

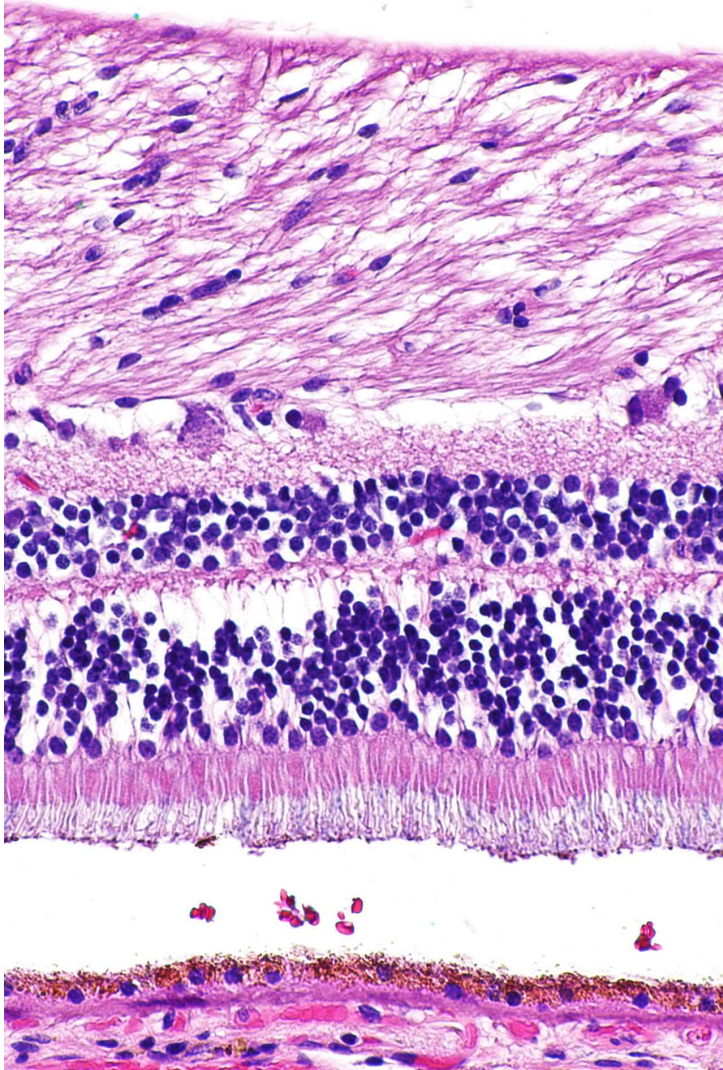
A Process Simulation Model for a Histopathology Laboratory

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Introduction



- Histopathology = microscopic examination of (stained) tissue samples for study & diagnosis of disease
- Especially critical for cancer screening
- Yet:
 - NHS England, 2022/3 – only **70.2%** of cancer diagnoses delivered within **4** weeks
 - Compare with guideline: **90%** within **10** calendar days (Royal College of Pathologists, 2013)
- [Brown 2004](#): **80%** of lab time is non-value-adding
 - (Case study of histopathology lab in Leicester, UK)

Process Simulation

- Identify bottlenecks in the process
- Estimate key performance indicators (KPIs)
 - Mean turnaround time (TAT) (reception to diagnostic report)
 - % specimens with TAT < n days
 - Mean lab TAT (excludes reporting stage)
 - % specimens with Lab TAT < n days
- Scenario analysis
 - E.g., staffing levels / shift scheduling



Outline

1. Overview of modelling process
2. Simulation model features
3. Results and scenario analyses
4. Discussion



Process modeling

- What are the steps? What order? How long do they take?
- Obtain data from:
 - Workshops
 - Lab observations (shadowing)
 - Review of Standard Operating Procedure (SOP) documents
- For simplicity:
 - Focus on core processes
 - Focus on standard stains (hematoxylin and eosin / H&E)



Process Modelling (2)

- Object Hierarchy
 - Specimen
 - Block
 - Slide
- Tasks operate on any of the above, or batches of these
- Specimens collated before transfer between stages
 - Easier tracking

Stages:

1. Specimen reception
2. Cut-up
3. Processing
 1. Decalcification (sometimes)
 2. Processing machine
 3. Embed & trim
4. Microtomy
5. Staining
6. Labelling
7. Scanning
8. Block/quality check
9. *Reporting*

