

AAPM Grand Challenges Symposium

**The AAPM Working Group
on Grand Challenges**

S.G. Armato III



Disclosures

- Royalties and licensing fees for CAD technology from the University of Chicago
- Consultant, Aduro Biotech, Inc.

Challenges

- Allows comparison of different algorithms
 - all playing by the same rules,

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Challenges

- Allows comparison of different algorithms
 - all playing by the same rules,
 - on a common set of images,
 - reporting standardized output,
 - with uniform performance assessment.

Challenges

- Controlled environment
 - database of images
 - training/testing paradigm
 - performance evaluation
 - scoring method
 - reporting structure

How to Host a Challenge

- Central organization

How to Host a Challenge

- Central organization
 - time, effort, resources

How to Host a Challenge

- Central organization
 - time, effort, resources, and a crystal ball



How to Host a Challenge

- Challenges will not succeed without groups willing to “accept the challenge”

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- “Build it and they will come”



How to Host a Challenge

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- “Build it and they will come”
- Organizers must “build it SO they will come”

How to Host a Challenge

- Challenge must be considered reputable
 - worthy of participants’ time and effort
- Challenges should offer an incentive
 - conference panel or co-authorship

How to Host a Challenge

- Challenge must be considered reputable
 - worthy of participants’ time and effort
- Challenges should offer an incentive
 - conference panel or co-authorship
 - fame and fortune

AAPM in the Challenge Arena

CT Low-Dose Challenge

Low-dose CT for the detection and classification of metastatic liver lesions: Results of the 2016 Low Dose CT Grand Challenge

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Farhana Khan

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Shuai Leng, Kyle L. McMillan, Gregory J. Michalak, Kristina M. Nunez, Lifeng Yu, and

Joel G. Fletcher

Department of Radiology, Mayo Clinic, Rochester, MN 55905, USA

(Received 28 October 2016; revised 11 May 2017; accepted for publication 11 May 2017;

published 12 October 2017)

CT Low-Dose Challenge

A fast method to emulate an iterative POCS image reconstruction algorithm

Gengsheng L. Zeng¹

Department of Engineering, Weber State University, Ogden, UT 84408, USA

Department of Radiology and Imaging Sciences, University of Utah, Salt Lake City, UT 84108, USA

A deep convolutional neural network using directional wavelets for low-dose X-ray CT reconstruction

Eunhee Kang,^{*} Junhong Min,^{*} and Jong Chul Ye²

Bio Imaging and Signal Processing Lab., Dept. of Bio and Brain Engineering, KAIST, Daejeon, Korea

Low-dose CT reconstruction using spatially encoded nonlocal penalty

Kyungsang Kim, Georges El Fakhri, and Quanzheng Li¹

Gordon Center for Medical Imaging, Massachusetts General Hospital and Harvard Medical School, 125 Nashua Street 6th floor,

Suite 660, Boston, MA 02114, USA

SPIE-AAPM-NCI LUNGx Challenge

- Computerized lung nodule classification
 - Armato et al. JMI, 2015.
 - Armato et al. JMI, 2016.
- Collaboration among SPIE, AAPM, and NCI
 - University of Chicago, University of Michigan, and Oak Ridge National Laboratory

SPIE-AAPM-NCI PROSTATEx Challenges

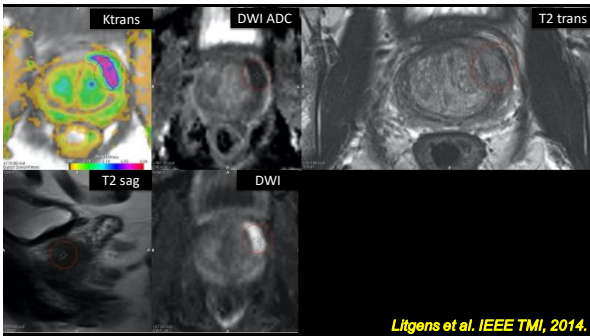
- PROSTATEx: SPIE Medical Imaging 2017
- PROSTATEx-2: AAPM Annual Meeting 2017

SPIE-AAPM-NCI PROSTATEx Challenges

- PROSTATEx: SPIE Medical Imaging 2017
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- Same data: multi-parametric prostate MR scans
- Two diagnostic tasks

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- Same data: multi-parametric prostate MR scans
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SPIE-AAPM-NCI PROSTATEx Challenges

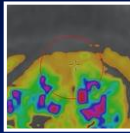
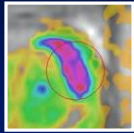
- PROSTATEx must not bias PROSTATEx-2

