

# THE FORMULA SHEET

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## X-ray Production & Interaction

k-shell binding energies (keV)

Mo	Rh	Ag	W	Iodine	Barium
20	23	25	70	33	37

\*k-edge = k-shell binding energy

characteristic energies (keV)

Mo	Rh
18 & just below 20	20 and just below 23

- PE absorption  $\propto Z^3/E^3$
- Compton Scatter  $\propto 1/E$
- Prob(PE) = Prob(Compton) @ 25 keV soft tissue @ 40 keV bone
- HVL = thickness needed for transmitted x-ray beam intensity to be 50% of initial beam intensity ex.) 3 HVL attenuation =  $(1/2) \times (1/2) \times (1/2) = (1/8)$  => 1/8th of incident beam is transmitted. Caution: don't confuse absorbed vs. transmitted.

## Projection Radiography

- 15% ↑ in kVp increases efficiency of x-ray production, requiring only half the initial mA·s to keep the same total number of photons.

Film-Screen	Indirect Digital	Direct Digital	CR
screen + film	CsI scintillator	amorphous selenium	PSP

- assume AEC in effect when in doubt
- geometric magnification = SID/SOD

### 3 Determinants of Resolution

(1) Detector (pitch), (2) Motion (exposure time, compression paddles), (3) Effective Focal Spot Size (filament size, geometric blurring, anode angle)

## Fluoroscopy

- deterministic risk: AK in [Gy]
- stochastic risk: KAP = AK x area in [Gy·cm<sup>2</sup>]
- JC Sentinel Event if >15 Gy fluoro to single field @1m, operator gets 1/1000th of dose to patient safety: time, distance, shielding
- collimation decreases KAP
- magnification increases AK (II > FPD) and improves spatial resolution for II, and improves spatial resolution for FPD only when binning turned off.

## Mammography

- Target/Filter Combinations → → increasing breast density
- Mo/Mo, Mo/Rh, Rh/Rh, W/Ag
- focal spot: 0.1mm mag, 0.3mm reg
- PNL within 1 cm between CC & MLO
- ACR Phantom: 4.2cm, 50% glandular
- (fiber, spec groups, masses)
- Old: (6,5,5), passing (4,3,3)
- New: 🍌 (6,6,6), passing (2,3,2)

- Breast MR: 0.1 mmol/kg gad, perform during days 7-13 of menstrual cycle, first delayed phase image within 2 minutes of contrast injection

## Computed Tomography (CT)

- beam pitch =  $\frac{\text{distance table traveled in one rotation}}{\text{beam thickness}}$
- beam pitch > 1: ↓ dose, worse resolution
- beam pitch < 1: ↑ dose, better resolution
- keep pitch < 2; "pitch" = "beam pitch"
- Cardiac CT: prospective gating target 50-60 bpm. retrospective gating ↑ dose, but allows diast+ syst imaging separately, hence EF can be calculated.
- CNR  $\propto \sqrt{\text{voxel volume}}$
- 16cm head phantoms, 32cm body phantoms; these phantoms underestimate dose to peds, and overestimate dose to obese (fat protects organs)
- CTDIvol =  $(2/3 \text{ periph} + 1/3 \text{ central})/\text{pitch}$ , in [Gy]
- DLP = CTDIvol x scan length, in [Gy·cm], approximates cancer risk using conversion factor

## Ultrasound

- $c_{\text{soft tissue}} = 1540 \text{ m/s}$ ;  $c_{\text{bone}} > c_{\text{soft tissue}} > c_{\text{fat}}$
- $c = \text{TF} \times \lambda$ , where TF = transducer freq. (in Hz),  $\lambda$  wavelength (in meters)
- axial res = SPL/2, where SPL ≈ 2λ
- "axial res" here is smallest distance resolvable between two features. "Better" is smaller number.
- FOV<sub>axial</sub> =  $c \times \text{PRP} / 2$
- PRF = 1/PRP
- Don't confuse TF with PRP. ↑TF → better ax/lat/elev res + worse attenuation. ↑PRP → deeper axial FOV and slower screen refresh rate.
- lat/elev res best in focal zone, also ↑scan line density → better lateral res
- attenuation of sound in soft tissue (dB) = 0.5 dB/cm/MHz, where MHz is TF, cm is total distance traveled. Total distance (roundtrip) traveled by an ultrasound pulse is 4 cm for lesion of depth 2 cm.
- depth-resolution tradeoff: use the highest frequency that penetrates deep enough for you to see the structure you're interested in. ↑TF → better ax/lat/elev resolution but greater attenuation per cm of tissue depth
- Thermal Index (TI) - heating
- Mech. Index (MI) - cavitation risk

