

Adaptive Imaging: Review of techniques and update on state-of-the-art CT

Timothy P. Szczykutowicz, Ph.D. Associate Professor Radiology, Medical Physics, Biomedical Engineering

COI

TPS consultant GE Healthcare, supplies CT protocols under a licensing agreement to GE Healthcare, founder Protocolshare.org LLC, CAB and consultant to iMALOGIX, consultant Takeda Pharmaceuticals



Outline

- 1. Motivation for Patient Specific Imaging in CT: Cardiac
- 2. Heart Rate based Adaption
- 3. Motivation for Patient Specific Imaging in CT: Fluence modulation
- 4. Patient Specific Fluence Modulation



Human body

- It is not static
 - Blood is always moving through arteries/veins and perfusing through soft tissue
 - Heart is always moving
 - Lungs are always moving



Human body

- It is not static
 - Blood is always moving through arteries/veins and perfusing through soft tissue
 - Rate of blood movement is a function of cardiac output, local impedance (i.e. is there a plaque?)
 - > Opacification level is a function of blood pool volume



Anatomy and Physiology 2016 Rice University

ORIGINAL RESEARCH BRAIN

4D Digital Subtraction Angiography: Implementation and Demonstration of Feasibility

B. Davis, K. Royalty, M. Kowarschik, C. Rohkohl, E. Oberstar, B. Aagaard-Kienitz, D. Niemann, O. Ozkan, C. Strother, and C. Mistretta

ABSTRACT

BACKGROUND AND PURPOSE: Conventional 3D-DSA volumes are reconstructed from a series of projections containing temporal information. It was our purpose to develop a technique which would generate fully time-resolved 3D-DSA vascular volumes having better control and temporal evolution that which is available with CT or MD participation.



Journal List > Int J Biomed Imaging > v.2011; 2011 > PMC3166726



Int J Biomed Imaging. 2011; 2011: 467563. Published online 2011 Aug 28. doi: 10.1155/2011/467563 PMCID: PMC3166726 PMID: 21904538

Deconvolution-Based CT and MR Brain Perfusion Measurement: Theoretical Model Revisited and Practical Implementation Details

Andreas Fieselmann, ^{1, 2, 3, , *} Markus Kowarschik, ³ Arundhuti Ganguly, ⁴ Joachim Hornegger, ^{1, 2}, and Rebecca Fahrig ⁴

► Author information ► Article notes ► Copyright and License information Disclaimer



Human body

- It is not static
 - Heart is always moving

Speed			Ŧ
70	=	0.156586	
Millimeter / Second	\$	Miles per hour	\$

This may not seem fast, but consider we commonly use a RFOV of 25 cm and 512 voxels, that's a voxel size of 0.48 mm. In 1 second @ 70 mm/s...we will see blurring

Motivation for patient specific imaging in CT

Radiology

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Home > Radiology > VOL. 216, NO. 2 Cardiac Imaging

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In-Plane Coronary Arterial Motion Velocity: Measurement with Electron-Beam CT

Stephan Achenbach, Dieter Ropers, Jochen Holle, Gerd Muschiol, Werner G. Daniel, Werner Moshage

✓ Author Affiliations

Published Online: Aug 1 2000 https://doi.org/10.1148/radiology.216.2.r00au19457

\prec 🔧 Tools 🛛 < Share

Abstract

PURPOSE: To determine the speed of and changes in the speed of coronary arterial movement during the cardiac cycle with electron-beam computed tomography (CT).

MATERIALS AND METHODS: With electron-beam CT, 20 consecutive cross-sectional images were acquired at the mid right coronary artery (with 50-msec acquisition time, 8-msec intersection delay, 7-mm section thickness, and intravenous administration of 40 mL of contrast agent) in 25 patients. On the basis of the displacement of the left anterior descending, left circumflex, and right coronary arterial cross sections from image to image, movement velocity in the transverse imaging plane was calculated and was correlated with the simultaneously recorded electrocardiogram.