PROSTATEx

SPIE-AAPM-NCI Prostate MR Classification Challenge

PROSTATEx

Task: Distinguish between “clinically significant” and “not clinically significant” prostate lesions on MR



PROSTATEx

Task: Distinguish between “clinically significant” and “not clinically significant” prostate lesions on MR

Timeline:

- November 21, 2016: training cases available
 - 330 lesions with truth (binary clinical significance)
 - spatial location
 - zone (anterior fibromuscular stroma, peripheral, seminal vesicle, transition)

PROSTATEx

Task: Distinguish between “clinically significant” and “not clinically significant” prostate lesions on MR

Timeline:

- November 21, 2016: training cases available
 - 330 lesions with truth (binary clinical significance)
- December 12, 2016: test cases available
 - 208 lesions **without** truth
 - spatial location
 - zone

PROSTATEx

Task: Distinguish between “clinically significant” and “not clinically significant” prostate lesions on MR

Timeline:

- November 21, 2016: training cases available
 - 330 lesions with truth (binary clinical significance)
- December 12, 2016: test cases available
 - 208 lesions **without** truth
- February 15, 2017: SPIE Medical Imaging 2017

PROSTATEx

Task: Distinguish between “clinically significant” and “not clinically significant” prostate lesions on MR

- submit single real number on $[0, 1]$ for each lesion representing likelihood of clinical significance
 - performance assessment: ROC analysis

PROSTATEx-2

SPIE-AAPM-NCI Prostate MR Gleason Grade Group Challenge

PROSTATEx-2

Task: Develop quantitative MR biomarkers for the determination of Gleason Grade Group in prostate cancer

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Timeline:

- May 15, 2017: training cases available
 - 112 lesions with truth (Gleason Grade Group)
 - spatial location
 - zone (anterior fibromuscular stroma, peripheral, seminal vesicle, transition)

PROSTATEx-2

Task: Develop quantitative MR biomarkers for the determination of Gleason Grade Group in prostate cancer

Timeline:

- May 15, 2017: training cases available
 - 112 lesions with truth (Gleason Grade Group)
- June 5, 2017: test cases available
 - 70 lesions **without** truth
 - spatial location
 - zone

PROSTATEx-2

Task: Develop quantitative MR biomarkers for the determination of Gleason Grade Group in prostate cancer

Timeline:

- May 15, 2017: training cases available
 - 112 lesions with truth (Gleason Grade Group)
- June 5, 2017: test cases available
 - 70 lesions **without** truth
- August 1, 2017: AAPM Annual Meeting 2017

PROSTATEx-2

Task: Develop quantitative MR biomarkers for the determination of Gleason Grade Group in prostate cancer

- submit ordinal value on [1, 5] for each lesion representing Gleason Grade Group
 - performance assessment: quadratic-weighted kappa

2019 Digital Pathology Challenge

SPIE-AAPM-NCI CellularityX Challenge

SPIE-AAPM-NCI CellularityX Challenge

- Develop algorithms to automatically assess **cellularity** in pathology whole slide image patches
- **Cellularity** assessed as
 - a score and a categorical classification
 - 0 (normal)
 - 1–30 (low cellularity)
 - 31–70 (medium cellularity)
 - 71–100% (high cellularity)

SPIE-AAPM-NCI CellularityX Challenge

- Reference standard: clinical assessment of cellularity by two pathologists

SPIE-AAPM-NCI CellularityX Challenge

Stay tuned for further details

AAPM Working Group on Grand Challenges

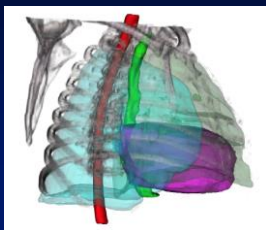
AAPM Working Group on Grand Challenges

- Mission 1
 - Identify “hot topic” challenge tasks and organize internally

AAPM Working Group on Grand Challenges

- Mission 1
 - Identify “hot topic” challenge tasks and organize internally
- Mission 2
 - Solicit challenge proposals from the community
 - Assist with resources and expertise

Auto-Segmentation for Thoracic Radiation Treatment Planning Challenge



AAPM Working Group on Grand Challenges

Proposals seek:

- title of challenge (including proposed nickname)
- background on the issue to be addressed
- statement of the unmet need to be served

AAPM Working Group on Grand Challenges

Proposals seek:

