The 2007 Recommendations of ICRP Dr Jack Valentin, Scientific Secretary, ICRP

• International Commission on Radiological Protection ICRP: Who, why, what?

• The 2007 Recommendations

Justification (political) – optimisation – limits & constraints The exposure situation Include non-human species

About ICRP



ICRP, an Independent Registered Charity

Established to advance for the public benefit the science of Radiological Protection,

in particular by providing recommendations and guidance on all aspects of protection against ionising radiation.

<u>Structure of ICRP, 2005 – 2009</u>

Main Commission

Chair: Dr L-E Holm, SE

12 other members



Scientific Secretariat Dr J Valentin, SE

C1- Radiation Effects *Dr R J Preston*, *US*

C2- Doses from Radiation Exposure *Dr H Menzel, CH*

C3- Protection in Medicine *Dr C Cousins, UK*

Task Groups

C4- Application of ICRP Recommend:s *Dr A Sugier, FR*

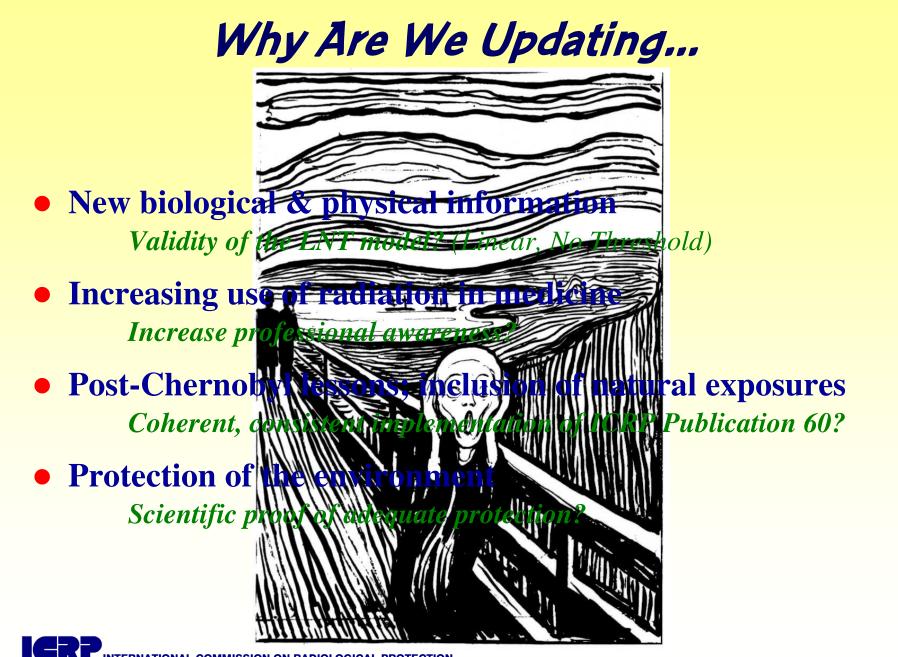


C5-Prot. of the Environment *Prof J Pentreath*, *UK*

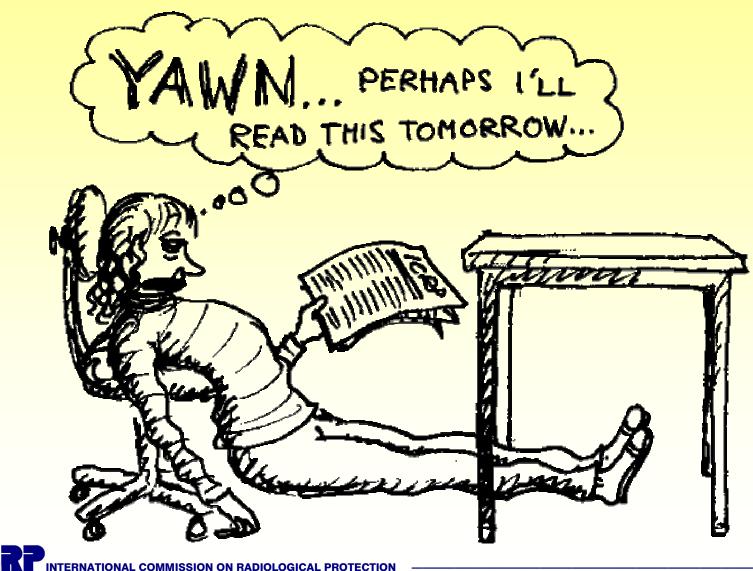
INTERNATIONAL COMMISSION ON RADIOLOGICAL PROTECTION

