LIMNOLOGY AND OCEANOGRAPHY

### ASLO e-Lectures



# Radioactivity in the Marine Environment

**Understanding the Basics of Radioecology** 

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#### What is Marine Radioecology?

Marine radioecology examines how radioactive substances interact with the marine environment and the various mechanisms and processes that influence radionuclide migration in the marine ecosystem.

This field of study includes aspects of field sampling, design of field and laboratory radiotracer\* experiments, the development of predictive simulation models, and dose assessments to humans and biota.

\* Includes radionuclides and radiolabeled compounds

Courtesy of Scott Fowler

- 1. Overview of how marine biota uptake radionuclides from marine systems
- 2. Biogeochemical transfer and transport pathways of radionuclides in the marine environment
- 3. Radiation dose and exposure
- 4. Environmental protection: Understanding ecological risk assessment and management

1. U-Th series radionuclides – of primordial origin, occur naturally on land and in ocean, and produce a series of "daughter" radionuclides via radioactive decay.

Examples: <sup>238</sup>U, <sup>234</sup>Th, <sup>210</sup>Pb, <sup>223, 224, 226, 228</sup>Ra and <sup>222</sup>Rn.

2. Cosmogenic Radionuclides – continuously being created by cosmogenic rays that interact with materials in the atmosphere and on Earth.

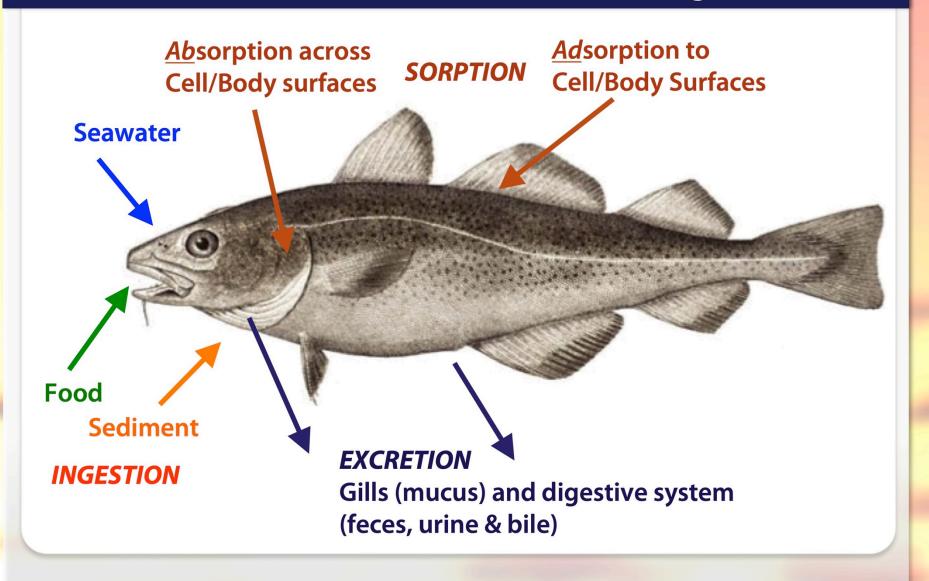
Examples: <sup>3</sup>H, <sup>14</sup>C, <sup>7</sup>Be

3. Anthropogenic radionuclides – continuously being produced by humans

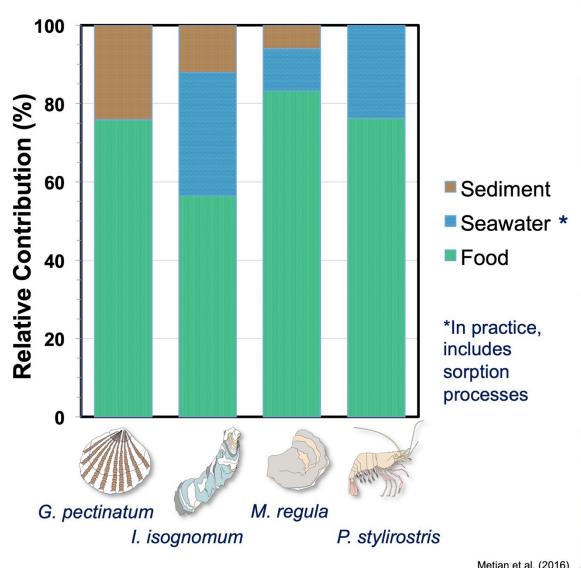
Examples: <sup>3</sup>H, <sup>14</sup>C, <sup>90</sup>Sr, <sup>137</sup>Cs, <sup>129</sup>I, <sup>238, 239, 240</sup>Pu

Note: Some radionuclides have both cosmogenic and anthropogenic sources (e.g., <sup>3</sup>H and <sup>14</sup>C).

#### 1. Radionuclide bioaccumulation in marine organisms?



#### 1. Radionuclide bioaccumulation in marine organisms?



**Ingestion:** Relative contributions (%) of three uptake pathways (sediment, seawater, and food) to the total bioaccumulation of <sup>134</sup>Cs in marine organisms.

Metian et al. (2016)

#### **Terminology**

**Concentration Factor (CF)\*:** Ratio of the radionuclide activity in a specific organism relative to the radionuclide concentration *in ambient sea water*.

$$CF = \underline{Bq \ g^{-1} \ wet \ weight \ of \ organism}}$$

$$\underline{Bq \ g^{-1} \ sea \ water}$$

**Transfer Factor (TF)\*:** Ratio of radionuclide activity in a specific organism relative to the radionuclide concentration *in sediment or food*.

TF =  $\underline{Bq g^{-1} wet weight of organism}$  $\underline{Bq g^{-1} sediment or food}$ 

#### **Uses:**

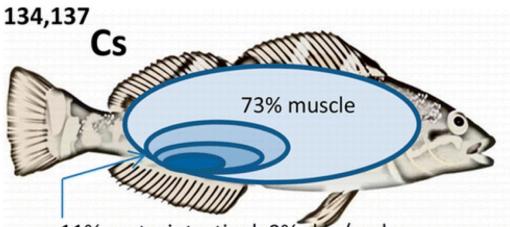
- Compare relative bioavailability of different radionuclides to a given organism
- Compare ability of different organisms to accumulate a given radionuclide
- <u>Allow the development of models</u> that can predict the resulting radionuclide activity in an organism from known seawater activities.
- To identify potential "sentinel organisms" of radionuclide contamination.

Carvalho et al. (2018)



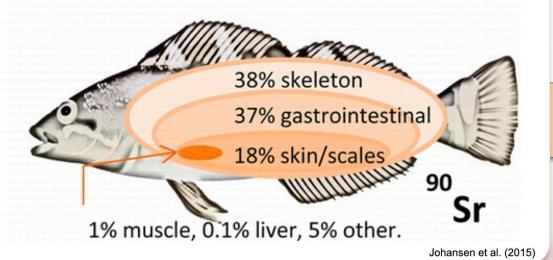
<sup>\*</sup>Assumes input = output and that the radionuclide of interest exists in soluble forms in seawater

#### Distribution of <sup>134,137</sup>Cs and <sup>90</sup>Sr in fish tissue\*



<sup>137</sup>Cs is analog to <sup>40</sup>K

11% gastrointestinal, 9% skin/scales, 3% skeleton, 2% gills, 1% liver, 1% testes/ovary.

