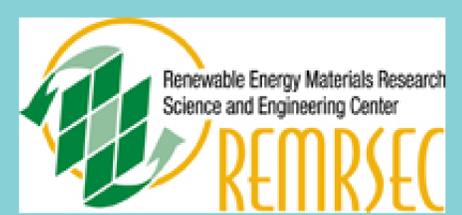




Spatial Frequency Modulated Imaging of Real-Time Laser-Matter Interaction

Confocal, Potentially Phase Sensitive Microscope with Reticule Modulation Mask



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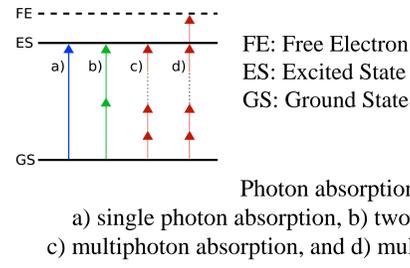
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Abstract

Laser modification of materials or biological systems would benefit from imaging systems that are able to quantify the interaction in real-time. One of the important constraints for such an imaging system is that it must be robust against optical scattering, as interactions may take place deep within a scattering material.

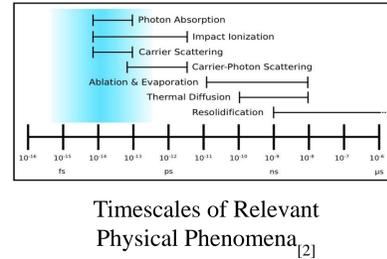
Here, for the first time, we demonstrate confocal spatial frequency modulation imaging that can show real-time changes and make two and three dimensional images of the material. Single element detection is used to aid in mitigating scattering effects of the optical signal, and an 800nm excitation wavelength enables detection down to millimeter depths in glass and plastic scattering systems.

Multi-Photon and Infrared Microcopy

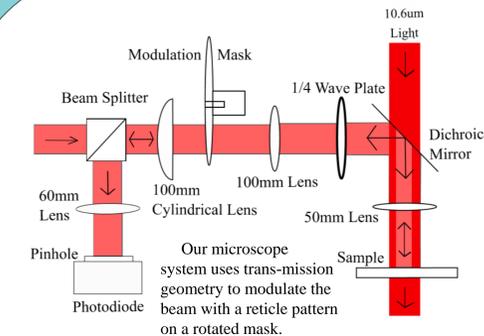


Infrared microcopy is useful because it is less damaging to biological materials_[3]. Shorter wavelengths, like UV, have enough energy to ionize the sample which leads to photobleaching, while longer wavelengths, like infrared, do not.

- Minimal heat deposited to material during removal
- Minimal interaction with material outside the focal volume
- Extremely high intensities ($> 10^{12}$ W/cm²) possible
- Sub-micron resolution over tens of cubic centimeters possible_[11]



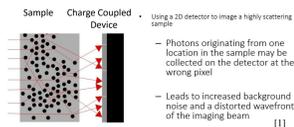
SPAtial Frequency Modulated Imaging (SPIFI)



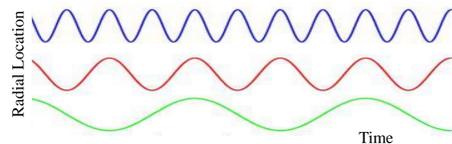
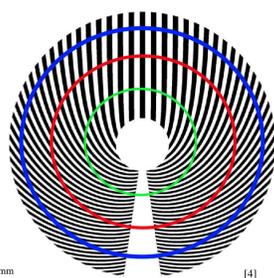
Background

- In SPIFI the mask modulates the amplitude of the light at a specific frequency in the beam based on its location on the radius of the mask.
- Taking the Fourier Transform of the signal from the detector maps the signal back to the horizontal location on the mask and sample which eliminates scattering ambiguity.

