daughter of ^{126}Sn) ^c	8.63	8.58	4.34E-7
^{126}Sb (direct and daughter of ^{126m}Sb) ^d	1.29E4	154	5.44E-5
^{22}Na	1.01E3	0.711	6.22E-3
^{84}Rb ^e	24.2	0.11	2.31E-5
^{90}Y (daughter of ^{90}Sr)	7.90E-3	2.77E-4	1.80E-8
^{102}Rh (direct and daughter of ^{102m}Rh) ^f	35.9	0.044	2.39E-4
^{102m}Rh ^g	69.9	0.060	5.45E-4
^{124}Sb	1.62E5	417	0.509

Basic Detector Parameters for Phase I Demonstrator

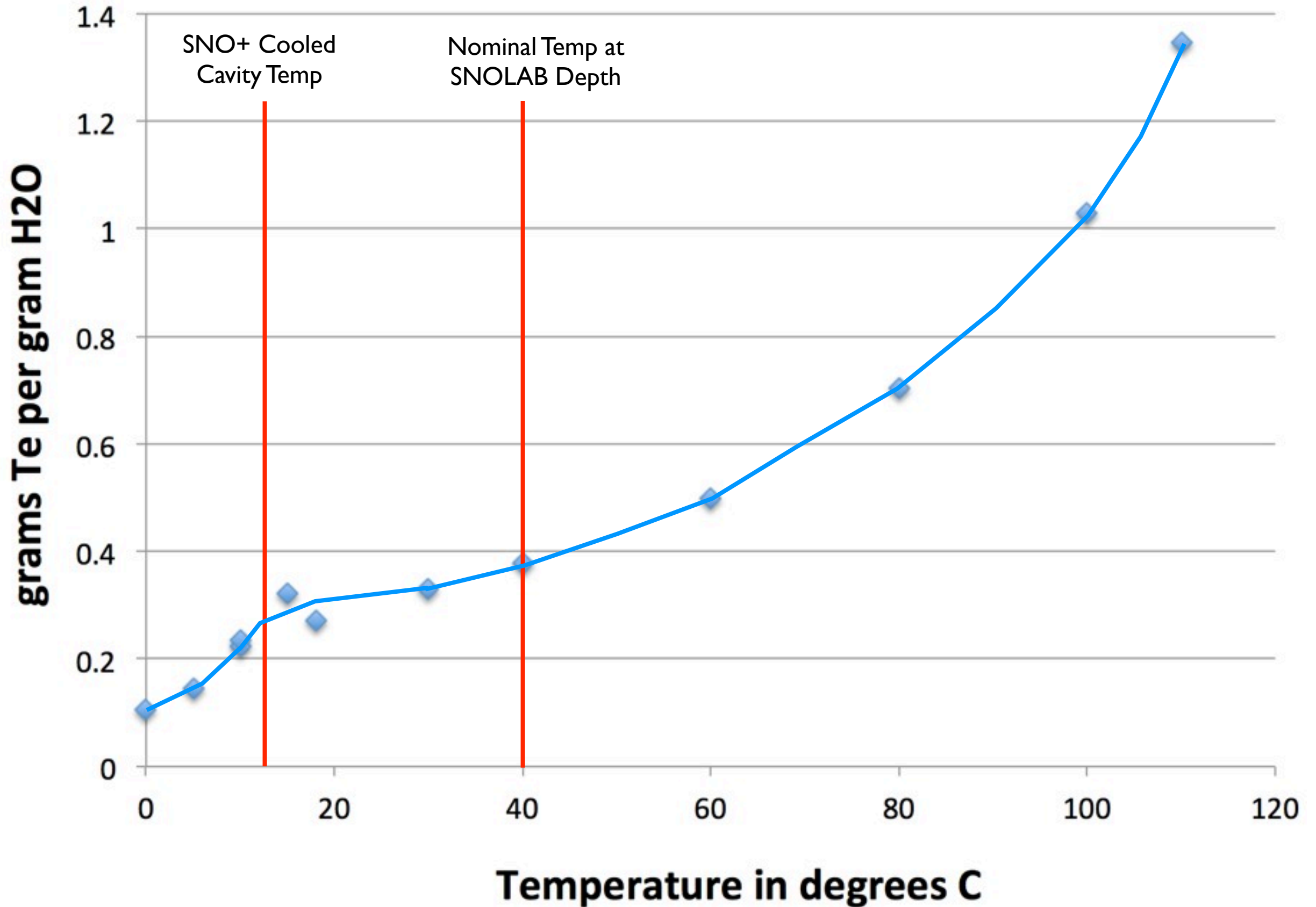
- Light Level: ~200-300 pe/MeV, depending on final optics and choice of secondary shifter. Assume 200 pe/MeV for this talk.
- Loading Level: 0.3-0.5% (0.8-1.3 tonnes ^{130}Te), depending on final Te system resources. Assume 0.3% for this talk.
- Fiducial Volume: 20-30%, depending on light level, loading fraction and final backgrounds. Assume 20% for this talk
(R=3.5m, ~10 times current K-Z fiducial volume)