Lab turnaround time

Total turnaround time



Simulation Model

- Discrete-event simulation model
 - Implemented in Arena
 - Entities created, worked on, disposed by navigating a flowchart
 - Flowchart blocks include Seize/Delay/Release/Batch/Set Attributes/etc.
 - Input/Output blocks for connecting to Excel spreadsheets
 - Model parameters
 - Simulation results (KPIs)
 - Plots for work-in-progress counters (overall / per-stage)



Selected Model features



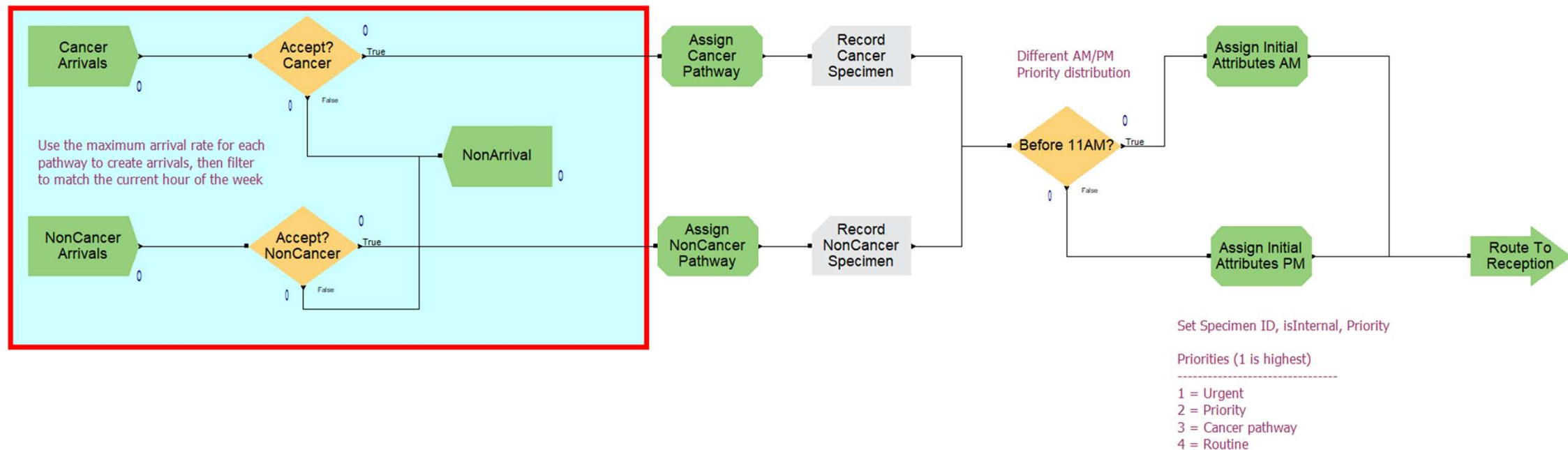
Attributes of a Specimen

- Unique ID#
- Priority: 1-4, critical/priority/routine cancer/routine non-cancer
- Start/End timestamps for each stage
- Internal/External request
 - Affects duration of reception tasks
- Cut-up category
 - Affects cut-up duration and staff requirements
- Block and slide type
 - Affects processing machine duration, etc.
- # of blocks (created during cut-up stage)



