



UKAEA FISPACT-II applications: material activation, damage and waste

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United Kingdom Atomic Energy Authority

FISPACT-II workshop

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[Grant number EP/P012450/1]



Applications

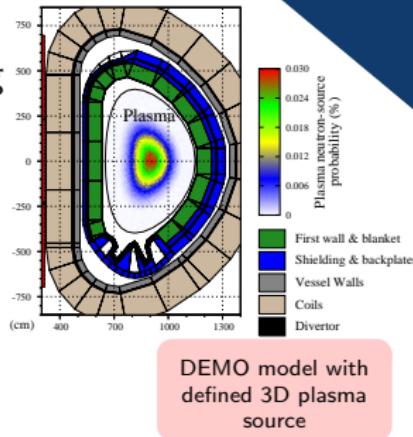
- Integrated assessment of helium embrittlement
- DEMO fusion power plant activation and waste analysis
- Material response database
- Primary damage spectra (& emitted particle spectra)

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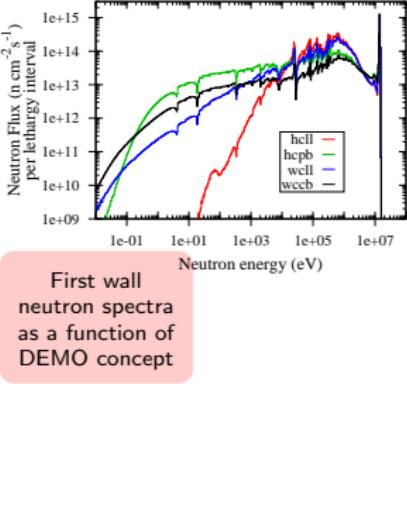
Integrated assessment: Outline

- Neutron transport modelling results for DEMO (MCNP)
- Neutron-induced transmutation (FISPACT-II)
 - ▶ helium production
- Helium embrittlement model and critical lifetimes

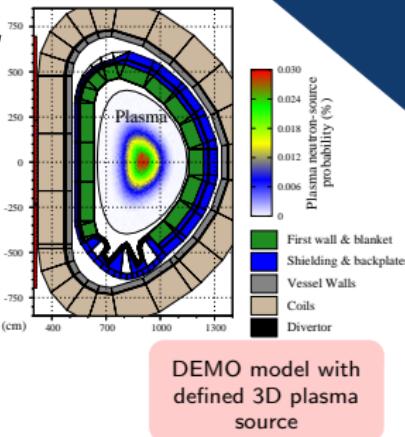


Gilbert, Dudarev, Nguyen-Manh, Zheng, Packer, Sublet
J. Nucl. Mater. **442** (2013) S755
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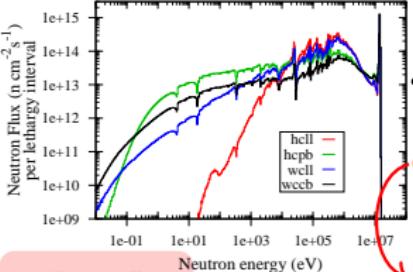


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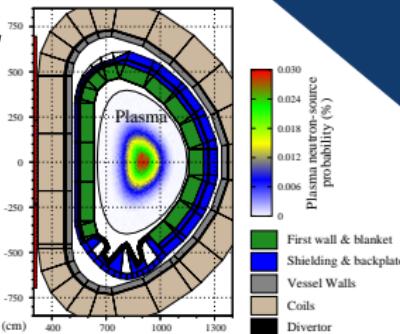
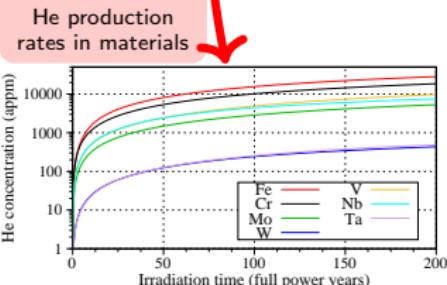


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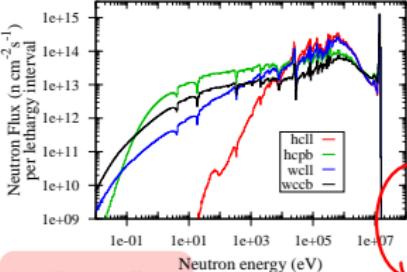
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DEMO model with defined 3D plasma source

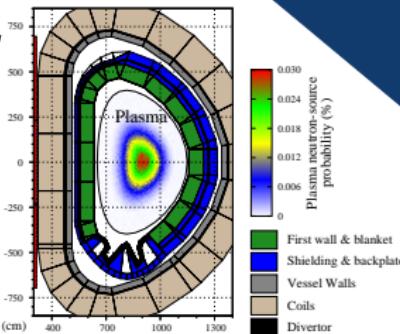
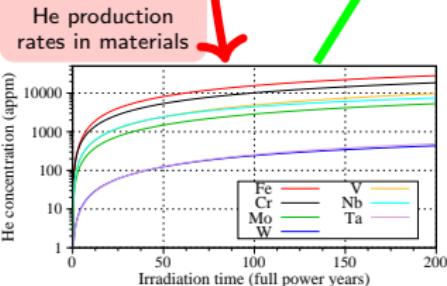
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Integrated assessment: Outline



First wall
neutron spectra
as a function of
DEMO concept

- Neutron transport modelling results for DEMO (MCNP)
- Neutron-induced transmutation (FISPACT-II)
 - ▶ helium production
- Helium embrittlement model and critical lifetimes

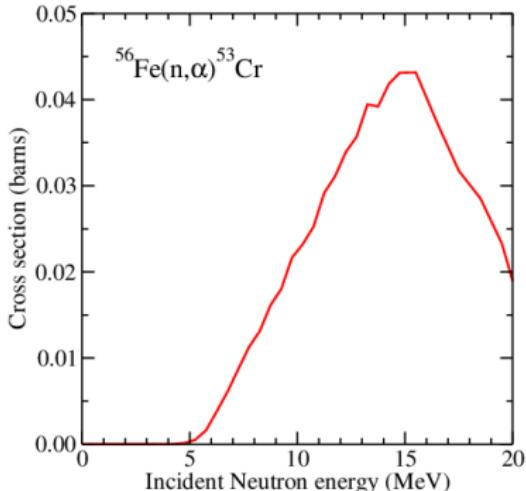


DEM0 model with
defined 3D plasma
source

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Helium production

- Helium production is a particular problem for fusion because of the generally higher neutron energies – many of the He producing reactions have thresholds
- Material comparison under identical DEMO hcpb conditions
- 3 fpy under outboard equatorial FW armour irradiation:

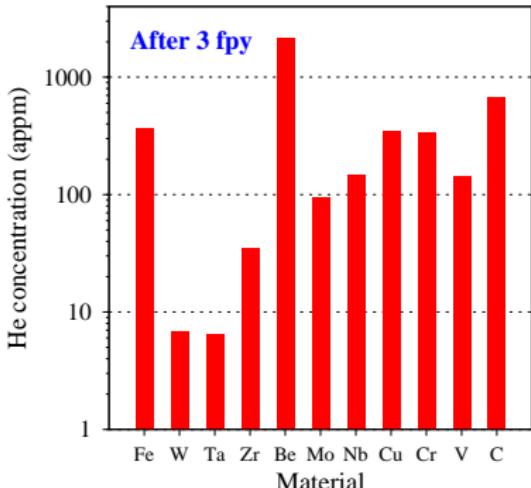


- FISPACT-II inventory calculations:
- Helium production highest in Be (~ 710 appm/fpy)
- More than an order of magnitude lower in Fe (~ 120 appm/fpy)
- Only ~ 2 appm/fpy in W

appm=atomic parts per million; fpy - full power years

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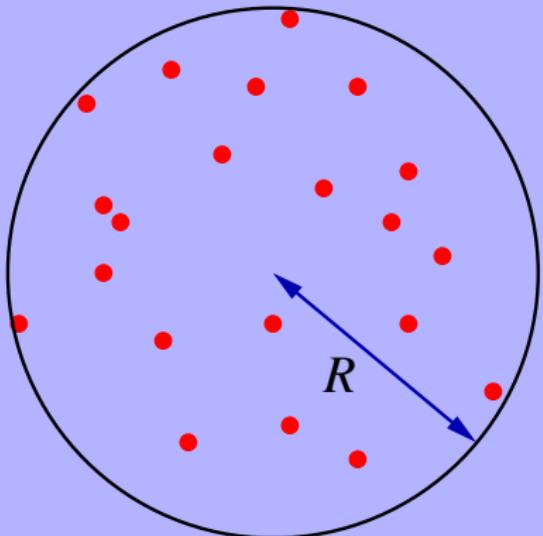
appm=atomic parts per million; fpy - full power years

Modelling: He embrittlement of GBs

Simple modelling of grain boundary (GB) failure

1. Number of He atoms in spherical grain:

$$N_{\text{He}} \approx \frac{4}{3}\pi R^3 n G_{\text{He}}$$



Assumptions:

- All helium atoms produced migrate to grain boundary
 - ▶ traps neglected
 - ▶ most valid for small grains

G_{He} = bulk concentration
(time evolution from inventory calcs.)

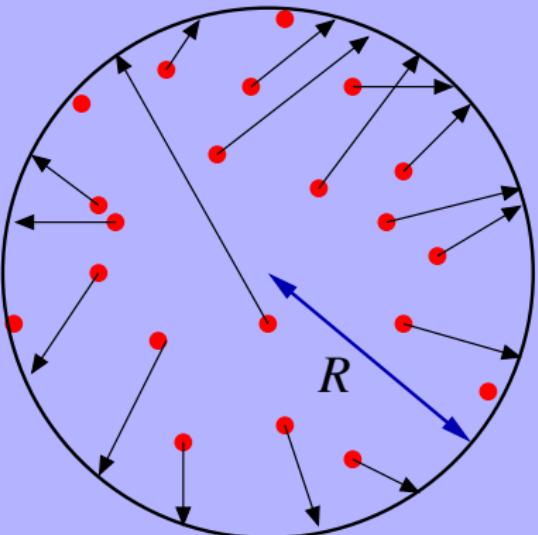
n = atom density

Modelling: He embrittlement of GBs

Simple modelling of grain boundary (GB) failure

2. All He atoms move to GB: – ∴ surface total \equiv bulk total

$$4\pi R^2 \nu_{\text{He}} = \frac{4}{3}\pi R^3 n G_{\text{He}} \Rightarrow \nu_{\text{He}} = \frac{R}{3} n G_{\text{He}}$$



Assumptions:

- All helium atoms produced migrate to grain boundary
 - ▶ traps neglected
 - ▶ most valid for small grains

G_{He} = bulk concentration
(time evolution from inventory calcs.)

