Dosimetry of dental CBCT

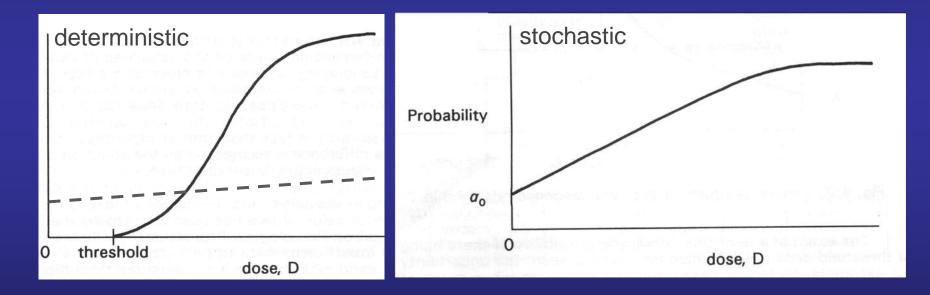






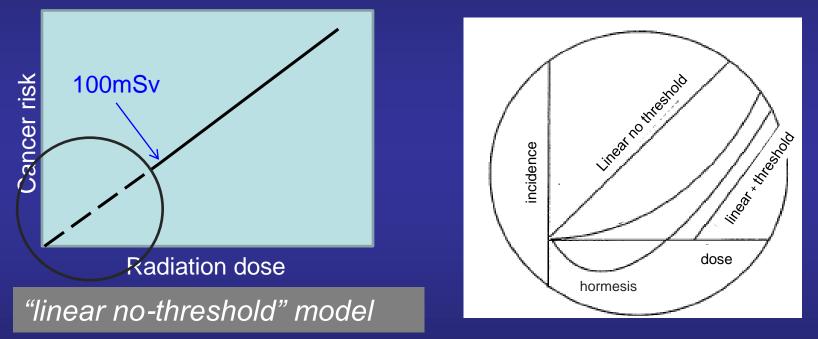
X-rays \rightarrow ionizing radiation

\rightarrow deterministic and stochastic effects



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Why dosimetry? Cancer risk? International Commission on Radiological Protection¹ French Academies Report, 2005²

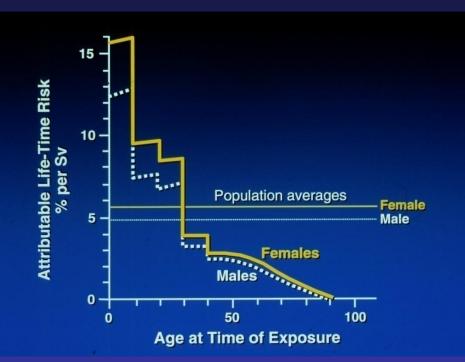


¹ICRP103: The 2007 Recommendations of the International Commission on Radiological Protection . Annals of the ICRP Vol 37 (2007)

²French Academies Report, 2005. La relation dose-effet et l'estimation des effets cancérogènes des faibles doses de rayonnements ionisants.



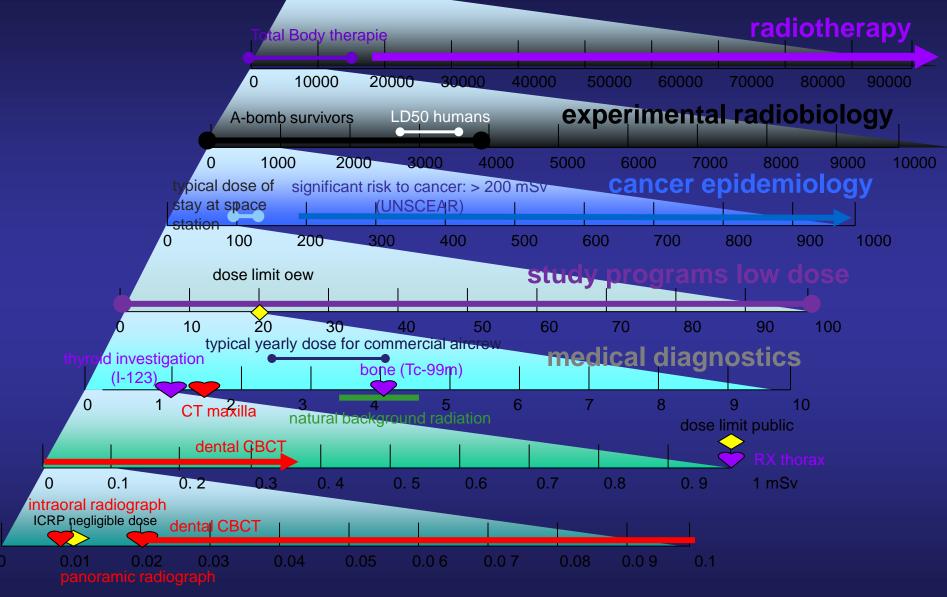
Why dosimetry? Risk of cancer and age



Age group (years)	Multiplication factor for r		
<10	× 3		
10-20	x 2		
20-30	x 1.5		
30-50	× 0.5		
50-80	× 0.3		
80+	negligible risk		

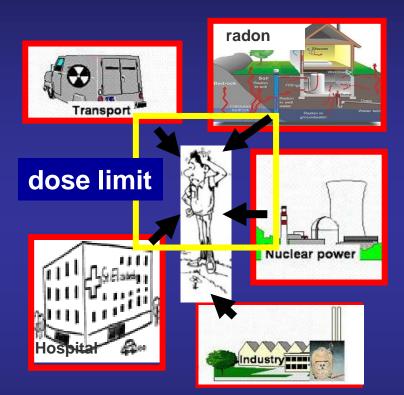
SEDENTEXCT

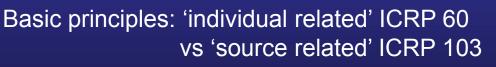
Why dosimetry? Effective dose ranges and dental investigations

