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Exciton reactions on carbon nanotubes: an experimental testbed for critical dynamics¹ JEREMY ALLAM, Advanced Technology Institute and Department of Physics, University of Surrey, Guildford GU2 7XH, United Kingdom

The one-dimensional coalescing random walk $(X + X \to X)$ is a paradigmatic reaction-diffusion system due to both its exact solvability [1,2] and the experimental observation of nonclassical kinetics at asymptotically long times [3]. The solvability rests on the anticommutative property of intersecting trajectories of particles that react *instantly* and at *short range*: however, the validity of these assumptions in real systems has not previously been tested by experiment. We have shown that excitonexciton recombination (fusion) on carbon nanotubes provides a platform for quantitative studies of critical kinetics in a simple non-equilibrium system [4]. Under high excitation density we observed a crossover in the exciton density *n* between regimes of classical $(n \propto t^{-1})$ and anomalous $(n \propto t^{-1/2})$ scale invariance as predicted by renormalization group [5] and approximate [1] theories, arising from a finite reaction probability of ≈ 0.2 per encounter. At long times the exciton density and hence a demonstration of universality extending over both classical and critical regimes. Under low excitation, the early kinetics followed a Smoluchowski-Noyes form $dn/dt \propto n^2 t^{-1/2}$ rather than the asymptotic $dn/dt \propto n^3$, providing direct evidence for the spatial self-ordering that precedes critical scale invariance. We studied the re-emergence of microscopic detail at the classical crossover, which is abrupt and nonmonotonic due to competition between temporal and spatial averaging of critical fluctuations (i.e. finite reaction rate and range). It appears that real-world experiments will require more complete descriptions of the interactions than is available in existing models.

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