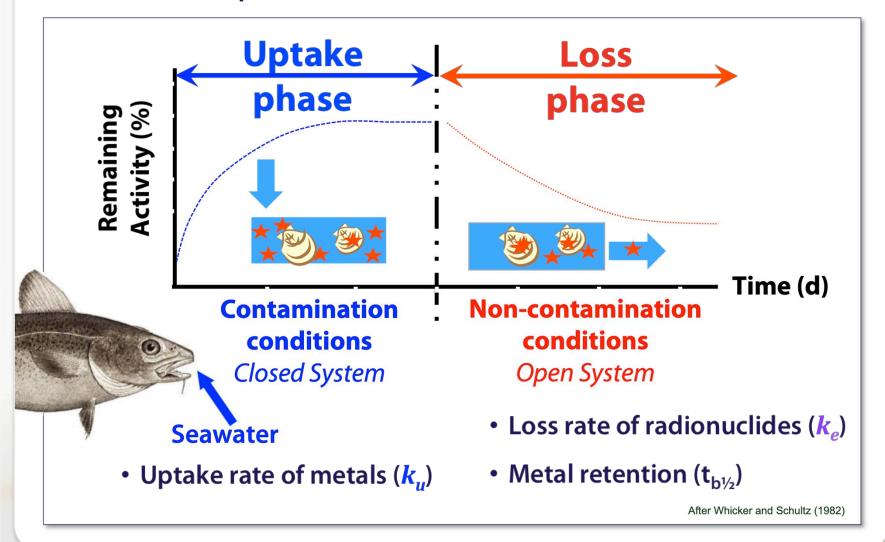


<sup>90</sup>Sr is analog to Ca

\* 2 years after the Fukushima accident

#### **How to determine the CF: (Lab experiments)**

Model development of radionuclide contamination via seawater



#### How to determine the CF

#### **Uptake phase: Depends on time (kinetic parameters)**

$$CF_t = k_u t$$
 (During uptake phase,  $k_u$  dominates)

$$CF_t = CF_{SS}(1 - e^{-k_e t})$$
, where at steady state  $CF_{SS} = \frac{k_u}{k_e}$ 

#### Where:

 $CF_t$  - the concentration factor at time t (d)

 $CF_{SS}$  - the concentration factor at steady state

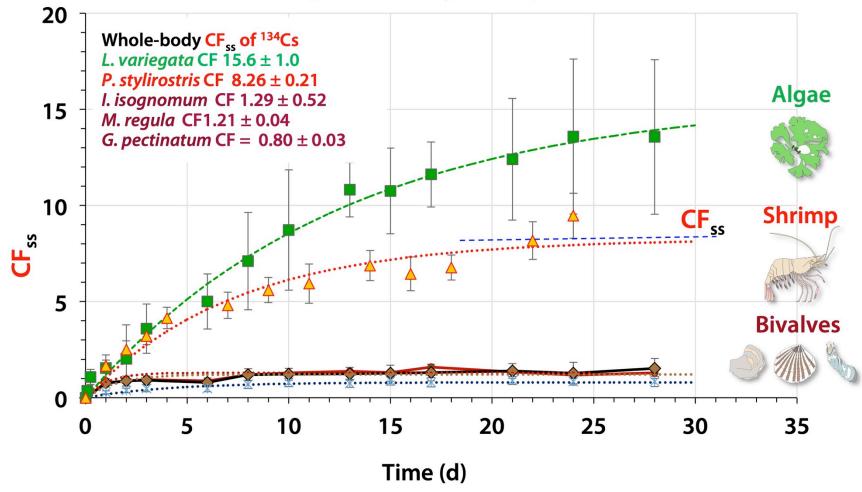
 $k_u$  - uptake rate constant (d<sup>-1</sup>)

 $k_e$  - loss rate constant (d<sup>-1</sup>)

After Whicker and Schultz (1982)

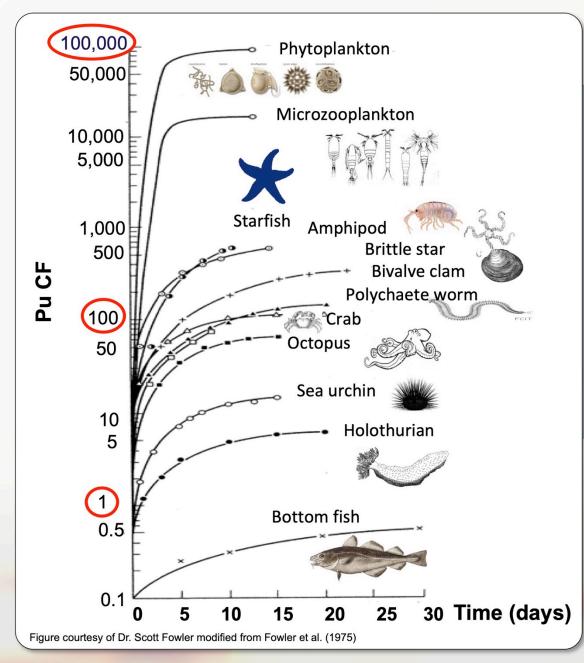
#### How to determine the CF

### Example: Uptake kinetics ( $k_u$ ) of dissolved <sup>134</sup>Cs ( $t_{\frac{1}{2}}$ = 2.06 y) during 24 - 28 days of exposure



Metian et al. (2016)





Species dependent uptake kinetics of plutonium from seawater by various marine organisms

Note the variability in  $k_u$ 

## Effect of Temperature on the CF (specifically $k_u$ ) of $^{60}$ Co, $^{241}$ Am and $^{134}$ Cs from water by brown macroalgae



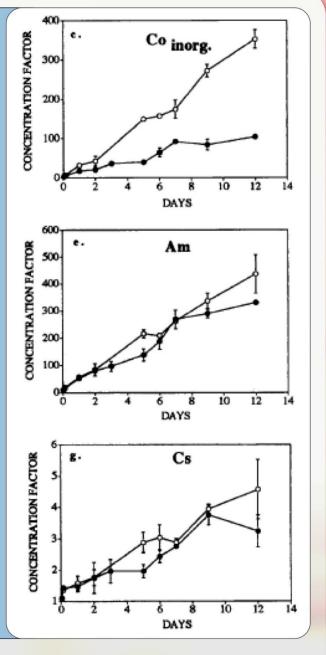


Image: http://www.seaweed.ie/descriptions/fucus\_vesiculosus.php

12°C ( ° )

2°C(•)

(Boisson et al, 1997)



#### Loss (depuration) phase: kinetic parameters (k<sub>e</sub>)

Loss kinetics are used to determine the biological half-life  $(t_{b\frac{1}{2}})$  and can be modeled using a *one component* exponential model:

$$A_t = A_0 e^{-\mathbf{k_e}t}$$

#### Where:

 $A_t$  - Remaining activities at time t (%)

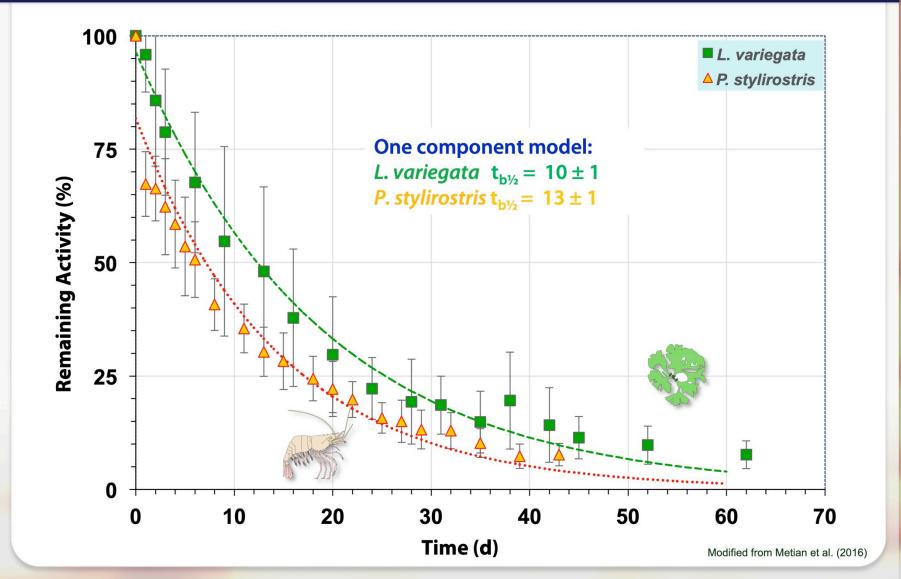
 $A_0$  - Remaining activities at time 0 (%)

