

UNIVERSITY OF CALIFORNIA

Abstract

- A light field microscope is capable of capturing 3D information within a single shot, but it loses lateral resolution. [1]
- A combination of a fluorescence microscope with a diffuser is a proposed solution with the goal of obtaining better 3D resolution than a light field microscope equipped with a microlens array. [2]
- To demonstrate this idea, Zemax, Python, and Matlab are combined to compare the simulation results of different diffusers and microlens arrays.
- With this research, fluorescence beads are investigated, then experiments on living organisms, such as zebrafish are conducted.
- The goal is to eventually utilize this technique in improving the diagnosis and treatment of human diseases

Why Fluorescence?



Figure 1: The excited electrons jump from level 2 to level 3. The higher the electron goes more energy is needed. Later, the electrons on level 3 loses photons of energy and reverts back to its original ground state. [3]

- Fluorescent materials absorb light and UV radiation (i.e. x-rays, UV light)
- Emits wavelength than the originally one.
- A longer wavelength causes less scattering, allowing penetration inside the brain.

Comparison using Zemax

Software to replicate a live scenario optical design simulation

Diffuser – Random distribution of concave and convex regions Micro Lens Array – Multiple small lenses

Fluorescence Microscope with a Diffuser

Fluorescence Light Field Microscope (equipped with a micro lens array)



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Image

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More than 1 point source = Blur

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From the above comparison, we conclude that using a diffuser relieves the field of view constraint on the optical system. In the situation of having a huge field of view the diffuser would give us a better reconstruction than a microlens array. Thus, concluding that a microlens array would work best for this instant.

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Results



Diffuser Result



Sensor Data



Microlens Array Results

Conclusion

References

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