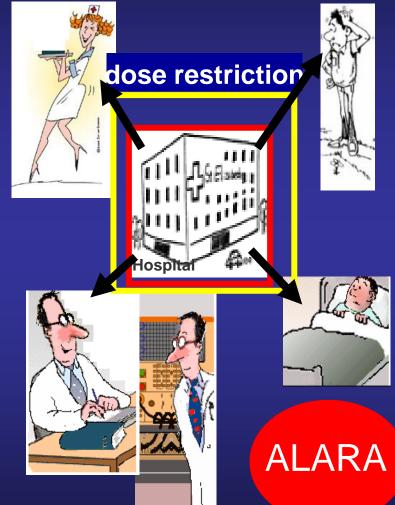


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The ICRP basis for radiation protection: patient and personnel protection policy European Directive 97/43 Euratom

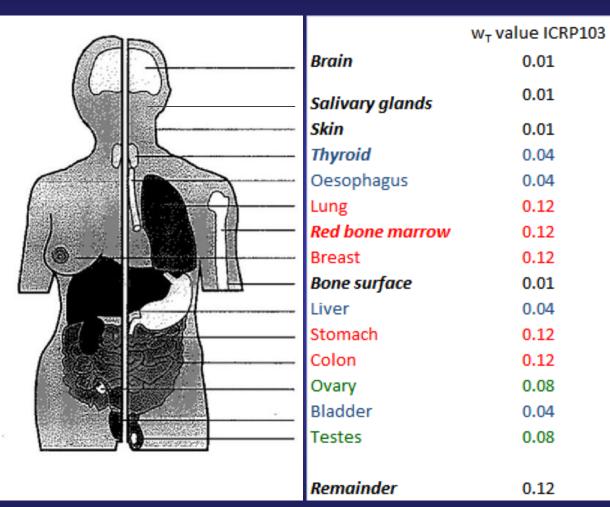








Patient dose and technical dose characterization





0.01

0.01

0.01

0.04

0.04 0.12

0.12 0.12

0.01

0.04

0.12

0.12

0.08

0.04

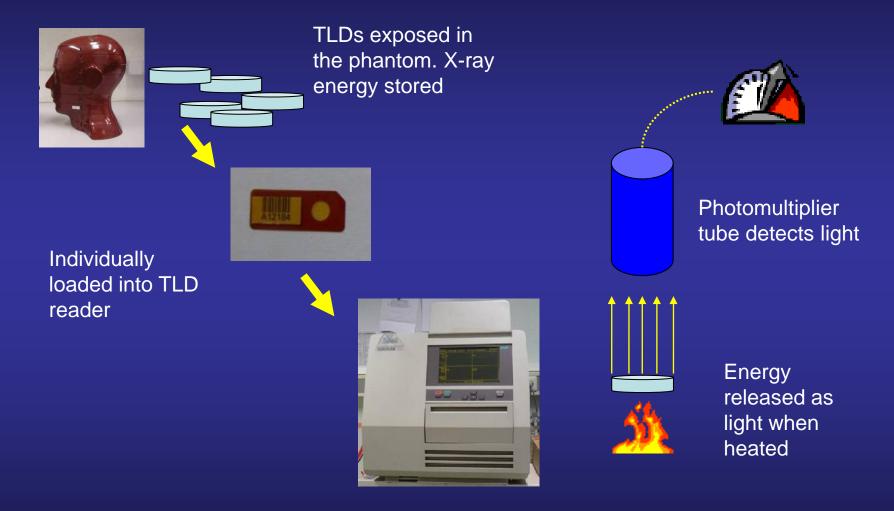
0.08

0.12



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Patient dose and technical dose characterization





Patient dose and technical dose characterization

- effective dose is influenced by technical parameters as mAs and FOV
- definition of standard dose index that can be measured in a physical phantom during a routine medical physics quality control check
- candidates:
 - cfr. medical CT: Computed Tomography Dose Index,

DLP (Dose Length Product) DAP

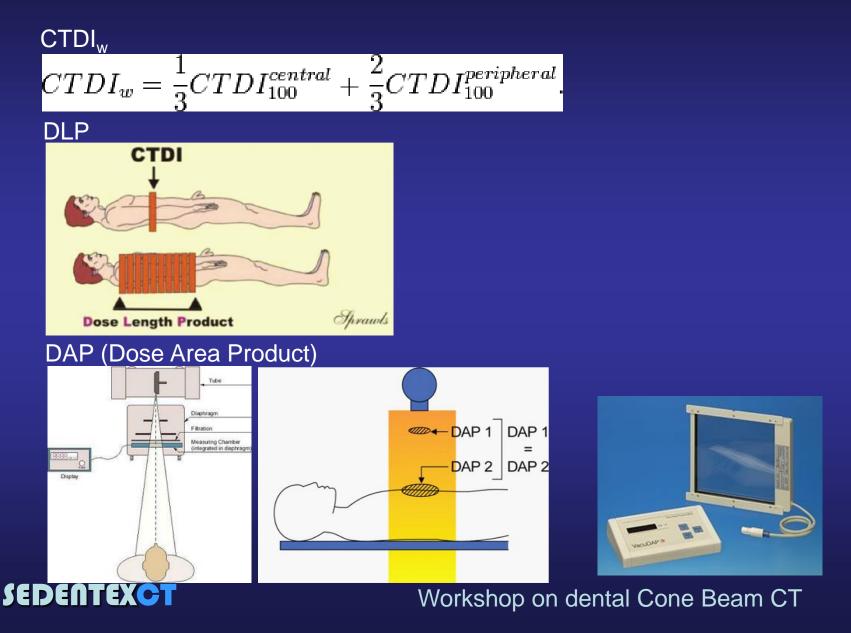
DOSE CTDI Sprawls r50mm $CTDI_{100} =$ $D_a(z)dz$ 50mm





Workshop on dental Cone Beam CT

CTDI



- determine conversion factors between technical quantities and effective dose:

the more dose determining factors that are included in the dose index, the more uniform the conversion factors will be:

dose conversion factors from mAs to effective dose will be different for different CBCTs and imaging protocols

dose conversion factors from a dose index including the influence of both mAs and FOV can possibly be the same for different CBCTs and imaging protocols



