

# General structure of experiments for comfortable laboratory automation

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## Introduction

For large-scale laboratory automation, dynamic experiment management software that adapts to various types of experiments is essential. However, most automation software is designed for specific research or experimental methods(1-3). In addition, no experimental structure exists for the purpose. To address this, **we define a flexible experimental structure** capable of representing dynamic progress and develop the General Experiment Management Software (**GEMS**) to manage it.

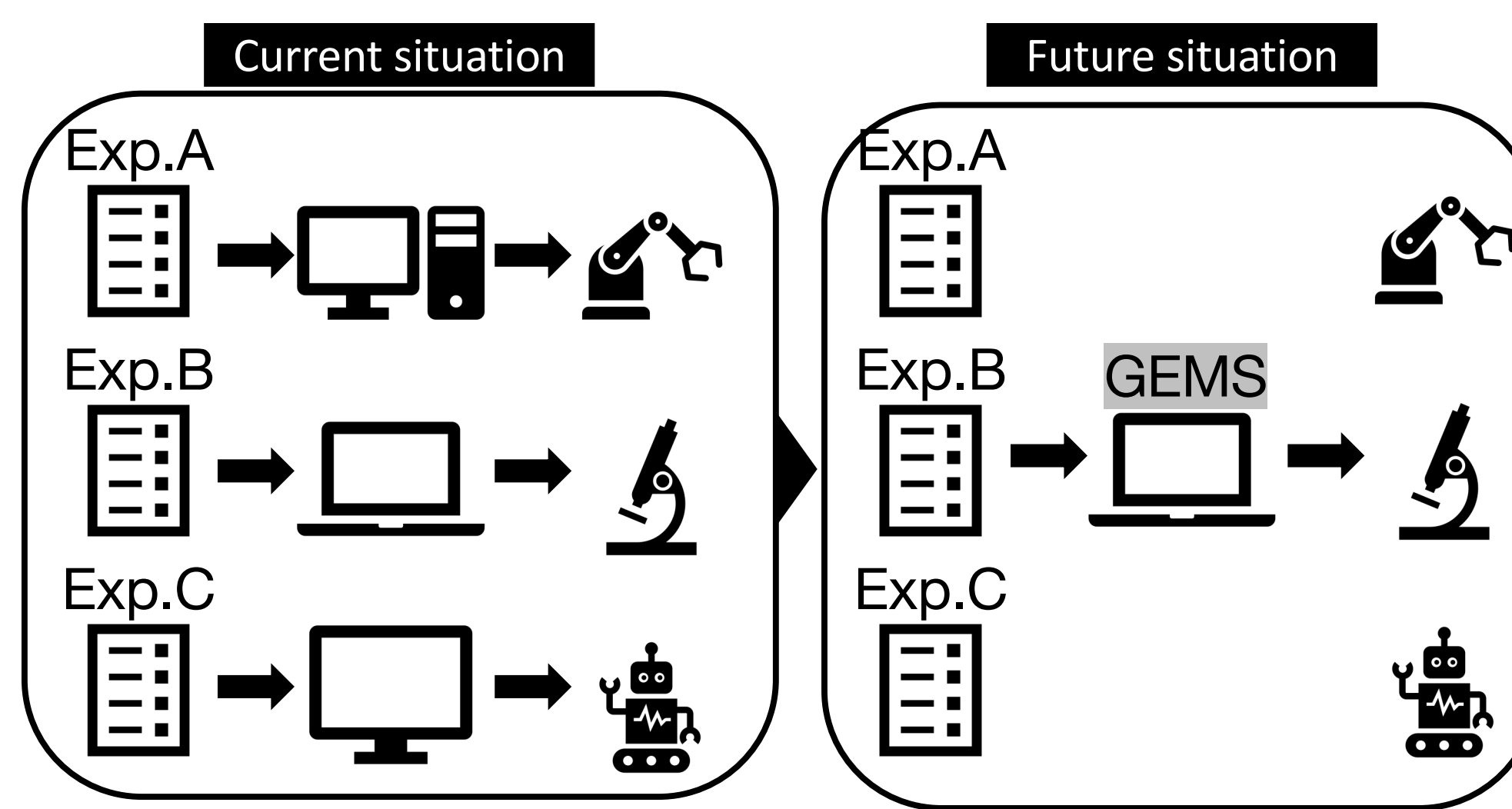


Fig.1 Current and future situation

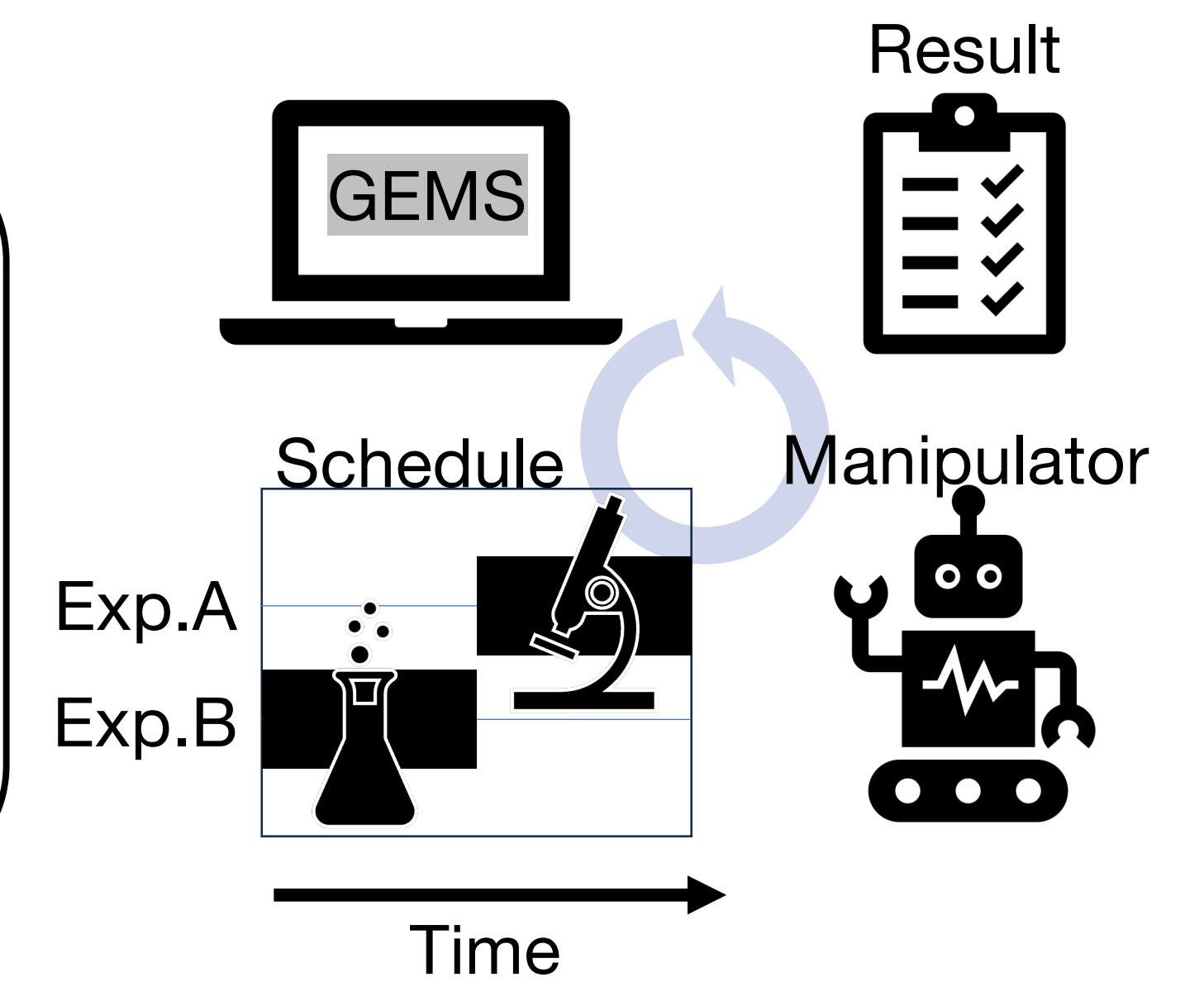


Fig.2 Autonomous experiments with GEMS.

## Methods

### General structure of an experiment

- Experiments consist of multiple states.
- Specific tasks, and a formula for calculating the optimal execution time, are defined for each state.
- Transition rules between states are predefined for each state.