 $k_{\rho}$  - Loss rate constant (d<sup>-1</sup>)

$$T_{b\frac{1}{2}} = In(2)/k_e$$

After Whicker and Schultz (1982)

### Example: $^{134}$ Cs ( $t_{\frac{1}{2}}$ = 2.06 y) depuration kinetics when maintained for 43 – 62 d in clean seawater



#### Loss (depuration) phase: kinetic parameters (k<sub>e</sub>)

In some cases, the loss phase should be described by a *two component* model that may reflect differences in specific tissues where the radionuclide has accumulated.

$$A_t = A_0(N_1^* e^{-ke1^*t} + N_2^* e^{-ke2^*t})$$

Where:  $A_t$  - Remaining activities at time t (%)

 $A_0$  - Remaining activities at time 0 (%)

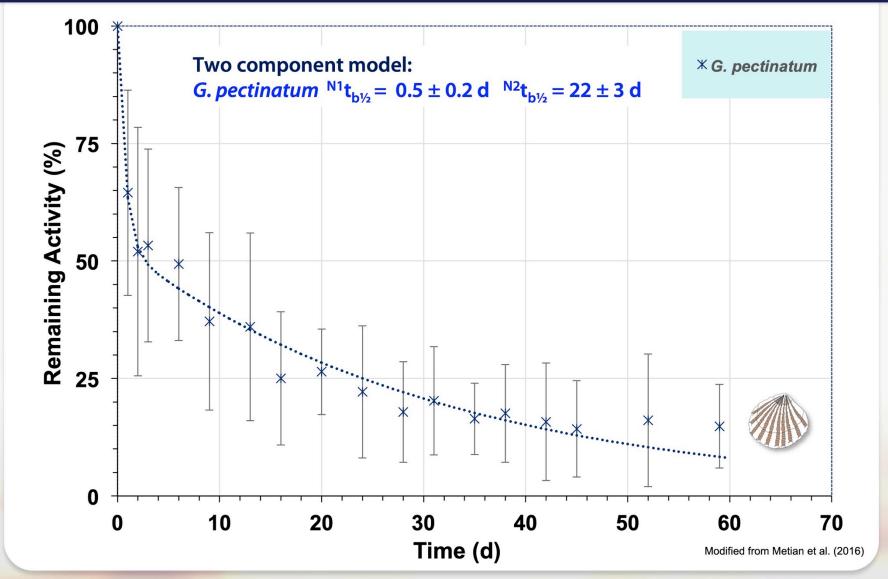
 $k_e$  - Loss rate constant (d<sup>-1</sup>) for a specific component N<sub>1</sub> and N<sub>2</sub>

 $N_1$  and  $N_2$  - Denote differences in the loss rates with respect to specific components (e.g., tissues vs. shell) within an organism, where  $N_1 + N_2 = 1$ 

Biological half-lives (Tb<sub>1</sub> & Tb<sub>2</sub>) can be calculated from the corresponding depuration rate constant ( $k_{e1}$  &  $k_{e2}$ ) according to the relation Tb=ln(2)/ke

After Whicker and Schultz (1982)

### Example: $^{134}$ Cs ( $t_{\frac{1}{2}}$ = 2.06 y) depuration kinetics when maintained for 43 – 62 d in clean seawater



### **CF**<sub>ss</sub> of selected radionuclides within different taxonomic groups

Organism	<sup>137</sup> Cs	<sup>239+240</sup> Pu	<sup>241</sup> Am	<sup>210</sup> Po
Macroalgae	50	4,000	8,000	1,000
Phytoplankton	20	200,000	200,000	70,000
Zooplankton	40	4,000	4,000	30,000
Decapod crustaceans	50	200	400	20,000
Bivalve Molluscs	60	300	1 000	20,000
Cephalopods	9	50	100	20,000
Teleost fish	100	100	100	2,000

IAEA (2004)



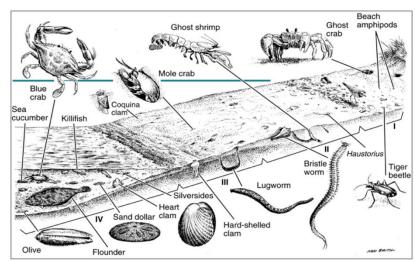
**Seawater** 

#### **TF** of radionuclides accumulated from contaminated sediments

Organism	Uptake Time (days)	<sup>239+240</sup> Pu	<sup>241</sup> Am	<sup>137</sup> Cs	<sup>60</sup> Co
Worms	11 - 50	0.0016 -0.002	0.0009 -0.003	0.2	0.06
Clams	40 - 50	0.006 - 0.01	0.004 - 0.02		
Isopod	40 - 50		0.006-0.032		
Amphipod	14	0.10	0.11		

Fowler et al. (1997), IAEA (1985)





### Assimilation Efficiencies\* of selected radionuclides within different taxonomic groups

\*AE = The fraction of ingested radionuclide via food that is absorbed and retained in body tissues

Organism Food type	<sup>137</sup> Cs	<sup>239+240</sup> Pu	<sup>241</sup> Am	<sup>210</sup> Po
Zooplankton feeding on Phytoplankton	1	0.8 – 1	0.9 – 10	20 – 55
Decapod crustaceans feeding on artemia	1	10 – 60	8 – 58	5
Bivalve Molluscs feeding on phytoplankton	3 - 4	0.9	0.6 – 38	17
Cephalopods feeding on crabs	23 – 29	1	51 – 60	1
Teleost fish feeding on shrimp	42 – 95	0.1 – 1	0.7 – 6	5

Stewart et al. (2005); Fowler et al. (1975; 2005)



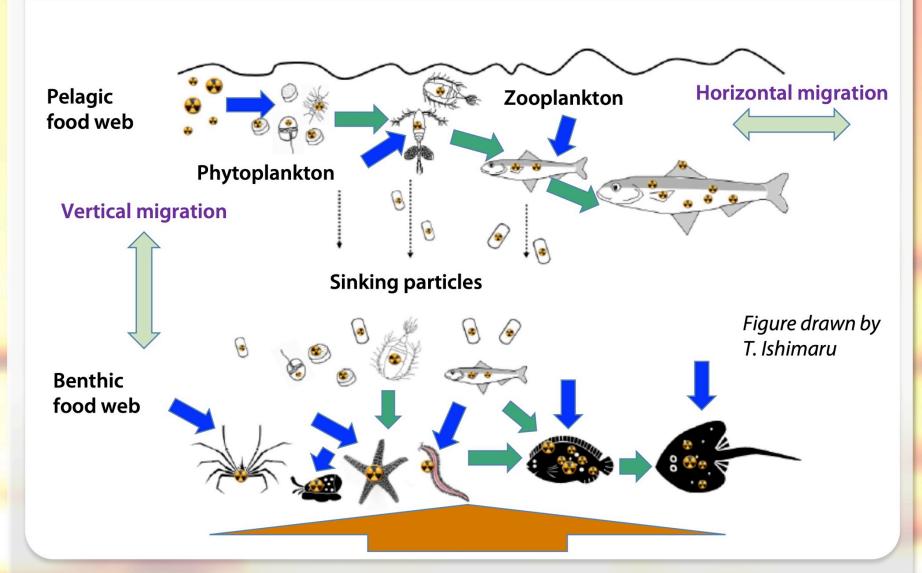


#### **Combined ingestion and excretion pathways**

Bulk <sup>137</sup>Cs activities in different *size classes* of male and female European Hake females males 2.0 <sup>37</sup>Cs (Bq kg<sup>-1</sup> dry weight) \*\* 1.5 \*\* 1.0 ns 0.5 0.0 25-29 30-34 35-39 40-49 50-59 >60 Size (cm) Harmelin-Vivien et al. (2012)

