

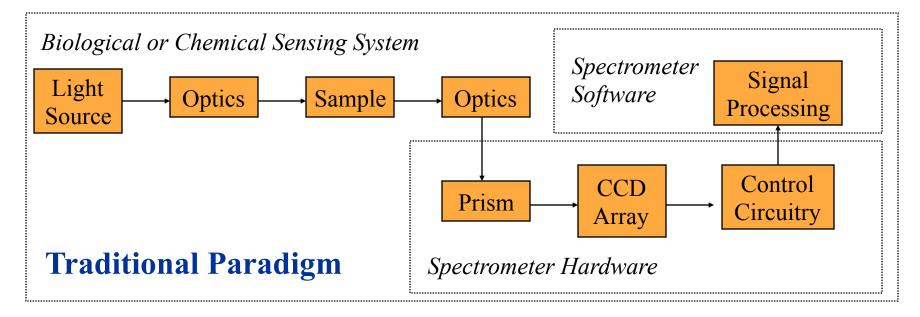


What's the problem?

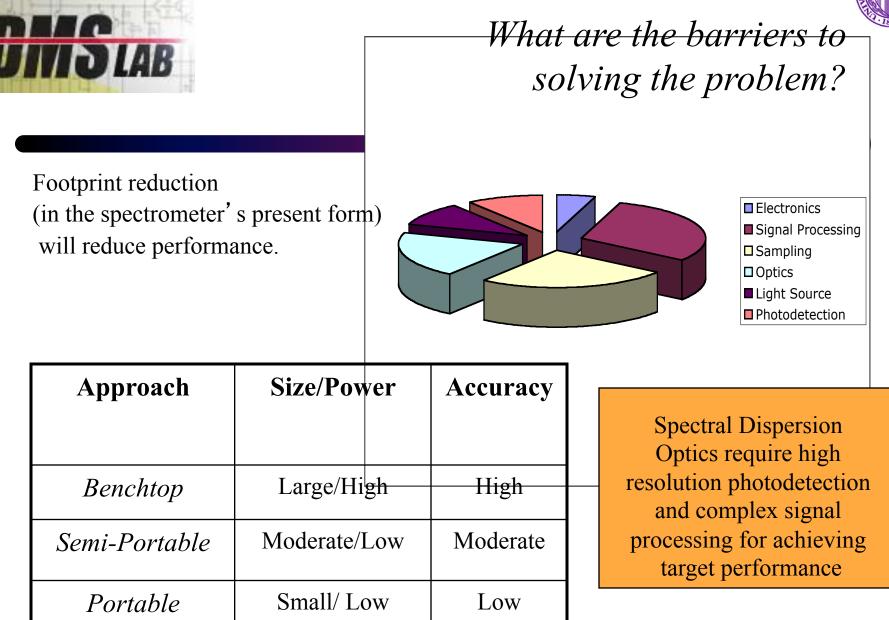
Reducing the cost/size/power envelope of Traditional Spectrometer Design is impossible

Ocean Optics sells the current market standard in portable spectrometers at \$4000-\$10,000 (configured)

Within reasonable expectations of performance.

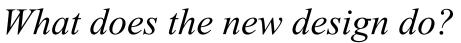




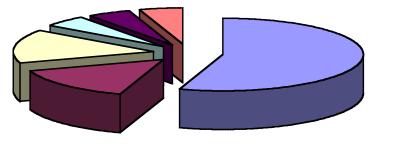








The new design paradigm enables scalable systems without compromising performance.



Electronics
Signal Processing
Sampling
Optics
Light Source
Photodetection

| Approach | Size/Power | /Size | Accuracy |
|-----------------|------------|-------|-----------|
| MMS | Small | | Moderate* |
| (Semi-Portable) | Low | | |
| MMS | Very Small | Very | Moderate* |
| (Portable) | Low | | |

The burden of spectral dispersion is moved to the light source and its control electronics, which are (far) more readily miniaturized.

