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Model Based Dose Calculation Algorithms for Photons in External Radiation Therapy

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Objectives

1. To provide an educational review of the physics and techniques behind model based algorithms e.g. convolution/superpositioning models.
2. To review the methods used to improve the simulation efficiency i.e. pencil beam and collapsed cone convolutions.
3. To briefly review the vendor codes currently used for clinical treatment planning.
4. To briefly review the potential clinical implications of accurate calculated dose distributions.



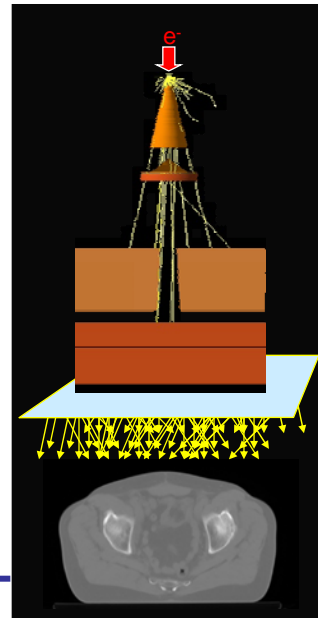
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The Problem

- Modelling the linac
 - Energy fluence
 - Source models
 - Monte Carlo
- Modelling of dose in patients
 - Interpolation and correction of measured data
 - Fluence to dose modelling
 - Monte Carlo



Remember

- Photons are indirect ionizing radiation
- Produces electrons through interaction
 - Pair production, Compton and photo
- Electrons deposit energy by ionization
- Thus keeping track of electrons is highly important for accurate dose calculations



Photon Fluence via TERMA to Absorbed Dose using Convolution

D - Dose distribution
 T - TERMA distribution
 K - dose deposition kernel



$$D(x) = \int T(x') \cdot K(x-x') dx'$$

$$D(\mathbf{r}) = T(\mathbf{r}) \otimes K(\mathbf{r})$$

This idea was explored by several papers at the ICCR 1984

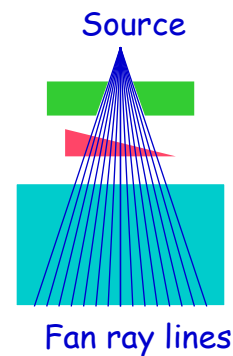


What is TERMA?

- Ray-tracing Total Energy Released in Mass (TERMA)
- Similar to determining effective or radiological depth

$$T(E, z) = \frac{\mu_E}{\rho} \cdot \Phi(E, z) \cdot E = \frac{\mu_E}{\rho} \Phi_0(E, 0) \cdot e^{-\mu_E \cdot z_{eq}} \cdot E$$

$$z_{eq} = \frac{1}{\rho_{water}} \int_0^z \rho(z') \cdot dz'$$



Dose deposition kernel - *K*

0.5 cm → Primary

10 cm → First scatter

10 cm → Total

1.25 MeV

Phys. Med. Biol., 1988, Vol. 33, No. 1, 1-20. Printed in the UK

5 MeV

15 MeV

Courtesy A. Ahnesjö

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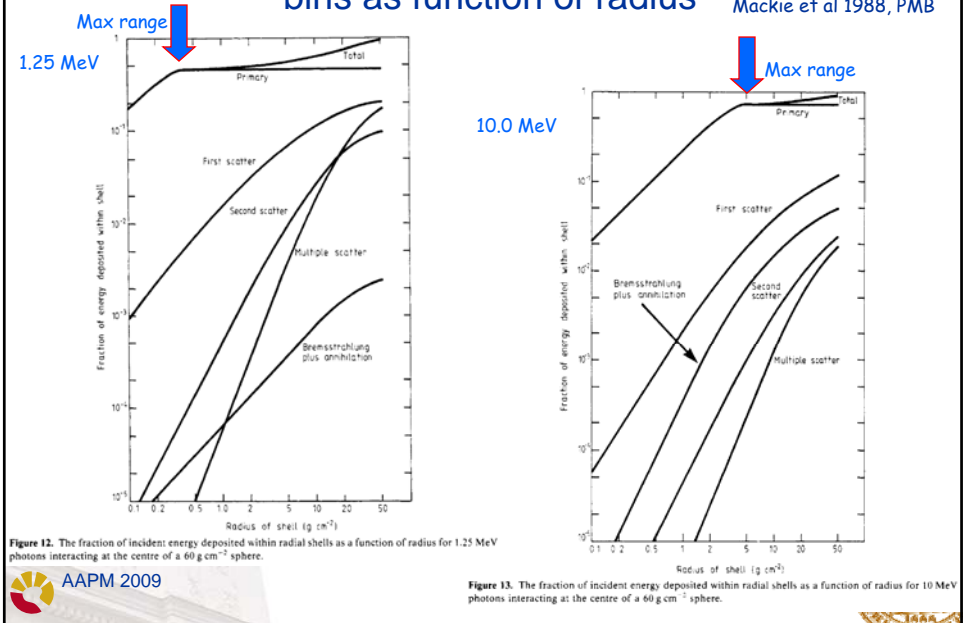
Generation of photon energy deposition kernels using the EGS Monte Carlo code

