

Artificial Intelligence in Medical Imaging for Precision Medicine

Vaanathi Sundaresan

Biomedical Image Analysis Laboratory Department of Computational and Data Sciences (CDS) Indian Institute of Science, Bangalore

vaanathi@iisc.ac.in





भारतीय विज्ञान संस्थान





Talk overview

- Medical imaging in Precision medicine
- Al Applications in imaging for precision medicine
 - Individual subject-level
 - Population-level demographic factors-driven
- Challenges in AI tool development
 - Data/label limitations
 - Domain shifts across centres and scanners
 - Data sharing and privacy
- Opportunities/future directions





What is precision medicine?

- Personalized care
 - subject's medical history / lifestyle factors
 - Disease biomarkers
 - Genetic factors
- Patient complains about mood changes and gait disturbances
 - Could be symptoms of many disorders
 - Knowing additional information could help (e.g., hypertension, lesions in specific regions, smoking/stress etc.)



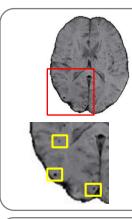
Medical imaging in precision medicine

- Early diagnosis
 - Routine scans, incidental findings
- Differential diagnosis
 - Differentiating diseases with similar symptoms
- Screening and treatment planning
 - Longitudinal analysis, response to drug administration
- Monitoring and predictive analysis
 - Disease severity prediction, pre- vs post-surgery analysis
- Data-driven and hybrid decision making
 - Using scans and *imparting domain* (clinical) knowledge/factors
- Further understanding of disease conditions
 - Large-scale epidemiological studies, *population-level modelling*





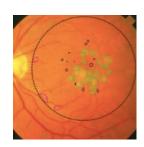
Al application in imaging for precision medicine



Early diagnosis

Detection of cerebral microbleeds

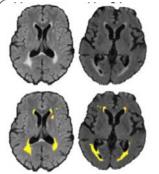
Early sign of neurodegeneration of various cerebrovascular diseases

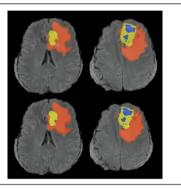


Screening & monitoring

Screening diabetic retinopathy and macular degeneration

Scoring based on early, intermediate and late signs





Extraction of imaging biomarkers for diseases

Segmentation of white matter hyperintensities and brain tumor lesions – estimate the lesion load

Useful to investigate their distribution and rating to get more information on their incidence.

V Sundaresan et al. NeuroImage, 2019a V Sundaresan et al. ISMRM, 2020 V Sundaresan et al. Medical Image Analysis, 2021 V Sundaresan et al. OMIA, MICCAI, 2015 V Sundaresan et al. IEEE EMBC, 2015 V Sundaresan et al, OHBM, 2018, 2019

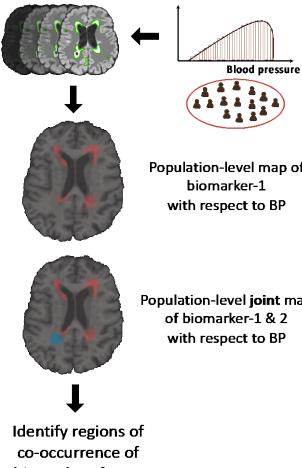


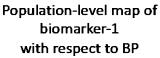


Demographic factors-driven precision diagnosis

Modelling the distribution of anomalies within a population

Useful in obtaining a population-level prior for specific diseases





Population-level joint map of biomarker-1 & 2 with respect to BP

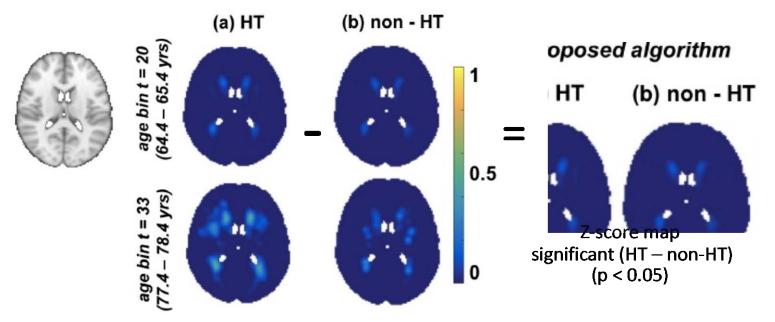
biomarkers for BP





Modelling white matter lesion distribution within a population

biobank^{**} Largest brain imaging database, currently ~40,000 subjects scanned, with the target of 100,000 subjects. Big data - total database size: 500 TB.







