

MR-guided Laser Ablation in Oncology

R. JASON STAFFORD, PH.D.

DEPARTMENT OF IMAGING PHYSICS

THE UNIVERSITY OF TEXAS

MDAnderson ~~Cancer~~ Center



Introduction

- MR-guided laser ablation is seeing increased use in brain, spine, prostate, liver and other organs.
- MR-guidance plays a critical role in assuring a safe and effective treatment, particularly in its role in providing temperature imaging feedback.
- Here we overview the MR-guided laser ablation process to identify potential pitfalls physicists should be aware of which could negatively impact the safety and efficacy of treatment delivery.



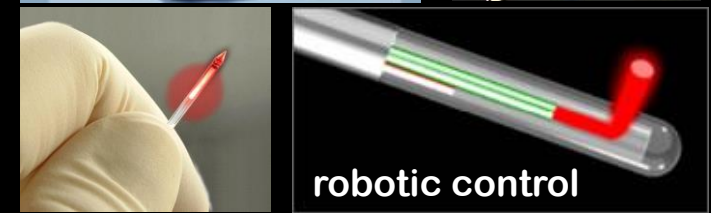
Thermal Ablation Modalities: Laser

- Cryoablation
- Radiofrequency
- Microwave
- Ultrasound
- **Laser**

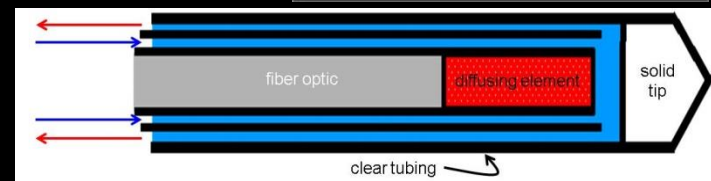
Compact CW or pulsed diode lasers



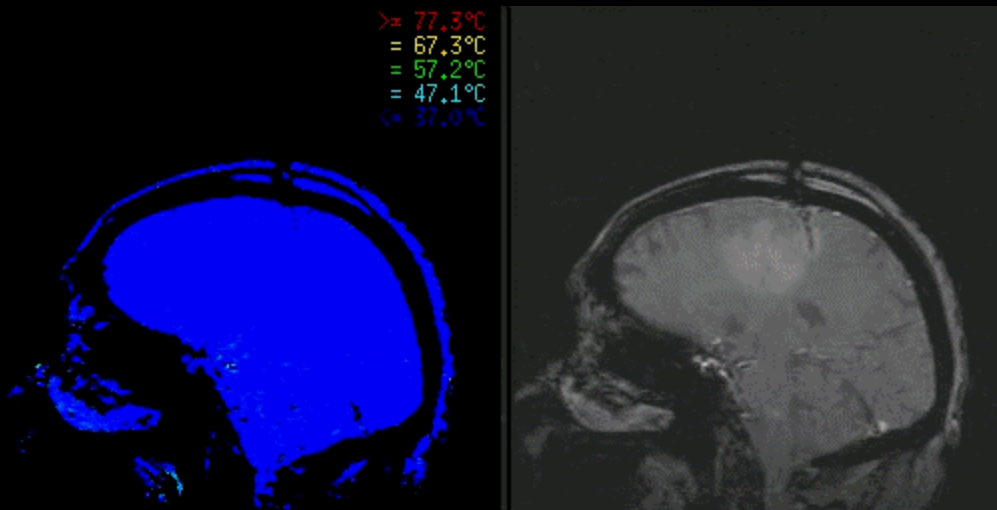
Diffusing or side fire fiber optics



Actively cooled catheters



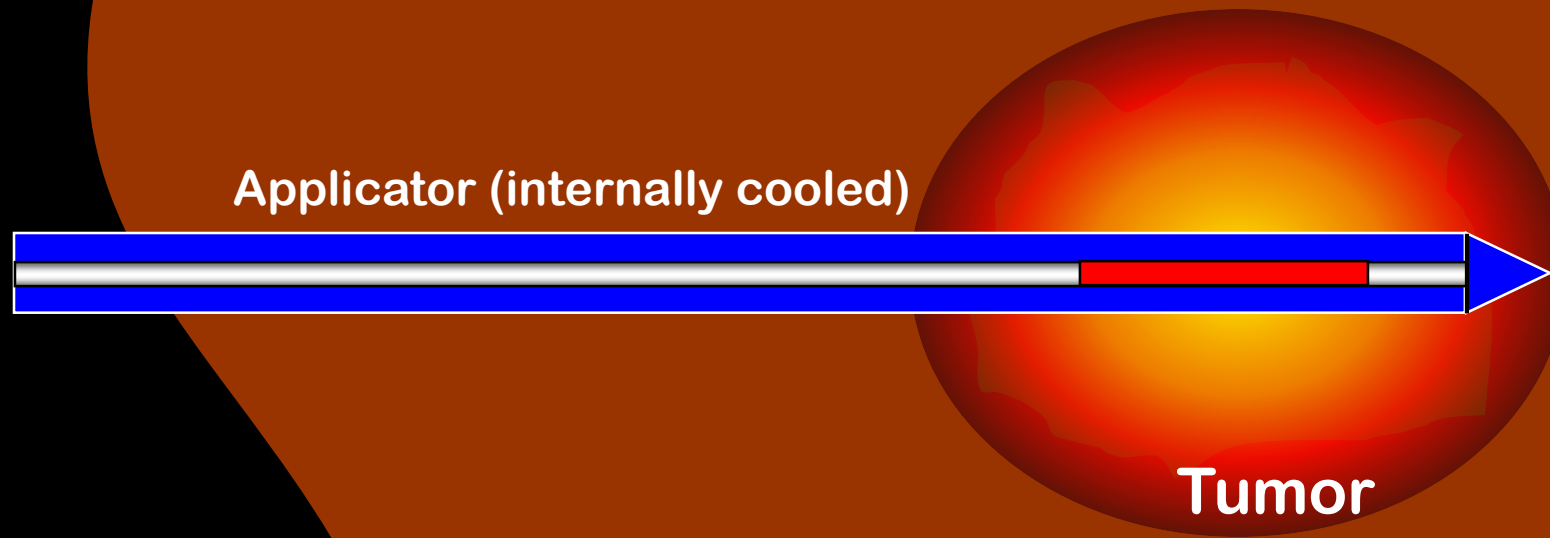
1.6mm (16 gauge)
saline-cooled catheter



Stafford RJ, et al, Crit Rev in Biomed Eng, 2010.



Overview of laser ablation



Tissue Parenchyma



Overview of laser ablation

$$\rho c \frac{\partial T(q_i, t)}{\partial t} = \nabla \cdot [(k \nabla T(q_i, t))] + \rho_b C_b V (\kappa - 1) (T - T_a) + P(q_i, t)$$

heat diffusion
heat convection
heat source

HEAT SOURCES (laser, ultrasound, RF, microwave, etc) :
 $P = \text{absorbed power density (W m}^{-3}\text{)} = \text{SAR} \cdot \rho$

SAR = Specific Absorption Rate (W/kg)

$g = \text{anisotropy factor}$
 $\mu_s = \text{optical scattering}$
 $\mu_a = \text{optical absorption}$

HEAT CONDUCTION (diffusion):

$T = \text{temperature (Kelvin)}$
 $c = \text{specific heat of material (J kg}^{-1}\text{ }^\circ\text{C}^{-1}\text{)}$

$\rho = \text{density (kg/m}^3\text{)}$
 $k = \text{thermal conductivity of tissue (W m}^{-1}\text{ }^\circ\text{C}^{-1}\text{)}$

HEAT CONVECTION (effects of perfusion):