The 2007 Recommendations of ICRP





ICRP 1990 Rec's: Logical But Complex



Aims of the Revision

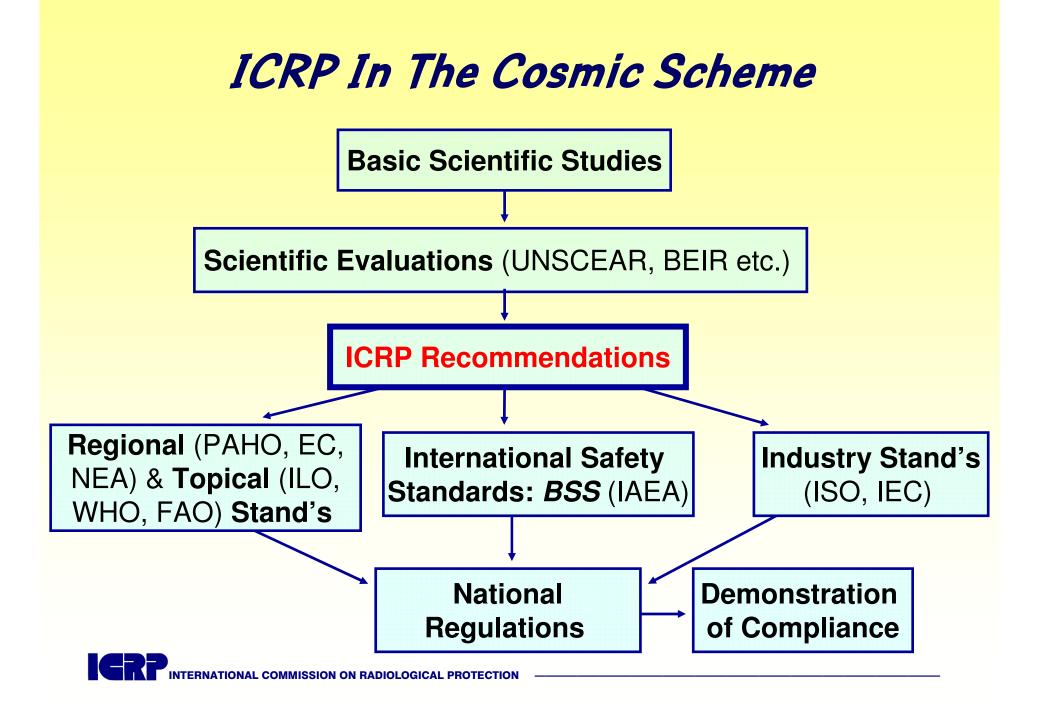
- Take account of new science
- Feed back experience of current radiation safety standards
- Improve & streamline the presentation
- Use an open, transparent process (9 years gestation!)
- Maintain as much stability as is consistent with the new information

To Get the Recommendations...

- **Buy printed or electronic copies** *IRPA Associated Societies are eligible for a discount*
- Developing countries: free download at HINARI
- Junior staff: coming summary in JRP
- For all of this, see www.icrp.org
- Or translate them (an Italian version is ready!)

