

# Assessment of environmental consequences of the normal operations of the ESS facility

- status report -

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ARIA'17 / Lund, Sweden

## **Environment - Regulatory aspects**

**EUROPEAN SPALLATION** 

SOURCE

Swedish legal requirements : SSMFS2008:23, SSMFS2010:2 limited/no validity for ESS New Swedish regulations in 2018 Euratom BSS 2013, ICRP & IAEA recommendations SSM2015-127 Licensing conditions ESS GSO

Unified methodology under development: PREDO project

						Facility dos	e hudget to Reference person
Dose constraints	Linac	Target	Instruments	Waste facility	<b>X</b> = maior	No partition RELEASES <	ning of direct radiation vs releases < 40 $\mu$ Sv/y as stray radiation dose=
Emission into Air	AA 🗙	τα 🗙	IA X	WA 🗙	<b>x</b> = minor	<b>10μSv/y</b>	
Direct Radiation	АВ 🗙	тв 🗙	IB X	WB X	Contribution	Radiation P ESS Radiati	rotection Design Criteria on Safety Officer (RSO)=>
Activation Gr. water	AC X	TC <b>X</b> ?	IC _	WC _		Requiremen	nts (design, construction, operations)
Discharge into drains/evaporator	AD X	TD X	id 🗙	WD X		Ste	p#1 screening approach => E < 0.1 μSv/
					ESS Tota	al <sup>Step</sup>	p#2 realistic approach
Operation & maintenance	0.03	0.01	0.005	0.005	0.05 mSv/y	H1	=> 10 μSv/y
Expected events	0.05	0.35	0.05	0.05	0.1 mSv/eve	ent <b>H2</b>	SSM 2013/1525
Non-expected events	1.00	1.00	1.00	1.00	1 mSv/even	t <b>H3</b>	All isotopes
Design Basis Accidents (DBA)	20.00	20.00	20.00	20.00	20 mSv/eve	nt <b>H4</b>	giving contribution
Beyond Design Basis Accidents (BDBA)	50.00	50.00	50.00	50.00	50 mSv/eve	nt <b>H5</b>	$\blacktriangleright$ 1% to release

ALARA: Justification & Optimization + BAT Swedish Environmental Code (DS 2000:61)

## De minimis dose < 10 $\mu$ Sv/y => facility is deemed to be justified and optimized

Dose constraint is not a dose limit, but is used as the starting value for the optimization of radiation protection. (SSMFS:14-2480, 2014)

This means that the actual doses to members of the public, via the best possible technique and optimization of radiation protection are, for most cases, expected to be significantly lower



# Airborne releases general considerations



## GRADED APPROACH defining Source Term (ST):

1.**SCREENING**: conservative calculations (IAEA SRS19) for *selection of the radionuclides potentially radiation safety important* Breakdown\_of radionuclides -> calculate DF -> Quantification of ST (Bq) (calculations/estimates) -> Ranking & selection via screening analysis using screening dose rate 0.1 μSv/y per nuclide & summed dose rate of all radionuclides screened out < 1 μSv/y => i) important nuclides tbt via realistic analysis; ii) screened-out nuclides;

2.REALISTIC APPROACH (PREDO method) => realistic wind dispersion and realistic radioecological models.

#### Total Dose = $\Sigma$ realistic dose + $\Sigma$ dose of the radionuclides screened out

#### Threshold values:

Radioisotopes with  $T_{1/2} < 10$  sec will not be included in the analysis.

The traveling time from the stack until the closest potential Reference Group is about 2 min.

No unique threshold value of radioactivity= > Graded approach

#### **Criteria for selection of the radionuclides of the source terms**:

- are significant in terms of radiological impact;
- are significant in terms of quantity of radioactivity discharged, whether or not are significant for radiological impact;
- have long radioactive half-lives, that may persist and/or accumulate in the environment and that may contribute significantly to the dose.

#### Additional processes used for derivation of releases to the stack:

•Ventilation system rate (VR);

•"Filter" effectiveness (F) | "filter" is used here as generic for all devices able to: filter, absorb and/or delay the isotopes from their production to the stack. •Radioactive decay (I).

