

# The Promise of Quantitative Imaging

 $\Lambda \sim$ •Patient stratification in order to decide on alternative Predict treatments •Analysis of heterogeneity within and across lesions (can Virtual assess varying pharmacokinetics, receptor status, proliferative/apoptotic rates, ...) Biopsy •Early prediction of treatment response During •Basis for modifying therapy Tx •Monitoring for Treatment Efficacy After Тх •Longitudinal monitoring and evaluation (can be done before Follow-up then after treatment, substituting for longitudinal tissue biopsy) Quantitative Imaging, Radiology 25 se for Multi-Stakeholder Participation aging, Radiology 258:906-914, 2011

# Modality-Independent Issues

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## General quantitative imaging challenges

- Lack of detailed assessments of sources of bias and precision
- Lack of standards (acquisition, analysis, and reporting)
  - · Varying measurement results across vendors and centers
- Little support from imaging equipment vendors
- No apparent competitive advantage (reimbursement) or regulatory requirements
  - Varying measurement results across vendors
  - Varying measurement results across time for any particular vendor
- Highly variable quality control procedures
  - QC programs, if in place, are typically not specific for *quantitative* imaging
    - Varying measurement results across centers

# Modality-Independent Issues

General quantitative imaging challenges (continued)

- Cost of QIB studies (comparative effectiveness)
- Radiologist acceptance
  - · QIBs are not a part of radiologist education & training.
  - · The software and workstations needed to produce and interpret QIB results are typically not integrated into the radiologists' workflow.
  - · Clinical demand on radiologists is high --- "time is money"
  - · There are few guidelines for QIB reporting.
- Resource availability
  - · Technologists trained in advanced, quantitative, protocols
  - · Physicists and/or imaging scientists, data processing capabilities, etc.

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### Single-vendor, single-site studies:

## Key Challenges

- Acquisition protocol optimization
  - · Scan mode and acquisition parameter optimization for:

    - contrast response and CNR
      temporal resolution (for dynamic imaging)
    - spatial resolution anatomic coverage
  - · Application specific phantoms needed for initial
  - validation scans and ongoing quality control
    - phantom acquisition and data analysis protocols
    - established frequency of assessment and data reporting
- Mechanism for detecting and addressing changes in
- measured response due to system upgrades (Quality Control)
  - Vendors focused on competitive advantage in radiology, not on quantitative imaging applications and maintaining signal response characteristics over time

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#### Single- to multi-vendor studies:

#### Key Challenges

- Acquisition protocol harmonization
  - Scan mode and acquisition parameter selection for matched:
    - contrast response and CNR

    - validation scans and ongoing quality control

## · Can be achieved, but requires substantial effort at start up

- Vendors focused on competitive advantage in radiology, not on quantitative imaging applications and maintaining signal response
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# Single- to multi-<u>center</u> studies:

- Acquisition protocol

# Key Challenges

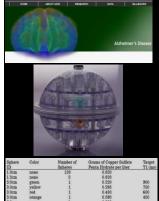
- Harmonization across centers <u>and</u> vendors
- Distribution and activation of protocols
  - Distribute/load electronically
  - Provide expert training and initial protocol load/test
  - Develop / utilize local expertise
- Compliance with protocol
  - Local radiologists, technologists
- Widely varying quality control
  - Ranging from specific for a given imaging biomarker, to ACR
    accreditation, to none
  - Even if QC program is in place, it may not test parameters relevant to the study
- "Scanner upgrade dilemma"
- Data management and reporting
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#### Data Analysis:

## Key Challenges

- Data analysis implementation strategies are often as variable as acquisition strategies
- Choice of model must match data acquisition strategy, *e.g.*, temporal resolution of the acquired data
- Analysis parameters must be standardized, *e.g.*, choice of model, ROI definition, *etc*.
- To facilitate testing/validation of various analysis packages, readily available, standardized test data and analysis results are needed:
  - Digital reference objects
  - Physical phantoms and standardized acquisition protocols and data analysis software
  - Publicly available test/retest human subject data and associated metadata

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# ADNI

- · Multicenter, multivendor study
- Optimized pulse sequence / acquisition parameters for each platform
- MagPhan/ADNI phantom scan at each measurement point
- Access to vendor gradient correction parameters
- With corrections for gradient nonlinearities and optimized acquisition strategies, spatial accuracies of <0.3 mm can be obtained over a ~180 mm spherical volume

# Quantitative MR Imaging Initiatives

| Consensus Group Guidelines |                  |                                                                                |
|----------------------------|------------------|--------------------------------------------------------------------------------|
|                            | NCI:             | RIDER and Academic Center Contracts<br>Imaging Response Assessment Team (IRAT) |
|                            | RSNA:            | Quantitative Imaging Biomarkers Alliance (QIBA)                                |
|                            | ISMRM:           | Ad Hoc Committee on Standards for Quantitative MI                              |
|                            | NCI:             | U01-Funded Quantitative Imaging Network (QIN)                                  |
|                            | ACRIN:<br>(CQIE) | NCI Centers of Quantitative Imaging Excellence                                 |
|                            | Core Labs        |                                                                                |
|                            |                  |                                                                                |