National Photonics Initiative (NPI) Cancer Moonshot Task Force: Medical Imaging Used to Accelerate Progress

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Introduction:

Eight of the ten focus areas, recommended by the Cancer Moonshot Blue Ribbon Panel and implemented under the direction of the National Cancer Institute, require the support of medical imaging.

At a workshop entitled, "Improving Early Detection of Cancer and Response to Therapies through Imaging Technologies" in Rockville, Maryland on April 12, 2017, the NPI convened stakeholders including the medical imaging industry, academia, government agencies and patient advocates to identity key issues and propose solutions for more effective use of medical imaging to accelerate the

Challenges/Opportunities:

Developed as part of the Precision Medicine Initiative, the national IT "cancer knowledge network," which aims to house and integrate genomic information from tumors with clinical response data and outcomes information, does not currently include medical imaging data – data that is digital and ideally suited for machine learning on "big data."

Imaging data provides a wealth of information, which can be computer extracted to yield "radiomics," e.g., quantitative descriptors of tumor size, shape and heterogeneity, allowing image features to be related to protein and gene expression through artificial intelligence and deep learning. Imaging and radiomics datasets could be shared on the cancer knowledge network and "data-mined" to advance medical discoveries and accelerate translation of innovations into clinical practice.







Response to

therapy

Registries for

accelerating

research,

developing

medical

evidence

Biomarker in

clinical studies

translation of innovations into clinical practice.

The Unique Value of Medical Imaging:

Tumor heterogeneity and the sampling problems associated with repeated tumor biopsies renders incomplete information. Cancer medical imaging is essential to overcome limitations to biopsies, and with machine learning, can potentially yield virtual digital biopsies, which are noninvasive, repeatable and cover the entire tumor.

Today and in the future, imaging drives diagnosis and decisionmaking in efficient and effective cancer patient care.

Examples:



MR/ultrasound fusion-guided biopsy showing position of 1 biopsy core



PET/CT Monitoring Response to Adjuvant Immunotherapy in High-Risk Pediatric Sarcomas. RIGHT: FDG- PET scans taken at presentation of 24 years of age showing disseminated Ewing Sarcoma involving kidney, bone, bone marrow and lungs. She received standard cytotoxic therapy followed by immunotherapy. The image on the left demonstrates FDG-PET scans taken at 4.5 years following presentation. She remains free of disease with no evidence of recurrence. From Merchant, MS et al., Clin. Cancer Res., 22, 13; 3182 2016.





Also, given variations in image acquisition protocols, developments to standardize across vendors is needed so that imaging data can be compared and utilized across different clinical sites for purposes of research and medical evidence development.

Recommendations:

Collaboration is required to develop application-specific standards and protocols to accelerate contributions of medical imaging to the cancer knowledge network.

Medical data registries need to be developed to accommodate medical imaging datasets for rapid collection, computer image analyses and sharing that will accelerate patient benefit and quality outcomes assessments across sites and providers.



Datamining of Radiomics-Genomics Associations in Breast Cancer Discovery. LEFT: Heatmap representation of statistically significant associations between radiomic phenotypes and transcriptional activities of some cancer-related genetic pathways. In the heatmap, genetic pathways are rows and radiomic phenotypes are columns. From Zhu, Y et al., Nature Scientific Reports 5:17787, 2015.

When used as a quantitative biomarker, imaging can provide surrogate endpoints for accelerating clinical trials. Because of its importance in screening, diagnosis and monitoring response to therapy, imaging data needs to be included in registries, databases and the "cancer knowledge network" as a necessary accelerant for translating innovation into clinical practice.

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Patient stakeholders need the ability to contribute to and access their own medical images and clinical diagnostics with privacy guidelines in order for the cancer knowledge network to evolve into a database that facilitates translational research and clinical evidence development.

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