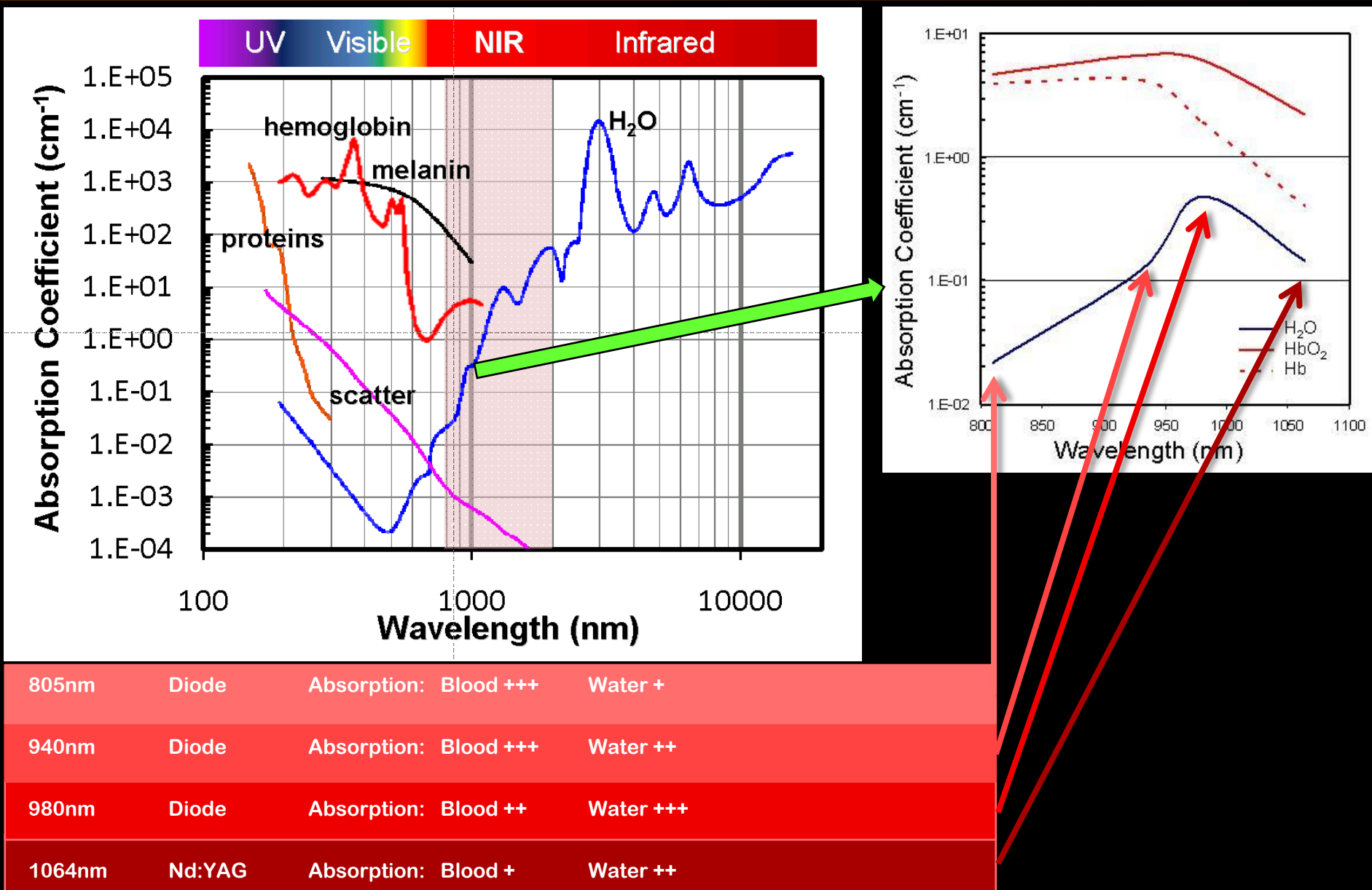
$\rho_b = \text{blood density (kg m}^{-3}\text{)}$
 $V = \text{volume flow rate per unit volume (s}^{-1}\text{)}$

$C_b = \text{specific heat of blood (J kg}^{-1}\text{ }^\circ\text{C}^{-1}\text{)}$
 $\kappa = \text{dimensionless convection scale factor}$

Tissue thermal conductivity largely governs conductive heat transfer
Tissue perfusion largely governs convective heat transfer in tissue



Overview of laser ablation



Overview of laser ablation

$$\rho c \frac{\partial T(q_i, t)}{\partial t} = \nabla \cdot [(k \nabla T(q_i, t))] + \rho_b C_b V (\kappa - 1) (T - T_a) + P(q_i, t)$$

- Laser ($\lambda \sim 800\text{--}1064 \text{ nm}$)
- Photon absorption
- SAR $\sim \mu_a \cdot \phi$

Applicator (internally cooled)

penetration $\sim \text{mm}$



Tumor

Higher absorption
Higher scattering

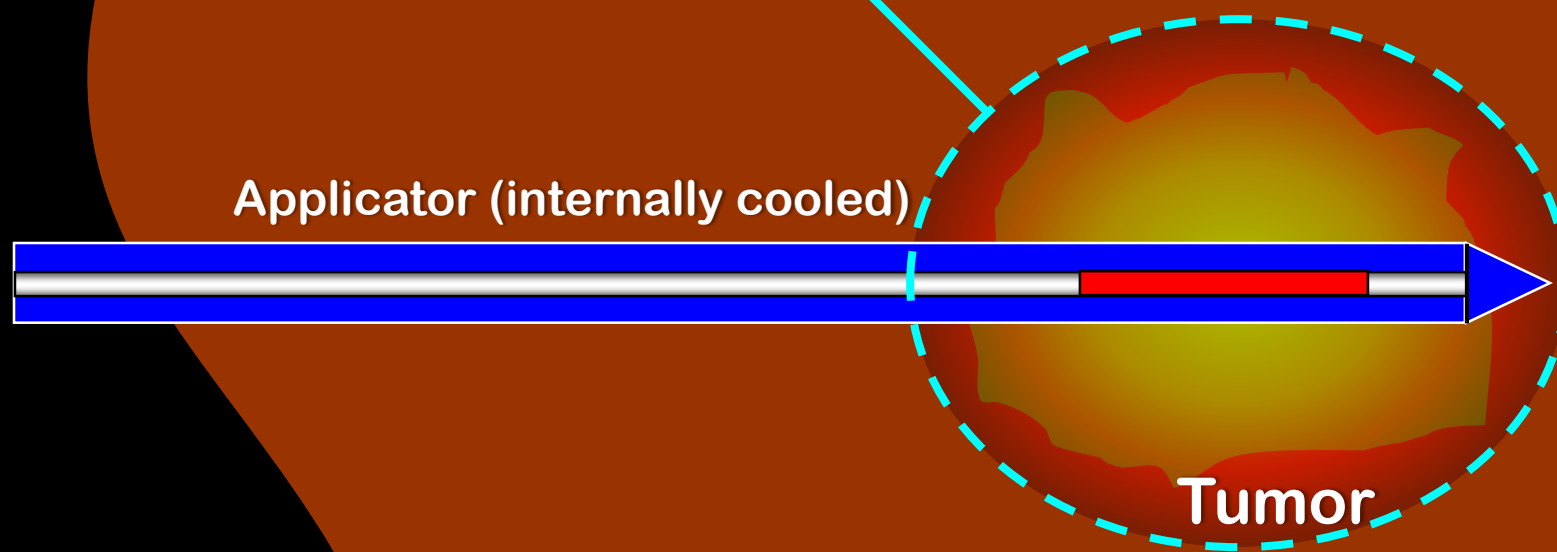
- more heating
- less penetration
- more local absorption
- more heating

Tissue Parenchyma



Overview of laser ablation

$$\rho c \frac{\partial T(q_i, t)}{\partial t} = \nabla \cdot [(k \nabla T(q_i, t))] + \rho_b C_b V (\kappa - 1) (T - T_a) + P(q_i, t)$$



Heat conduction: Lesion boundary based on extent of thermal diffusion beyond area of direct heating vs thermal convection.

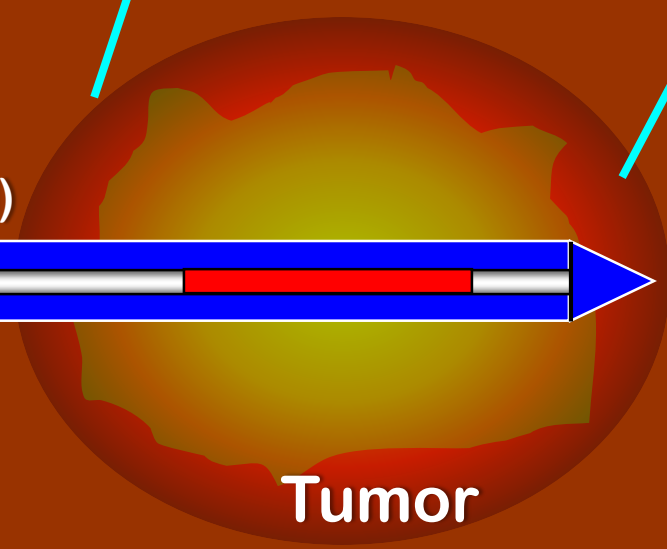
Tissue Parenchyma



Overview of laser ablation

$$\rho c \frac{\partial T(q_i, t)}{\partial t} = \nabla \cdot [(k \nabla T(q_i, t))] + \rho_b C_b V (\kappa - 1) (T - T_a) + P(q_i, t)$$

Applicator (internally cooled)



Tumor

Heat convection: Heated blood in perfused tissue is replaced by arterial blood at body temperature.