•No retention

#### **<u>Two cases of release rates were considered:</u>**

-pessimistic release, that assumes the maximum that the facility can potentially release (no credit to the filtration equipment at the main stack); -optimal release asuming that all abatement equipment works optimally

#### **Two main scenarios**:

**1. chronic long-term release** corresponding to a normal operation of the facility: i) uniform release during 50 years of the facility operation and ii) averaged weather conditions

**2.** short-term release : short-term planned interventions on the facility, when significant amounts of radioactivity may be released at once and the most adverse and unchanging weather conditions assumed.

# ESS site plan

ESS



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# Release points & locations of the reference groups

# Main Target Station f Stack High Bay f A2T High Bay f Monolith Roms f Monolith Roms f Mainela.ene@esss.ce d f

## **Airborne Release**

#### Two main scenarios:

1. chronic long-term release corresponding to a normal operation of the facility: i) uniform release during 50 years of the facility operation and ii) averaged weather conditions

**2. short-term release** : short-term planned interventions on the facility, when significant amounts of radioactivity may be released at once and the most adverse and unchanging weather conditions assumed.

# Atmospheric release Source term from Linac



stack On main

stack

yes

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	Activity to stack*	Screening ap	proach SRS19
Radionuclide	(Bq/y)	Dose factors	Annual effective dose**
	6.245.07	(SV/y per Bq/y)	(SV/y)
H-3	6.21E+07	1.73E-18	1.07E-10
Be-7	6.33E+05	2.09E-16	1.32E-10
Be-10	3.48E-02	6.29E-14	2.19E-15
C-11	4.09E+12	5.27E-19	2.16E-06
C-14	1.06E+07	1.24E-16	1.32E-09
N-13	7.70E+12	6.02E-19	4.63E-06
0-14	7.88E+11	1.86E-21	1.47E-09
0-15	8.29E+12	2.27E-19	1.88E-06
0-19	1.45E+09	1.32E-17	1.91E-08
F-18	8.53E+09	2.09E-18	1.79E-08
Ne-23	3.50E+09	9.36E-20	3.27E-10
Ne-24	7.12E+08	2.01E-23	1.43E-14
Na-22	9.21E+01	8.20E-14	7.56E-12
Na-24	2.11E+05	8.19E-17	1.73E-11
Na-25	1.85E+06	3.18E-19	5.87E-13
Mg-27	2.27E+06	3.81E-19	8.66E-13
Mg-28	3.00E+04	4.97E-16	1.49E-11
Al-26	4.56E-04	6.60E-12	3.01E-15
Al-28	9.93E+06	4.78E-19	4.74E-12
Al-29	4.62E+06	2.03E-18	9.38E-12
Si-31	1.25E+06	4.24E-19	5.29E-13
Si-32	6.09E-01	9.47E-13	5.77E-13
P-30	2.87E+06	2.84E-19	8.15E-13
P-32	9.30E+04	1.65E-15	1.53E-10
P-33	3.78E+04	2.61E-16	9.87E-12
P-35	3.63E+06	1.92E-19	6.95E-13
S-35	1.46E+04	2.28E-15	3.32E-11
S-37	1.14E+07	1.72E-20	1.97E-13
S-38	1.55E+06	4.00E-18	6.18E-12
Cl-34	3.73E+09	1.78E-18	6.64E-09
Cl-36	9.90E+01	7.32E-15	7.25E-13
Cl-38	1.30E+11	1.68E-18	2.19E-07
Cl-39	1.87E+11	1.94E-18	3.63E-07
CI-40	4.64E+10	2.08E-21	9.67E-11
Ar-39	4.48E+05	1.45E-21	6.49E-16
Ar-41	1.24E+12	7.59E-19	9.42E-07
K-38	6.89E+09	1.75E-18	1.21E-08
K-40	2.05E-03	2.13E-13	4.37E-16
Total	2.25E+13		1.03E-05
No filter effec	:t:		
Total	2.26E+13		1.16E-05



calculated to a hypothetical group located at 350 m from the release point.

