

Linked topological colloids in a nematic host

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r s r f l s . f l s r l s r d m



Serius =



r m n n n r n , n d n m n n s . n s ,
 l n - k n r n s (33) h m n n s f l l
 l d l d n l s d m r . E r m n l s d f l
 m l n s f l n f l r l s n d d f l l s n r m s f l
 l d d r n f l d d n l d t l s d l s l
 l r n m n s f l r n s m m r f l L C s s , . d s
 n s n n n l n d l n m s (34), s l l s
 n d d r s f l s f l m r s s m s n r n l l
 n r l n f l r n s (3, 7, 8, 19, 35-37). r n d d t l r
 l r n s n l d s l n l s s n d s d f l s r r l s
 d r m s , s l l s l n f l n m n , k n r s s s , -
 r l , s r f l n n r n l f l n k d l l d l m n n s , n d
 r n l f l d s n r l m (35-37). L n k d m l m n n r -

laser light. The 3PEF-PM fluorescence intensity exhibits a strong well-defined dependence (25) on the orientation of linear polarization of the excitation beam relative to (). 3PEF-PM images, comprised of 3D stacks of optical slices, such as the ones shown in the SI Appendix, Figs. S12 and S13 and [Movies S3–S5](#), reveal orientations and relative positions of linked rings as well as the corresponding locations and configurations of topological defects accompanying them. Close analysis of 3PEF-PM stacks reveals dependence of 3D ()-structures on boundary conditions and topology of colloids. Optical videomicroscopy and holographic laser tweezers (25) probe elastic interactions between the linked rings. Additionally, high-power beams of laser tweezers allow for locally

a 170- μm -thick coverslip, spaced by 50- μm -thick Mylar strips. As the beam focus was translated through the monomeric fluid, we always began polymerization at the substrate-fluid interface to effectively anchor the structure while it is being drawn. Arrays of particles were then detached from substrates by gentle sonication and dispersed into LCs. As-manufactured particles impose tangential boundary conditions for (), but some of them were treated with DMOAP for perpendicular ones (8).

Optical Imaging and Laser Manipulation. Director structures are studied using a combination of conventional polarizing optical microscopy and a 3D non-linear imaging technique dubbed “three-photon excitation fluorescence polarizing microscopy” (3PEF-PM) (25), which is based on fluorescence of LC molecules excited through three-photon absorption of femtosecond infrared