xMINT: A Multimodal Integration Transformer for Xenium Gene Imputation

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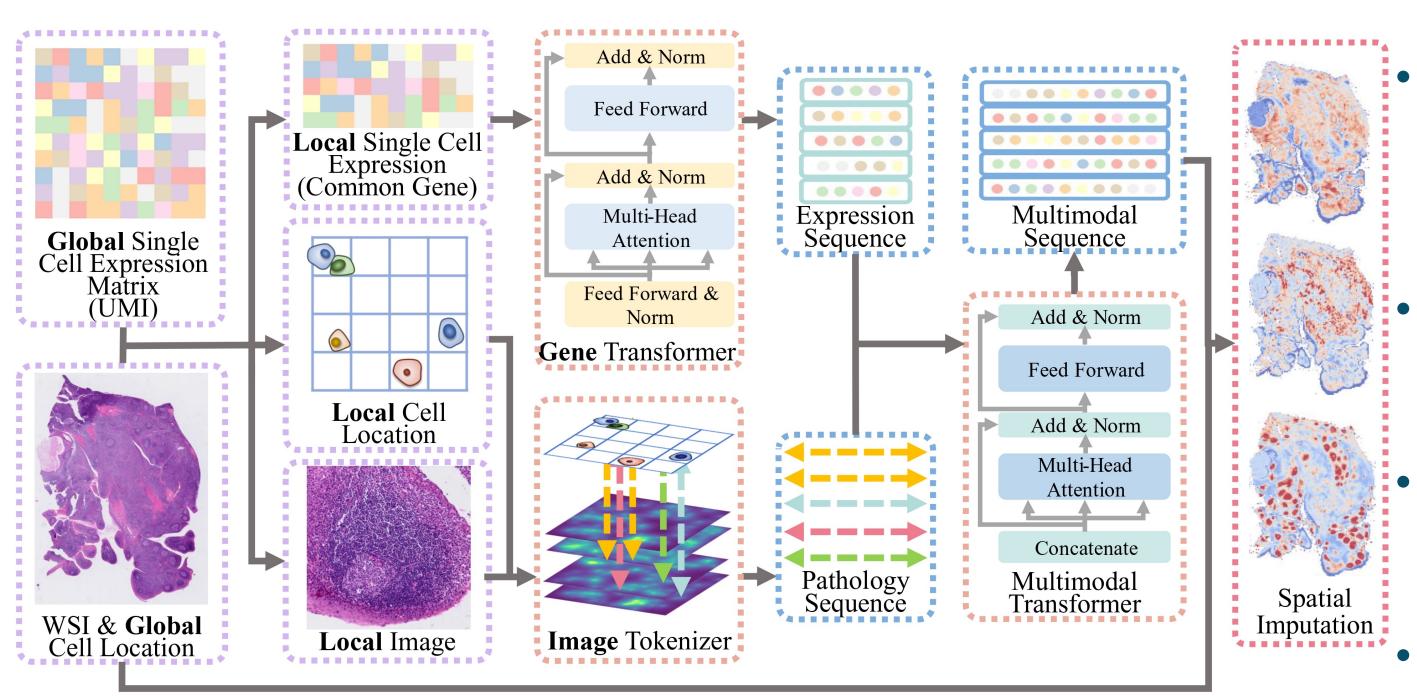
Background

- Gene Sequencing Limitations: Xenium technology sequences only around 500 genes, limiting comprehensive analysis.
- Technological Discrepancies: Integration of Xenium data with scRNA-seq is problematic due to differences in sequencing technologies.
- Data Underutilization: Current methods do not leverage available pathology images, which can provide critical insights into cell types and states.

Contribution

- Multimodal Integration: xMINT uses a Transformer-based model to integrate gene expression data with high-resolution pathology images, enhancing gene imputation accuracy.
- Enhanced Accuracy: Achieves better imputation accuracy than existing scRNAseq-based methods by utilizing additional visual information from pathology images.

