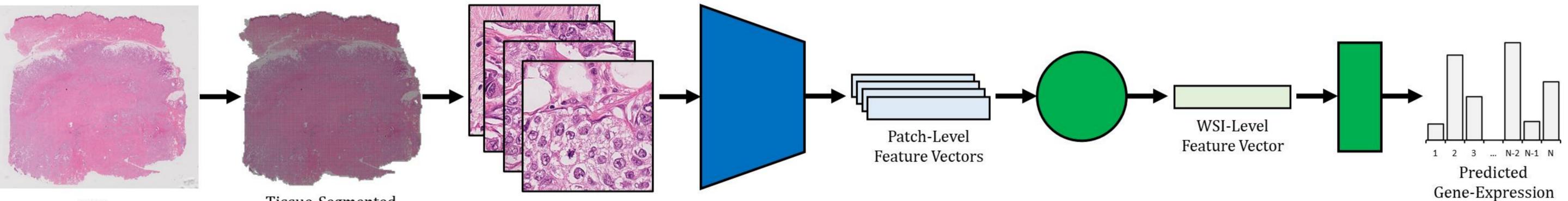
Evaluating Deep Regression Models for WSI-Based Gene-Expression Prediction

Fredrik K. Gustafsson¹, Mattias Rantalainen^{1, 2}

¹Department of Medical Epidemiology and Biostatistics, Karolinska Institutet, Stockholm, Sweden