ν_{He} = surface density

Modelling: He embrittlement of GBs

Simple modelling of grain boundary (GB) failure

3. GB destabilize when E of inserted He equals E of surfaces:

$$E_{\text{He}}^{\text{insrt}} \nu_{\text{He}}^c \approx 2\varepsilon_{\text{surf}}$$

Material	He insert. energy (eV) [†] – $E_{\text{He}}^{\text{insrt}}$	Surf. energy (Jm ⁻²) [*] – $\varepsilon_{\text{surf}}$	Critical He conc. at GBs (cm ⁻²) – ν_{He}^c
Fe	2.77	2.4	1.08×10^{15}
Cr	2.68	2.3	1.07×10^{15}
Mo	1.91	3.0	1.96×10^{15}
Nb	1.60	2.7	2.11×10^{15}
W	1.61	3.5	2.71×10^{15}
V	2.30	2.6	1.41×10^{15}

* Averages of experimental data reported in: L. Vitos *et al.*, 1998, *Surf. Sci.*, **411** 186–202

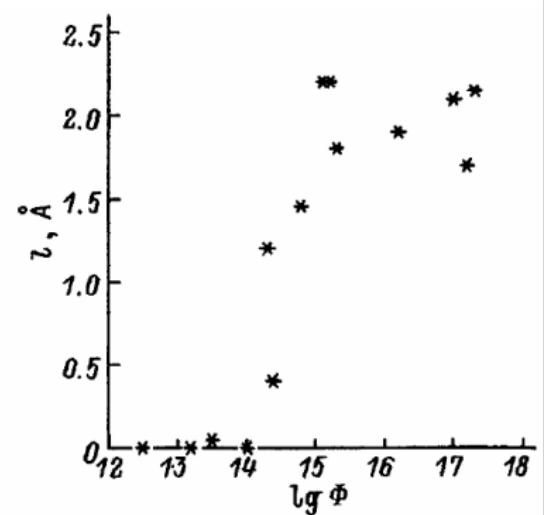
† DFT values – Gilbert *et al.* *J. Nucl. Mater.* **442** (2013) S755

Modelling: He embrittlement of GBs

Simple modelling of grain boundary (GB) failure

3. GB destabilize when E of inserted He equals E of surfaces:

$$E_{\text{He}}^{\text{insert}} \nu_{\text{He}}^c \approx 2\varepsilon_{\text{surf}}$$



- Experimental confirmation:
 - ▶ Helium irradiated W bicrystals
 - ▶ Expansion of grain boundaries at He fluence of 10^{14} – 10^{15} ions cm^{-2}
 - ▶ our ν_{He}^c value: 2.71×10^{15}

Gerasimenko, Mikhaĭlovskiĭ,
Neklyudov, Parkhomenko, and
Velikodnaya

Tech. Phys. **43** (1998) 803

Modelling: He embrittlement of GBs

Simple modelling of grain boundary (GB) failure

4. Critical bulk He concentration:

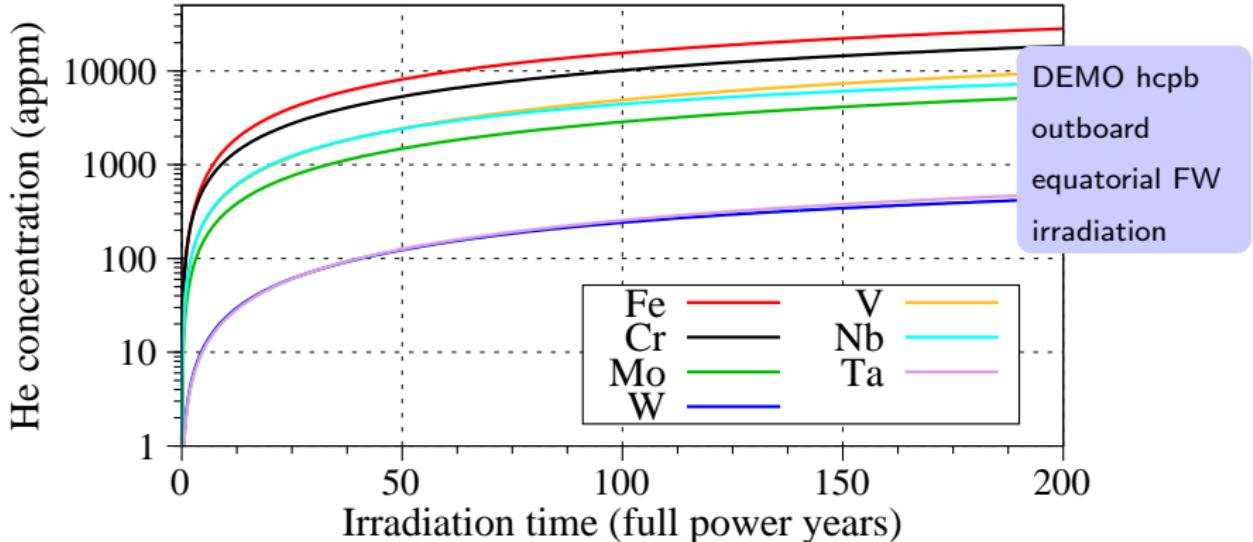
$$G_{\text{He}}^c = \frac{3}{Rn} \nu_{\text{He}}^c$$

Material	ν_{He}^c (cm^{-2})	n (cm^{-3})	G_{He}^c (appm)
Fe	1.08×10^{15}	8.5×10^{22}	764.6
Cr	1.07×10^{15}	6.4×10^{22}	771.9
Mo	1.96×10^{15}	5.5×10^{22}	1833.8
Nb	2.11×10^{15}	6.3×10^{22}	2275.2
W	2.71×10^{15}	1.2×10^{23}	2582.1
V	1.41×10^{15}	4.7×10^{22}	1172.2

- Assumed Grain size of $R = 0.5\mu\text{m}$
- G_{He}^c varies with $1/R$ appm – atomic parts per million

FISPACT-II results: helium production rates

- standard output during irradiation in FISPACT-II simulation
- Estimated “time to destabilize” t_{He}^c based on inventory calculations



appm = atomic parts per million

He embrittlement: Critical times

- t_{He}^c for irradiations in outboard equatorial DEMO FW armour position assuming linear grain size of $0.5 \mu\text{m}$

Element	ν_{He}^c (cm^{-2})	G_{He}^c (appm)	t_{He}^c (fpy)			
			hcll	hcpb	wcll	wccb
Fe	1.08×10^{15}	764.6	9	6	8	7
Cr	1.07×10^{15}	771.9	7	7	7	7
Mo	1.96×10^{15}	1833.8	58	63	59	62
Nb	2.11×10^{15}	2275.2	45	47	46	46
W	2.71×10^{15}	2582.1	700+	597	700+	700+
V	1.41×10^{15}	1172.2	25	25	25	25

- For some elements, such as W, the He production rates are so low, and/or the ν_{He}^c is so great that the critical lifetimes are many hundreds of years
- only for $t_{\text{He}}^c \lesssim 10$ is this failure mechanism likely to be of concern

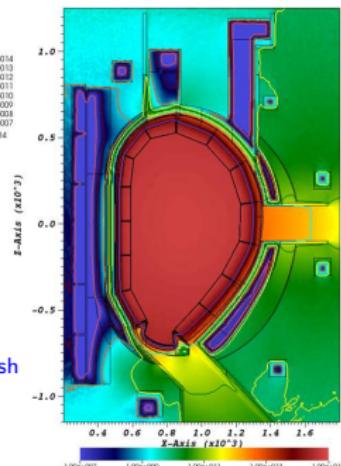
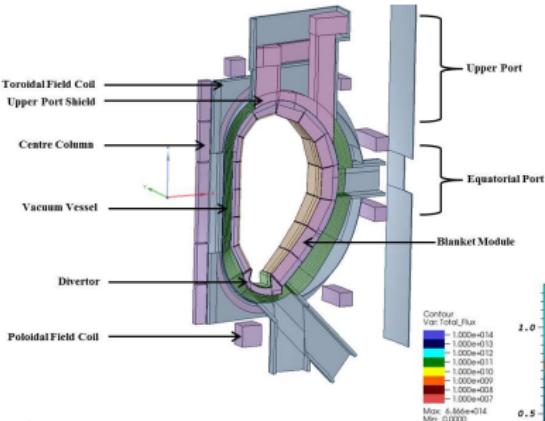
appm = atomic parts per million, fpy = full power year

Applications

- Integrated assessment of helium embrittlement
- DEMO fusion power plant activation and waste analysis
- Material response database
- Primary damage spectra (& emitted particle spectra)

Application: DEMO activation & waste predictions

- (European) designs for a demonstration fusion power plant (DEMO) are still conceptual
- But inventory calculations have a role to play in guiding choices relating to
 - ▶ shielding requirements
 - ▶ maintenance schedules
 - ▶ radioactive waste generation
 - ▶ (active) cooling needs
- FISPACT-II is well-suited to such analyses
 - ▶ easily script-able for 1000s of simulations
 - ▶ fast (using binary data files)
 - ▶ standardized output formats for easy post-processing



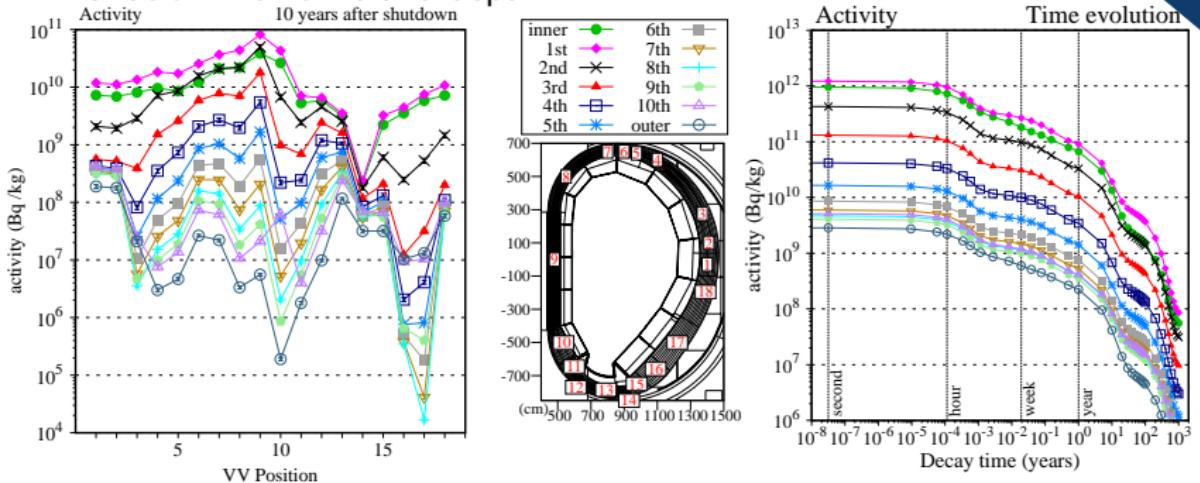
Input neutron flux mesh

from Monte Carlo

- Material composition (& hence activity) of each cell evolved in time according to a DEMO operational scenario
 - ▶ 22-year, 2-phase plan
 - ▶ one blanket replacement – 1-year shutdown
 - ▶ 3 divertor replacements – 8-month shutdowns

FISPACT-II automation

- e.g. for the vacuum vessel (VV) at end-of-life
- variation with time and depth in VV



- Output used to define waste (& recycling) class of component as a function of time

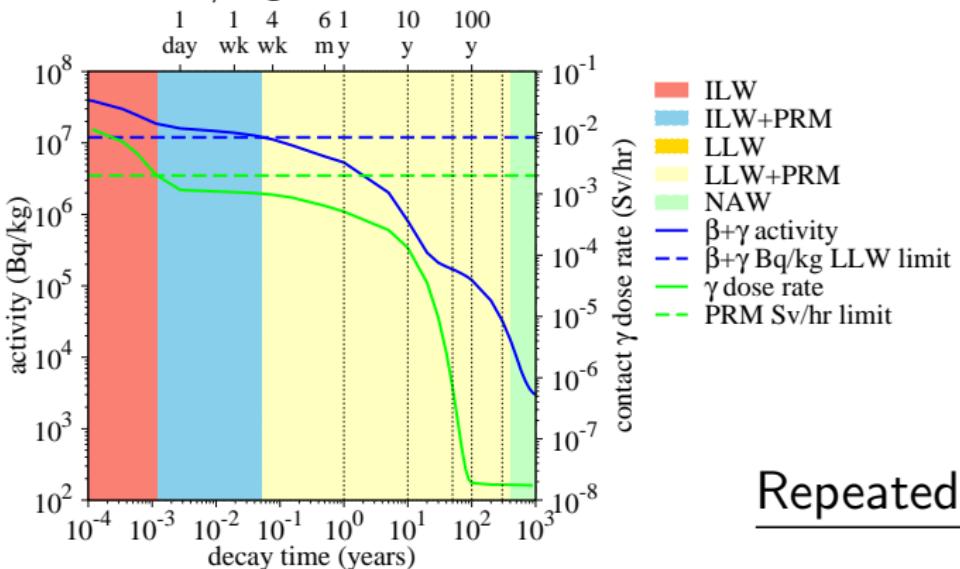
Waste classification

- Preliminary waste classes based on IAEA structure and UK limits*:
 - 1) **NAW** – (non-active waste)
 - IAEA clearance index less than 1
 - 2) **LLW** – (low-level waste)
 - α activity less than 4 MBq kg^{-1} and combined β and γ activity of less than 12 MBq kg^{-1}
 - 3) **ILW** – (intermediate-level waste)
 - activities above LLW limits [high-level waste assumed to be zero]
- Recycling assessment:
 - ▶ component considered as being Potentially Recyclable Material (PRM) if contact γ dose is below 2 mSv hr^{-1}
 - (contact dose is a standard result, but also a conservative overestimate)
- replaced components (i.e. blankets and divertor) included in waste assessment
- 100% tritium removal assumed for blanket components