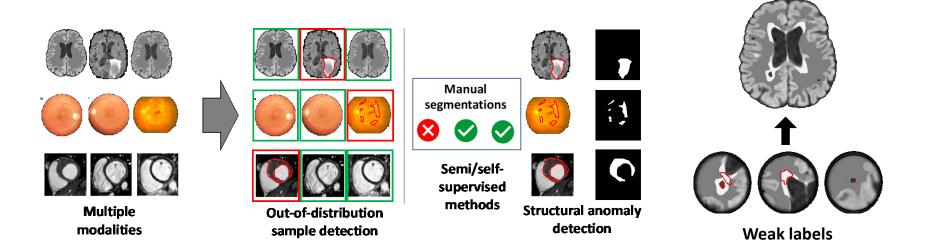
Challenges in Al tool development





Data/label limitations

- Medical data is sensitive and difficult to obtain, have to be anonymized
- Manual annotation of medical imaging is time-consuming and expensive
- Self-/semi-supervised techniques

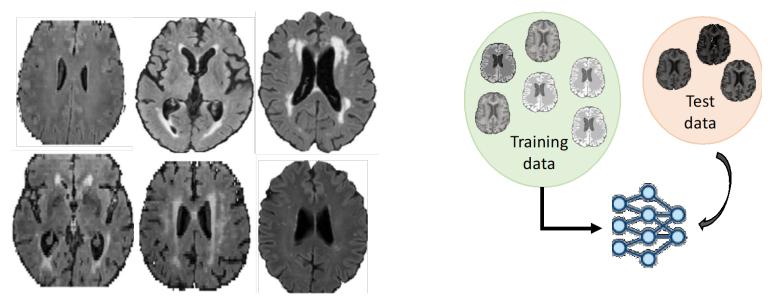






Data domain shifts in multicentric data

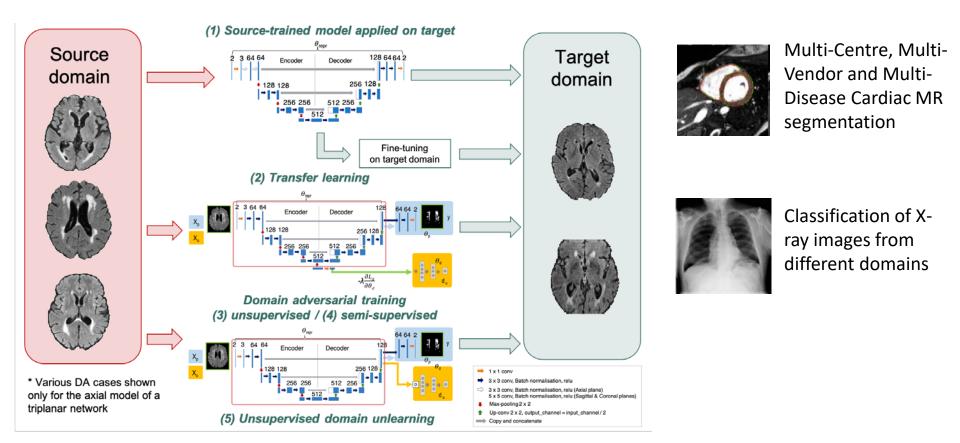
Domain adaptation: Transfers knowledge from source to target domain by leveraging domain invariant features to reduce domain shift.







Comparison of domain adaptation (DA) techniques for neurodegeneration lesion segmentation



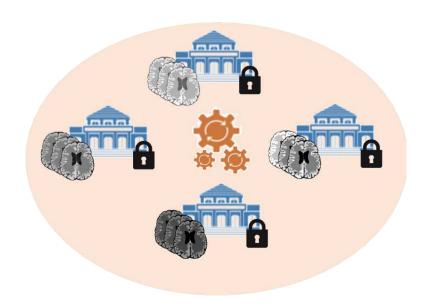


Data sharing and privacy protection

- Federated learning:
 - Preserves data privacy
 - Model training done individual centres rather than sharing data
 - Aggregate model parameters centrally from individual models

- Emerging areas:

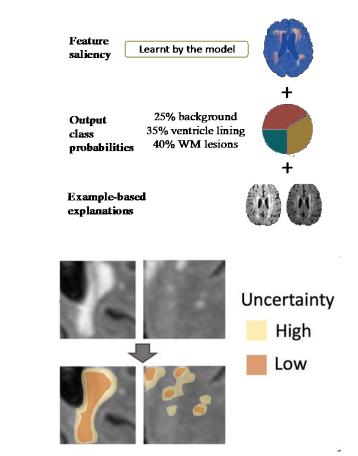
- Addition of unseen centres in FL, cross-modal domain adaptation, missing modalities





Opportunities/future directions

- Interpretability and Trust explainability
 - Including clinicians in the loop
- Out-of-distribution detection
 - Open-world anomaly detection (openset recognition)
 - Integration within a federated learning set-up
 - Anomaly informed precision diagnosis (incidental findings)
- Accurate extraction of underexplored biomarkers
 - Disentanglement of source (e.g., pathological vs scanner-induced)
 - Characterization and reporting





THANK YOU!

IISc, Bangalore

Biomedical image analysis Laboratory

Ramanujam N Shreyas H R Gargi G Collaborators University of Oxford WIN analysis group Mark Jenkinson Ludovica Griffanti Steve Smith Karla L. Miller Nicola K. Dinsdale IBME group Alison Noble Vicente Grau Ana IL. Namburete

Collaborators Politechnico di Milano Valentina Bordin Ilaria Bertani University of Modena and Reggio Emilia Giovanna Zamboni University of Siena Marco Battaglini Nicola De Stephano Giordano Gentile University of Nottingham Stam N. Sotiropoulos Robert A. Dineen Dorothee P. Auer Nikola Sprigg



भारतीय विज्ञान संस्थान