

# EXCITONIC FINGERPRINT OF ATOMICALLY THIN 2D MATERIALS

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As truly two-dimensional materials with a weak dielectric screening, monolayer transition metal dichalcogenides (TMDs) show a remarkably strong Coulomb interaction giving rise to the formation of tightly bound excitons. In addition to the optically accessible bright excitonic states, there is also a variety of optically forbidden states including excitons exhibiting a non-zero angular momentum or a non-zero center-of-mass momentum.

Here, we present the excitonic fingerprint in optical absorption and differential transmission spectra based on a microscopic approach that combines the Wannier equation with TMD Bloch equations. We show the appearance of a pronounced Rydberg-like series of excitonic transitions with binding energies in the range of 0.5 eV [1]. We investigate the microscopic origin of their homogeneous linewidth including radiative and phonon-assisted non-radiative relaxation channels [2, 3]. We predict a significant disorder-induced coupling of bright and dark excitonic states offering a strategy to circumvent optical selection rules and make dark states visible in optical spectra [4]. Finally, we study exciton valley dynamics including Coulomb-driven intervalley coupling mechanisms between the high symmetry K and K' points [5].

*Keywords:* 2D materials; dark and bright excitons

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