T. R. Mackie¹*, A. F. Bielajew¹, D. W. O. Rogers² and J. J. Battista³

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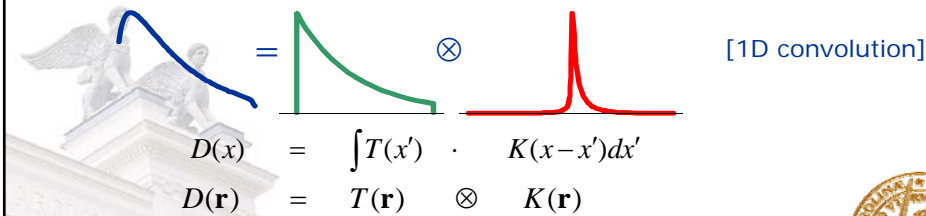
The fraction of energy deposited within radial bins as function of radius

Mackie et al 1988, PMB

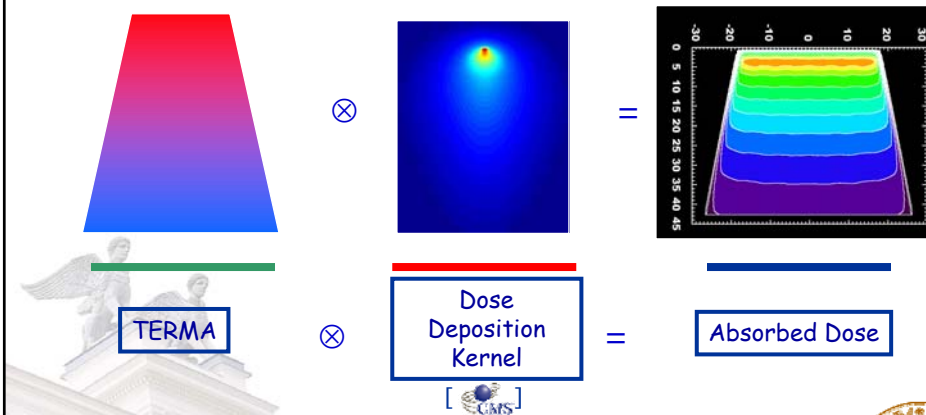


Convolve!

- Apply the dose kernel to each TERMA point
- Integrate over the whole volume i.e. a convolution



Convolution in 2D



Convolution is efficiently solved by Fast Fourier Transform techniques



Example: Point kernel convolution - CMS

- Re-sampling of Mackie's kernels to Cartesian coordinates
- Fast Fourier Transform (FFT) solution
- Two separate calculations:
 - A **primary kernel** for which the calculation is performed at high-resolution but over a small region – **high gradient – short range**
 - A **scatter kernel**, where the calculation is performed at a lower resolution but over a larger area – **low gradient – long range**
 - Time saving of about 65 % by this technique



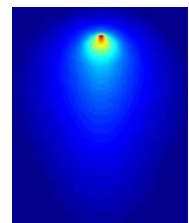
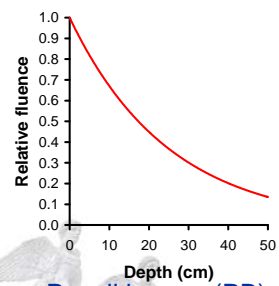
Limitations of convolution

- Kernels are not invariant in space
 - Energy distribution varies with position in beam
 - Beam softening laterally
 - Beam hardening longitudinally
- Kernels vary with density
- Divergence leading to tilted kernels
- Pre-calculated kernels won't make it!!!
- FFT not suitable – analytical methods must be used – time consuming
- Approximate methods required



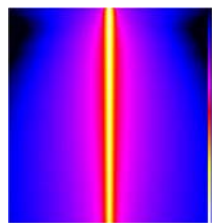
1st approximation Pencil Beam

- Reduce the dimensionality of the problem by pre-convolving in the depth dimension



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- => Pencil beams (PB)
- Superposition of pencil beams in 2D => Faster
- Creation by: De-convolution or differentiation from measurements or by Monte Carlo methods

Use of fast Fourier transforms in calculating dose distributions for irregularly shaped fields for three-dimensional treatment planning¹⁾
Radhe Mohan and Chen-Shou Chui
Department of Medical Physics, Memorial Sloan-Kettering Cancer Center, New York, New York 10027
(Received 18 June 1988; accepted for publication 22 October 1988)