Available knowledge at the start of SEDENTEXCT

• Few dosimetric studies

≠ anthropomorphic phantoms

≠ TLDs

Limited number of TLDs & CBCT devices

- Few studies investigating CTDI and alternative indices for modern MSCT scanners
- No dose simulation studies for dental CBCT
- No reports on scattered dose in dental CBCT



SEDENTEXCT dosimetry

KUL Leuven, UNIMAN Manchester, NKUA Athens, VU Vilnius, MAHOD Malmö, LTO Leeds

- Dental CBCT dose index
- Assessment of dental CBCT dose (adult & paediatric phantom, patients)
- Monte Carlo simulation framework for optimisation
- Personnel dose



Measurement of the dose distribution in anatomical phantoms and subsequent calculation of effective dose



Phantom dose

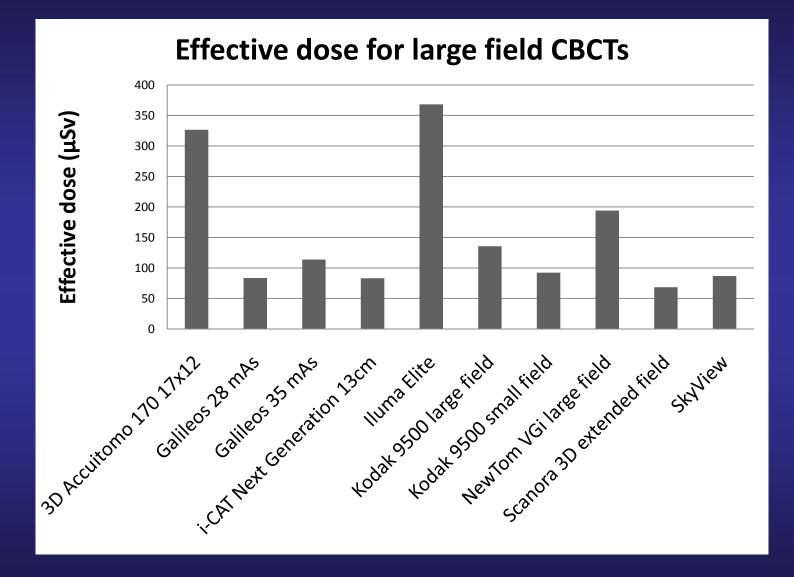
Adult (Alderson) and paediatric (ATOM) phantoms