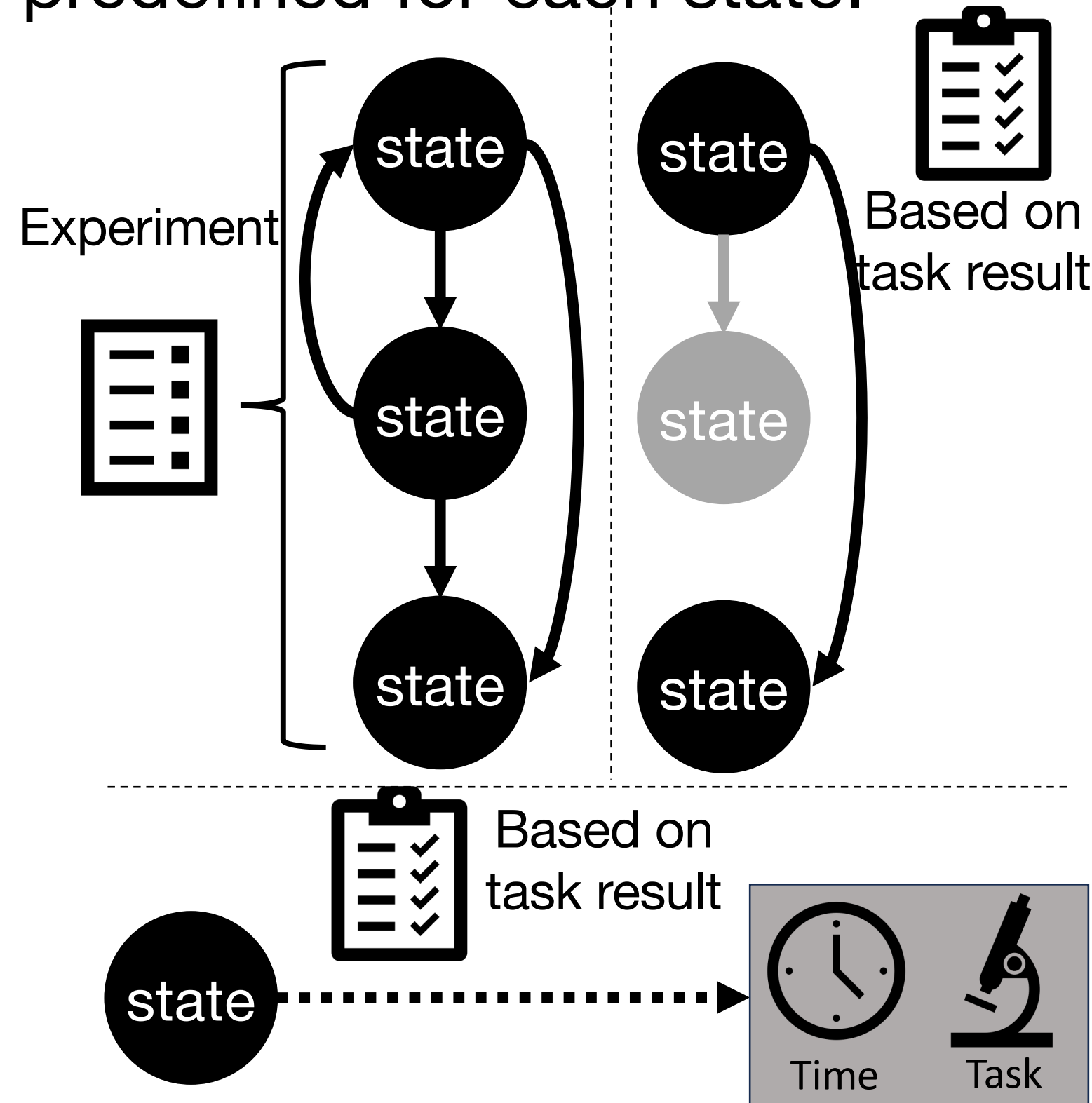


Fig.3 General structure of an experiment

### Software configuration

- Transition Manager updates the state of the experiment(s).
- Task Generator calculates the optimal time of the next task.
- Task Scheduler schedules all the tasks.

