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Title

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Permalink

https://escholarship.org/uc/item/8bh109hx

Journal

Biomacromolecules, 16(7)

ISSN

1525-7797

Authors

Das, Saurabh Miller, Dusty R Kaufman, Yair et al.

Publication Date

2015-07-01

DOI

10.1021/acs.biomac.5b00759

Peer reviewed

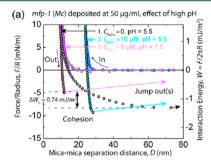


Correction to "Tough Coating Proteins: Subtle Sequence Variation Modulates Cohesion"

Saurabh Das, Dusty R. Miller, Yair Kaufman, Nadine R. Martinez Rodriguez, Alessia Pallaoro, Matthew J. Harrington, Maryte Gylys, Jacob N. Israelachvili,* and J. Herbert Waite*

Biomacromolecules, 2015, 16 (3), 1002-1008. DOI:10.1021/bm501893y

igure 3 caption corrected. Corrections highlighted in bold.



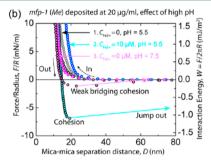


Figure 3. pH dependence of Fe³⁺-mediated cohesion between two symmetric mfp-1 (Mc) films. Representative force vs distance plot showing the interaction between two symmetric mfp-1 (Mc) films deposited at 50 μ g/mL in 0.1 M sodium acetate buffer, pH 5.5, 0.25 M KNO₃, and 1 mM bis—tris with $C_{Fe^{3+}} = 0$ (gray) and 10 μ M (blue) at pH 5.5. The cohesion between the mfp-1 (Mc) films was preserved after increasing the pH to 7.5 (magenta). (b) Representative force vs distance plot showing the interaction between two symmetric mfp-1 (Mc) films deposited at 20 μ g/mL in 0.1 M sodium acetate buffer, pH 5.5, 0.25 M KNO₃, and 1 mM bis—tris with $C_{Fe^{3+}} = 0$ (gray) and 10 μ M (blue) at pH 5.5. The surfaces showed a weak bridging cohesion ($W_c < 0.2 \text{ mJ/m}^2$) after increasing the pH to 7.5 (magenta). It should be noted that $C_{Fe^{3+}}$ is the concentration of ferric cation in the bulk solution between the surfaces. Flushing with buffer at pH 7.5 removes iron from the bulk solution, however, to the extent that the preadsorbed protein films already had some bound Fe³⁺, the Dopa—Fe³⁺ complexes will be present in them.