

# CT Contrast Parameters for the Medical Physicist

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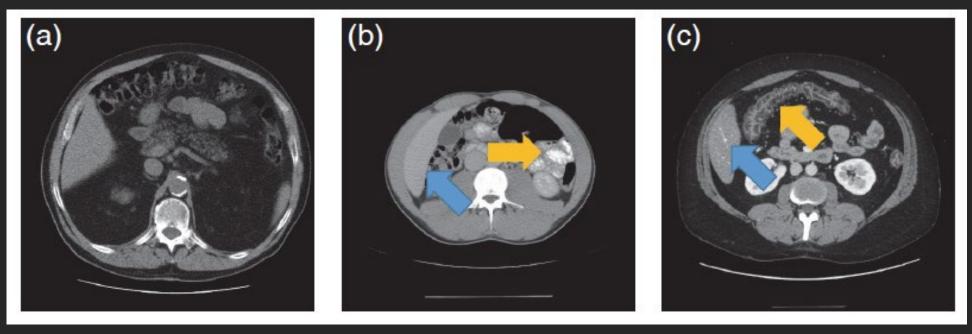
\_\_\_\_@Prof\_TimStick

# **Conflict of Interest**

- TPS supplies CT protocols to GE Healthcare under a licensing agreement, TPS is a consultant and on an advisory board to GE Healthcare, TPS receives research support from GE Healthcare
- TPS is on the MAB of Imalogix LLC
- TPS is a consultant to AstoCT LLC, and cybermed.ai (DBA RadFlow), AiDoc, iSchemaView
- TPS receives book royalties from Medical Physics Publishing

No contrast

Positive Oral and IV contrast (parenchymal phase) Negative Oral and IV contrast (liver arterial phase)



"The CT Handbook: Optimizing Protocols for Today's feature-rich scanners" By Tim Szczykutowicz. Medical Physics Publishing 2020

#### CT CONTRAST 101 CO2 gas

(a)

(c)

contrast agent

Same pt no CO2 (b)

Note: the localizers even show the CO2

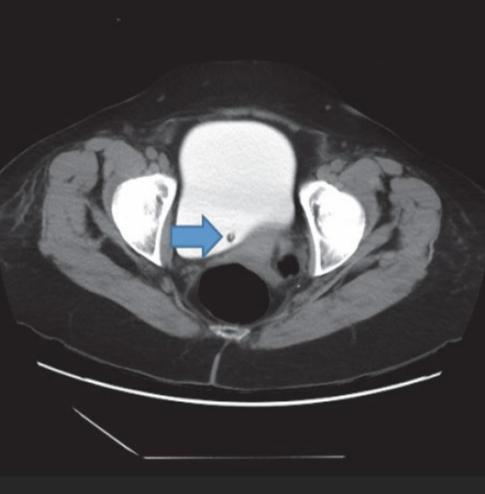
"The CT Handbook: Optimizing Protocols for Today's feature-rich scanne

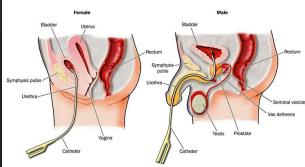
Both scans have positive oral agent



Positive oral contrast introduced via Foley catheter

(arrow shows catheter, don't confuse this with an artifact)



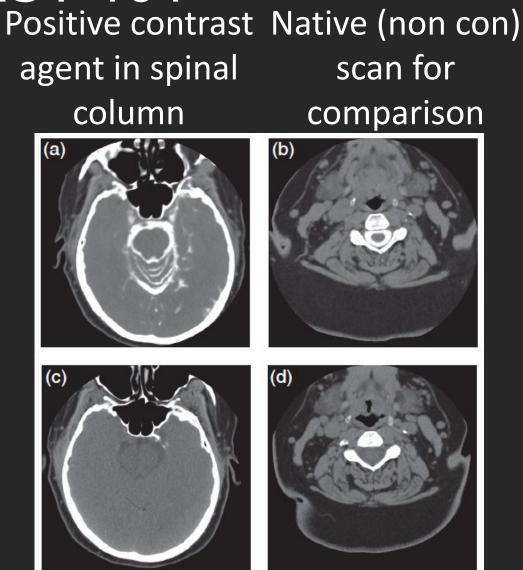


https://www.ausmed.com/ cpd/articles/urinarycatheter

"The CT Handbook: Optimizing Protocols for Today's feature-rich scanners" By Tim Szczykutowicz. Medical Physics Publishing 2020

Introducing contrast agent into the spinal column is called CT Myelography

(into joint space is called arthrogram)



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#### General Guidance

Patient with moderate or severe allergy to lodine based contrast

- . Give these patients Barium Readicat, or perform the scan without oral contrast. Consult the attending radiologist for guidance.
- If the patient is going to surgery post imaging, then give them lohexol, surgery prefers this, please consult the attending radiologist for approval.

Outpatients: 1 dose = 4mL of lohexol in 200 mL of clear liquid.