Significance of male-female difference: ns = not significant, \* p < 0.05, \*\* p < 0.01

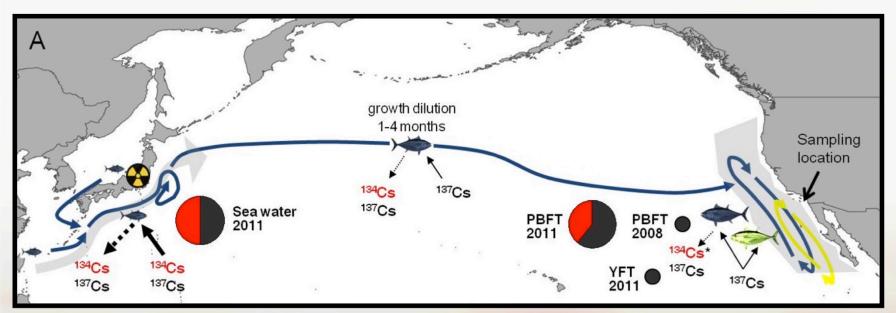
### 2. Biogeochemical transfer and transport pathways of radionuclides in the marine environment



#### **Horizontal Transport**

Here, Pacific Bluefin Tuna (PBFT) transported <sup>134</sup>Cs during their migration across the Pacific Ocean after having been contaminated near Japan by the releases of the Fukushima Dai-ichi nuclear power plant.

Note that Yellowfin Tuna (YFT) contained none of the short lived <sup>134</sup>Cs, confirming that transport was not by water movement alone, but by active horizontal transport by the PBFT.

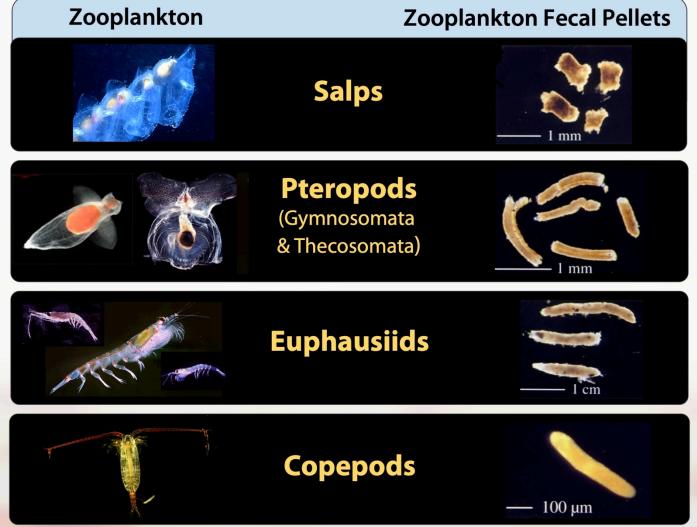


Madigan et al., 2012



#### **Vertical Transport**

Production of fecal pellets is an efficient way for transporting contaminants to the seabed



ASLO e-lectures

#### **Vertical Transport**

Concentrations of selected radionuclides (Bq kg<sup>-1</sup> dry) in plankton and sinking particulate matter in the water column

Organism	<sup>239+240</sup> Pu	<sup>241</sup> Am	<sup>210</sup> Po	<sup>137</sup> Cs	<sup>134</sup> Cs
Phytoplankton	0.35 – 1.0	0.052 - 0.22	21 – 61	-	-
Copepods Fecal pellets	0.05 – 0.48 1.2 - 3.4	0.022 - 0.12 1.0 - 2.7	126 617	*34 ± 7 *6300 ± 1000	*22 ± 6 *3400 ± 600
Euphausiids Fecal Pellets Molts	0.015 3.6 0.18 ( <b>90%)</b>		40.7 648 – 1000 13.3 ( <b>2.5%)</b>	1 1	
Salps Fecal pellets	- 8.9	- 5.0	260 583 - 750	-	-
†Japanese Copepods (Pre – Fukushima)	-	-	÷	0.42 ± 0.15	·-
**Japanese Copepods (Post – Fukushima)	-	-	1	0 - 56	0 - 46

Courtesy Scott Fowler

<sup>\*</sup> Samples are Chernobyl fallout-enriched (Fowler et al., 1987)

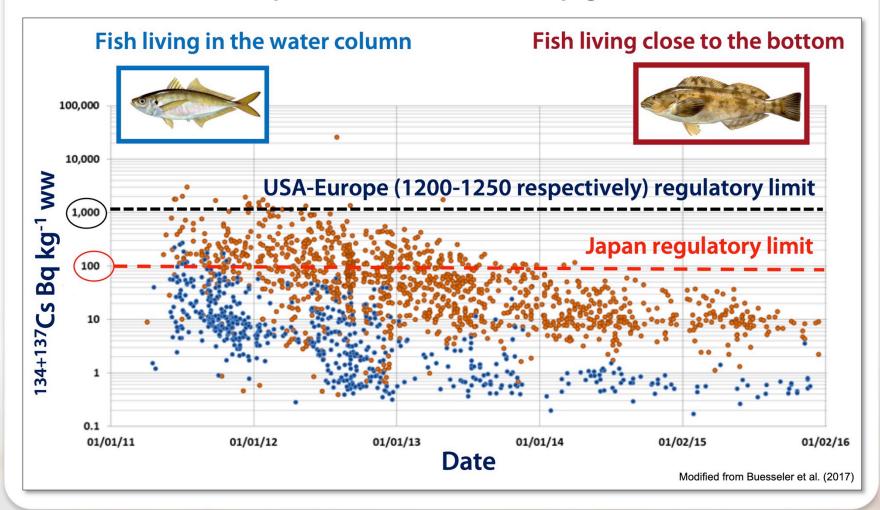
<sup>†</sup> from Tateda (1998)

<sup>\*\*</sup> from Buesseler et al. (2012)

<sup>( ) =</sup> Percent of euphausiid whole body radionuclide concentration contained in the molt

#### **Vertical Transport**

Following accidental release from Fukushima Dai-ichi Nuclear Power Plant Example: radiocesium in fish (Bq kg<sup>-1</sup> ww)

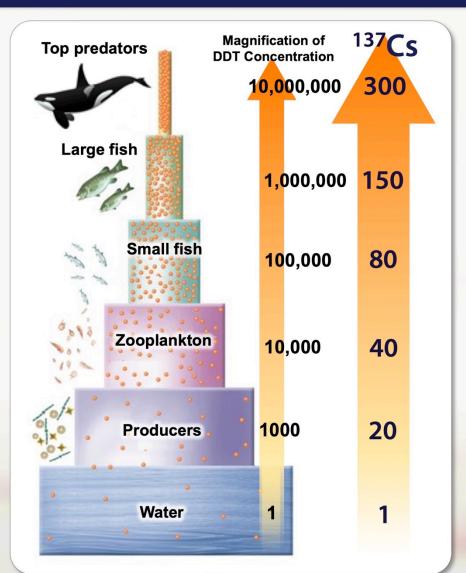


### Transport within food chains: Do radionuclides biomagnify in marine food chains?

Biomagnification = increasing concentrations in organisms at successively higher levels in the food chain.

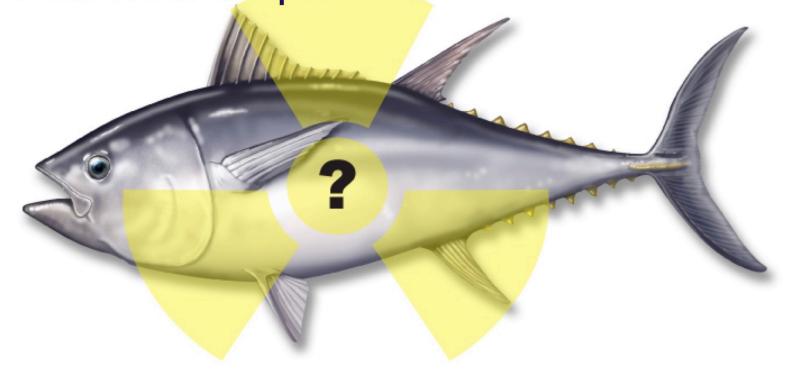
A limited number of substances magnify in marine food chains. For example: mercury, PCBs, and DDT.