By using realistic wind dispersion and realistic radio-ecological models it is expected the conservative screening dose result reported will be substantially lesser.  $^6$ 

1.8 - 1.9

12

2

Tunnel access

# ATMOSPHERIC PATHWAY Target Station



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Breakdown of radionuclides with potential of releasing into the ambient air Two main ways of ST generation are considered:

- spallation and activation of the air & cooling water and
- contamination of air with gaseous and volatile elements as well as dust (erosion and/or corrosion products).
- **3 important contributors to radioactive releases in locations where HVAC will be implemented:**



## HeL scheme |Helium purification system (HePS)



# **ATMOSPHERIC PATHWAY** Target Station | HeL Elements released from W target to HeL



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Gases	Volatile	es Semi- volatiles	Others ejected spallation products (particles)	Three ma responsi	Three main mechanisms responsible for releasing of the		
Noble	Halogen	s: As, Se, Sb,	Alkali metals: Li, Na, Rb;	radionuc	lides from the	W target	
gases:	I, Br, F, (	Cl Te, Ru, P,	Alkaline earths: Be, Mg, Ca, Sr, Ba;	within th			
Ne, Ar,		S, Cs, K	Boron group: B, Al, Ga, Te, In;	• sputt	ering or direct	ejection	
Kr, Xe			Transition metals: Sc, Ti, V, Cr, Mn, Fe,	(dom	inant),		
			Co, Ni, Cu, Zn, Y, Zr, Nb, Mo, Tc, Rh, Pd,	• diffus	sion,		
0, C*, N	Н		Ag Cd Hf Ta W Re Os	ablat	ion.		
			Metalloids: Be Si Ge Bh Sn Sh		1		
					RFs		
			Gd Th Dy Ho Fr Tm Yb Lu	Species	Ejection fraction & Diffusion	Conservatism	
*C0 or C0					(E. Pitcher, 2015)	Factor*	
				Noble gases (He-Ne-Ar-Kr-Xe)	3E-4; 1E-4; 3E-5	5	
Class of radio	nuclides	"Filter" type within	HePS	н	1	1	
Noble Gases, N Cl	, O, C, F, n	no		Volatiles (F-Cl-Br-I)	3.00E-04	10	
Semivolatiles	d	lust & aerosol special f	ilters	Semi volatiles	1.00F-04		
(As, Se, Sb, Te, Cs, K)	Ru, P, S, b ti	out with potential to es he filters and to be get	cape tered.	(C thru Zr)	1.001-04	5	
Non-volatiles ( metals) ejected target & W dus	mainly d l from t	lust & aerosol special f	ilters.	Semi-volatiles (Nb thru Os)	3.00E-05	2	

\*a conservatism factor applied to account for the calculations uncertainty.



Differential equations governing the system:  $\frac{dA_T}{dt} = c(1 - R_S) - (\lambda + D + \varepsilon_F + \varepsilon_H)A_T$  $\frac{dA_H}{dt} = A_T (D + \varepsilon_H) - (\lambda + G + L) A_H$  $\frac{dA_r}{dt} = LA_H - (\lambda + r)A_r$  $A_{out}(t) = r A_r(t)$  $A_{out} = r \int_0^T A_r(t) dt$  $A_{out} = r \int_{0}^{T} A_{rH}(t) dt = P N r \left[ a - \frac{a - b}{k \Gamma} \right] (1)$ 

Target activity	(1)	c Rg
Helium gas phase activity	(2)	λ D ε"
HVAC room activity	(3)	ε <sub>⊧</sub> G L
Activity to the stack	(4)	r T
Annual released activity	(5)	
$(-e^{-k_{1}T}) - \frac{b}{k_{2}T} (1 - e^{-k_{1}T})$	<sup>k2 T</sup> )]	

where:

Rele Rs

Operation time

**Compartment model for developing the** analytic expressions governing radioactivity releases from helium cooling loop

		Parameters used				
		Release Fraction Iodine	RF	3.00E-03	1	
		Getter_lodine (1/sec)	G	3.00E-05	1	
		Leakage (1/sec)	L	9.26E-09	1	
		HVAC_rate (1/sec) room: D02.115.4003	r	1.39E-03	1	
		Irradiation time (sec)	T <sub>irrad</sub>	1.80E+07		
		Radionuclide	T <sub>1/2</sub> (sec)	λ (1/sec)	Annual production in target (Bq)	Annual release (based on Eq. 11 & parameters above) (Bq)
		I 117	1.38E+02	5.02E-03	9.29E+13	1.11E+05
		I 118	8.20E+02	8.45E-04	1.21E+14	2.39E+06
		l 118*	5.10E+02	1.36E-03	5.51E+12	5.57E+04
		I 119	1.15E+03	6.05E-04	1.45E+14	4.42E+06
		120	4.86E+03	1.43E-04	1.85E+14	2.70E+07
		I 120*	3.18E+03	2.18E-04	1.41E+12	1.37E+05
		I 121	7.63E+03	9.08E-05	1.78E+14	3.84E+07
		I 122	2.18E+02	3.18E-03	2.10E+14	5.53E+05
		l 123	4.75E+04	1.46E-05	1.90E+14	1.17E+08
		I 124	3.61E+05	1.92E-06	7.17E+11	6.22E+05
		I 125	5.20E+06	1.33E-07	1.66E+14	1.53E+08
		I 126	1.12E+06	6.16E-07	6.43E+11	5.82E+05
		I 128	1.50E+03	4.62E-04	3.27E+11	1.38E+04
		I 129	4.95E+14	1.40E-15	1.17E+03	1.08E-03
		I 130	4.45E+04	1.56E-05	1.56E+10	9.39E+03
		I 130*	5.40E+02	1.28E-03	7.13E+09	7.84E+01
		131	6.93E+05	1.00E-06	9.79E+09	8.75E+03
		I 132	8.26E+03	8.39E-05	3.05E+09	7.01E+02
		I 132*	5.02E+03	1.38E-04	5.46E+08	8.20E+01
		I 133	7.49E+04	9.25E-06	1.33E+09	9.33E+02
_		l 133*	9.00E+00	7.70E-02	1.56E+08	9.97E-04
		I 134	3.15E+03	2.20E-04	1.56E+08	1.50E+01
		I 134*	2.21E+02	3.13E-03	6.82E+02	1.84E-06
		I 135	2.37E+04	2.93E-05	7.80E+07	3.57E+01
		I 136	8.34E+01	8.31E-03	1.96E+00	9.35E-10
	I	I 136*	4.69E+01	1.48E-02	1.82E+00	2.93E-10
		TOTAL				3.44E+08
re	:					
r	adionucli	de prod. rate	per radion	uclide		Bq/sec
R	elease fr	action sputtering	per radion	uclide (3E-	3 lodine )	
ra	adiation	decay constant	per radion	uclide		1/sec
d	iffusion r	ate	per radion	uclide		1/sec
unfilterable dust form, rate			1.67E-16			1/sec
filterable dust form rate			1.67E-11			1/sec
	unificati-	aust ionn. I dte	JE E			1/200
p	urnicatio		5E-5			T\Sec
le	eakage ra	ite	9.3E-9			1/sec
٧	'ent. syst	. exchange rate	per room	(1.4E-3 as e	example)	1/sec



sec

1.8E+7

# ATMOSPHERIC PATHWAY Target Station | **Hot Cells**



**1. Tungsten Dust** (pasted-up on the surfaces of the structures and pipes) => spread =>HVAC=> environmnent

Ablation rate 3E-4/year formation of W dust

$$A_{1y} = \frac{q_s}{\lambda} \left[ 1 - e^{-\lambda T_{irrad}} \right]$$

at the end of the 5<sup>th</sup> annual run:

$$A_{5y} = A_1 \sum_{i=1}^{5} \left[ e^{-\lambda T_1} \right]^{i-1}$$

 $RF_{H3}^{60degC} = 1E-4$ 

## 2. Stainless Steel Dust

$$RF_{H3}^{60degC} = 0.5$$



 $q_s = P e f_{pu}$  is the plated up rate of the W dust derived using: P = annual production, e = ablation rate,  $f_{pu}$  = fraction and  $T_{irrad}$  is the operation time

annual release to the environment:

$$A_{HC_W} = \frac{A_5 r}{\lambda + r} \left[ 1 - \frac{1}{T_{HC}(\lambda + r)} \left( 1 - e^{-(\lambda + r)T_{HC}} \right) \right]$$

ST from ActCs: short term release

-				dismantling
	Total from ActC	6.25E+11	6.24E+11	during
	Total dust	5.65E+08	1.70E+05	Release
1	W dust	1.06E+08	3.17E+04	6.24E+11
	SS dust	4.59E+08	1.38E+05	2.01E+08
	Nuclide class	release	release	зН
		i costituistic	Optimu	





# **Atmospheric releases**

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# Annual effective dose to the reference Averaged Man Results from Linac, HeL and Hot Cells