A pencil beam model for photon dose calculation
Anders Arvidsson
Department of Radiation Physics, Karolinska Institute, Stockholm, Sweden and Rikshä. Å. Box 1704
S-141 86 Ejnarby, Sweden
Mikael Sauer and Auro Trapp
Rikshä. Å. Box 1704, S-141 86 Ejnarby, Sweden
(Received 29 October 1990; accepted for publication 1 July 1991)

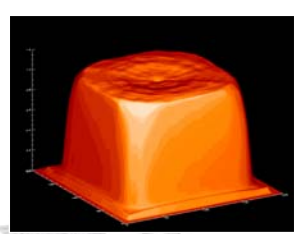
Experimental determination of the dose kernel in high-energy x-ray beams
Oleker P. Cederberg and Bernd E. Björngård
Department of Radiation Oncology, Roger Williams Medical Center, Brown University, Providence, Rhode Island 02912-0332
Timothy C. Zhu
Department of Radiation Oncology, Mankin Cancer Center, University of Florida, Gainesville, Florida 32610-6187
(Received 26 June 1995; accepted for publication 12 December 1995)

Extraction of pencil beam kernels by the deconvolution method
Chen-Shou Chui and Radhe Mohan
Department of Medical Physics, Memorial Sloan-Kettering Cancer Center, New York, New York 10027
(Received 15 June 1987; accepted for publication 14 December 1987)

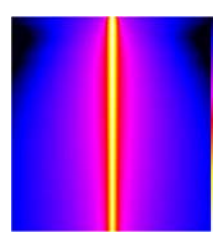
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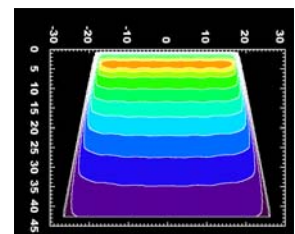
Illustration of Pencil Beam superpositioning (convolution)



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Energy fluence

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Dose
Deposition
Kernel

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Absorbed Dose



Example: Pencil beam model – Nucletron OMP

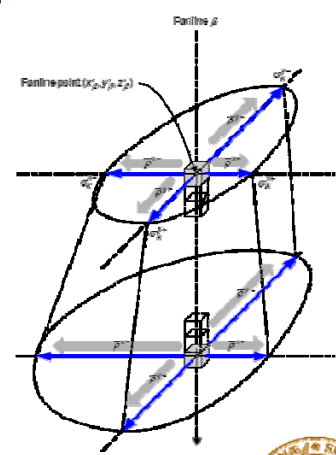
- Pencil beams based on MC calculated point kernels, integrated and fitted to a limited number of depth doses
- Separates “primary” and “scatter” dose
- Heterogeneities handled via effective path length – **only longitudinal** scaling
- Extensive beam modelling

Nucletron (former MDS Nordion and Helax-TMS)



Example: Pencil beams model - Eclipse

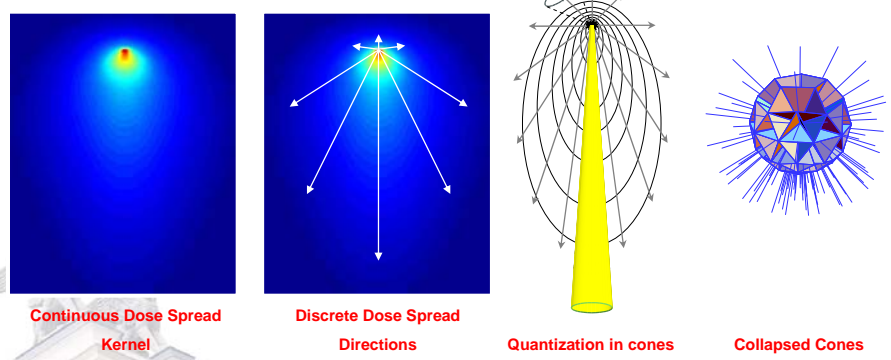
- Uses pencil beams extracted from measurements (SPB) or from Monte Carlo calculation (AAA)
- Heterogeneities handled via effective path length – longitudinal
- AAA adds a scaling of the spread of the pencil based on density – **lateral**
- AAA also have an extensive beam modelling



Analytical Anisotropic Algorithm



2nd approximation Collapsed cone convolution



Collapsed cone convolution of radiant energy for photon dose calculation in heterogeneous media
 Anders Ahnesjö
 Department of Radiation Physics, Karolinska Institute and University of Stockholm, Box 4021, S-141 81
 Stockholm, Sweden
 (Received 15 August 1988, accepted for publication 3 May 1989)

Collapsing removes the inverse square law – only exponential attenuation is left

Best Med Phys paper 1989

Kernels are discretised

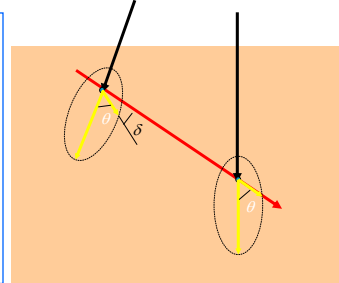
Number of collapsed cones or directions

- Sufficient density of cones to distribute energy to all voxels
 - Not possible but at least while the energy is significant
 - ~100 (Mackie et al, 1996 Summer school)
 - Voxels will be missed at large distances – very low energy contribution
- 128 CC are used in CMS (48 for the fast version)
- 106 CC are standard in OMP