- Abd CT: 2 doses (400 mL of oral contrast)
- Abd/Pel CT: 4 doses (800 mL of oral contrast)
- Last dose on CT table: 8mL of lohexol in 200 mL of clear liquid.
  - X-Large cups = 800 mL-(1x q 1 hour)
  - Large cups = 400 mL-(2x q 30)
  - Small cups = 200 mL- (4x q 15)
  - · Patients have their choice of to mix the lohexol with lemonade or water.
- Billing = All oral contrast (including COT)

Inpatients: 1 dose = 4mL of lohexol in 200 mL of clear liquid.

- · Abd CT: 2 doses q 30 minutes
- Abd/Pel CT: 4 doses q 30 minutes
- Billing = None (Patient is billed on the floor)

#### Bariatric Oral

- If the patient is a bariatric post-op patient they will not drink up on the floor. Rather, they will get one 2x concentrated dose on CT table: 8mL of lohexol in 200 mL of clear liquid. If you have questions
  please ask the protocolling radiologist.
- Billing = 8 ml of lohexol





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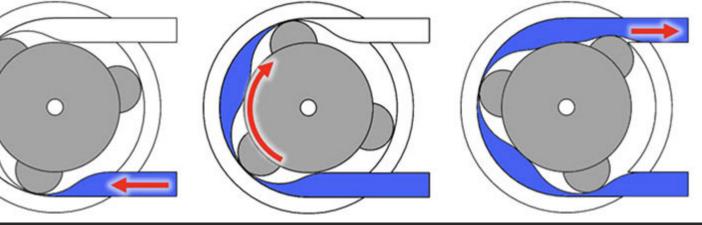






Piston based pump... simple, the plunger pushes the agent out

Peristaltic pump, the rotating action pushes agent along a flexible tube



https://dienerprecisionpumps.com/positive-displacement-pumps/

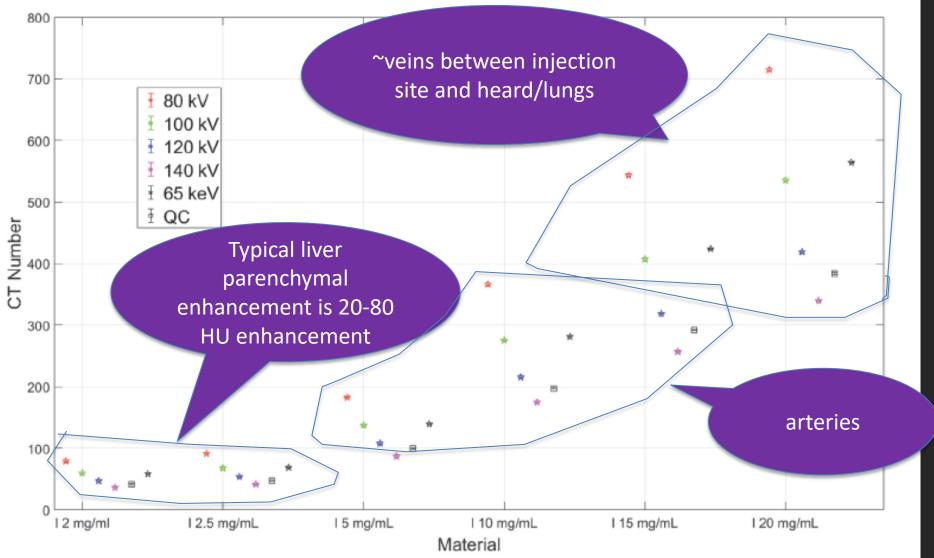
#### Air/CO2

Generates CT number ~-1000, "usually doesn't get diluted"

#### Used for CTC

#### lodine

Workhorse for CT. Usually it will always be diluted by blood (IVC administration) or water (oral) or urine (catheter injection)



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Material	80 kV	100 kV	120 kV	140 kV	N/A
Water	0	0	0	0	[-4 4]**
Air	-1,000	-1,000	-1,000	-1,000	[-1005 -995]**
Fat	-152*	-111*	-89*	-69*	[-100 -80]**
Brain	47*	43*	39*	37*	
Soft Tissue	62*	58*	54*	52*	
Solid Cortical Bone	3,760*	2,590*	1,940*	1,330*	[≈200 > <b>1</b> 000]**
Pure Calcium	9,570*	5,960*	3,950*	2,090*	-
Pure Iodine	405,000*	267,000*	180,000*	93,200*	
Iodine Contrast	See footnote a	See footnote a	See footnote a	See footnote a	
Relative Iodine Enhancement <sup>b</sup>	1.68	1.27	1	0.826	
Relative Iodine Enhancement <sup>e</sup>	1.70	1.28	1	0.81	
Kidney					[20 40]**
Pancreas					[30 50]**
Blood					[50 60]**
Liver					[50 70]**
PMP					-200***
Low-Density Polyethylene					-100***
Polystyrene					-35***
Acrylic					120***
Delrin®					340***
Teflon®					990***

