

# Doses from tritium



EU Seminar, November 2007

Effective dose

Dose coefficients for HTO and OBT

RBE, DDREF and  $w_R$

OBT in Cardiff Bay fish

## A protection device

Allows summation of doses from different radionuclides and external dose

Use for regulatory purposes for comparison with dose limits / constraints

Relates to stochastic effects only

→ total detriment of  $7\% \text{ Sv}^{-1}$  (6% in new recommendations)

## Not individual specific

- reference biokinetic and dosimetric models, and defined  $w_R$  and  $w_T$  values, are used to calculate reference dose coefficients for protection purposes

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## Use of individual monitoring data

- better assessment of intake / exposure for the calculation of effective dose to a reference person

# Calculation of equivalent and effective dose



1. Absorbed dose in tissues / organs, Gy

2. Equivalent dose in tissues / organs, Sv

- Gy x  $w_R$

$$H_T = \sum_R w_R D_{T,R}$$

3. Effective dose, Sv

- equivalent dose x  $w_T$

$$E = \sum_T w_T H_T$$

# Weighting factors



Equivalent dose in tissues / organs, Sv

$w_R$	1	for low LET radiations
	20	for $\alpha$ particles

Effective dose, Sv

$w_T$	0.01	bone surface, skin
	0.05	bladder, breast, liver, oesophagus, thyroid remainder
	0.12	bone marrow, colon, lung, stomach
	0.2	gonads

# Tissue weighting factors



<i>Current</i>	0.01	bone surface, skin
	0.05	bladder, <b>breast</b> , liver, oesophagus, thyroid, <b>remainder</b>
	0.12	bone marrow, colon, lung, stomach
	0.2	<b>gonads</b>
<i>New</i>	0.01	bone surface, skin, brain, salivary glands
	0.04	bladder, liver, oesophagus, thyroid
	0.08	<b>gonads</b>
	0.12	bone marrow, colon, lung, stomach, <b>breast, remainder</b>



# Sex-specific detriment



Applying to ages from 0 - 85 at exposure

Overall detriment is about 40% greater in females than males

Differences include:

Colon                    x 0.4 in females cf. males

Liver                    x 0.5

Lung                    x 2.0

Thyroid                x 4.3

Breast                one-quarter of total detriment in females

# Age-specific cancer risks

- *life-time attributable risk*

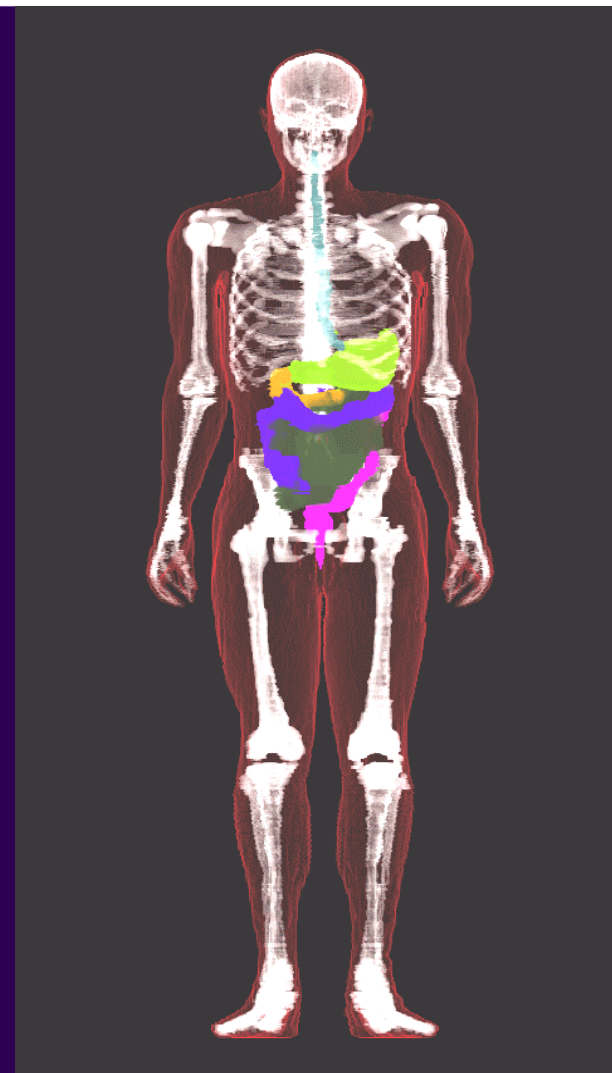
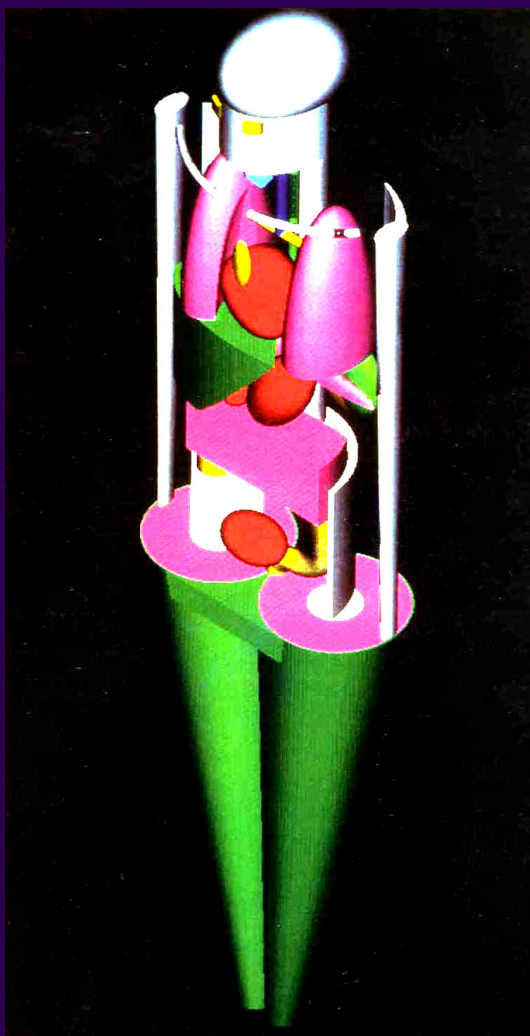


Cases per 10<sup>6</sup> exposed to a single dose of 10 mGy (BEIR VII)

