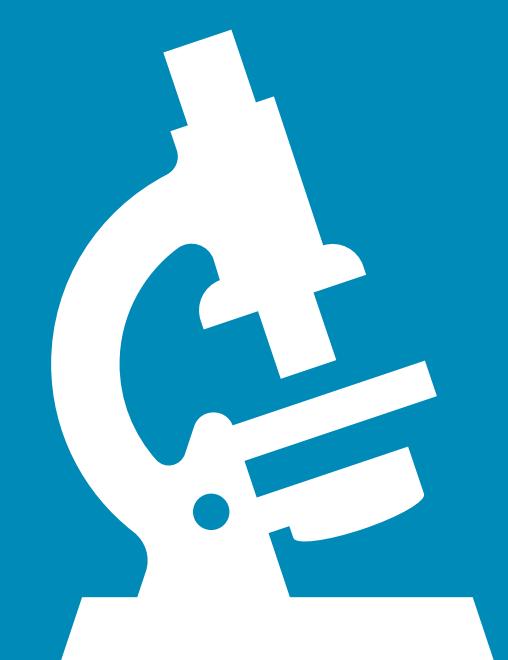


# Computer Vision Meets Microbiology:

## Deep Learning Algorithms for Classifying Cell Treatments in Microscopy Images



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### INTRODUCTION

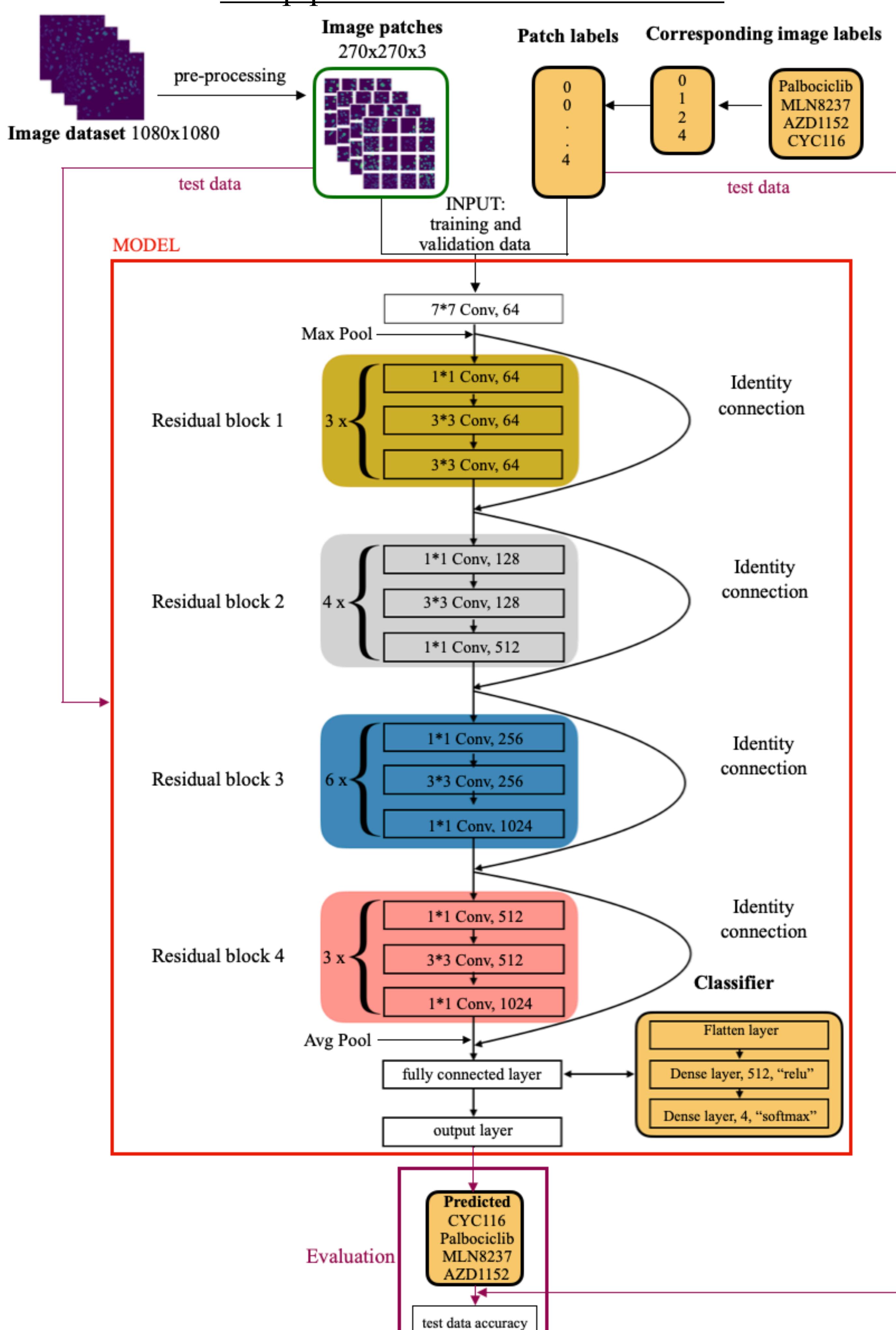
**Cell classification** is one of the most complex challenges in biomedical research that has significant importance to personalised medicine, cancer diagnostics and disease prevention.