RESULTS: The velocity of in-plane coronary arterial motion varied considerably during the cardiac cycle. Peaks were caused by ventricular systole and diastole and by atrial contraction. The mean velocity was 46.6 mm/sec \pm 12.5 (SD). The mean velocity of right coronary arterial movement (69.5 mm/sec \pm 22.5) was significantly faster than that of the left anterior descending (22.4 mm/sec \pm 4.1) or the left circumflex coronary artery (48.4 mm/sec \pm 15.0). The lowest mean velocity (27.9 mm/sec) was at 48% of the cardiac cycle.

CONCLUSION: The lowest velocity of coronary arterial movement, which displays considerable temporal variation, was at 48% of the cardiac cycle.

Human body

- It is not static
 - Heart is always moving



Motivation for patient specific imaging in CT



NIH-PA Author Manuscript

NIH Public Access **Author Manuscript**

Published in final edited form as: IEEE Trans Med Imaging. 2006 March ; 25(3): 369-375.

Displacement and Velocity of the Coronary Arteries: Cardiac and

Respiratory Motion

Guy Shechter,

The Laboratory of Cardiac Energetics, National Institutes of Health, (NHLBI), Department of Health and Human Services (DHHS), Bethesda, MD 20892 USA, and the Department of Biomedical



Fig. 3.

Velocity of the left coronary tree during the cardiac contraction. The plot shows individual results for seven patients (thin lines) and the mean velocity (thick line). Velocities are plotted at (a) the LM ostium, (b) the LM bifurcation, (c) a point on the LAD 5 cm from the LM ostium, and (d) a point on the LCx 5 cm from the LM ostium.

Fig. 2.

Displacement of the RCA origin during a tidal breathing cycle. The plots show individual results for four patients (thin lines) and the mean displacement (thick line). One-dimensional displacements are with respect to the patient's (a) left-right, (b) inferior-superior, and (c) posterior-anterior axes. Positive displacements are toward the left, inferior, and posterior, respectively. Since end-expiration was the reference state, the shape of the 3-D magnitude displacement curve (d) has a minimum at end-expiration ($\rho = 0$), and a maximum near endinspiration ($\rho = \pm 1$).

• Human body

- It is not static
 - Lungs are always moving...even during breath holds!



Figure 3. (a) During inhalation, the diaphragm contracts, the abdomen is forced down and forward, and the rib cage is lifted. (b) The intercostal muscles also contract to pull and rotate the ribs, resulting in increasing both the lateral and anterior-posterior (AP) diameters of the thorax. [Reproduced from reference 226: J. B. West, *Respiratory Physiology: The Essentials*, Figures 3a,3b. © 1974, with permission from Lippincott Williams, and Wilkins.]



Figure 7.54 Example of a chest wall motion under "breath-hold" condition.



AAPM Report 91 "The management of respiratory motion in Radiation Oncology" "Computed Tomography" by Hsieh

Figure 7.53 Example of a respiratory motion curve measured at the chest wall under "normal" breathing condition.

for each conort of subjects. The motion is in three dimensions (51, A1, EK).						
Observer	Direction					
0	SI	AP	LR			
Barnes ⁸⁵ : Lower lobe	18.5 (9-32)					
Middle, upper lobe	7.5 (2-11)					
Chen et al.84	(0-50)					
Ekberg et al.26	3.9 (0-12)	2.4 (0-5)	2.4 (0-5)			
Engelsman et al.28:						
Middle/upper lobe	(2-6)					
Lower lobe	(2-9)					
Erridge et al.101	12.5 (6-34)	9.4 (5-22)	7.3 (3–12)			
Ross ⁷⁶ : Upper lobe		1 (0-5)	1 (0-3)			
Middle lobe		0	9 (0–16)			
Lower lobe		1 (0-4)	10.5 (0-13)			
Grills et al.91	(2-30)	(0-10)	(0-6)			
Hanley et al.77	12 (1-20)	5 (0-13)	1 (0-1)			
Murphy et al.87	7 (2–15)					
Plathow ²²⁰ : Lower lobe	9.5 (4.5-16.4)	6.1 (2.5-9.8)	6.0 (2.9–9.8)			
Middle lobe	7.2 (4.3-10.2)	4.3 (1.9–7.5)	4.3 (1.5-7.1)			
Upper lobe	4.3 (2.6-7.1)	2.8 (1.2-5.1)	3.4 (1.3–5.3)			
Seppenwoolde et al.67	5.8 (0-25)	2.5 (0-8)	1.5 (0-3)			
Shimizu et al.52		6.4 (2-24)				
Sixel et al.92	(0-13)	(0-5)	(0-4)			
Stevens et al.66	4.5 (0-22)					

 Table 2. Lung tumor-motion data. The mean range of motion and the (minimum-maximum) ranges in millimeters for each cohort of subjects. The motion is in three dimensions (SI, AP, LR).