Chapter 1: History, development, structure





Chapter 2: Aims and scope

Primary Aim of Our Recommendations

To *contribute to* an *appropriate* level of protection for people

and the environment

without *unduly* limiting the *desirable human activities* that may be associated with radiation exposure

The Principles of Protection

Source-related, in all exposure situations:

• Justification

More benefit than detriment

• Optimisation of protection Dose and risk constraints to (a) increase equity, (b) consider multiple sources

Individual-related, in *planned* exposure *situations*

• Application of dose limits Except medical exposure of patients

Utilitarian ethics Judge actions by the consequences	
Justification Do more good than harm	
Optimisation Maximise good > harm	

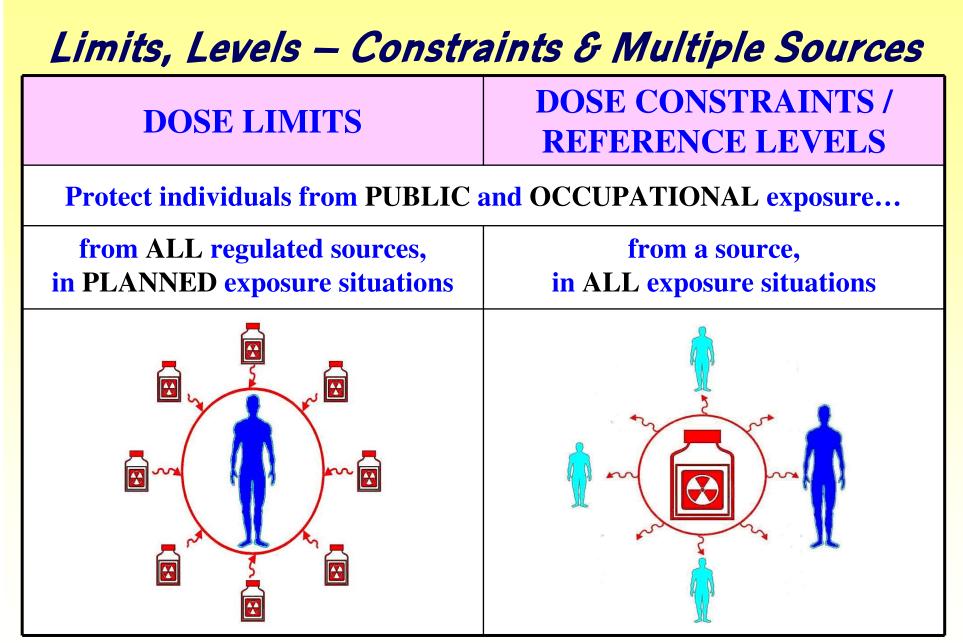
Utilitarian ethics	Deontological ethics
Judge actions by the consequences	Some duties are imperative
Justification	Limitation
Do more good than harm	No individual unduly harmed
Optimisation Maximise good > harm	

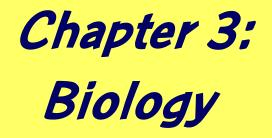
Utilitarian ethics	Deontological ethics
Judge actions by the consequences	Some duties are imperative
Justification	Limitation
Do more good than harm	No individual unduly harmed
Optimisation Maximise good > harm	Dose constraints Increased equity = emphasise the individual





Limits, Levels – Constraints & Multiple Sources **DOSE CONSTRAINTS / DOSE LIMITS REFERENCE LEVELS Protect individuals from PUBLIC and OCCUPATIONAL exposure...** from ALL regulated sources, in PLANNED exposure situations (\mathbf{x})







Deterministic and Stochastic Effects

These words will still be the default terms!

• Deterministic

Harmful, mostly late, tissue reactions

• Stochastic

Cancer and heritable disease

Cancer probability now based on incidence, not mortality LNT: Scientifically plausible but not unambiguous (cf. Central Limit Theorem, i.e., valid at the population level)

Heritable Disease

- Induced mutation rates: based on mouse studies Induced genetic effects not demonstrable in man! Human spontaneous mutation rates used to estimate Doubling Dose
- **Probability of heritable risk was over-estimated in 1990** *Particularly for multifactorial diseases*
- Nominal probability coeff/s: 2 generations only Based on UNSCEAR 2001, agrees with BEIR VII
 1990 calculation to equilibrium – assumptions not sustainable Risk after 2 generations small, no substantial difference 2 – 10 generations

Thus, no significant underestimation of genetic risk

Epigenetic Responses to Radiation

• Genomic instability: Damage expressed after several cell generations

Why and how does it happen? Does it really affect normal cells? If it does, does it change the total risk assessed epidemiologically?