²MedTechLabs, BioClinicum, Karolinska University Hospital, Solna, Sweden



WSI

Tissue-Segmented & Patched WSI

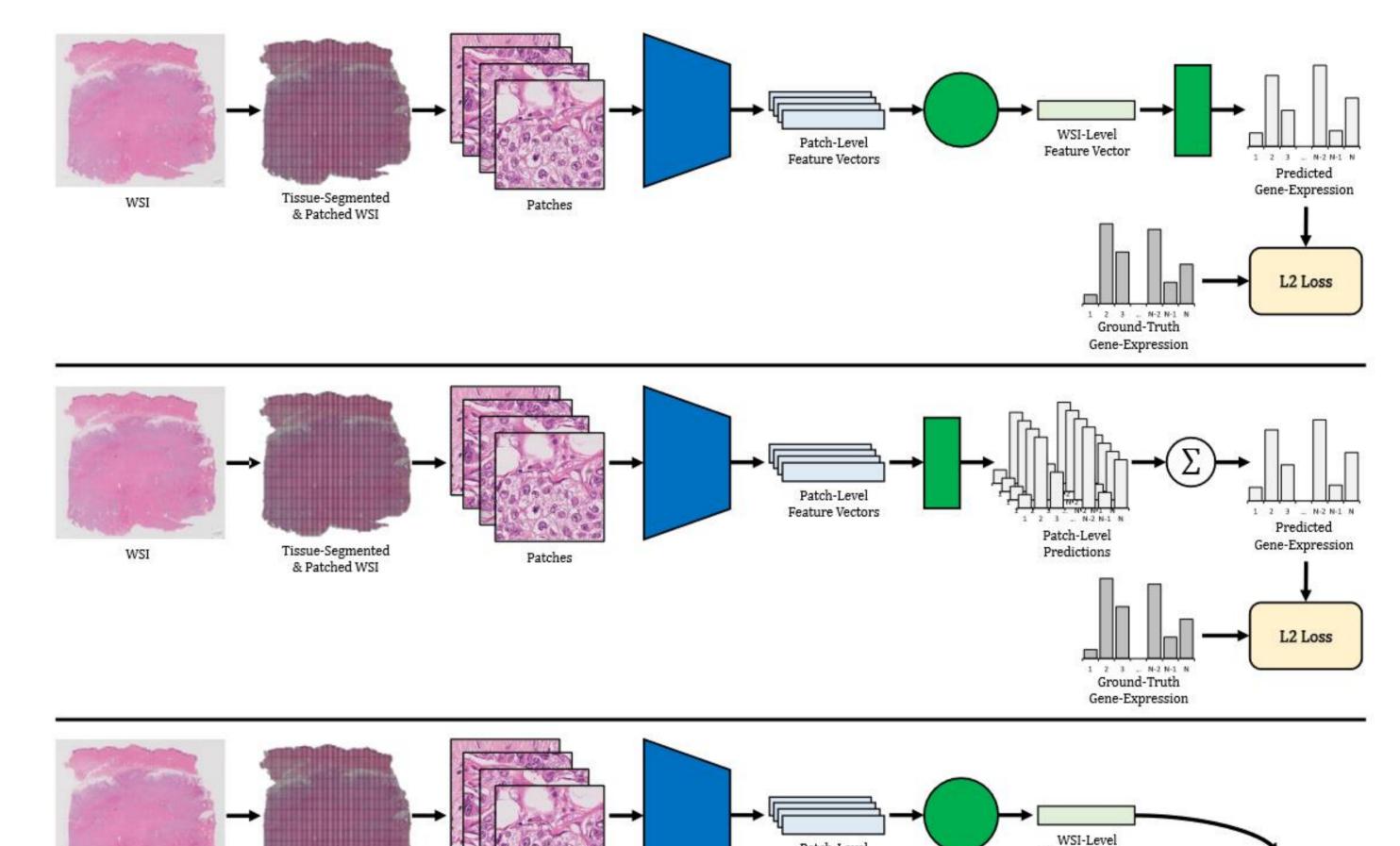
Patches

Introduction

Prediction of mRNA gene-expression profiles directly from routine WSIs using deep learning models could potentially offer cost-effective and widely accessible molecular phenotyping. While such WSI-based gene-expression prediction models recently have emerged, the highdimensional nature of the corresponding regression problem offers numerous design choices which remain to be analyzed in detail.

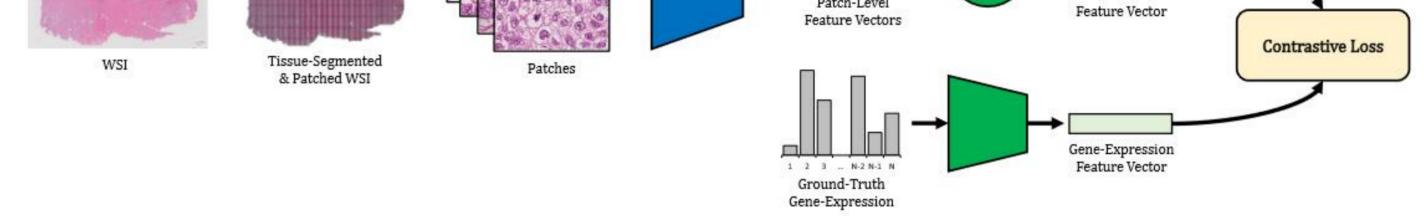
This study provides recommendations on how deep regression models should be trained for WSIbased gene-expression prediction.

