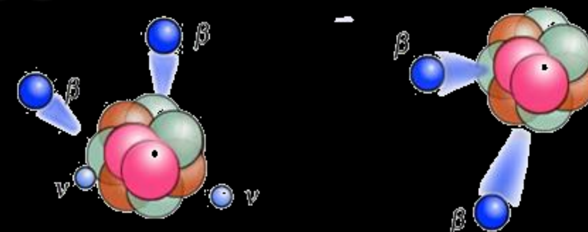
SNO+
Neutrinoless Double Beta Decay with an Organic Scintillator

Szymon Manecki, Queen's University

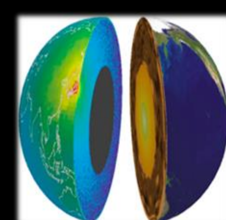
CAP Congress, May 31st, 2017

SNO+ Physics Goals

- Neutrinoless Double Beta Decay of ^{130}Te
- Low Energy Solar Neutrinos
- Reactor Antineutrinos
- Geo-Neutrinos
- Supernova-ν



- Three stages:
  - Water phase
  - Liquid scintillator phase
  - Te-loaded liquid scintillator



Double Beta Decay

- Are neutrinos their own anti-particles?
  - 2νββ (Dirac)

    $(A, Z) \longrightarrow (A, Z + 2) + 2e^{-} + 2\nu_e$

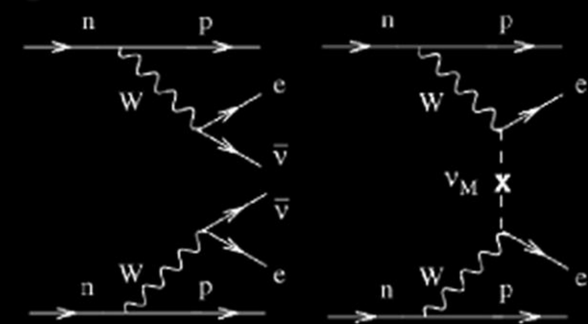
    ~ 10^{18}-10^{21} years
  - 0νββ (Majorana)

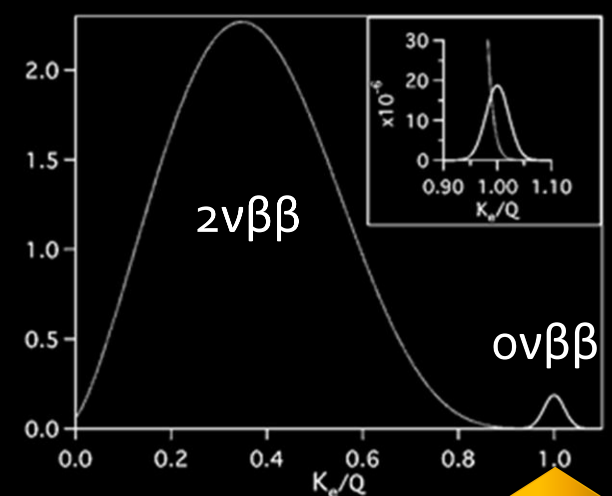
    $(A, Z) \longrightarrow (A, Z + 2) + 2e^{-}$

    > 10^{25} years
  - With the Mass Mechanism:

$$(T^{0\nu}_{1/2})^{-1} = G^{0\nu} \cdot |M^{0\nu}|^2 \cdot \langle m_{\beta\beta} \rangle^2$$

$$\langle m_{\beta\beta} \rangle^2 = |\textstyle\sum_i U^2_{ei}\, m_{\nu i}|^2$$





D.B.D. experiments need good energy resolution, low backgrounds, and large amounts of isotope.

SNO+ Detector

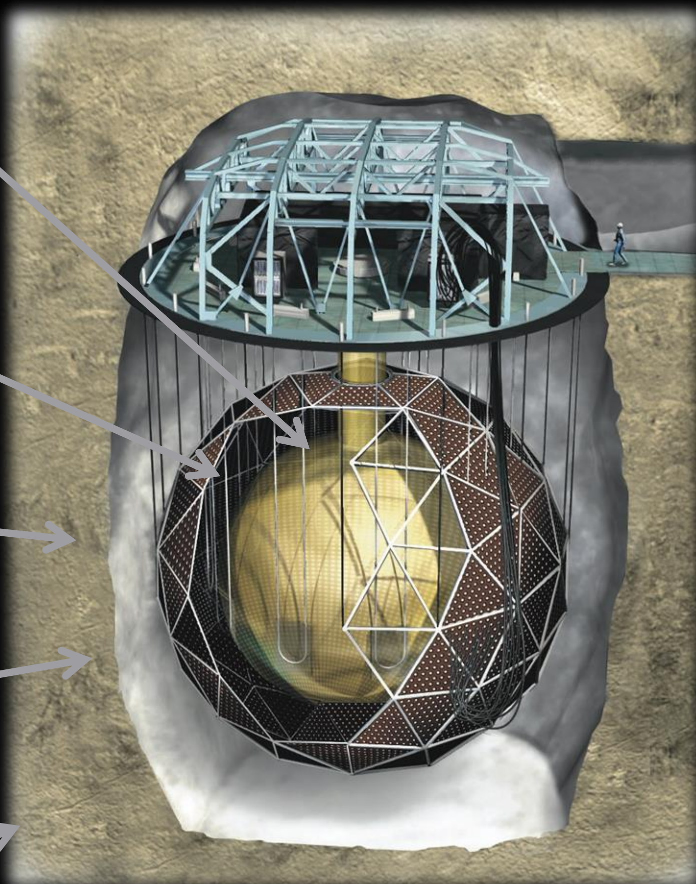
PMT Support Structure
~9400 PMT, 54% Coverage

Acrylic Vessel (AV)
Φ=12m, thickness=5cm

Light water (H_2O) shielding
- 1700t internal
- 5300t external

Urylon Liner/Radon Seal

Norite Rock

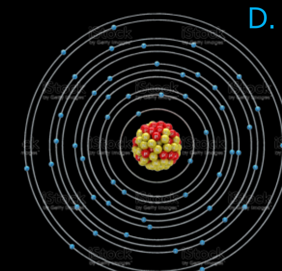


Linear Alkylbenzene (LAB)

R. Ford Talk

+

D. Bartlet Poster



Tellurium (^{130}Te)

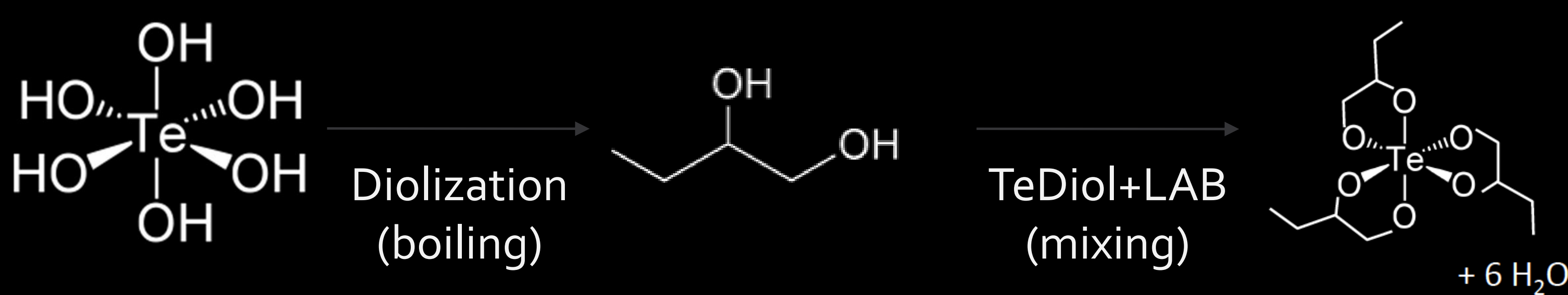
0νββ LS Requirements

- Reach high tellurium concentration
  - 0.5% Te in 780 tonnes of scintillator
- Preserve good optics of the cocktail
  - Transparency, Scattering, Light Yield
- Maintain high purity of the scintillator
  - U/Th reduction factor
  - Cosmogenic activation

The TeDiol Complex

- **Tellurium loading in Linear Alkyl Benzene**
  - **Through direct mixing in of an organometallic complex of Tellurium**

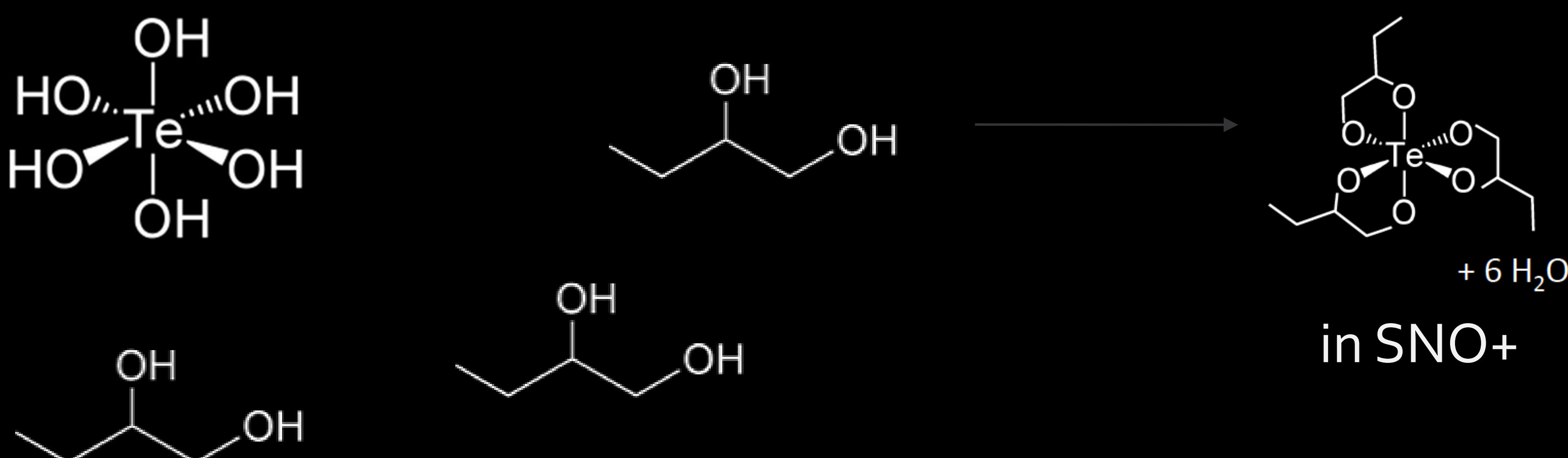
- **Butane-Diol based Te complex ("TeDiol"):**