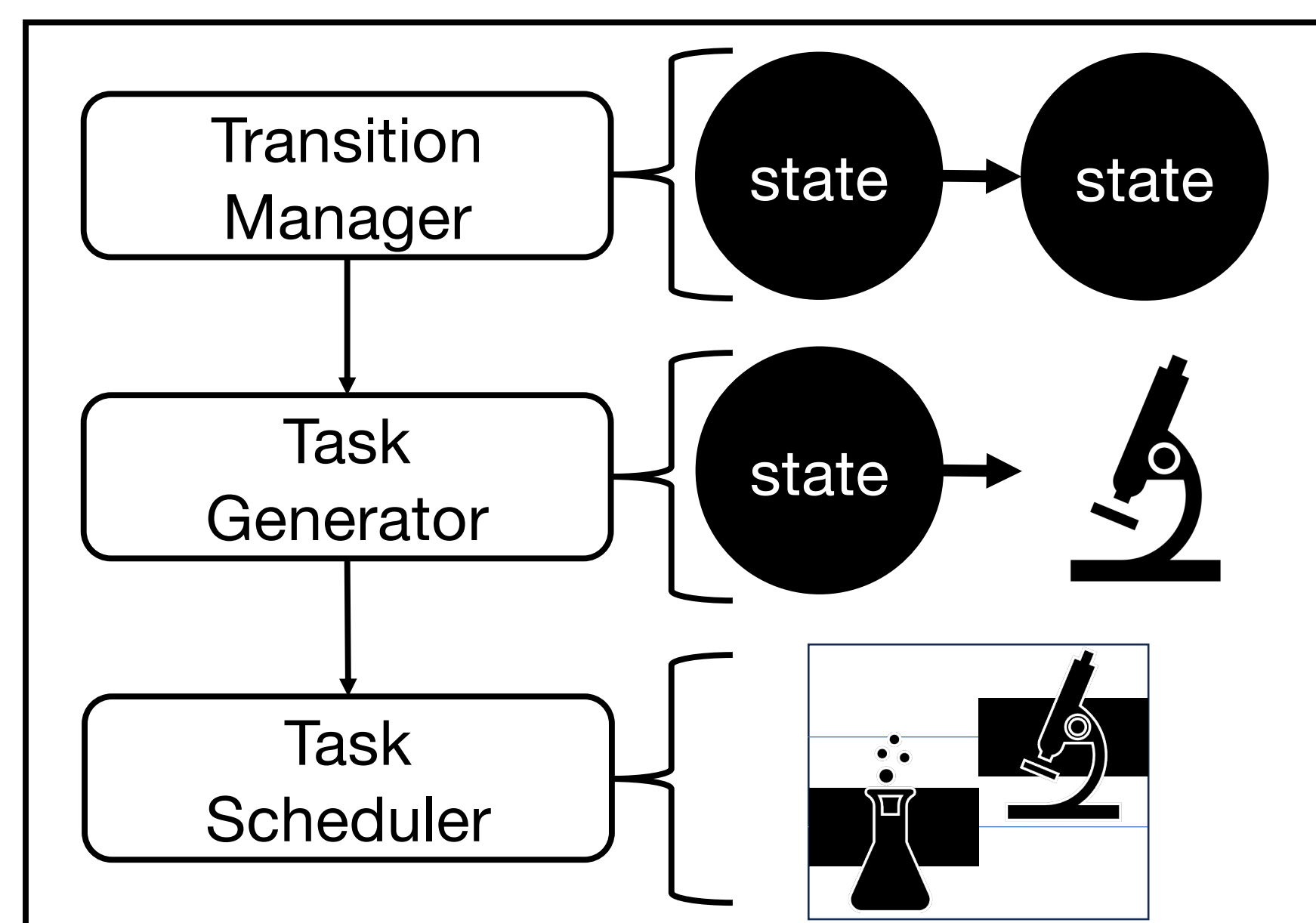


Fig.4 GEMS configuration

### Settings of simulation experiment

[Maintenance of multiple cell lineages]

- Each cell line has a unique growth curve.
- All cell lines start in the same initial state.
- The optimal passage time is when the cell count reaches 0.3 and 0.7 for iPS cells and HEK293A cells, respectively.
- Regular task: restocking reagents.
- Priority: passages of iPS cells over HEK293A cells.