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$$\mu(E) = a_1 \times \frac{1}{E^3} + a_2 \times f_{KN}(E)$$

$$Z \text{ is 5-7 for soft tissue}$$

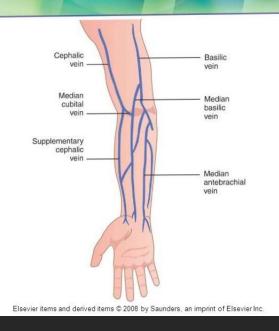
$$a_1(x, \gamma) = C \times \rho_e \times Z^{n-1},$$

$$a_2(x, \gamma) = \rho_e,$$

Become familiar with these equations... and you can understand every "lower kV equals dose reduction for CTA" paper ever written

#### 1. Inject via antecubital vein

#### **Antecubital Veins**



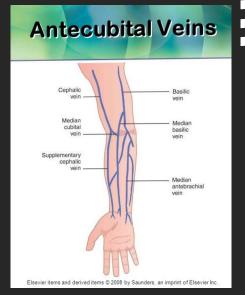
3. Blood goes to rightventricle and goes outpulmonary artery to lungs

4. Returns via pulmonary vein and goes to left side of heart to be pushed out aorta via left ventricle.

5. First stop is supplying heart via coronaries. Then head via carotid and vertebral vessels.

# 2. Blood goes into SVC and then into right atrium

#### 1. Inject via antecubital vein



Min: -48 Max: 2137 (HU) Average: 958.75 StdDev: 580.81 (HU) Area: 49.18 mm² Perim: 26.61 mm

> 1. Vein coming from injection site 958 HU

Min: 331 Max: 785 (HU) Average: 516.71 StdDev: 132.65 (HU) Area: 32.73 mm<sup>2</sup> Perim: 20.70 mm

2. SVC 516 HU

3. Pulmonary arteries 389 HU

Min: 356 Max: 429 (HU) Average: 389.45 StdDev: 15.45 (HU) Area: 105.00 mm<sup>2</sup> Perim: 36.85 mm

50 m

Min: 257 Max: 377 (HU) Average: 322.01 StdDev: 26.67 (HU) Area: 152.73 mm<sup>2</sup> Perim: 44.01 mm

4. Descending aorta 322 HU

Numbers increase with ROI locations via path blood follows from injection site → blood/I mix HU goes down as we move from site because of blood mixing diluting agent

Contrast time of arrival will vary from person to person, but assuming an antecubital injection, values will generally be in the range of 7 to 10 seconds for the pulmonary artery, 12 to 15 seconds for the ascending aorta, 15 to 18 seconds for the abdominal aorta, and 30 to 40 seconds for hepatic parenchyma.



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Another reference Bae, K. T., J. P. Heiken, and J. A. Brink. (1998). "Aortic and hepatic contrast medium enhancement at CT. part i. prediction with a computer model." *Radiology* 207(3):647–55.



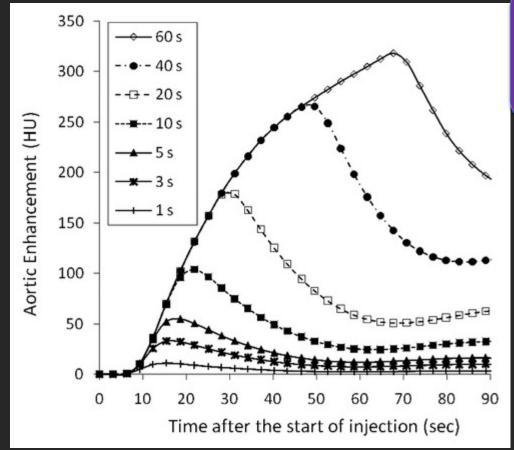
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#### Intravenous Contrast Medium Administration and Scan Timing at CT: Considerations and Approaches