Telluric Acid   Butanediol   in SNO+

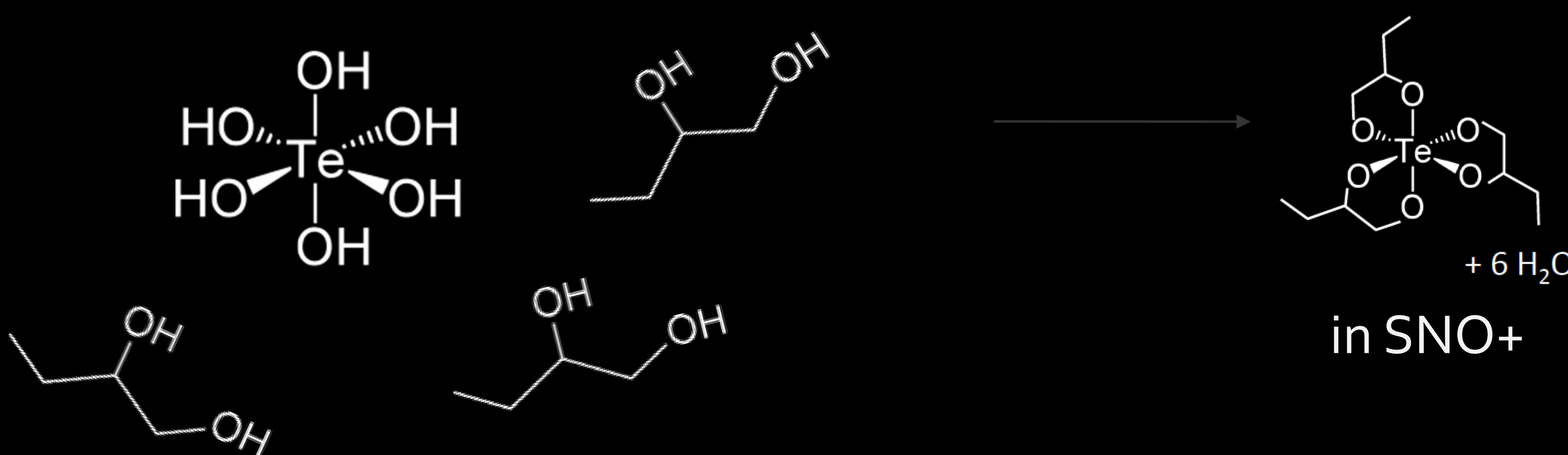
The TeDiol Complex

- Tellurium loading in Linear Alkyl Benzene
  - Through direct mixing in of an organometallic complex of Tellurium
- Butane-Diol based Te complex (“TeDiol”):



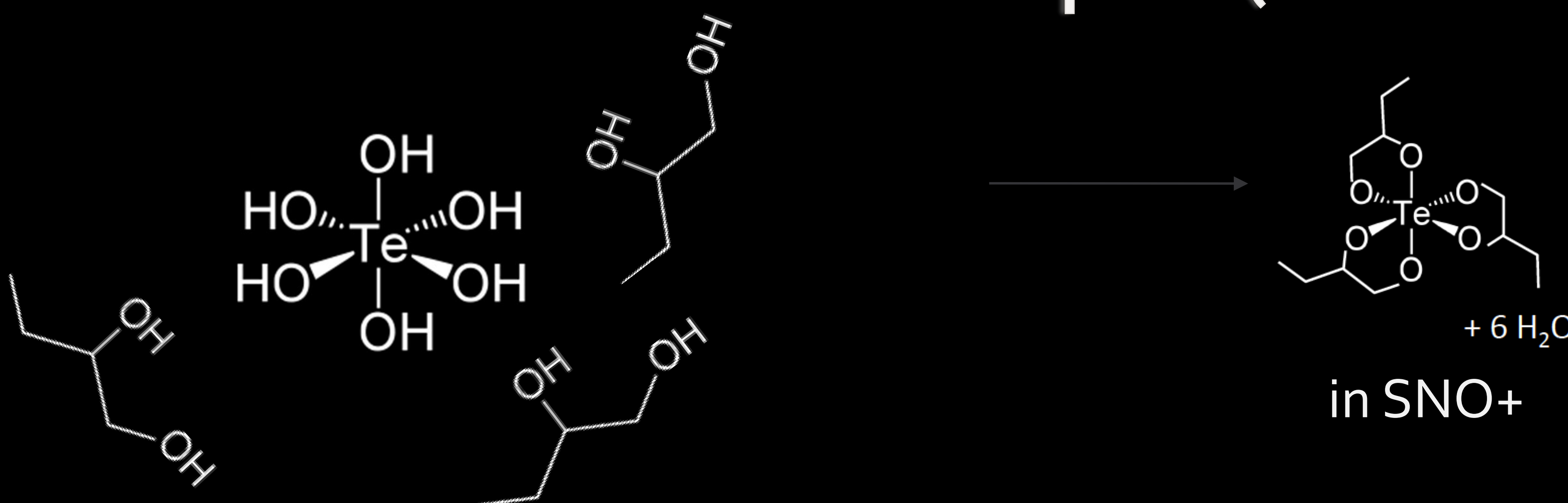
The TeDiol Complex

- Tellurium loading in Linear Alkyl Benzene
  - Through direct mixing in of an organometallic complex of Tellurium
- Butane-Diol based Te complex ("TeDiol"):



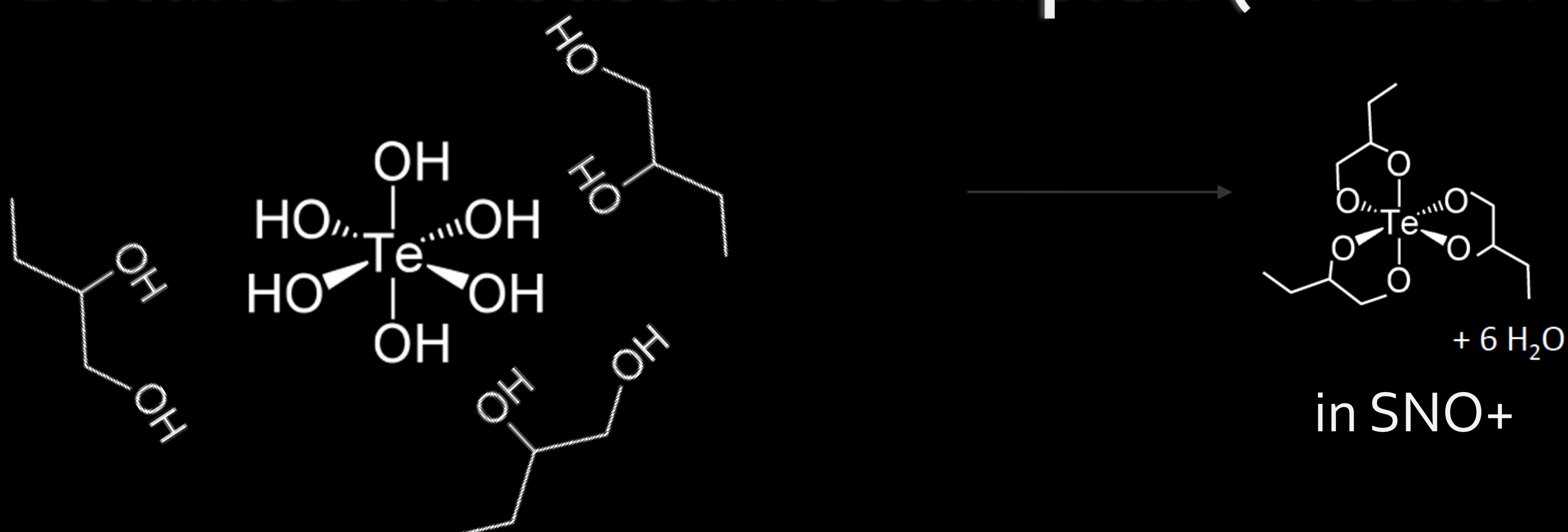
The TeDiol Complex

- Tellurium loading in Linear Alkyl Benzene
  - Through direct mixing in of an organometallic complex of Tellurium
- Butane-Diol based Te complex (“TeDiol”):



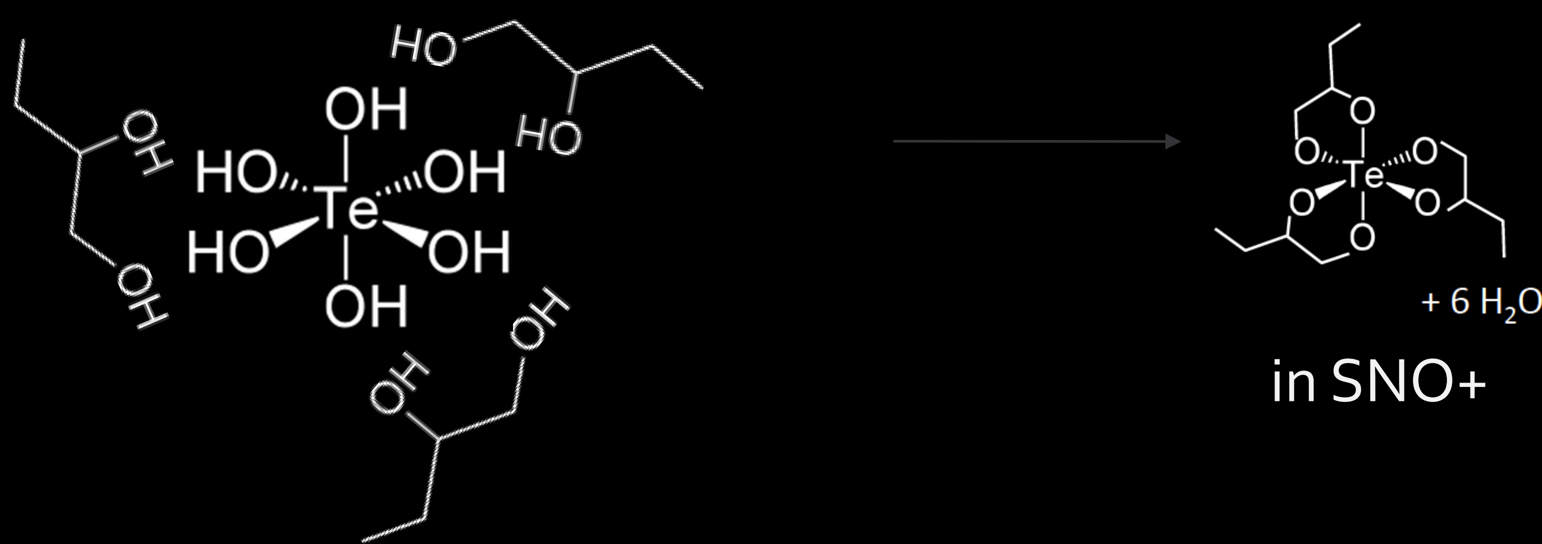
The TeDiol Complex

- Tellurium loading in Linear Alkyl Benzene
  - Through direct mixing in of an organometallic complex of Tellurium
- Butane-Diol based Te complex (“TeDiol”):



