

Nanoscale dynamics in ultrafast relaxation from radiation damage in SiO₂

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Abstract

Ionisation dynamics on the nanoscale seed the processes that govern relaxation to macroscopic equilibrium in irradiated matter. Therefore, understanding the conditions that underpin this transition is critical in a wide range of applications from healthcare to radiation science. Recently we have demonstrated that laser driven ion accelerators can provide an ultrafast tool for studying this inherently multiscale regime with temporal resolution < 0.5 ps [1,2]. Here we demonstrate that it is possible to interrogate these ultrafast processes in real-time by contrasting how recovery scales with interaction dimensionality for different ionising species. We employ single-shot optical streaking to track the decay time constant, τ_c , of free carriers in SiO₂ due to picosecond-scale (ps, 10^{-12} s) pulses of X-rays and protons from a single laser-driven accelerator. Exploiting the nanoscopically heterogeneous density of SiO₂ aerogels, our results reveal a strong scaling of τ_c with average density (ρ_{av}) for proton interactions. Conversely, τ_c is observed to be nearly independent of ρ_{av} for X-ray interactions. Cross referencing these observations with novel numerical simulations exposes the delicate interplay between the nanoscale conditions and emergent mechanisms, namely electron-phonon interaction, that drive ultrafast recovery in irradiated bulk SiO₂.

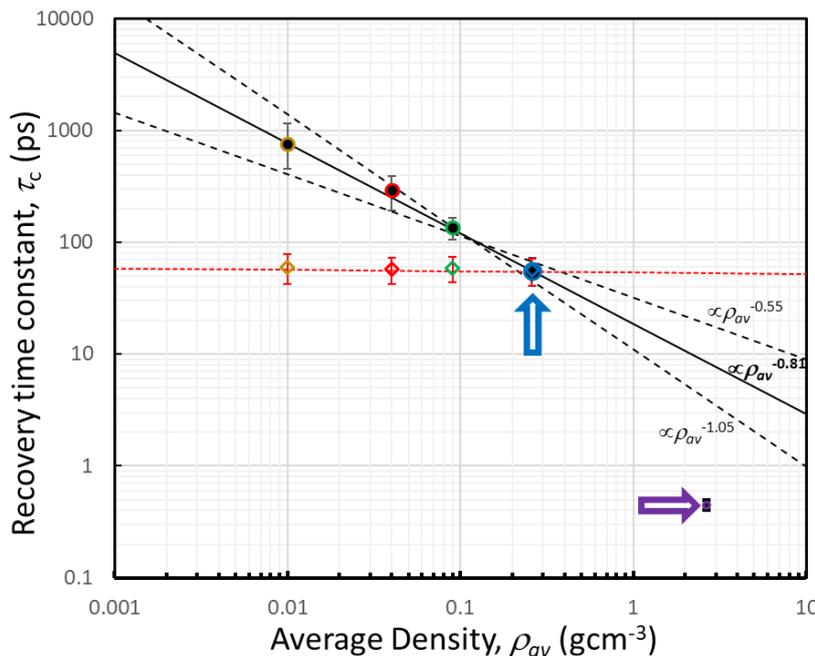


Figure 1. The scaling of the recovery time constant for ps scale pulses of X-rays (red dotted line) and protons (black solid line) interacting in SiO₂ ($\rho_{av} = 2.66$ gcm⁻³) and silica aerogels ($\rho_{av} = 0.01 - 0.26$ gcm⁻³)

References

- [1] Dromey, B., et al., *Nat. Comms.*, **2016**, 7, 10642
- [2] Senje, L., et al., *App. Phys. Letts.*, **2017**, 110, 104102