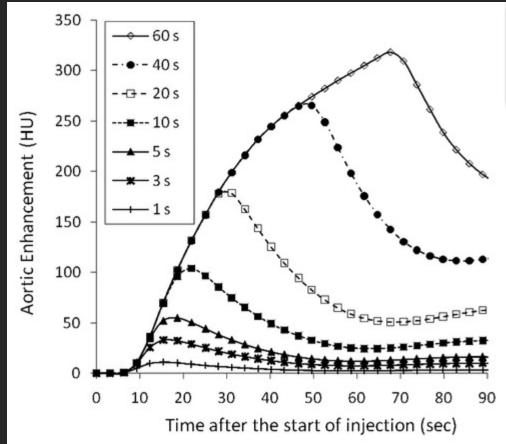
Kyongtae T. Bae 🖾

✓ Author Affiliations

Published Online: Jul 1 2010 https://doi.org/10.1148/radiol.10090908

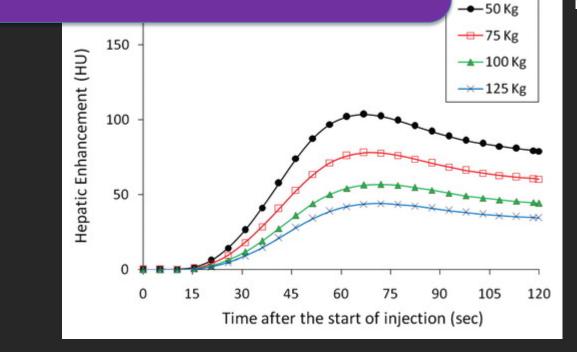


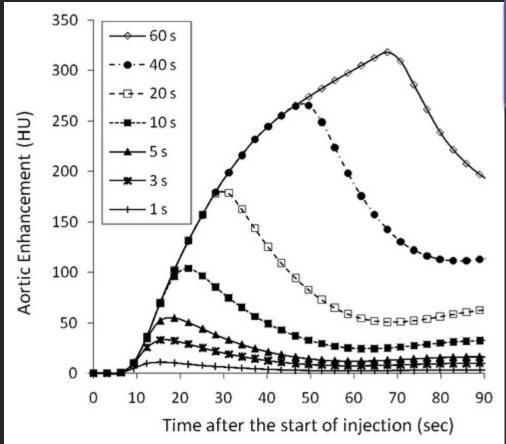
Longer and longer injections push CT enhancement up and delay peak enhancement



Longer and longer injections push CT enhancement up and delay peak enhancement

> Bigger people have more blood... which dilutes contrast agent

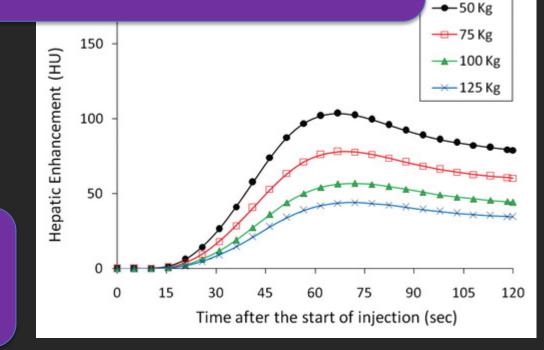




Arterial enhancement is "in and out" faster relative to parenchymal

Longer and longer injections push CT enhancement up and delay peak enhancement

> Bigger people have more blood... which dilutes contrast agent



Equations Governing the Major Facets of Contrast Delivery

 $Volume_{arbitrary strength}(ml) = \frac{Strength_{reference}(mg I / ml)}{Strength_{arbitrary}(mg I / ml)} Volume_{reference strength(ml)} [See Table 8.2]$   $Volume(ml) = Duration(s) \times Injection flow rate(ml / s) [See Table 8.3]$   $Total iodine load (mg I) = Contrast concentration (mg I per ml) \times Contrast volume(ml)$   $Scan delay = Time to optimal enhancement - \frac{1}{2}Scan duration$ 

 $Scan speed (mm/s) = \frac{Collimation(mm) \times Pitch}{Rotation time(s)}$ 

 $Scan \, duration \, (s) = \frac{Scan \, range \, (mm)}{Scan \, speed \, (mm \, / \, s)}$ 

 $= \frac{Scan \, range \, (mm) \times Rotation \, time \, (s)}{Collimation \, (mm) \times Pitch}$ 

Contrast volume as a function of patient weight and contrast strength is shown in Table 8.3 for routine abdominal parenchymal enhancement.

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#### Equations Governing the Major Facets of Contrast Delivery

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Total iodine load (mg I) = Contrast concentration (mg I per ml) × Contrast volume (ml)

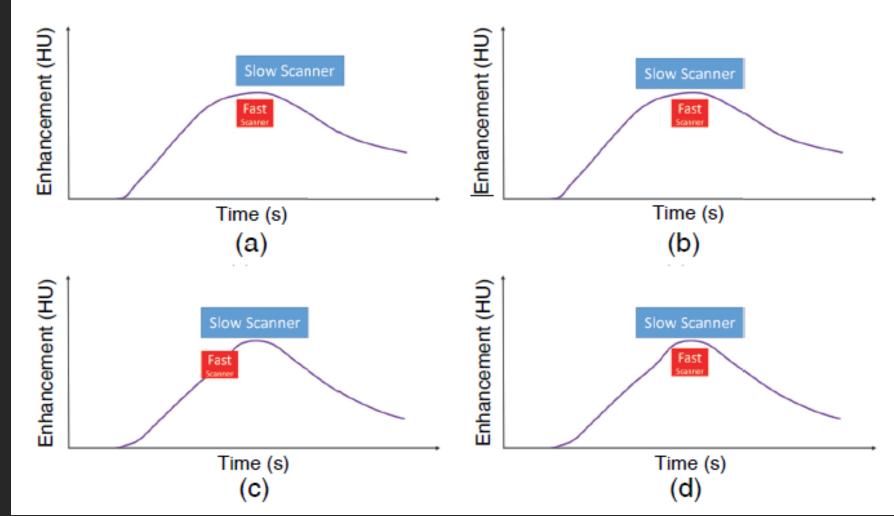
#### Iodine load versus contrast volume

All volumes are not created equal... a lower volume of high concentration agent can deliver the same total lodine as a larger volume of less concentrated agent Want more enhancement? Increasing volume can give more enhancement, but change contrast timing