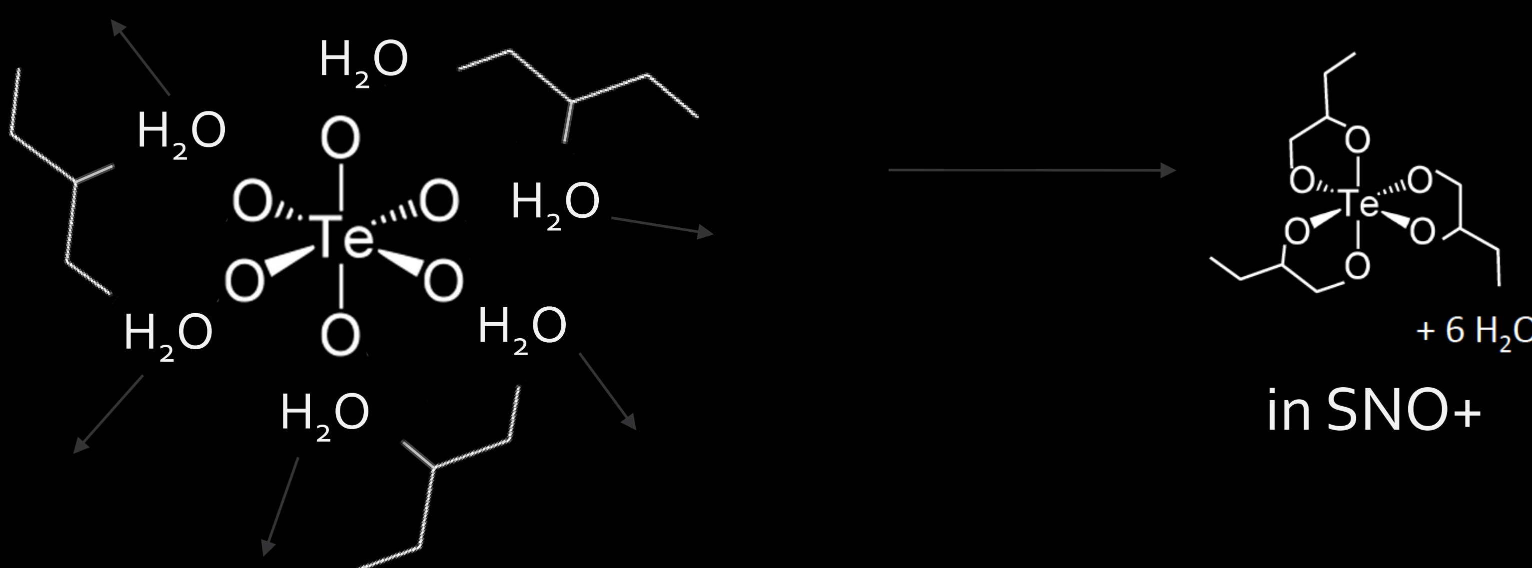
The TeDiol Complex

- Tellurium loading in Linear Alkyl Benzene
  - Through direct mixing in of an organometallic complex of Tellurium
- Butane-Diol based Te complex (“TeDiol”):



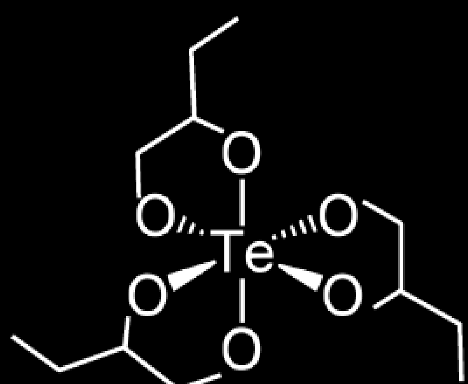
The TeDiol Complex

- Tellurium loading in Linear Alkyl Benzene
  - Through direct mixing in of an organometallic complex of Tellurium
- Butane-Diol based Te complex (“TeDiol”):



The TeDiol Complex

- Tellurium loading in Linear Alkyl Benzene
  - Through direct mixing in of an organometallic complex of Tellurium
- Butane-Diol based Te complex ("TeDiol"):

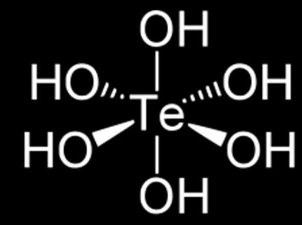


+ 6 H_2O

TeDiol+LAB
(mixing)

in SNO+

Telluric Acid Production

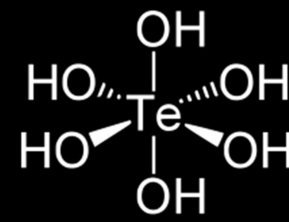


- Te extracted from mine (depth ~ 300 m) in April 2014
  - Visit to the production site prior to start of processing
  - QA/QC tests on samples from each barrel before approval to send to SNOLAB

3.8 tonnes of $Te(OH)_6$, corresponding to ~2.1 tonnes Te, or ~0.26% Te loading

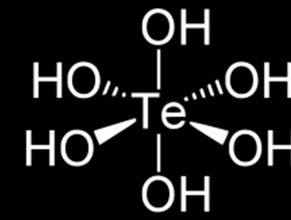
- Shipped to SNOLAB (January 7th 2015)
  - Transported underground on January 19th 2015
  - Testing one sample from one of the barrels to cross-check previous results

Telluric Acid Purification



- The purification technique relies on solubility of TeA in water based on pH
  - $Te(OH)_6 \leftrightarrow Te(OH)_5O^- + H^+$
    in-soluble soluble
  - Insoluble contamination
    - Dissolve in water, and filter
  - Soluble contamination
    - Force TeA to recrystallize by adding Nitric Acid, let it precipitate out, and drain the "dirty" liquid
  - The process can be made tellurium selective

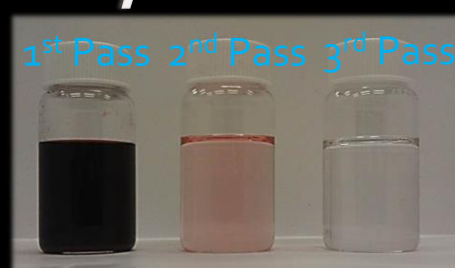
Telluric Acid Purification



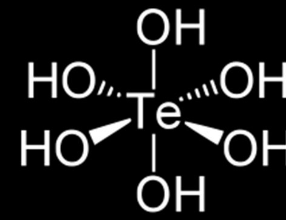
- 0.5% Tellurium Target levels:
  - 1.3×10^{-15} g/g in ^{238}U (3×10^{-8} Bq/kg)
  - 5×10^{-16} g/g in ^{232}Th (1.2×10^{-9} Bq/kg)
    - (raw Te ~10^{-11} g/g U/Th, 10^{-4} Bq/kg)
- Cosmogenic contamination from activation on Te
  - ^{60}Co, ^{110m}Ag, ^{126}Sn, ^{88}Zr, ^{88}Y, ^{124}Sb
    - Rejection needed 10^4-10^5

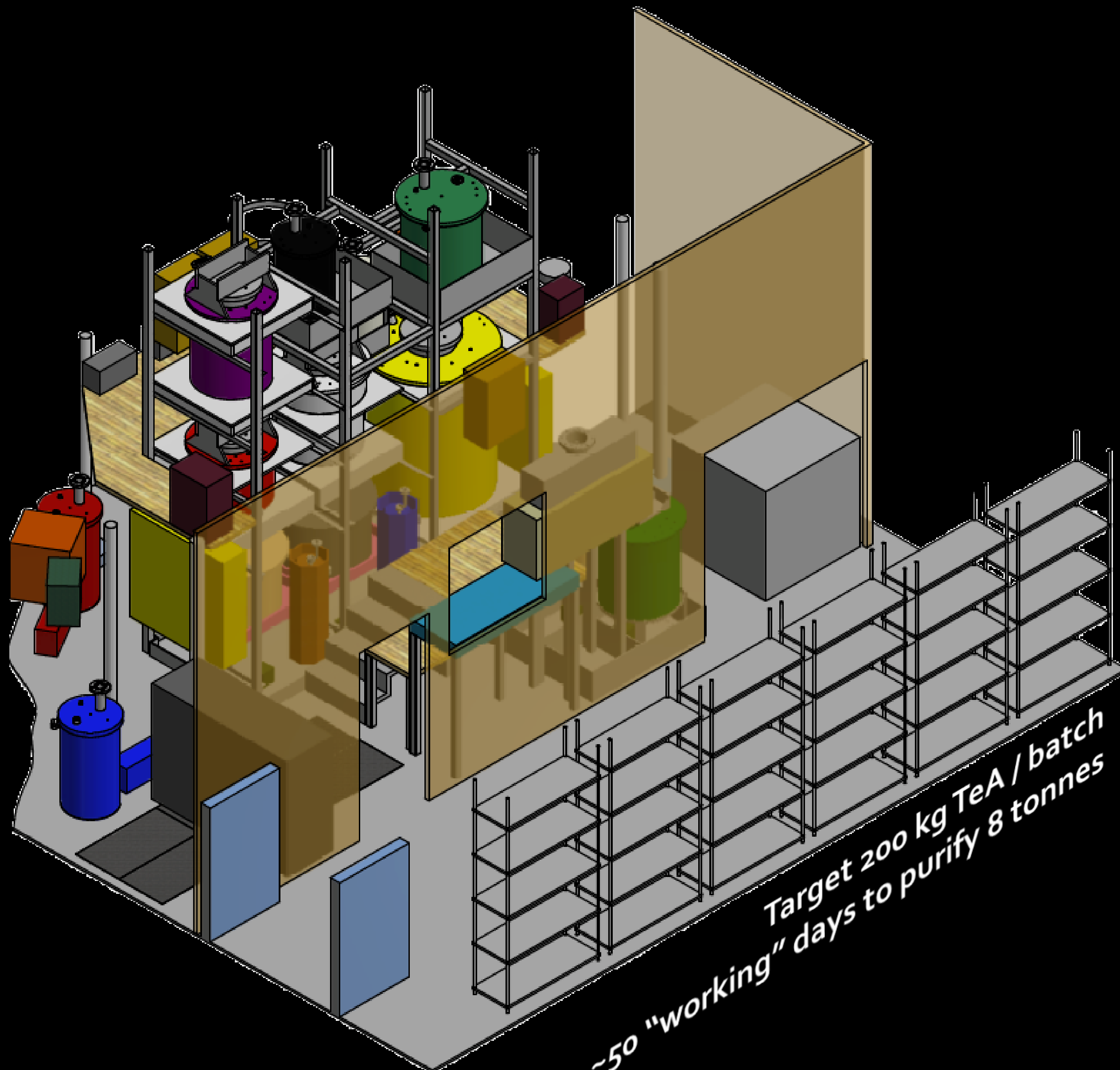
Isotope	t_{exp}=1 yr
^{22}Na	15309
^{26}Al	0.048
^{42}K	565
^{44}Sc	102
^{46}Sc	43568
^{56}Co	2629
^{58}Co	25194
^{60}Co	6906
^{68}Ga	37343
^{82}Rb	18047
^{84}Rb	11850
^{88}Y	390620
^{90}Y	823
^{102}Rh	276189
^{102m}Rh	133848
^{106}Rh	1534
^{110m}Ag	69643
^{110}Ag	939
^{124}Sb	3101138
^{126m}Sb	240
^{126}Sb	358996