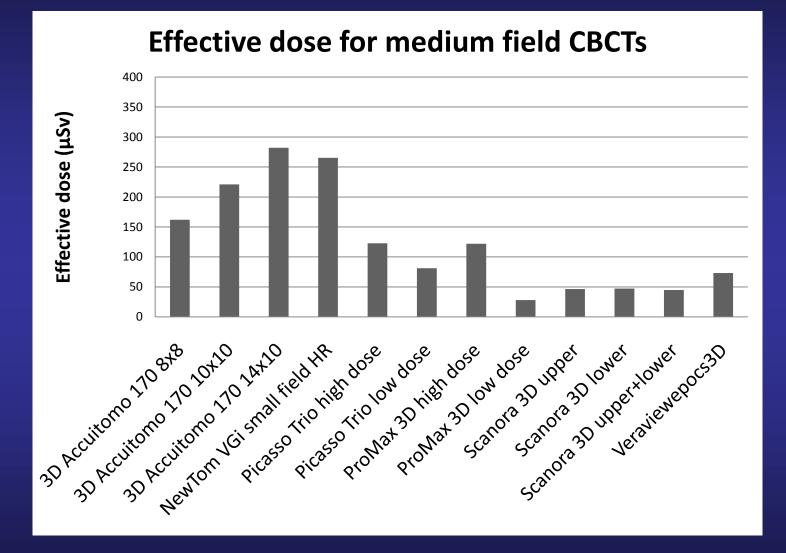




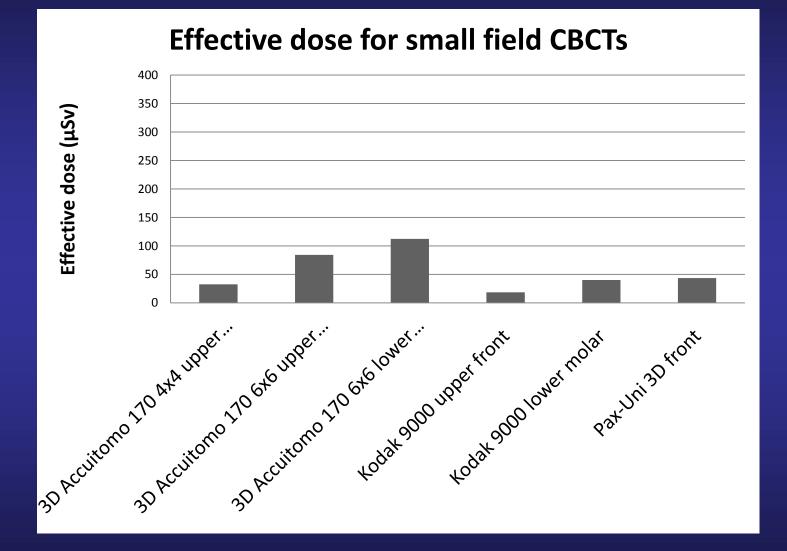




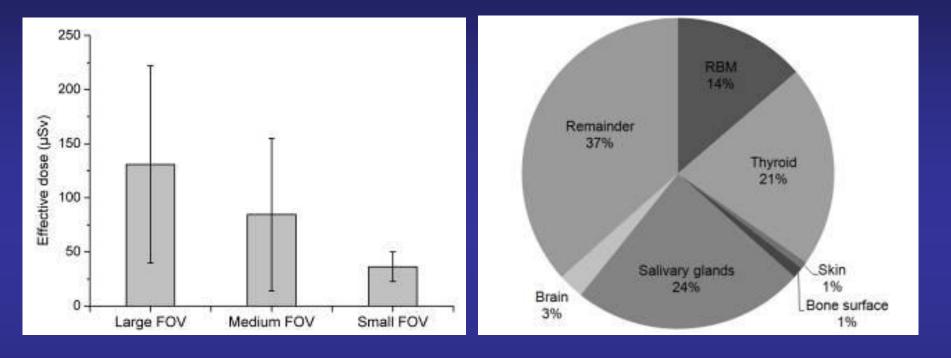








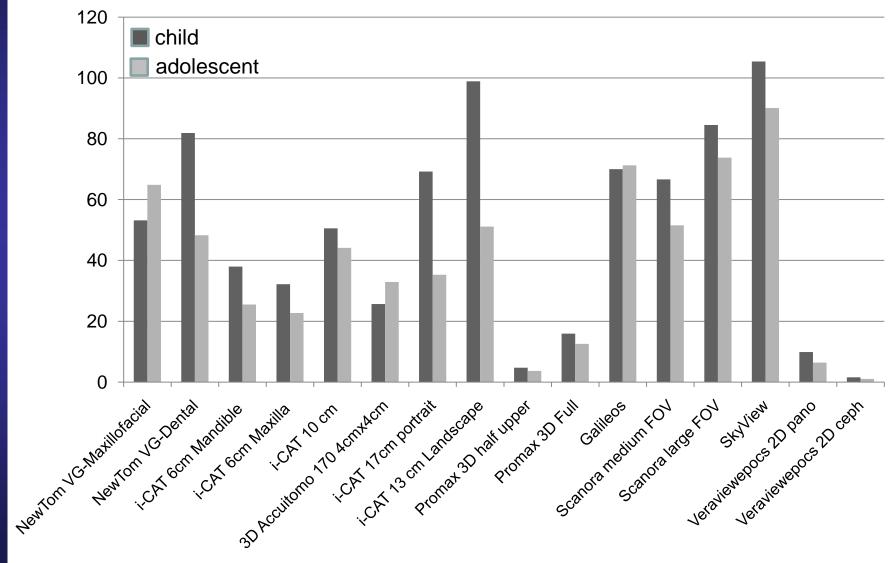






Phantom dose: paediatric

Effective dose (µSv) for paediatric phantoms



Phantom dose

Conclusions

- Wide dose range (~20 μ Sv to 400 μ Sv: 20-fold)
- Main contributors: thyroid gland, salivary glands, 'remainder' (oral mucosa, extrathoracic airways)
- Clear effect of FOV size & position on effective dose
- Other differences between doses: interplay with image quality!
- Optimised ("customised") patient dose: FOV selection based on region of interest + exposure selection based on image quality requirement



In vivo dose measurements

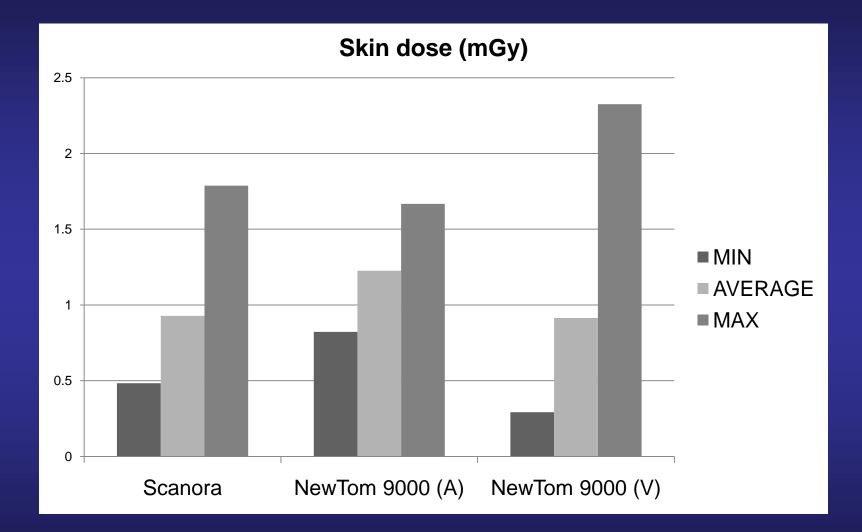


In vivo dose

	KUL Scanora 3D		VU NewTom 9000		NKUA NewTom 9000	
Clinical indication	# Patients	Age	# Patients	Age	# Patients	Age
Implant placement	43	13-61	30	20-68	15	28-62
Orthodontic planning	4	10-13	0	/	1	13
Impacted teeth	8	10-20	43	10-83	10	18-33
Maxillofacial trauma/ tumors/ development abnormalities	1	20	29	11-49	0	/
Sinus visualisation	4	35-60	42	22-76	0	/
Others	10	10-54	0	/	8	24-62



In vivo dose





In vivo dose

Conclusions

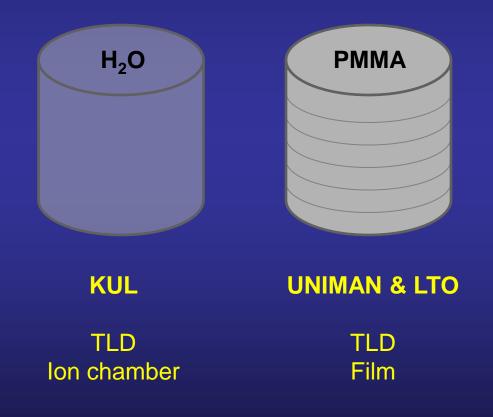
- Wide range of in vivo doses due to difference in exposure and patient size
- Manual adjustment of exposure parameters ↔ Pre-set exposure protocols based on patient size ↔ Automatic exposure control



Development of a standardised dose index to characterise dose distribution in dental CBCTs