**Relative to benchtop systems at \$50,000+ unit cost*



The Alternative Spectrometer Design

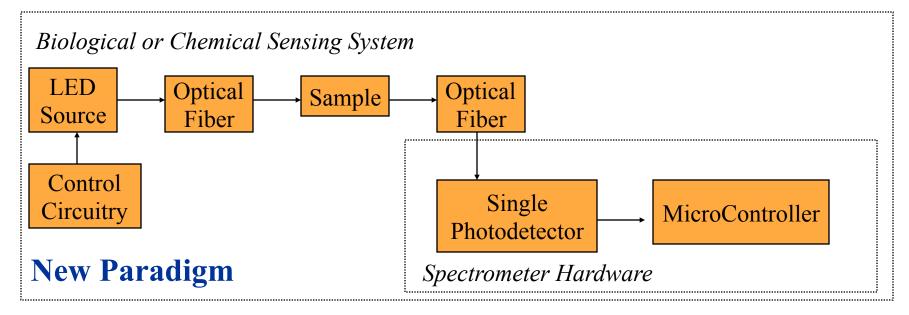
Spectral Dispersion is accomplished with:

Precise, high resolution manipulation of LED spectral emission

Rather than

Via Optical Means

Resulting in one light source/one photodetector systems with minimal optics



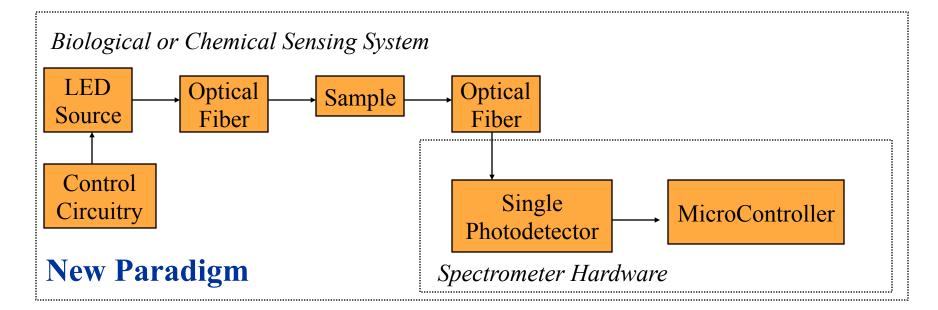




Alternative Spectrometer Design Making it Competitive

Relevant Questions:

- 1. Does LED-based Spectral Manipulation really work?
- 2. Does this design work well in a practical context?
 - Application #1: Fluorescence Analysis
 - Application #2: Surface Plasmon Resonance
- 3. Business Strategy

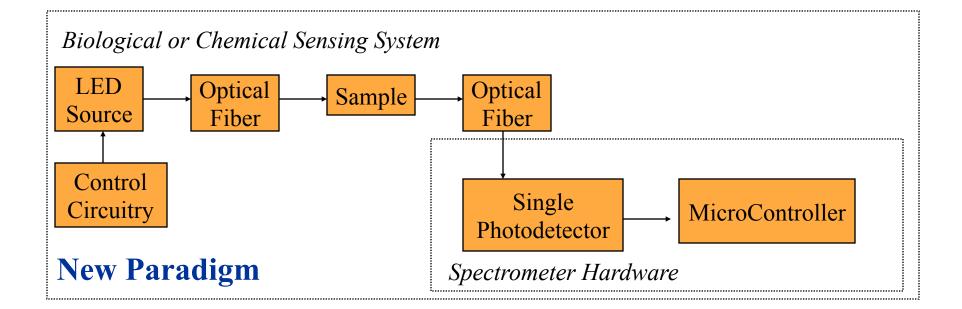






Alternative Spectrometer Design Making it Competitive

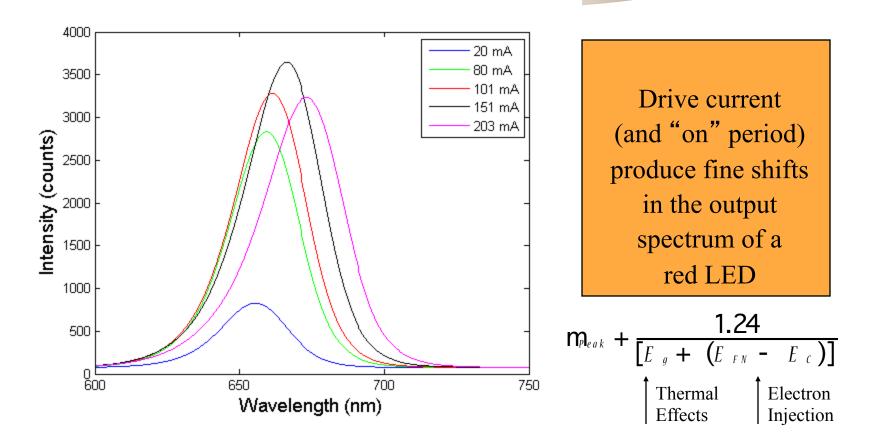
1. Does LED-based Spectral Manipulation really work?