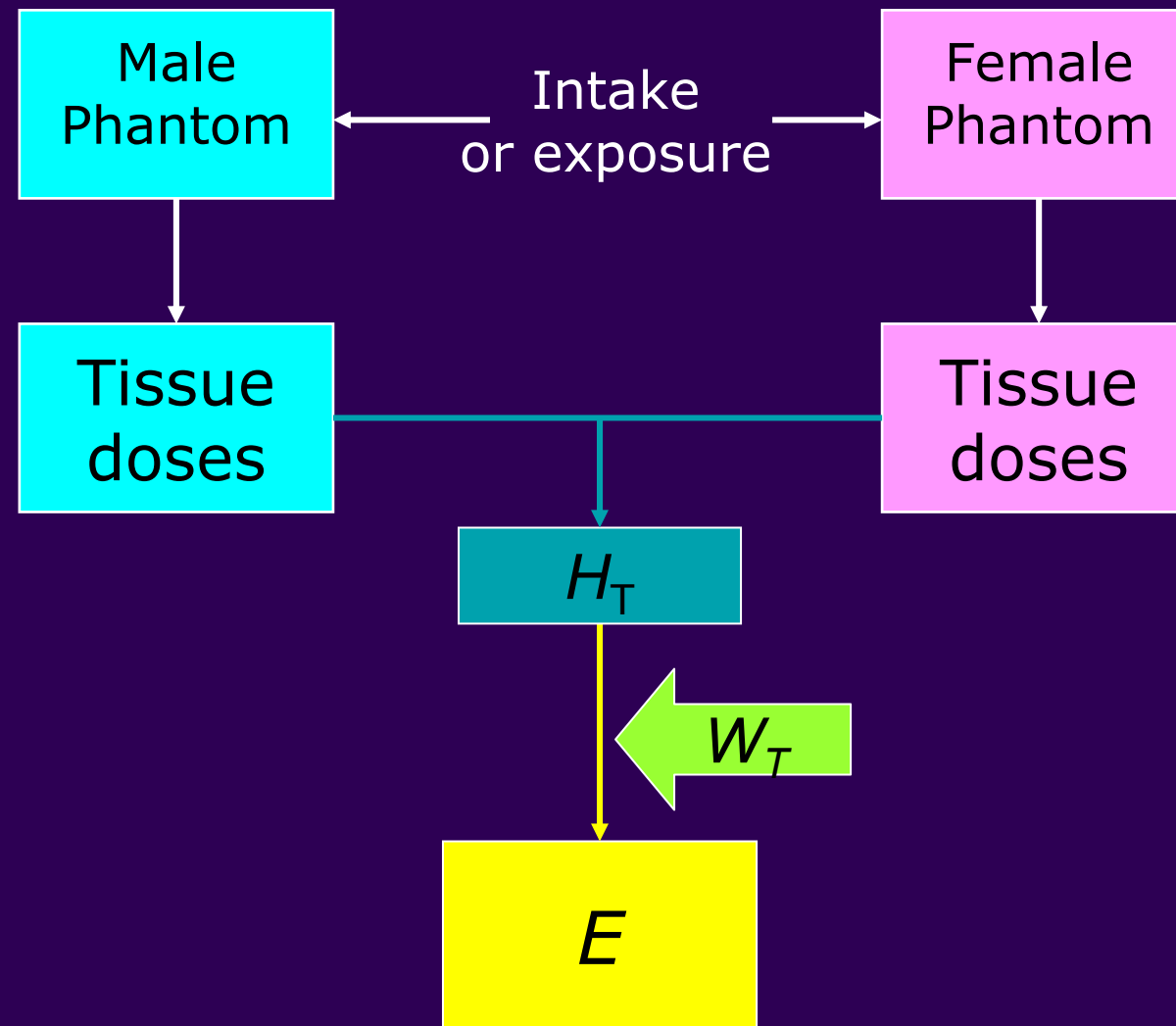
Cancer site	Age at exposure, years					
	Males			Females		
	0	20	60	0	20	60
Breast	-	-	-	1171	429	31
Colon	336	173	94	220	114	62
Liver	61	30	14	28	14	7
Lung	314	149	89	733	346	201
Thyroid	115	21	0.3	634	113	1
Leukaemia	237	96	82	185	71	57
All cancers	2563	977	489	4777	1646	586

# Dosimetric phantoms

- MIRD and Voxel



# Sex averaging in calculation of Effective Dose



# Forms of tritium



Tritiated Water ( HTO )

Organically-bound Tritium ( OBT )

- Non-exchangeable binding of  $^3\text{H}$  to C atoms in organic molecules

Includes  $^3\text{H}$  in carbohydrates

lipids

proteins

nucleic acids

# ICRP biokinetic assumptions



HTO, OBT ingestion, inhalation as vapour

Absorption to blood = 100%

Uniform distribution in all body tissues

Two components of retention, corresponding to HTO (A) and OBT (B) in body tissues

	A,%	B,%
Intakes as HTO	97	3
OBT	50	50

# ICRP biokinetic assumptions - retention half-times



Age	Half-time, d	
	A	B
3 months	3.0	8
1 year	3.5	15
5 years	4.6	19
10 years	5.7	26
15 years	7.9	32
Adult	10.0	40

# ICRP tritium ingestion dose coefficients



Age	Committed effective dose Sv Bq <sup>-1</sup> x 10 <sup>11</sup>	
	HTO	OBT
3 months	6.3	12
1 year	4.8	12
5 years	3.0	7.3
10 years	2.3	5.7
15 years	1.8	4.2
Adult	1.8	4.2