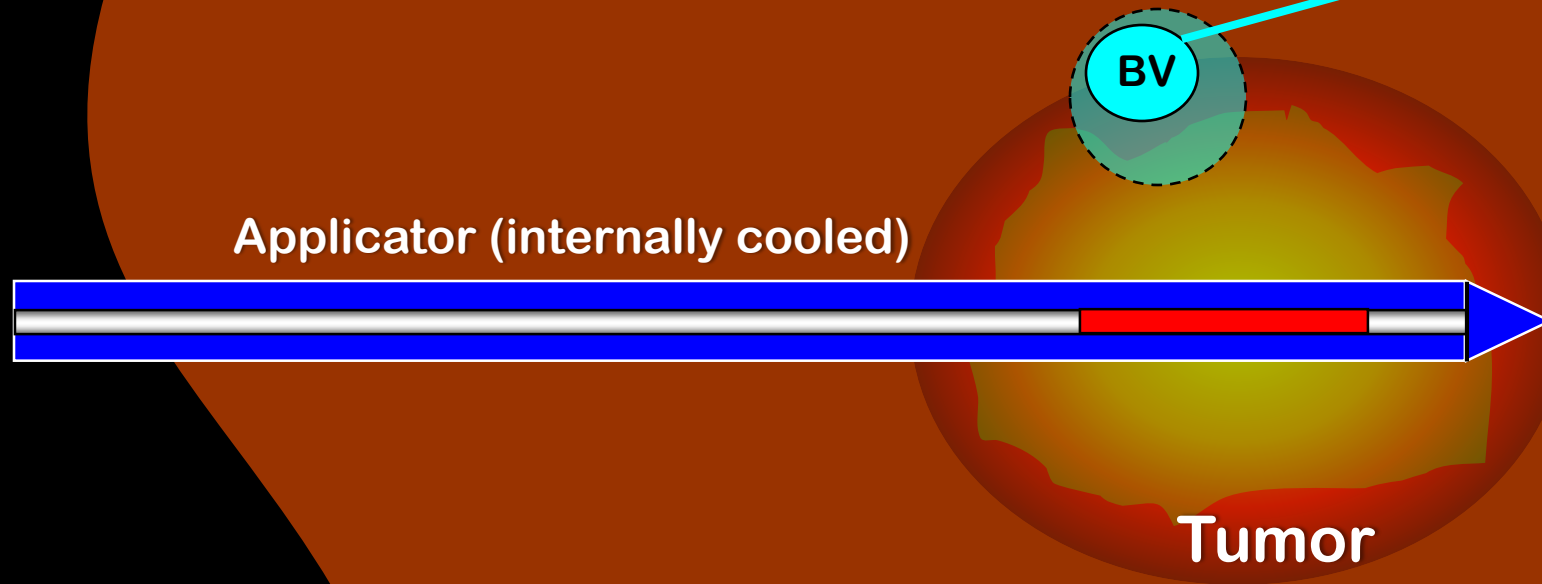
Perfusion in lesion destroyed by ablation.

Large lesions: heat deposited over larger OAR region



Overview of laser ablation

$$\rho c \frac{\partial T(q_i, t)}{\partial t} = \nabla \cdot [(k \nabla T(q_i, t))] + \rho_b C_b V (\kappa - 1) (T - T_a) + P(q_i, t)$$



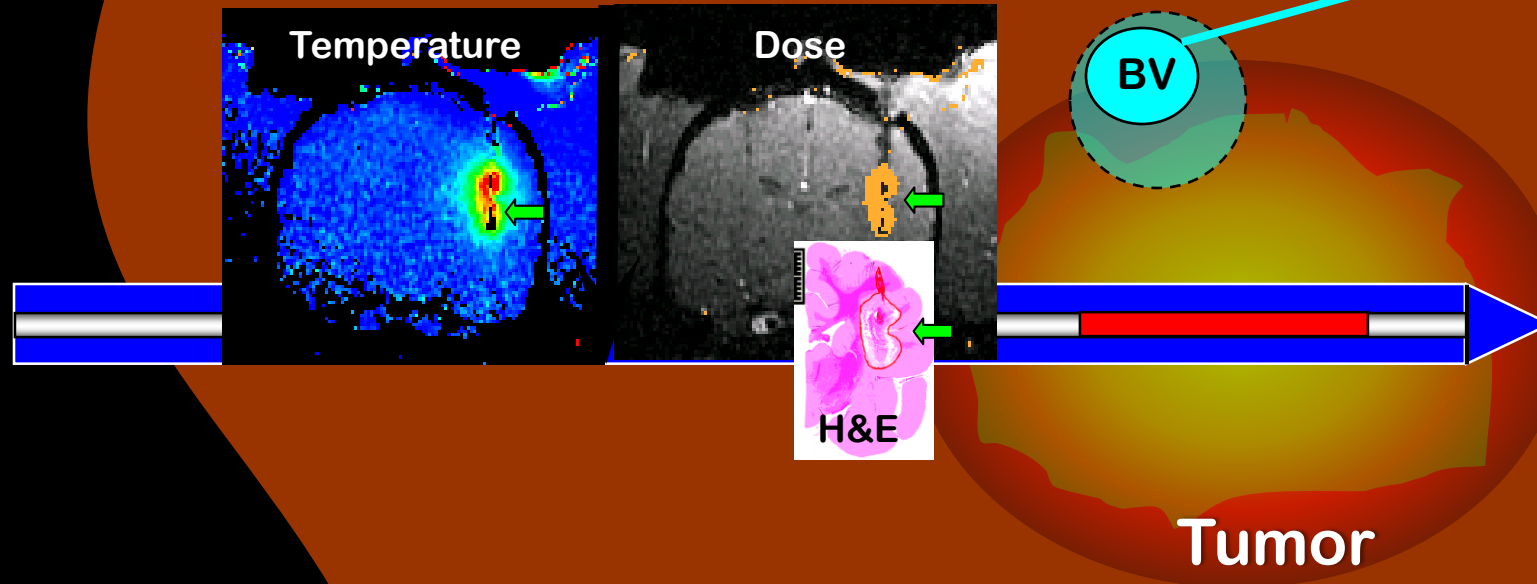
Heat convection: Large vessels or areas of ventilation can rapidly disperse heat resulting in a 'heat sink' effect.

This can result in cold spots.



Overview of laser ablation

$$\rho c \frac{\partial T(q_i, t)}{\partial t} = \nabla \cdot [(k \nabla T(q_i, t))] + \rho_b C_b V (\kappa - 1) (T - T_a) + P(q_i, t)$$



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Overview of laser ablation

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Modality interactions with tissue (redux):

Applicator cooling is a powerful method for counteracting adverse propagation and absorption issues that facilitates increased efficiency in energy delivery.

Applicator **[internally cooled]**



Tumor

Cooling functions as a much stronger BV 'heat sink'.