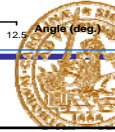
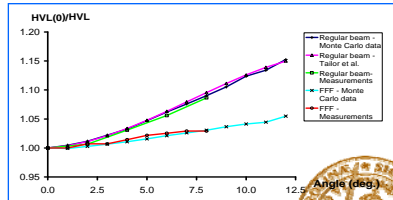
Implementation issues

- Accounts for
 - Heterogeneities
 - Kernels scaled for different tissues
 - Lateral energy transport
 - Beam Hardening and Off-axis spectrum softening
 - Included in Ray Trace process
 - Tilt of kernels
 - Included in Transport



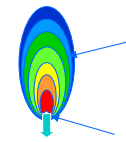
Polyenergetic Spectrum accounted for by weighted sum of monoenergetic kernels calculated by Monte Carlo

Weights determined by comparison with measured data



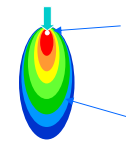
Examples: Collapsed cone

- Philips - Pinnacle
 - Polyenergetic weighted kernels, total energy
 - Off-axis/tilting considered during TERMA
 - Collecting dose or dose point of view
- CMS - XiO
 - Two kernels, Primary electron dose and scattered photon dose
 - No Off-axis/tilting
 - Collecting dose or dose point of view
- Nucletron – Oncentra MasterPlan
 - Two Kernels are used:
 - One for Collision Kerma into Primary Dose
 - One for 'Scerma' into Phantom Scatter Dose
 - Kernels parameterised and fitting parameters stored for run time
 - Off-axis/tilting
 - Recursive dose collect/deposit model along parallel lines



These are 'iso-scatter' lines.

They link points producing equal scatter to here.



Primary interaction point

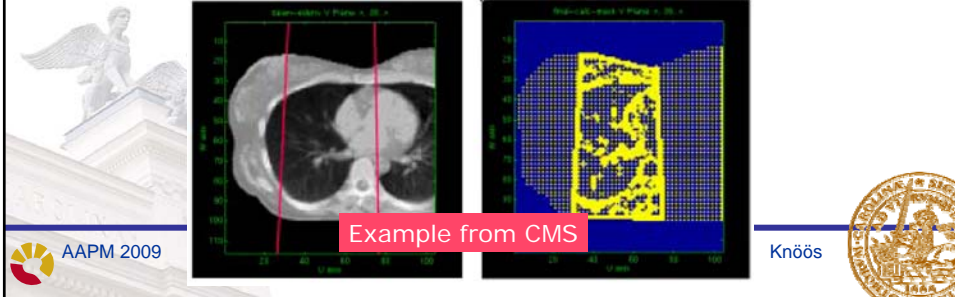
These are 'isodose' lines

From Deshpande, Philips



Further approximation

- Multigrid solution (CMS - XiO)
 - Only calculate dose using superposition at points where it is necessary, and at all other points use interpolation to get a reasonable estimate of dose
- Adaptive CCC (Philips - Pinnacle)
 - Only performs convolution at every 4th point in the TERMA array
 - Gradient search performed on TERMA array
 - Dose in between is interpolated if gradient low (i.e TERMA doesn't change much)
 - Convolution performed at every point if TERMA gradient high



