

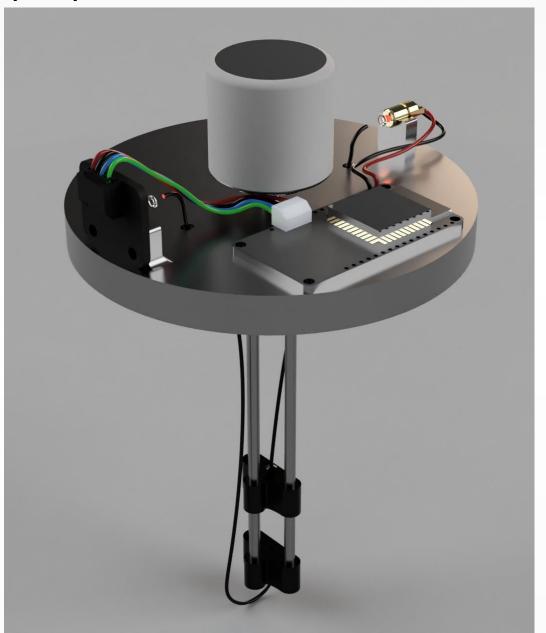
## Design & Development of a Novel Spectrophotometry Based Sensor for Cyanobacterial Growth Monitoring

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Industry standard benchtop spectrophotometry is time-consuming, expensive, has low data throughput, and is prone to user error [1]. Spectrophotometric sensors measure light attenuation/absorbance, which correlates to the bacterial concentration [2]. The biggest drawback, however, is that non-linear attenuation and multiple scattering events limit these sensors to a fixed range of concentrations [3]. This project looked at developing an improved spectrophotometry sensor for photobioreactor integration, enhancing experimental power and increasing research accessibility.

#### **Initial Design**

The initial design was a variable-pathlength fiber optic sensor. This approach reduces multiple scatter events without dilution, as absorbance and optical pathlength are proportional [4].



Pathlength adjusted using:

- Piston-displacement equation and Hill Climb optimisation autocalibration [5]. Error is shown in **figure 2.**
- Motor controlled by an ESP32 microcontroller.
- A luminosity sensor recorded transmitted light [6].

Figure 1: CAD render of initial design.

PMMA optical fibers were used as LASER waveguides for their IR transmittance, and cost. Allows for in-situ sensing, ambient filtering (**figure 3**) and improving signal/noise ratio.

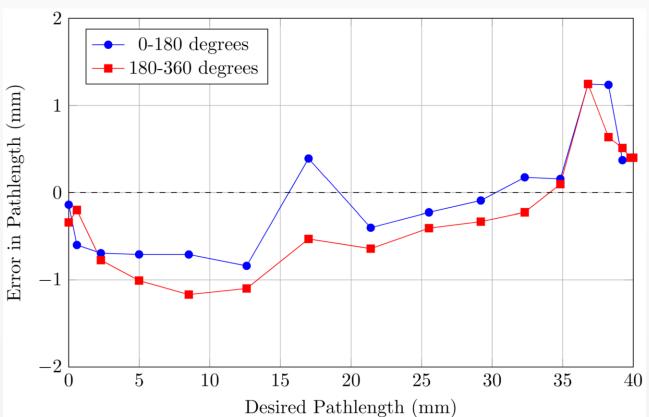


Figure 2: Variable pathlength error analysis for full motor rotation.

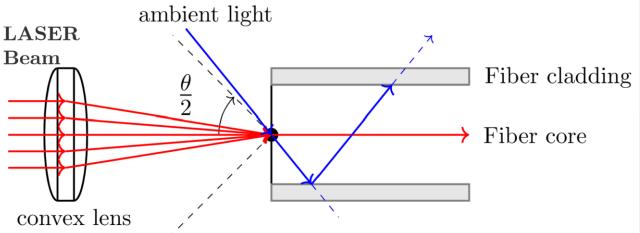


Figure 3: LASER-to-fiber optic coupling.

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Figure 4: Showcase at Prototypes For Humanity Expo and competition, Dubai 2023.

#### **Post-Mortem Analysis**

The device in **figure 1** featured at the Prototypes for Humanity Expo in Dubai (**figure 4**), attracting interest from academics and BioTech investors. It can run continuously, adjusting pathlength as culture density increases. Main limitations are:

- 1. 650nm LASER photopigment overlap
- 2. Relative size and weight
- 3. Coupling pathlength and optics

#### Improved Design

The second design has; a 780nm LASER, reducing pigment overlap and improving sensitivity; a collimated fixed pathlength; a novel cyclic LASER intensity measurements (CLIM); custom sensing element.