Begin just prior starting ablation

Set MRTI reference prior to turning on

Should be kept on post ablation to dissipate heat
(minimize risk to surrounding OAR)

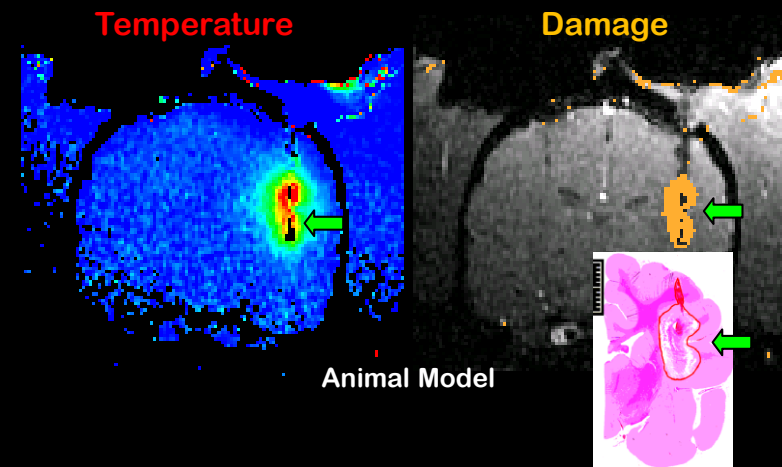
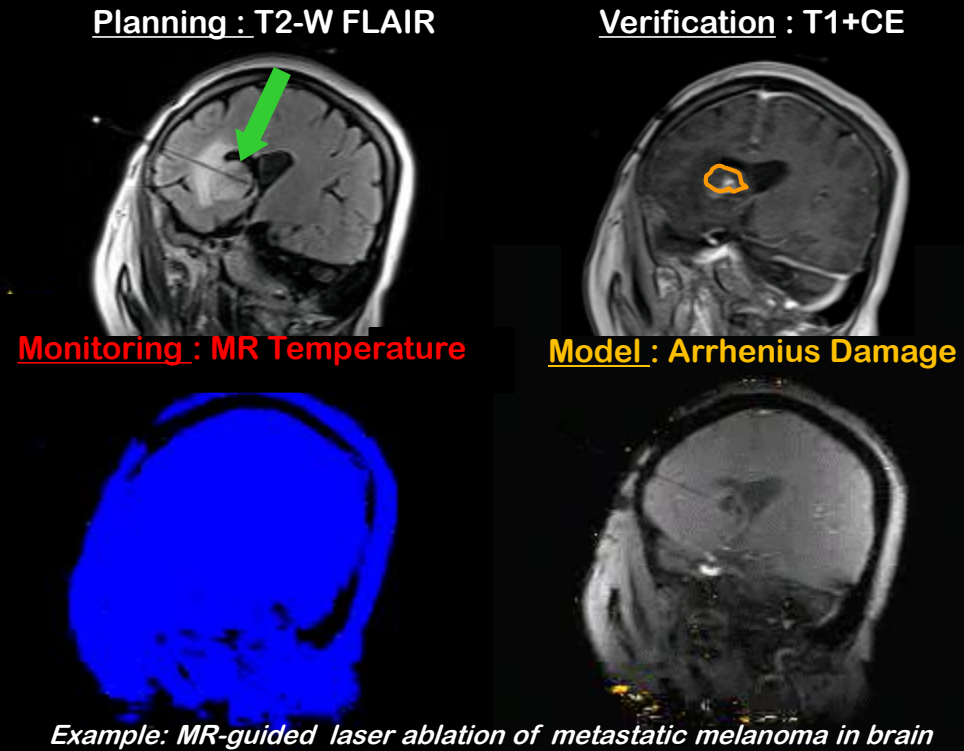
Careful not to cool tissue below baseline if setting a new reference

Tissue Parenchyma



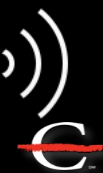
MR-guidance in thermal ablation

- MRI-guidance useful for
 - planning
 - navigation & targeting
 - **monitoring & control**
 - verification
- Synergy with **biological and physical modeling & simulation**
- Endgame
 - ‘close the loop’
 - increase safety + efficacy
 - **facilitate minimally invasive approaches previously not considered safe or effective**



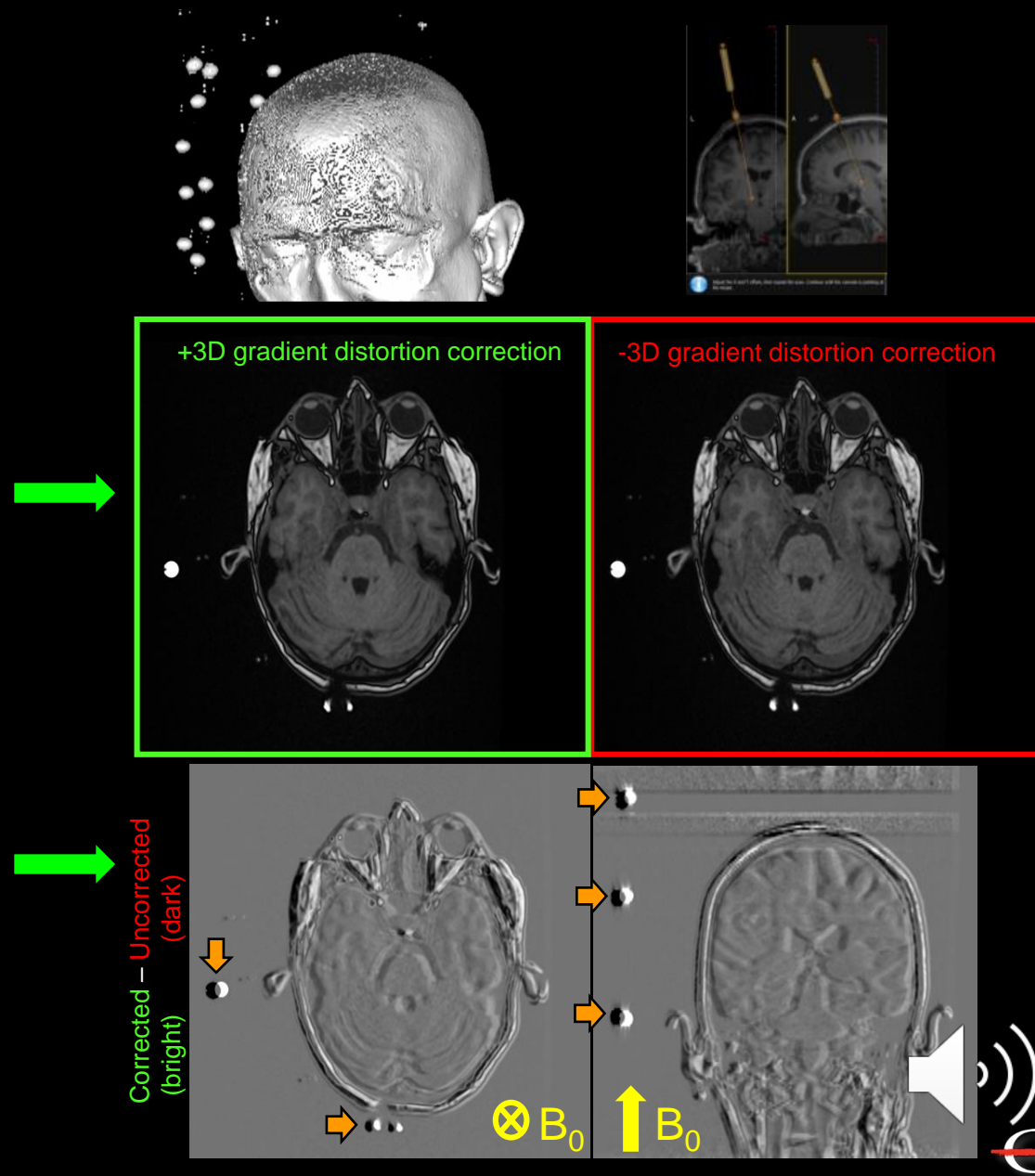
MR-guidance in thermal ablation

- Potential to use MRI for
 - planning
 - navigation & targeting
 - monitoring & control
 - verification
- Lesion & OAR visualization and delineation critical
- Targeting accuracy
 - Geometric distortion
 - MR-guided navigation systems
- MR protocols: CNR & resolution
- Must verify device placement vs plan