Summary of Models/Algorithms

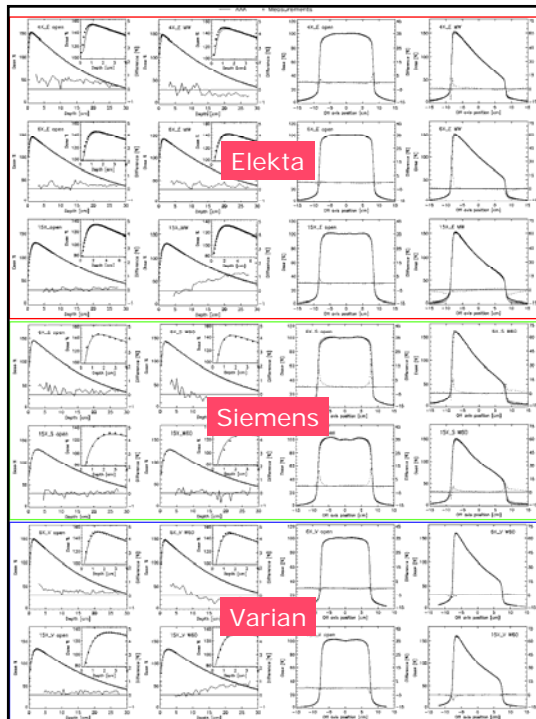
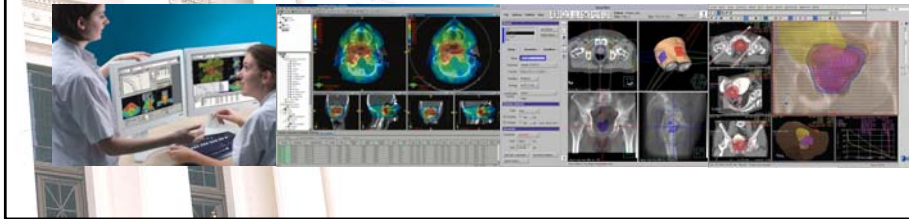
- Inhomogeneities are handled by scaling the kernels rectilinearly with electron density according to the theorem by O'Connor 1957
- **Type a** – Models primarily based on EPL scaling for inhomogeneity corrections.
 - Eclipse/SPB, OMP/PB, PPLAN, XiO/Convolution
 - **LONGITUDINAL** scaling
- **Type b** – Models that in an approximate way consider changes in lateral electron transport
 - Pinnacle/CC, Eclipse/AAA, OMP/CC, XiO/Superpositioning.
 - **LONGITUDINAL** and **LATERAL** scaling





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Performance of Convolution Models



AAA-PB model in Eclipse 24

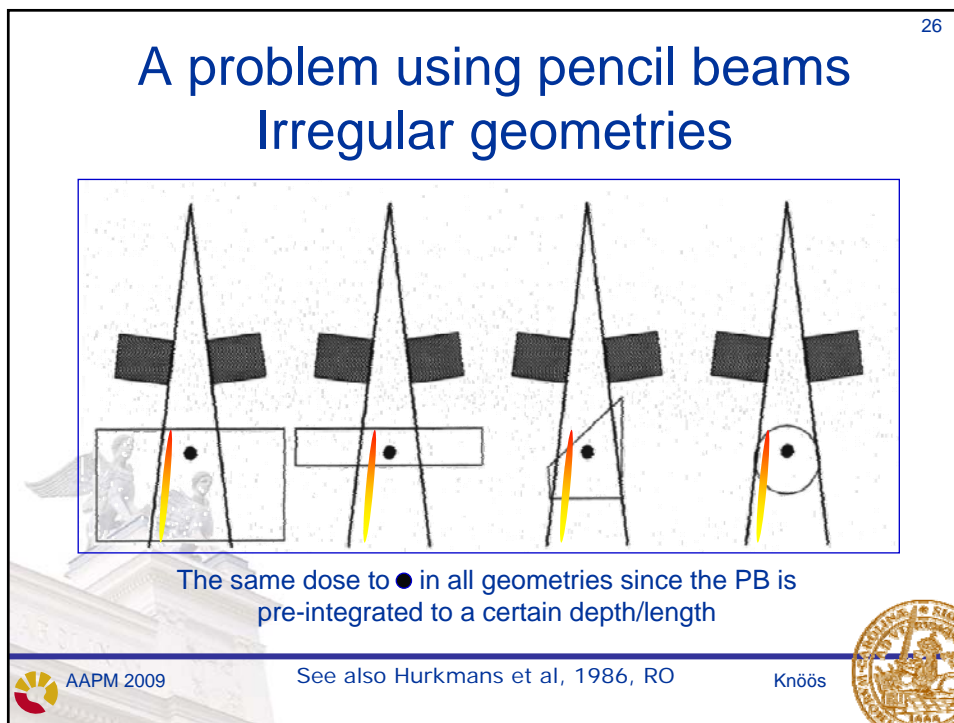
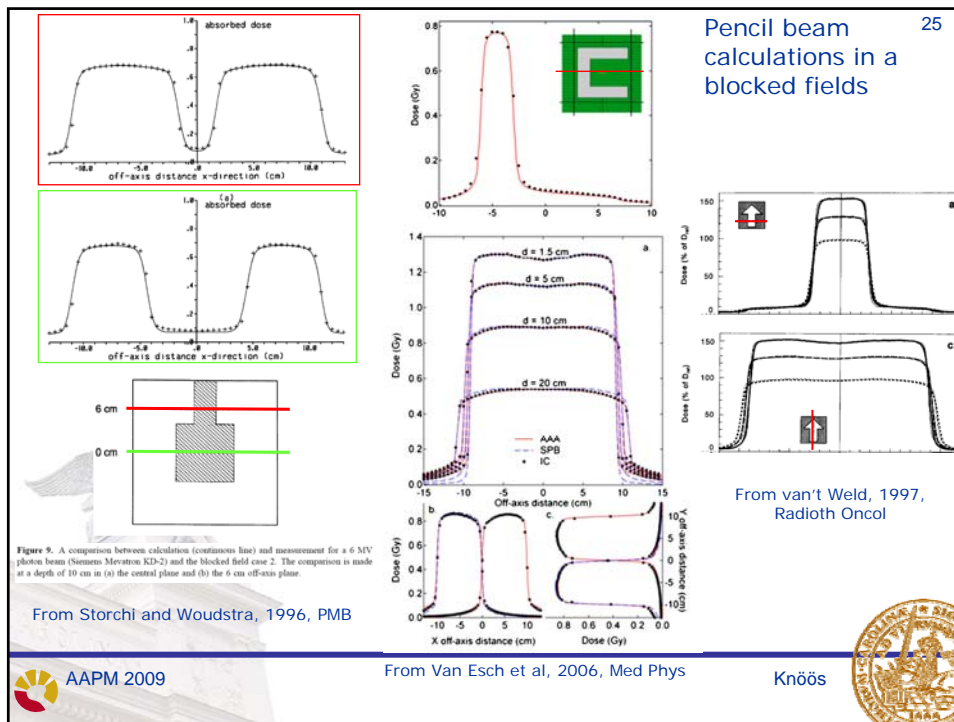
Carefully implemented algorithms together with accurate beam models works for most linacs

