

Ultrafast, Time-Resolved Dynamics of Carriers, Spins, and Phonons in Solid-State Materials Studied by Femtosecond Optical Spectroscopy

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Femtosecond optical pump-probe spectroscopy is the most versatile and most frequently used method for time-resolved studies of the ultrafast nonequilibrium phenomena in solid-state materials. The technique uses femtosecond laser pulses for both optical excitation and probing, simultaneously providing a timing gate for measured events, and acts as a precise clock to control the generation and detection of tested events. In all pump-probe experiments, absorption of pump photons changes optical properties of a tested material by one or several optically-induced effects, such as band-filling, electron-hole recombination, phonon or thermal relaxation, electron-phonon interaction, spin polarization, Cooper-pair breaking in superconductors, etc. Next, a second beam (probe beam) of much weaker intensity and time-delayed with respect to the pump, is incident on the material overlapping the pump-excitation region. The reflected and/or transmitted probe signal is collected with a photodetector and the corresponding normalized differential reflectivity and/or transmissivity change as a function of the pump-probe time delay is a measure of the nonequilibrium dynamics of the excited subsystem in the studied material. Jitter-free time resolution of ~100 fs or better is routinely obtained in the sampling-type, pump-probe measurements. We will review the current state-of-the-art and the advancement of the femtosecond pump-probe spectroscopy and demonstrate its unique usefulness in studying nonequilibrium, time-resolved dynamics of carriers, spins, and coherent phonons in condensed-matter systems varying from semiconductors to spintronic materials and superconductors.

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