Does LED Spectral Manipulation Work?

Controlled Spectral Shifts up to 40nm are possible.





Fluorescence Analysis is one of the most common uses of the spectrometer, AND

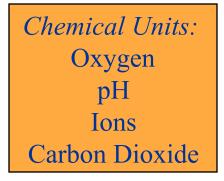
Portable fluorescence analysis is essential to:

- In-Situ,
- Point of Care,
- Continuous,
- Distributed,

Monitoring of Biological and Chemical Systems

Biological Units: Proteins Plankton Bacteria DNA/RNA

Biological Events: Metabolism Respiration Efficacy Reproduction







- Of the two most common means of fluorescence analysis: Steady-State Lifetime Analysis
- This system demonstration focuses on Steady-State

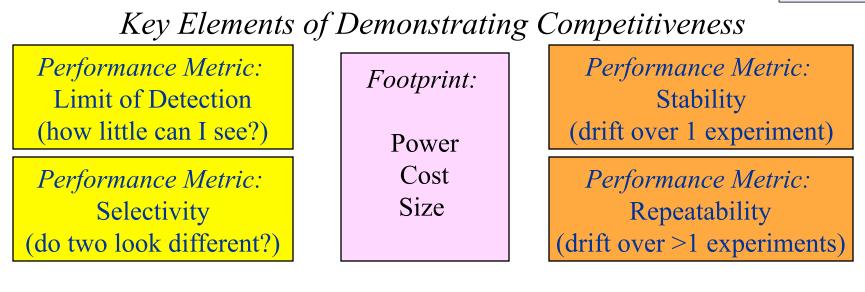
Orthogonal Measurements

Incoming Light

Detection Circuitry

Emitted Light

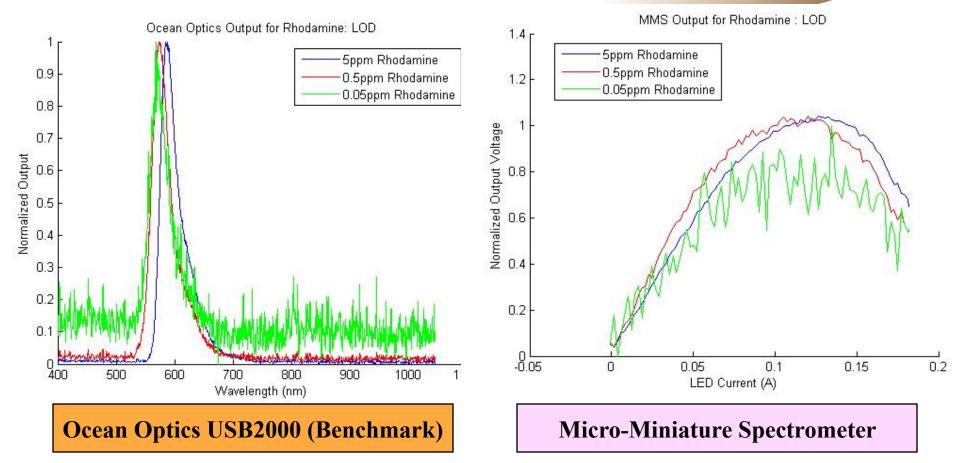
Sample







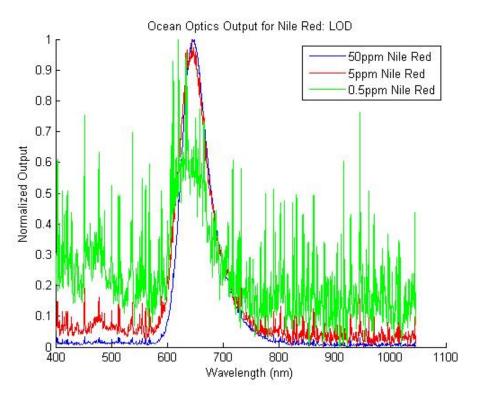
Limits of Detection (Rhodamine)