# Proposed ICRP assumptions - occupational intakes

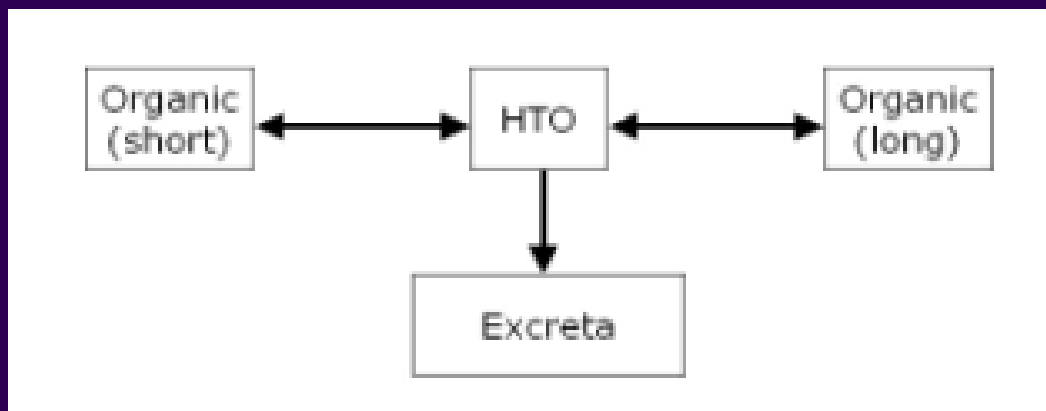


## Dose coefficients and interpretation of bioassay

*Taylor ; Rad. Prot. Dosim. 105, 225 ; 2003*

Intakes of **HTO**: 99% 10d, 0.98% 40d, 0.02% 350d

Dose coefficient:  $1.7 \times 10^{-11}$  Sv Bq<sup>-1</sup>



# Comparative doses to adult members of the public



Radionuclide	Comparative doses	
	Ingestion	Inhalation
Tritium (HTO)	1	1
Carbon-14	30	30
Strontium-90	1,600	1,300
Iodine-131	1,200	400
Caesium-137	720	260
Plutonium-239	14,000	2,800,000

# ICRP dose coefficients - fetus / infant



Ingestion during pregnancy and lactation

Radionuclide	Ratio of offspring : adult dose	
	Pregnancy	Lactation
Tritium (HTO)	1.7	1.2
Phosphorus-32	10	0.7
Calcium-45	11	2.7
Strontium-90	1.5	0.6
Iodine-131	1.1	2.4
Caesium-137	0.4	0.3
Polonium-210	0.1	0.4
Plutonium-239	0.04	0.0004

# Uncertainties in data used in tritium dose coefficients



*Harrison, Khursheed, Lambert; Rad. Prot. Dosim. 98, 299; 2002.*

5 - 95% range for uncertainty on central values  
for population groups

Absorption to blood

Incorporation into OBT in body tissues

Retention half-times

Relative biological effectiveness ( RBE )