\S HLW (high-level waste) assumed to be zero

Typical evolution

- For a cell/region in the outboard Vacuum Vessel

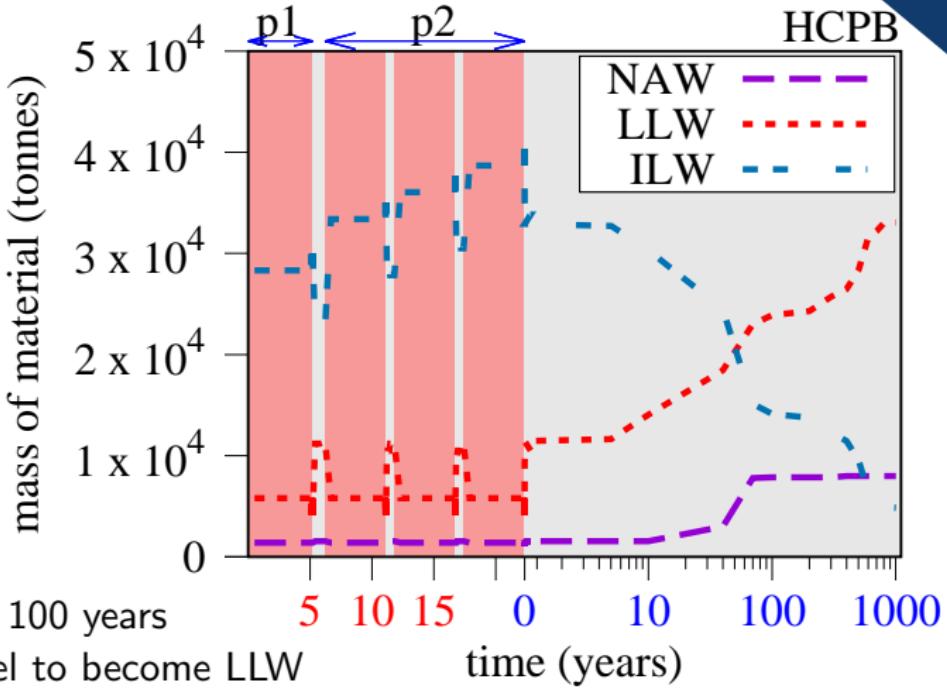


Repeated for all cells

- Never exceeds α -activity limit for LLW (true in general in fusion)
- Satisfies the $\beta + \gamma$ -activity LLW limit after 20 days of decay cooling
- Meets the PRM limit sooner (after ~ 10 hours)

Total reactor waste results

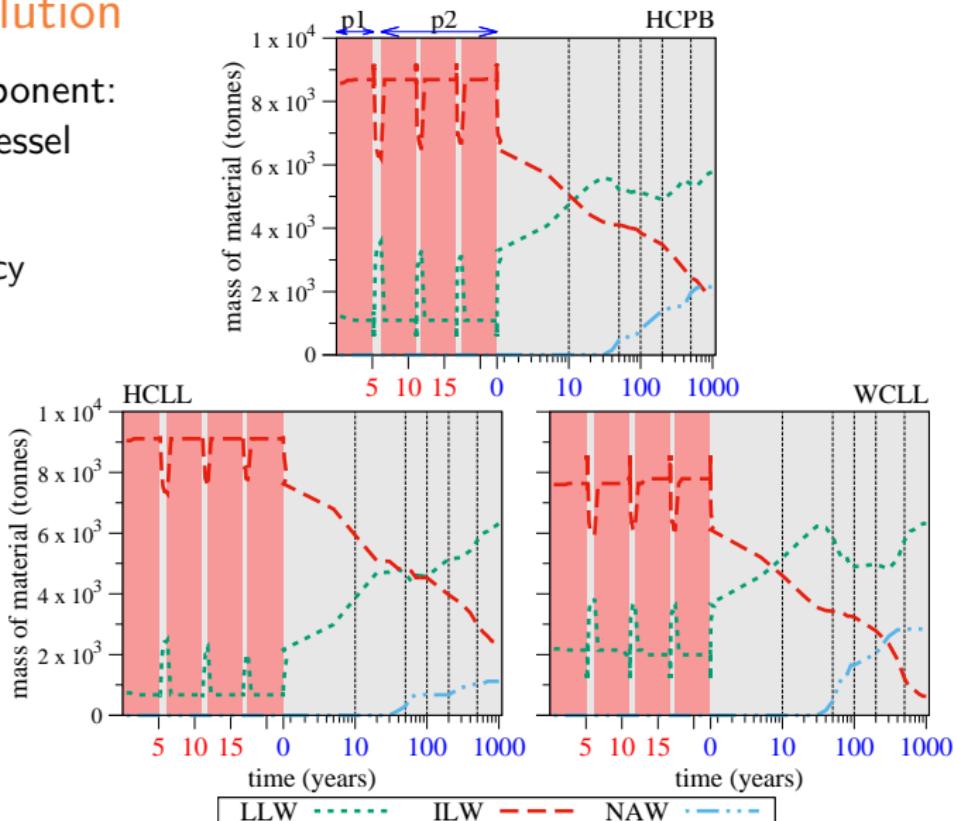
- jumps in waste masses due to new components
- mainly ILW in first few decades after shutdown
- & very little NAW at any time
- takes more than 100 years for most of vessel to become LLW (dominated by blanket module masses)



blue time labels are post final shutdown;
red during operational life;
red shading is for periods of irradiation;
grey shading during shutdown.

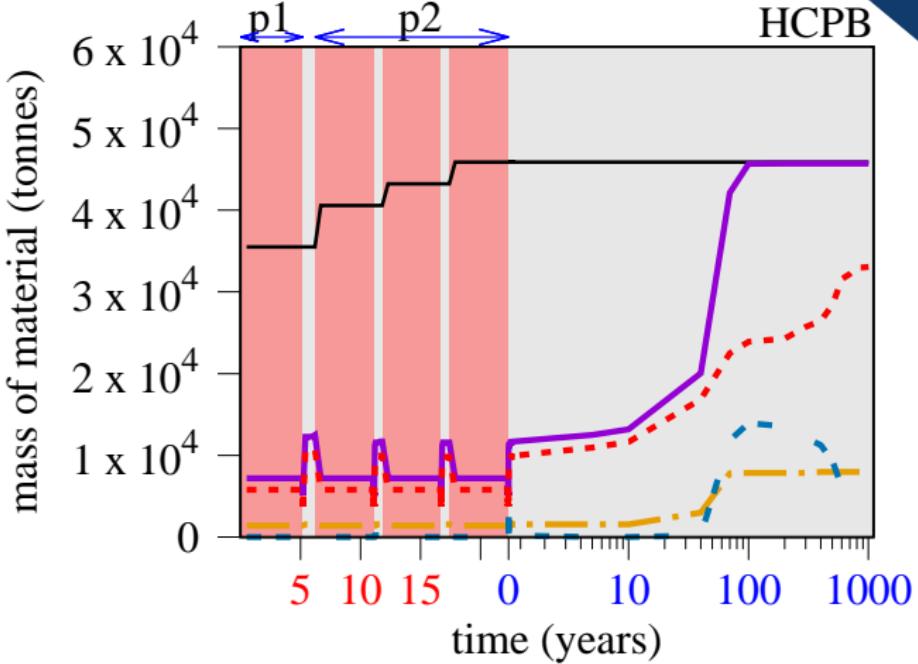
Waste Evolution

- For whole component:
e.g. Vacuum Vessel
- slightly lower shielding efficiency in HCLL design
 - no water/Be moderation
- ⇒ higher neutron fluxes in VV
- ⇒ later cross-over to majority LLW



Total reactor recycling results

- entire design (including VV+IVCs) becomes potentially recyclable within 100 years despite waste classification



total	—	RM-LLW	- - -
RM	—	RM-ILW	- - - -
RM-NAW	- - -		

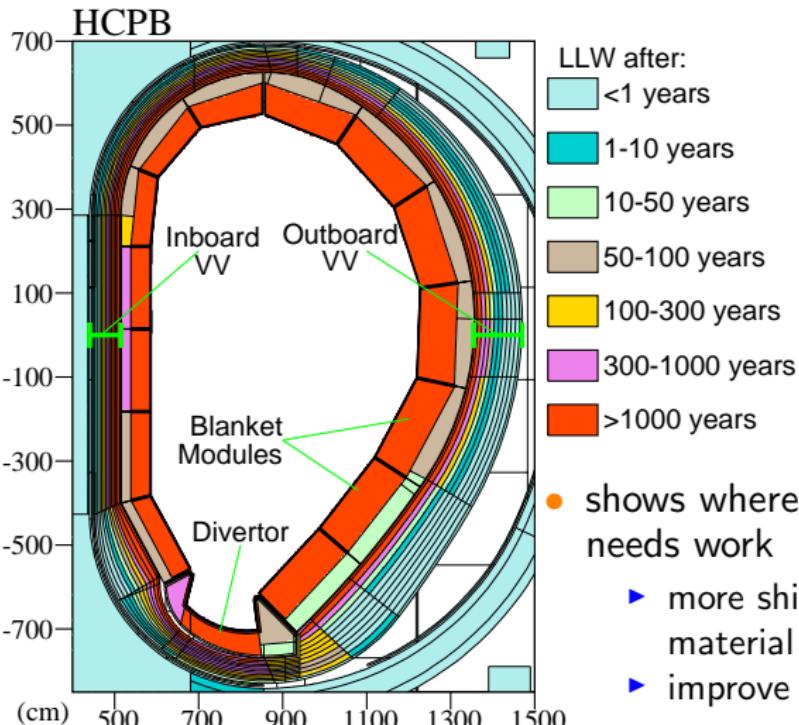
IVCs – in-vessel components

Time variation in class - entire DEMO reactor

- including pie-chart showing waste distribution)

Time to LLW plot

- re-processing of results to predict disposal criteria time-limits



- shows where design and modelling still needs work
 - more shielding, change to starting material compositions ...
 - improve detail (resolution) in model

Applications

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- DEMO fusion power plant activation and waste analysis
- **Material response database**
- Primary damage spectra (& emitted particle spectra)

Application: material response database



CCFE-R(16)36
August 2016

Mark R. Gilbert
Jean-Christophe Sublet

**Handbook of activation,
transmutation, and radiation damage
properties of the elements simulated
using FISPACT-II & TENDL-2015;
Magnetic Fusion Plants**

785 pages
~60000 calculations

www.ccfc.ac.uk

- Reference document of typical activity and burn-up responses of the elements to neutron irradiation is a useful tool during material selection exercises for nuclear components
- Results for all naturally occurring elements from H to Bi, with separate reports for:
 - ▶ predicted DEMO and ITER conditions
 - ▶ fission conditions (PWR, FBR, HFR)
- Available to download from
<http://fispact.ukaea.uk>

Scoping calculations for each element:

① Tabulated activation response

- ▶ % contributions of important radionuclides to various radiological quantities as a function of cooling time after irradiation

② Graphs of activation response after irradiation

- ▶ decay evolution of total activity, heat, and γ -dose under three different spectra & compared to Fe + indicative dominant nuclide contributions
- ▶ nuclide contributions as a function of time

③ Importance diagrams

- ▶ spectrum independent mapping of important radionuclides in the neutron-energy vs. decay time phase-space

