

# **MR Brain Protocols**



Making Cancer History<sup>®</sup>

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### **Routine Brain Protocol**

#### **ACR Requirement:**

Sag T1 (or 3DT1) AX T2 AX FLAIR AX DWI

+C if breakdown of BBB AXT1+C

Slice <= 5mm, gap <= 2.5mm

- Sagittal T1W
- Axial DWI
- Axial T2W
- Axial FLAIR
- Axial T1W
- Post contrast Axial T1W
- Post contrast 3DT1W

### **Protocol Consideration**

• Signal to Noise Ratio (SNR)

$$SNR \propto x \Delta y \Delta z \frac{\sqrt{N_x N_y NEX}}{\sqrt{BW}} B_0 f$$

- Contrast to Noise Ratio (CNR)
- Resolution
- Scan time

## **Tissue Characters**

#### •*Tissue Characters at 1.5T*\*

Tissue	ρ	T1 (msec)	T2 (msec)
CSF	1.0	4500	2200
WM	0.65	600	80
GM	0.8	950	100
FAT	0.9	250	60

Magnetic Resonance Imaging, Haacke, Brown, Thompson and Venkatesan, p. 457



## Pulse Sequences

• Spin Echo Sequence (SE)

$$S_{SE} \propto 
ho_{1_{H}} e^{-TE/T_2} \left(1 - e^{-TR/T_1}\right)$$

- Gradient Echo Sequence (GRE)  $M_{xy}(t) = M_0 \frac{(1 - e^{-TR/T_1})e^{-TE/T_2^*} \sin \alpha}{(1 - e^{-TR/T_1} \cos \alpha)}$
- FSE,SSFSE belongs to SE family
- SPGR, MPRAGE belongs to GRE family
- Advanced sequences: EPI, Propeller,

#### Spiral, ASL

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http://199.116.233.101/index.php/Pulse\_Sequences





### Let TE ==> 0 and TR << $\infty$ , then S ~ S<sub>0</sub>(1 - e<sup>-TR/T1</sup>)

T1 Weighted images, TR ~500msec. Short T1 substances will be bright (FAT), long T1 substances will be hypointense (CSF).

 $\bullet$ 



#### • Let TE > 0 and $TR ==> \infty$ , then

 $S \sim S_0 e^{-TE/T2}$ 

T2 Weighted images, TE ~100msec. Short T2 substances will be dark, long T2 substances will be hyperintense (CSF).



• Let TE==>0 and TR ==>  $\infty$ , then S  $\sim$  S<sub>0</sub>

Proton density weighted image – not popular!

### T1-Weighted image protocol





SE, AX or SAG TR = 500-700 msec TE = min (10 – 20 msec) FOV = 220 – 240 mm, no FC Matrix = 256x192 (1.5T), (352x224) (3T) slice thickness = 5.0 mm, gap = 1.0mm Phase encoding dir = RL BW = 20.83 (31.25, 3T) kHz, 1NEX 180 mm SI coverage Scan time ~ 2- 4 minutes

Gray matter – dark gray White matter – lighter gray CSF – black Fat - white

# T2-Weighted image



FSE, AX TR > 3500 msec TE = 80 - 140 msec, etl >= 16 FOV = 220 - 240 mm Matrix = 288x224 (1.5T), (352x256) (3T) slice thickness = 5.0 mm, gap = 1.0mm Phase encoding dir = RL, BW = 20.83 (41.67, 3T) kHz, 2NEX Scan time ~ 2- 4 minutes

Gray matter – light gray White matter – darker gray CSF – bright Fat – less bright than CSF (T2 increase due to elimination of J-coupling between adjacent fat protons).

Andrew Murphy, <u>https://radiopaedia.org/articles/spin-echo-sequences?lang=us</u> Henkelman RM, et al, J Magn Reson Imaging 1992; 2:533-540

# **FLAIR** Image



 $TI = T1_{csf} * \ln \frac{2}{(1 + e^{-TR/T_1})}$ T1<sub>csf</sub> = 4500 msec, same for 1.5T and 3T



- FLAIR FLuid Attenuated Inversion Recovery.
- IR + FSE
- IR flip Mz to –Mz
- Long TI null the CSF
- Long TR Mz recovery
- Long TE T2W image.

TR = 9000 – 12000 msec (TR >> T1csf) TI = 2250 – 2500 msec (auto) TE = 80 – 140 msec (T2W) FOV = 220 – 240 mm Matrix = 288x192 (1.5T), (288x224) (3T) slice thickness = 5.0 mm, gap = 1.0mm Phase encoding dir = RL, etl >= 16 BW = 20.83 (31.25, 3T) kHz, 1NEX Concatenation >= 2 (interleave) Scan time ~ 2- 4 minutes FLAIR image is almost in all Brain protocols.



Concats = 1





GFR < 30, contrast < 24 hrs



# **FLAIR Artifacts**

#### CSF not suppressed

- in-flow from neighboring slices
- TR/TI sensitivity, not correct TI
- Recent GBCA with low GFR issue
- Subtle reduced T1 due to high tissue O2 concentration during general anesthesia
- Motion sensitive to motion due to long TR, TI and TE
  - Traditional uncontrollable motion
  - Occasional motion (movement)



Propeller MRI

Multiple blades to oversample the center of k-space to reduce motion.