10kg pilot-scale plant operated successfully
Final design ~200 kg TeA/batch under construction

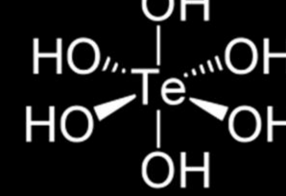


Telluric Acid Purification

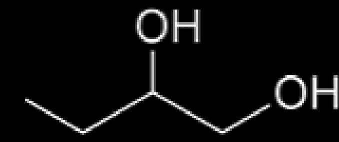




Telluric Acid Purification



The Diol Assay



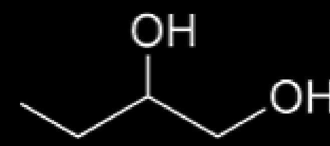
- Identified distributor in Japan, Kowa-Co.
  - High quality and affordable (8 tonnes needed)
  - $^{14}C/^{12}C$ to confirm its non-biogenic origin
    - Accelerator Mass Spectrometry at uOttawa:
      - Sample #1: $(14.3 \pm 1.2) \times 10^{-16}$ Blank #1: $(26.0 \pm 7.4) \times 10^{-17}$
      - Sample #2: $(4.8 \pm 1.2) \times 10^{-16}$ Blank #1: $(2.5 \pm 1.2) \times 10^{-17}$

The André E. Lalonde AMS Laboratory

University of Ottawa
25 Templeton Street
Ottawa, ON K1N 6N5
Canada

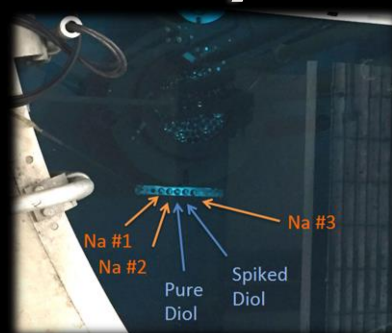
Radiocarbon@uOttawa.ca

The Diol Assay



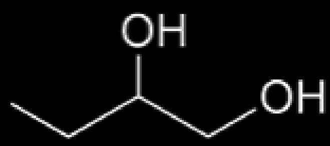
- Gamma-ray spectrometry
  - High Purity Ge (HPGe) detector at SNOLAB
    - ^{238}U < 3.13 ppb
    - ^{232}Th < 0.26 ppb
    - ^{40}K < 386.56 ppb

- Neutron Activation Analysis
  - NAA at UC Davis
    - ^{238}U < 0.3 ppb
    - ^{232}Th < 3.3 ppb
    - ^{nat}Na ~ ppm -> a fraction of which is ^{22}Na

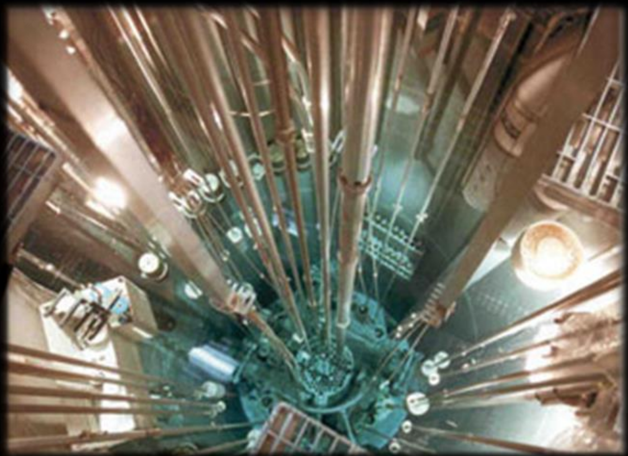


Experiment or Laboratory	Total (2005 - Today)
SNO	11
SNO+	125
SNOLAB	81
EXO	19
MiniCLEAN	56
DEAP	133
HALO	13
PICASSO	9
DM-ICE / DRIFT	23
COUPP / PICO	92
DAMIC	15
NEWS-SNOLAB	1
Total	578
Calibrations & Tests	118

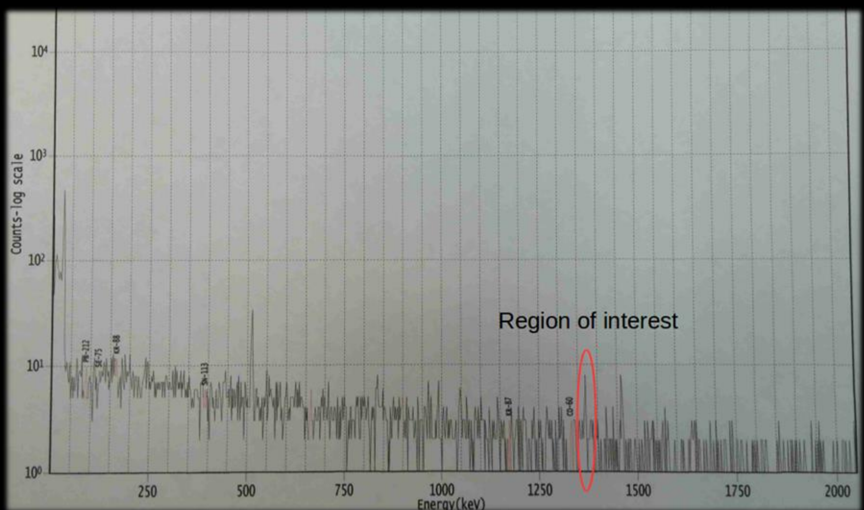
The Diol Assay



- **Tracing sodium contamination with NAA**

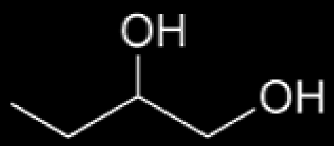


TRIGA-type research reactor in Sacramento, owned and operated by UC Davis



Na (2.2 ± 1.0) ppb

The Diol Purification

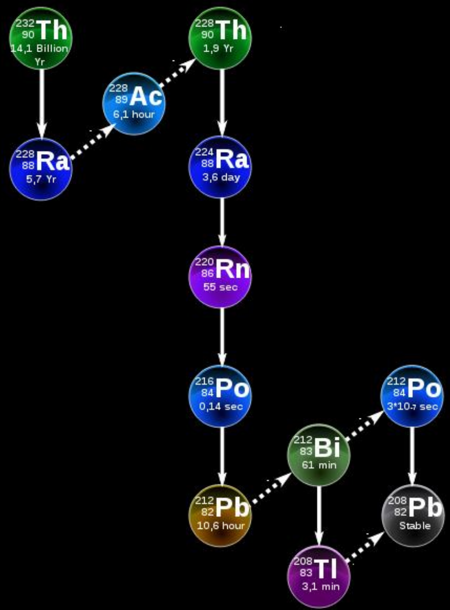


- Bench-top distillation with radio spikes
  - ^{228}Th spike in 1,2-Butanediol
    - Low T (70 °C, 80 mTorr)

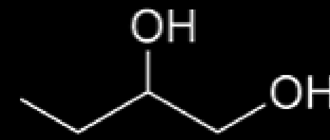
	Initial activity mBq/g	Distillate activity mBq/g	Reduction factor
^{228}Th	72	<0.014	>5100
^{224}Ra	72	<0.013	>5500

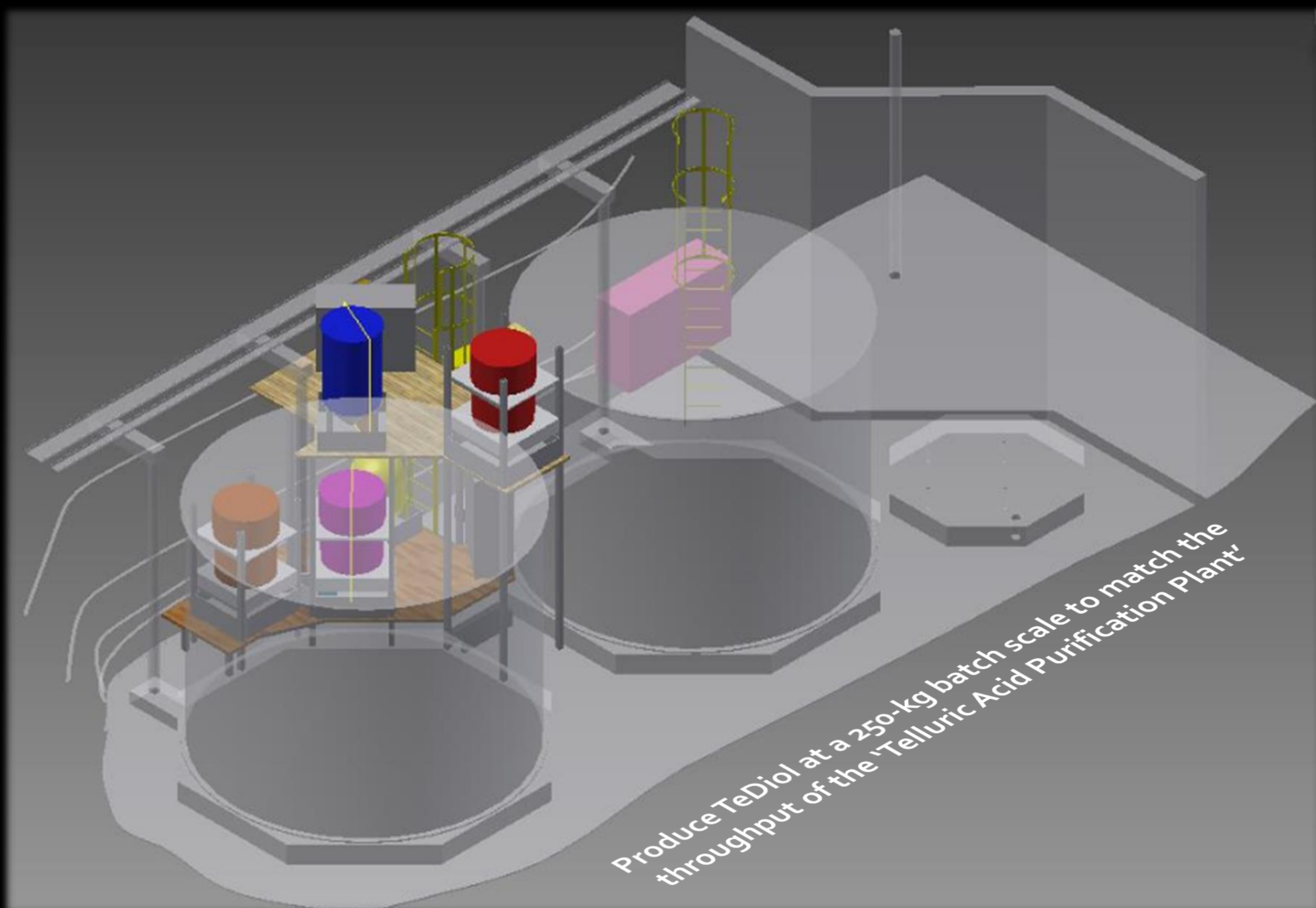
    - High T (170 °C, 225 Torr)