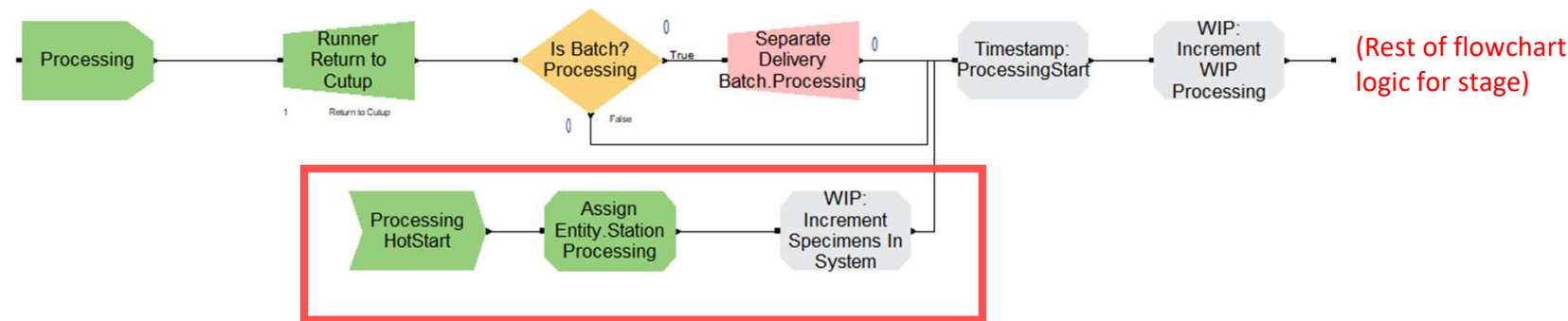
Specimen Arrivals

- Excel table → Arena schedule
- Hourly rates, defined over one week (cyclical)
- Separate rates for cancer/non-cancer pathway specimens
- Rejection sampling used to create time-varying Poisson processes



Simulation hot-start

- Bootstrap initial simulation state
- Read specimens from Excel file (one sheet per stage)
- Each sheet contains, for each specimen, timestamps for completed stages + attribute values set so far
- Related: **early exit** points for studying parts of simulation model in **isolation**
 - Triggered manually by modifying simulation settings



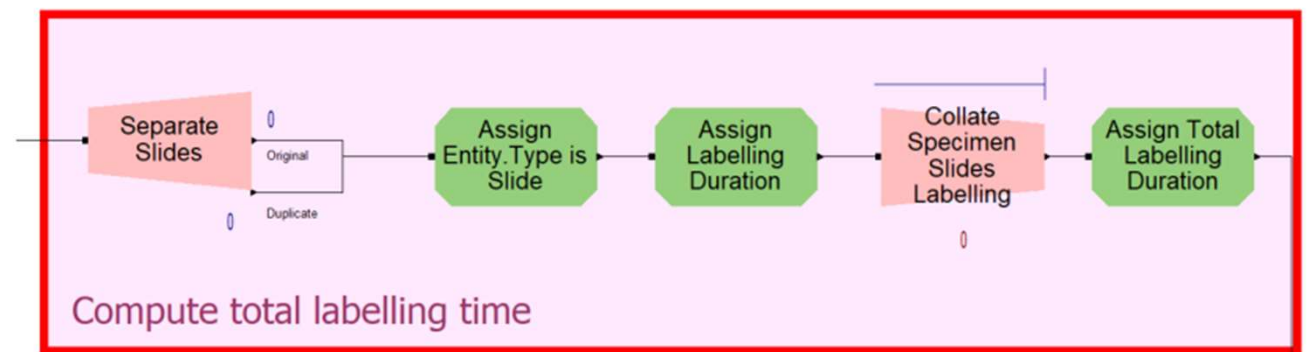
Task durations

- Three-point estimates → triangular distribution
- Exception: machine tasks
 - Constant duration
- For machine tasks:
 - Loading time
 - Machine time (unstaffed)
 - Unloading time
- Can be defined per specimen/block/slide/batch
- Batching/collation times not explicitly defined; combined with preceding task



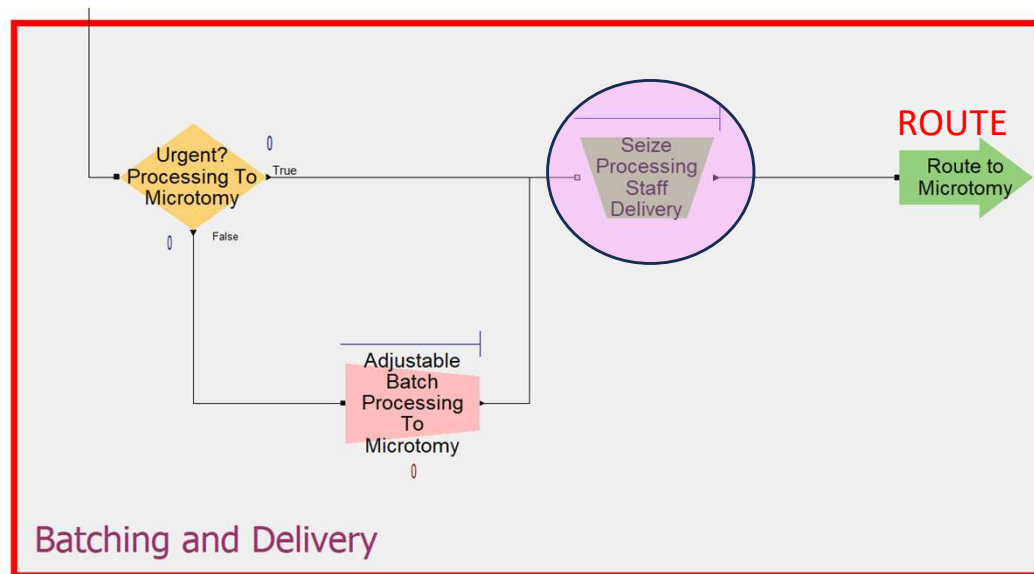
Task duration based on # of subcomponents