④ Transmutation response

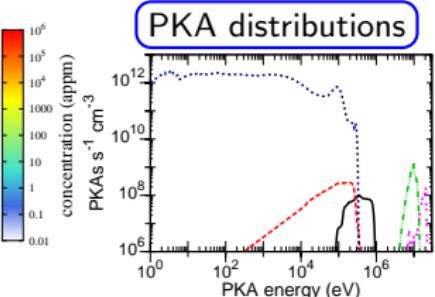
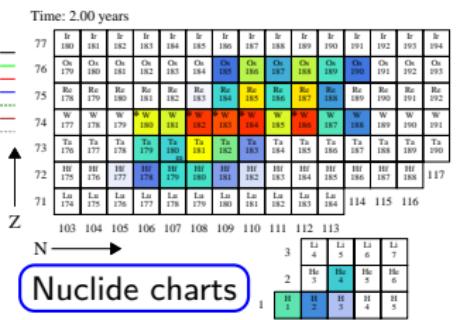
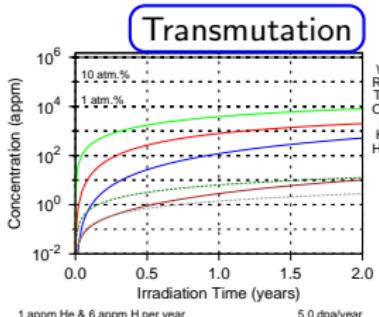
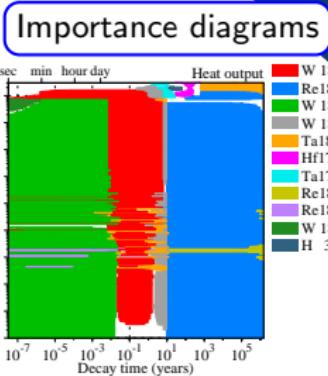
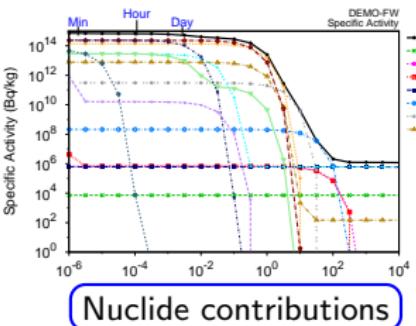
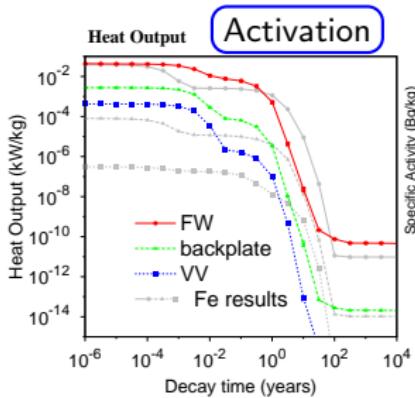
- ▶ time evolution under irradiation of initially-pure elemental composition
- ▶ nuclide concentration maps

⑤ Primary knock-on atom (PKA) distributions

- ▶ $t_{irr} = 0$ spectra plotted as both elemental and isotopic sums

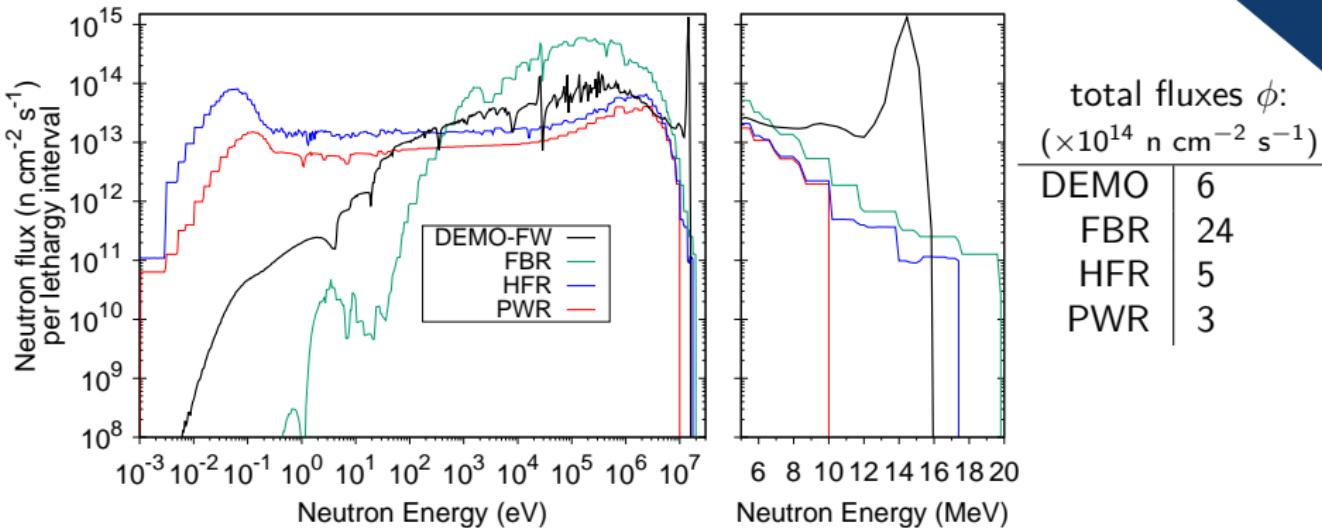
⑥ Reaction pathways

- ▶ major production pathways for important radionuclides at four characteristic neutron-energy ranges



Neutron irradiation fields

- Monte Carlo simulations of a fusion DEMOnstration power plant & typical fission reactors

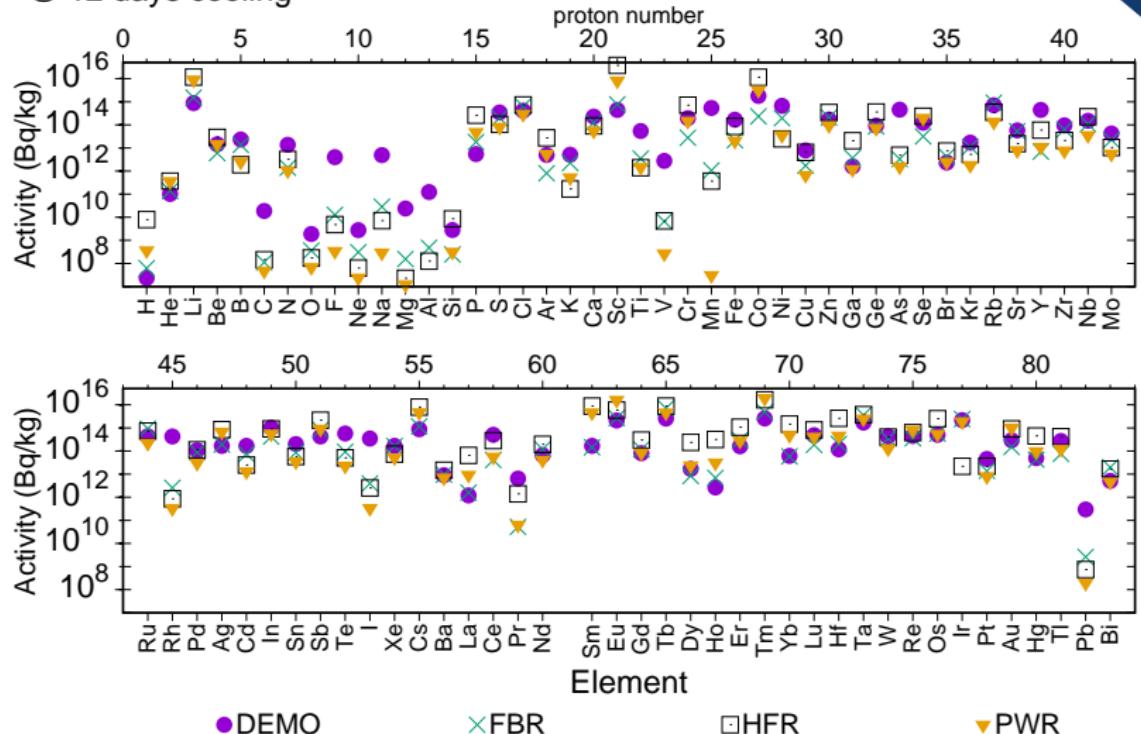


- fusion spectrum in first wall (FW) dominated by 14 MeV peak
- well-moderated (averaged) fission spectra don't have such dominant peaks but can have tails that explore the 14 MeV region of fusion

FBR – superphenix Fast Breeder Reactor
 HFR – High Flux Reactor, Petten
 PWR – Pressurized Water-cooled Reactor

Activation – fusion vs. fission

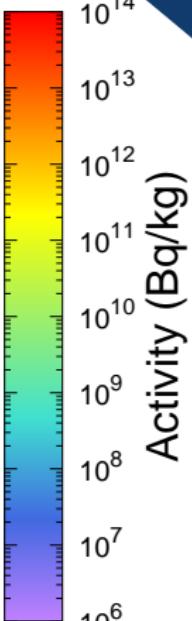
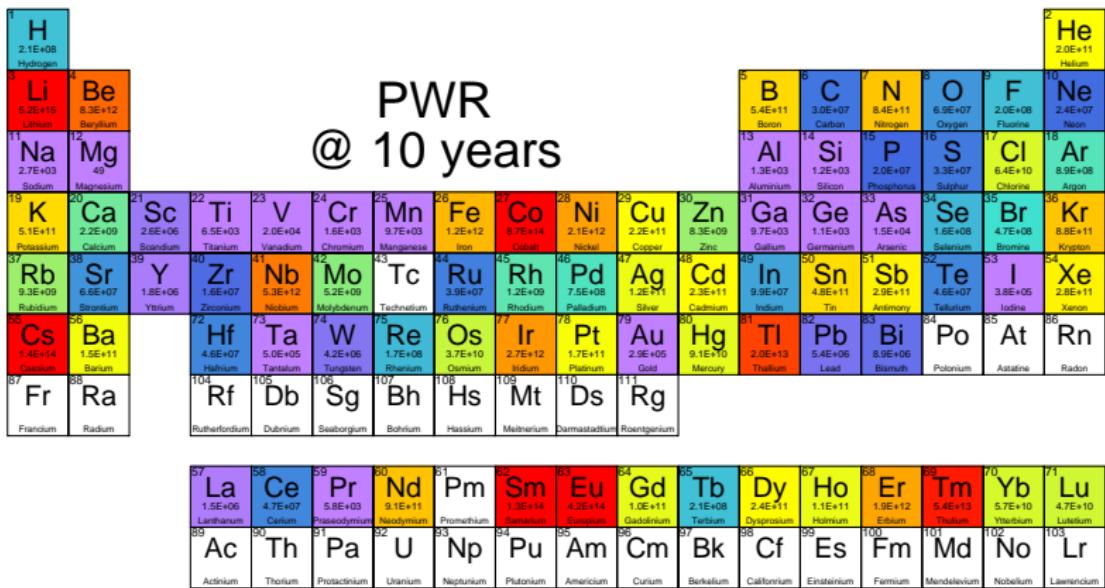
- time snapshot of activity during decay cooling[§]:
 @ 12 days cooling



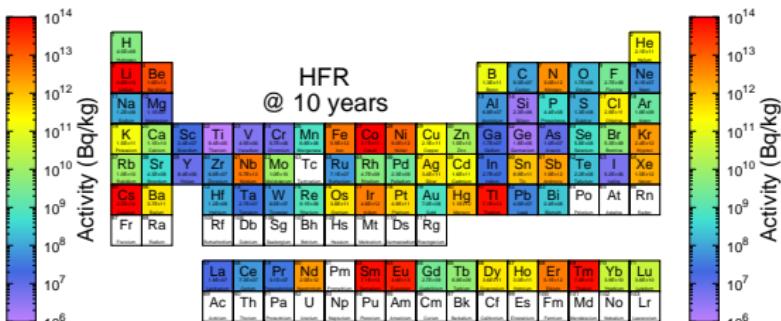
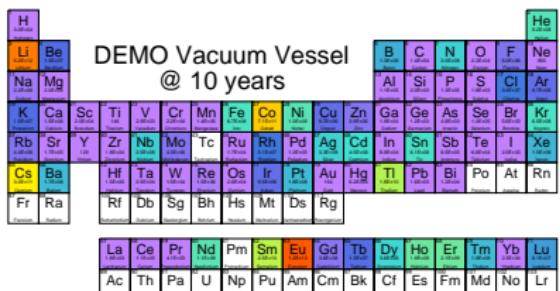
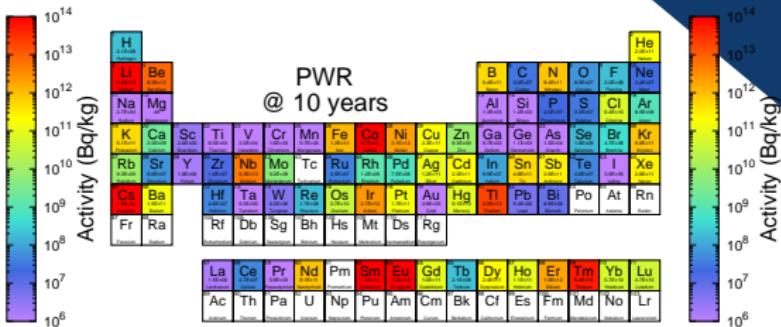
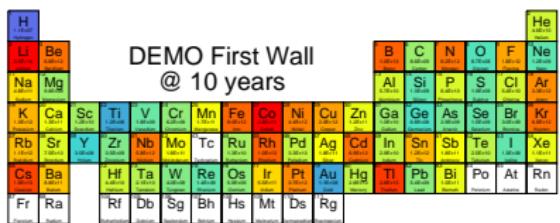
[§]after 2 years of continuous irradiation