Cesium and Polonium are radionuclides that have *limited* biomagnification.



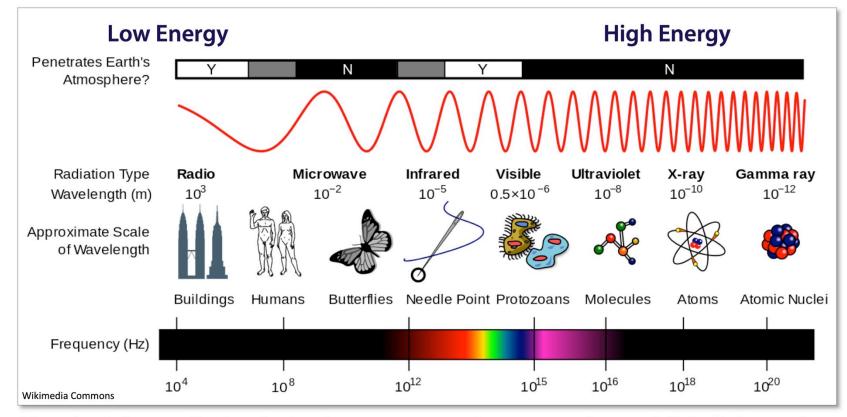
#### **Toxic radionuclides in marine food chains**

Once radionuclides are released into the environment and incorporated into marine organisms, how do we assess *potential impacts* on the environment and *risks* associated with seafood consumption?



#### 3. What is Radiation?

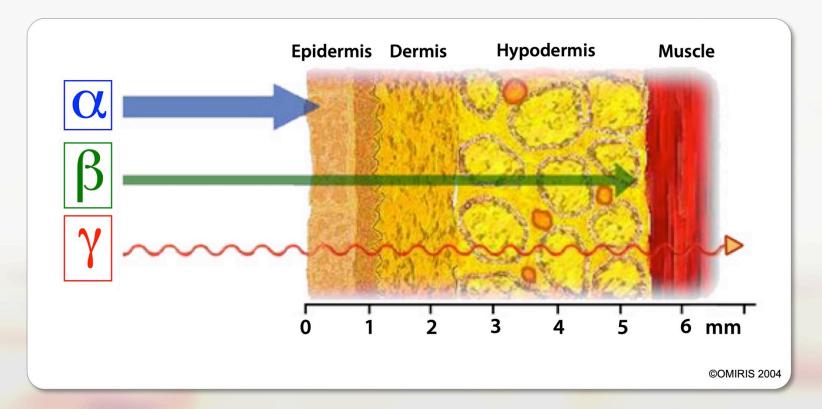
Radiation is energy in the form of high-speed particles (or electromagnetic waves or photons). It can be ionizing or non-ionizing.



Non-ionizing radiation lacks the energy to alter atoms (e.g., visible light and microwaves). Ionizing radiation has enough energy to change normal cellular functioning. Ionizing radiation may cause cells to die or transform into a cancerous cell. It is these particles that are emitted from a radioactive nuclide.

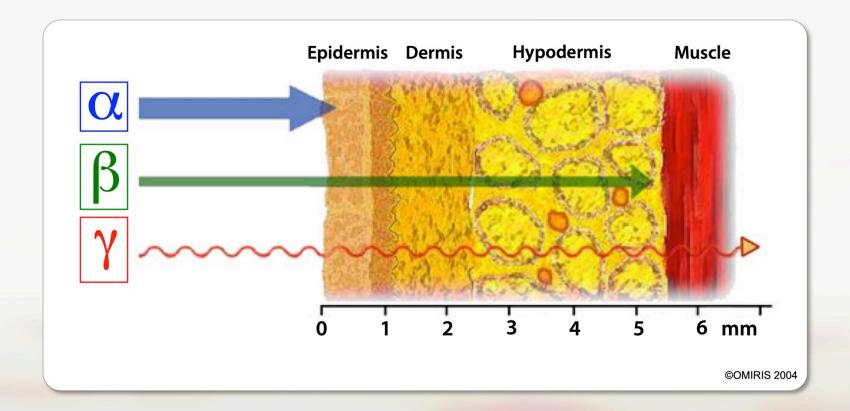
#### Radiation types we primarily focus on:

Alpha (a) particles: Most densely ionizing, but least penetrating. This means that cells can be protected or shielded from damage by alpha particles by clothing. Even the dead outer layer of your skin will protect you from damage from alpha particles. However, if alpha emitters are inhaled or ingested or get into a cut on the skin, they can cause damage to cells. As alpha particles are emitted inside the body, the surrounding cells are damaged.



#### Radiation types we primarily focus on:

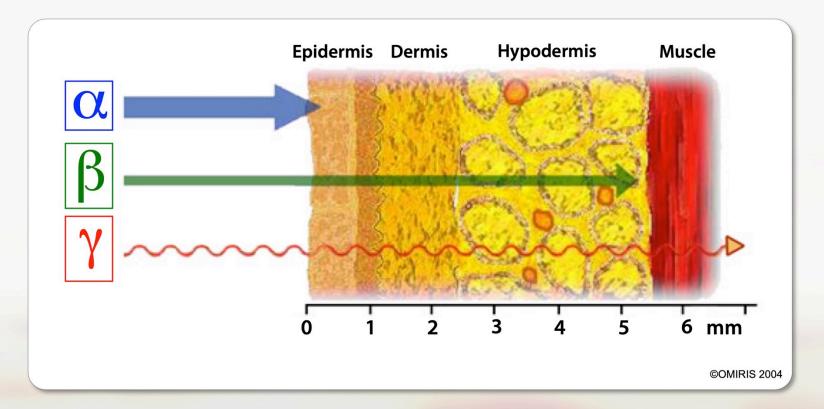
**Beta** ( $\beta$ ) **particles:** More energetic. Can travel several feet through air. Are stopped with denser materials such as wood, glass, or aluminum foil.



#### Radiation types we primarily focus on:

#### **Gamma** (γ) rays:

High-energy electromagnetic energy waves and the most penetrating type of radiation. Cells must be shielded from gamma rays with concrete, lead or steel. Not all may do cellular damage, but they must interact with the material to do so.



#### **Development of International Standards**

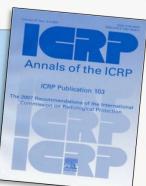
**SCIENCE**Doses and effects

UNSCEAR www.unscear.org



PRINCIPLES
Philosophy and policy

**ICRP** www.icrp.org



**STANDARDS** 

**Nuclear Safety and Security** 

**IAEA**www.iaea.org

IAEA Safety Standards
for protecting people and the environment

Radiation Protection and
Safety of Radiation Sources:
International Basic
Safety Standards
Safety Standards
Safety Standards
Safety Standards
Safety Standards
Safety Standards
No. GSR Part 3

No. GSR Part 3



#### **Radiation Dose Concepts**

 Adsorbed dose (Gray, Gy) is the absorption of radiation per unit mass of tissue. 1 Gy = 1 Joule per kg

#### For humans only:

• Equivalent dose (Sievert, Sv) adjusts for biological damage by different types of ionizing radiation ( $\alpha$ ,  $\beta$ ,  $\gamma$ ) using a weighting factor ( $W_R$ )

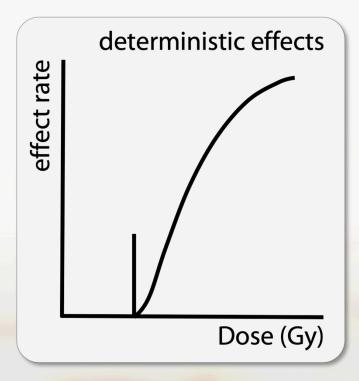
$$\mathbf{w}_{R} = 1 (\beta, \gamma, X) \text{ and } 20 (\alpha)$$

 Effective dose (Sievert, Sv) is the tissue-weighted sum of the equivalent doses in all specified tissues and organs of the human body.