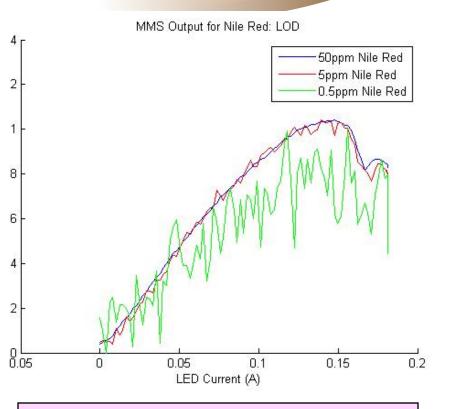




Limits of Detection (Nile Red)



Ocean Optics USB2000 (Benchmark)

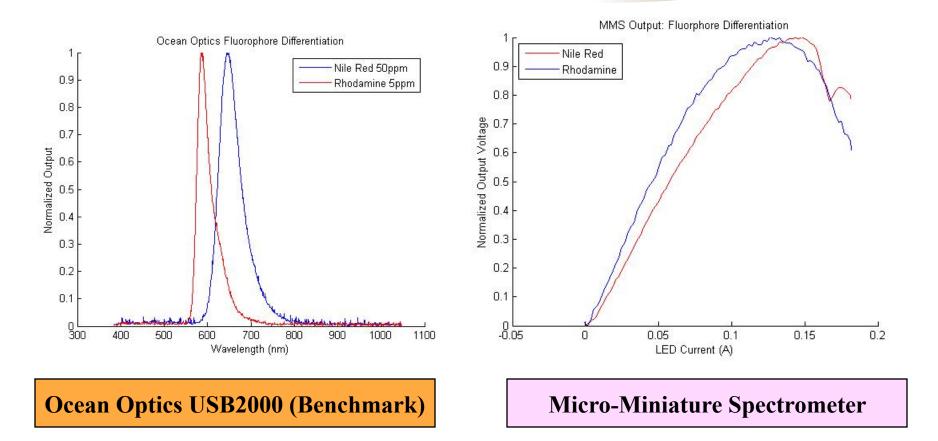


Micro-Miniature Spectrometer





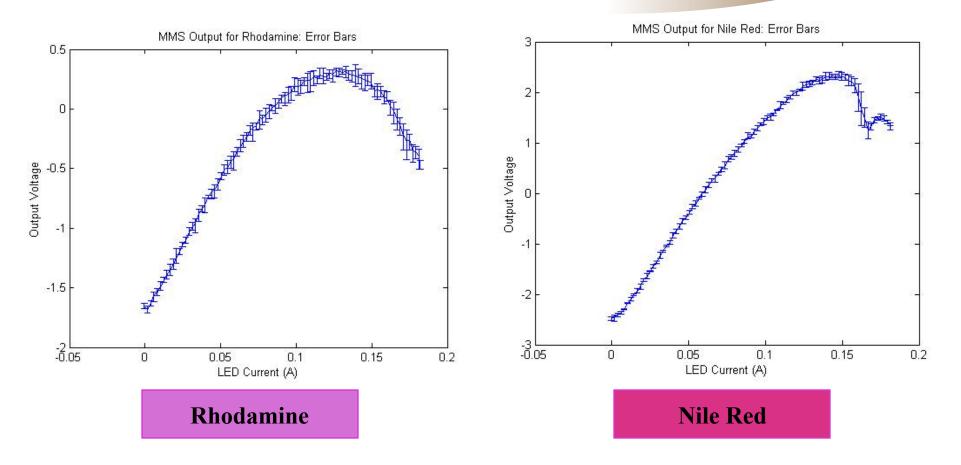
Selectivity (Rhodamine B & Nile Red)







Repeatability (Rhodamine B & Nile Red)







YES Performance Summary

| System | Limit of Detection | Selectivity | Stability | Repeatability |
|-------------------------|--|-------------|------------------|------------------|
| Ocean Optics USB2000 | 0.5 ppm (Nile Red) 0.05 ppm (Rhodamine) | Yes | 2.02%/ 5 min* | 5.06%/ 5 Days |
| MMS | 0.5 ppm (Nile Red) 0.05 ppm (Rhodamine) | Yes | 3.22%/ 5 min* | 6.63%/ 5 Days |

*Combination of drift, EMI, and fluctuations in sensing medium