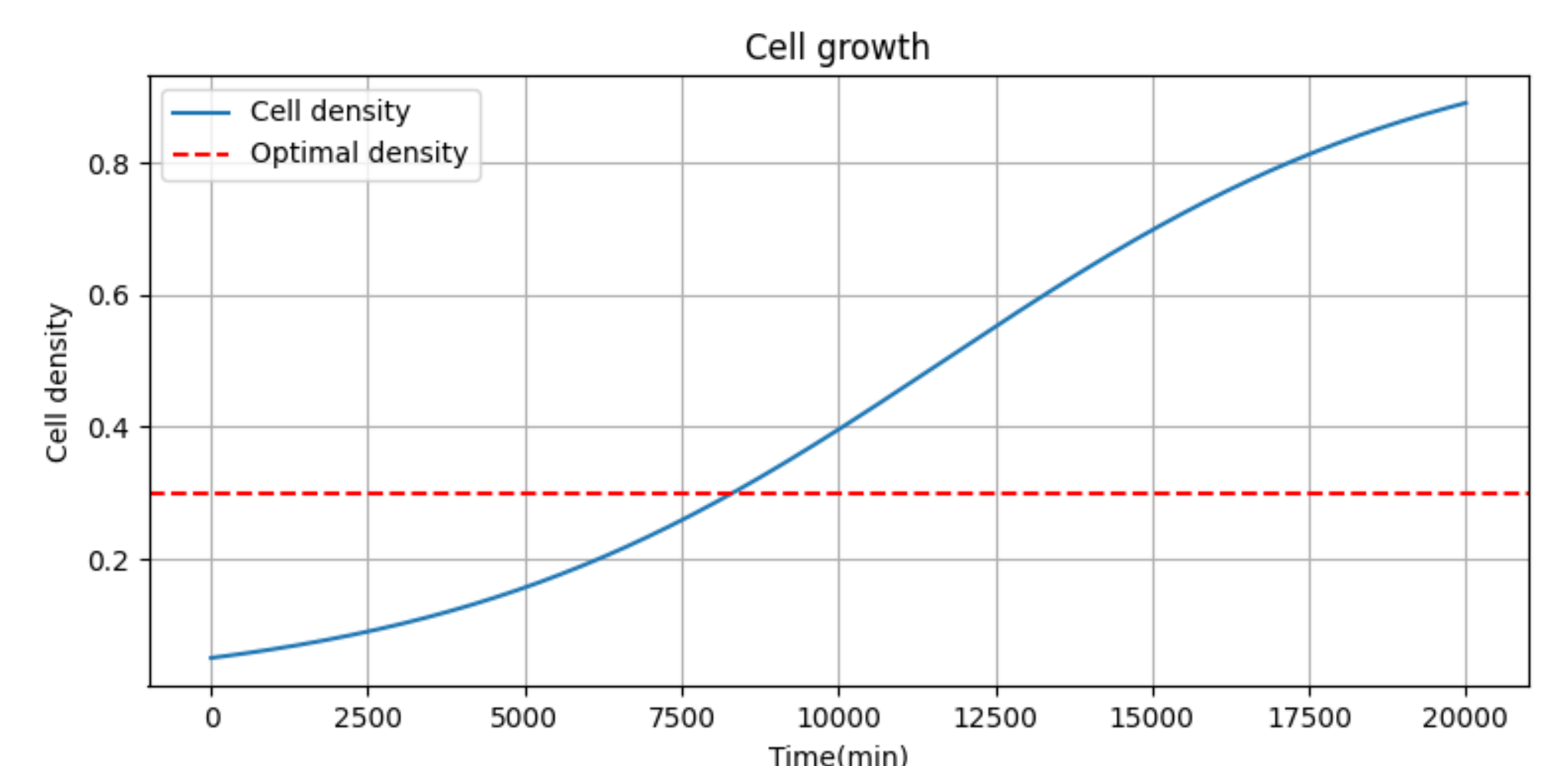


Fig.5 Cell proliferation and optimal passage timing

## Results

### Simulation result

- Maximum and minimum densities of iPS cells: 0.32 and 0.29.
- Maximum and minimum densities of HEK293A cells: 0.86 and 0.57.
- Culture timing of iPS cells maintained within  $\pm 0.02$  of target density.
- GEMS could consider the priority successfully.
- GEMS proceeded with the experimental group with reagent restocking without affecting passages.