• Bystander signalling: Damage to non-irradiated cells in an irradiated cell population

Why and how does it happen? Does it change the total risk assessed epidemiologically?

• Important for understanding - currently no way to assess effect on risk – major effect unlikely

Adaptive Response to Radiation

- A priming dose sometimes confers increased resistance against a second dose
 - Not a universal feature
 - Considerable variation, usually transient
 - Mechanistic knowledge fragmentary
 - Not evident at 'protection' doses
 - No consistent evidence of reduction of adverse health effects
- Thus, scientifically important but at present not relevant for radiological protection

Females and Males

Buzzardbros.com

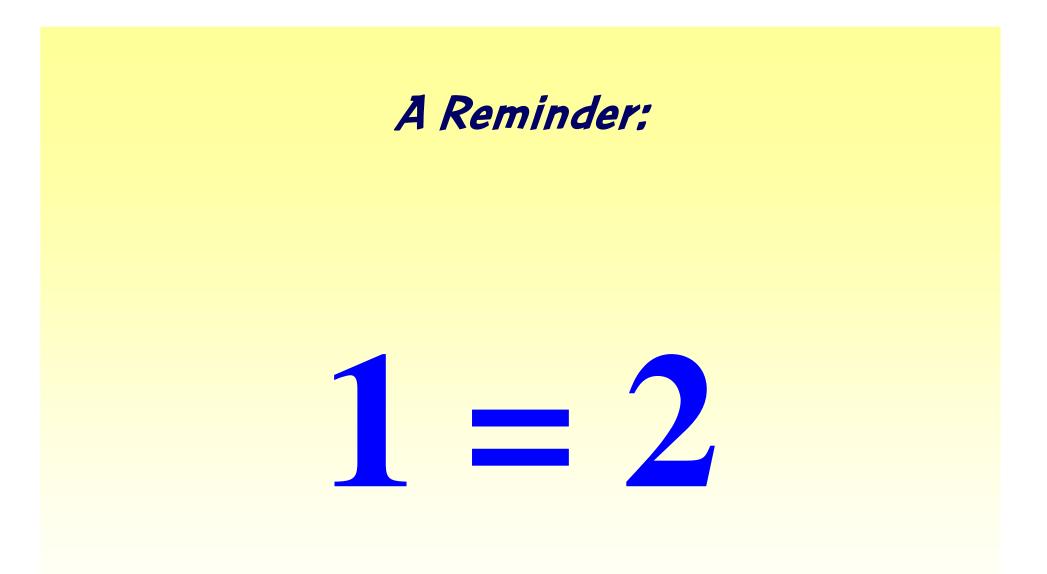
Nominal risk estimates for protection Individual retrospective assessments require specific information

the average achieves adequate protection for both sexes A value judgement, based on science

EVERY DAY I LIVE WITH FEAR BUT OCCASIONALLY I LEAVE HER AND GO Regluces discrimination DRAG RACING

Nominal Probability Coefficients (% Sv⁻¹)

Exposed popula- tion	Cai	ncer		table ects	То	tal
		ti çal ,p				,2007
the Whole	6.0	l risk (5.5 still ap	1.3	0.2	~5% 7.3	5.7
Adult	4.8	4.1	0.8	0.1	5.6	4.2





Does ICRP Over- Or Underestimate Risk?





Chapter 4: Physical quantities

Radiation Weighting Factors, w_R

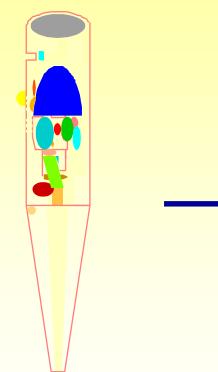
Type and energy range	Publication 60	2007	
Photons, all energies	1	1	
Electrons and muons, all energies	1	1	
Protons	5	2	
Alpha particles, fission fragments, heavy nuclei	20	20	
Neutrons	Stepwise function	Continuous function	
		<10 keV, 2.5	