	Initial activity Bq/g	Distillate activity μBq/g	Reduction factor
^{228}Th	1.94	7 ± 1	280 000
^{224}Ra	1.94	13 ± 5	150 000

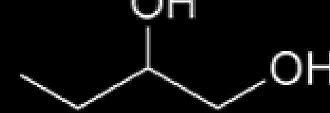


The TeDiol Plant

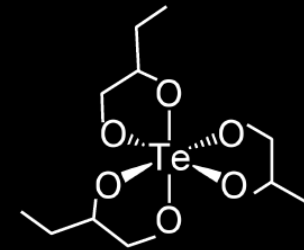




The TeDiol Plant



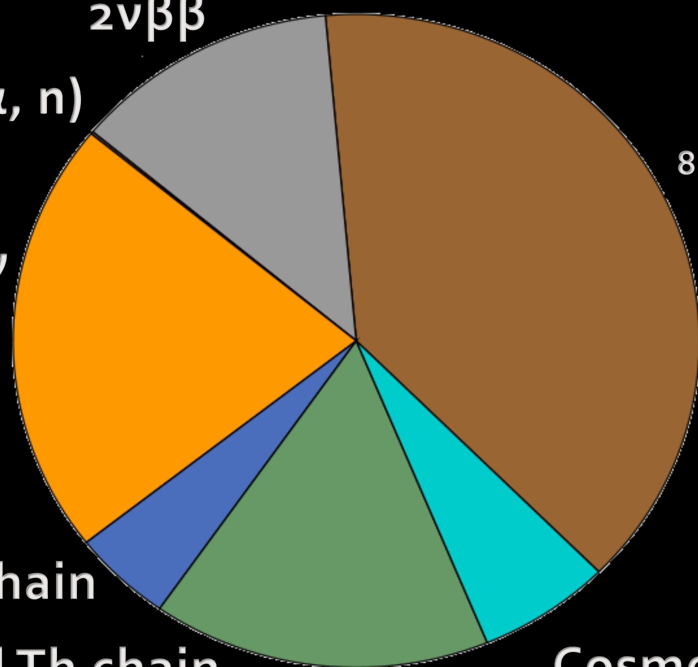
Backgrounds Budget



(α, n)
Alpha-capture on $^{13}C/^{18}O$
Neutrons produced
Capture of thermal neutrons
Delayed coincidence tag

^{8}B ν ES
^{8}B solar neutrinos:
Flat spectrum
Constrained by SNO/SK data
Limited by resolution

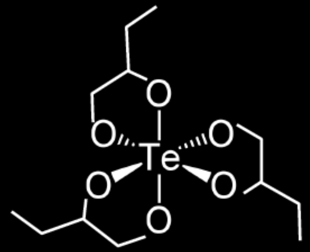
External γ
From AV, ropes, water, PMTs
Fiducial volume (20%) cut
50% extra rejection multi-site cuts



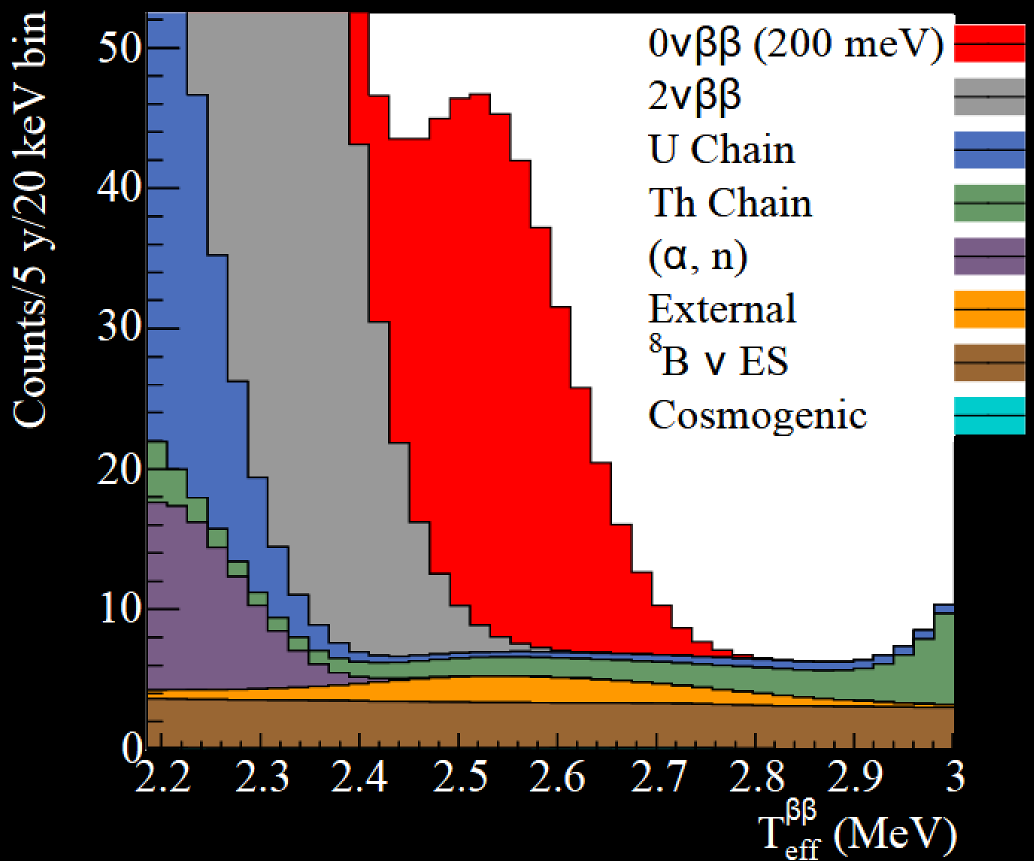
Cosmogenics
Mitigation: purification
+ "cool-down" UG
< 1ev/yr in RoI-FV
Further reduction if
needed: multi-site events

Internal U/Th
$^{214}BiPo$, $^{212}BiPo$
B-α delayed coincidence tagging
100% rejection in RoI
In-window trigger: x50 rejection

0νββ Sensitivity



- 1.3 tonnes of ^{130}Te in LAB (at 0.5% ^{nat}-Te)



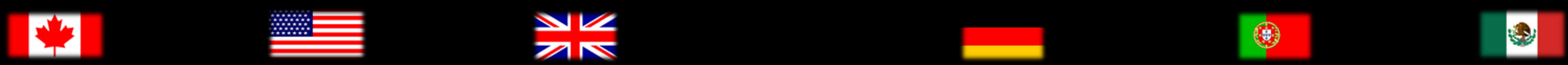
- [-0.5; +1.5] σ around $Q_{\beta\beta}$
- 400 NHits/MeV (~4% ΔE)
- Fiducial Volume: 20% total

	$T_{1/2}$ [yr]	$m_{0\nu\beta\beta}$ [meV]
1 yr	$8x10^{25}$	75.2
5 yrs	$1.96x10^{26}$	38 – 92

oνββ Schedule

- 2017-2018
  - Scintillator plant commissioning
  - Scintillator fill
    - Unloaded scintillator phase (short)
    - Evaluation of backgrounds for oυββ
  - Commissioning of the Tellurium plant(s)
- 2018-2019
  - Tellurium loading
  - Begin oυββ phase