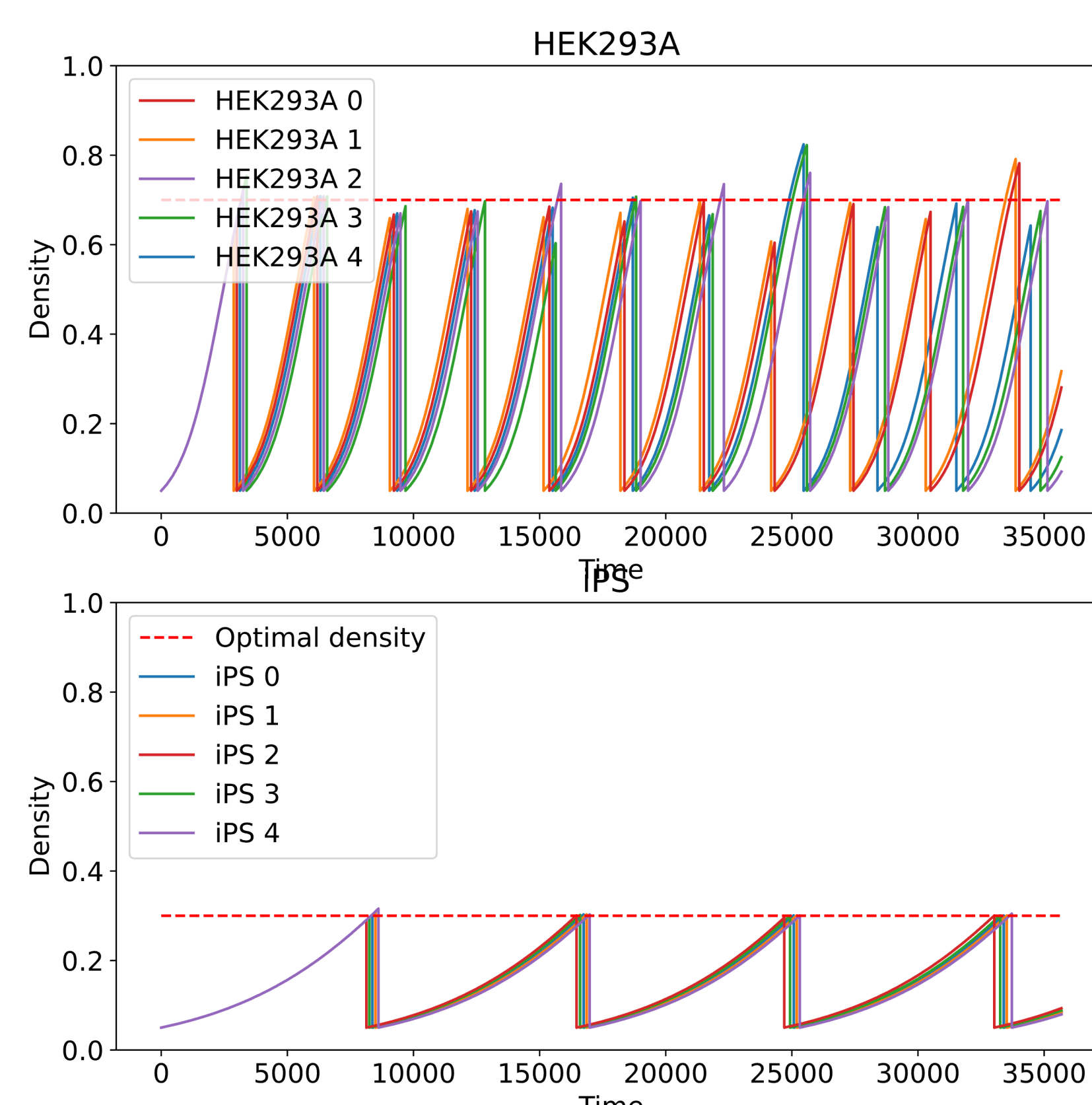
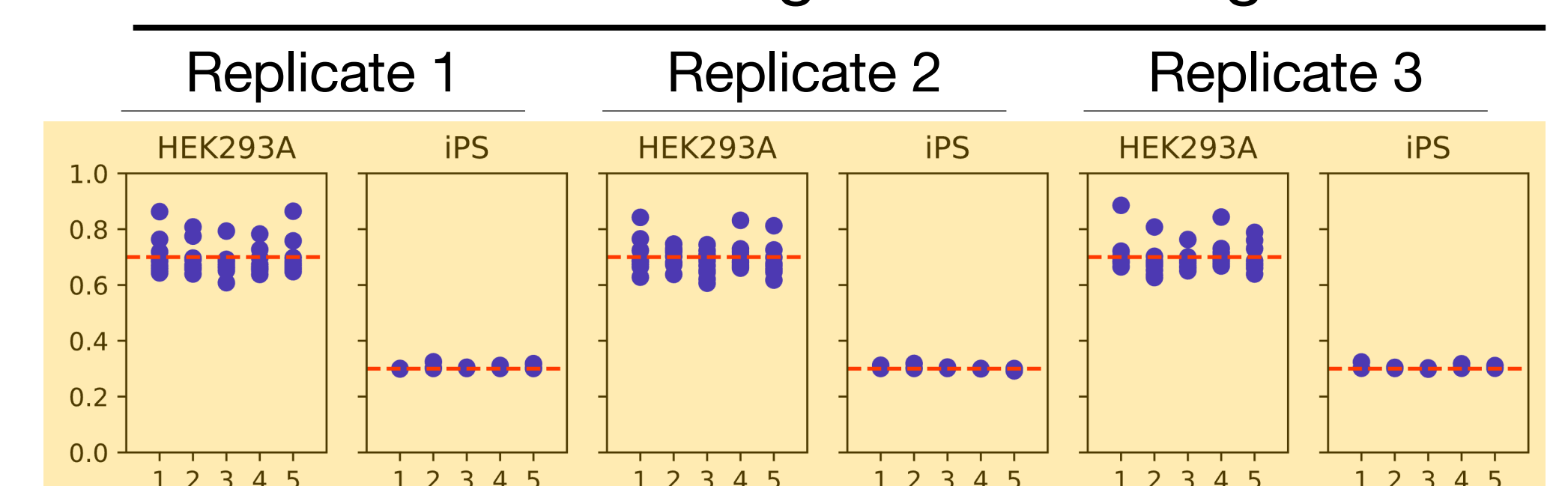