Tissue Weighting Factors, w_T

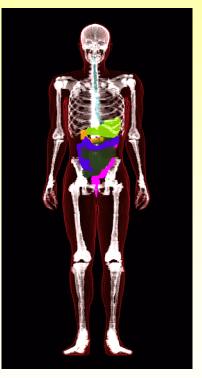
Tissue	w _T	∑ ₩ _T
Bone-marrow, breast, colon, lung, stomach, remainder tissues (13/14)	0.12	0.72
Gonads	0.08	0.08
Bladder, oesophagus, liver, thyroid	0.04	0.16
Bone surface, brain, salivary glands, skin	0.01	0.04

New Reference Phantoms

MIRD Phantom Voxel Male and Female Phantoms



Fetus; child: in preparation



New dose coefficients in 2008 🙂

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The Use of Effective Dose (E)

- For compliance and prospective planning
- Not for detailed retrospective dose and risk assessments after exposure of individuals particularly not for patients (old, unhealthy population)
- Not for epidemiological studies (at least not for risk assessment)

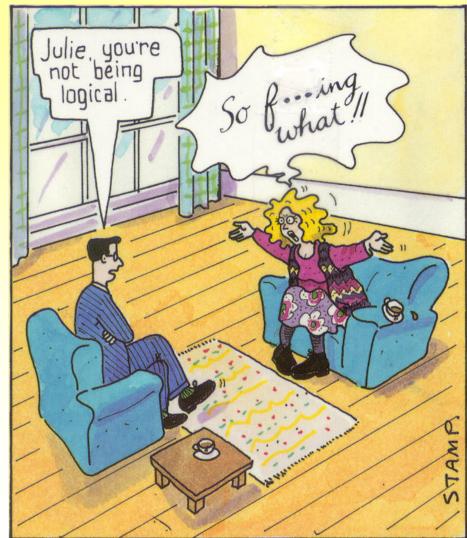
The Use of Collective Dose (S)

- For optimisation
- For comparing technologies and protection options
- Not for epidemiologic risk assessment Inappropriate to use it in risk projections based on epidemiology
- Not for predicting number of cancer deaths due to trivial exposures to large populations An unreasonable, unintended, incorrect use of collective dose

Collective Dose: Logical, But Is It Right?

Equates many small doses to few large doses...

Are 500 road traffic casualties just as bad as 500 plane crash victims?



Chapter 5: System of protection, man



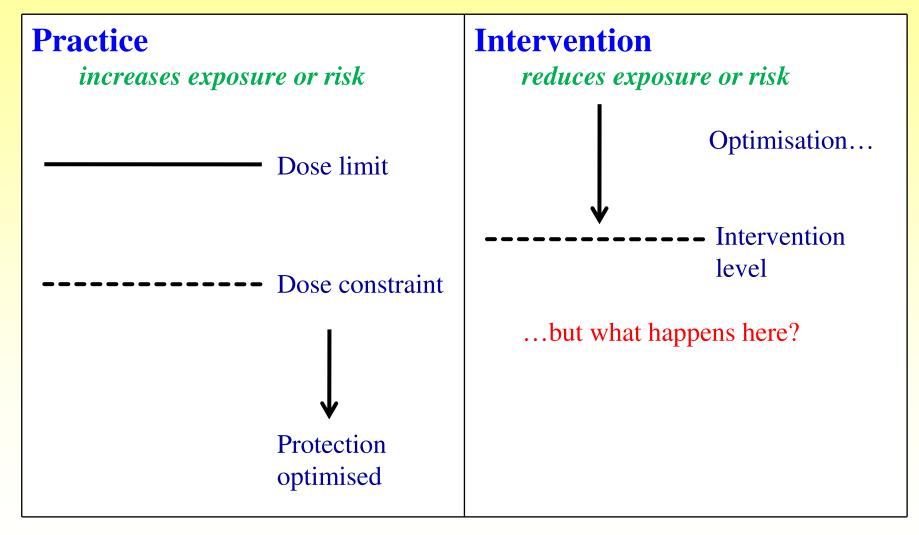
In 1990, a Process-Based Approach

Practice Intervention reduces exposure or risk increases exposure or risk

In 1990, a Process-Based Approach

Practice increases exposure or risk	Intervention reduces exposure or risk
Dose limit	
Dose constraint	
Protection optimised	