#### **Different types of Effects: Deterministic versus Stochastic**

Deterministic effects: Severity increases with the dose. Radiation dose levels, or "thresholds", are determined, below which, effects from

radiation exposure are absent. Generally they are short term effects and specific to the *individual*.



#### **Dose Threshold for Deterministic Effects\***

Tissue	Total Acute Dose Threshold (Gy)		
Lens of eye			
Detectable opacities	0.5–2	>1 year	
Cataract formation	5	>1 year	
Skin			
Skin reddening	3–6	1–4 weeks	
Temporary hair loss	4	2-3 weeks	
Skin death and scarring	5–10	1–4 weeks	
Testes			
Temporary sterility	0.15	3-9 weeks	
Permanent sterility	3.5-6	3 weeks	
Ovaries			
Permanent sterility	2.5-6	< 1 week	
Gastrointestinal			
Mucosa lining loss	6	6–9 days	
Bone marrow			
Reduction of blood cell production	0.5	1–2 months	
PSE \$15504.0500700035			

\* 1% incidence level based on ICRP publication 103 (2007)

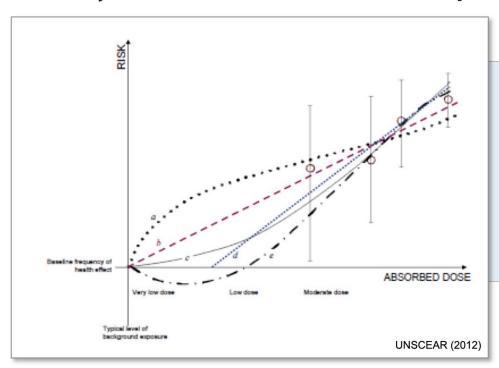
From Peck and Samei (2017)



#### Different types of Effects: Deterministic versus Stochastic

#### **Stochastic effects:**

Severity of stochastic effects is independent of the dose.



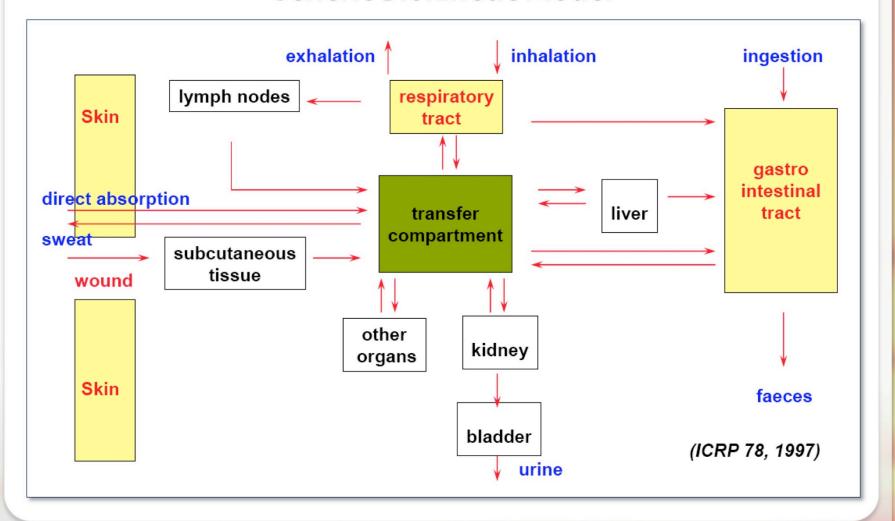
#### **Possible Effects:**

- 1) Cancer (e.g., long-term results from smoking)
- 2) Hereditary defects (e.g., Down Syndrome)

There is no threshold, and the probability of having effects is proportional to the dose adsorbed at the population level. In other words, depending on the conditions of exposure, effects may or may not occur.

# How do you determine the **Effective Dose** to a human?

## **Generic Biokinetic Model**



## How do you determine the **Effective Dose** to humans?

$$E = \sum_{T} W_{T} H_{T} = \sum_{T} W_{T} \sum_{R} W_{R} D_{T,R}$$

$$E = \sum_{\mathbf{T}} W_{\mathbf{T}} H_{\mathbf{T}} = \sum_{\mathbf{T}} W_{\mathbf{T}} \sum_{\mathbf{R}} W_{\mathbf{R}} D_{\mathbf{T},\mathbf{R}} \qquad \text{(Sv)}$$

$$E = \sum_{\mathbf{T}} W_{\mathbf{T}} H_{\mathbf{T}} = \sum_{\mathbf{T}} W_{\mathbf{T}} \sum_{\mathbf{R}} W_{\mathbf{R}} D_{\mathbf{T},\mathbf{R}} \qquad \text{(Sv)}$$

$$Radiation quality$$

$$Tissue radiosensitivity$$

- Based on a linear, no threshold (LNT) dose-risk relationship
- For stochastic effects only
- E and  $H_T$  are not directly measurable and are model-derived

ICRP, 2007

## Tissue Weighting Factor: $(W_T)$

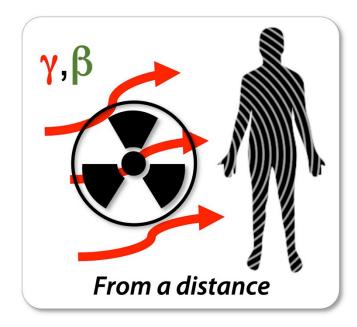
Tissue	$W_T$	$\sum W_T$
Bone-marrow (red), colon, lung, stomach, breast, remainder tissues	0.12	0.72
Gonads	0.08	0.08
Bladder, oesophagus, liver, thyroid	0.04	0.16
Bone surface, brain, salivary glands, skin	0.01	0.04

Reminder:  $w_R = 1 (\beta, \gamma, X)$  and 20 ( $\alpha$ )



ICRP, 2007

## **Irradiation versus contamination**



#### Irradiation

Body is exposed to external radiation (i.e., <u>via water</u>, soil/sediment and air)



## **Contamination**

The radioactive substance is directly on the skin (external) or within the body (internal)

# Let us examine one pathway as an example and major concern for Marine Systems: Seafood ingestion

$$D_{eff-ing} = \sum_{i} \sum_{j} A_{i,j} Q_{j} DC_{i}$$

#### Where:

 $D_{eff-ing}$   $A_{i,j}$   $Q_{j}$   $DC_{i}$ 

Effective dose by ingestion
Radionuclide i massic activity in foodstuff j
Consumption rate of foodstuff j
Dose coefficient for radionuclide i