- Example: slide labelling
- Ensure all slides of specimen are labelled together, but duration depends on number of slides
- Steps
 1. Split Specimen into correct # of slides
 2. Assign random task duration for each slide based on distribution
 3. Collate slides based on Specimen ID#
 4. Assign total duration by summing across all slides in collated group
- Works because Arena can retain attributes of group members in a collation, while still treating collation as ordinary specimen object

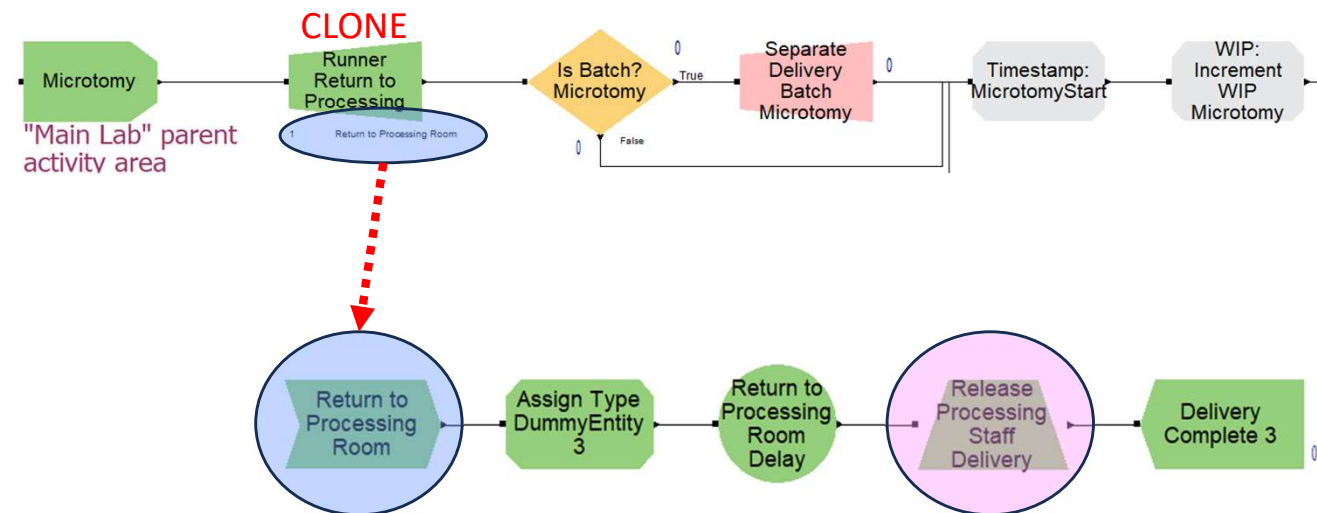


Delivery between stages

- Originating stage
 - Urgent specimens are unbatched
 - Other specimens are **batched**, but with a time limit
 - **ROUTE** block has an associated delay (read from Excel file)



- Receiving stage
 - **CLONE** block triggers runner's return trip
 - Specimen continues on **main** path
 - Delivery batches **separated**



Time-triggered batch jobs (1)

- Example: machine processing of wax blocks
 - Series of chemical processes
 - Except for urgent specimens, left to run overnight
 - Each batch contains up to 36 or 300 wax blocks depending on type
 - Need **time-based gate** to delay machine load / start
- Need to:
 - Batch **specimens** based on how many **blocks** can fit in a job
 - Blocks from same specimen **cannot** be split across batches
 - Release **incomplete** batches before end-of-day



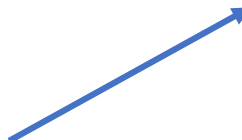
Time-triggered batch jobs (2)