Site Observer Deep Shallow Suramo et al.74 20 (10-30) Pancreas 43 (20-80) Bryan et al.75 20 (0-35) ----Weiss et al.89 Liver 13 +/- 5 ----Harauz et al.90 14 ----Suramo et al.74 25 (10-40) 55 (30-80) Davies et al.68 10 (5-17) 37 (21-57) Suramo et al.74 Kidney 19 (10-40) 40 (20-70) Davies et al.68 11 (5-16) ----Wade⁸⁰ 17 Diaphragm 101 Korin et al.79 13 39 Davies et al.68 12 (7-28) 43 (25-57) Weiss et al.89 13 +/- 5 ----Giraud et al.78 35 (3-95) ---Ford et al.86 20 (13-31) ----

Table 3. Abdominal motion data. The mean range of motion and the (minimum-maximum) ranges in millimeters for each site and each cohort of subjects. The motion is in the superior-inferior (SI) direction.

Breathing mode

AP: anterior-posterior; LR: left-right; SI: superior-inferior.

AAPM Report 91 "The management of respiratory motion in Radiation Oncology"

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• CCTA clinical reality

Patients will present with a wide range of heart rates

 There exists and optimal time post contraction which varies as function of HR → we define image time using a relative time post R peak or an absolute time post R peak



https://clinicalgate.com/physics-of-cardiac-computedtomography/

Heart Rate based Adaption

(d)



Motion artifact is highly dependent on phase/cycle location

(e)

Description Springer Link

European Radiology

Heart Rate based Adaption



Borut Marincek, Philipp A. Kaufmann, Hatem Alkadhi 🖂

coronary angiography with 64-slice CT

LC)

Optimal image reconstruction intervals for non-invasive

Cardiac First Online: 13 May 2006

European Radiology

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<u>BMC Med Imaging</u>. 2013; 13: 5. Published online 2013 Feb 1. doi: <u>10.1186/1471-2342-13-5</u>

PMCID: PMC3570333 PMID: 23375107

Defining the mid-diastolic imaging period for cardiac CT – lessons from tissue Doppler echocardiography

James M Otton,^{1,2} Justin Phan,² Michael Feneley,^{1,2} Chung-yao Yu,¹ Neville Sammel,¹ and Jane McCrohon^[21,2]

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Abstract



Heart Rate based Adaption



Heart Rate based Adaption



F	IR		
Lower Limit	Upper Limit	Peak 1	Peak 2
30	50	75	
51	60	75	
61	65	70 – 80	
66	70	70- 80	
71	85	70- 80	40 - 55
86	100	40 - 55	
101	200	40 - 60	