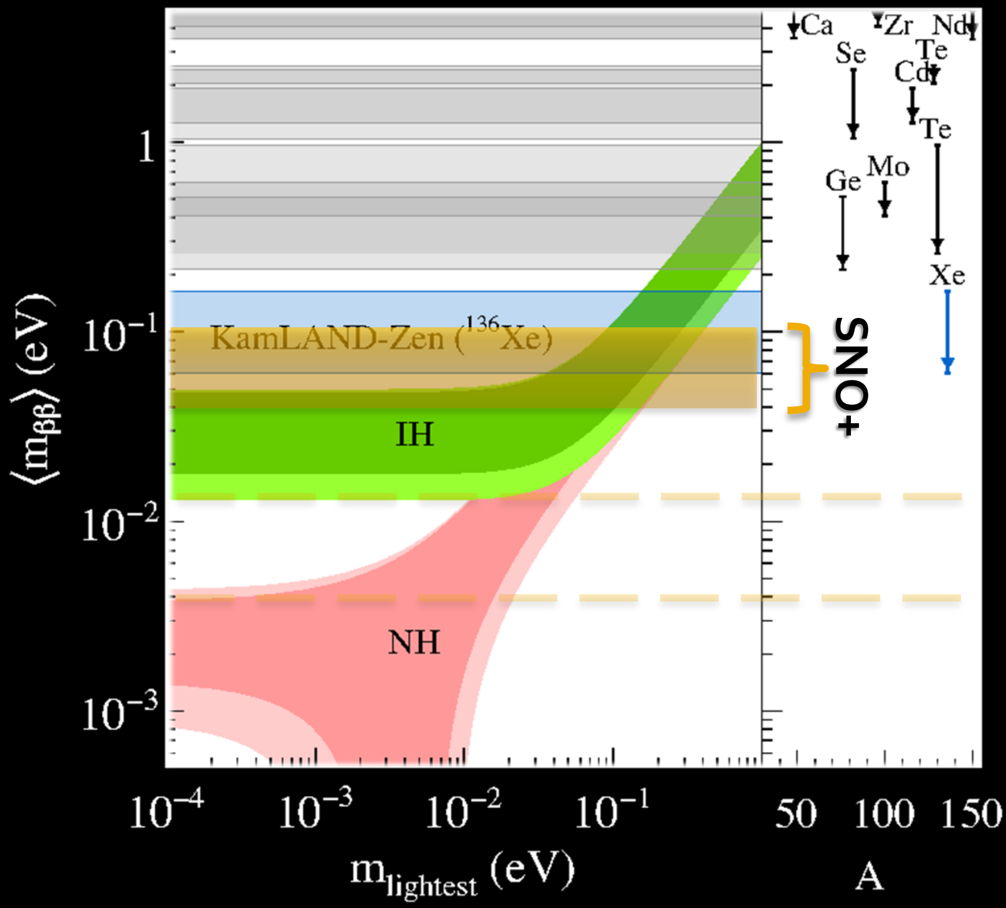
SNO+ Collaboration



Backup

0νββ Sensitivity

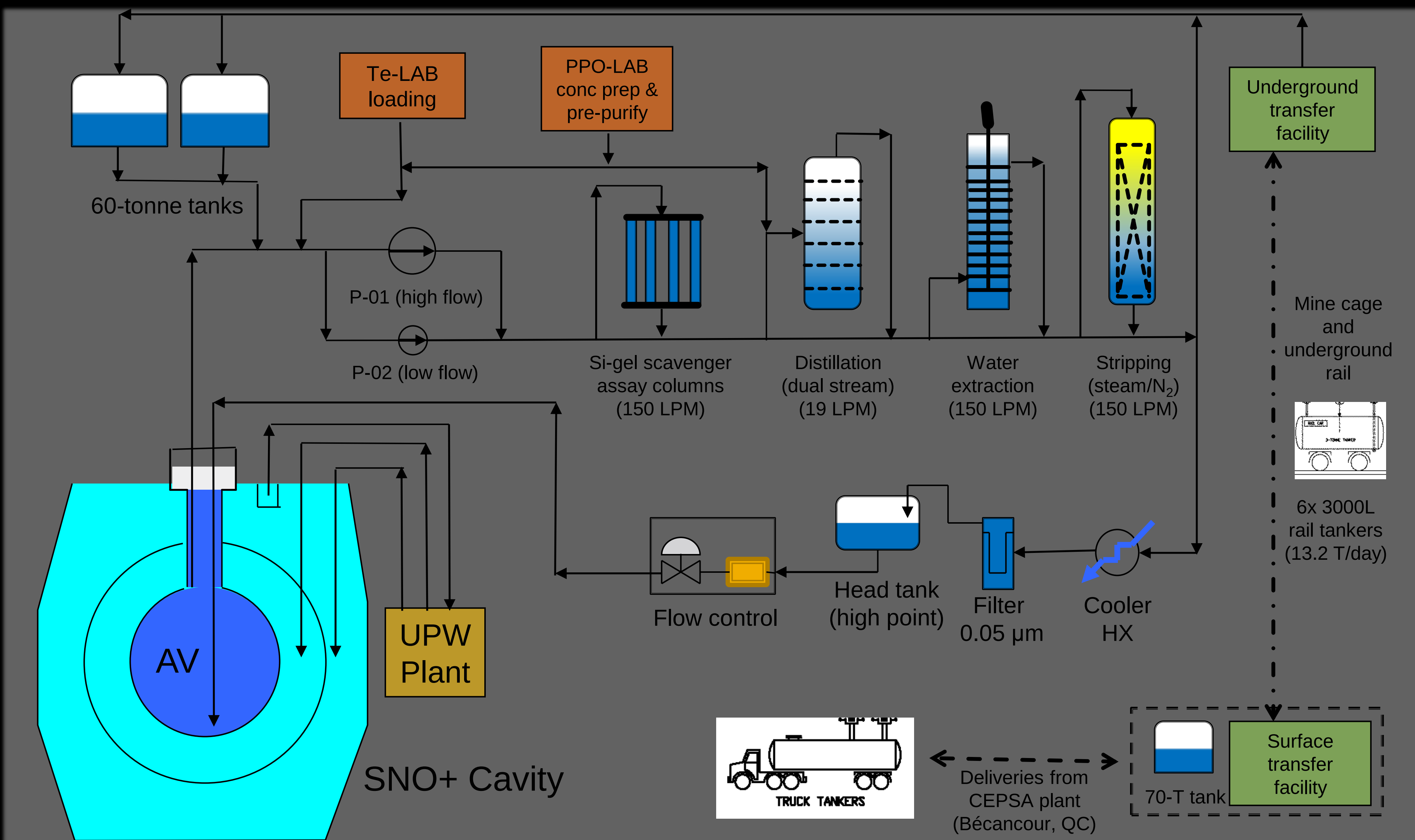
- 1.3 tonnes of ^{130}Te in LAB (at 0.5% $^{nat-}$Te)



- [-0.5; +1.5] σ around $Q_{\beta\beta}$
- 400 NHits/MeV (~4% ΔE)
- Fiducial Volume: 20% total

	$T_{1/2}$ [yr]	$m_{0\nu\beta\beta}$ [meV]
1 yr	8×10^{25}	75.2
5 yrs	1.96×10^{26}	38 – 92

Scintillator Purification Plant



Scintillator Purification Plant

- Multi-stage distillation
  - Dual-stream PPO distillation
  - Removes heavy metals
  - Improves UV transparency
- N_2 / steam stripping
  - Removes Rn, Kr, Ar, O_2
- Water extraction
  - Removes Ra, K, Bi
- Metal scavenging
  - Removes Bi, Pb
- Microfiltration
  - Removes dust
  - Target Levels
    - ^{85}Kr: 10^{-25} g/g
    - ^{40}K: 10^{-18} g/g
    - ^{39}Ar: 10^{-24} g/g
    - U: 10^{-17} g/g
    - Th: 10^{-18} g/g

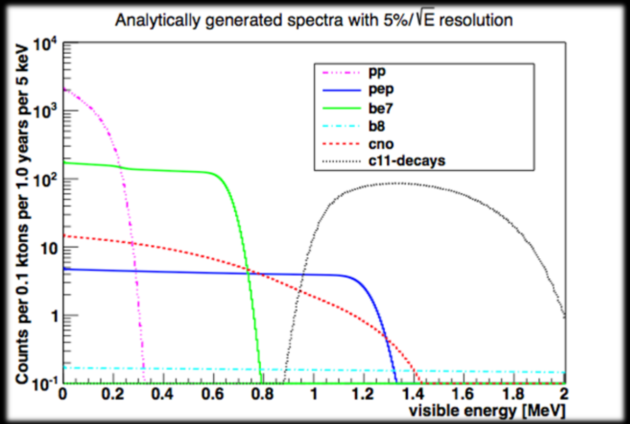
Backgrounds Budget

- Current sensitivity studies assume that in the background budget, solar neutrinos would be the dominant factor
  - $2\nu\beta\beta$ spectrum "leaks" into the ROI [8.5/23.2 c-yr]
    - Improved energy resolution with good optics
  - External backgrounds (^{208}Tl ,^{214}Bi) [3.5/23.2 c-yr]
    - Minimized with proper fiducialisation, and PSD
  - Internal backgrounds and detector response
    - U/Th [3.8/23.2 c-yr] and cosmogenics [0.1/23.2 c-yr] reduced by purification & cooling
    - Bi-Po/(α,n) tagged with space-time coincidence
    - ^{210}Po-$2\nu\beta\beta$/^{210}Bi-$2\nu\beta\beta$ pile-up events reduced based on PMT-hit time distribution
    - Apply the "source-in – source-out " approach
  - Flat ^{8}B (ES) e^- normalized to known flux [7.2/23.2 c-yr]

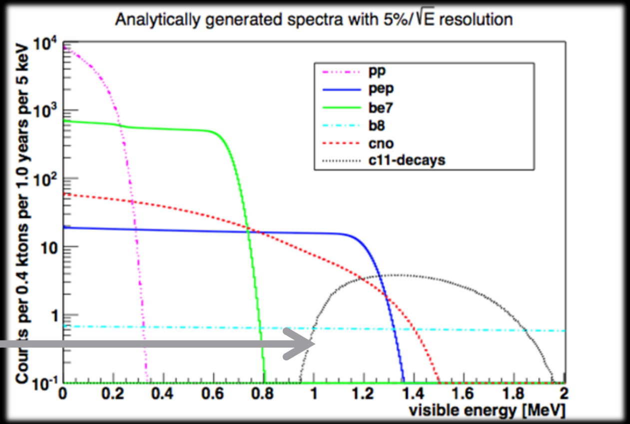
SNOLAB Facility

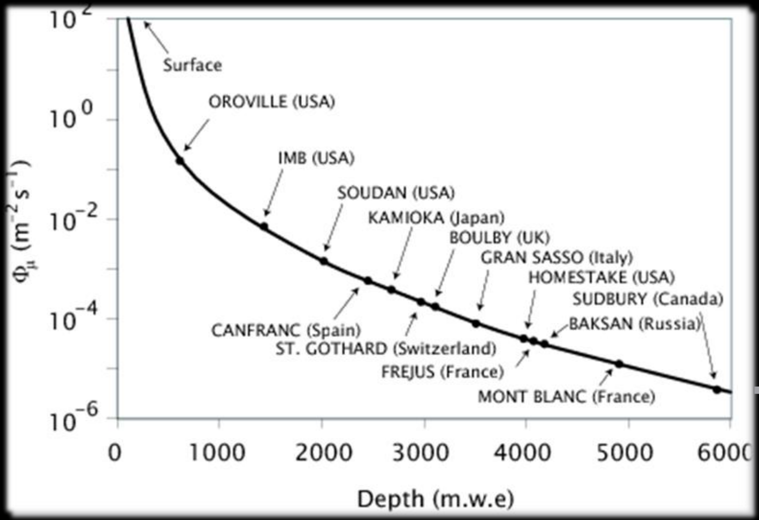
- Depth = 2070 m (6000 m.w.e.)
- 60 muons/day in SNO+
- 10,000 sq ft class-2000 clean room