Increasing concentration will increase enhancement and usually wont change timing

Scan delay = Time to optimal enhancement  $-\frac{1}{2}$  Scan duration

Scan delay = Time to optimal enhancement  $-\frac{1}{2}$  Scan duration



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$$Scan speed (mm/s) = \frac{Collimation (mm) \times Pitch}{Rotation time (s)}$$

$$Scan duration (s) = \frac{Scan range (mm)}{Scan speed (mm / s)}$$

$$= \frac{Scan range (mm) \times Rotation time (s)}{Collimation (mm) \times Pitch}$$

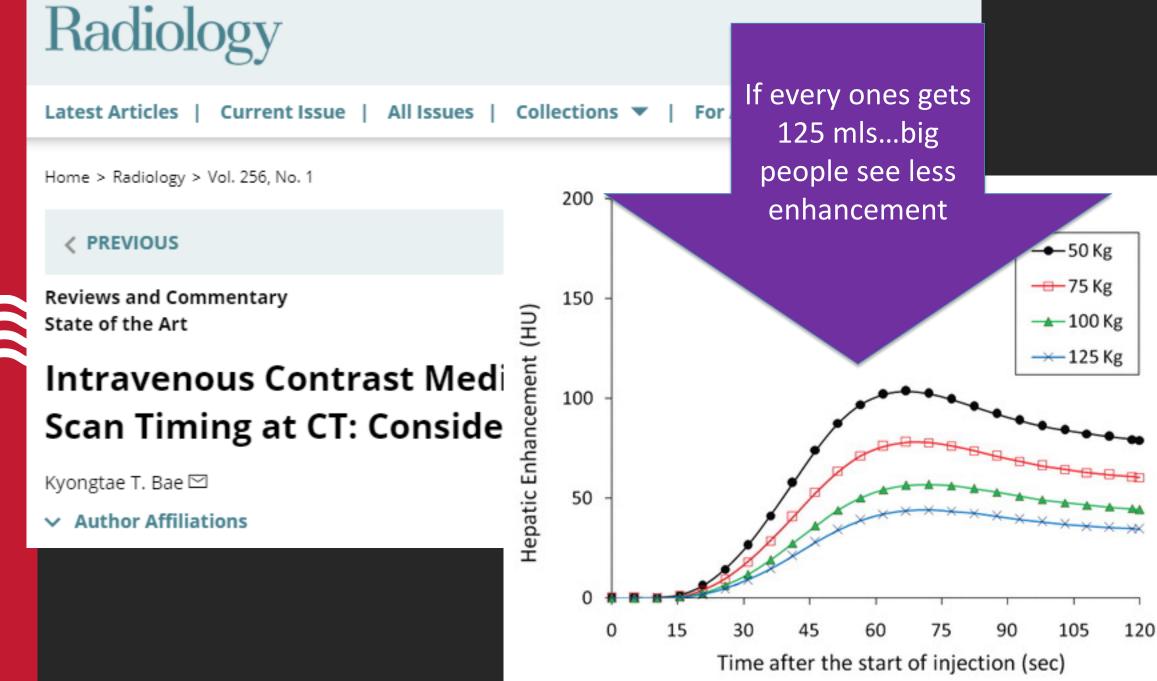


# @Prof\_TimStick's Actionable information

- As a physicist, you should "own" scan duration. You can get scan length from dose report (DLP/CTDIvol).
  - Yes I know about overscanning. It doesn't matter here...
- As a physicist, you can suggest scanner specific scan delays differences based on scan speed.
  - Faster scanners usually need longer delays
- Physicists are good at multiplying and dividing numbers... so do site specific volume calcs for your sites having different strength agents.

# CT Contrast 201...graduating to the clinic ;)





 Typical IV contrast prescription. Most sites around the world will have increases in I contrast with weight.

Example CTPA (PE) contrast prescription

#### IV Contrast Parameters

#### Patient weight < 140 kilos.(Less than 300 lbs.)

- 100 mL lohexol (Omnipaque) 300 MG/ML @ 5 mL/sec
- 10 mL Sodium Chloride 0.9% @ 5 mL/sec

#### Patient weight 140-160 kilos.(300-350 lbs.)

- 100 mL lopamidol (Isovue 370) 370 mgl/ml @ 5 mL/sec
- 10 mL Sodium Chloride 0.9% @ 5 mL/sec