**AIM:** to explore potential of deep learning to automate the classification of microscopy cell images into four cell treatments: Palbociclib, MLN8237, AZD1152, and CYC116.

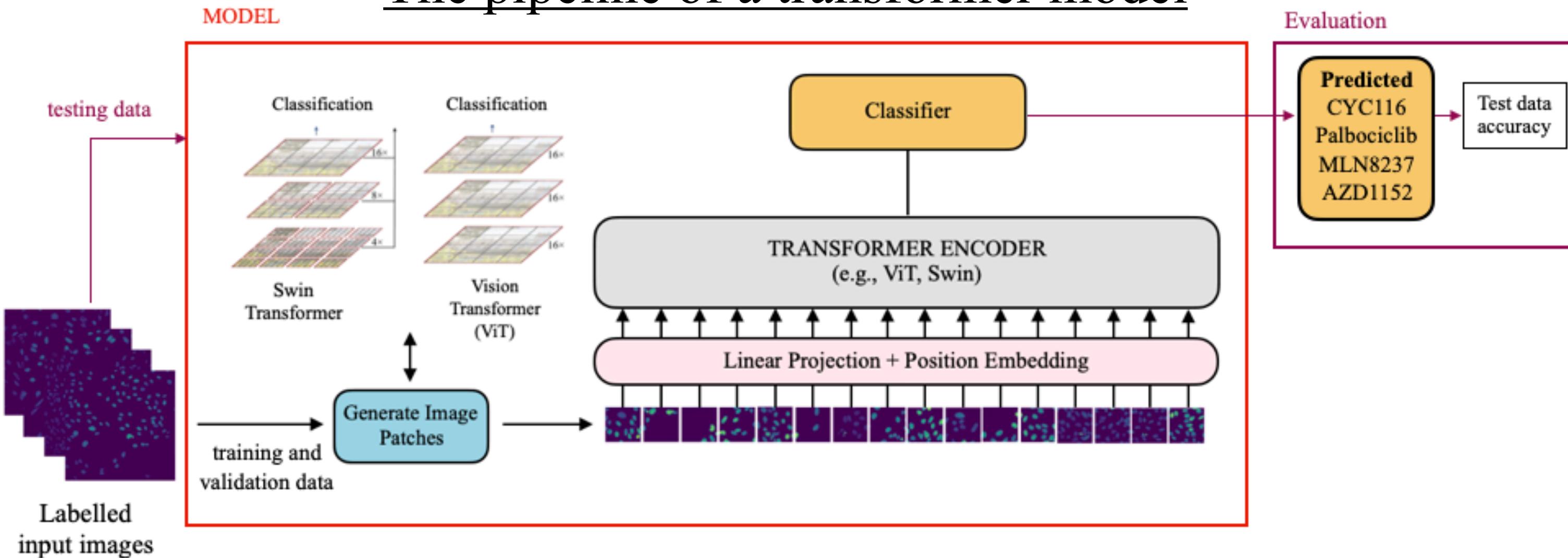
### METHODS

Three pre-trained state-of-the-art deep learning models, such as ResNet50, ViT and Swin Transformer were utilised to automatically classify bright-field and fluorescent microscopy images across single and multi channels.

#### The pipeline of the ResNet50 model



#### The pipeline of a transformer model



### DATASET

The dataset consists of 696 images. There are 4 cell treatment classes: Palbociclib contains 192 images; MLN8237, AZD1152 and CYC116 classes contain 168 images each.

Each image of the dataset consists of 7 channels: 5 fluorescence channels and 2 bright-field channels, examples are shown below:

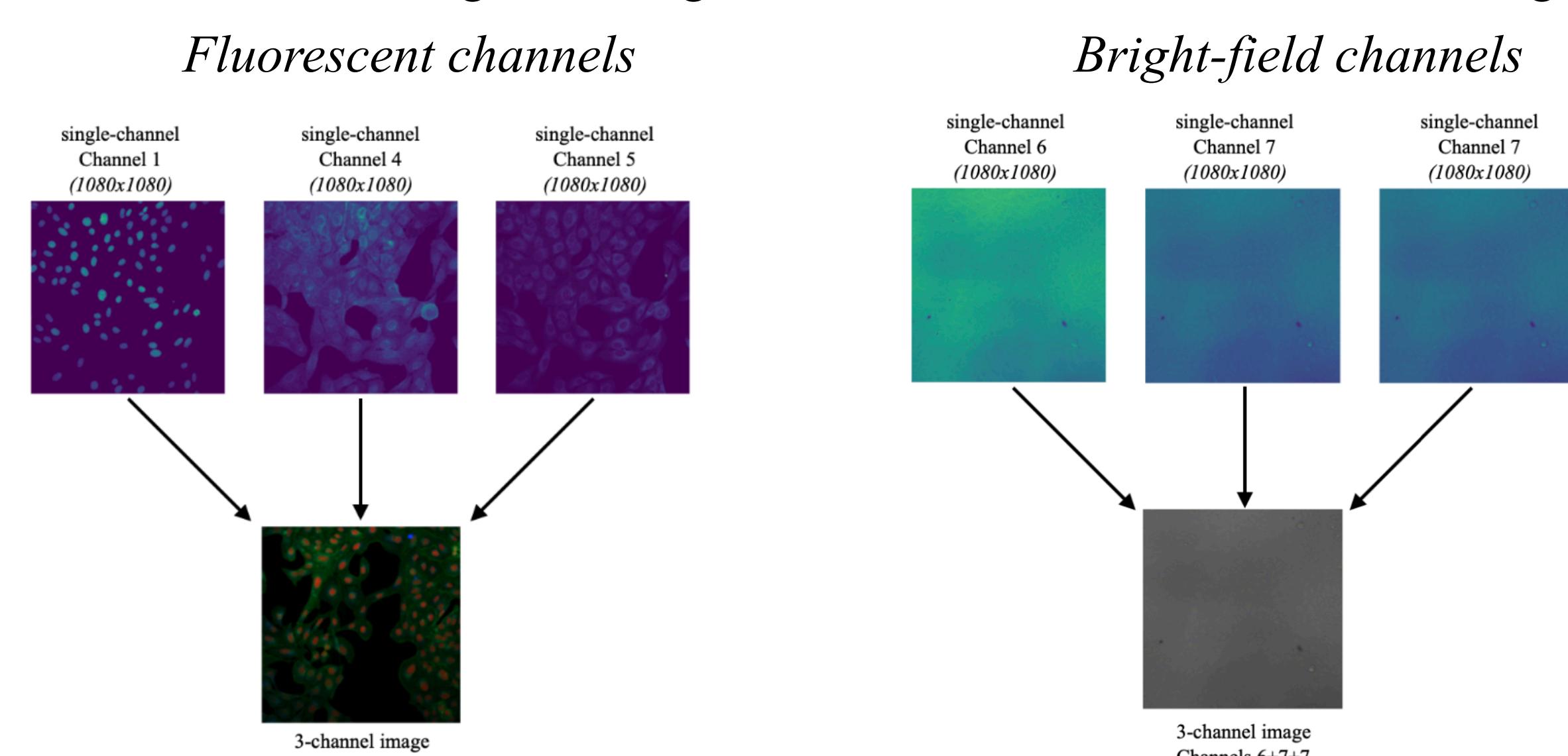
| Cell treatments | Channels |      |      |      |      |      |      |
|-----------------|----------|------|------|------|------|------|------|
|                 | Ch 1     | Ch 2 | Ch 3 | Ch 4 | Ch 5 | Ch 6 | Ch 7 |
| Palbociclib     |          |      |      |      |      |      |      |
| MLN8237         |          |      |      |      |      |      |      |
| AZD1152         |          |      |      |      |      |      |      |
| CYC116          |          |      |      |      |      |      |      |

### EXPERIMENTAL RESULTS

Approach 1: Apply DL models to classify cell treatments on single-channels.

Approach 2: Apply DL models to classify cell treatments on three-channels.

Process of combining three single-channels into one three-channel image



Approach 3: Apply DL models to classify cell treatments on many-channels (> 3)

The results of 1st and 2nd experimental approaches, since 3rd approach is on training

| Channel               | Fluorescent                       |      |      |      |      | Brightfield |      |      |             |
|-----------------------|-----------------------------------|------|------|------|------|-------------|------|------|-------------|
|                       | 1                                 | 2    | 3    | 4    | 5    | 1+4+5       | 6    | 7    | 6+7+7       |
| Experimental approach | 1st                               |      |      |      |      |             |      |      | 2nd         |
| Model                 | Accuracy on test image data, in % |      |      |      |      |             |      |      |             |
| ResNet                | 80.0                              | 75.0 | 69.0 | 77.0 | 77.0 | 84.0        | 59.0 | 61.0 | 52.0        |
| ViT                   | 78.0                              | 34.0 | 27.0 | 39.0 | 27.0 | 38.0        | 27.0 | 27.0 | 27.0        |
| Swin                  | 79.0                              | 48.0 | 26.0 | 44.0 | 35.0 | <b>86.0</b> | 27.0 | 27.0 | <b>59.0</b> |

### CONCLUSION & FUTURE WORK

- The highest accuracy achieved on 3-channel fluorescent images was **86%** by Swin Transformer.
- The highest accuracy achieved on 3-channel bright-field images was **59%** by Swin Transformer. While ResNet has achieved **61%** accuracy on 1-channel bright-field images.
- This necessitates further exploration of DL models for classification of single- and multi-channel bright-field microscopy images.