Method



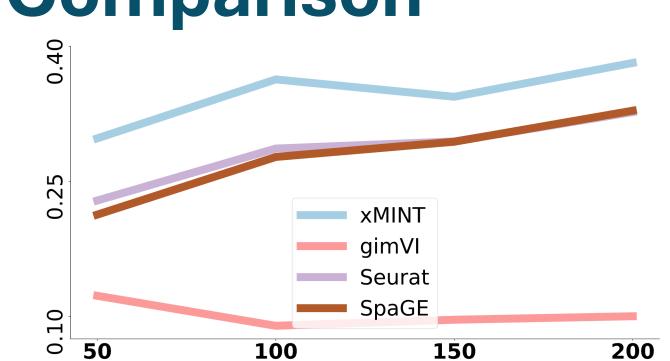
- **Initial Division**: Whole Slide Image is divided into local regions.
- Sequence Generation: Within each local region, sequences are generated using Morphology features & RNA sequencing values.
- **Concatenation**: The pathology sequence and expression sequence are concatenated.
- Multimodal Transformer: The concatenated sequences are input into a multimodal transformer.
- **Output:** The multimodal transformer outputs imputed gene sequences.

Parameter Number & Computation Time

f	Ехр.	Head	Layer	Parameters	Time (s/batch)
256	4	4	4	16,799,760	1.03
256	8	8	8	53,564,432	1.09
128	4	4	4	4,338,064	0.56
128	8	8	8	13,545,360	0.60

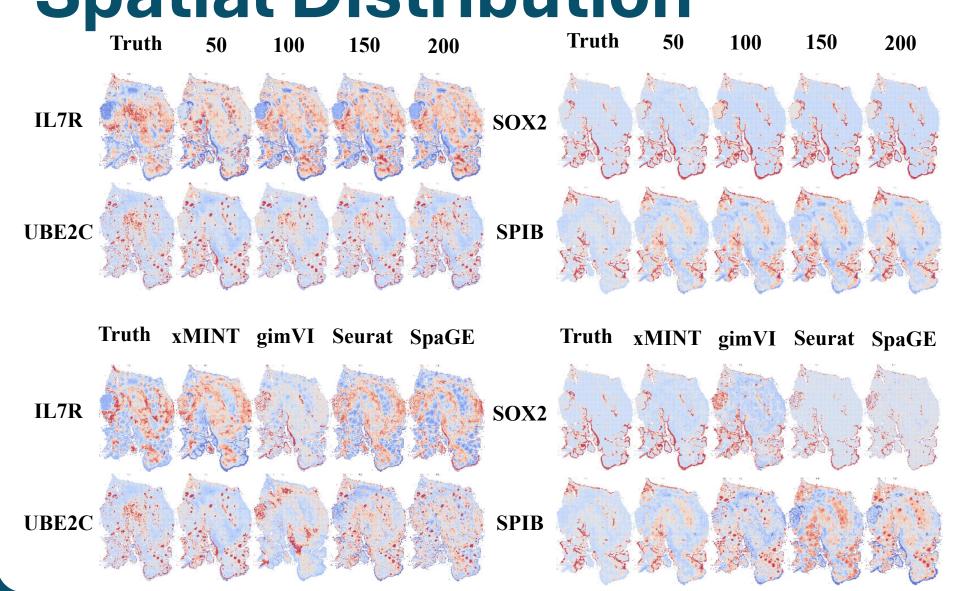
- **Exp.**: Forward expansion factor.
- **Head**: Number of attention heads.
- Layer: Number of transformer layers.
- f: Embedding dimension.

Comparison

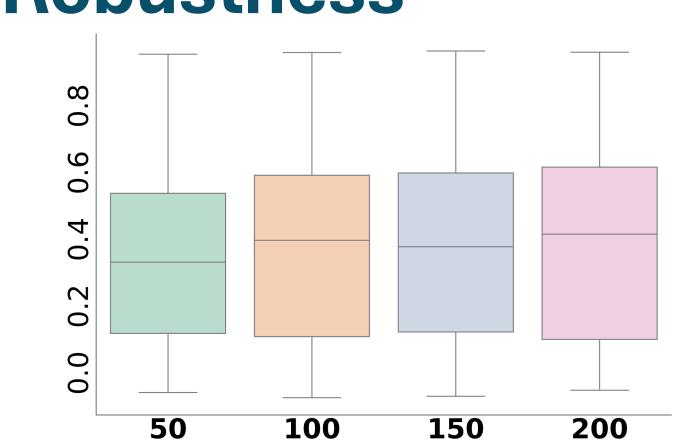


Median spearman correlation, with 50, 100, 150, 200 shared genes.

Spatial Distribution

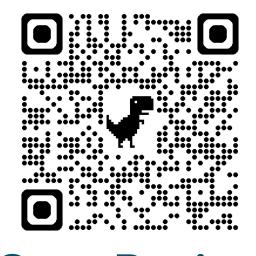


Robustness



Spearman correlation distribution, with 50, 100, 150, 200 shared genes.





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