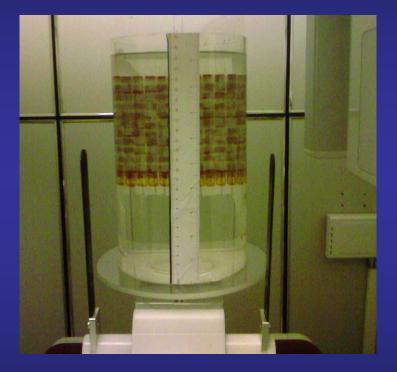


Assessment of dose distribution





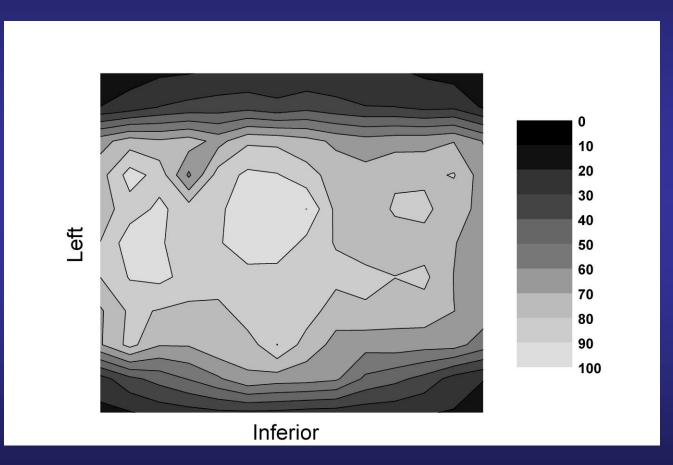
Assessment of dose distribution





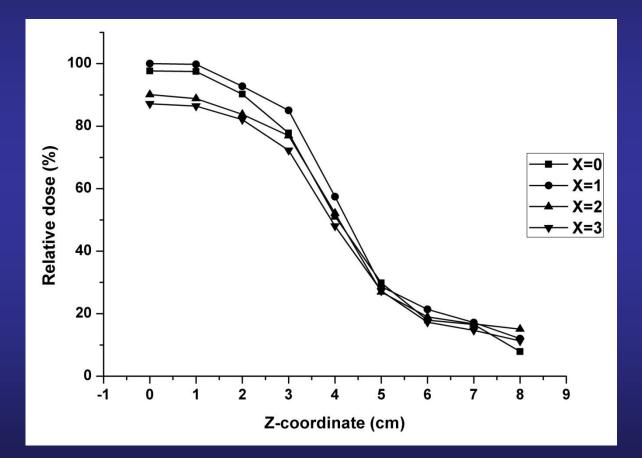


Scanora 3D, TLDs in 'coronal' plane





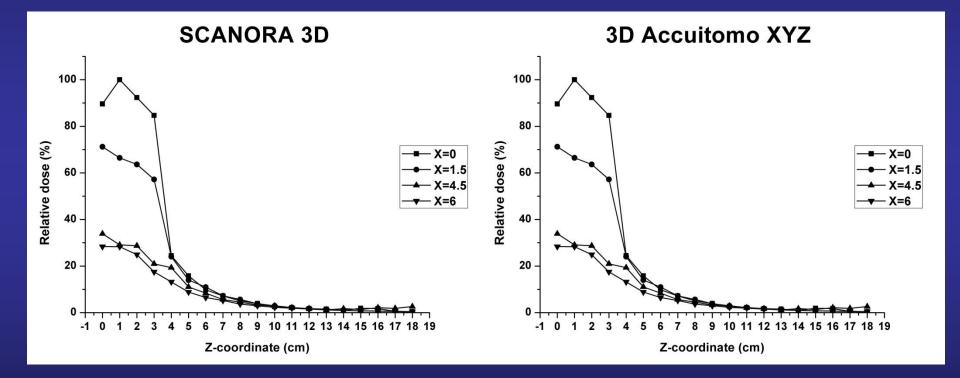
Scanora 3D, ion chamber, along z-axis



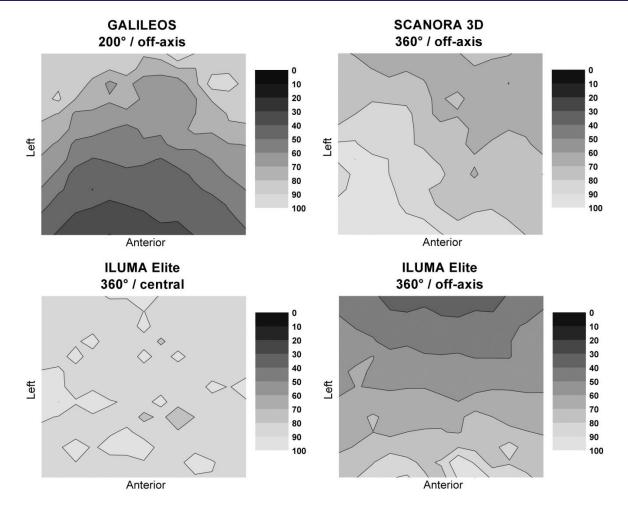
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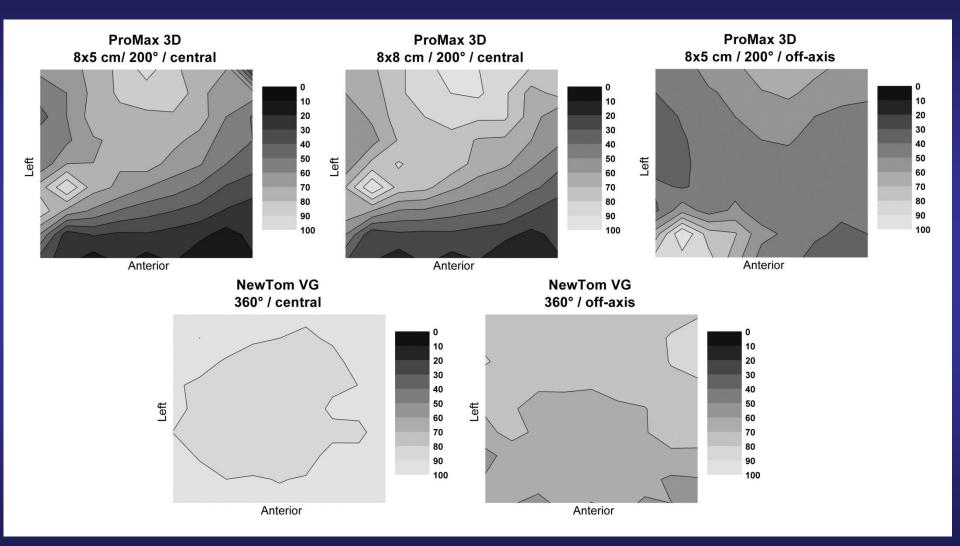
TLDs, along z-axis