# Uncertainties in tritium dose coefficients



## Probability Distributions on Ingestion Dose Coefficients for Adults, Sv Bq<sup>-1</sup> x 10<sup>11</sup>

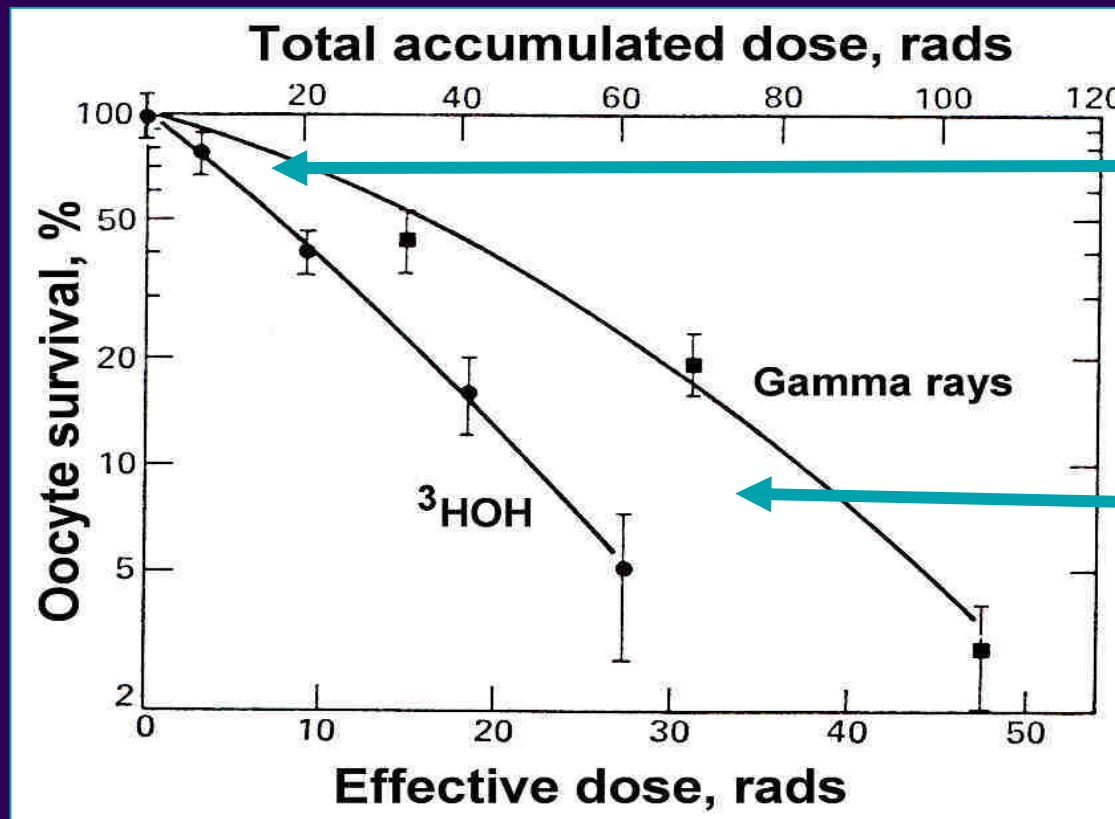
Form	Range, %			ICRP
	5	50	95	
HTO	2.1	3.9	6.6	1.8
OBT	3.9	8.7	20.0	4.2

*Strictly, Gy Bq<sup>-1</sup> x RBE*

# *In utero / germ cell sensitivity*



HTO ingestion /  $^{60}\text{Co}$  gamma ray exposure of mice throughout pregnancy and 14d lactation - oocyte survival in offspring