Auto Gating Configuration								
Profile: CCTA Profile Name: GE CCTA Rename								
Variable Beat to Beat Threshold 6 Irregular Threshold 2 Highly Irregular Threshold 4								
Average Heart Rate (BPM)								
	3	5	1	61				200
	Stable	0.25 e/mt 75-75% 100% HRV: 0.5 AG Center Phase	0.35 s/rat 76-75%, 100%, SSF 40-30%, 100% 5-95%, 20% HRV: 0.8 AG Center Phase	0.35.9/rot 70-80%, 100%, SSF 40-90%, 100% 5-95%, 20% HRV: 1.0 AG Center Phase, SSF	0.28.5/rpt 70-80% 100%, SSF 49-30.5%, 100% 5-95%, 20% HRV: 1.0 AG SmartPhase, SSF	0.20 s, ot 70-80%, 100%, SSF 40-55%, 100%, SSF 5.05%, 20% HRV: 1.0 AG SmartPhase, SSF	0.90.55%. 40-55%. 40-55%. 5-95%. 5-95%. HRV: AG SmartPhase, SSF	0.00 stat 40-60% 100%, SSF 5-95%, 20% HRV: 1.0 AG SmartPhase, SSF
Rhythm	Variable	0.35 s/rot 76-75%, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 3.0 AG Center Phase	0.35 s/rot 76-75%, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 5.0 AG Center Phase	0.35 s/rot 70-80%, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 5.0 AG Center Phase, SSF	0.28 s/rot 70-80%, 100%, SSF 40-56%, 100% 5-95%, 20% HRV: 5.0 AG SmartPhase, SSF	0.28 s/rot 40-80%, 100%, SSF 40-56%, 100% 5-95%, 20% HRV: 5.0 AG SmartPhase, SSF	0.28 s/rot 40-55%, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 5.0 AG SmartPhase, SSF	0.28 s/rot 40-60%, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 5.0 AG SmartPhase, SSF
H	Irregular	0.35 s/rot 76-75%, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.5 AG/2 Scans Center Phase	0.35 s/rot 75-75%, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.8 AG/2 Scans Center Phase	0.35 s/rot 70-80%, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 1.0 AG/2 Scans Center Phase	0.28 s/rot 70-80%, 100%, SSF 40-56%, 100% 5-95%, 20% HRV: 1.0 AG/2 Scans SmartPhase, SSF	0.28 s/rot 40-80%, 100%, SSF 40-56%, 100% 5-95%, 20% HRV: 1.0 AG/2 Scans SmartPhase, SSF	0.28 s/rot 200-400ms, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.0 AG/2 Scans SmartPhase, SSF	0.28 s/rot 175-376ms, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.0 AG/2 Scans SmartPhase, SSF
	Highly Irregular	0.35 s/rot 260-700ms, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.0 AG/2 Scans Center Phase	0.35 s/rot 225-500ms, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.0 AG/2 Scans Center Phase	0.35 s/rot 200-450ms, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.0 AG/2 Scans SmartPhase	0.28 s/rot 200-450ms, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.0 AG/2 Scans SmartPhase, SSF	0.28 s/rot 200-400ms, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.0 AG/2 Scans SmartPhase, SSF	0.28 s/rot 200-400ms, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.0 AG/2 Scans SmartPhase, SSF	0.28 s/rot 175-376ms, 100%, SSF 40-55%, 100% 5-95%, 20% HRV: 0.0 AG/2 Scans SmartPhase, SSF

Scan Settings HR: 30 - 50; Stable

Slowest Allowable Rotation Speed 0.35 s/rot							
Acquisition Part 1 🗹 75 - 75 🕅 🕶 mA 100 % Widen for SSF 📃							
Acquisition Part 2 🔲 40 - 55 🔭 mA 100 % Widen for SSF 📃							
Acquisition Part 3 🔲 5 - 95 🔭 mA 20 % Widen for SSF 📃							
HR Variation Type Scale Factor HR Variation Amount 0.5							
Adaptive Gating 🗹 🛛 Repeat Acquisition 🔲							
Secondary Recon Phase Type Center Phase (1) Secondary Recon SSF 							
Display Alert None							
Alert Message							

Heart Rate based Adaption



CCTA clinical reality

Patients will present with heart rate variability

• patients with irregular HRs (afib), or with PVCs

Now we need to talk about data collection.

Stable HR has HR variation ~ 5 bpm, variable >5 bpm

Image from 128-slice Dual Source CT: How Does it Work and What Can it Do? By Bruesewitz et al. Mayo Clinic 2010 RSNA



Heart Rate based Adaption

Heart rate range	Gantry speed	Pitch
30 to 42 BPM	0.4 sec	0.18
43 to 49 BPM	0.4 sec	0.20
50 to 59 BPM	0.4 sec	0.23
60 to 69 BPM	0.4 sec	0.26

Example table of scan pitches as a function of HR

<u>"Step and shoot" axial/sequential mode</u> All vendors will just widen the time the x-rays are on for a given bed position or collect multiple spins one location if an irregular beat is detected