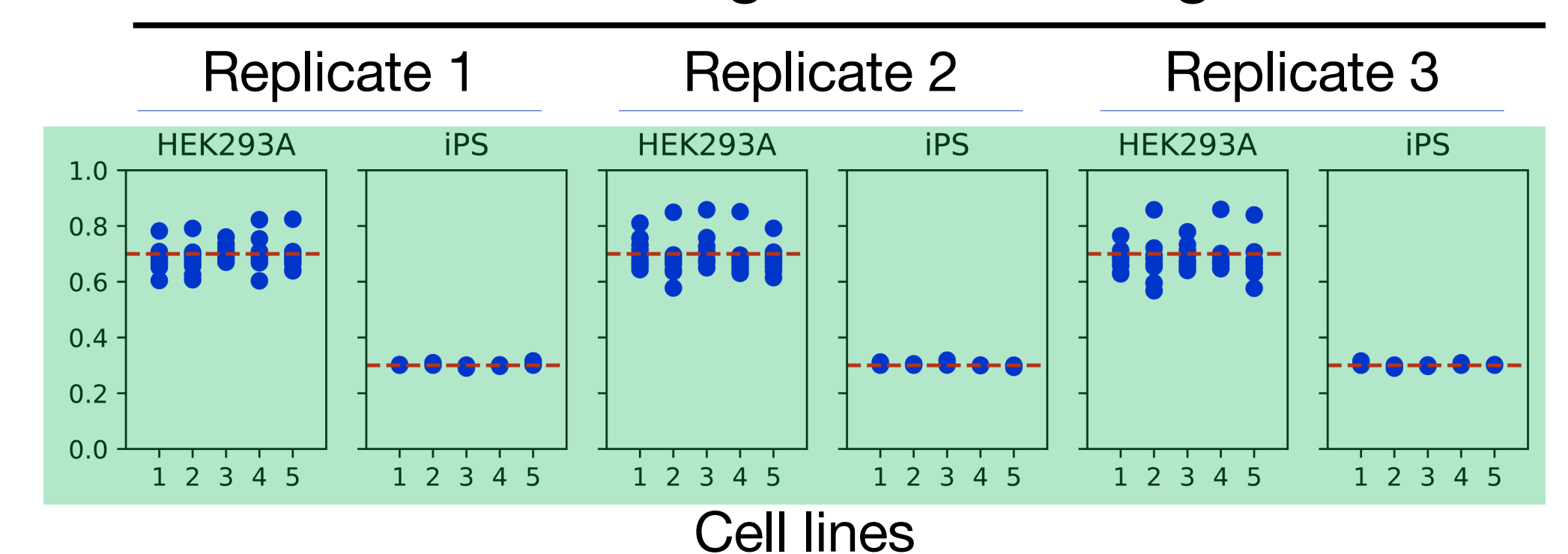