- Gamma-analysis, calc-meas
- Inside field after buildup
- Less than 0.5 % of points outside 3 mm/1 %
- One implementation 0.7 %

Cozzi et al, 2008, Z Med Physik

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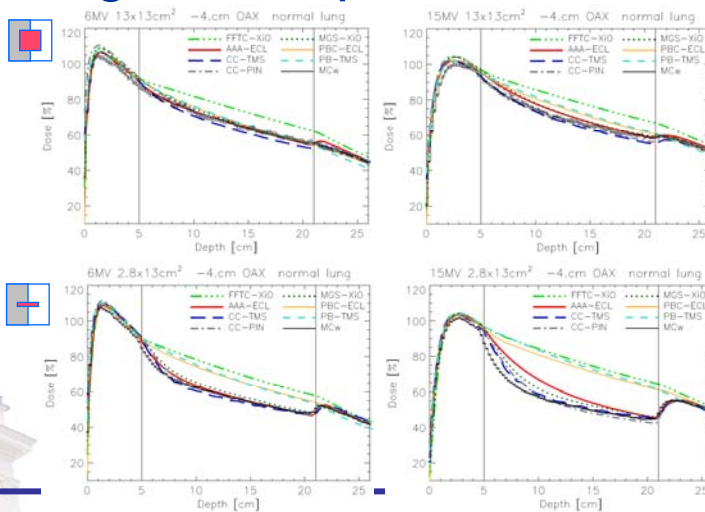
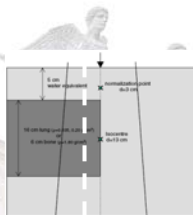
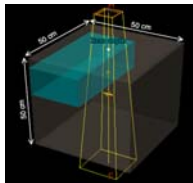


Convolution methods in homogeneous water

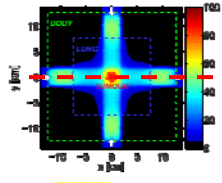
- Differences in beam modelling (not part of this SAM)
 - Head scatter
 - Electron contamination
 - Wedges/Blocks
 - MLC
- May lead to slightly different accuracy
- Basically all models perform well in water
 - Point, pencil or collapsed cone implementations