High pitch method "Flash mode"

and cover the entire heart in a single

beat using a helical/spiral scan mode

Slow pitch helical/spiral mode

are detected you would use a lower pitch

Wide axial/sequential mode

Your detector can cover the entire heart, so you sit over the heart with ECG trace on scan when you predict your target phase will be Scan for a longer time when the HR is high or you anticipate an irregular beat

Image from 128-slice Dual Source CT: How Does it Work and What Can it Do? By Bruesewitz et al. Mayo Clinic 2010 RSNA

Dual Source cCTA Imaging Guidelines



Flash / Turbo Flash Mode			Adaptive Cardio Sequence (Prospective Gating)		Retrospective ECG Gating	
Heart Rate	Flash: <65bpm Force: <78bpm			Any Heart Rate		Any Heart Rate
HR Stability	Stable			At high HR's need to be stable		Any Wave Form
Contrast Amount	Physicians discreti	on	Sc	can time X injection rate plus 10 cc's	Scan tim	e x injection rate plus 10 cc's
Injection Rate	5 to 6 ml/s (faster is b	etter)		5 to 6 ml/s (faster is better)	5 to	o 6 ml/s (faster is better)
Rotation Time	Flash/Drive: 0.28, Force: 0.25			Flash/Drive: 0.28, Force: 0.25	Flash/Drive: 0.28, Force: 0.25	
CARE kV	ON			ON	ON	
Quality ref mAs	Use Default			Use Default	Use Default	
CARE Dose 4D	ON			ON		ON
Iterative Reconstruction	ON			ON		ON
Patient Dose	Less than 1 mSv (10	0kV)		Depends on ECG pulsing range	Depends on ECG pulsing range, (min dose available)	
When to use	Rule out CAD			Routine with HR variability	Atrial Fibrilation, very difficult case ECG editing capability	
Cardiac Function	N/A			Yes	Yes Yes	
Beta Blockers	Physicians discretion			Physicians discretion	Physicians discretion	

- Limitations of the gated Flash mode:
 - The heart rate must be regular. To scan using a high pitch, the table requires about 1 sec to be accelerated. The scanner must accurately predict the timing of future R-waves in order to synchronize the xray-on time with the diastolic phase.
- Limitations with wide axial scanners and prospective scanning
 - Dose reduction potential is reduced
 - Need to widen gating window
 - Need to "double scan" if a messed up beat is detected

- So in summary
 - There exist multiple modes for cardiac scanning
 - There are optimal modes for different heart rates
 - There are optimal modes for different degrees of HR irregularities
 - There are different target gating windows for different heart rates
 - There are different strategies for dealing with HR irregularities
- Current state of the art in cardiac scanning will have the scanner picking the mode and target window for the operator
- Honestly, the peer reviewed literature is not the place to understand state of the art in cardiac scanning, get your apps person to send your vendors whitepapers and user manuals for cardiac mode.

Outline

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Human body

- It varies in size
 - Region to region
 - Angularly
 - Patient to patient
 - As a function of position

Human body

- It varies in size
 - Region to region
 - head vs. pelvis



46 cm



Multiple orders of magnitude changes will occur in detector signal due to body region size changes!

Left is non log, right is log plot of detector signal as a function of soft





- Human body
 - It varies in size
 - Angularly

> projection through lateral direction longer than AP/PA for all regions except the head



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DOI: 10.1002/acm2.12223

IMAGING PHYSICS

Evaluation of AAPM Reports 204 and 220: Estimation of effective diameter, water-equivalent diameter, and ellipticity ratios for chest, abdomen, pelvis, and head CT scans

Christiane S. Burton¹ | Timothy P. Szczykutowicz²

TABLE 2 Elliptical ratio (LAT/AP) calculation for patients of routine adult (abdomen and pelvis), adult chest ([†]subset of adult chest), and pediatric cases.

