# Explainable AI for Digital Histopathology



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### Motivation

Processing high resolution digital histopathology images with AI is difficult:

- Large images: The images are so large that conventional machine learning cannot be used.
- Lack of labels: Labelling these images is very expensive annotating individual cells is time consuming and requires a trained clinician.
- **Detail required**: If the images are down-sampled, their important details are lost, so accurate predictions cannot be made.

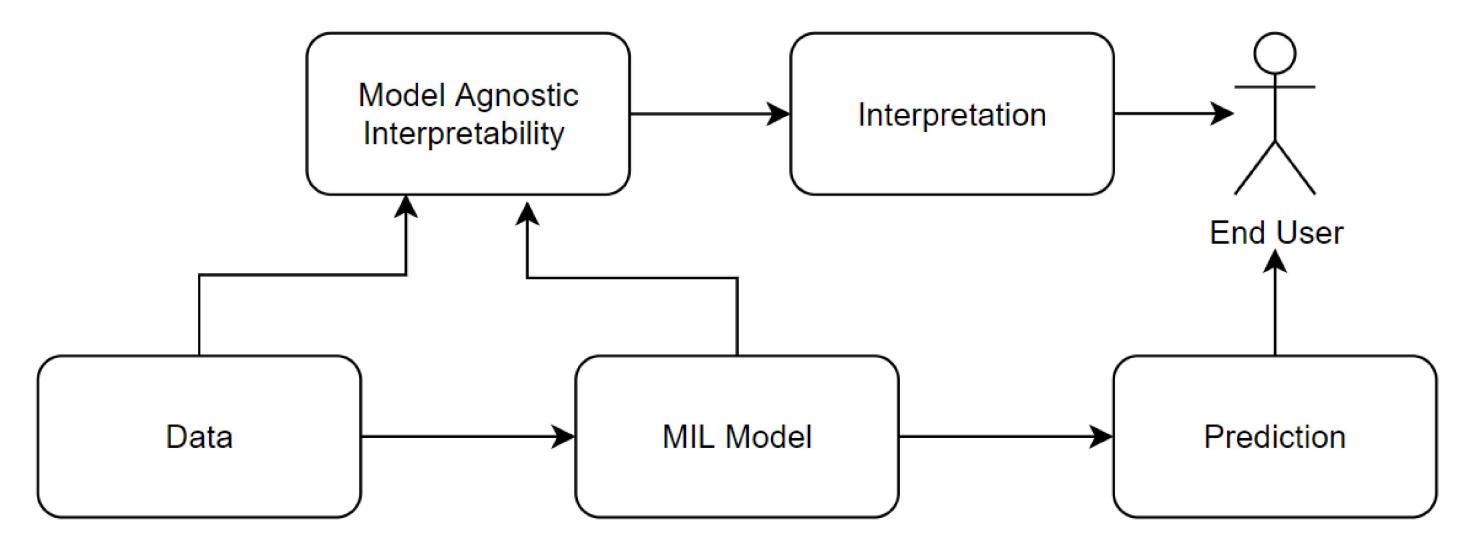
As such, we have to break the whole slide images into patches and use a technique known as **Multiple**Instance Learning (MIL).

#### Our work investigates:

- How MIL models make their decisions. Which patches are the important ones and what outcomes do they support?
- The role of explainable AI in digital histopathology. Can we understand how a classifier makes decisions?

## **Explainable AI for MIL**

Explainable AI allows both technical and non-technical users to better understand how an AI system makes decisions, which helps facilitate trust.



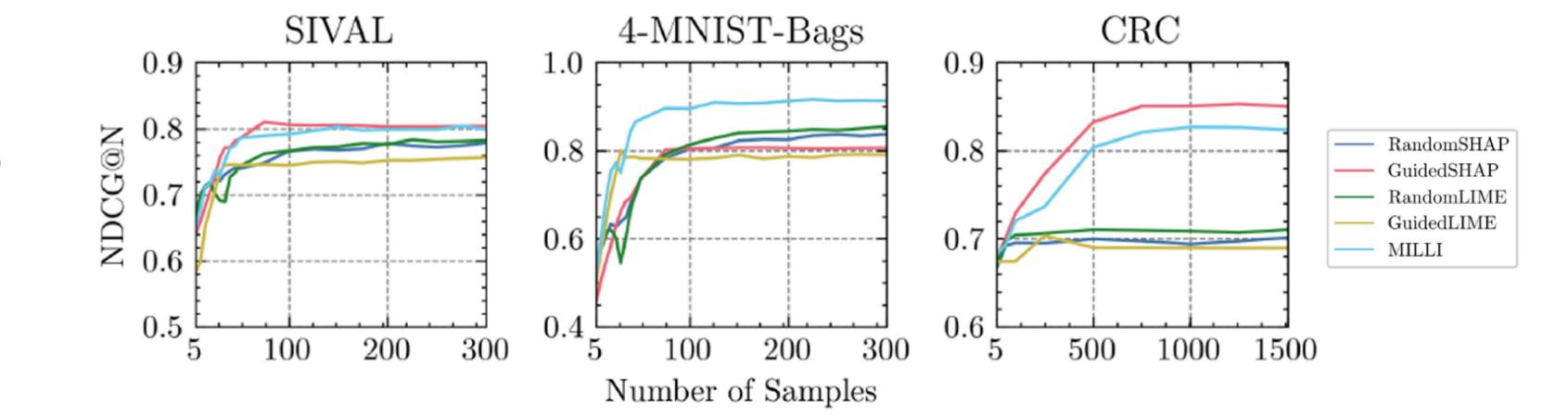
Our approach, Multiple Instance Learning Local Interpretations (MILLI), determines the importance of instances by analysing the effect of removing patches from the original image:

 Patch contribution: Different patches contain different data and will contribute to different outcomes. By removing patches and observing any changes in prediction, we can determine which outcomes they contribute to.