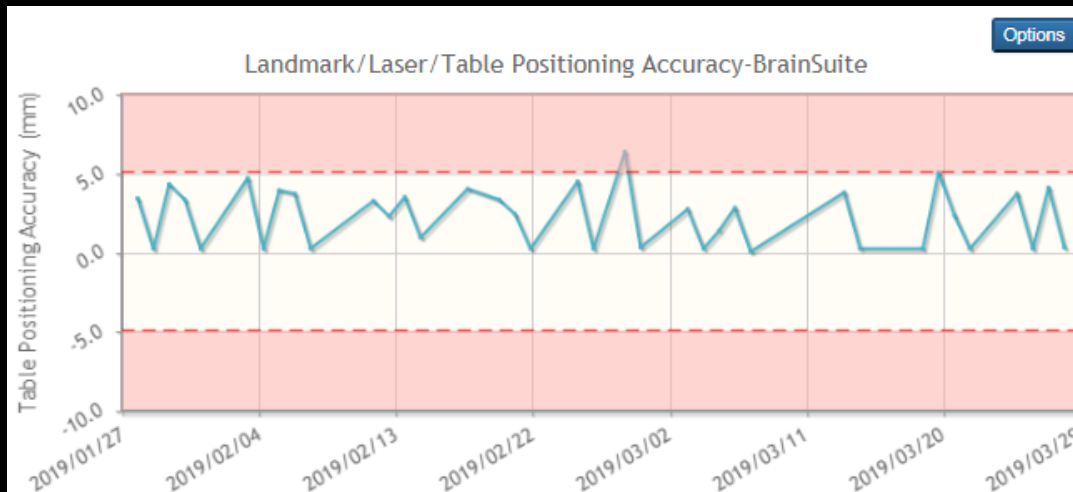
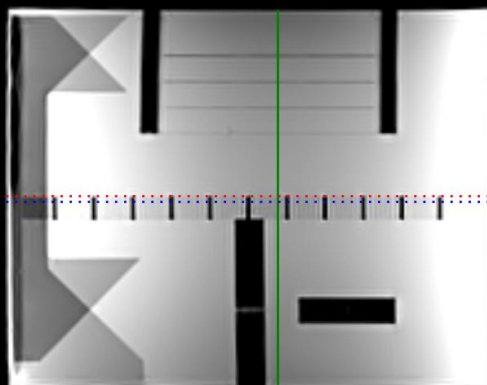
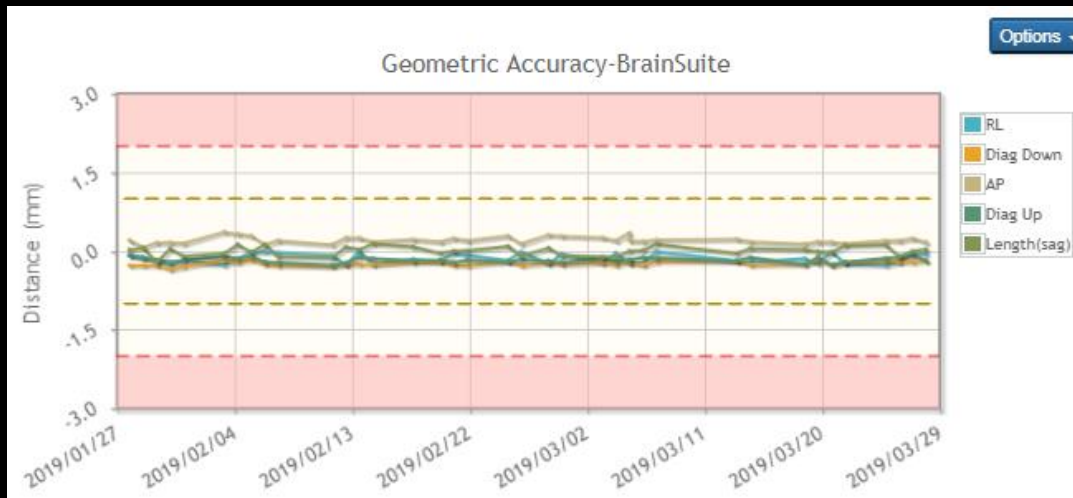
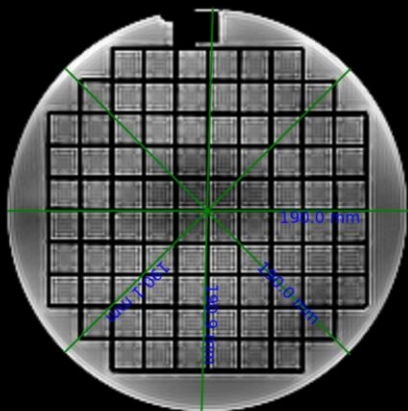


Geometric distortion in MRI

- Field inhomogeneity (δB_0)
 - 2D vs 3D
 - higher rBW
 - higher resolution
 - shimming
- Non-linear gradients
 - software correction
 - 2D vs 3D
- Both effects worsen away from magnet isocenter
 - often this is where navigation fiducials are

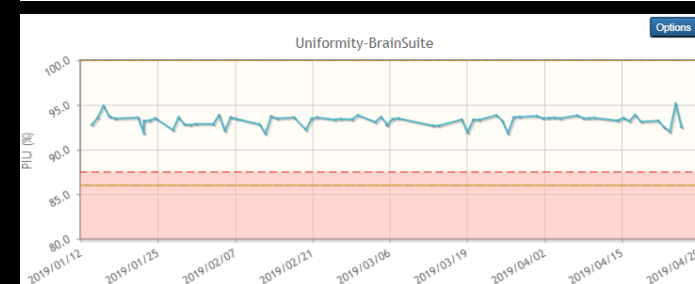
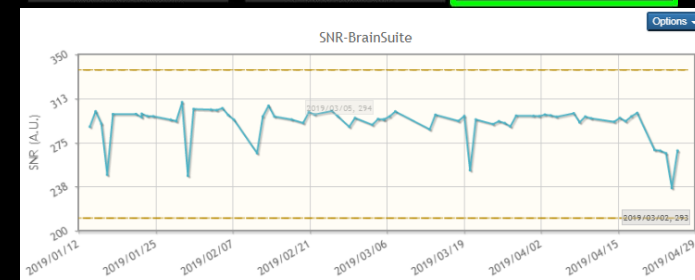
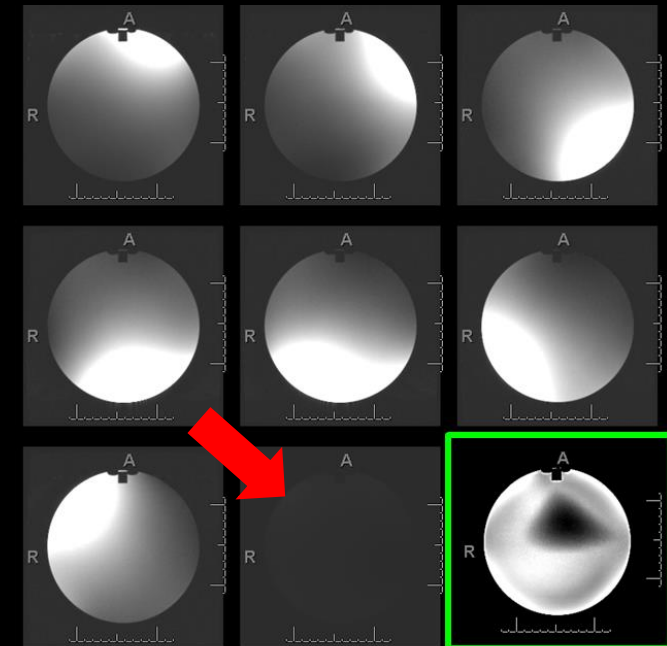


Daily & Periodic MRI Quality Assurance



Daily & Periodic MRI Quality Assurance

- Periodic equipment performance evaluations mimic recommendations of ACR program
 - daily instead of weekly QA
 - tighter geometric distortion constraint
 - test coil used for procedures
 - **ARTIFACT ASSESSMENT**
- Tight control over software/hardware upgrades that may impact therapeutic equipment performance
- Approve/review vendor PM

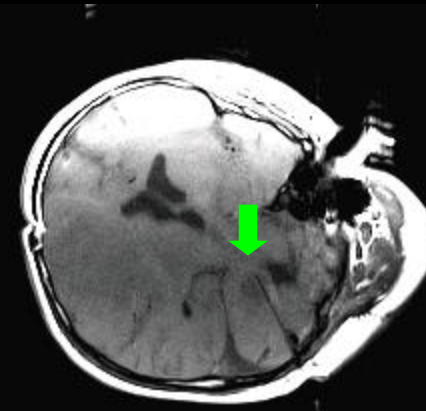


MR-guidance in thermal ablation

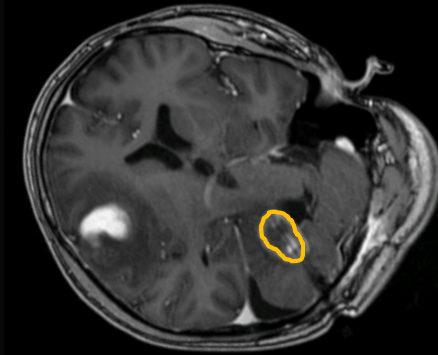
- Potential to use MRI for
 - planning
 - navigation & targeting
 - monitoring & control
 - verification