Data set	Mean ellipticity ratio (LAT/AP)	Std. Dev.	Min–Max
Adult abdomen pelvis	1.48	0.22	1.20–1.94
Adult chest	1.60	0.23	1.21–2.07
Adult shoulder †	2.28	0.22	1.08-3.34
Adult thorax ^{\dagger}	1.51	0.21	1.09–1.98
Adult abdomen only †	1.38	0.20	1.16–1.73
Pediatric abdomen pelvis	1.53	0.30	1.07–1.75
Adult head	0.85	0.08	0.83–0.87

vide data for the future e-scale confirmation of the ith size surrogate data for ports, and additionally proerent body regions. pur analysis including data I pediatrics. We calculated id in AAPM Reports 204/

WILEY



Kids have big heads... that is why they are cute and why we treat them as adults after they turn 7

Human body

- It varies in size
 - Patient to patient
 - newborn to bariatric adult



Imaging bariatric patients is difficult. Rule of thumb in CT is we need 2x dose for every 4 cm of tissue to maintain image noise. So if we like the scan of a 30 cm

person, to get to 50 cm we need 5x dose doublings. If we were using 300 mA for the 30 cm scan, then we need 1,500 mA for the bariatric person!

Human body

- It varies in size
 - Patient to patient
 - newborn to bariatric adult

2 months old



Extremely fit 27 year old*



<u>"white</u> out" bariatric patient



*its younger me 🙂

Human body

- It varies in size
 - As a function of position
 - > prone versus supine, with respiratory state diaphragm/liver move multiple cm which changes attenuation

Same patient, three different positionings and three different attenuation distributions







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ratio

Ellipticity

- State of the art mA modulation
 - Adjustment of tube output as a function of z axis position, or both as a function of z position and gantry angle
 - Can reduce dose up to 60% in some cases¹





Image from *Computed Tomography* by Willi A Kalender 2005 Phantom Size mA modulation can handle size and ellipticity changes

Image from: McCullough, C.H. *et al. "CT Dose Reduction and Dose Management Tools: Overview of Available Options"* Radiographics **26** 2006

¹Kalender *et al.* "Dose reduction in CT by anatomically adapted tube current modulation II: Phantom experiments" Med. Phys. **26** 1999

- Current state of the art in mA modulation is organ dose modulation
 - Reduces mA when the tube is directly irradiating a radio-sensitive organ
 - Typically, ~30% dose reduction possible



Previous state of the art, angular mA modulation



- Patient Specific Fluence ModulationDixon, M. T., Loader, R. J., Stevens, G. C., & Rowles, N. P. (2016). An evaluation of organ dose modulation on a GE optima CT660-computed tomography scanner. Journal of applied clinical medical physics, 17(3), 380-391.
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That was mA modulation, what about kV?



Optimal Beam Energy Selection in CT

• Key references for understanding the optimal beam energy in

Application- and patient size-dependent optimization of x-ray spectra for CT

Willi A. Kalender,^{a),b)} Paul Deak,^{a)} Markus Kellermeier,^{a)} Marcel van Straten,^{a),c)} and Sabrina V. Vollmar^{a)}

Institute of Medical Physics, University Erlangen-Nürnberg, Henkestr. 91 91052 Erlangen, Germany

(Received 21 August 2008; revised 20 November 2008; accepted for publication 6 January 2009; published 25 February 2009)

Although x-ray computed tomography (CT) has been in clinical use for over 3 decades, spectral optimization has not been a topic of great concern; high voltages around 120 kV have been in use since the beginning of CT. It is the purpose of this study to analyze, in a rigorous manner, the energies at which the patient dose necessary to provide a given contrast-to-noise ratio (CNR) for various diagnostic tasks can be minimized. The authors used cylindrical water phantoms and quasianthropomorphic phantoms of the thorax and the abdomen with inserts of 13 mm diameter

Automatic selection of tube potential for radiation dose reduction in CT: A general strategy

Lifeng Yu,^{a)} Hua Li, Joel G. Fletcher, and Cynthia H. McCollough Department of Radiology, Mayo Clinic, Rochester, Minnesota 55905

(Received 13 July 2009; revised 23 October 2009; accepted for publication 27 October 2009; published 10 December 2009)

Purpose: To optimize radiation dose efficiency in CT while maintaining image quality, it is important to select the optimal tube potential. The selection of optimal tube potential, however, is highly dependent on patient size and diagnostic task. The purpose of this work was to develop a general strategy that allows for automatic tube potential selection for each individual patient and each diagnostic task.