Annual Dose  $(\mu Sv/y)$  to the Average Adult and contributions for the two analyzed release cases. (Main stack) Effective dose (µSv/y) **Released activity SCREENING** REALISTIC Component (Bq/y) **APPROACH APPROACH** Optimal Pessimistic Optimal Pessimistic Optimal Pessimistic 2.25E+13 Linac 2.26E+13 1.16E+01 1.02E+01 2.84E-01 2.82E-01 **Helium Loop** 1.20E+01 5.57E-02 5.57E-02 Iodine 1.35E+09 1.35E+09 1.20E+01 Continuous release 5.94E+07 1.98E-03 other halogens 5.94E+07 1.98E-03 1.98E-03 1.98E-03 6.24E+11 H-3 1.08E+00 1.08E+00 5.60E-03 5.60E-03 6.24E+11 7.33E+03 7.33E+03 7.53E-09 7.35E-09 3.83E-09 3.83E-09 gases(O, N, CO) 1.28E-03 6.60E+09 6.60E+09 1.28E-03 1.28E-03 1.28E-03 Noble gases 8.17E+01 2.76E-07 2.45E-07 Metaloids 1 2.72E+05 9.20E-04 3.26E-07 Metaloids 2 5.29E+07 1.59E+04 2.06E-01 6.19E-05 1.02E-04 6.20E-05 6.32E+11 1.33E+01 Total 6.32E+11 1.31E+01 6.47E-02 6.47E-02 bunker 2.33E+13 2.31E+13 2.49E+01 2.33E+01 3.48E-01 3.47E-01 Total @ stack

u	Dismantling in ActC							
terr	ase	W dust	6.24E+11	6.24E+11	1.26E+00	1.08E+00	2.94E-02	2.92E-02
ort	ele	SS dust	6.60E+08	2.01E+08	7.44E-01	5.71E-04	2.18E-03	6.54E-07
shi	2	Total @ stack	6.25E+11	6.24E+11	2.01E+00	1.08E+00	3.16E-02	2.92E-02

#### Maximum release each 5 years

 TOTAL @ stack
 2.39E+13
 2.37E+13
 2.69E+01
 2.44E+01
 3.80E-01
 3.76E-01

Pessimistic release = abatement equipment @ stack =>down

Optimal release = abatement equipment @ stack works optimally



#### Total dose rate and contributions from different exposure pathways (Average Adult, optimal release rate)

PN	Annual	Cloud	Deposited		
<b>NIN</b>	dose rate,	immersion	radionuclid	Water inges	Food ingestion
Total	3.5E-01	92	1	0	7
N-13	1.1E-01	100	0	0	0
C-11	6.2E-02	98	1	0	0
0-15	4.5E-02	100	0	0	0
Ar-41	2.7E-02	100	0	0	0
I-125	1.7E-02	5	0	0	95
H-3	5.6E-03	19	0	22	60
Cl-39	5.4E-03	97	2	0	0
Cl-38	3.9E-03	98	1	0	0
P-32	2.1E-07	14	0	0	86
Be-7	5.8E-08	7	29	0	64
S-35	4.0E-08	7	0	0	93
W-185	1.0E-08	1	0	0	99
Hf-172	2.1E-09	12	52	0	37
Co-60	1.3E-11	5	81	0	15
Mn-54	6.0E-12	3	59	0	37
Co-58	3.6E-12	9	32	0	59

# ATMOSPHERIC PATHWAY Waste facility: source term



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The waste facility (WF) is a treatment and interim storage facility for radioactive waste. It is complementary to Hot Cells (HC).

## Main conditioning processes with potential airborne release:

- Cementation of the waste, such as spent ion exchange resins and activated/contaminated liquid waste;
- Treatment of the waste waters prior to discharging to the sewage system;
- Other treatment/conditioning processes (such as cutting), as well as the temporary storage of the obsolete getters and W dust bags from the Target Station releasing gases, volatiles and/or dust that will be extracted via HVAC through the WF stack to the atmosphere.

#### Processing activities to be performed periodically during campaigns => short-term release via a WF stack

## ST to the atmosphere can be estimates based on:

- Initial radioactive inventory of the waste prior to the treatment;
- Effectiveness of the treatment in terms of releasing (the percentage from the total radioactive inventory of the waste to be treated that will be released through the WF stack into the environment);
- Frequency and duration of the campaign (cementation, water treatment, other with release to the stack).