Expected Average Spectra of Contributing Backgrounds for Two Live Years of Data



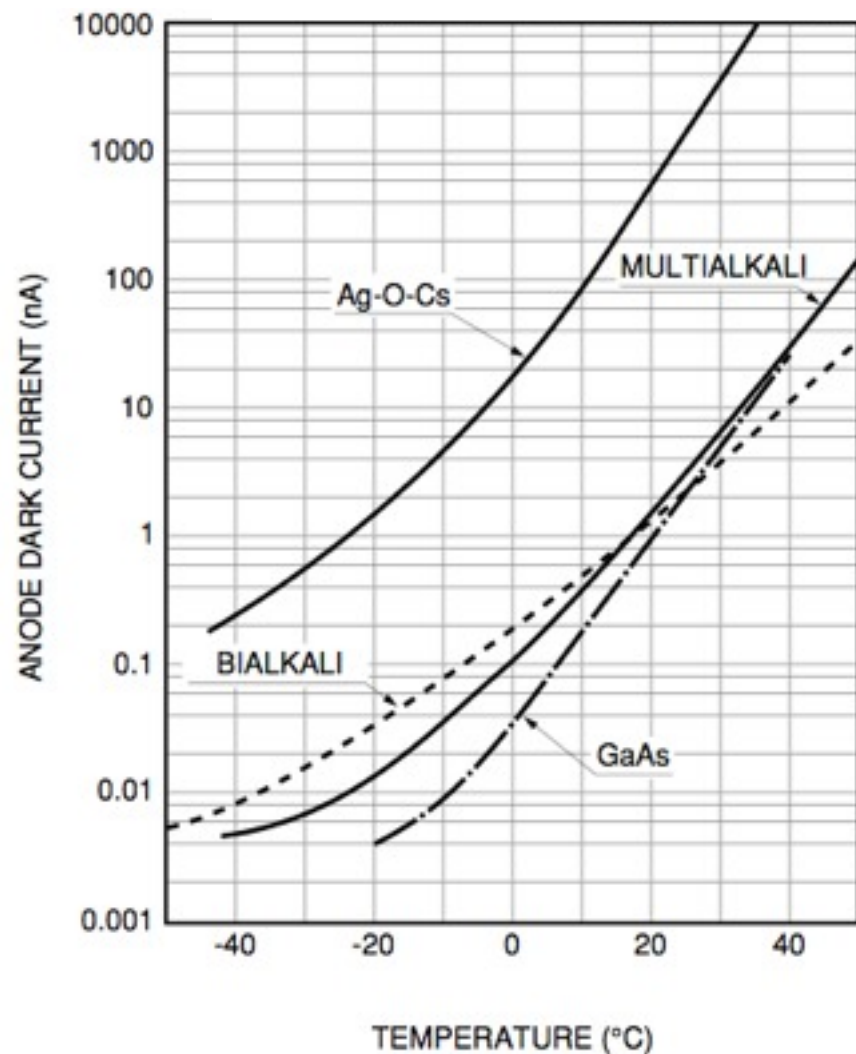
Pushing
Beyond
SNO+

Te Solubility (telluric acid in water)