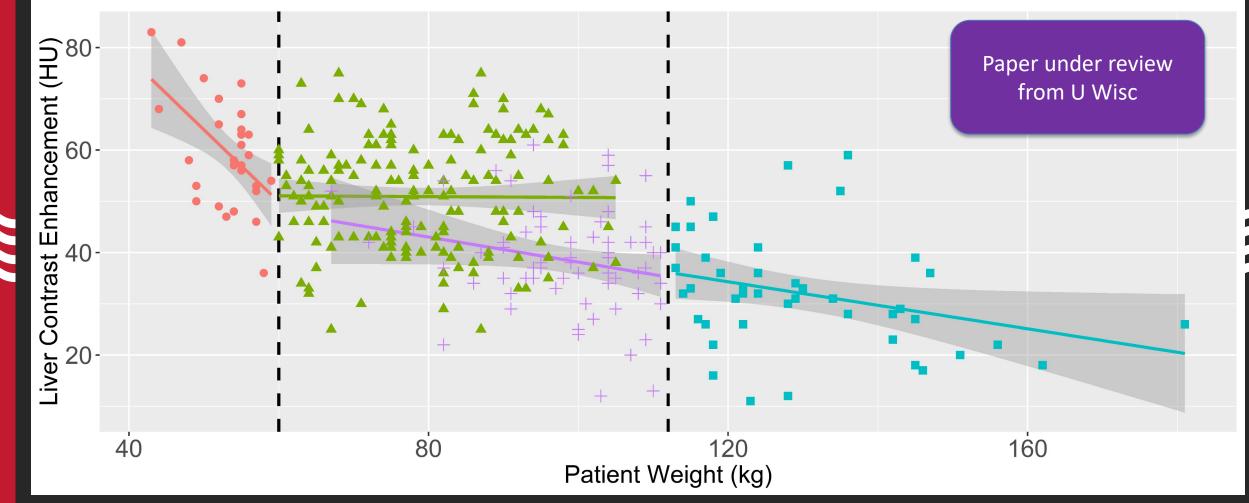
#### Patient weight > 160 kilos.(More than 350 lbs.)

- 150 mL lopamidol (Isovue 370) 370 mgl/ml @ 5 mL/sec
- 10 mL Sodium Chloride 0.9% @ 5 mL/sec

Example routine parenchymal phase torso contrast prescription

Patient Weight (Ibs)	Contrast Volume (ml or cc)		
130 and less	80 (minimum amount to load)		
140	86		
150	92		
160	98		
165	101		
170	104		
175	107		
180	110		
190	116		
200	122		
210	129		
220	135		
230	141		
240	147		
250 and larger	150 (max amount to load)		

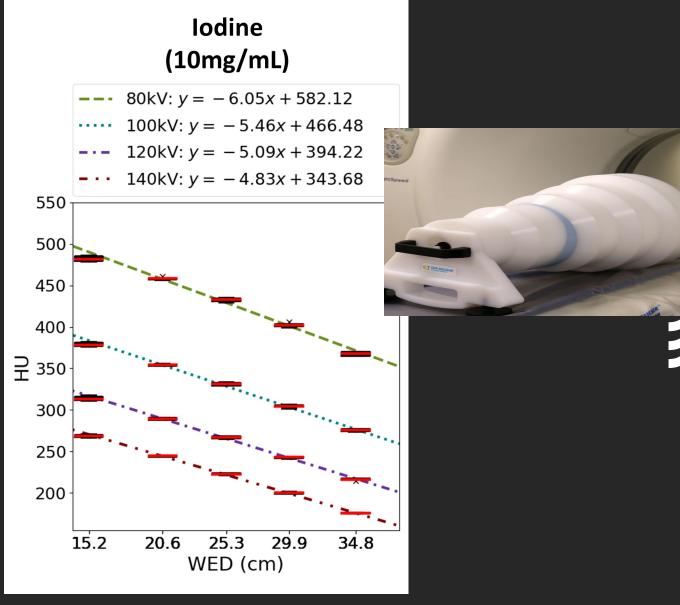
Patient Cohort 🚽 <60kg, 120kV 📥 60-112kg, 120kV 📥 >112kg, 140kV 🔚 60-112kg, 140kV



Smaller people see big increase in enhancement (opposite effect and rationale for large people) Within black lines we use weight based dosing

Large patient see quick fall off in enhancement due to blood volume increase and beam hardening

- Just due to beam
   hardening, we see a
   HUGE reduction in CT
   number with increasing
   patient size
  - -5 HU per cm of WED!

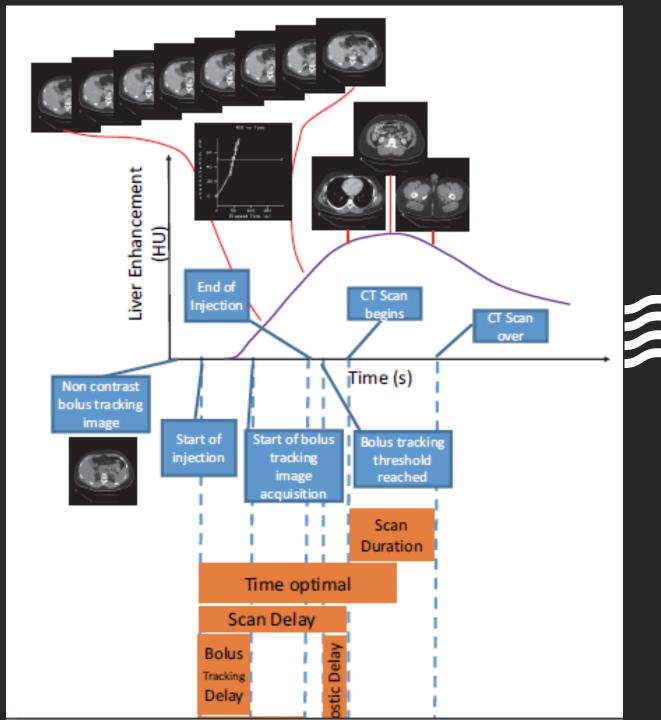


**Parameterizing Size-Based Variations in CT Number** AAPM ePoster Library. Rose S. 07/12/20; 302594; BReP-SNAP-I-36 Topic: Multi-detector CT