Disadvantages of Non-Spatial Frequency Modulation for Imaging



Modulation Mask

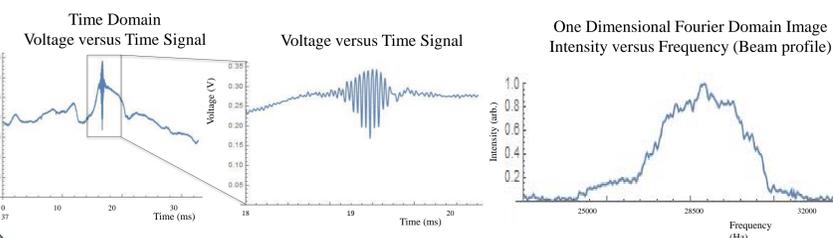


System Details

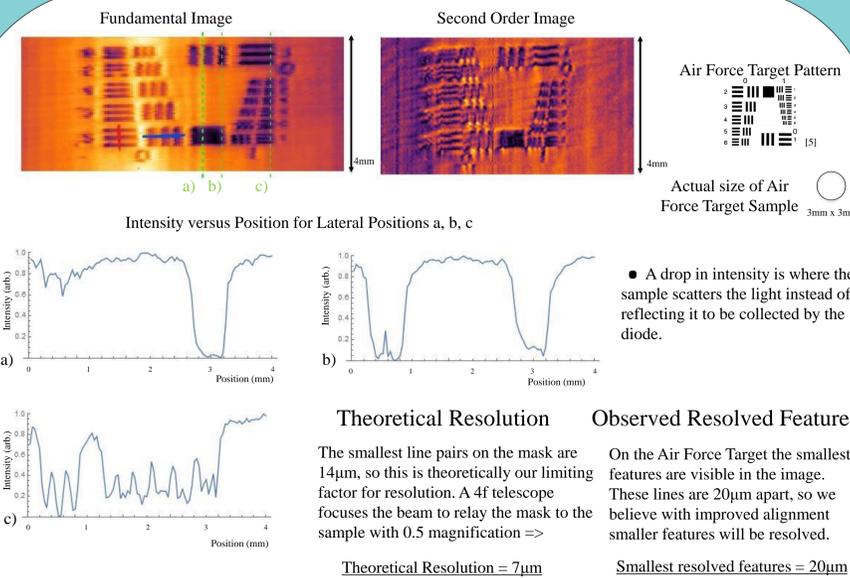
- 800nm Ti:Sapphire laser
- Average power of 78 mW
- Repetition rate of 76M Hz
- Pulse length of 80 fs
- Half-power spot diameter 13mm
- Mask rotates at 1600 rpm
- Dichroic mirror (transmits 10μm light, reflects 1μm)
- Dichroic used to integrate CO₂ laser
- Pinhole diameter of 600μm
- DET 100A Si Biased photodiode

Data Collection

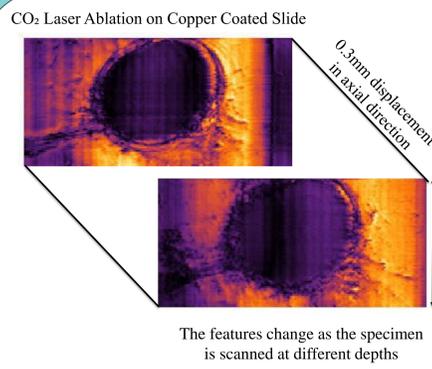
- Track the voltage vs. time signal from the photodiode.
- Take the fast Fourier transform to get the corresponding frequency of the modulated signal and its intensity at that frequency.
- The intensity vs. frequency graph shows a one dimensional image of the beam on the sample.
- By stepping the beam over the sample we create a two dimensional image of the sample's surface.



Resolution



Confocal System

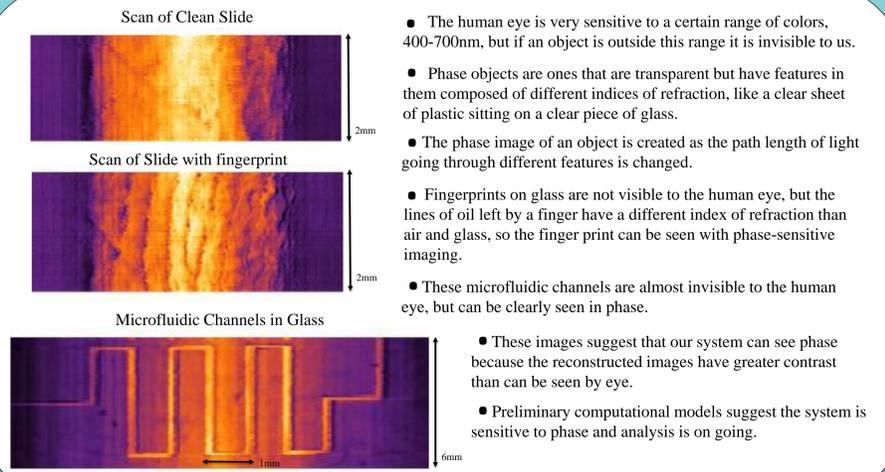


- In confocal microscopy illumination of the specimen is limited to a focused spot or line which is then scanned around the sample to get the full image.
- When this spot passes to the detector it is again limited by an exit pinhole to get rid of scattered light and signal not from the focus_[3].

Properties of Confocal Microscopes

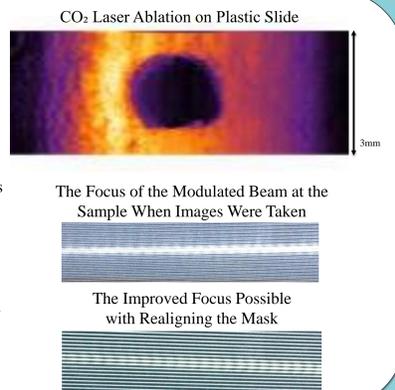
- Can be used in transmission or reflection geometries
- Detailed cross-sectional imaging is possible because signal from the sample not in the focus plane quickly drops off
- Improved signal to noise ratio
- Multiphoton absorption allows for the generation of images with different contrast mechanisms
- Creates much improved imaging of thick, light scattering materials_[3]

Phase Imaging



Conclusion

- We have demonstrated a novel new imaging modality that enables real-time characterization of laser ablation from a 10.6μm CO₂ laser. The resolution of the system was tested through imaging a series of line-pair targets fabricated in-house by our femtosecond laser micromachining system.
- Line-pairs down to 20μm were observed, which is greater than the calculated image resolution for the system. Improved alignment methods should enable diffraction-limited imaging.
- Finally, we imaged a fingerprint on acrylic and microfluidic channels in glass, which are both essentially phase targets. The image contrast obtained suggests that the system is capable of generating contrast based on phase differences in the target. We are working on new models of the microscope to help explain the apparent phase contrast.



References and Acknowledgements

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[3] Pawley, J., Ed. *Handbook of Biological Confocal Microscopy*, Third Edition; Springer Science+Business Media, LLC: New York, 2006.

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[5] Original USAF Resolution Charts, ProSciTech, https://proscitech.com/?navaction=show_page&chapter

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