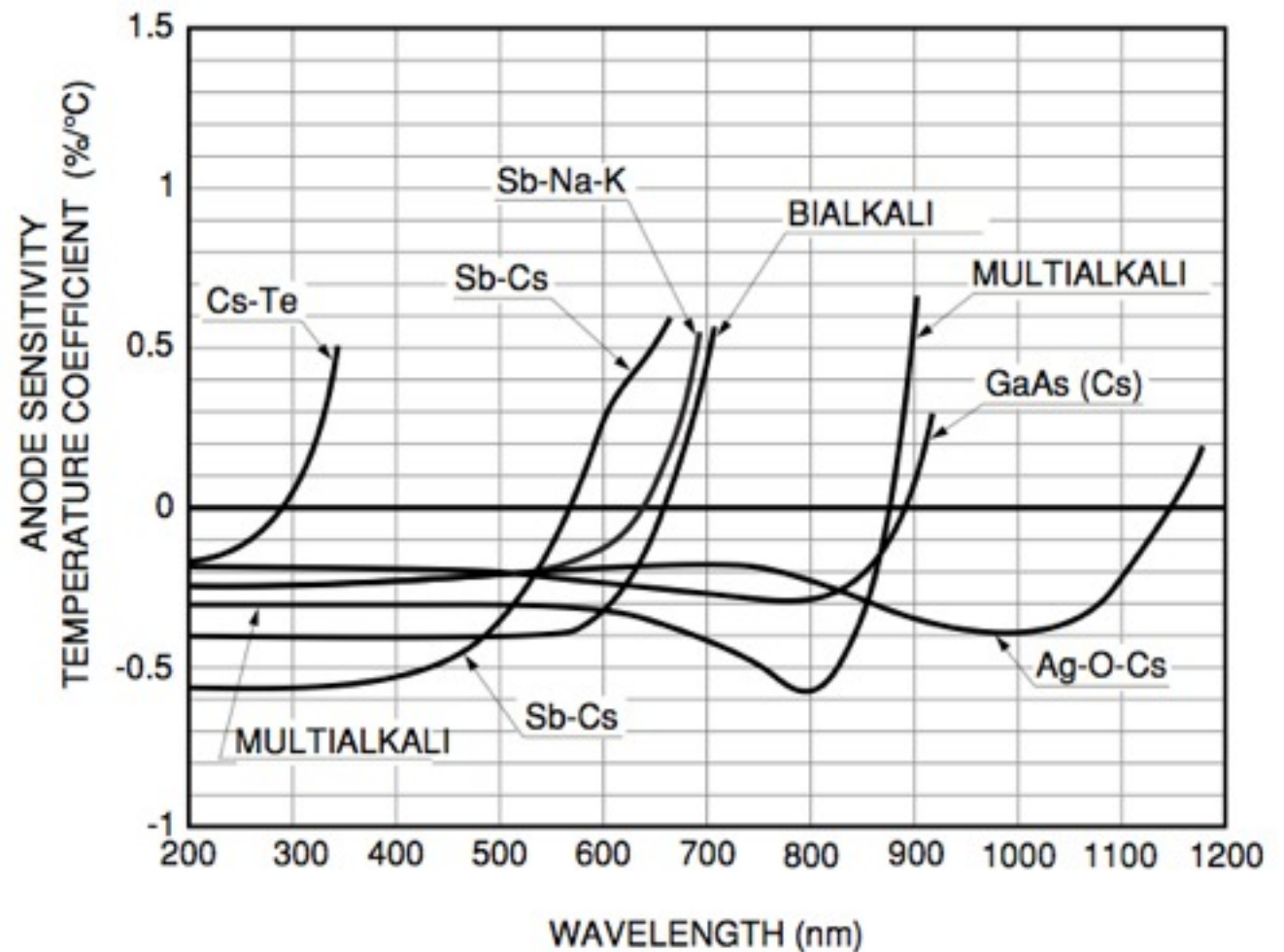


PMT performance at higher temperatures

(Hamamatsu PMT Handbook, 2007)