In 1990, a Process-Based Approach



2007, Exposure Situation: Planned / Emergency / Existing



Inappropriate to plan to allow higher exposures

How to Select Constraints / Reference Levels

• 100 – 20 mSv

Direct benefit. Information, training, dose monitoring. Example: Radiological emergencies

• 20 – 1 mSv

Direct or indirect benefit. Information, training, dose monitoring or assessment. Examples: Occupational exposures in planned situations, radon in dwellings

• Less than 1 mSv

Societal benefit. Dose assessment. Example: Public exposures in planned situations

The Collective Dose in Optimisation

- A key parameter (at least in occupational protection), but we usually also need to know
 - average dose, number exposed, range, etc
- Perhaps give more weight to
 - a few large doses than to many small doses
 - doses now than to doses in the far future

Dose Limits for Planned Exposure Situations

• They remain the same as in 1990!

PUBLIC	OCCUPATIONAL			
1 mSv in a year	20 mSv per year, averaged over defined 5-year periods			
In special circumstances, an average of 1 mSv per year	100 mSv in 5 years, and less than 50 mSv in one year			
averaged over defined 5-year periods				

Chapter 6: Implementation

Planned Exposure Situations

Occupational exposure

Constraint usually set by operator (small operators may need guidance) Transient/itinerant workers need special attention

• Public exposure

Constraints usually set by regulator About 0.3 mSv in a year appropriate 0.1 mSv in a year if prolonged exposure

Potential Exposures

- Workplace accidents
 - Number of people affected is speal Letriment = health rise to those directly expo
- Large disasters Nymber of people affected on the brige Detriment also includes contominated long, food restrictions, etc
- Exposures in the far future, e.g. from waste repositories Considerable uncertainties Dose calculations useful to compare protection options but not to project detriment

Assessment of Potential Exposures

- Everybody is responsible for safety, incl. security Particularly important to remember outside the nuclear fuel cycle
- Risk constraints: guide optimisation of protection against risk (probability of death) = Prob (accident) * Prob (death | accident dose)
- ICRP continues to recommend established generic constraints:

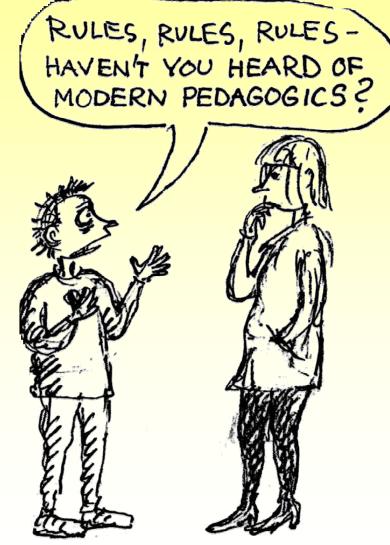
Potential exposure of workers: 2 10⁻⁴ per year Potential exposure of the public: 1 10⁻⁵ per year

Existing Exposure Situations: Radon

• Upper level of dose: 10 mSv (radon/progeny equilibrium) Upper level of activity conc. retained for continuity National regulators can [should] set lower constraints 'There is now evidence for what we thought all the time' No real difference UNSCEAR-ICRP; an ICRP TG statement in preparation

Situation	Reference level
Domestic dwellings	600 Bq m ⁻³
Workplaces	1500 Bq m ⁻³

Regulatory Philosophy





Chapter 7:

Medical exposure of patients



Special Features of the System of Protection

• Justification in medicine

Benefit and risk apply to the same person (patient)

• Optimisation in medicine

Diagnostic Reference Levels, not constraints Radiation therapy, maximise PTV but minimise other dose

 Dose limits Do not apply to patients

Justification or Indiscriminate Referral



'Buy Our CT, Earn \$ 2,163,000 in 5 Years'



Minimum space requirement is 175 sq. ft.

Many private practices have imited space available. However, most physicians are surprised at how fitte space is actually needed to site a CT in their efficient. The declarad Sparents in office CT scanters are do signed with the smallest possible footprint and have easy siting equilements— no matter where your practice is located.