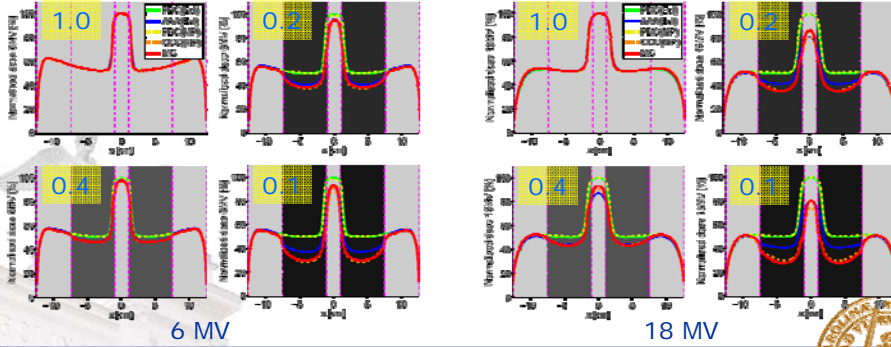
Comparison in inhomogeneous phantoms



Common implementation vs MC



Small solid lesion in low density lung tissue e.g. stereotactic treatment



6 MV

18 MV



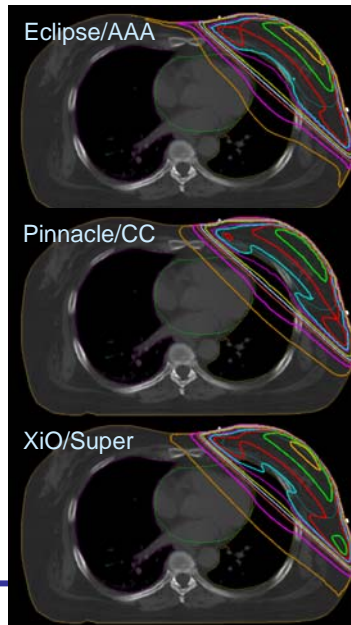
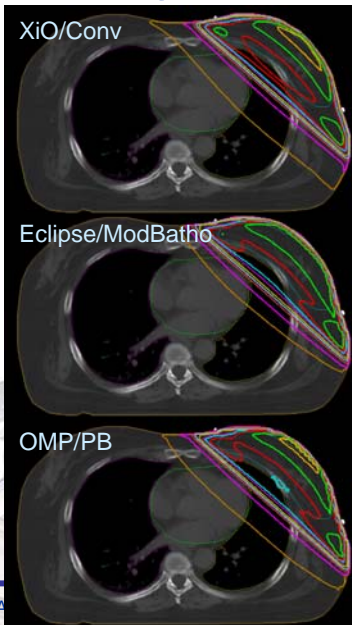
From Lasse Rye Aarup et al, RO, 2009

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Tangential treatment of breast