## A conservative envelope of radionuclides with potential of releasing from the WF is proposed:

- Radionuclides with potential to be contained in the waste water, produced via activation or contamination during ESS operations;
- Radionuclides of the structural material dust, that were selected based on their radiological impact, as representative isotopes of both: i) corrosion products bound in the ion exchange matrixes and ii) potential dust to be released during the cutting operations;
- Radionuclides of the W dust, as some of the potential contaminated waste from the target station monolith may be hosted in WF, directly or after temporarily HC storage.





The realistic Dose Factors were used to derive discharged limits by dividing the assumed dose target 10  $\mu$ Sv/y by the Dose Factor





# WATER PATHWAY

Source Term

According to SSM, none of the Swedish restrictions on liquid surface discharges: (SSMFS2008:23, 2008) applicable for nuclear power plants (NPP)s or (SSMFS: 2010:2, 2010) applicable for other type of facilities is valid for ESS.

ESS has to agree with wastewater treatment plant (Va Syd Källby Avlopps & Reningsverk) company in Lund the conditions for discharging.

## Breakdown of radionuclides with potential of discharging into the sewage system:

## A conservative envelope of 72 radionuclides potential contained in the waste water (ESS-0028551) :

i) activation of the water (Short lived radionuclides decay inside the close circuits & long lived ones such <sup>7,10</sup>Be are removed by the ion-exchangers);

Activated water of Main Cooling Circuits (MCC) is removed only if the chemistry of the water is damaged or from radiation protection reasons the continuous addition of fresh water within MCC is necessary. The replaced water from the circuit may be discharged after treatment if the activities of the constituents are below the allowed discharged surface water limits.

## ii) contamination of water with corrosion products;

## iii) contamination with W dust.

(Ion-exchangers clean also activated corrosion products from the metallic pipes & W dust)

Selection of the isotopes was done taking into consideration the existing measurements and experience in management of the wastewater of other spallation facilities, such LANSCE, USA (Borden M., 2014), ISIS, UK (Boyer F., 2011), (Masterson P., 2014), FERMILAB, USA (Vaziri K., 2014), CERN Switzerland (Vojtyla P, 2005).

The treatment, method within Waste Facility of the waste water contributes essentially in defining the ST of discharges. Experience in Swedish NPPs (Hoglund A., 2015) is the use of an evaporator in order to treat the wastewater. In the resulted water to be discharged into environment remains less than 1 Bq/kg of gamma emitters.

Because quantification of the ST is not known & No assumptions on classes of radionuclides (gross beta, gross gamma, others) are available it was decided to derive the ESS specific discharge limits for all 72 radionuclides.

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SOURCE

## Discharge of radioactive substances to the sewage system

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#### The assessment accounts for exposure to

- 1) workers in the sewage plant that are exposed to radionuclides in the non-treated water or sewage sludge and
- 2) the general public exposed, either directly (through consumption) or indirectly (though use of water for irrigation or for feeding cattle).

The realistic Dose Factors were used to derive discharged limits by dividing the assumed dose target 10 µSv/y by the maximum of the Dose Factor for workers and general public

## Waste facility outlet:

## Derived Reference Discharge Limits for waste water (Bq/y) -72 radionuclides



# GROUNDWATER PATHWAY: Source Term & annual dose



Activity of nuclides produced in the 1<sup>st</sup> meter thick soil layer around the ESS tunnel wall at shutdown after 40 years of continuous operation.

Rn*	T <sub>1/2</sub>	Total accumulated inventory at 40 y (Bq)	Average production rate, Bq/y	Leaching & H <sub>2</sub> O content (%)
H-3	12.33 y	2.10E+10	1.32E+09	100 & 16
Be-7	53.12 d	7.40E+09	3.52E+10	
Na-22	2.60 y	4.37E+09	1.17E+09	15 & 16
Na-24	14.96 h	5.01E+10	2.03E+13	
P-32	14.26 d	1.61E+09	2.86E+10	
S-35	87.32 d	5.27E+08	1.53E+09	
Ca-45	162.61 d	4.31E+07	6.71E+07	
Mn-54	312.3 d	1.62E+09	1.31E+09	
Sc-46	83.79 d	9.75E+08	2.94E+09	
Fe-55	2.73 y	1.40E+10	3.55E+09	
Zn-65	244.26 d	1.87E+06	1.94E+06	