- challenge methodology and logistics
 - source and numbers of cases for training/testing
 - parameters to be held fixed or to be “matched” across cases
 - reference standard

AAPM Working Group on Grand Challenges

Proposals seek:

- challenge methodology and logistics
 - proposed challenge hosting platform
 - proposed data hosting platform
 - performance assessment metric(s)
 - publication plan
 - plan for data persistence

AAPM Working Group on Grand Challenges

Proposals seek:

- WGC assistance requested (logistical, financial)

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We want to help you “build it SO they will come”



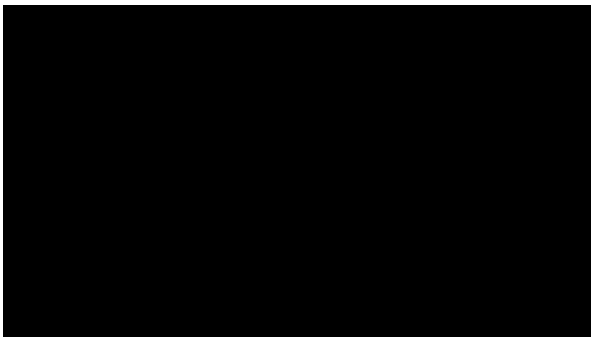
Sparse-View Reconstruction Challenge for 4D Cone-Beam CT (SPARE Challenge)

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- Participants developed algorithms to reconstruct cone-beam CT images from sparsely sampled projection data

Sparse-View Reconstruction Challenge for 4D Cone-Beam CT (SPARE Challenge)

- Participants developed algorithms to reconstruct cone-beam CT images from sparsely sampled projection data
- Organized by Andy Shieh, The University of Sydney
 - with WGGC support





Dataset

- MR scans of each patient contained
 - Transaxial and sagittal T2-weighted images
 - K_{trans} images (computed from dynamic contrast-enhanced images)
 - Apparent diffusion coefficient (ADC) images (computed from diffusion-weighted images)
- Thumbnail of each lesion for visual reference

Litgens et al. IEEE TMI, 2014.

Rules

- Train with training set without restriction
- No human intervention with test set, except for manual or human-supervised delineation of prostate boundary or gross lesion margin
- Participants not allowed to withdraw from the Challenge after test set groupings submitted

Challenges

- Friendly competition
- Educational
- Participants deserve much credit

Challenge Considerations

- Database

Challenge Considerations

- Database
 - How many cases? balance effort/cost vs. power
 - Technical parameter consistency
 - slice thickness
 - contrast enhanced
 - Lesion characteristics consistency
 - size
 - histology

Challenge Considerations

- Database
- Instructions

Challenge Considerations

- Database
- Instructions
 - Acceptable level of human input
 - Appropriate use of test set
 - Don't assume

Challenge Considerations

- Database
- Instructions
- Clinical data

Challenge Considerations

- Database
- Instructions
- **Clinical data**
 - limit to DICOM header (age, gender)?

Challenge Considerations

- Database
- Instructions
- Clinical data
- **Commitment by participants**

Challenge Considerations

- Database
- Instructions
- Clinical data
- **Commitment by participants**
 - anonymity

Challenge Considerations

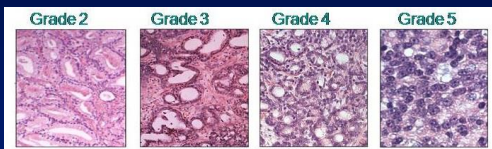
- Database
- Instructions
- Clinical data
- Commitment by participants
- **Epilogue**

Challenge Considerations

- Database
- Instructions
- Clinical data
- Commitment by participants
- **Epilogue**
 - image data persistence
 - publish?

Gleason Score (Grade)

- Reflects the appearance of cancer cells from biopsy
 - 1 = similar to normal prostate cells
 - 5 = very abnormal



<http://urologyrbh.blogspot.com/2011/06/prostate-biopsy.html>

Gleason Score (Grade)

- Reflects the appearance of cancer cells from biopsy
 - 1 = similar to normal prostate cells
 - 5 = very abnormal
- Two ratings assigned
 - one to the most dominant cell appearance
 - one to the second-most dominant appearance
- The two ratings are combined: Gleason score [2-10]

Gleason Score (Grade)

- Clinically meaningful binning of Gleason scores

Gleason Grade Group	Gleason Score
1	≤ 6
2	7 (3 + 4)
3	7 (4 + 3)
4	8
5	9 or 10

Epstein et al. Am J Surg Pathol, 2016.