Sitting Both the SOMATOM Spirit and the SOMATOM Emotion are satting the industry benchmark with their compact designs and small footprints (as small as 17.5 square foot, making them idea to she in a private practice. Both systems are air cosed, which eliminates the necessity for an exits water chierer typicary found with other CT products. The actual instantion can be done easily within one or two days.

A Siement project manager will be glad to visit your site and assess whether the selected CT system can be installed in a room or power modifications are necessary. Literally any existing room can be modified to meet the criteria of a CT room and most medical buildings are ady have the necessary power supply.

Quick Check#3: Potential CT room available? Yes No What is the size of that room/area in square feet?

Take a manyert to rock around Bo you have a room or space where you can envision your new CT scanner? In-office CT can be a significant new source of practice revenue, Let us show you how.

Siemens Financial Services offers you the textibility to easily finance or wate-your CT equipment as part of our one-stop shopping toution. Your Siemens thrandal analyst will provide you with a customized business por torma, based on your actual practice numbers... the of charge.

Affordability Both the SOMATOM Splitt and the SOMATOM Endotes an uniquely affordable solution. The SOMATOM Split offers a new level of cost-left othernes is aching the beakeven point faster than conventional scanters, thus machineling your return on investment. This gives practices with initiad budgets and low patient volume the opportunity to invest in their own CT bechnology. That's affordability

The SOMATOM Emotion combines the best of bothwords, clinical performance and low life cycle cost, to make it an cutotanding investment for mid-site and large practices with a higher patient throughput or broad patient examination mits

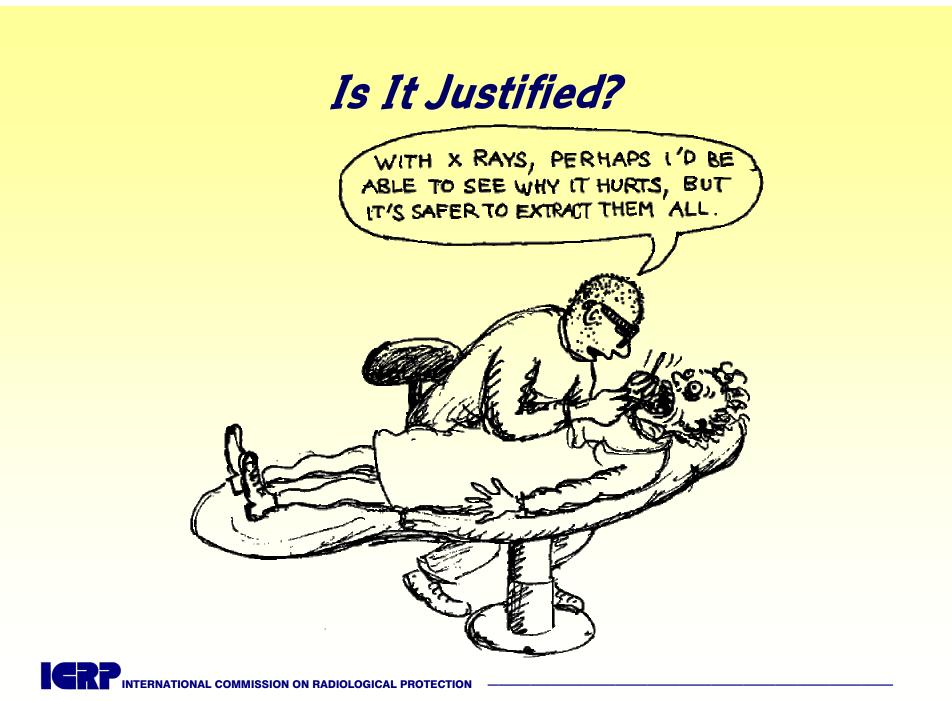
Our CT system prices include: Shipping, Installation, Resting one-year warranty and our unique Lile Customer Care Services such as Application Training, Application Hot-Line Support, and much more