## Nuclear Medicine

- Activity in Becquerel (Bq) is decays per second
- 1 mCi = 37 MBq
- $1/(\text{effective } t_{1/2}) = 1/(\text{physical } t_{1/2}) + 1/(\text{biological } t_{1/2})$

Radioisotope Physical Half-Lives					
Tc99m	I-123	I-131	Ind, Thal, Ga	Xe-133	F-18
6 hrs	13 hr	8 d	3 d	5 d	110 min

Gamma Energies (keV)					
Tc99m	I-123	I-131	Ga-67	Xe-133	F-18
140	159	365	~100, 200, 300, 400	81	511

- Activity Exponential Decay:  $A(t) = A_0 \exp(-\lambda t)$
- λ is decay constant, A<sub>0</sub> is initial activity at t=0
- Half-life (t<sub>1/2</sub>) can be calculated using  $t_{1/2} = 0.693/\lambda$
- Activity remaining at N half-lives is  $A_0(1/2)^N$
- Milk the Mo-Tc generator q24 hrs x 5 days
- Mo breakthrough - 700 keV photons
- Aluminum breakthrough - color strip

### Gamma Camera QC

- daily: extrinsic uniformity, photopeak
- weekly: intrinsic uniformity, bar phantom (linearity, spatial resolution)
- monthly: center of rotation (COR)
- quarterly: Jaszczak phantom

### Dose Calibrator QC

- Constancy (daily)
- Linearity (quarterly)
- Accuracy (annually)
- Geometry (on repair)

## MRI

- Larmor Freq = (42 MHz/T) x B
- Longitudinal (T1) relaxation - spin-lattice
- Transverse (T2) relaxation - spin-spin

Weighting	TR	TE
T1	short	short
T2	long	long
Proton Density	long	short

- spin-echo TR: short (<500ms), long (>1500ms)
- spin-echo TE: short (<30ms), long (>150ms)

Scan Time =  $(\text{NEX}) \times (\# \text{ slices}) \times \text{TR} \times \text{N}_y$   
turbo factor

- NEX = number of repetitions (a.k.a. NSA)
- N<sub>y</sub> = # phase encoding steps
- turbo factor = 1 for regular spin-echo

$\text{SNR} \propto \text{voxel volume} \times \sqrt{\frac{\# \text{ measurements}}{\text{receiver bandwidth}}}$

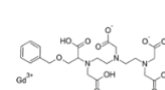
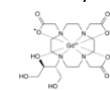
- N<sub>x</sub> : # freq encoding steps, N<sub>y</sub> : # phase encoding steps
- # measurements = N<sub>x</sub> x N<sub>y</sub> x NEX

$\text{voxel volume} = \left( \frac{\text{FOV}_x}{N_x} \cdot \frac{\text{FOV}_y}{N_y} \cdot \Delta \right)$

Δ : z-axis slice thickness

- ↑ voxel volume → worse resolution, ↑SNR
- ↑ N<sub>y</sub> → ↑ pixels in phase encoding axis, ↑ scan time
- ↑ receiver bandwidth → ↓ chem shift Type 1, ↓SNR, ↓ scan time (from shorter echo)
- Change TE to remove chemical shift Type 2
- FDA limit whole-body SAR: 4 W/kg for 15 min

Macrocytic vs Linear



- macrocytic better for chelation, decreased NSF risk
- NSF risk by ACR Group: 2 (low, GFR screening optional), 1 (high, avoid for GFR<30, dialysis, AKI), 3 (insufficient info – treat like 2).

## Radiation Dose Units & Nucs Safety

See [Radiology Simplified](#)