FLAIR - motion

#### Propeller FLAIR

# T2\*-Weighted image

- GRE Sensitive to tissue susceptibility
- GRE signal decay with T2\*
- TR is long (>700msec)
- TE is long for GRE (15 25 msec)
- FA is small (20)
- 3D SWI better resolution, minIP

#### T2\*, GRE

Long TR (> 700msec, 1.5T, >600 3T) Long TE (25 msec, 1.5T, 15msec 3T) Small FA (20) FOV = 220 – 240 mm Matrix = 256x192 slice thickness = 5.0 mm, gap = 1.0mm Phase encoding dir = RL, BW = 15.63 (31.25, 3T) kHz, 1NEX

SWI – 3D FGRE TR = 25 – 50msec TE = 20 -40msec FA = 15 - 20 FOV = 220 – 240 mm Matrix = 320x256 Slice thickness <2.0mm, mIP Mag, Phase image

T2\*W

minIP

# Diffusion-Weighted Image (DWI)



Single shot SE-EPI (most popular) TR > 5000 msec TE = min (~70) FOV = 220 – 240 mm Matrix =128x128 slice thickness = 3.0-5.0 mm, gap = 0 Phase encoding dir = AP BW = 250 (Max) kHz, shortest esp b-value = 0, >= 1000, 3 directions, 1NEX Scan time < 1.0 min It is essential for Brain study, acquired first

$$b = (\gamma G \delta)^2 \left( \Delta - \frac{\delta}{3} \right) \quad S/S_0 = e^{-bADC}$$
$$ADC = \frac{\lambda_1 + \lambda_2 + \lambda_3}{3}$$



Alexander AL, Neurotherapeutics. 2007

## Post contrast T1W





#### Paramagnetic Contrast Agents

- Gadolinium (Gd) is most common in MR imaging.
- Gd is toxic must be tightly chelated
- Affect both T<sub>1</sub> and T<sub>2</sub> relaxation time, with the dominant effect being shortening of the T<sub>1</sub> relaxation time (at routine clinical dose).
- Gd is very bright on T1W images. It is very useful in vascular images (MRA) and breakdown of blood-brain barrier.

http://mriquestions.com/does-gd-affect-t2.html

# T1+C with Fat Sat

- Fat has short T1 bright on T1W
- Confused with contrast uptake
- Fat sat increases diagnostic accuracy and improves lesion delineation
- Fat suppression increases scan time, also depends on local field homogeneity





T1W + C

T1W +C, FS

### Post Contrast T1W Artifacts

• Most post contrast T1W image artifacts are non-uniform fat suppression, flow artifacts in the posterior fossa, motion.



Fat Sat fail

No Flow comp

Flow Comp

# Post Contrast 3D T1W imaging



Maximum signal at Ernst angle  $\alpha_{F} = \cos^{-1} (e^{-TR/T1})$ 

Markl M, Leupold J, JMRI, 35:1274-1289 (2012).

 Fast GRE SNR is a function of flip angle (FA), TR, TE

- Gradient spoiled, T2/T1-contrast
- RF-spoiled, T1 contrast, low SNR/CNR clinical (FSPGR, FLASH).
- Balanced SSFP, truly RF and gradient balanced, T2/T1 contrast, (FIESTA, true FISP).

3D FSPGR

 Optimum T1W contrast happen at flip angle large than Ernst angle. For FSPGR, FA = 20 shown to have best tissue contrast.

AX, 3D FSPGR, FA = 20 TR = min (<10msec),TE = min (close to in-phase) slice thickness = 1.4mm, gap = -0.7mm FOV = 240 mm, matrix =256x256 Phase encoding dir = RL, 1NEX BW = 20.83 (41.67 3T) kHz, scan time ~ 4min Reformat to 1mm slice on AX, SAG, COR



- IR based BRAVO/MPRAGE has excellent GM and WM contrast
- Use with caution for post contrast
- It could miss small metastases



AX SE T1+C, FS, 6:32 COR SE T1+C, 3:16





SAG SE T1+C, 2:15

AX SE, flow artifact



AX 3D T1+C, 3:44

Cor reformat

Sag reformat

**3D FSPGR** 

### **Other Options**

- GRE T1W –uncooperative patient
- DTI fiber tracking
- ASL Perfusion
- T2\*W/SWI calcification, hemorrhages
- DIXON option for fatsat
- FLAIR+C LMD

### Advanced Techniques

- MR Spectroscopy (MRS)
- Dynamic Contrast Enhanced imaging (DCE)
- Dynamic Susceptibility Contrast (DSC)
   Imaging
- fMRI (EPI) BOLD Imaging

# Summary

	TR (msec)	TE (msec)	TI (msec)
T1W	Short (500)	Short (Min, 10)	N/A
T2W	Long (>3500)	Long (>80)	N/A
FLAIR	Very Long (>8000)	Very Long (>100)	Very Long (>2200)
DWI	Long (>5000)	Min (~ 70)	N/A
T1W + C, FS	Short (500)	Short (10)	N/A