Fig.6: Simulation results - Cell proliferation (left), Passage densities (right)

### Without reagent restocking



### With reagent restocking



## Future plan

### 1. Simulate other experiment

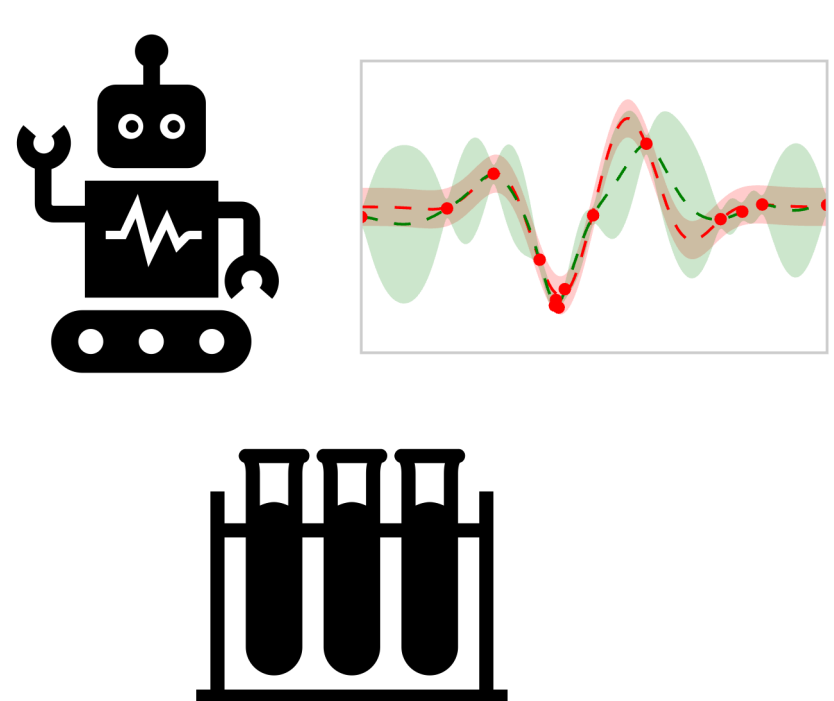


Fig.7 Colour water optimisation

### 2. Real-world experiment

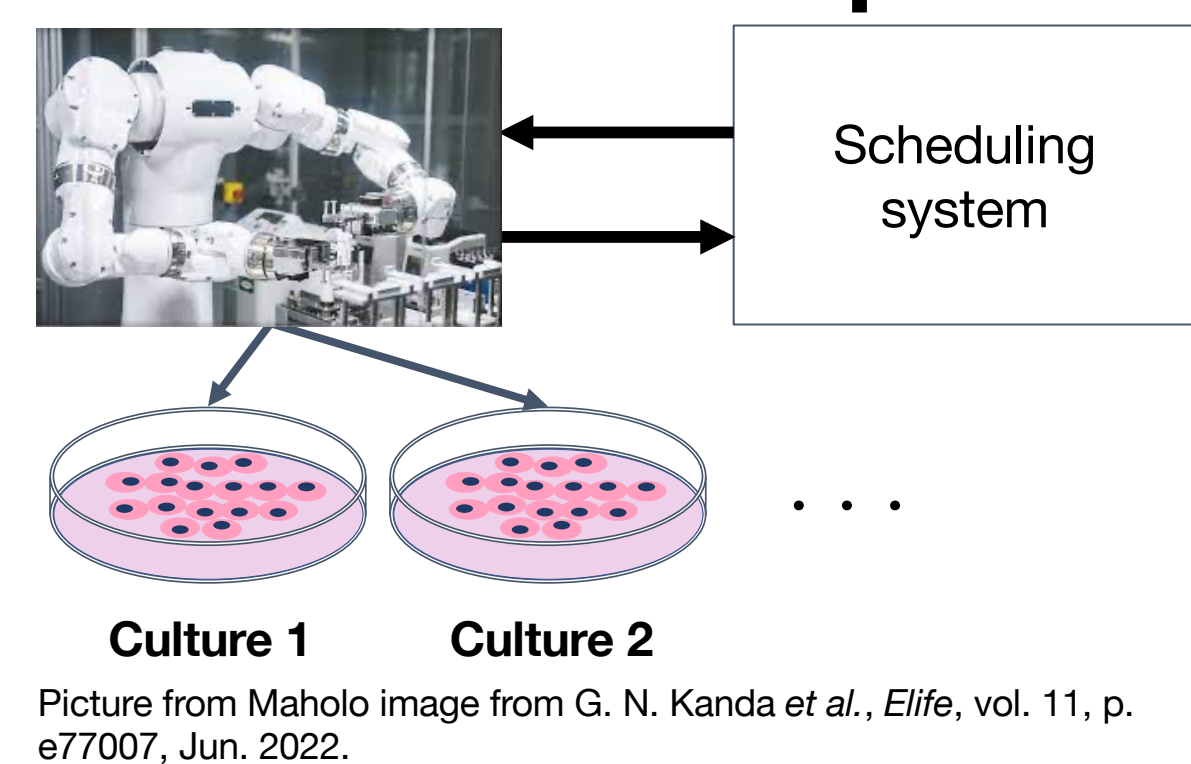