Activation summary – periodic table plot

- total becquerel/kg activity from each element at 10-years following a 2-year PWR irradiation:



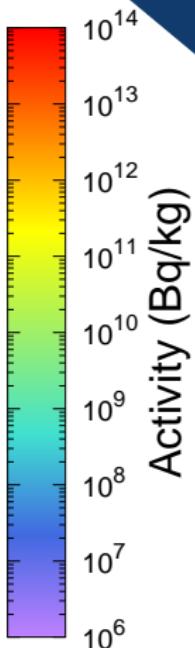
Activation summary – periodic table



Activation summary – animated

- total becquerel/kg activity from each element as a function of time following 2-year irradiation

**DEMO First Wall
@ 3.2E-08 years**



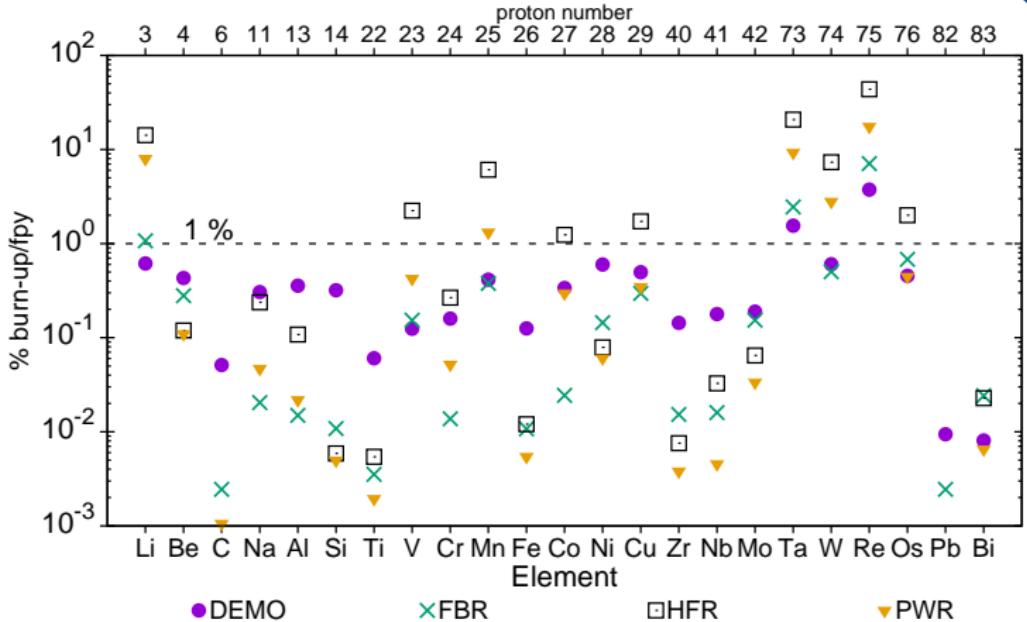
1 H Hydrogen $1.9\text{E}+07$	2 He Helium $8.2\text{E}+10$
3 Li Lithium $8.5\text{E}+14$	4 Be Beryllium $1.2\text{E}+14$
5 Na Sodium $3.5\text{E}+14$	6 Mg Magnesium $3.2\text{E}+14$
7 K Potassium $2.0\text{E}+13$	8 Ca Calcium $2.3\text{E}+14$
9 Rb Rubidium $1.1\text{E}+15$	10 Sr Strontium $2.0\text{E}+14$
11 Cs Cesium $1.1\text{E}+15$	12 Ba Barium $2.0\text{E}+14$
13 Fr Francium $1.1\text{E}+15$	14 Sc Scandium $9.0\text{E}+14$
15 Hf Hafnium $7.4\text{E}+14$	16 Ti Titanium $2.0\text{E}+14$
17 Ta Tantalum $2.0\text{E}+14$	18 V Vanadium $2.0\text{E}+14$
19 Cr Chromium $2.0\text{E}+14$	20 Mn Manganese $1.4\text{E}+15$
21 K Potassium $2.0\text{E}+13$	22 Fe Iron $2.3\text{E}+14$
23 Zr Zirconium $4.0\text{E}+14$	24 Co Cobalt $4.0\text{E}+14$
25 Nb Niobium $5.3\text{E}+14$	26 Ni Nickel $7.5\text{E}+14$
27 Tc Technetium $3.2\text{E}+14$	28 Cu Copper $6.2\text{E}+14$
29 Ru Ruthenium $6.0\text{E}+14$	30 Zn Zinc $3.4\text{E}+14$
31 Rh Rhodium $3.2\text{E}+15$	32 Ga Gallium $1.3\text{E}+15$
33 Pd Palladium $1.1\text{E}+15$	34 Ge Germanium $3.8\text{E}+14$
35 Ag Silver $3.6\text{E}+15$	36 As Arsenic $5.8\text{E}+14$
37 Cd Cadmium $4.7\text{E}+14$	38 Se Selenium $3.4\text{E}+15$
39 In Indium $7.2\text{E}+15$	39 Br Bromine $3.8\text{E}+14$
41 Sn Tin $2.8\text{E}+14$	40 Kr Krypton $3.2\text{E}+15$
43 Tl Thallium $1.7\text{E}+15$	41 I Iodine $7.3\text{E}+14$
45 Pb Lead $8.7\text{E}+13$	42 Xe Xenon $4.4\text{E}+14$
47 Bi Bismuth $7.3\text{E}+12$	43 Po Polonium $8.4\text{E}+12$
49 At Astatine $2.2\text{E}+15$	44 Rn Radon $8.6\text{E}+11$
51 La Lanthanum $6.2\text{E}+13$	52 Ce Cerium $6.8\text{E}+14$
53 Pr Praseodymium $7.2\text{E}+14$	54 Nd Neodymium $4.1\text{E}+14$
55 Pm Promethium $1.0\text{E}+15$	56 Sm Samarium $3.8\text{E}+15$
57 Eu Europium $3.9\text{E}+14$	58 Gd Gadolinium $3.9\text{E}+14$
59 Tb Terbium $2.8\text{E}+15$	60 Dy Dysprosium $3.2\text{E}+14$
61 Ho Holmium $3.6\text{E}+15$	62 Er Erbium $8.1\text{E}+14$
63 Tm Thulium $2.2\text{E}+15$	64 Yb Ytterbium $2.1\text{E}+14$
65 Lu Lutetium $2.4\text{E}+15$	66 Md Mendelevium $1.0\text{E}+15$
67 Ac Actinium $1.0\text{E}+15$	68 Th Thorium $1.2\text{E}+14$
69 Pa Protactinium $1.0\text{E}+14$	70 U Uranium $1.0\text{E}+15$
71 Np Neptunium $9.3\text{E}+14$	72 Pu Plutonium $1.0\text{E}+15$
73 Am Americium $9.4\text{E}+14$	74 Cm Curium $1.0\text{E}+15$
75 Bk Berkelium $9.7\text{E}+14$	76 Cf Californium $1.0\text{E}+15$
77 Es Einsteinium $9.9\text{E}+14$	78 Fm Fermium $1.0\text{E}+15$
79 Md Mendelevium $1.0\text{E}+15$	80 No Nobelium $1.0\text{E}+15$
81 Lr Lawrencium $1.0\text{E}+15$	

Activation summary – animated

- total becquerel/kg activity from each element as a function of time following 2-year irradiation

Transmutation – fusion vs. fission

- % burn-up per full-power year (fpy) for selected elements:

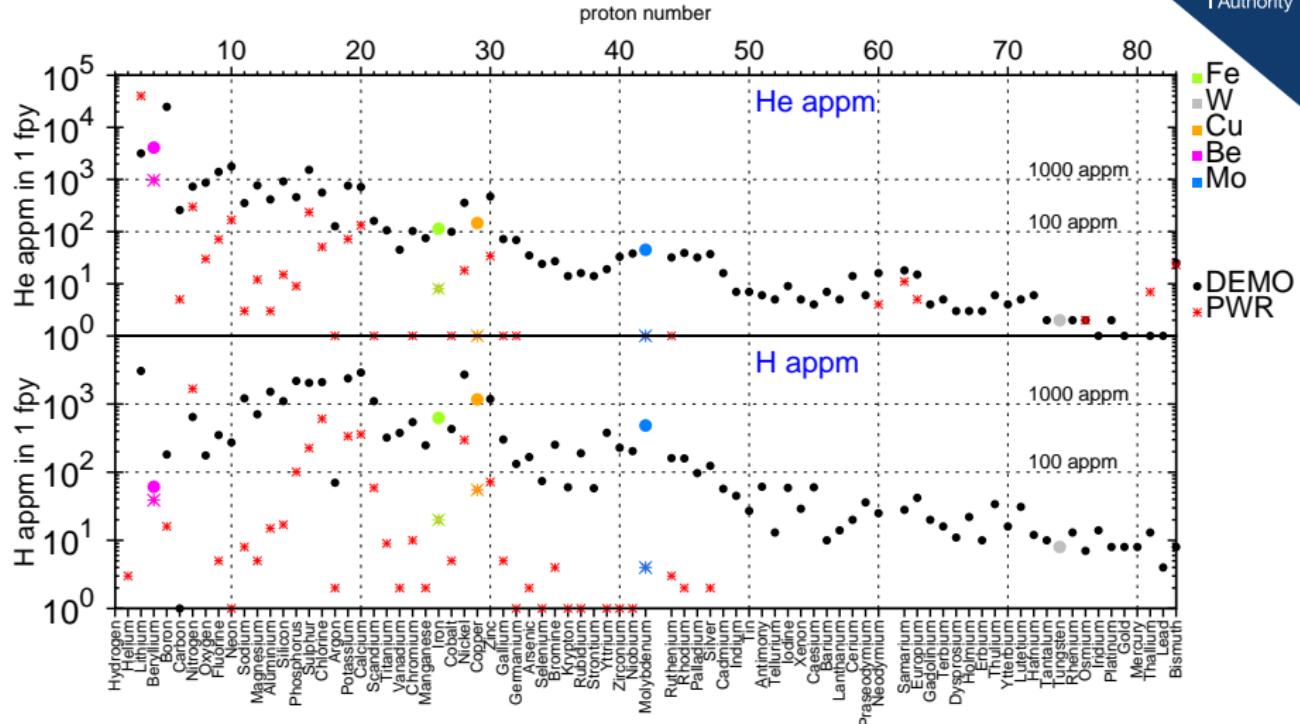


- Fission spectra are more-moderated ("softer") than fusion & combined with high-flux in HFR produces greater burn-up rates in many elements[§] ⇒ spectrum modification (via shielding) required to match fusion in experimental campaigns

[§]due to higher neutron capture rates

Gas production – fusion vs. fission

- Gas appm after 1 fpy in DEMO first wall and PWR:



- gas production is significantly higher under fusion neutrons

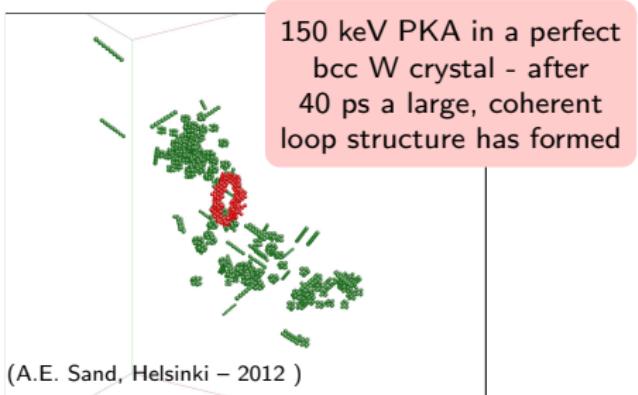


Applications

- Integrated assessment of helium embrittlement
- DEMO fusion power plant activation and waste analysis
- Material response database
- Primary damage spectra (& emitted particle spectra)

Application: PKA distributions

- Motivation: a full description of the initial damage events (the primary knock-on atoms or PKAs) under irradiation is a necessary input to modelling and simulation of radiation damage
 - PKA energy.vs.flux distributions provide more information than simpler damage-dose measures (e.g. dpa)
- Complete evaluation requires all possible nuclear reaction channels to be considered:
 - elastic & inelastic scattering, and non-elastic reactions ((n, p) , $(n, 2n)$, etc.)
 - including secondary emitted (light) gas particles - α (${}^4\text{He}$), protons (${}^1\text{H}$), etc, as PKAs



PKA – primary knock-on atom

Additional need

- emitted particle spectra for α , protons, neutrons, γ , etc.
- could be used in FISPACT-II to compute transmutation from secondary particles

PKA – primary knock-on atom



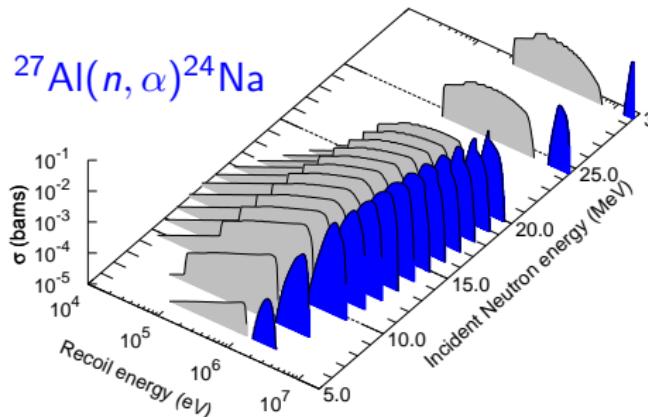
Calculation method (1)

- From raw pointwise nuclear data (TENDL)
- Neutron interaction recoil matrices $M^{x \rightarrow y} \equiv \{m_{ij}^{x \rightarrow y}\}$ calculated using NJOY[§] (via GROUPR)
 - ▶ $m_{ij}^{x \rightarrow y}$ is the recoil cross section (in barns) for a recoil energy E_i of daughter y resulting from an incident neutron energy E_j on parent x
- for each target (parent) isotope x there will be a set of $M^{x \rightarrow y}$
 - ▶ (at least) one for each reaction channel
- & each reaction may have more than one associated matrix
 - ▶ for example, (n,α) will produce a heavy recoil and an α -particle (${}^4\text{He}$ nucleus)
- recoil matrices can also be produced for charged particle interactions

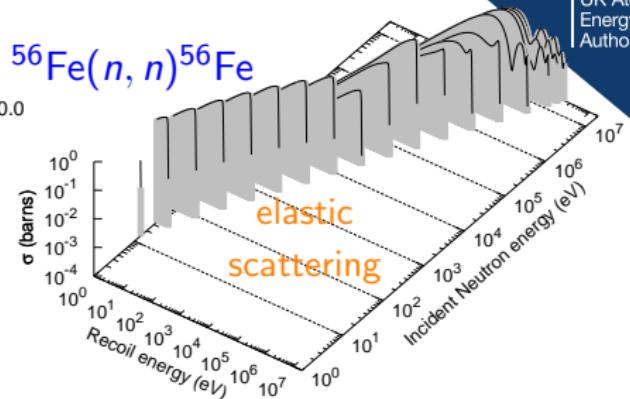
[§]the nuclear data processing system developed at LANL

$M^{x \rightarrow y}$ matrices & vectors

$^{27}\text{Al}(n, \alpha)^{24}\text{Na}$



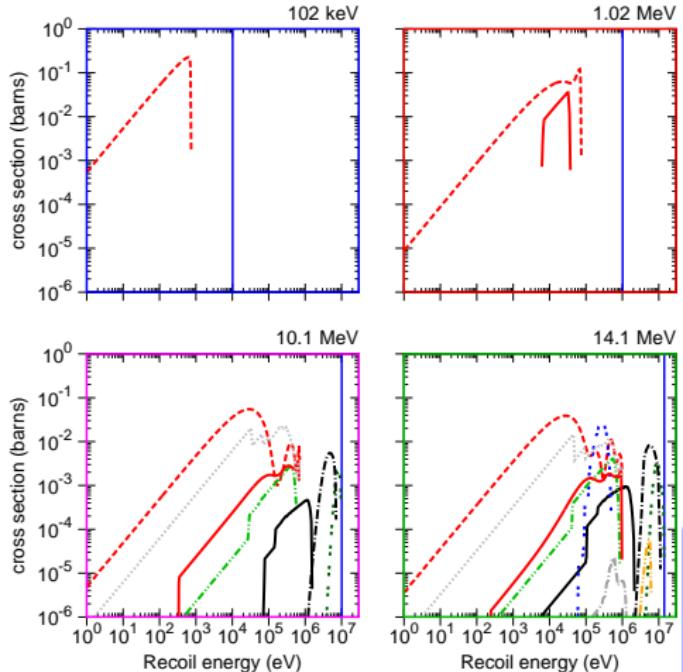
$^{56}\text{Fe}(n, n)^{56}\text{Fe}$



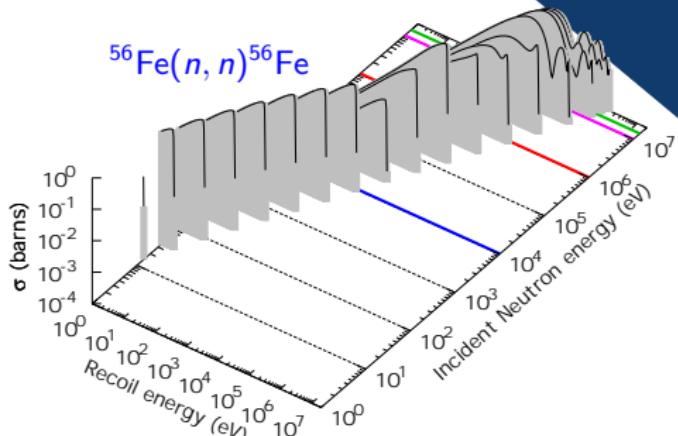
- Non-elastic nuclear reactions can produce two recoiling species
- The main (heavy) residual and a higher energy light (gas) particle
- On ^{27}Al the threshold (n, α) reaction produces ^{24}Na (grey) recoils and ^4He particles (blue)

- The primary reaction channel is normally simple elastic scattering
- Produces recoils for all incident neutron energies
- Other channels only become important at higher energy...

$M^{x \rightarrow y}$ matrices & vectors – ^{56}Fe



reaction channels	
—	$(n,\alpha)^{53}\text{Cr}$
- - -	$(n,\alpha)^{52}\text{Cr}$
- · -	$(n,n)^{56}\text{Fe}$
- · -	$(n,n')^{56}\text{Fe}$
- · -	$(n,p)^{1}\text{H}$
- · -	$(n,\alpha)^4\text{He}$
- · -	$(n,\alpha)^{55}\text{Fe}$
- · -	$(n,p)^{56}\text{Mn}$
- · -	$(n,\alpha)^4\text{He}$



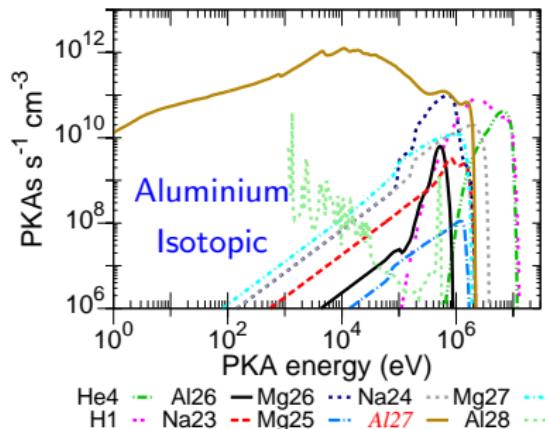
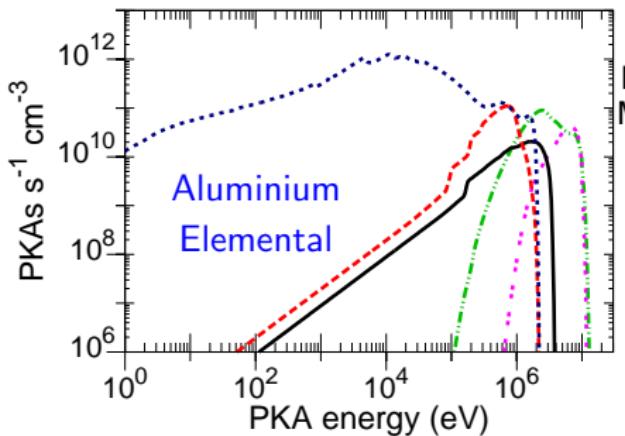
- In ^{56}Fe , at low incident energy only elastic and inelastic channels are open
- Number of contributing channels increases at higher energies (MeV range)

Calculation method (2)

- Collapsing each recoil matrix with a neutron irradiation spectrum $\{\phi_j\}$ gives the recoil-energy spectrum $R^{x \rightarrow y}(E)$:

“PKA-spectrum”
 under $\equiv R^{x \rightarrow y}(E) \equiv \{r_i^{x \rightarrow y}\} = \left\{ \sum_j m_{ij}^{x \rightarrow y} \phi_j \right\}$
 neutron irradiation

- processing (& summing) done with SPECTRA-PKA[§] utility

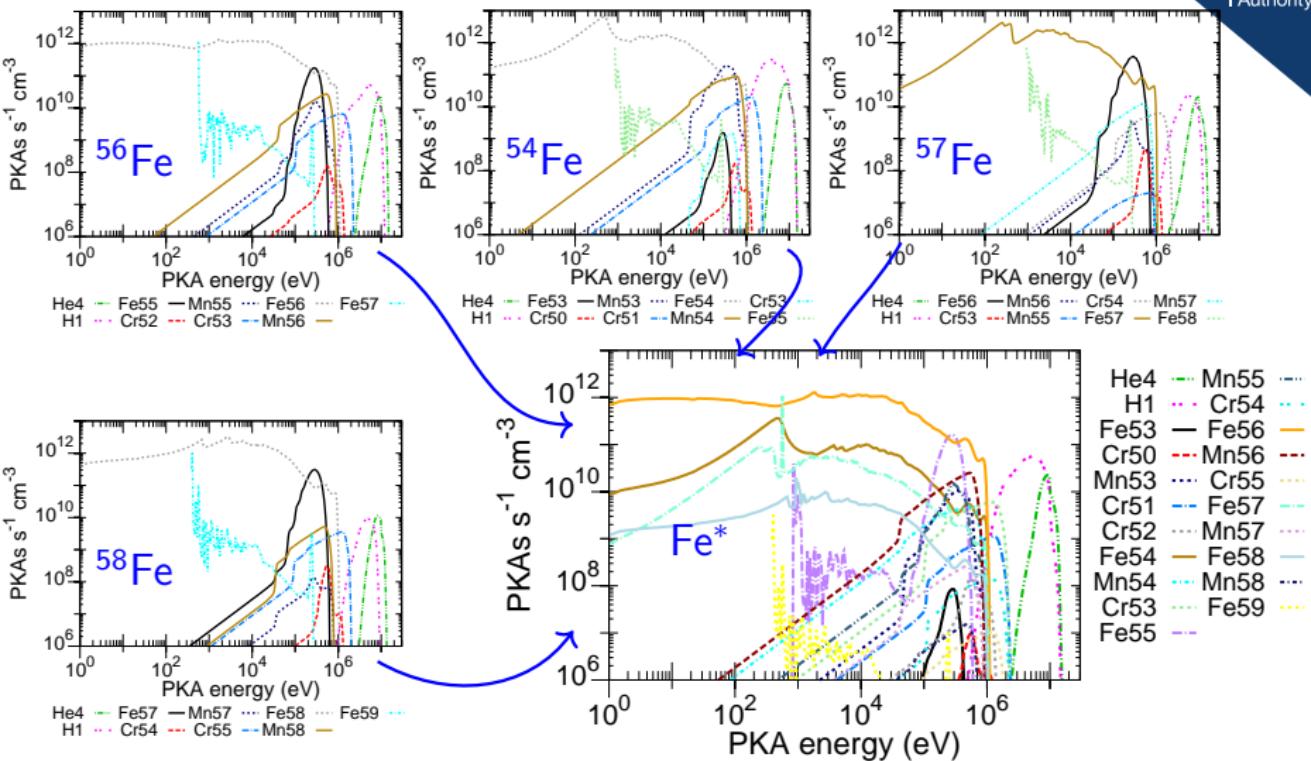


- PKA spectra of pure Al under DEMO conditions
- many different recoil species (even with just Al27 target)