Methods: The authors propose a general strategy that allows automatic adaptation of the tube potential as a function of patient size and diagnostic task, using a novel index of image quality, "iodine contrast to noise ratio with a noise constraint (iCNR_NC)," to characterize the different

Optimal Beam Energy Selection in CT



Medical Physics, Vol. 36, No. 3, March 2009

Optimal Beam Energy Selection in CT



https://www.edu-quip.co.uk/prod/27573/matrix-springer-3-way-see-saw

	Canon	GE	Philips	Siemens		
mA AEC	^{SURE} Exposure 3D	smartmA	DoseRight	CAREDose 4D		
kV AEC	SURE kV	kV Assist	{}**	CARE kV		
Organ sparing	OEM*	ODM	{}	X-CARE		
Quality target	SD	NI	DRI	Quality ref.mAs		
Quality target to dose	$SD\propto rac{1}{\sqrt{CTDI_{vol}}}$	$NI \propto \frac{1}{\sqrt{CTDI_{vol}}}$	$CTDI_{vol} \propto (1.12)^{DRI-24***}$ $CTDI_{vol} \propto$ $(0.88)^{24-DRI****}$	Quality ref.mAs ∝ CTDIvol		
Quality target to noise	SD≈ Image Noise	NI ≈ Image Noise	DRI $\propto 24 - \frac{\ln(image \ noise)}{2\ln(1.12)} ***$ DRI $\propto 24 + \frac{\ln(image \ noise)}{2\ln 0.88} ****$	Quality ref.mAs $\propto \frac{1}{\sqrt{Image Noise}}$		
Quality Target with patient size	Non linear, not adjustable	constant	Non linear, not adjustable	Non linear, adjustable (strength setting)		
Organ Region boost	Allows you to set different SD for different scan regions	{}	Liver Brain DoseRight	{}		
*In CT fluoroscopy mode only **Philips recommends performing a dual energy scan **** For dose decrease below reference patient DRI						

Table 1.1: A summary of x-ray fluence modulating methods. Values in parenthesis represent ideal parameter values assuming the system could be optimized beyond the prototype stage according to the authors of the respective works.

Method	Separable Modulation Function	Used Clinically	Temporal Modulation	Spatial Modulation ^a	Applicability to FFMCT ^b
bowtie filtering 13	no	yes	none	infinite	low
piston driven mask equaliza- tion ^{17–21}	yes	no	$3~{\rm min}~({<}5~{\rm sec})$	0.159 cm	low
DBA (older version) $^{22-31}$	yes	no	$\frac{120~{\rm min}~(1~{\rm min})^{~22}}{3.5~{\rm min}.^{~23}}$	$0.3~\mathrm{mm}^{22}$	low
$rotating$ aperture 32,33	no	no	2.5 ms	none	possible
ROI filter ^{34,35}	no	no	none	fixed ROI	possible
z-axis collima- tion ¹³	no	yes	none	variable	possible
wedges ³⁶	no	yes	none	infinite	possible
slot scanning ^{37–40}	yes	yes	9 sec. ³⁸	$3.5~{\rm cm}^{38}$	$\operatorname{possible}^{c}$
Volume of Inter- est 41,42	yes	no	${\approx}150~{\rm ms}^{41}$?	$low-high^d$
${ m mA~modulation}^8$	no	yes	90% in 1 \sec^8	none	possible
$\mathrm{SBDX}^{43,44}$	yes	no	21-62 ${ m ms}^{43}$	$16~\mathrm{mm}^{\mathrm{e}43}$	high
electronic bowtie ⁴⁵	yes	no	?	?	high
${\rm TomoTherapy^{TM}f_{46}}$	yes	not for FFMCT	${\approx}3070~\mathrm{msec}^{47}$	$6.25 \mathrm{~mm}^{\mathrm{g}}$	high
DBA-FFMCT ⁴⁸	yes	no	?	?	high

^aDimension of modulation at the modulator plane, not at the object or image plane unless otherwise noted. ^bApplication to non-separable FFMCT, in the opinion of the author.

