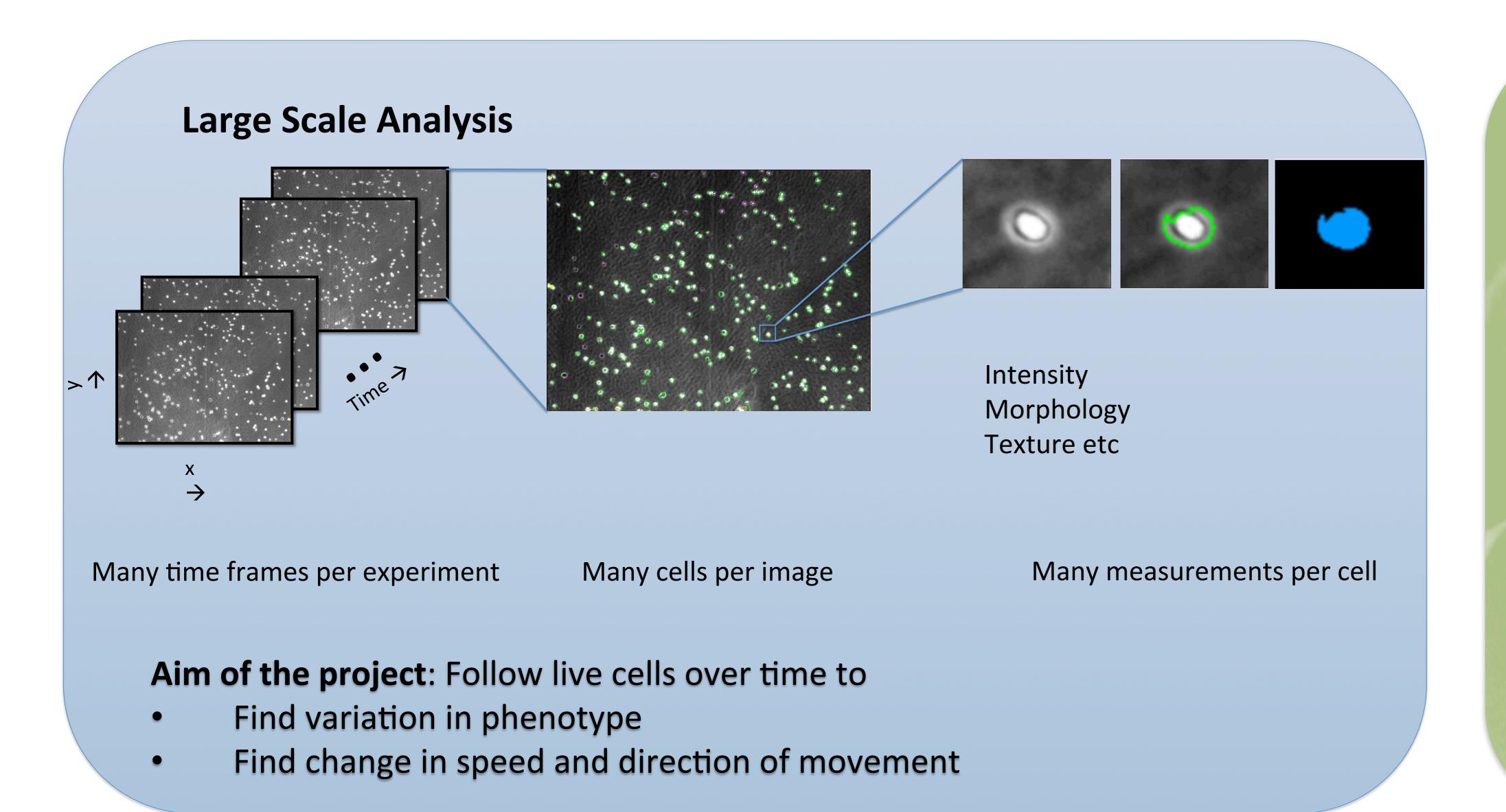
# Large-Scale Analysis of Live Cells

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#### Introduction

Obtaining quantitative data from live cells is of great importance for understanding the cellular and molecular processes. A recurring task in many experiments is the tracking of large number of cells and the analysis of their spatiotemporal behavior. Live cell experiments using automated imaging systems generate a huge amount of data that is difficult for a human observer to analyze.