# BOLUS TRACKING

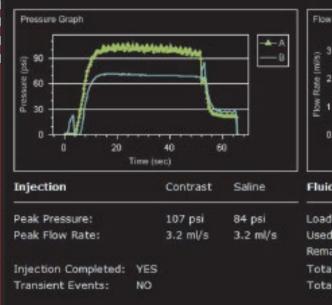
Allows for CT protocol acquisition tuned to patient specific cardiac output and flow dynamics

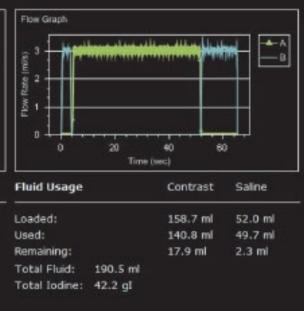
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#### Administered 140.8 ml of 300 mg/ml using P3T Abdomen.

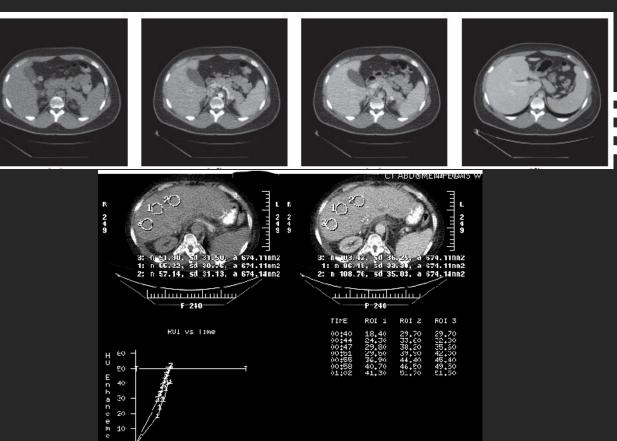
P3T Abdomen							
Programmed		ml/s ml		Actual		ml/s	ml
1	8	3.0	10	1	8	2.8	9.8
2	Hold			2	Hold		
3	Α.	3.0	141	з	A	3.0	140.8
4	в	3.0	40	4	в	3.0	39.9
Tat	tal Contrast (A):	141 ml		To	tal Contrast (A):	140.8 m	E.
Tot	tal Saline (B):	50 ml		To	tal Saline (B):	49.7 ml	
Del	ay:	None					
Pre	ssure Limit:	300 psi					





 $\checkmark$ 

#### Bolus tracking time series



Elepsed Time = 01:04

-10

EO 100 150 Elapsed Time (s) 35

Fil



# @Prof\_TimStick's Actionable information

- If your site isn't using weight-based dosing...they will see large changes in enhancement with weight (and are probably wasting a lot of money)
- If your site isn't using bolus tracking... also badness, not enough time today to get into that. But sick patients really benefit from BT as they usually need longer (tens of seconds) delays w.r.t. healthy people