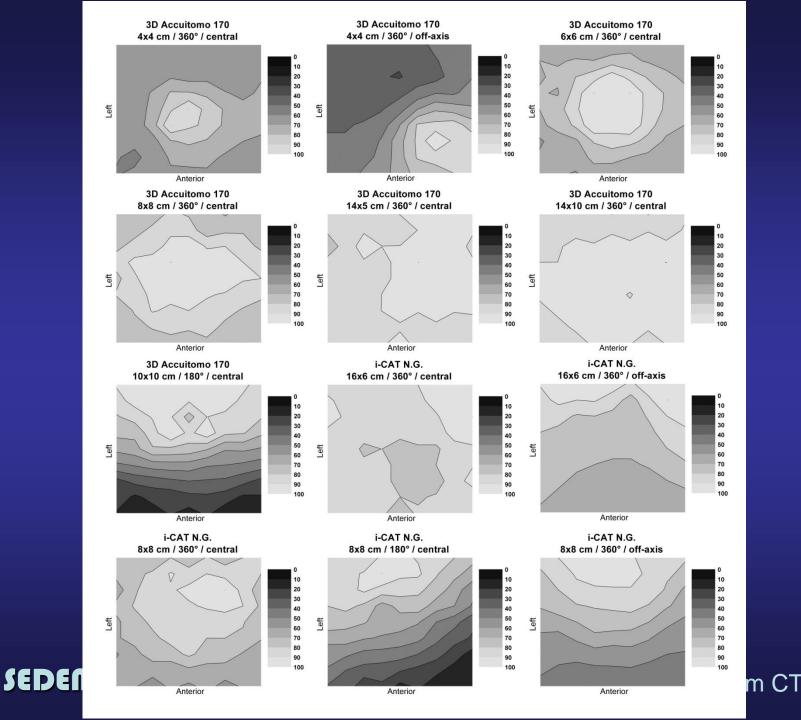
Dental CBCT dose index TLDs, 'axial' plane



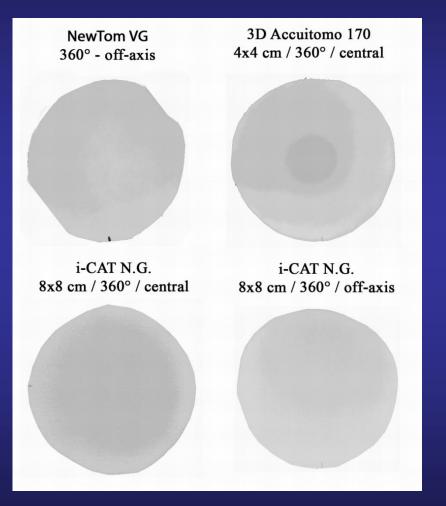
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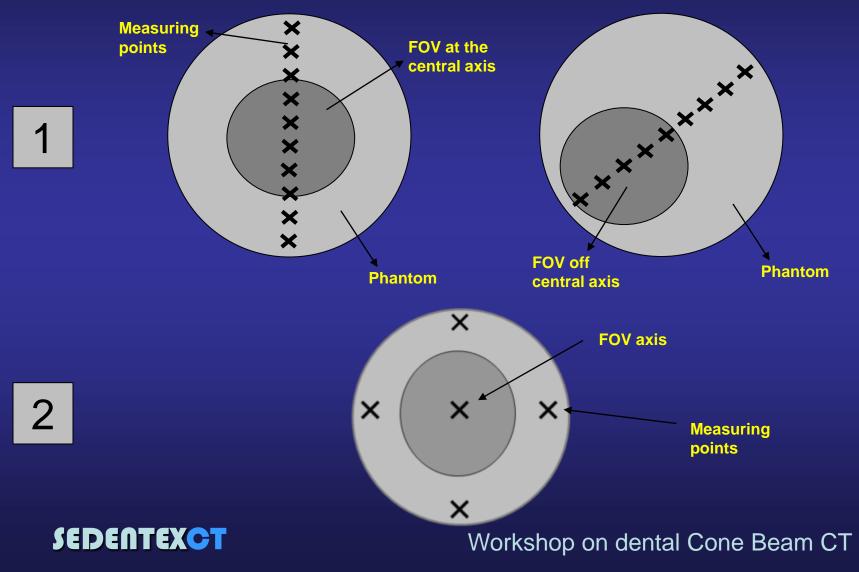


Film measurements, 'axial' plane





Dose distribution \rightarrow dose index definition



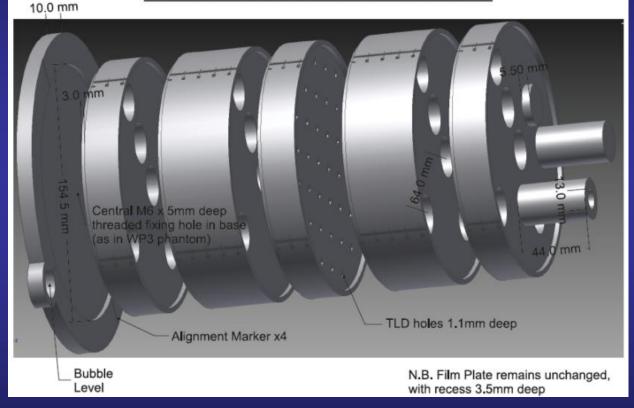
Dose Area Product (DAP)?