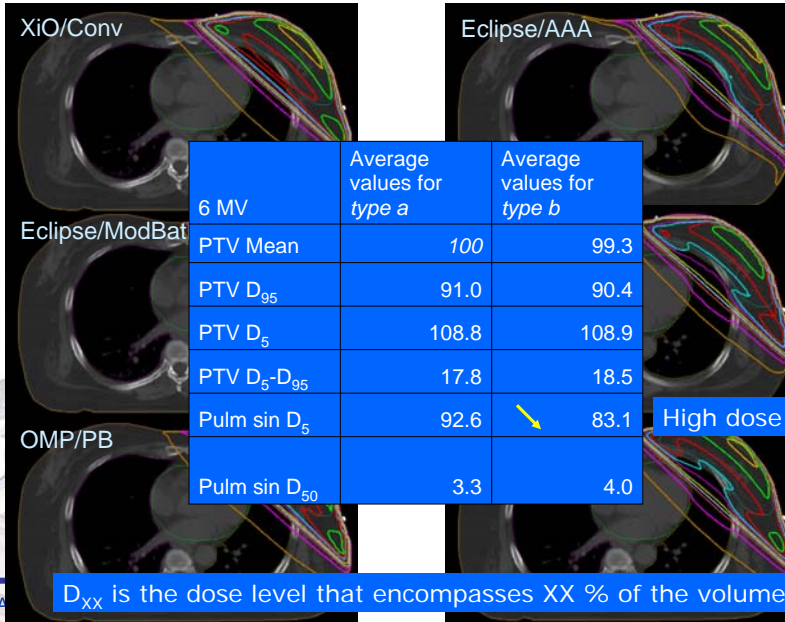
Knöös et al, 2006, PMB



Tangential treatment of breast

31

Knöös et al, 2006, PMB

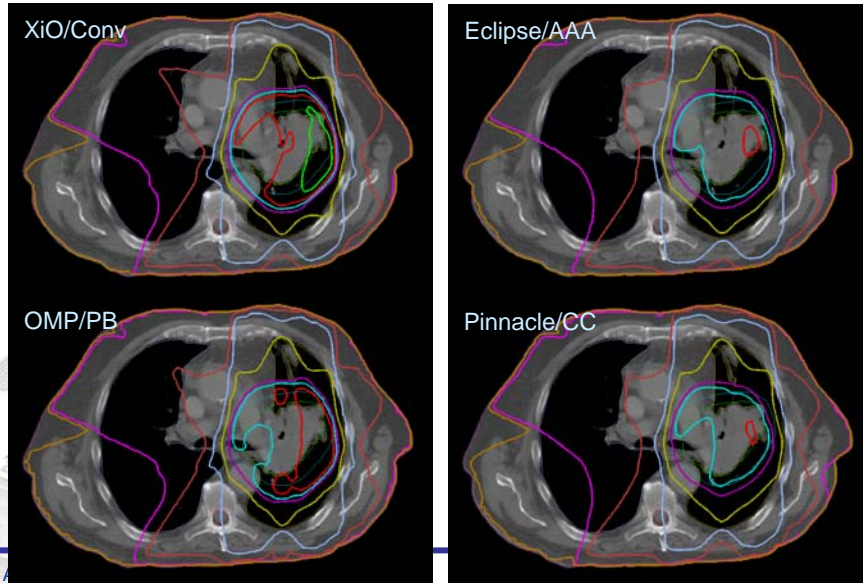


D_{xx} is the dose level that encompasses XX % of the volume

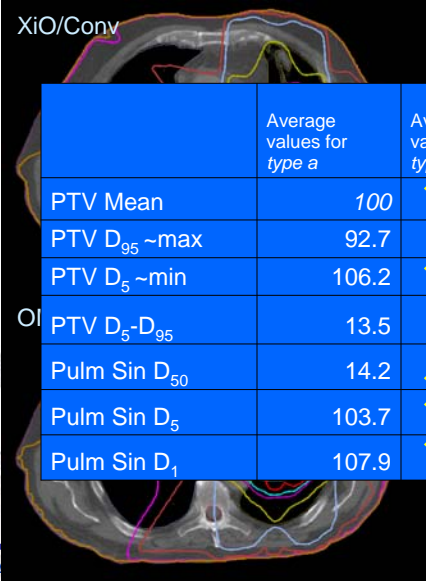
5 field 18 MV – lung

32

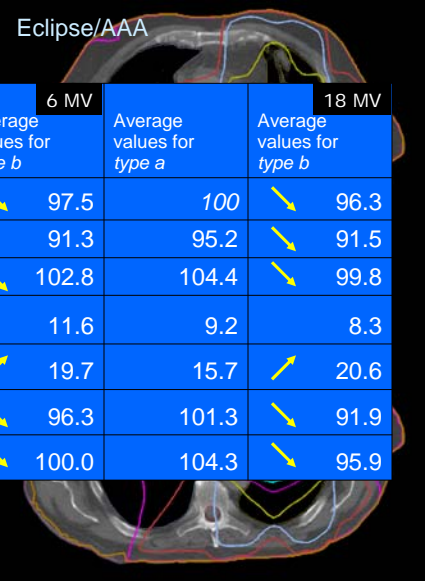
Knöös et al, 2006, PMB



5 field 18 MV – lung



XiO/Conv

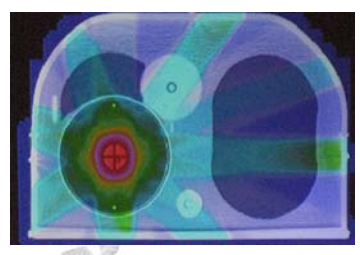


Eclipse/AAA

	Average values for type a	6 MV		18 MV	
		Average values for type b	Average values for type a	Average values for type b	
PTV Mean	100	↘	97.5	100	↘
PTV D ₉₅ ~max	92.7		91.3	95.2	↘
PTV D ₅ ~min	106.2	↘	102.8	104.4	↘
PTV D ₅ -D ₉₅	13.5		11.6	9.2	
Pulm Sin D ₅₀	14.2	↗	19.7	15.7	↗
Pulm Sin D ₅	103.7	↘	96.3	101.3	↘
Pulm Sin D ₁	107.9	↘	100.0	104.3	↘

Knöös et al., 2006, PMB

Results from RPC thorax phantom



- 15 cases planned with **type a**
 - 84% ± 16% of the pixels met the criteria (5%/5mm)
- 30 cases planned with **type b**
 - 99% ± 4% of the pixels met the criteria (5%/5mm)

AAPM 2008 TU-C-AUD B-3 P Alvarez et al



Conclusions – Dose changes

- Prostate
 - non-significant
- H&N
 - none (depending on accuracy of scatter integration) and air cavities (air or low dense water)
- Breast
 - slightly lower dose to breast and especially in lung in proximity to the target however larger irradiated lung volume
- Lung - PTV
 - 2-4 % lower average dose
 - Wider penumbra
- Lung (treated side)
 - 10 % lower dose to the highest irradiated parts of the lung
 - 5 % higher dose (15 => 20 %) to the lung (D_{50})
- Lung (healthy side)
 - Average dose identical (9.8-10.7 %)



Implications of introducing new and more accurate algorithms

- Significant changes in dose to target volumes and surrounding tissues especially when lung is involved
 - Consequences for assessment of dose-effect relationships
- Careful analysis of changes is required before adopting new algorithms
 - Retrospectively re-calculate plans when clinical outcome is known?
 - Construct new plans with older algorithms and re-calculate?
 - New plans with old prescriptions and new algorithms?
 - Optimize plans to the same biological effect on PTV and/or OAR?



Implications of introducing new and more accurate algorithms

- Significant changes in dose to target volumes and surrounding tissues especially when lung is involved
 - Consequences for assessment and effect
- Careful analysis of changes is required before adopting new algorithms
 - Do we need to produce new plans with older algorithms and re-calculate?
 - New plans with old prescriptions and new algorithms?
 - Optimize plans to the same biological effect on PTV and/or OAR?

Discussions are needed between physicists and oncologists to fully understand the effects and potential consequences



Conclusion

- Convolution methods are accurate
 - For low density regions – use models with lateral scaling
- Verification
 - Also Vendor's responsibility!
- Be careful when transferring to more accurate models but...
- **Important to do this!**