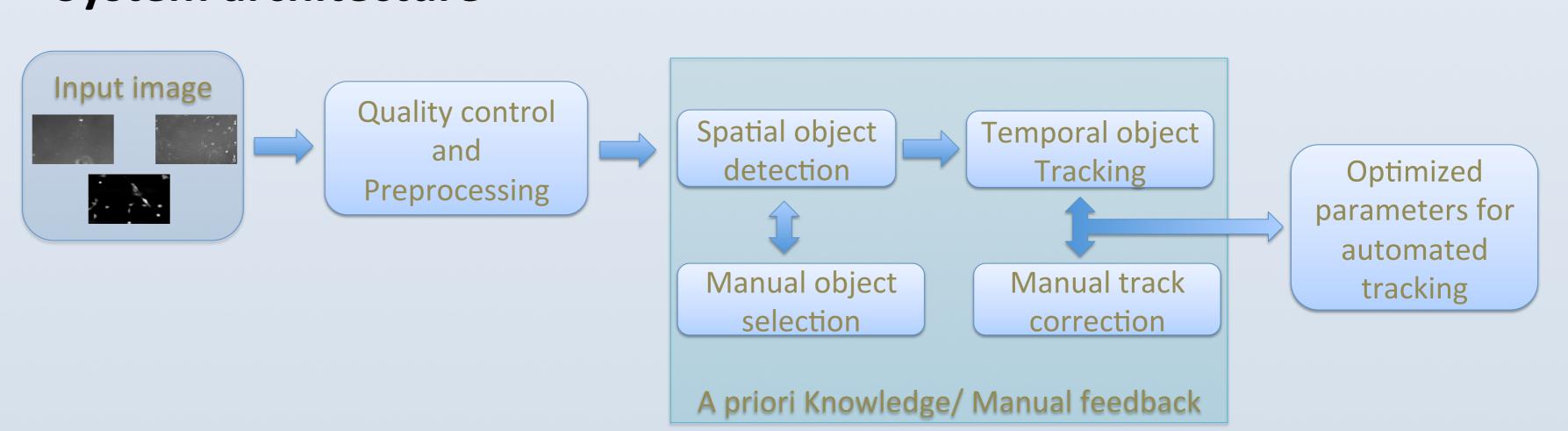
#### Issues:

- The investigators suffer from lack of userfriendly software and infrastructure in terms of computing and data storage.
- Need to know complex image analysis algorithms for application based parameter tuning which prohibits from routine usage.

### Novelty:

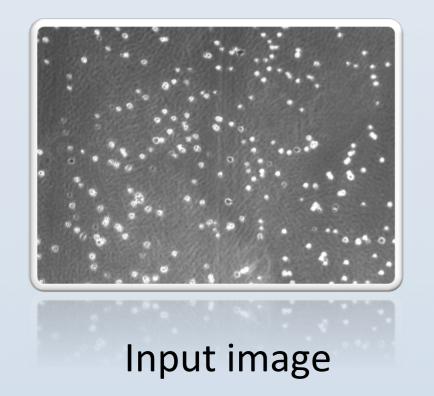
- Use the investigator's biological knowledge as feed-back to the system in the form of manual selection/training detection of objects of interest and manual editing of erroneous tracks
- Flexible automated high throughput system
- Reduce the time and algorithmic insight required by the investigator

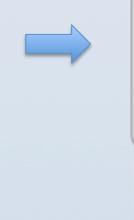
# System architecture

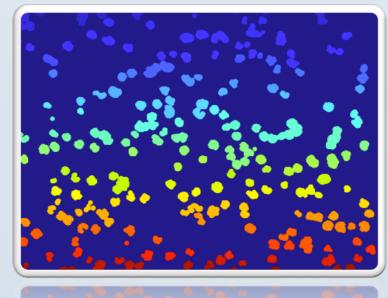


**Segmentation of neutrophils:** The time lapse microscopy images of neutrophils revealed that they are comparatively fast moving cells. This often causes them to come closer to each other, making it difficult to distinguish between two near by cells. For the tracking to be successful, it is important to identify all the cells present in the image. For this experiment local maxima is found at the scale of the cells and set as marker for segmentation. Thus close by cells were separated.

## **Initial results**







Initial segmentation

Segmentation of near by

cells





Markers detected on cells

Final segmentation result