\* Selection based on (Sullivan, 1992)

#### Two scenarios of radioactivity leaching from the soil:

- assumes an impermeable soil cover (or membrane), situated on the top of a tunnel, which will fail after the end of operations of the accelerator (the 40<sup>th</sup> year), leading to the leaching of the accumulated nuclides in the activated soil to groundwater;
- 2. assumes that there is **no protective cover present**, and a chronic release occurs through the operation life time of the facility (40 years).
- Groundwater transport is carried out in two steps:
- 1. Groundwater flow modeling using **MODFLOW** code and
- 2. Radionuclide transport modeling using the **NORMALYSA** model library implemented in Ecolego 6.0

#### Infiltration recharge Tunnel Elevation, m Well Activated soi 81 Surface soil (clay till) Water 1 m ļ fluxes 5 m Groundwater leve 74.7 m ~ 74.5-75 m 9n 72 Deep soil (sand) 2m70 300 m

## 2D cross-sectional groundwater flow model

#### Radionuclide activities (Bq/m<sup>3</sup>) in the water of the well @ 300 m

	Scenario 1		Scen		
Rn	Reference K <sub>d</sub> *	Conservative K <sub>d</sub> **	Reference K <sub>d</sub> *	Conservative K <sub>d</sub> **	
H-3	3.57E+04	3.57E+04	1.67E+04	1.67E+04	< 100 Bq/L
Na-22	1.45E-32	2.91E-11	1.28E-31	2.01E-10	(EU DWD
S-35	1.21E-88	1.10E-43	2.19E-87	1.75E-42	2015/1787)

\*K<sub>d</sub> taken from (Sheppard et al., 2011)

#### \*\* Kd (= Ref K<sub>d</sub>/10)

#### Annual dose rate (Sv/y)

	Scenario 1		Scenario 2		
Rn	Ingestion	Total	Ingestion	Total	
	of water		of water		
H-3	2.42E-07	2.45E-07	1.130E-07	1.15E-07	
Na-22	1.74E-41	3.22E-41	1.54E-40	1.54E-40	
S-35	3.51E-98	6.50E-98	6.34E-97	1.30E-40	18
Total	2.42E-07	2.45E-07	1.13E-07	1.15E-07	

# Locations on ESS site (selection of locations outside the fence)



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# Summary



**Q**Annual dose from the direct radiation is less than 10  $\mu$ Sv/y;

Maximum annual dose for continuous release to the main stack from Linac, HeL is less than 1 μSv/y and the major contribution (80%) is given: <sup>13</sup>N, <sup>11</sup>C, <sup>41</sup>Ar, <sup>15</sup>O, <sup>3</sup>H, <sup>125</sup>I, <sup>39,38</sup>Cl, <sup>32</sup>P, <sup>7</sup>Be, <sup>35</sup>S, <sup>185</sup>W, <sup>172</sup>Hf, <sup>60</sup>Co, <sup>54</sup>Mn, <sup>58</sup>Co;

 $\Box$ Maximum annual dose for short term release to the main stack is less than 0.1  $\mu$ Sv/y and the major contribution is given by <sup>3</sup>H release during dismantling;

The source term to the waste stack is not known therefore derived activity limits where derived for all 93 radionuclides with potential to be released;

The source term to the waste facility outlet is not known therefore derived activity limits where derived for all 72 radionuclide with potential of discharging into the sewage system;

**D**Maximum annual dose due to the migration of the contaminant with the groundwater is less than  $1 \mu Sv/y$  and is due to the drinking water contaminated with <sup>3</sup>H;

 $\Box$ Actual results of realistic dose calculations for airborne releases of radionuclides from the main stack show that annual doses to the public are well below the regulatory constraint of 100  $\mu$ Sv/y and even below the exemption level of 10  $\mu$ Sv/y. Obtained results shall be completed with remaining contributors.

This report is conceived according the knowledge that ESS staff has in this stage of the project. The current data are estimations subjected to evolution and update.

An environment monitoring program was defined and it will be implemented gradually during the commissioning of the ESS facility 20