- Monitor treatment progress
- High temp: device
- Low temp/dose: OAR
- Dose: prediction & control

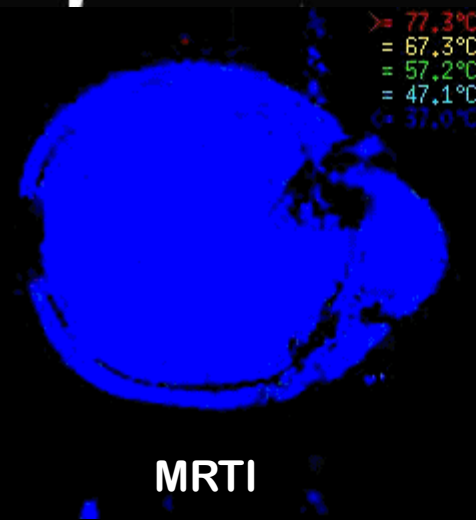
HIGH-GRADE ASTROCYTOMA



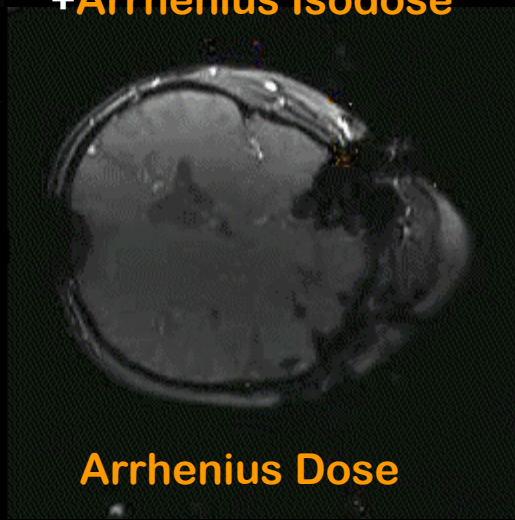
T1-W (fiber + tumor)



CE T1-W Post
+ Arrhenius Isodose



MRTI



Arrhenius Dose



MR Temperature Imaging for monitoring

- Diffusion
- T1-relaxation
- **PRF Shift**
- Linear shift in frequency (γB_0) of α (-0.01 ppm/°C) measured using fast gradient-echo phase-change ($\Delta\phi$)

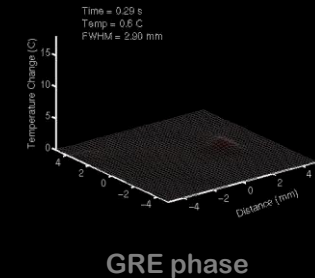
$$\Delta\phi = (2\pi \cdot \gamma B_0 \cdot TE) \cdot \alpha \cdot \Delta T$$

$$SNR_{\Delta\phi} = \frac{\Delta\phi}{\sigma_{\Delta\phi}} \propto TE \cdot SNR_{\text{magnitude}}$$

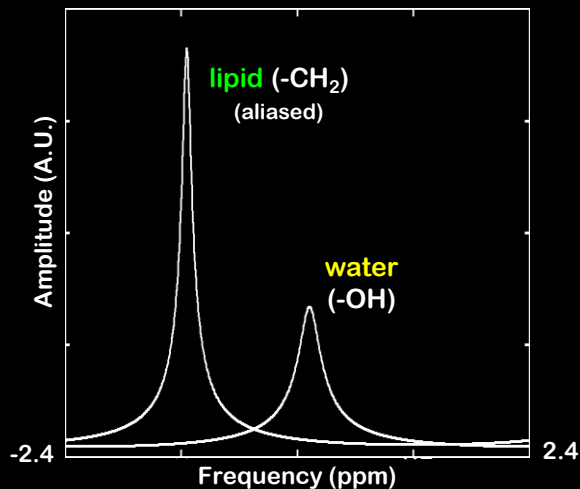
- $TE_{opt} \sim T2^*$
- optimize magnitude SNR for $\sigma_{\Delta\phi}$

- **Challenges**

- *relative* temperature changes
- **lipid** does not shift with temperature
- field drift, susceptibility, motion/flow



(water proton resonance frequency)



Fast chemical shift imaging
(1x1x3mm voxel)

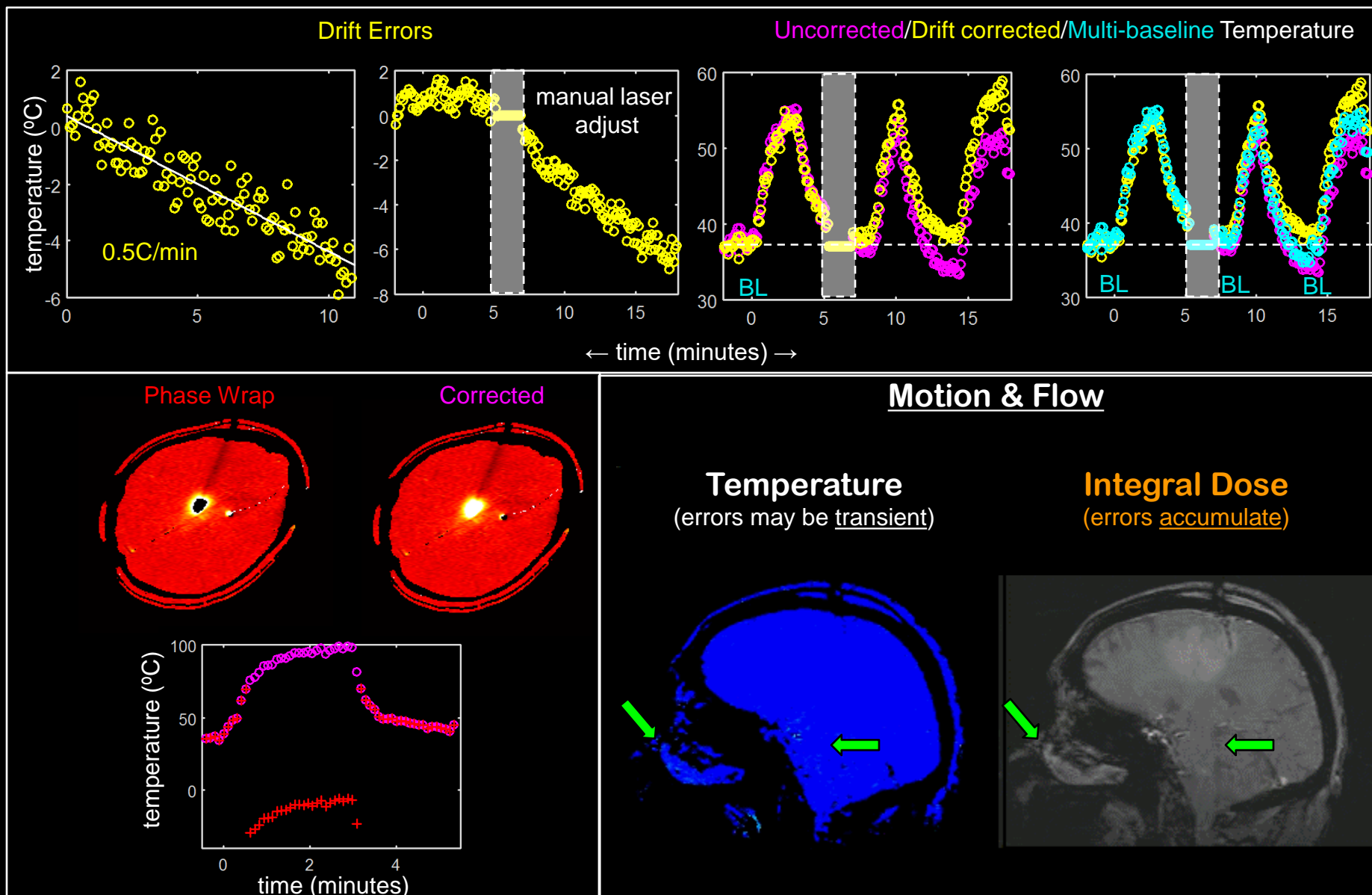


MR thermometry: managing key assumptions

- Phase-changes from temperature only
 - drift
 - magnetic susceptibility
 - phase-wrap
 - motion and/or flow
- Baseline temperature is known
 - remember only temperature **change** measured
 - must acquire reference prior to heating/cooling
 - drift correction: points cannot be in area of active heating or cooling
 - multi-reference: temperature must be back to baseline
 - leave cooling on after ablation to carry away heat
 - do not cool below baseline and then set reference
- Temperature sensitivity coefficient is known
 - lipid, magnetic susceptibility, blood, contrast agents