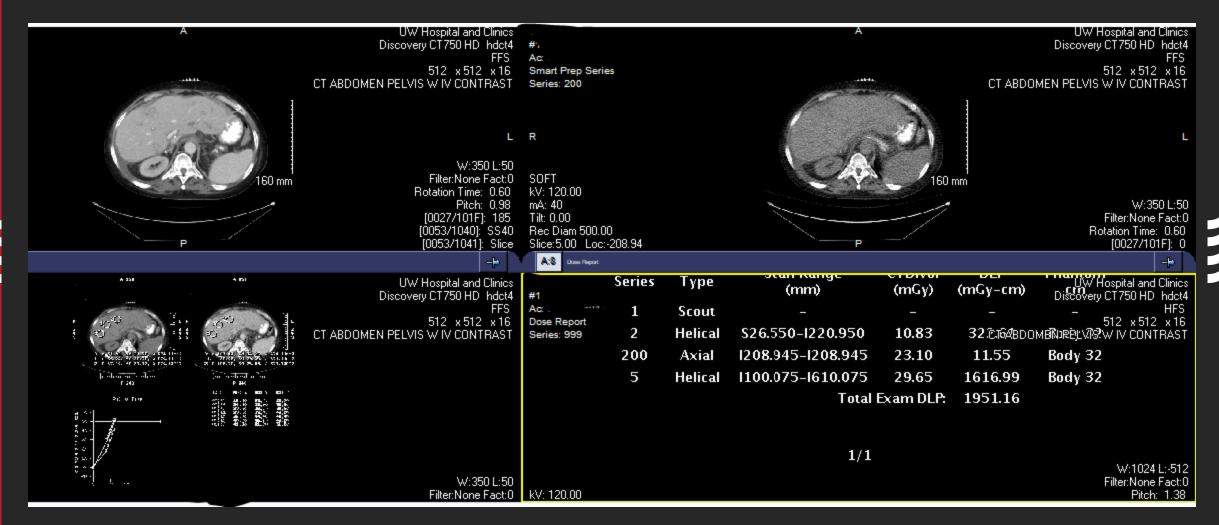


# @Prof\_TimStick's Actionable information

- Homework time!
  - Go to PACS and figure out the scan timing as I have done on the following slides
    - There will be site and vendor specific limitations and nuances you'll need to understand...

#### Parenchymal liver scan

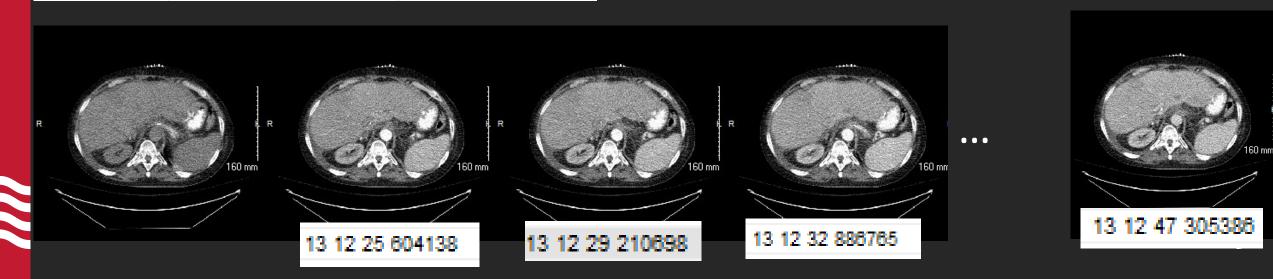
#### Baseline bolus tracking image



Scanner generated bolus tracking report

Dose slide. Series 200 on GE is the bolus tracking phase. (this study had a chest and a AP)

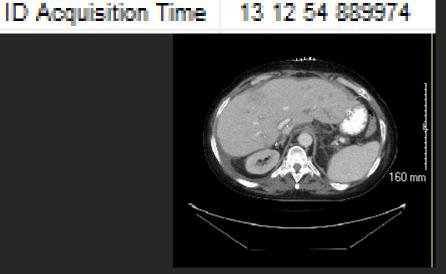
#### 0008 0032 ID Acquisition Time 13 11 26 504627



We have a story. At 1:11:26 the tech took the non contrast 0008 0032 baseline image for ROI placement

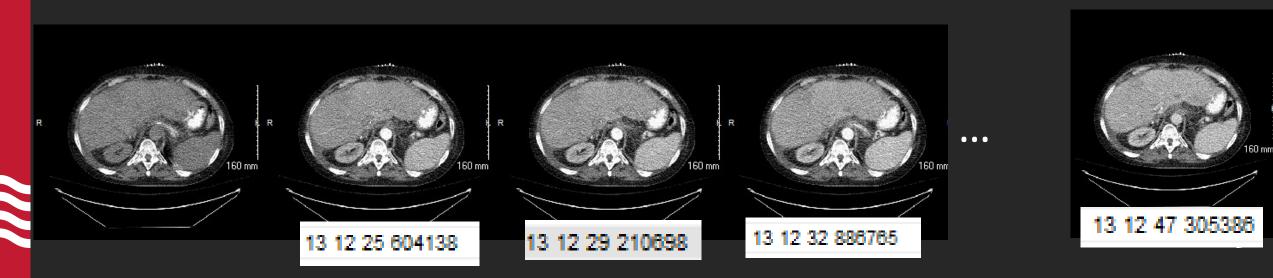
At 1:12:25 they started acquiring CINE images to monitor enhancement, taking an image every ~3-4 seconds... The pt hit enhancement threshold at 1:12:47

It took ~7 seconds to move from bolus tracking position to start of scan (1:12:54-1:12:47)



12 54 889974

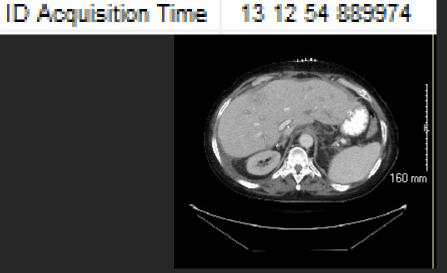
#### 0008 0032 ID Acquisition Time 13 11 26 504627



I know we use a 40 second delay before acquiring the first bolus tracking image, so we started injection at 1:12:25 – 40 seconds.

1:11:45 was injection start

So time of scan from injection start was 1:12:54 – 1:11:45 = 69 seconds





#### Please feel free to reach out to me with questions tszczykutowicz@uwhealth.org

@Prof\_TimStick