\rightarrow Validation of indices

SEDENTEX-CT WP2 PROTOTYPE 2 - B



Leeds Test Objects Ltd



 \rightarrow Validation of indices

 Measurement of indices in practice
Definition of correlation factors through Monte Carlo simulation



Dose index measurements for the 3D Accuitomo 170 Dose index me						lex measi	urements	s for the S	Scanora 3	3D
		Index1		Index 2			Index1		Index 2	
	Drotocol	(mGy)		(mGy)		Destand	(mGy)		(mGy)	
47.40	Protocol	AVG	STDEV	AVG		Protocol	AVG	STDEV	AVG	
17x12	90 kV, 5 mA, 17s, 360	8.8	0.4	8.8	14.5x7.5	24 mAs,	2.1	0.1	2.2	
	80 kV, 5 mA, 17s, 360	6.4	0.2	6.4		central				
	70 kV, 5 mA, 17s, 360	4.3	0.2	4.3	10x7.5	30 mAs,	2.2	0.1	2.3	
47.5	80 kV, 5 mA, 9s, 180	4.5	2.3	4.5		central				
17x5 14x10	90 kV, 5 mA, 17s, 360	7.5 8.6	0.4 0.3	7.4		30 mAs,	2.1	0.6	1.7	
	90 kV, 5 mA, 17s, 360			8.6 7.0		off-axis				
10x10	90 kV, 5 mA, 17s, 360 90 kV, 5 mA, 17s, 360	7.9 7.4	0.7 2.4	7.9 7.3	6x6	36 mAs,	1.8	0.4	2.0	
8x8	90 kV, 5 mA, 17s, 360 90 kV, 5 mA, 17s, 360	7.4 6.4	2.4	7.3 6.5		central				
020	80 kV, 5 mA, 17s, 360	0.4 4.8	2.3 1.8	0.5 4.9		36 mAs,	1.7	1.0	1.5	
	70 kV, 5 mA, 17s, 360	4.8 3.3	1.3	4.9 3.4		off-axis				
	90 kV, 5 mA, 9s, 180	2.9	0.6	3.4						
	80 kV, 5 mA, 9s, 180	2.3	0.0 0.5	2.5						
	70 kV, 5 mA, 9s, 180	1.5	0.3	1.8						
6x6	90 kV, 5 mA, 17s, 360	5.2	3.1	5.2						
0.00	80 kV, 5 mA, 17s, 360	4.0	2.4	3.9						
	70 kV, 5 mA, 17s, 360	2.8	1.8	2.7						
	90 kV, 5 mA, 9s, 180	2.0	0.2	2.7						
	80 kV, 5 mA, 9s, 180	1.6	0.2	2.0						
	70 kV, 5 mA, 9s, 180	1.0	0.1	1.4						
4x4	90 kV, 5 mA, 17s, 360	3.4	3.0	3.7						
	80 kV, 5 mA, 17s, 360	2.6	2.3	2.7						
	70 kV, 5 mA, 17s, 360	1.9	1.7	1.9						
	90 kV, 5 mA, 9s, 180	1.2	0.3	1.9						
	80 kV, 5 mA, 9s, 180	0.9	0.2	1.4						
	70 kV, 5 mA, 9s, 180	0.6	0.2	1.0						

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Conclusions

- dose distributions in homogenous phantoms have been measured for different CBCTs and settings
- 2 CBCT dose indices have been proposed in addition to DAP
- validation measurements have been performed



Development of mathematical models for dental CBCT dosimetry



- Development of computational models for dental CBCT dosimetry
- Calculation of conversion factors to effective dose using Monte Carlo (MC) simulations
- MC simulations are a class of computational algorithms that are based on repeated random sampling to compute the results
- MCNP5: a general purpose Monte Carlo N-Particle code



- Point sources with biased direction
- Arranged on a circle positioned at 10 degrees intervals
- A set of collimators to shape the beam
- ATOM 10 year old phantom
- ICRP male reference computational phantom
- ICRP female reference computational phantom (in progress)



→ Validation of the simulated CB geometry-Stage 1
• Simulated doses were compared against measured doses

- PMMA cylinder on a 3D Accuitomo 170 CBCT scanner
- Thermoluminescent dosimeters (<10% error)
- Full and half rotation CB geometry
- Range of tube voltages
- Range of FOVs
- Axis and off-axis

 % difference between simulated and measured normalised doses ranged between 7% to 19%



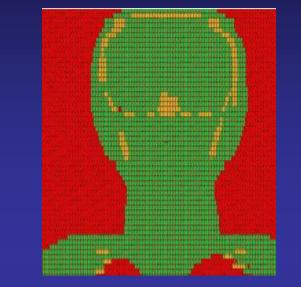
→ Validation of the simulated CB geometry-Stage 2
• Simulated doses were compared against measured doses

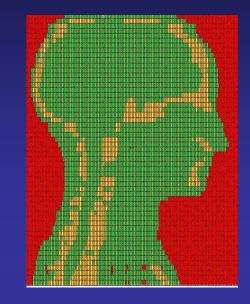
- ATOM 10 year old anthropomorphic phantom
- Thermoluminescent dosimeters (<10% error)
- Range of CBCT machines
- Range of FOVs
- Range of clinical examinations

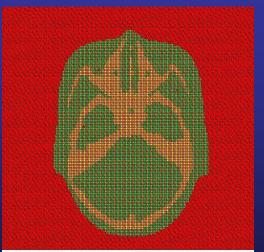
 % difference between simulated and measured effective doses ranged between 3% to 17%