MR thermometry: managing key assumptions



Thermal dose considerations

- Thermal damage is cumulative effect
 - isotherm characterization of bioeffects limited
 - high/low isotherms good for device/OAR protection
- Arrhenius models (Ω): damage as function of thermal exposure

$$\Omega = A \int_0^t e^{\frac{-E_a}{RT(\tau)}} d\tau$$

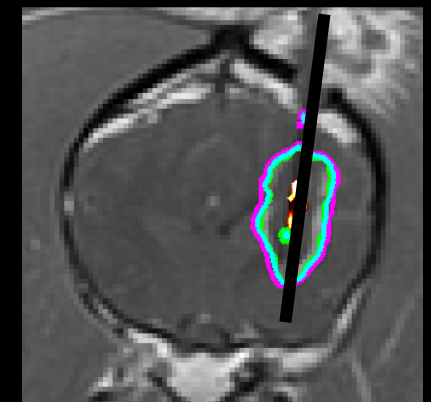
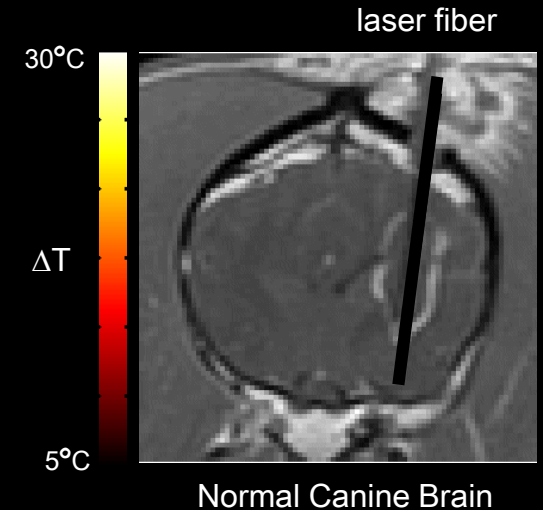
R = Universal Gas Constant
 A = Frequency Factor ($3.1 \times 10^{98} \text{ s}^{-1}$)
 E_a = Activation Energy ($6.3 \times 10^5 \text{ J}$)

Henriques FC, Arch Pathol, (1947)

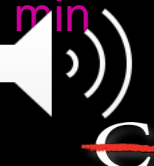
- Cumulative Equivalent minutes @ 43°C (CEM₄₃)
 - compare with hyperthermia exposure isoeffects

$$\text{CEM}_{43}(t_n) = \sum_{t=0}^{n \cdot \Delta t} R^{(43-T_n)} \cdot \Delta t, \text{ with } R = \begin{cases} 0.25 & T_n < 43^\circ\text{C} \\ 0.50 & T_n \geq 43^\circ\text{C} \end{cases}$$

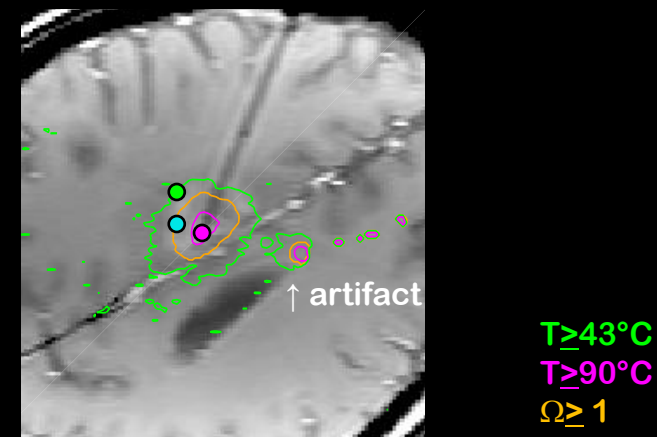
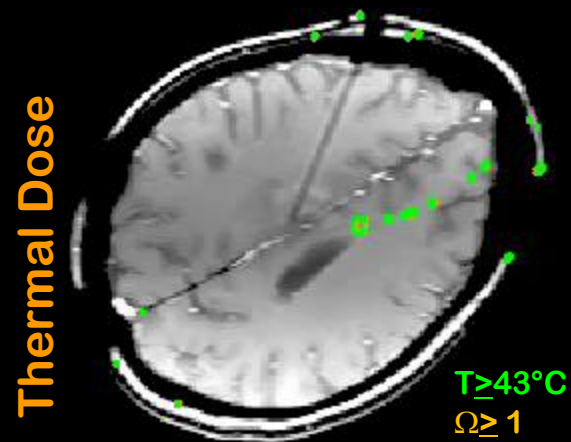
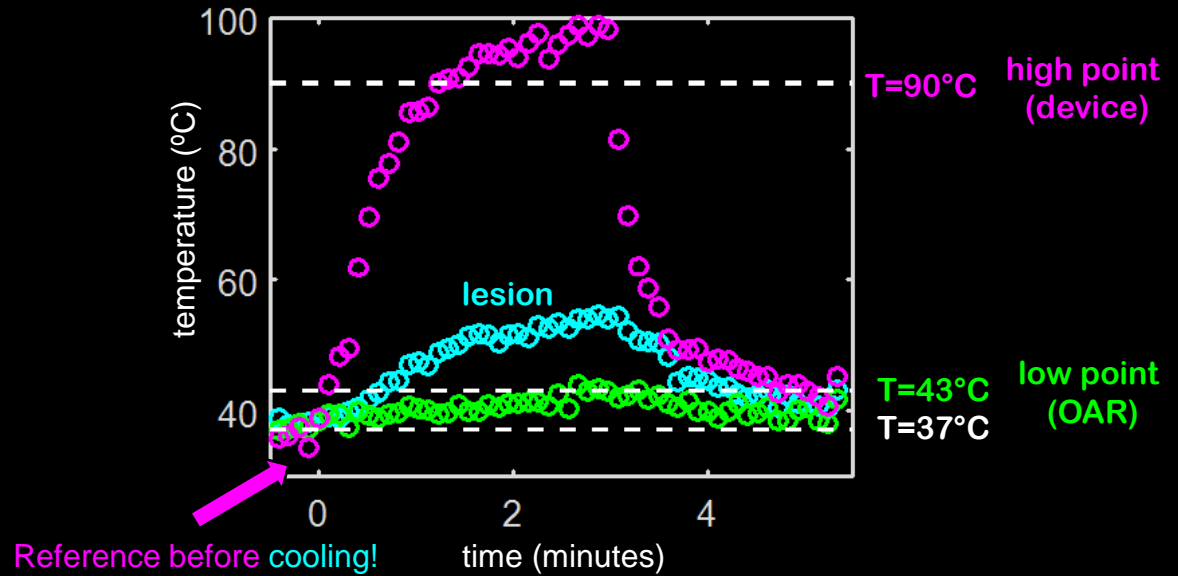
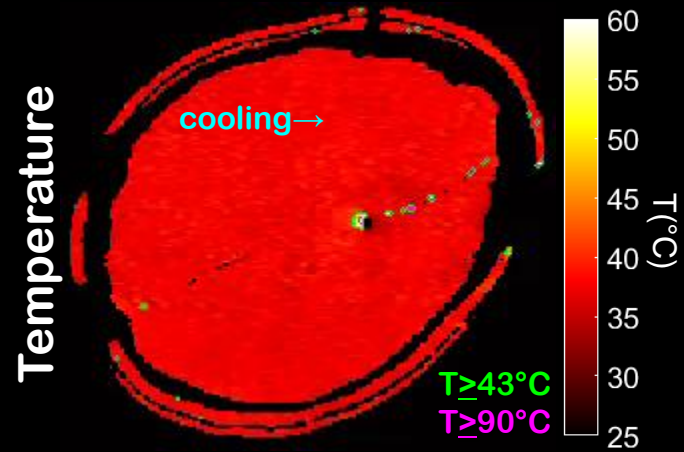
Sapareto SA, Dewey WC Int. J. Rad. Onc. Bio. Phys., 1984.; Damianou C, Hynynen K, JASA, 1994.