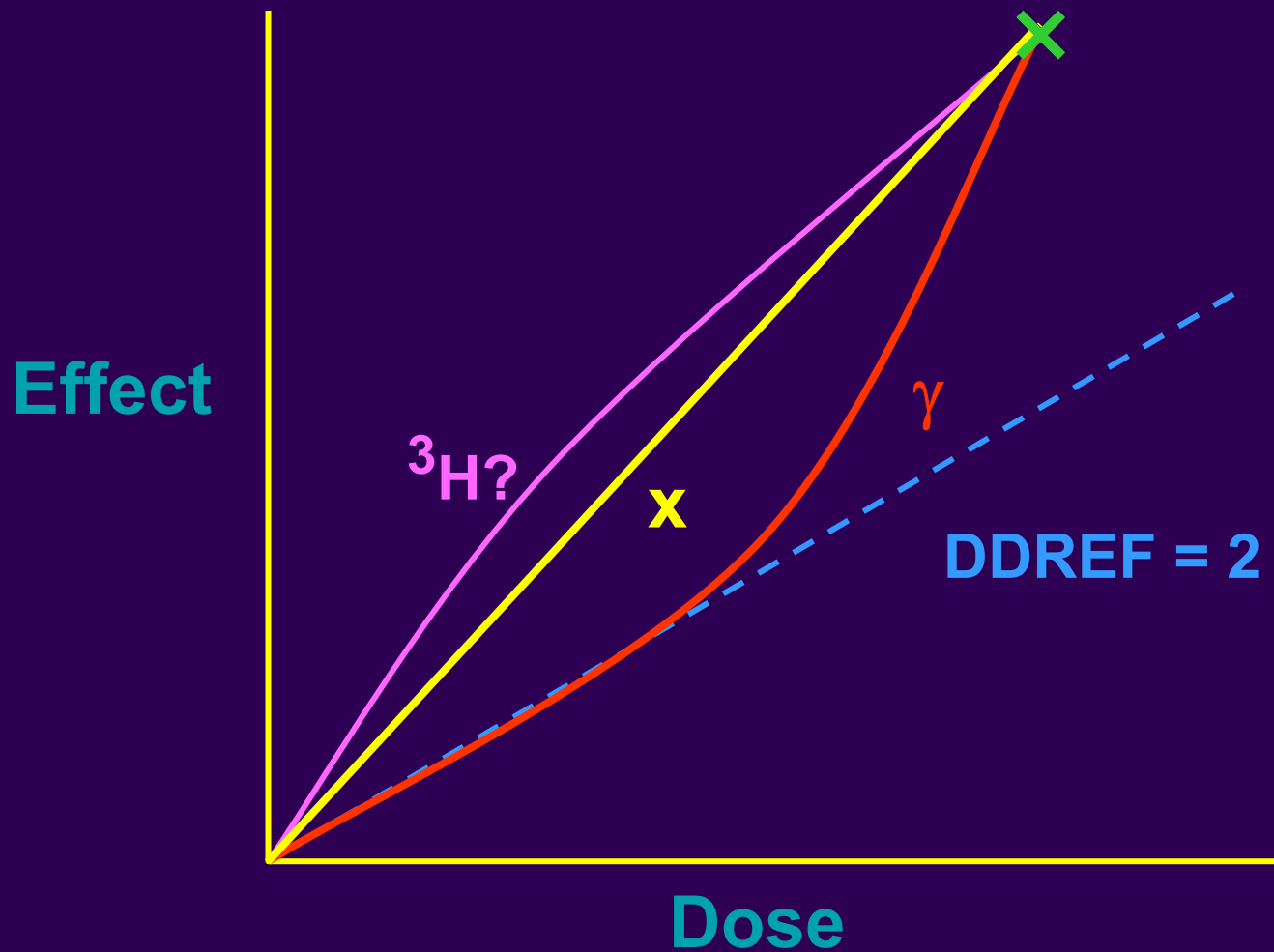


$\text{RBE}_M = 3$

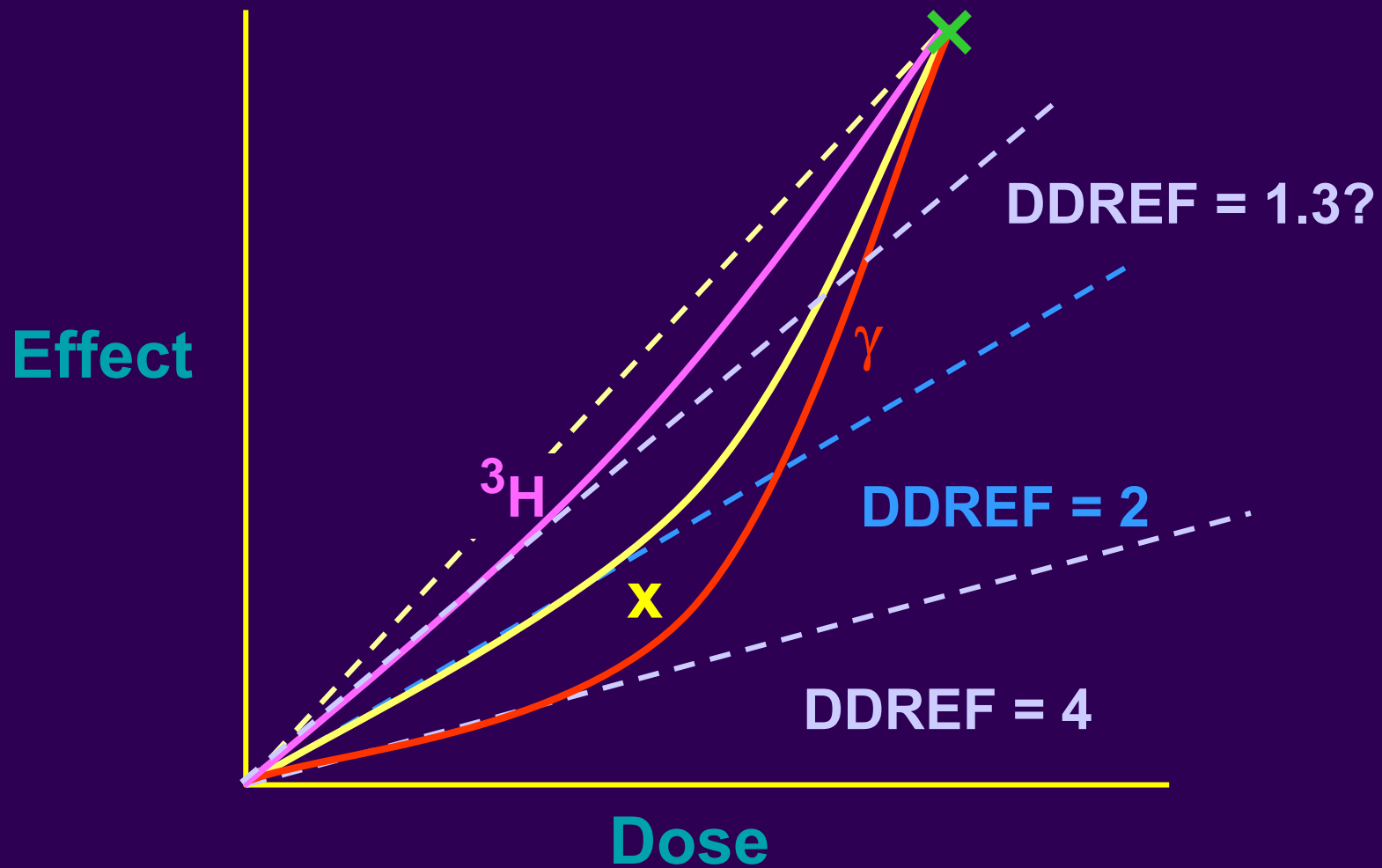
$\text{RBE} = 1.6$

*Dobson and Kwan  
Rad. Res. 66:615; 1976*

# DDREF and RBE



# DDREF and RBE





High tritium RBEs in systems with high DDREF

DDREF of 2 for cancer in humans implies RBE of  $\leq 2$

More complex treatment of  $w_R$  for low LET radiations ?

- inconsistent with intended use of effective dose
- implies greater knowledge of risk at low doses than is justified?

# OBT in Cardiff Bay fish



Whole body retention of tritium in rats after ingestion of freeze-dried flounder or HTO

## Rat data

HTO	0.97	3d	0.03	10d
Fish OBT	0.7	3d	0.3	25d

## Human

					Sv Bq <sup>-1</sup> x 10 <sup>11</sup>
HTO	0.97	10d	0.03	40d	1.8
OBT	0.5	10d	0.5	40d	4.2
Fish OBT	0.7	10d	0.3	100d	6.1

*Hodgson et al. J. Radiol. Prot. 25, 149; 2005*

# Conclusions



- Effective dose is a protection tool
- Metabolism of HTO understood  
OBT treated generically
- Uncertainties relatively small
- $RBE \leq DDREF$  ?
- Specific OBT may differ from ICRP