LNGS



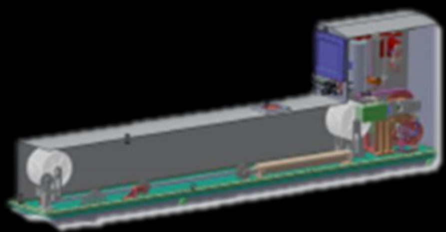
SNOLAB

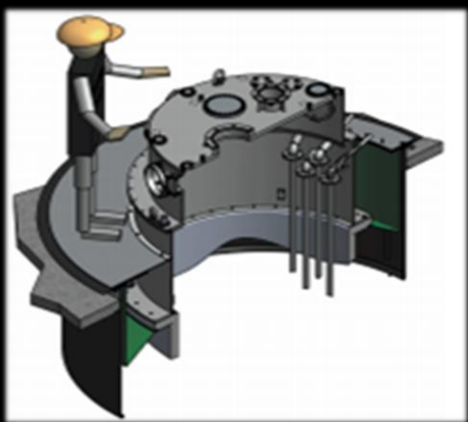


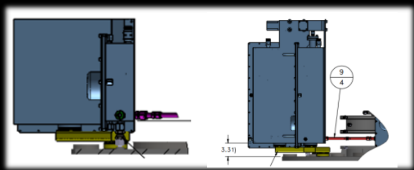


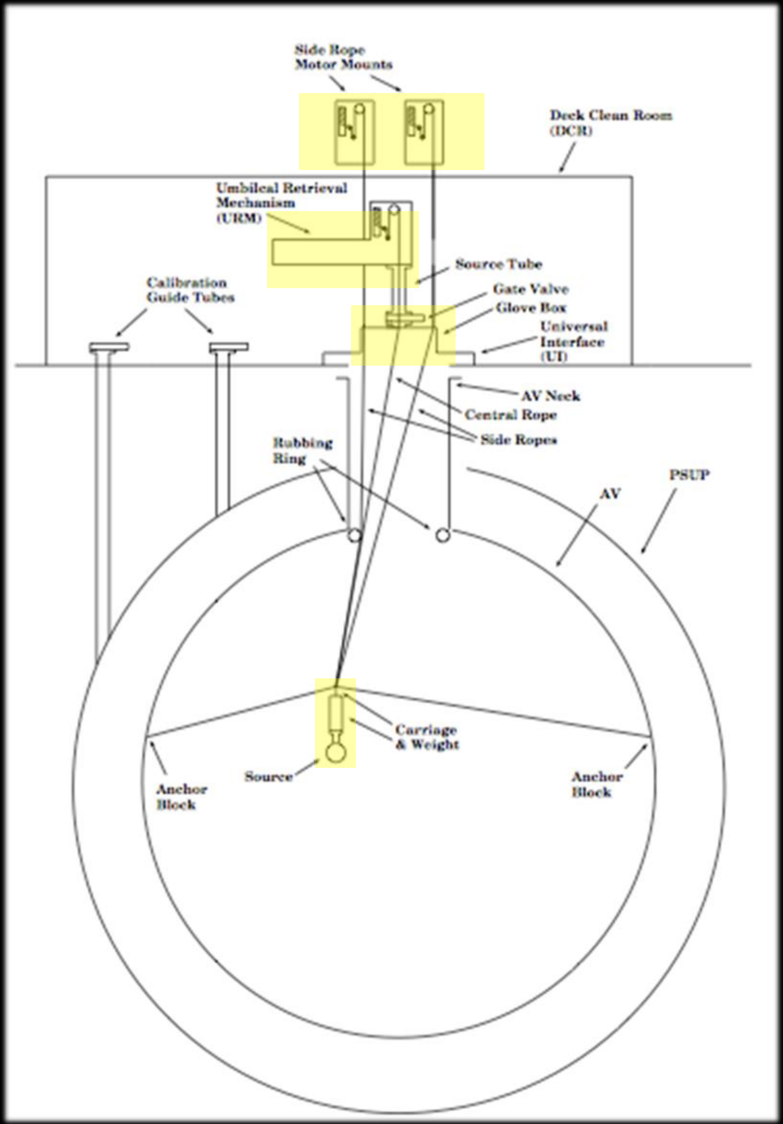
Calibration Hardware

New (Re)Design

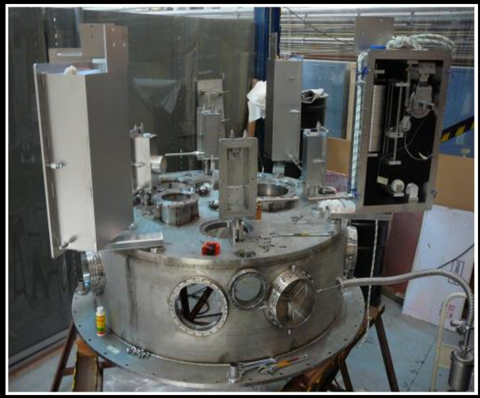








New Technology

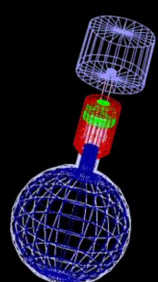


Calibration Sources

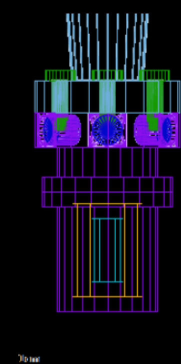
- **Need Double encapsulation**
  - Limitation for ^{222}Rn, ^{90}Y

- **Radioactive and optical sources α, β, γ, n, with laser injection laserball and Cherenkov**

LED

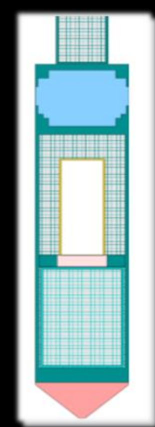


^{48}Sc

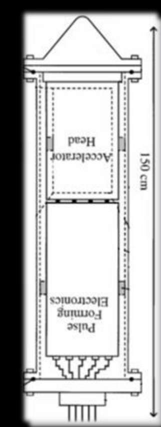


~7.12 s

$C^{16}N_2$



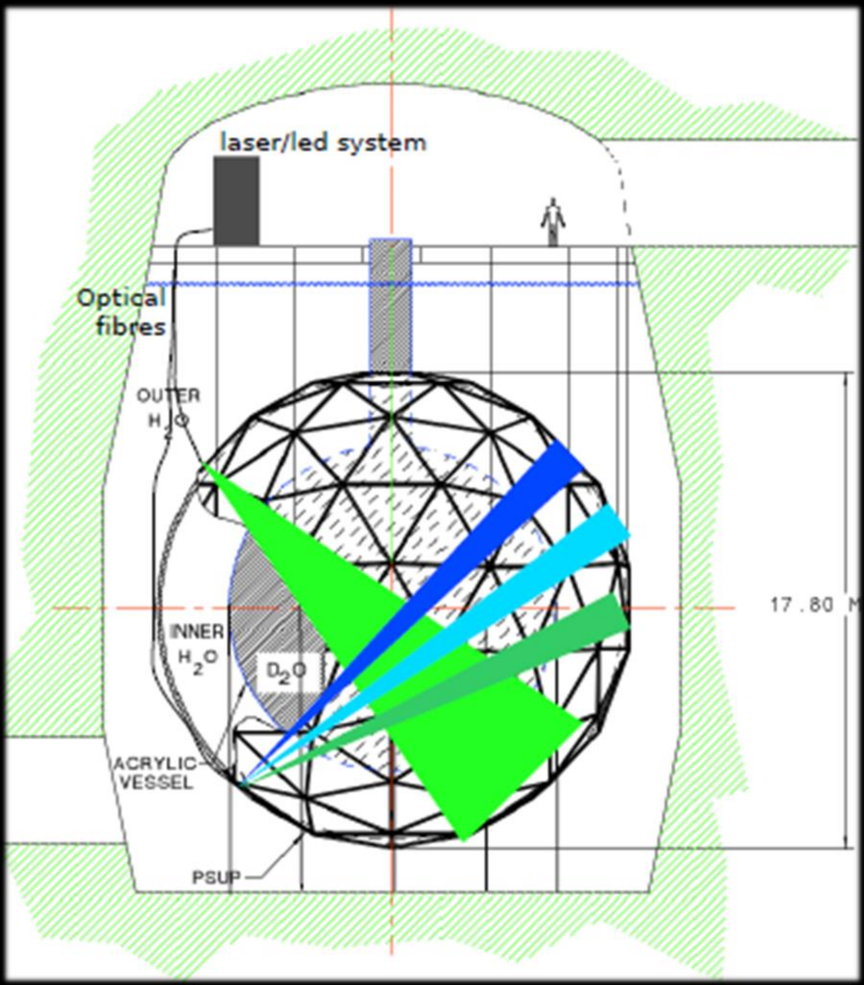
CO_2

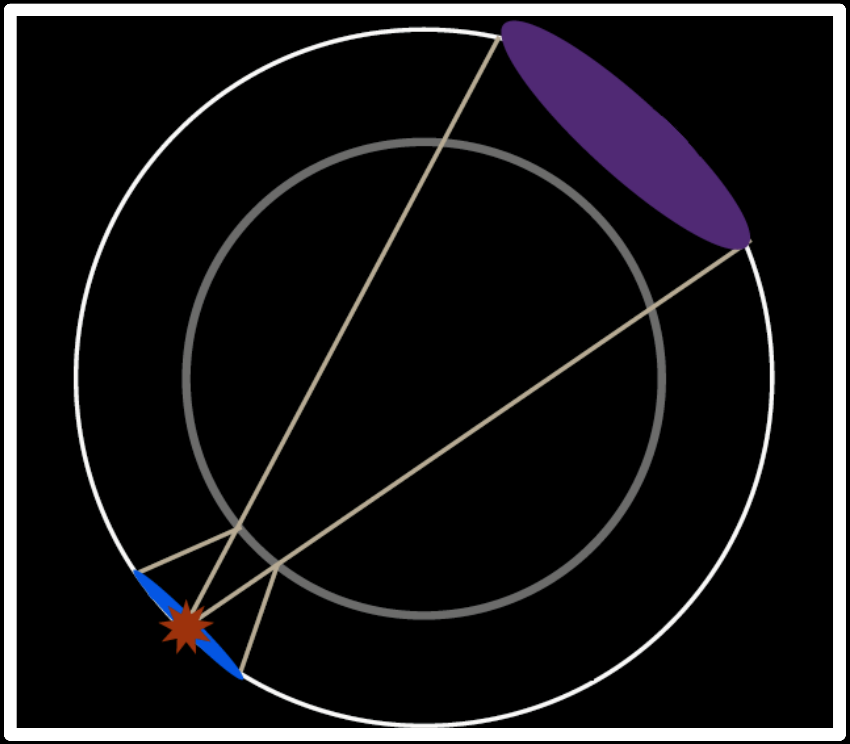


^{24}Na

Type	γ					β	α	n		
Src.	^{57}Co	^{60}Co	^{48}Sc	^{24}Na	^{16}N	^{90}Y(?)	^{214}Po(?)	n-p	n-^{12}C	n-Fe
MeV	0.1	2.5 (sum)	3.3 (sum)	4.1 (sum)	6.1	2.3	7.7	2.2	4.9	~7.5