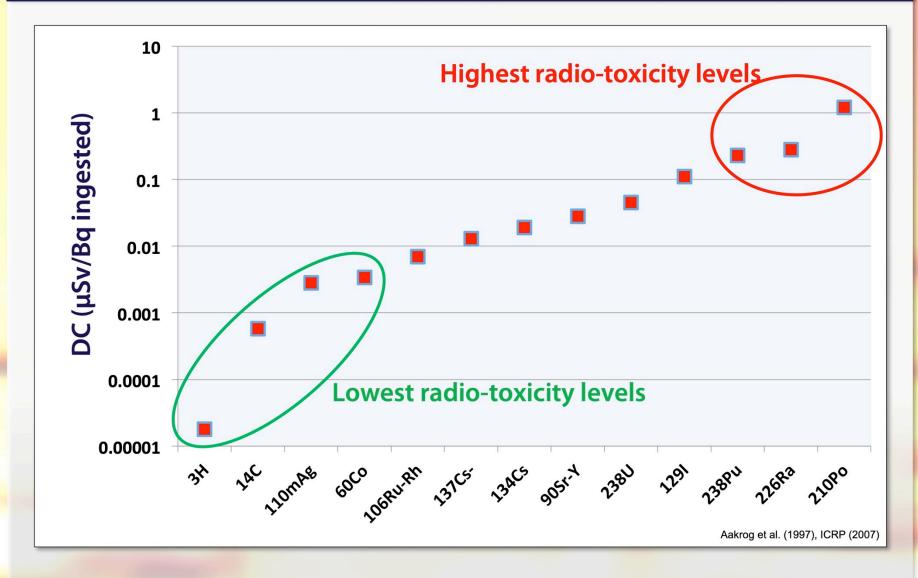
Sv y<sup>-1</sup>
Bq kg<sup>-1</sup>
kg y<sup>-1</sup>
Sv Bq<sup>-1</sup> ingested

#### **Specific Cases:**

The potassium concentration is kept constant by humans. The proportion of  $^{40}$ K to total K (specific activity: Bq kg $^{-1}$  of potassium) is also constant. So the  $^{40}$ K whole body activity is constant and leads to an effective dose of  $\sim 170~\mu Sv~y^{-1}$  for an adult (185  $\mu Sv~y^{-1}$  for a child).

For similar reasons,  $^{14}$ C activity is constant in the human body. This leads to an annual effective dose of  $\sim 12 \ \mu Sv/y$ .

# Dose coefficients (DC) for some naturally occurring and artificial radionuclides (ingestion pathway)



# After ingestion, how long do these radionuclides exist in the body (Effective t<sub>1/2</sub>)

# The effective half life (t<sub>1/2</sub>) represents the combination of the physical and biological half-lives.

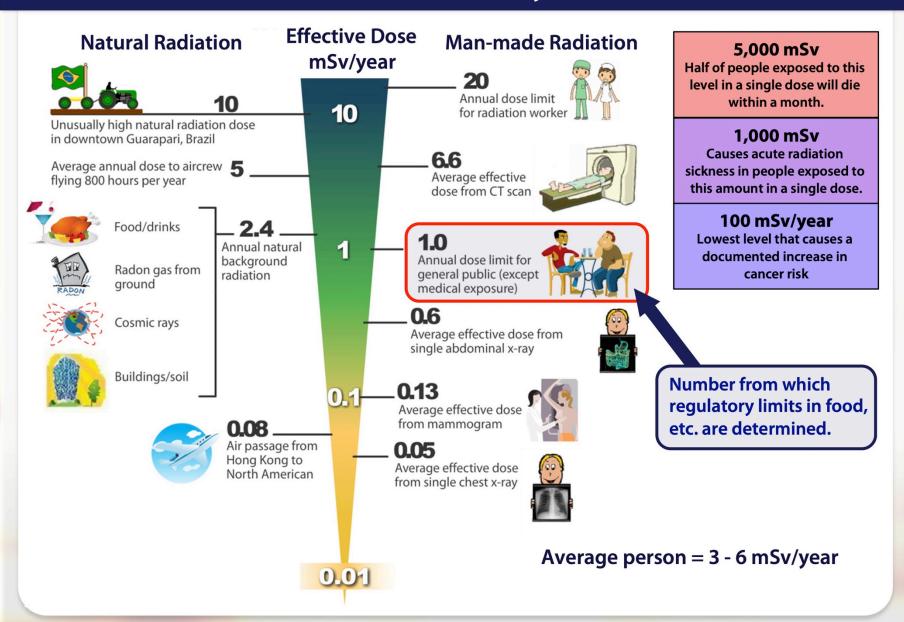
Radionuclide	t <sub>½</sub> (phys)	Part of the body considered	t <sub>½</sub> (eff)
<sup>3</sup> H	12.32 y	<b>Body Tissue</b>	12 d
<sup>14</sup> C	5730 y	Fat	12 d
<sup>32</sup> <b>p</b>	14.3 d	<b>Gastrointestinal Tract</b>	14 d
<sup>35</sup> <b>S</b>	87.5 d	Testis	76 d
<sup>60</sup> Co	5.27 y	<b>Gastrointestinal Tract</b>	0.75 d
129	$1.57 \times 10^3 \text{ y}$	Thyroid	140 d
131	8.02 d	Thyroid	7.6 d
<sup>137</sup> Cs	30.17 y	Total	70 d
<sup>210</sup> Po	138.4 d	Spleen	42 d
<sup>222</sup> Rn	3.8 d	Lung	3.8 d
<sup>226</sup> Ra	1600 y	Bone	44 y
<sup>238</sup> U	4.47 x 10 <sup>9</sup> y	Lung, Kidney	15 d
<sup>238</sup> Pu	$2.41 \times 10^4 \text{ y}$	Bone	64 y
<sup>241</sup> Am	432.2 y	Kidney	64 y

Radiotoxicity	Radionuclides and radioelements
Group I: Very High	<sup>90</sup> Sr, <sup>226</sup> Ra, <sup>238</sup> Pu
Group II: High	<sup>55</sup> Fe, <sup>210</sup> Bi, <sup>210</sup> Po
Group III: Medium	<sup>3</sup> H, <sup>14</sup> C, <sup>32</sup> P, <sup>35</sup> S, <sup>137</sup> Cs
Group IV: Low	<sup>42</sup> K, <sup>64</sup> Cu, <sup>85</sup> Kr

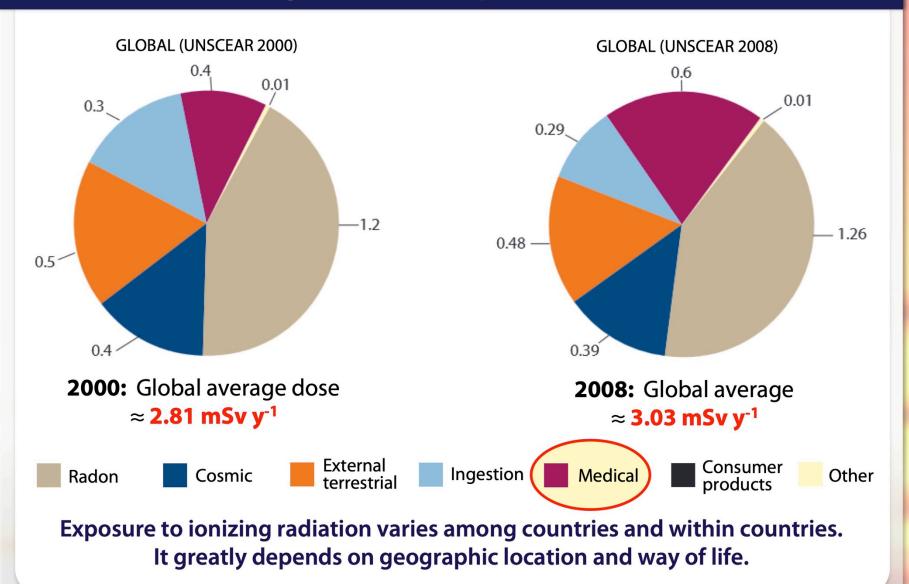
Depends on: Radiation emitted, mode of intake, amount, chemical properties/metabolic affinity, and the effective t<sub>1/2</sub>

Selected data from Leiser (2008) as adapted from ICRP (1993)