Evaluated Models

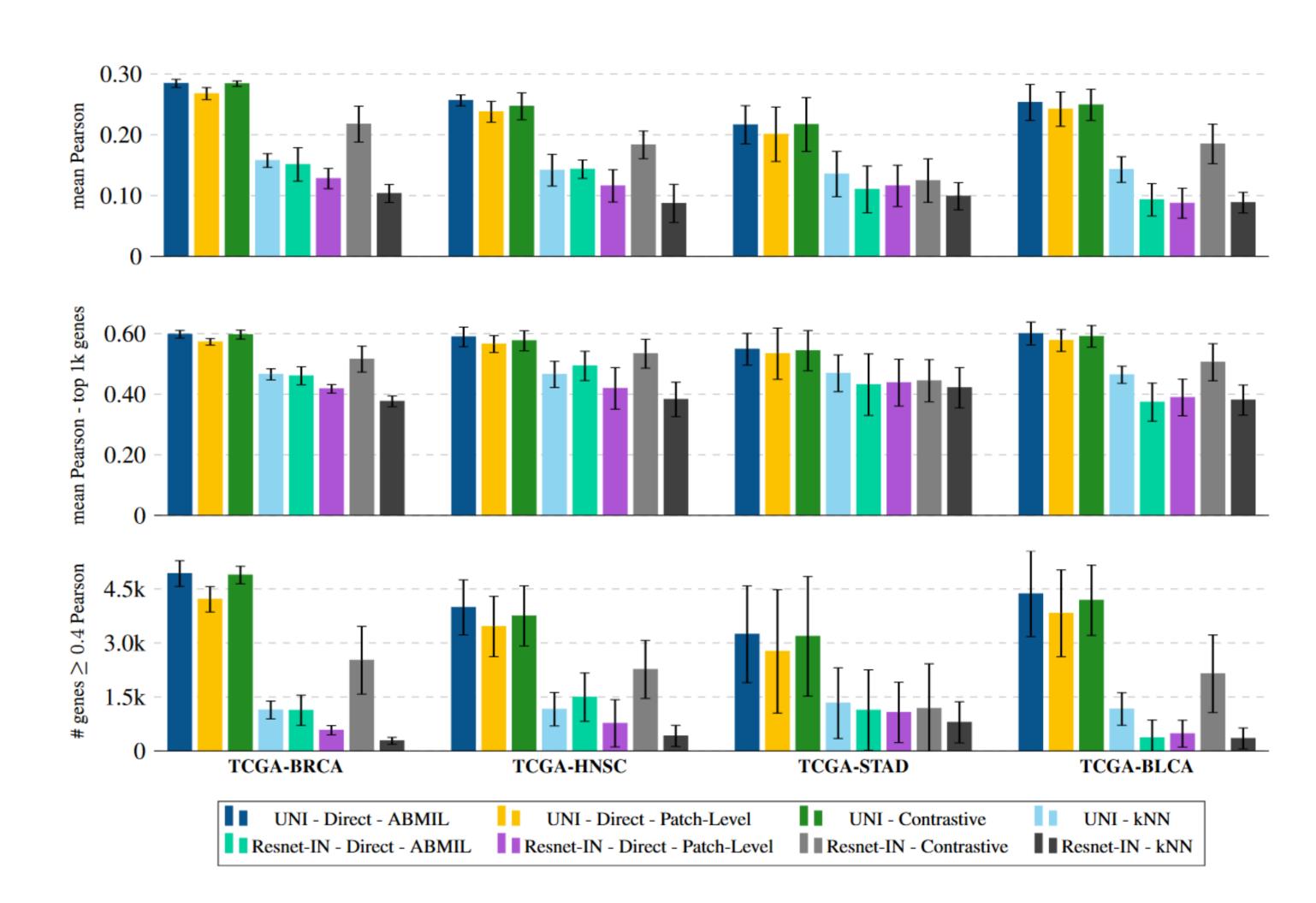


Main Takeaways

- Training regression models on top of UNI features gives accurate WSI-based models for gene-expression prediction (TCGA-BRCA: 4927 genes with Pearson corr. above 0.4, mean Pearson of 0.56 for PAM50 genes).
- 2. Despite conceptual differences, *Direct ABMIL* and Contrastive achieve very similar performance and should both be considered go-to regression models.
- 3. Training a single model to regress all 20530 genes is a computationally efficient and very strong baseline, this should be the starting point given any new dataset.



Top: Direct - ABMIL. Middle: Direct - Patch-Level. Bottom: Contrastive.



Results

4. Training one model for each individual gene incurs an extremely high computational cost yet achieves comparatively low accuracy.

Fredrik K. Gustafsson Karolinska Institutet www.fregu856.com