$\Omega \geq 1$
 $T \geq 57^\circ\text{C}$
 $\text{CEM}_{43} \geq 240 \text{ min}$

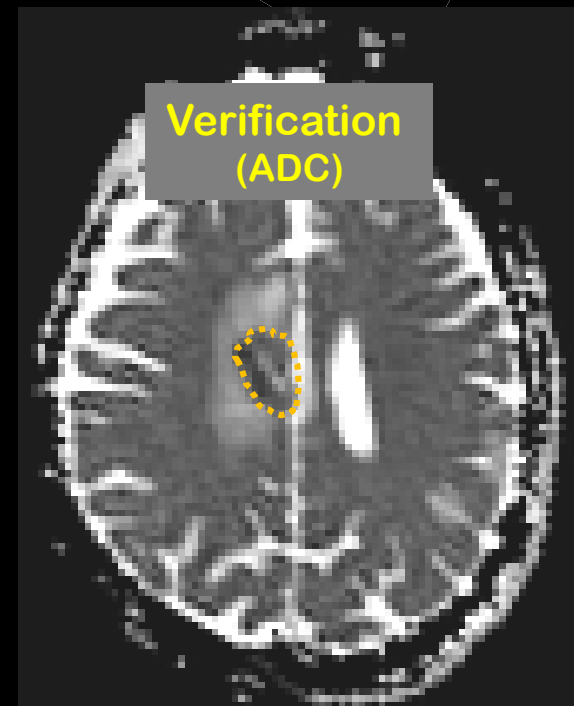
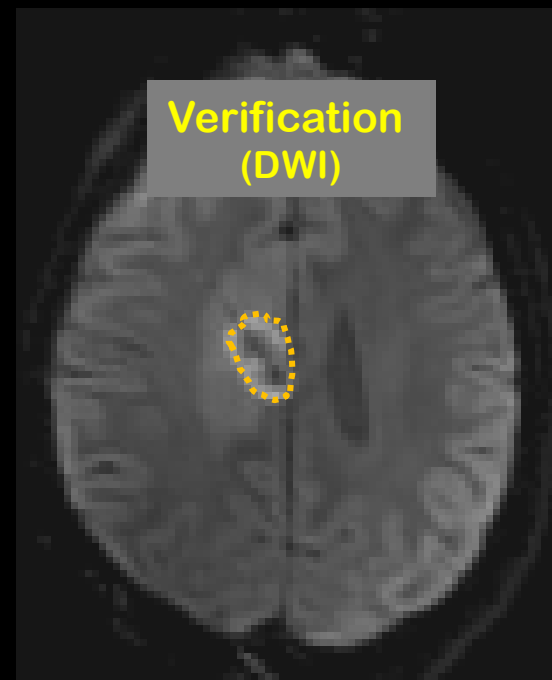
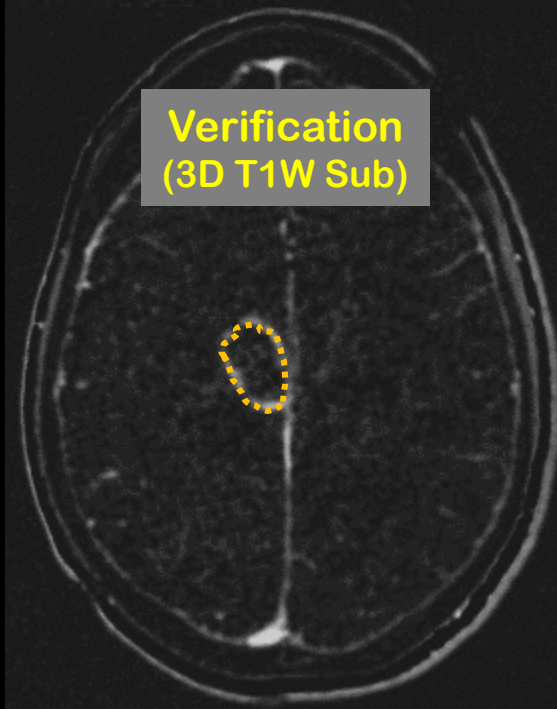
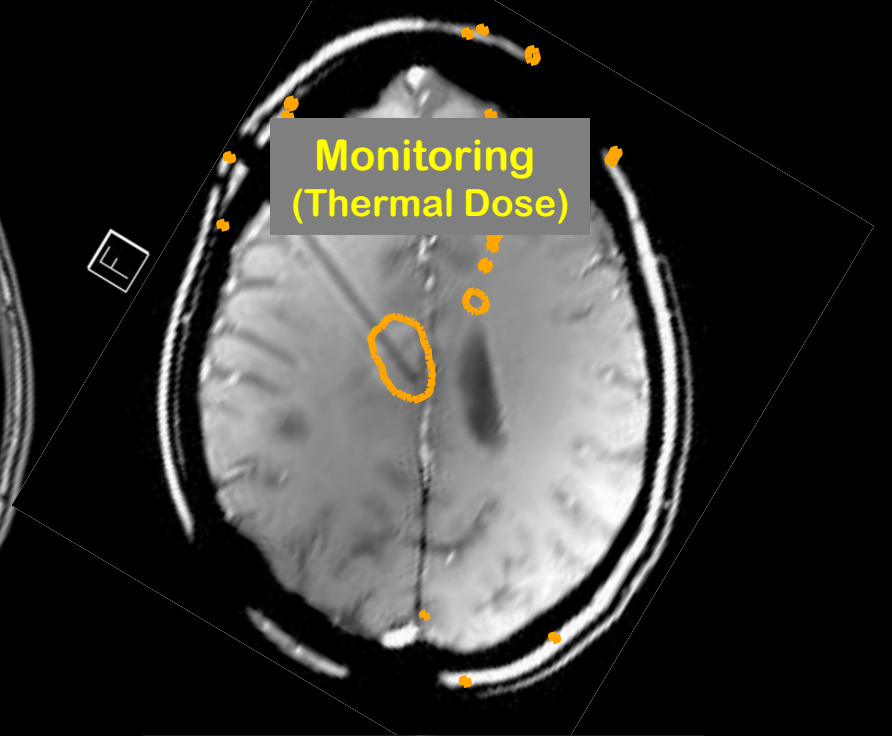
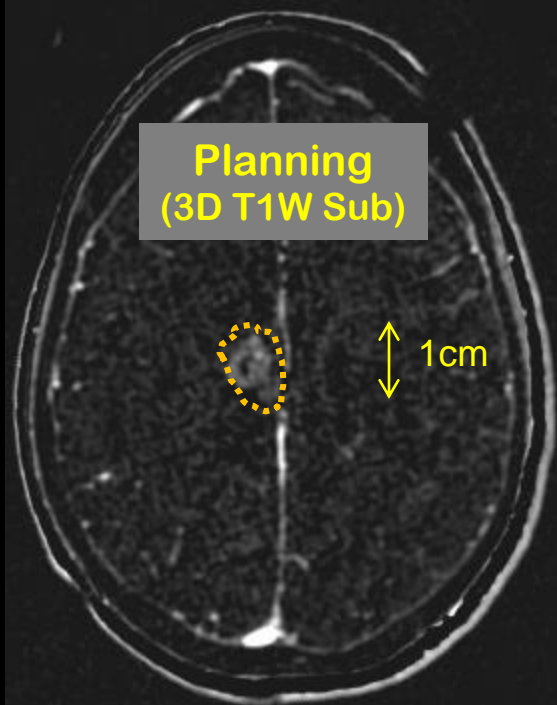


Putting it together: monitoring laser ablations



OAR: if you can't see it, you're not protecting it!





Conclusions

- Holistic understanding of technical laser ablation and MR monitoring process helps avoid pitfalls
 - basic physics, order of operations & procedures
- Some key considerations include:
 - patient & equipment selection
 - planning: size, needed exposure(s) and OAR
 - anatomy to be monitored and ability to perform MRTI over region
 - targeting: distortions and image-verification
 - monitoring: temperature/dose regulation points
 - when to turn cooling on/off
 - when to set/reset MR temperature references
 - MRI protocols vs laser vendor limitations
 - *temperature/anatomy overlay limitations*
 - *do not heat faster than can monitor*
 - **interpreting and avoiding temperature & dose artifacts**



**Thank you for
your time!**

Email: jstafford@mdanderson.org