## Radiation in daily life



## Worldwide average radiation exposure from various sources



## **Environmental Protection**

# **Ecological risk assessment and management**



# 4. Radiological protection and the environment: Changing perspectives from anthropocentric to ecocentric

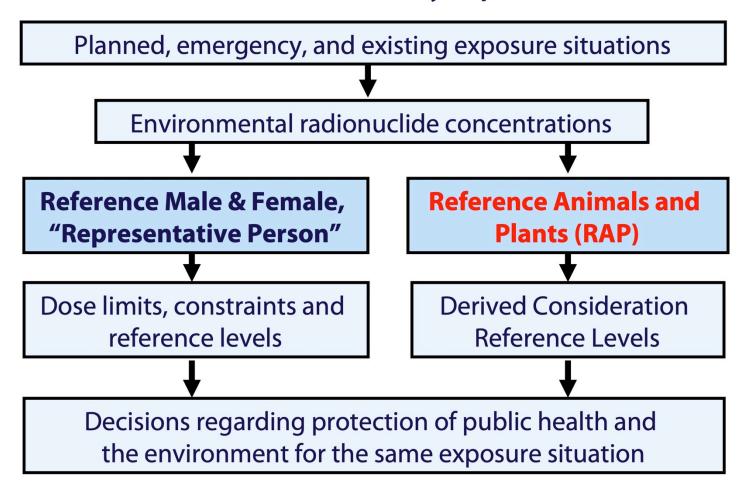
Initial focus was on humankind: It was considered that "if man is adequately protected, then other living things are also likely to be sufficiently protected." (ICRP, 1977)

Beginning in the late 1980s this assumption has been questioned. Due to increasing regulations regarding the protection of the environment, specific criteria for species other than man has reached a consensus by the International Community.

ICRP (2008)

## Similarity in approaches: Humans versus the Environment

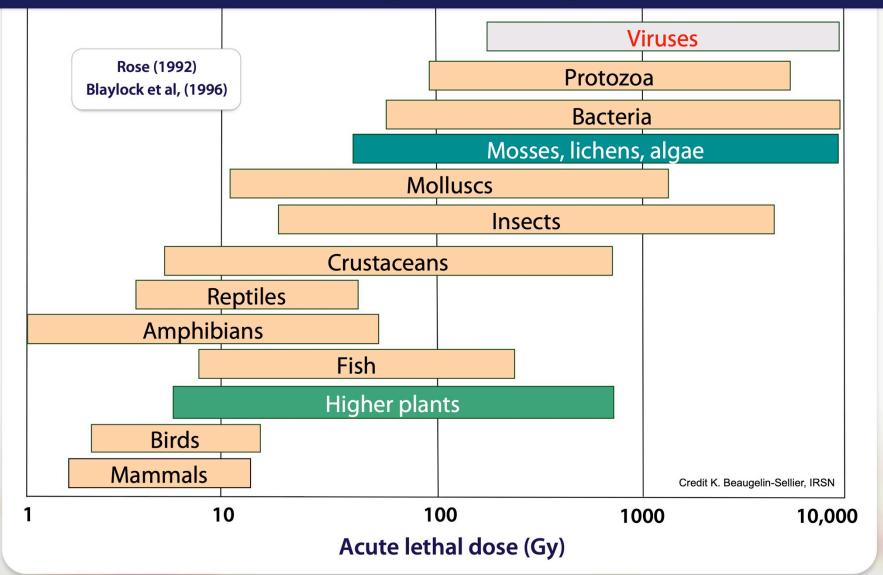
Schematic approach to the protection of both the public and the environment in relation to any exposure situation.



ICRP (2008. 2009); Pentreath (2009)

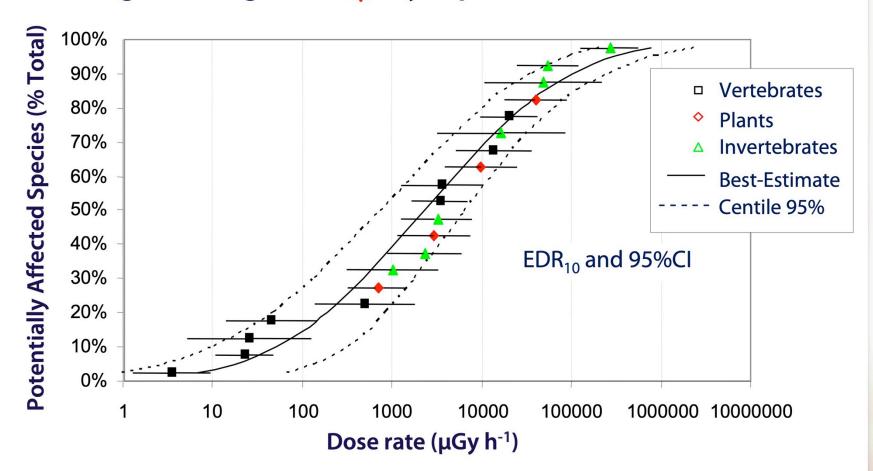


# What we know regarding the effects on organisms: Acute ionizing radiation exposure



## **Chronic Exposure**

# Species Sensitivity Distribution for a generic ecosystem and assuming chronic gamma ( $\gamma$ ) ray exposure



Garnier-Laplace et al. (2006, 2010)



## Radiological protection and the environment

- UNSCEAR (2008): Analyses of available data concludes that chronic irradiation at dose rates up to 400 μGy h<sup>-1</sup> to a small proportion of individuals in an aquatic population results in no detrimental effects at the population level
- ICRP (2008): Defines DCRLs (i.e., a range of dose rates within which it is likely to observe some deleterious effects) for a given Reference Animal or Plant (RAP).

Marine RAPs: Flatfish and seaweed: 42-420 μGy h<sup>-1</sup>

Crabs: 420-4200 **μGy h**<sup>-1</sup>

Various European Union programs have agreed on a **conservative Predicted No Effect Dose-Rate** of **10 \muGy h**<sup>-1</sup>, which is ultimately derived from the Species Sensitivity Distributions (see previous slide).

#### The ERICA tool

# The 1<sup>st</sup> European answer towards demonstration of environmental protection

**Environmental Risk from Ionising Contaminants: Assessment and Management** 

- The outcome of a suite of European research programs
- 2004-2007 (first release)

ERICA: Suggested screening benchmark: 10 μGy h<sup>-1</sup> (87.6 mGy y<sup>-1</sup>) to protect the structure and function of a generic ecosystem

#### **Development of the ERICA tool:**

- A free software approach to assessing the radiological risk to terrestrial, freshwater and marine <u>biota</u>
- Based on a reference organism approach based on biological effects on individual marine biota
- Updated in 2008, 2009, 2011, 2012, 2014 and 2016
- Website: <a href="https://wiki.ceh.ac.uk/display/rpemain/ERICA+Tool">https://wiki.ceh.ac.uk/display/rpemain/ERICA+Tool</a>

## **Summary**

# Radionuclides from a variety of sources are bioaccumulated by marine organisms

#### Bioaccumulation depends on:

- The marine organism (Species, trophic level...)
- The radionuclide (physical-chemical forms ...)
- Environmental parameters (temperature, salinity...)

Marine biota transport radionuclides horizontally (migration) and vertically (migration, feces). Some radionuclides may accumulate in higher trophic levels via grazing.

# Shark (fish-eating fish) Tuna (fish-eating fish) Herring (plankton-eating fish) Squid Fish larvae (zooplankton) Conchs (zooplankton) Diatoms (phytoplankton) Dinoflagellates (phytoplankton)

#### **Future Directions:**

- There is very little information on radionuclide uptake by plankton communities or in top predators, including mammals.
- Most of the data is derived from temperate climates. Larger geographical studies are needed.
- Comprehensive studies on radionuclide transfer in a food web are uncommon.

## Summary

# Radionuclides from a variety of sources are bioaccumulated by marine organisms

Doses to humans and biota *mainly* arise from natural sources.

Since 2000, the emphasis has been on the development of specific criteria for environmental protection.

#### **Future Directions:**

- More data is needed regarding the radiological chronic exposure effects on marine biota
- Refinement of dose-response relationships under natural versus laboratory conditions
- Environmental Risk Assessment needs to move from individual species to ecosystem (e.g., the impact on biodiversity index, trophic network structure, etc.)

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