

# Output Group

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## Part 1: Selection of Sensors

- What we can do right now!

## Part 2: Cloning/ Characterization

- Timeline

## Part 3: Imaging/ Modeling

# Part 1: Selection of Sensors

- Select Sensors: ***At least 5 promoters needed!***
- Promoters in Bioquant:
  - Elk 1 (ras-raf-MAPK signaling cascade)
  - PUMA (p53 up-regulated modulator of apoptosis)
  - ATM2 (DNA damage response)
  - Hypoxia (which one is the best?)
- Are there other groups at DKFZ or EMBL we could ask for promoters??

# Part 1: Selection of Output Signals

- Select Fluorescent Proteins to be used
  - Minimum of 3 colors
- Select useful Localizations Signals
  - Mitochondria, ER, Plasma membrane

## Part 2: Cloning/ Characterization

- Cloning (3-4 weeks)
- Stable cell lines (2-3 weeks)
- Characterization of Constructs (2-3 weeks)
- Modeling can start once a stable cell line is established

➤ **PARTS HAVE TO BE SUBMITTED BY END OF SEPTEMBER!!!**

## Part 3: Imaging/ Modeling

- As soon as we get a working system the modeling group can start working on the “Cell Profiler”
- Modeling can run in parallel:
  - promotor modeling / interface / CellProfiler / colour encoding

# Time Schedule

June

- Selection of Sensors
- Plasmid Design

July

- Cloning
- Characterization of Constructs

August

- Cloning cont.
- Generation of stable cell lines

September

- Cloning/ Characterization/ Stable Cell line cont.
- **Submit parts!!**

October

- Imaging
- Modeling - Toolbox