Signals triggered at 1pm, just before 4:30pm, and at 4:30pm



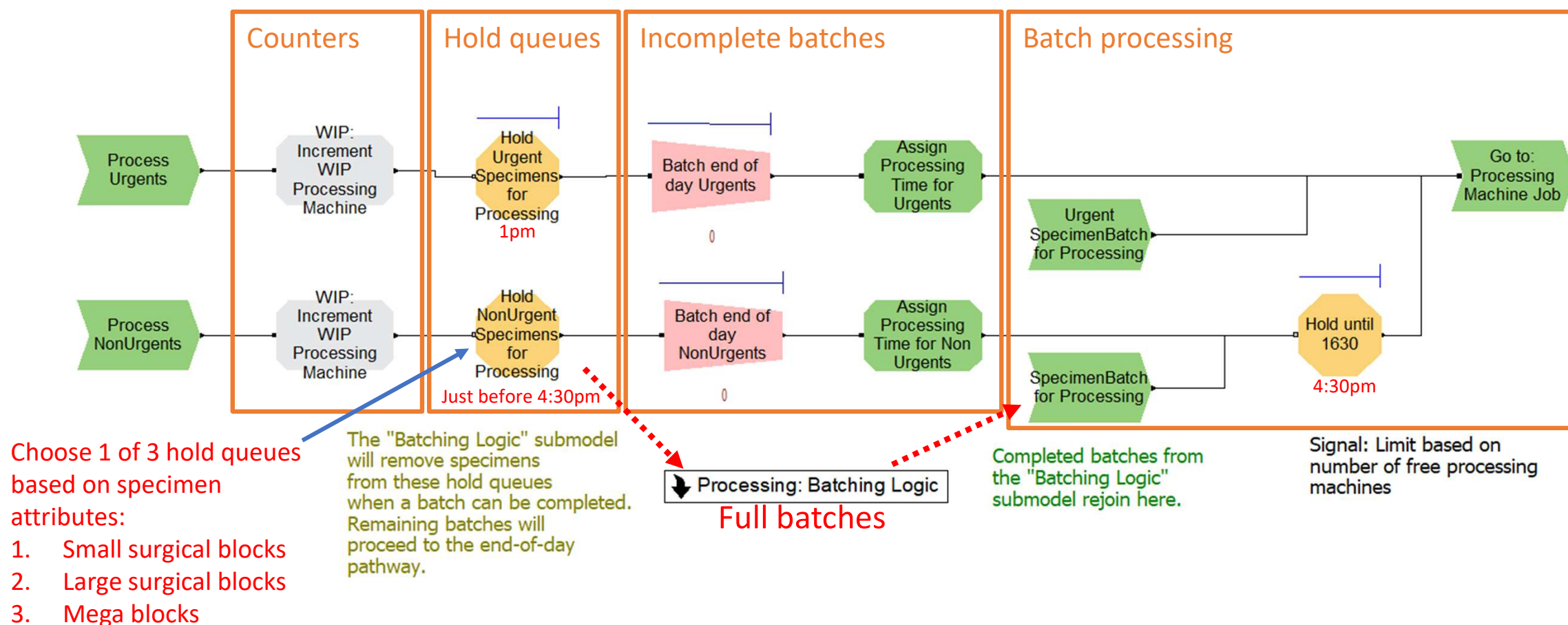
Choose 1 of 3 hold queues based on specimen attributes:

1. Small surgical blocks
2. Large surgical blocks
3. Mega blocks

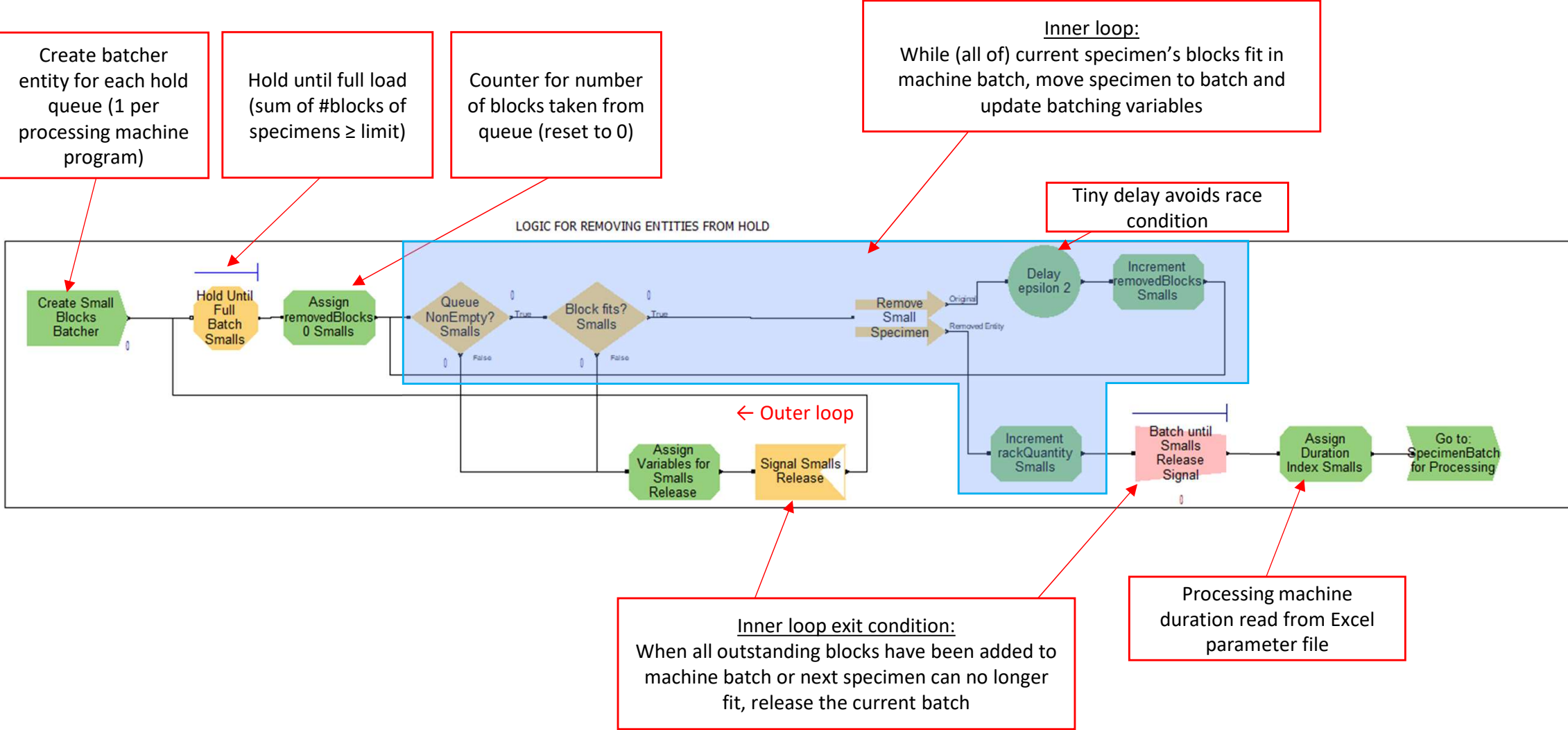


Time-triggered batch jobs (2)

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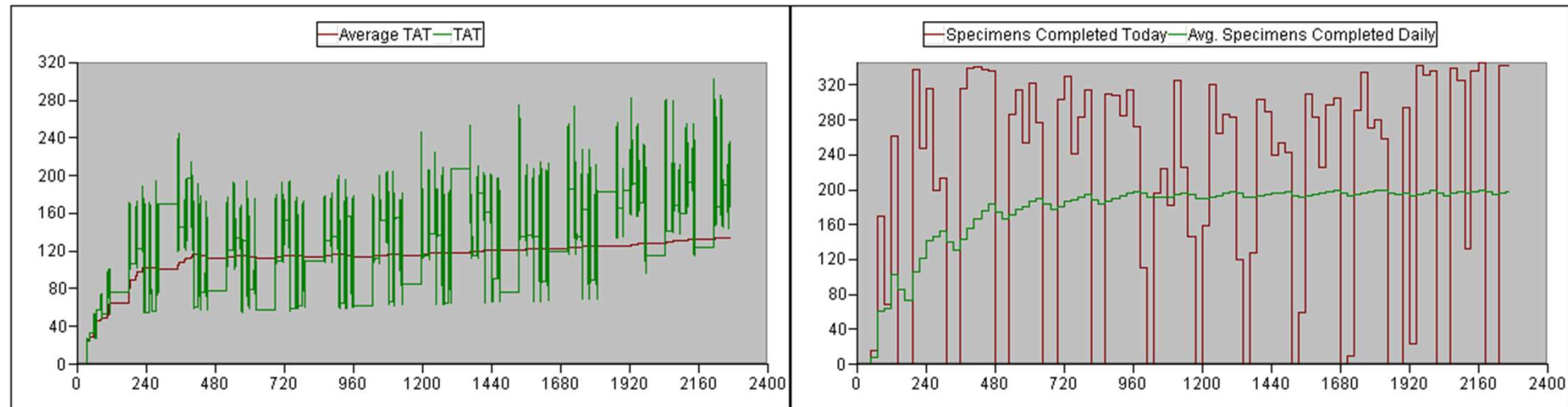
Time-triggered batch jobs (3)