[§]Gilbert et al., JNM 467 (2015) 121-134

PKA results – e.g. Fe

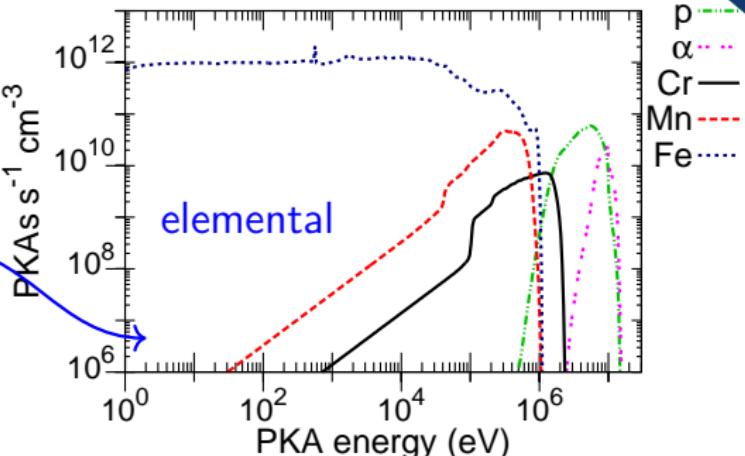
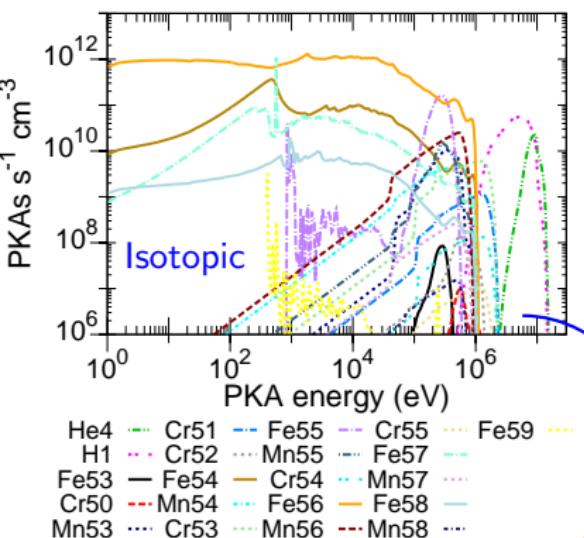
- Isotopic results for each nuclide in the material must be merged:



*5.845% ^{54}Fe , 91.754% ^{56}Fe , 2.119% ^{57}Fe ,
 0.282% ^{58}Fe (atomic percentages)

PKA results – e.g. Fe

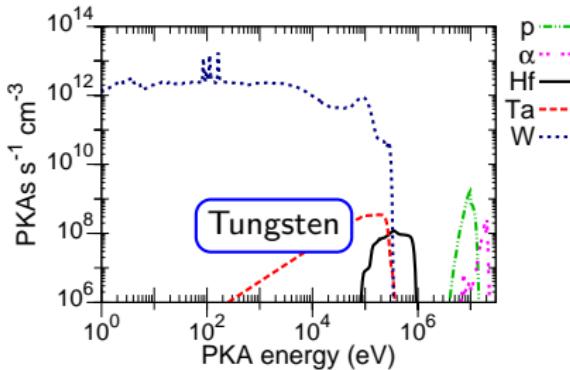
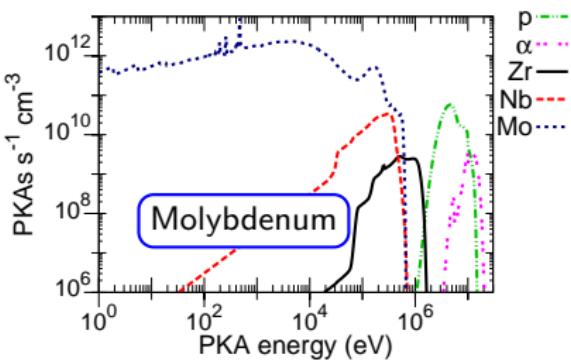
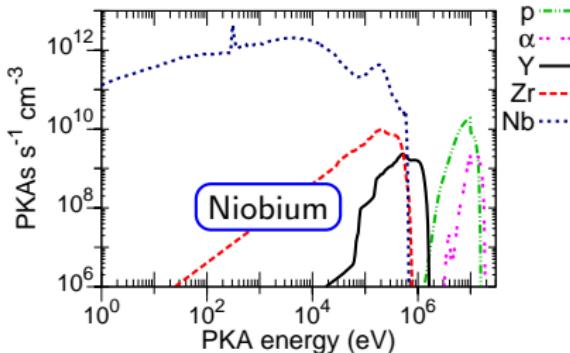
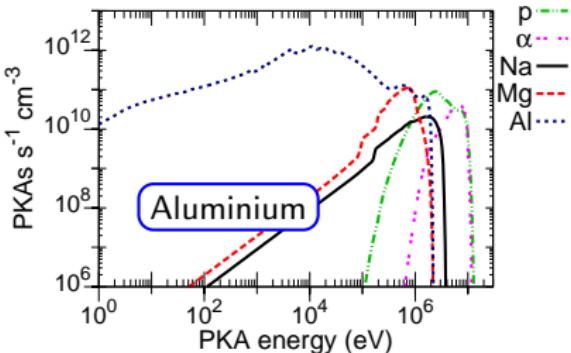
- For modelling it is more useful to sum the isotopic spectra:



- Main constituent usually dominates

- Total PKAs ($\text{s}^{-1}\text{cm}^{-3}$) in material can be calculated by summing (& integrating over) all elemental distributions
- here there are 4.33E+14 $\text{PKAs s}^{-1}\text{cm}^{-3}$ above 1eV (light particles excluded)
- the average PKA energy above 10 eV for the Fe+Mn+Cr spectrum is 18.8 keV

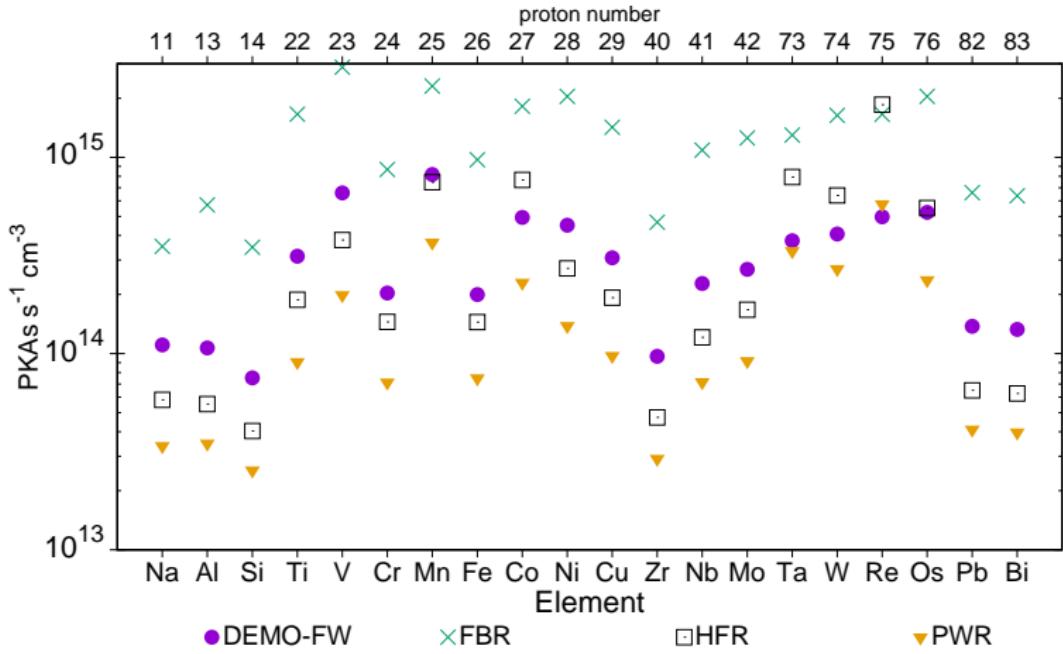
PKA distributions – elemental



- Under DEMO FW conditions

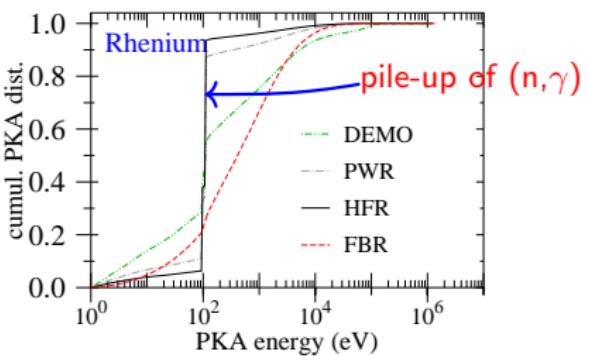
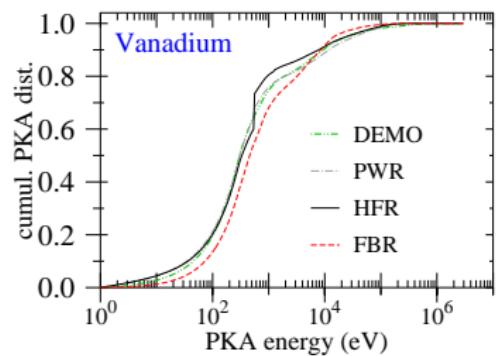
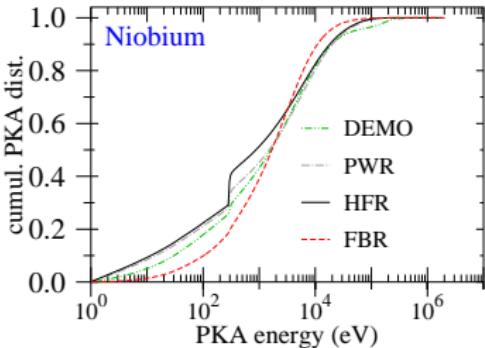
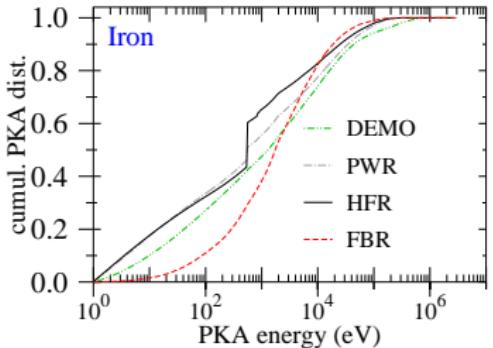
Total PKA rates – fusion vs. fission

- total heavy PKAs (H/He excluded) for selected materials



- FBR spectrum produces highest rates due to high, fast flux
- PWR the lowest

Cumulative (PKA) probability distributions



- FPR missing low and high energy parts due to peaked spectrum

Damage energy & dpa from SPECTRA-PKA

- The displacement energy in the NRT-dpa formula is normally calculated using the total damage kerma cross section
- However, with SPECTRA-PKA, it is possible to calculate the damage contribution as a function of reaction channel
 - a new & novel capability
- Standard LSS[§] formula to account for electronic loss and convert PKA energy into damage energy – including correct treatment of parent and daughter mass
- displacement-energy rate accumulated and summed using PKA rate at each damage energy. NRT formula can be applied to the total
- For a given reaction channel, the total displacement energy is:

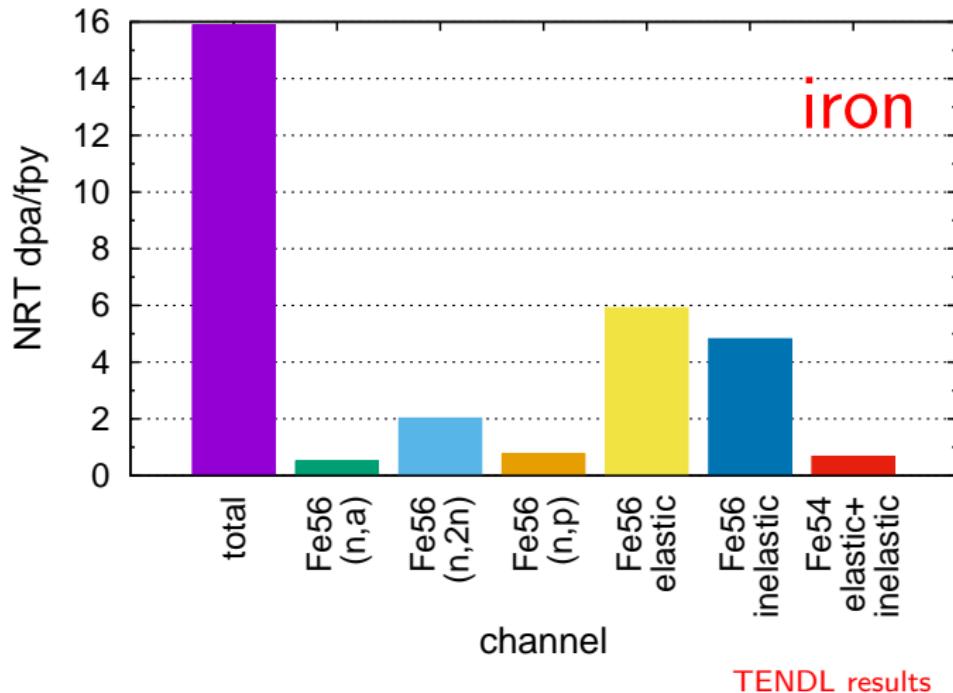
$$\sum_i T^{LSS}(E_i^{pka}) R_i^{pka},$$

where $T^{LSS}(E^{pka})$ is the LSS equivalent energy for PKA energy E^{pka} , and R_i^{pka} is number of PKAs at energy E_i^{pka}

[§]Lindhard, Scharff, Schiøtt, *Mat. Fys. Medd. Dan. Vid. Selsk.* 33 (1963) 1–42

Damage energy from SPECTRA-PKA

- Channel contributions to NRT dpa/fpy* for Fe[§] under DEMO FW conditions:



- Largest contributions are from scattering of ⁵⁶Fe (91.2 atm.%)
- ⁵⁶Fe(n,2n) also important (13%)
- total agrees with value obtained from standard approach – FISPACT-II & total damage kerma

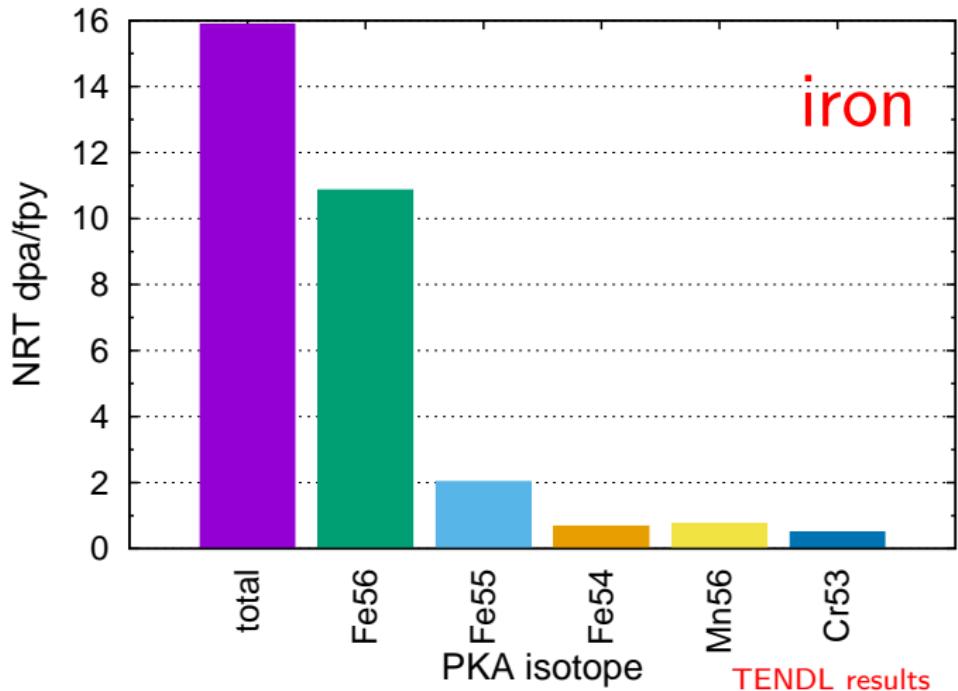
Gilbert and Sublet, JNM 504 (2018) 101-108

*fpy – full power years

$${}^{\S} E_d = 40 \text{ eV}$$

Damage energy from SPECTRA-PKA

- Nuclide contributions to NRT dpa/fpy* for Fe[§] under DEMO FW conditions:



- the per-channel dpas summed by daughter indicate the proportion of damage from different recoiling species
- dominated by Fe nuclides, but with a contribution from ⁵⁶Mn & ⁵³Cr

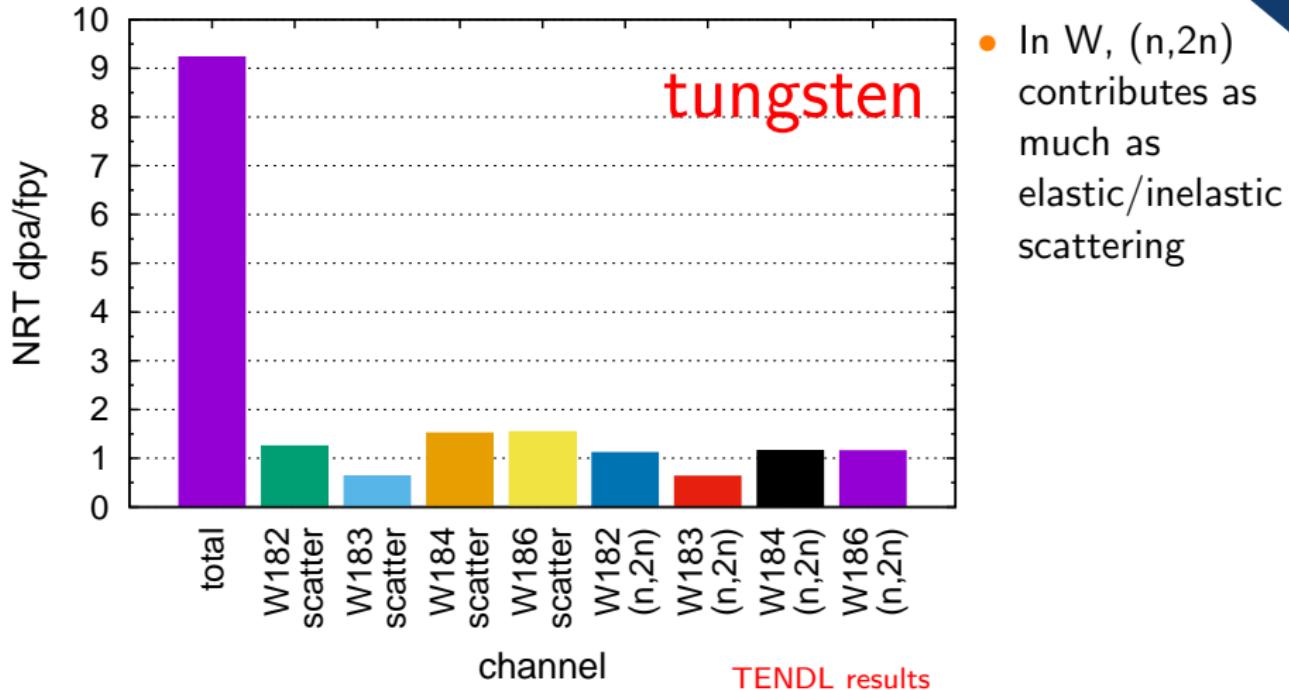
Gilbert and Sublet, JNM 504 (2018) 101-108

*fpy – full power years

[§] $E_d = 40 \text{ eV}$

Damage energy from SPECTRA-PKA

- Channel contributions to NRT dpa/fpy for W[§] under DEMO FW conditions:

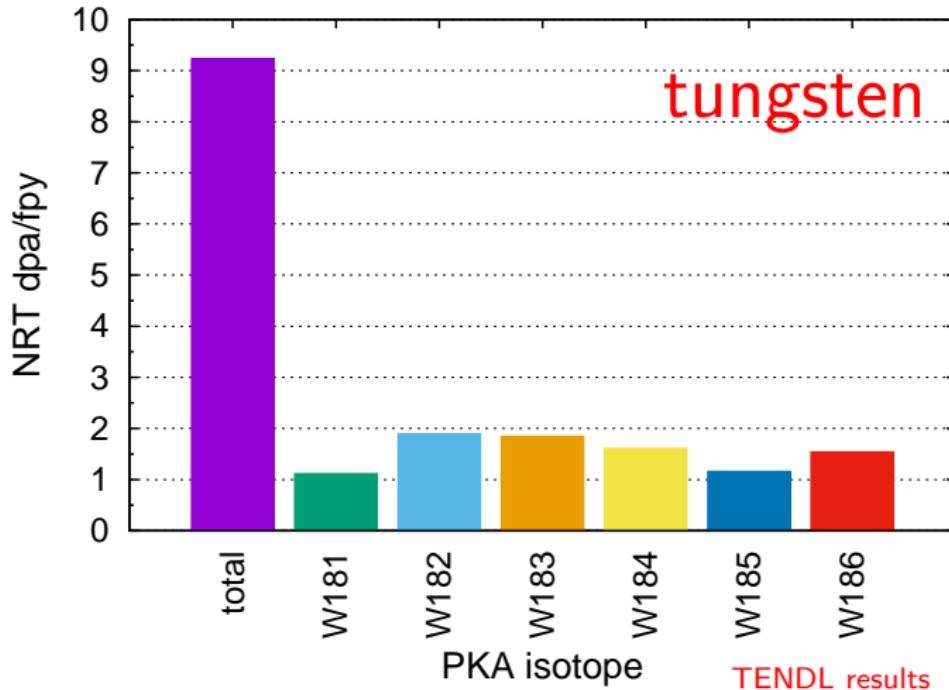


Gilbert and Sublet, JNM 504 (2018) 101-108

[§] $E_d = 55$ eV

Damage energy from SPECTRA-PKA

- Nuclide contributions to NRT dpa/fpy for W[§] under DEMO FW conditions:



- no significant contributions from reactions that produce isotopes of other elements

Gilbert and Sublet, JNM 504 (2018) 101-108

[§] $E_d = 55 \text{ eV}$

Running SPECTRA-PKA

- SPECTRA-PKA is now open-source and downloadable from:
<https://github.com/fispact/SPECTRA-PKA>
- code is command-line driven
- execution controlled by an input file of code words and values
(similar to FISPACT-II, but order does not matter)
`<location of SPECTRA-PKA>SPECTRA-PKA input.in`
- manual on the github explains use of all code-words & provide there is an example calculation...
 - ▶ including example plotting script to visualize results (with gnuplot)
- ...this is a good place to start
 - ▶ & example input can be easily modified to a specific material/spectrum of interest
 - ▶ if not already compiled on your system, the executable for SPECTRA-PKA can be easily created
 - navigate to `fispact/source/spectra-pka` and type `make`

Gilbert and Sublet, JNM 504 (2018) 101-108