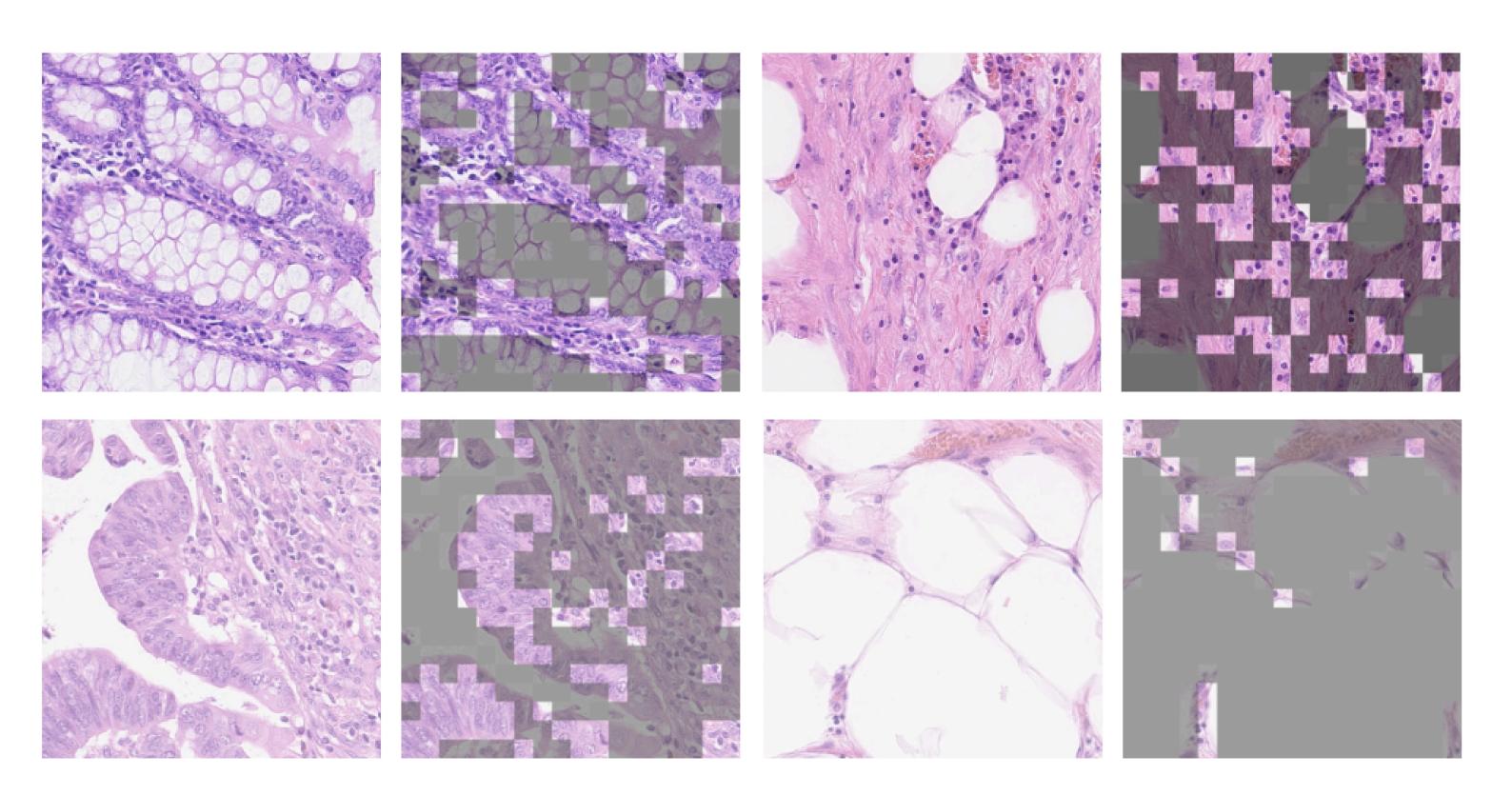




- Context aware: The patches cannot be considered in isolation, therefore we sample collections of patches to be removed together.
- **Optimisation**: It is expensive to determine the change in prediction for every possible combination of patches, therefore we **optimise our sampling** by first identifying the patches that are discriminatory and then biasing our sampling to choose more informative collections of patches.
- Improved efficiency: On average, compared to existing interpretability methods, MILLI is both more accurate and more efficient. It requires fewer samples to determine the contribution of patches, meaning it can generate accurate interpretations in a shorter amount of time.



## **Explainable Colon Cancer Detection**



We applied our methods to classifying tissue in colorectal cancer:

- Cell identification: Colorectal cancer originates from epithelial cells. Our approach means these cells can be found without requiring manual annotation, saving money and time.
  Furthermore, our approach also annotates other cell types as part of its explanations.
- **Better performance**: MILLI out-performed existing methods. Our interpretations were **30% more accurate** on average than existing approaches such as interpretable models (we found that these had poor general performance), or post-hoc methods such as LIME and SHAP.



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