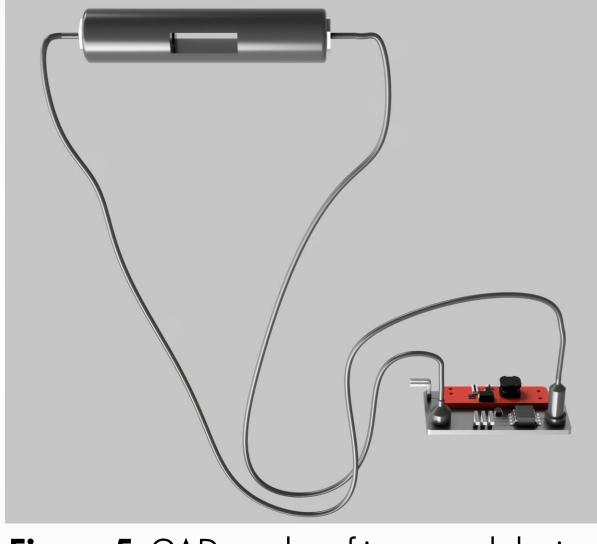
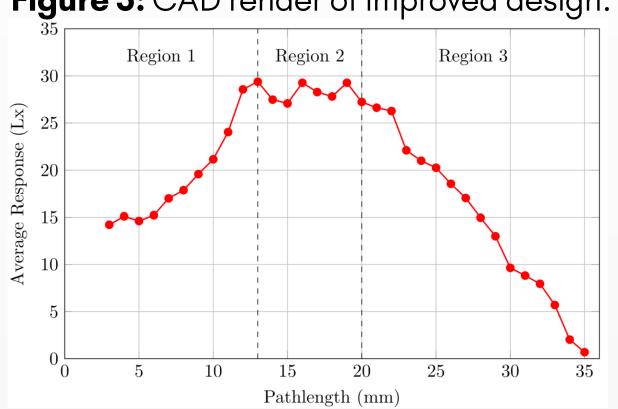


Figure 5: CAD render of improved design.



**Figure 6:** Region 1: converging. Region 2: collimated. Region 3: diverging.

Collimated pathlength ensures:

- No wasted incident light.
- Increasing bacterial cell contribution to absorbance reading.

The optimal pathlength was identified (**figure 6**, **Region 2**).

#### CLIMs:

- Enables dynamic sensitivity identification.
- Improved accuracy and sensitivity.
- Simplifies device by removing variable pathlength.

#### **Custom Sensing Element**

A transimpedance amplification (TIA) circuit (figure 7):

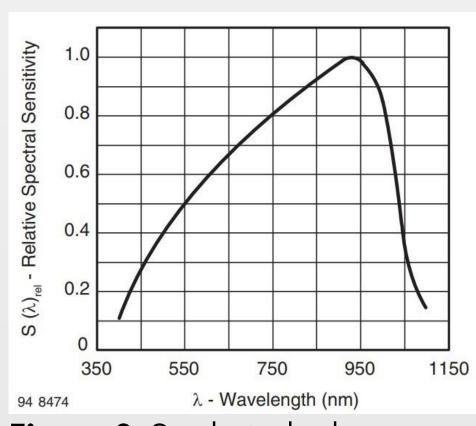
- Designed for silicon photodiode optimised for 780nm (**figure 8**).
- Fine-tuned sensing parameters for a reduced size.
- Output matches on-market hydroponic sensors.

Figure 7: Dual-stage TIA circuit.

### Conclusion

The sensor is a powerful and accessible tool for researchers and engineers, with:

- Higher accuracy
- Higher data throughput
- Reduced cost
- Easier integration with reactors



**Figure 8:** Si-photodiode sensitivity vs wavelength.

#### References

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- [2] K. Drescher, J. Dunkel, L. H. Cisneros, et al., "Fluid dynamics and noise in bacte-rial cell-cell and cell-surface scattering," vol. 108, doi:10.1073/pnas.1019079108/-/DCSupplemental.
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  [4] J. D. P. Peter Atkins, Atkins' Physical Chemistry, 8th ed. OUP Oxford, 2006, isbn: 10: 0198700725.
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