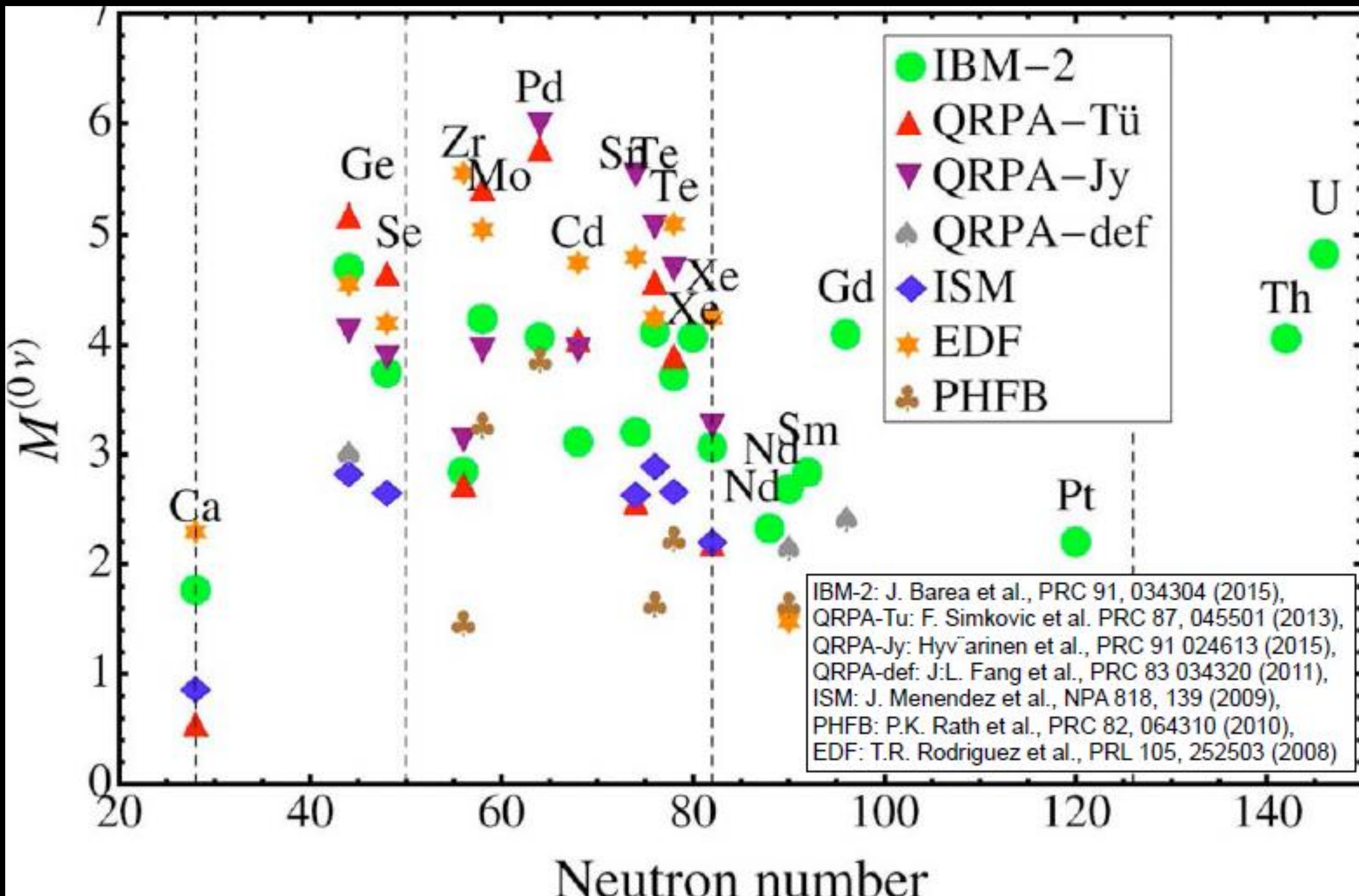
Optical Calibration





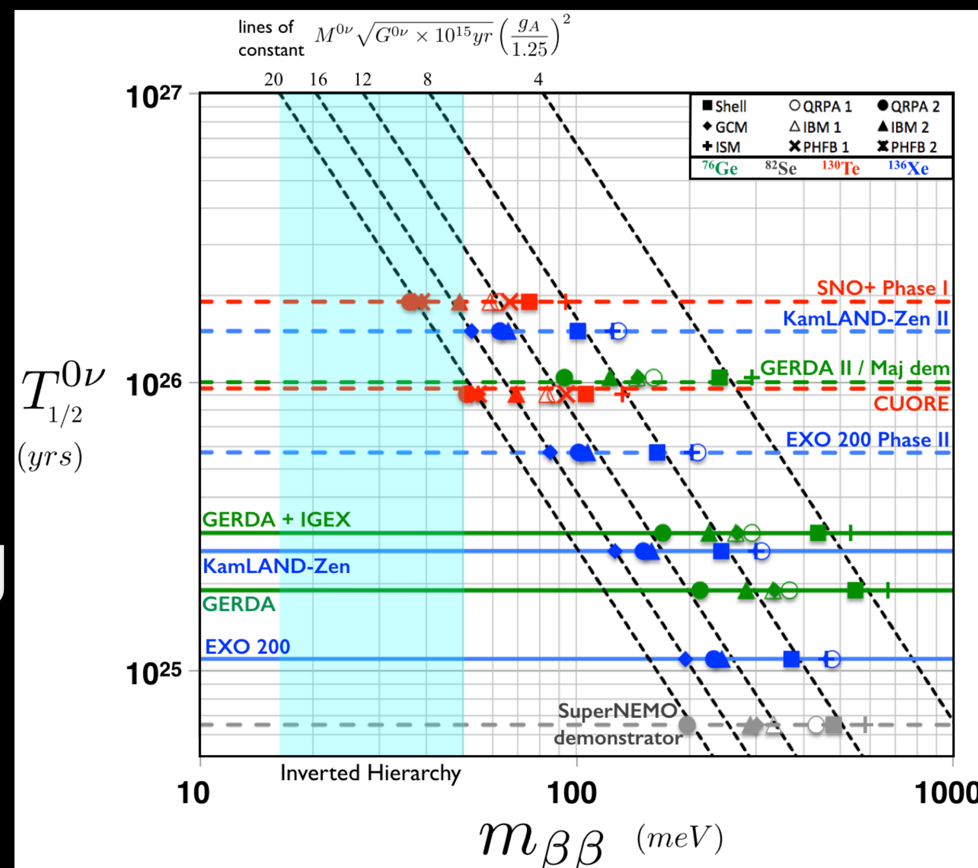
Light emitted from the support structure from 92 fibres installed between PMTs. Each gives 10E3-10E5 photons/pulse.

0νββ Isotope Selection

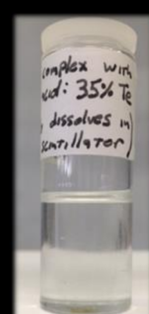


ovββ Sensitivity in Phase II

- Improve sensitivity by improving
  - Light yield and going to higher loading
    - Improve current technique
  - Higher QE PMTs
    - Improved concentrators
      - Coverage to 80%

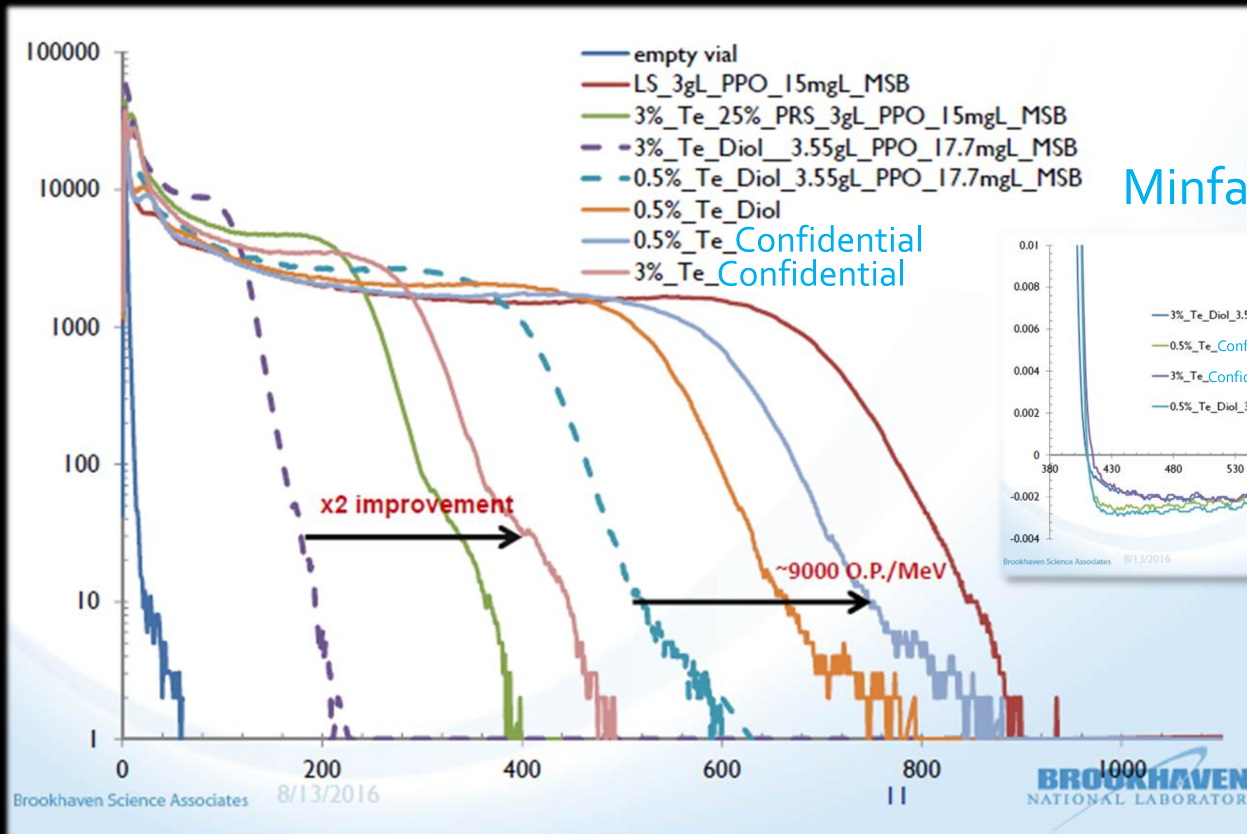


- Goal: 3% nat. Te loading
  - ~ 8 tonnes ^{130}Te
  - Higher QE PMTS
  - $T_{1/2}$ 0nbb ~ 10^{27} yr



ovββ Phase III R&D

- 2x the Light Yield and same absorption with alternative approach at 3%Te



Courtesy of Minfang Yeh of BNL