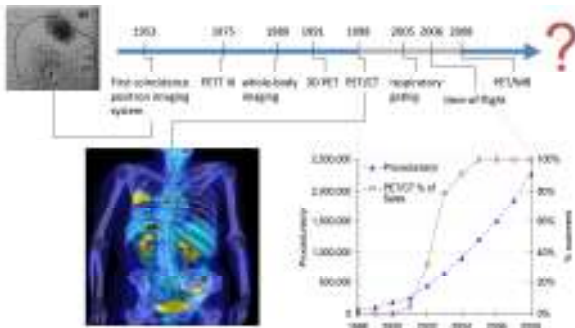


Quantitative FDG-PET/CT: Accelerating development of new therapies and improving assessment of response



PET-CT: A Proud History of Innovation



What's next? Quantitative PET to Characterize Disease Hallmarks

Drivers

- Clinical research, Clinical trials, and Drug discovery
- New molecular diagnostic agents
- Assessing individual response to therapy
- SUVs are now routinely reported, and are asked for, by referring physicians



Biomarkers To Quantify Hallmarks of Cancer



- New molecular diagnostic agents
- New uses for existing agents

Assist with increasing number of oncology targeted pharmaceuticals

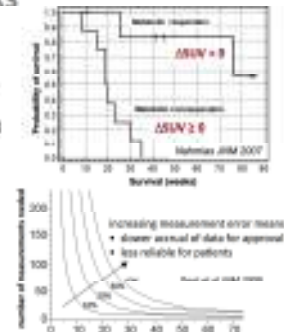
Treatment Population	2007	2008	2009	2010	2011	2012	2013	2014	2015
Cancer patients treated with Anti-angiogenesis treatment	2.8%	3.4%	4.0%	4.7%	5.3%	6.1%	6.7%	7.5%	8.3%



Quantitation Improves Characterization of Disease Hallmarks

Improve individual patient care

- Clinically proven detection and longitudinal quantitation for follow-up
- Moves imaging from diagnostics and staging to therapy assessment
- Accelerate adoption of new molecular diagnostics
- Write clinical trials of new therapies more effectively
- All tied to quantitative accuracy



QIBA FDG-PET/CT TC Projects

Title	PI	Status
Meta-analysis to analyze the robustness of FDG SUV changes as a response marker, post and during systemic and multimodality therapy, for various types of solid metastatic tumors.	D. Harkins, U of the Netherlands	Completed
QIBA FDG-PET/CT Digital Reference Object Project	K. Kinoshita, U of Washington	Completed
Analysis of SARC 13 Trial PET Data by PERCIST with linkage to Clinical Outcomes	S. Wahl, Johns Hopkins	Close to completion
Personnel Support for FDG PET Profile Completion	E. Pedersen, M&P, P. Goshay, U of Washington	Completed
Evaluation of the variability in Determination of Quantitative PET Parameters of Treatment Response Across Performance Sites and Readers	S. Wahl, Hopkins	Needs readers
Evaluation of FDG PET SUV Coefficients, Metrics and Response Criteria	J. Yap, Dana Farber	In Progress
Integration of Retrospective Reviews of 2-3 Groupings of Clinical Trial Datasets (This includes the current Harkins proposal) Will utilize the PERCIST analysis.	D. Harkins, Netherlands	In Progress

QIBA is an Active Sponsor in Regulatory Pathways that Leverage Collaboration



The QIBA Profile Provides Guidance for all aspects of quantitative FDG-PET/CT



What we've done and how you can participate

- ✓ Collection of recommendations for quantitative FDG-PET
- ✓ Presentation (joint with FNHI) to FDA
- ✓ NIBIB grant application
- ✓ Year 1 research targets
- Collaboration with UPICT on Protocols
- Bi-weekly telephone conferences
- Annual QIBA meetings and updates at RSNA and SNM
- Working visits with vendors
- Profile development
- Year 2 research targets
- o Profile testing and approval
- o Implementation of Profiles by QIBA and vendors)
- o Clinical use of Profiles

More information at www.rsna.org/research/qiba/