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Theoretical Description of the Self-Assembly Dynamics of DNA-functionalized Nanoparticles CHIA WEI HSU, Wesleyan University, FRANCESCO SCIORTINO, University La Sapienza, FRANCIS STARR, Wesleyan University — Nanoparticles (NP) functionalized with DNA strands can assemble into networks with unusual structure and phase behavior. Properties of such materials are known from computational studies, but a deeper understanding requires a theoretical description of the assembly process. As a step toward this goal, we show that the kinetics of self-assembly of a mixture of NP functionalized by two or three DNA strands can be quantitatively described by the Flory-Stockmayer (FS) theory of gelation. Below the percolation threshold for the DNA-linked NP network, the FS theory also accurately describes the cluster size distribution. These results rely on the formation of nearly loopless structures; a more general theoretical description remains to be developed. In addition, we show that at equilibrium, the diffusion of nanoparticles is connected to the formation of bonds through a simple relation.

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