^cSlot scanning can be thought of as a combination of mA modulation and x/z axis collimation in CT.

^dLow applicability to FFMCT when the goal is uniform detector fluence, high applicability when the goal is VOI. ^eDimension of modulation at the image plane.

 $^{\rm f}$ Use of this method for FFMCT has not been studied, but in the opinion of the author it is possible. $^{\rm g}$ Leaf size projected to isocenter.

Patient Specific Fluence Modulation

Circa ~2012, the concept of fluence modulated imaging in x-ray imaging was not new. Many approaches had been studied and were in clinical use.







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Bowtie filter works perfectly when

- Patient object is homogeneous and cylindrical
- Patient is perfectly centered on isocenter

Advantages

- Decrease dose and scatter by ~50%¹
- Increase noise uniformity²

Homogeneous and centered

Patient Specific Fluence Modulation



Can double surface dose! Can increase CTDI, by ~20%









"Computed Tomography Principles, Design, Artifacts, and Recent Advances" by Jiang Hsieh









Physics in Medicine & Biology

PAPER

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Experimental realization of fluence field modulated CT using

digital beam attenuation

T P Szczykutowicz¹ and C A Mistretta^{1,2,3} Published 20 February 2014 • 2014 Institute of Physics and Engineering in Medicine <u>Physics in Medicine & Biology, Volume 59, Number 5</u>

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Figures - References -

Article information

Abstract

Tailoring CT scan acquisition parameters to individual patients is a topic of much research in the CT imaging community. It is now common place to find automatically adjusted tube current options for modern CT scanners. In addition, the use of beam shaping filters, commonly called bowtie filters, is





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Physics in Medicine & Biology

PAPER

A prototype piecewise-linear dynamic attenuator

Scott S Hsieh^{1,2,4}, Mark V Peng³, Christopher A May³, Picha Shunhavanich³, Dominik Fleischmann¹ and Norbert J Pelc^{1,3} Published 10 June 2016 • © 2016 Institute of Physics and Engineering in Medicine Physics in Medicine & Biology, Volume 61, Number 13

Figures - Ret Article informa Abstract The piecewise-lin personalizing the have shown bene implementation.







No DBA



DBA





Hopkins group- noise uniformity better with FFMCT relative to static





bowtie



Can help us avoid situations like this, bad positioning caused large noise non uniformity



 Stanford group (Hsieh and Pelc) have looked at how FFMCT can help with photon counting CT via reducing dynamic range needs of detector





- What about for VOI imaging?
 - Need to predefine a SNR "prescription" for the image
 - Then the required wedge positions must be calculated
- Our first implementation
 - Max signal (minimum wedge thickness) to where ROI intersects wedgelet, minimum signal (maximum wedge thickness) otherwise



- Clinicians often do not need good image quality everywhere!
- Use FFMCT to provide region specific SNR enhancement/suppression

FFMCT implemented on a clinical MVCT scanner



Tomotherapy scanners already have a set of beam modulators, so we used them to do FFMCT!





Scan focused on "VOI 4" is shown here

 Toronto group looked at VOI imaging using different types of fluence modulation devices

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Compensator models for fluence field modulated computed tomography

Steven Bartolac 🗙, David Jaffray

First published: 18 November 2013 | https://doi.org/10.1118/1.4829513 | Cited by: 7

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SECTIONS

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- FFMCT can be thought of as a way to actually weight data during the collection process, in stead of post acquisition during reconstruction like one can do with noise weighting schemes in an iterative/non-linear reconstruction algorithm (D term below weights data based on its signal strength)
 - FFMCT would allow one to increase signal levels for highly attenuating rays OR
 - FFMCT would allow one to give up on highly attenuating rays and give them little to no dose

$$\hat{\mu} = \arg\min_{\mu} \left\{ \frac{1}{2} \left(p - A\mu \right)^T D \left(p - A\mu \right) + G(\mu) \right\}$$





Feel free to contact me at tszczykutowicz@uwhealth.org



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