

# Computerized Quantitative Imaging Assessment of Tumor Burden



Quantitative

image feature

analysis to

characterize

tumors and

treatment

response

Daniel L. Rubin, Sandy Napel, Edward Graves, Andrew Quon, George Fisher, Martin O'Connor, Debra Willrett, Andrew Evens Department of Radiology and Medicine (Biomedical Informatics Research), Stanford University

### **OBJECTIVES**

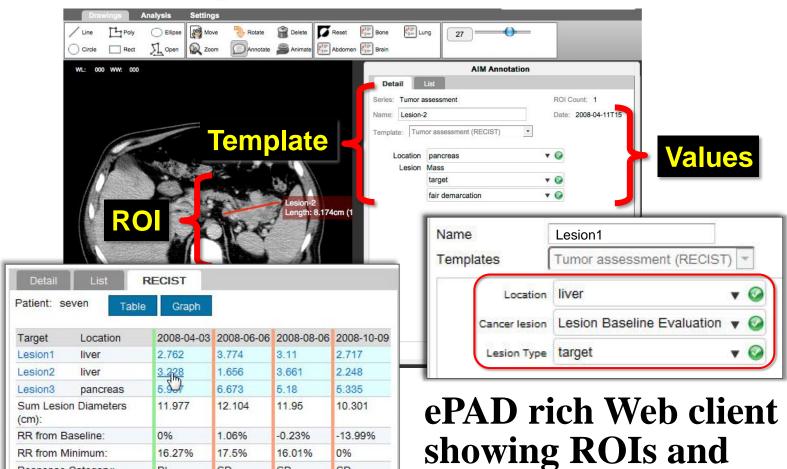
We are develooping a software framework built on caBIG technologies to standardize quantitative imaging assessment of tumor burden and to enable researchers to integrate and analyze a spectrum of quantitative imaging biomarkers to leverage quantitative imaging to better enable assessment of cancer and its treatment response. Our aims are (1) to create tools to reproducibly assess quantitative imaging features of tumor burden; (2) to develop methods to analyze quantitative image metadata and to help oncologists evaluate image-based quantitative criteria of treatment response; and (3) to evaluate the utility of our methods by applying them in two clinical trials and showing an improvement in response assessment in individual patients and patient cohorts.

Challenges we address:

- Poor reproducibility of image measurements
- Lack of *coordination and effective communication* between oncologists and radiologists in making quantitative imaging assessments
- No *standards* for collecting and using quantitative imaging data
- Lack of *tools* for recording image metadata to enable data sharing and data mining

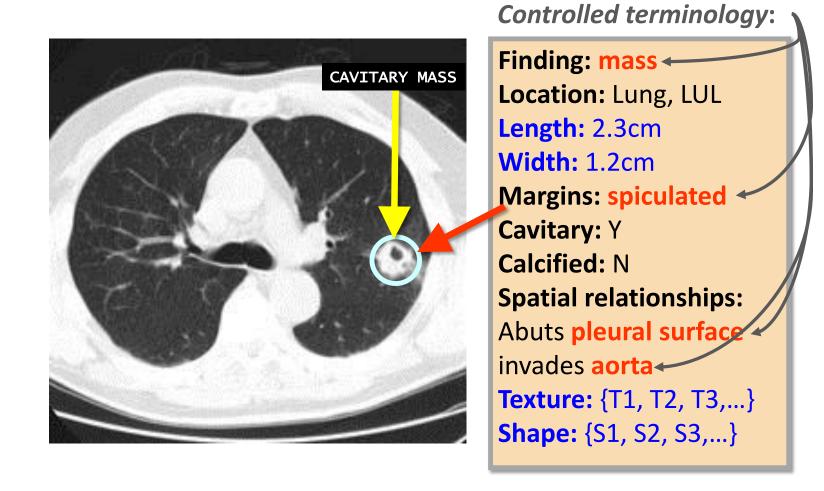
# **AIM 1: Tools and Algorithms**

ePAD (the <u>electronic Physician Annotation</u> <u>Device</u>) implements AIM in a rich Web client.



annotation template

#### ePAD enable semantic annotation of lesions:



Our System Architecture for quantitative imaging includes tools and resources for cancer researchers:

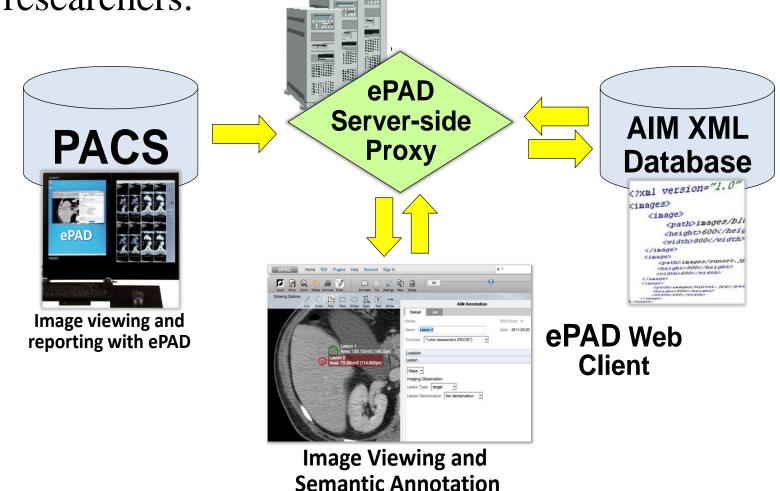
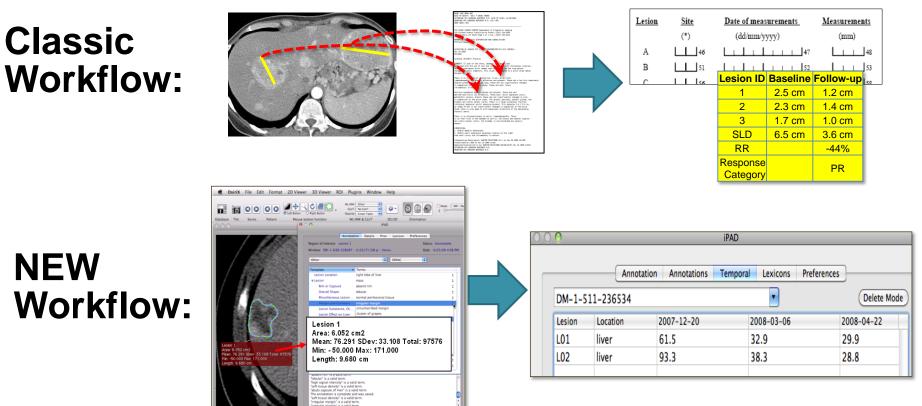
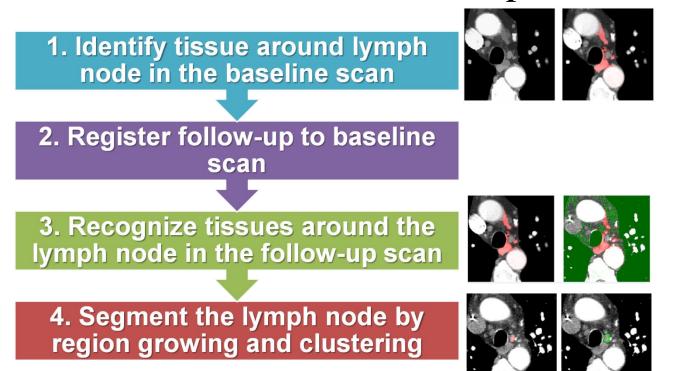


Image metadata storage and workflow for quantitative imaging is now more seamless, with separate data stores for images and their associated metadata, enabling new efficient workflow.



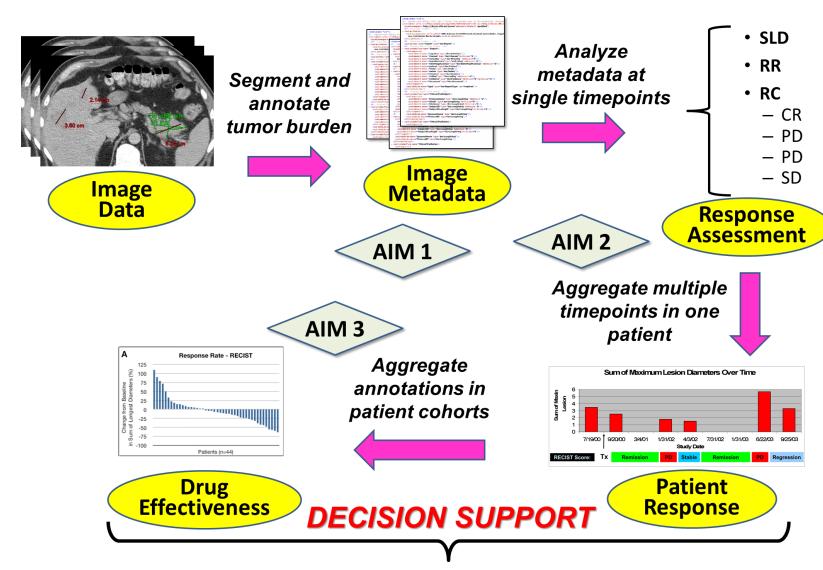
# **AIM 2: Analysis Algorithms**

Algorithms: Automated lesion identification and segmentation on follow up CT imaging. We use the baseline scan to automatically locate and segment cancer lesions on follow up studies.



Tools for **decision support**:

- 1. Resources for archiving/sharing images and image metadata
- 2. Data mining tools to discover alternative quantitative imaging biomarkers of cancer response
- 3. Decision support tools for evaluating patients and alternative treatments



- Tools to automate evaluation of tumor burden
- Quantitative image analysis methods to enable assessing tumor burden
- Application in clinical trials demonstrating value of methods/tools
- Pilot projects/engagement with other QIN sites to prove value to QIN

# AIM 3: Evaluation in Clinical Trials

Pixel and

**Neighbor Values** 

## Follicular lymphoma trial

- ECOG 2408 Randomized Phase II Trial of R-CHOP/R versus R-B-CHOP/R
- PI: Andrew Evens

Lesion Kinetic

**Image** 

- Study endpoints: CR rate after induction, DFS rate, TTP
- Response criteria: IHC criteria with six dominant lesions on CT plus PET

#### Colon Cancer trial

- Phase II Trial of Vandetinib with Capecitabine,
  Oxaliplatin and Bevacizumab
- PI: George Fisher, Stanford University
- Study endpoints: Response rates (RECIST 1.0), time to disease progression
- Response criteria: RECIST 1.0

### OPPORTUNITIES FOR QIN

- Tools for managing mage metadata.
- Resources for archiving images and metadata.
- Data mining tools to discover alternative quantitative imaging biomarkers of cancer response.
- Tools for decision support for treating individual patients (is the cancer responding?) and for evaluating alternative treatments (is the cohort response good?)

## **ACKNOWLEDGEMENTS**

Funding Support: NCI QIN U01CA142555