Results & Scenario Analyses

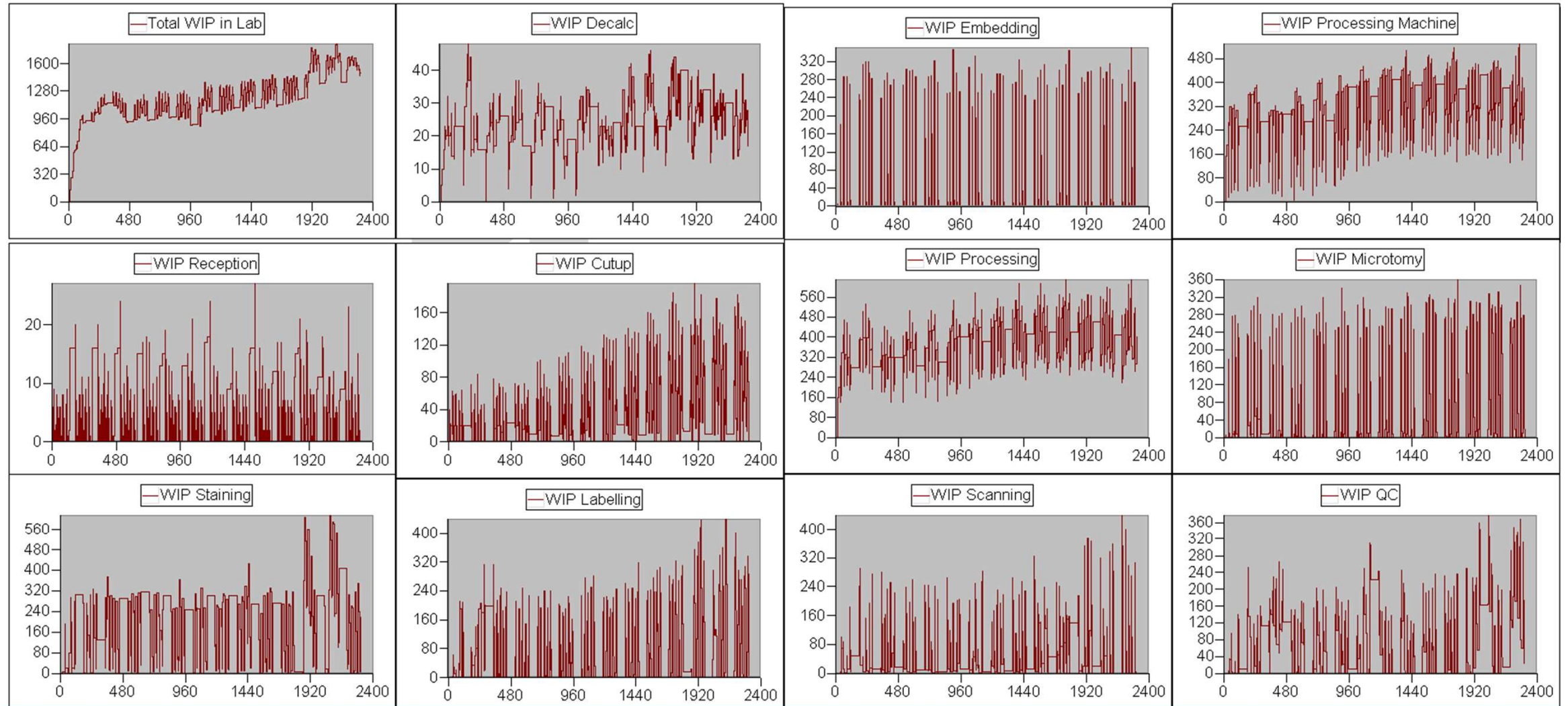


Plots: Turnaround time and throughput



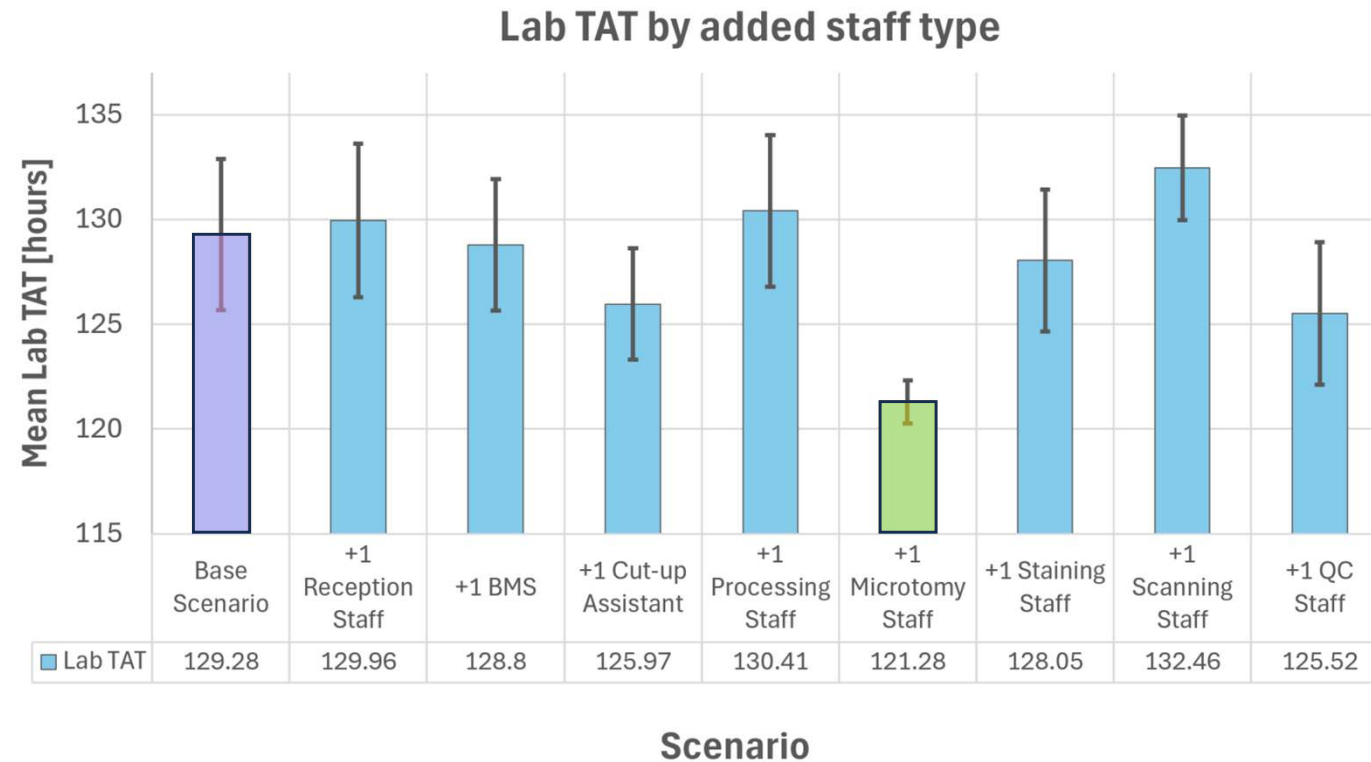
Mean TAT of about 140 hours; about 200 specimens completed daily

Plots: Work-in-progress



Scenario: adding one more staff

- Staff levels are defined in Excel for each staff type (nine types in model)
- Error bar shows 95% confidence interval of mean
- **+1 Microtomy staff only** scenario with significantly lower mean lab TAT compared to base scenario (no overlap of confidence interval)



Discussion



Summary

- Simulation model for KPI estimation, scenario analyses (what-if comparison)
- Can hot-start simulation with bootstrapped initial state
- Can handle batching with attribute-based batch size
 - i.e., total number of **blocks** among all **specimens** in batch
- Can handle simultaneous tasks
 - Delivery runners return to base while specimen continues along main path



Future work

- Improved arrival modelling
 - Current: time-varying Poisson processes (**single** arrivals)
 - **Batched** arrivals based on courier deliveries
 - **Federate** lab model with logistics model?
- Effect of specialities on cut-up, reporting stages
- Full implementation of specimen priorities
 - Currently, only urgent vs. non-urgent considered
- Data integration
 - Staff rotas
 - Building layout → travel time computation
 - Building state → e.g. if lift down / access door malfunctioning, need to take longer route (if one exists)
- Inventory management
 - Effect of low stock on processes / lab throughput



Thank you!

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