Fig.8 Real-world experiment Image

### 3. Extension of GEMS

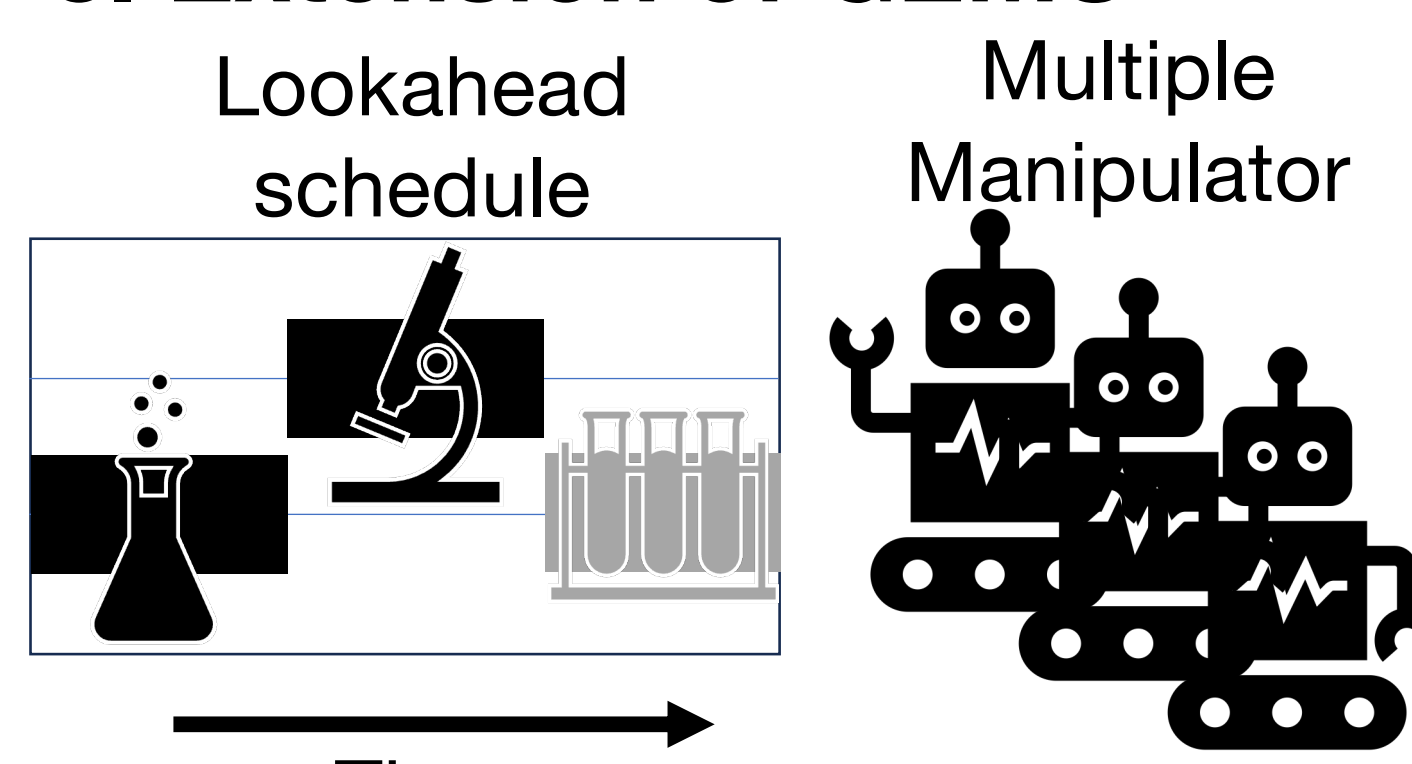


Fig.9 Extended experimental structure

## References

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- 2) I. Namatame et al., *SLAS Technol.*, Apr. 2023, doi: 10.1016/j.slast.2023.04.002.
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