Factor of ~15 increase in dark current in going from 12 to 40 °C for bialkali



Anode sensitivity changes by $(28\text{ °C})(-0.4) = -11\%$ in going from 12 to 40 °C for bialkali

Heating to higher temperatures would be nice, but probably requires some new technologies [high temp PMTs (SbNa2K) have lower QE]

Loading:

	Temp (°C)	Solubility (g/g)	Percent Natural Te Loading	
			5% H ₂ O	25% H ₂ O
Likely	12	0.27	1.35	6.75
Possibly	40	0.38	1.90	9.5
			Likely	Possibly

Difficult, requires R&D, but not completely mad

Use more soluble Te compounds? Alternative loading method (maybe use organotelluric compounds)?

Light:

Scale light yield from SNO+ and assume a factor of ~ 2 light loss at higher loading

(SNO PMTs: only ~ 13 - 15% peak efficiency (25% QE, 55% collec. eff.), $\sim 45\%$ photocathode coverage)

Light Yield Estimate at Higher Loading

Potential SNO+
Light Yield (pe/MeV)
at 0.3% loading

Boost from QE x Coverage

		3 (1.7x1.7)	4 (2x2)
Likely	300	450	600
Possibly	500 (not currently, but pure LAB+PPO yields ~ 550)	750	1000
		Likely	Possibly

Detector Scale:

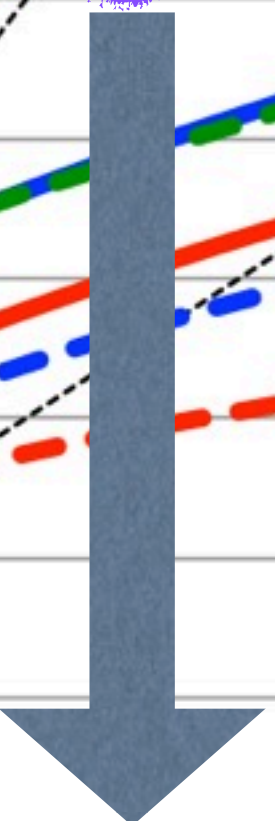
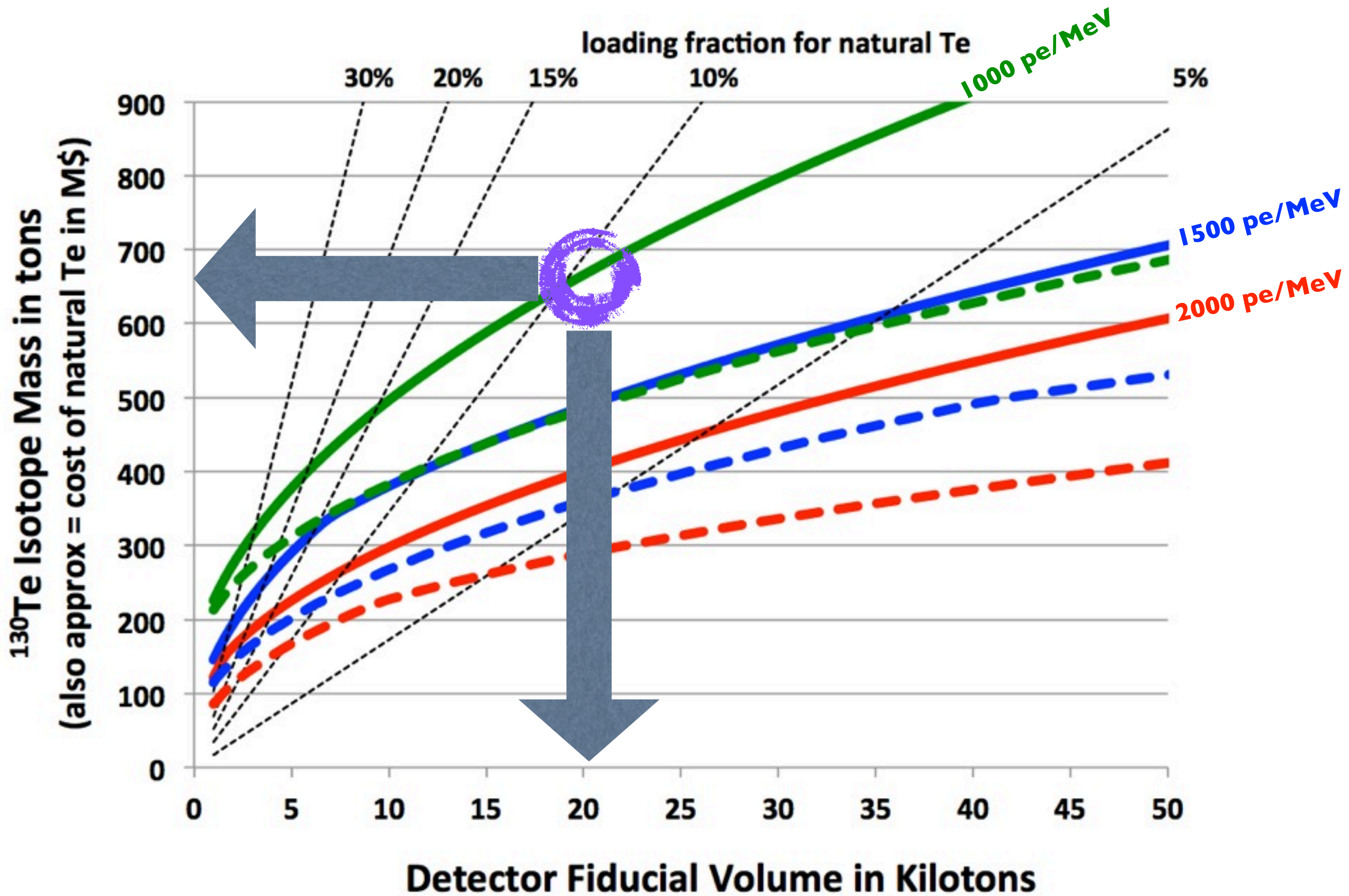
Absorption Length of Medium:

Spherical Volume ($d=\Lambda$, $\rho=1$):

20m	30m
4 kT	13.5 kT
Likely	Possibly

(LAB absorption
above 420nm)

However, light detection requirements favour multiple, small detectors... if finances allow.



Cherenkov Light?

Could this be picked out with time-separation and used to reduce ^8B and other backgrounds (perhaps even statistically test $0\nu\beta\beta$ model)?

(S. Biller, PRD 87, 071301(R), 2013)

need loaded scint with little absorption > 400 nm

Hand-Waving Argument

Cherenkov: $(8 \text{ pe/MeV}) \times (2) \times (2) \times (1/2) \times (2.5-0.5 \text{ MeV}) = 32 \text{ pe}$

scaling from SNO with water

assume boost to QE & coverage

assume half absorbed by scintillator

electron KE in ROI minus effective Cherenkov thresh

Scintillation: $(1000 \text{ pe/MeV}) \times (2.5 \text{ MeV}) \times (1/2) \times (5\text{ns}/25\text{ns}) = 250 \text{ pe}$

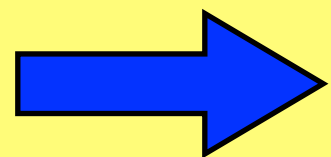
required light level

electron KE in ROI

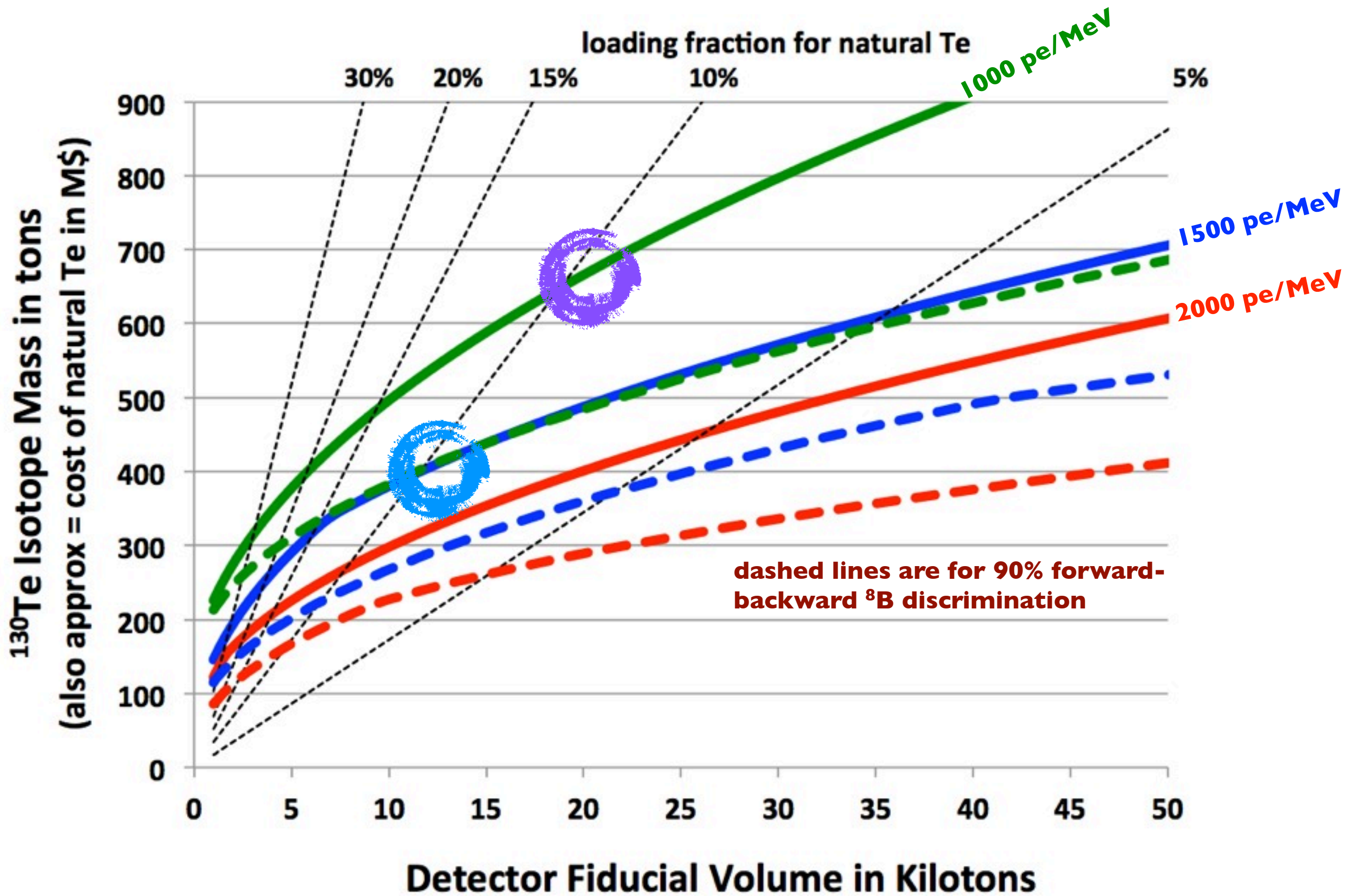
look for Cherenkov in "forward" hemisphere

assume 5ns prompt window and 25ns scint decay time

need Fluor with slow time const and high QY



$$32 / \sqrt{250} = 2\sigma \quad \text{"forward-backward" discrimination}$$



Conclusions:

- A practical normal hierarchy $0\nu\beta\beta$ experiment is **not** out of reach (“modest extrapolation”). With targeted R&D, construction for this could potentially even start within the next decade.
- Te-loaded liquid scintillator almost certainly offers the best (probably only) chance to achieve this.
- Current efforts from SNO+ look encouraging (if preliminary) and will provide an important test-bed for gaining a better understanding of the actual issues.
- There is still a lot to explore in the general development of this technology, which constitutes a worthy investment for the future programme.