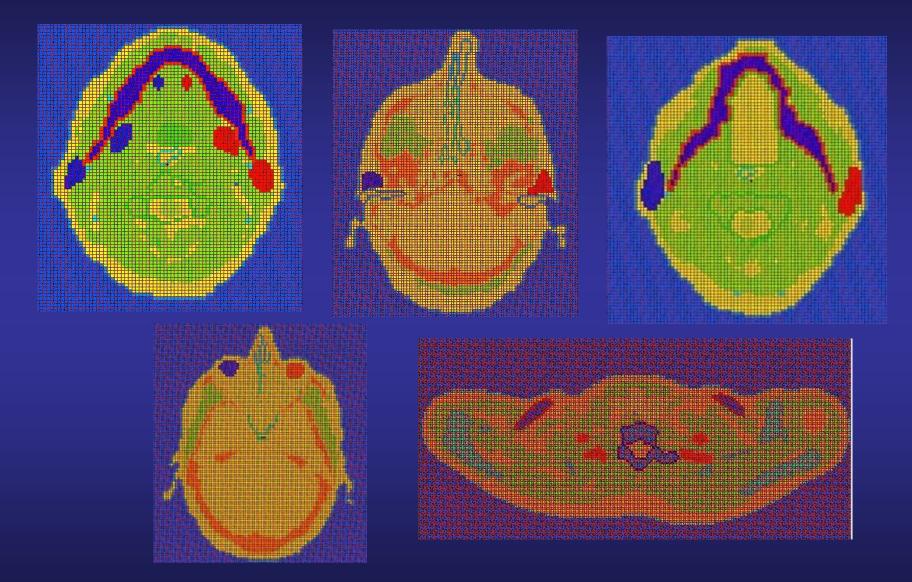














→ Calculation of conversion factors (mSv/mAs)

- Three computational phantoms
- Range of FOVs, tube voltages, imaging protocols
- Range of machines
 - i-CAT NG
 - Kodak 9000 3D
 - Kodak 9500 3D
 - NewTom VG
 - NewTom VGi
 - NewTom 5G
 - NewTom QR DVT 9000
 - Galileos Comfort
 - Galileos Compact
 - 3D Accuitomo XYZ
 - 3D Accuitomo 170
 - Planmeca Promax 3D (in progress)
 - Planmeca Promax 3D Max (in progress)

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3D Accuitomo 170- 10 year old phantom								
Imaging protocol	Tube voltage	Rotation	Conversion factor					
4Øcm x 4cm	70/80/90	360	0.00028/0.00030/0.0.00031					
Maxillary Canine								
10Øcm x 10cm	70/80/90	360	0.00248/0.00255/0.00261					
17Øcm x 5cm	70/80/90	180	0.00252/0.00254/0.00256					
Mandible								
i-CAT NG - ICRP male reference computation phantom								
Imaging protocol	Tube voltage	Rotation	Conversion factor					
16Øcm x 6cm	120	360	0.00130					
Maxilla								
16Øcm x 6cm	120	360	0.00189					
Mandible								
16Øcm x 13cm	120	360	0.00299					
Maxilla								



Conclusions

- a CBCT simulation model was set up and validated
- conversion factors from mAs to effective dose were calculated for a broad range of CBCTs
- relation between dose indices 1 and 2 and effective dose has been established but needs further elaboration
- the relationship with DAP needs to be stablished

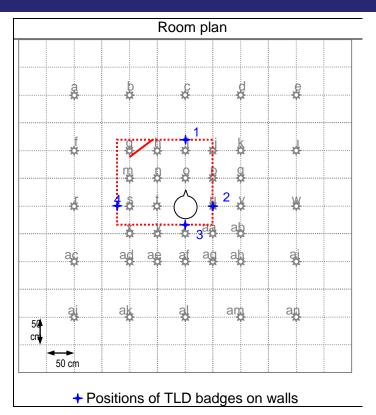


Measurements of scatter dose and radiation protection of personnel and helpers



Protocol

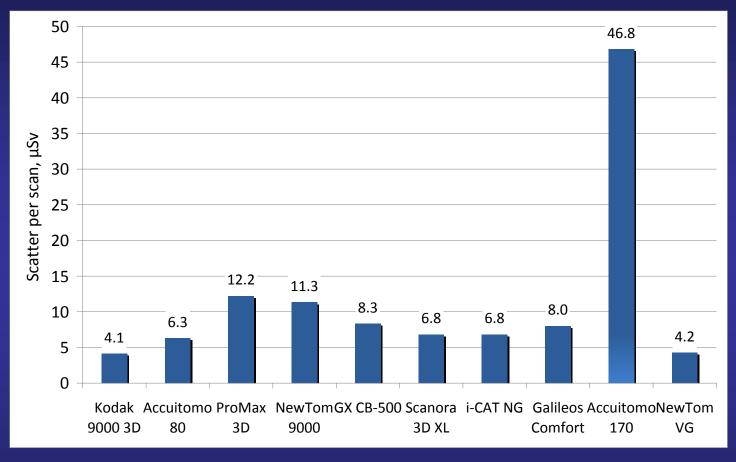
 Active measurements Using ionisation chamber or dose rate meter On standard measurement grid Passive dosimetry High sensitivity TLD badges In rooms where CBCT is sole X-ray source



SEDENTEXOT

Model	Manufacturer	Active measurements	Passive measurements
Kodak 9000 3D	Kodak	~	
Accuitomo 80	J. Morita	✓	✓
ProMax 3D	Planmeca	~	
NewTom 9000	QR	✓	✓
GX CB-500	Gendex	✓	
Scanora 3D XL	Soredex	✓	✓
i-CAT NG	Imaging Sciences International	~	
Galileos Comfort	Sirona	✓	
Accuitomo 170	J. Morita	✓	✓
NewTom VG	QR	√*	





Active



Shielding requirements

- For small FOVs, low workload and low occupancy of adjacent areas, additional shielding may not be required
- In most circumstances, 0.5-1.5mm Pb equivalence is required dependent on:
 - Average scatter per scan
 - Distance
 - Workload
 - Dose constraint



Conclusions

- Max scatter varies 4.1 to 46.8 μSv per scan
- Distance unlikely to be sufficient protection, shielding required up to about 1.5mm lead
- Advice of a qualified expert required
- Base calculations on maximum scatter and double expected workload
- Manufacturers should provide scatter data
- Keep scatter and workload under review
- Further work needed on workload and to assess new units

SEDENTEXCT

Work beyond SEDENTEXCT?

Further focus on paediatric dose

'Mathematical' dose optimisation (simulations, reconstruction algorithms)

Long-term quality control of CBCT devices



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