	Procedure s Per Day	Days Per Mosth	Average CPT	income	FMVL Cost	ROI" Per Month	RDI for 5 Yests	
•	1.0	20	\$220	\$7,950	\$7,950	Break Even	Break Even	
	5	20	\$220	\$22,000	\$7,950	\$14,050	\$943,000	
	10	20					\$2,163,000	
	17	vertation vertation vertation	- Realt SC Inter Career Ity our Sile It	MATCH S	Spirit con	ilgaration ary Will	cosed on o	
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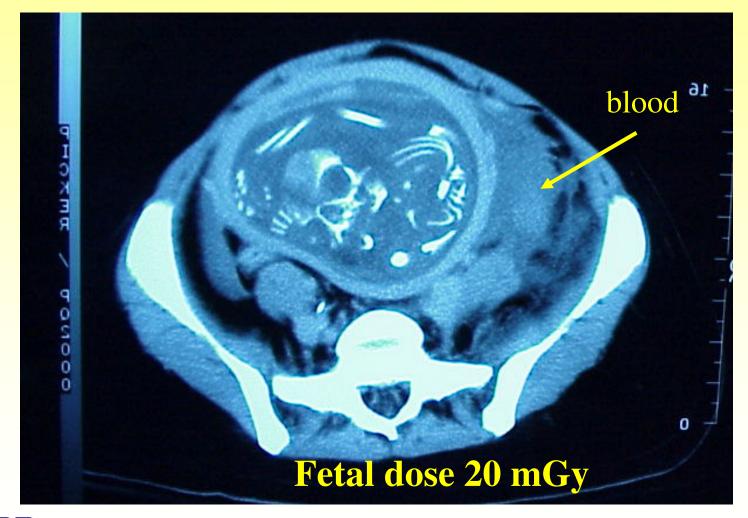
We here you to concurate your bottom rine. Let us know your investment needs.

	Procedures Per Day	Days Per Month	Average CPT	Income	FMVL Cost	ROI* Per Month	ROI for 5 Years
A	1.8	20	\$220	\$7,950	\$7,950	Break Even	Break Even
В	5	20	\$220	\$22,000	\$7,950	\$14,050	\$843,000
с	10	20	\$220	\$44,000	\$7,950	\$36,050	\$2,163,000

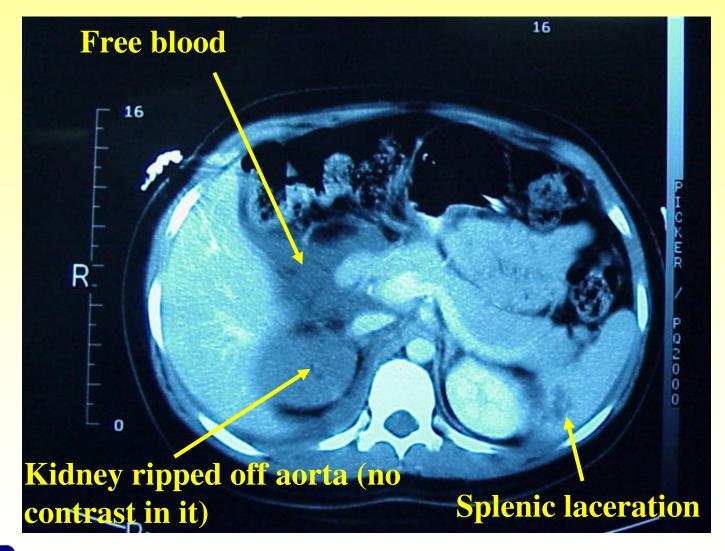
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Justification in Medicine...



...3 Min. Exam, Then OR (both survived!)



Some Medical Is Justified – But...





Chapter 8: Protection of the environment



Why Protect Other Species?



Why Protect Other Species?

- *NOT* driven by concerns of existing radiation hazards
- Fills a conceptual gap Science to show if the environment is adequately protected - and methods to improve protection if required
- Further guidance will be provided

To Summarise, ICRP is...

- Retaining the fundamental principles of protection
- Clarifying how they apply to sources and the individual
- Changing focus from process (practice/intervention) to exposure situation (planned/emergency/ existing)
- Extending the concept of source-related constraints to all situations
- Updating weighting factors and detriment
- Maintaining the current dose limits

2007 Recommendations

