

Supplementary Material

Combining Polymer and Cyclodextrin Strategy for Drug Release of Sulfadiazine from Electrospun Fibers

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S1: Histograms

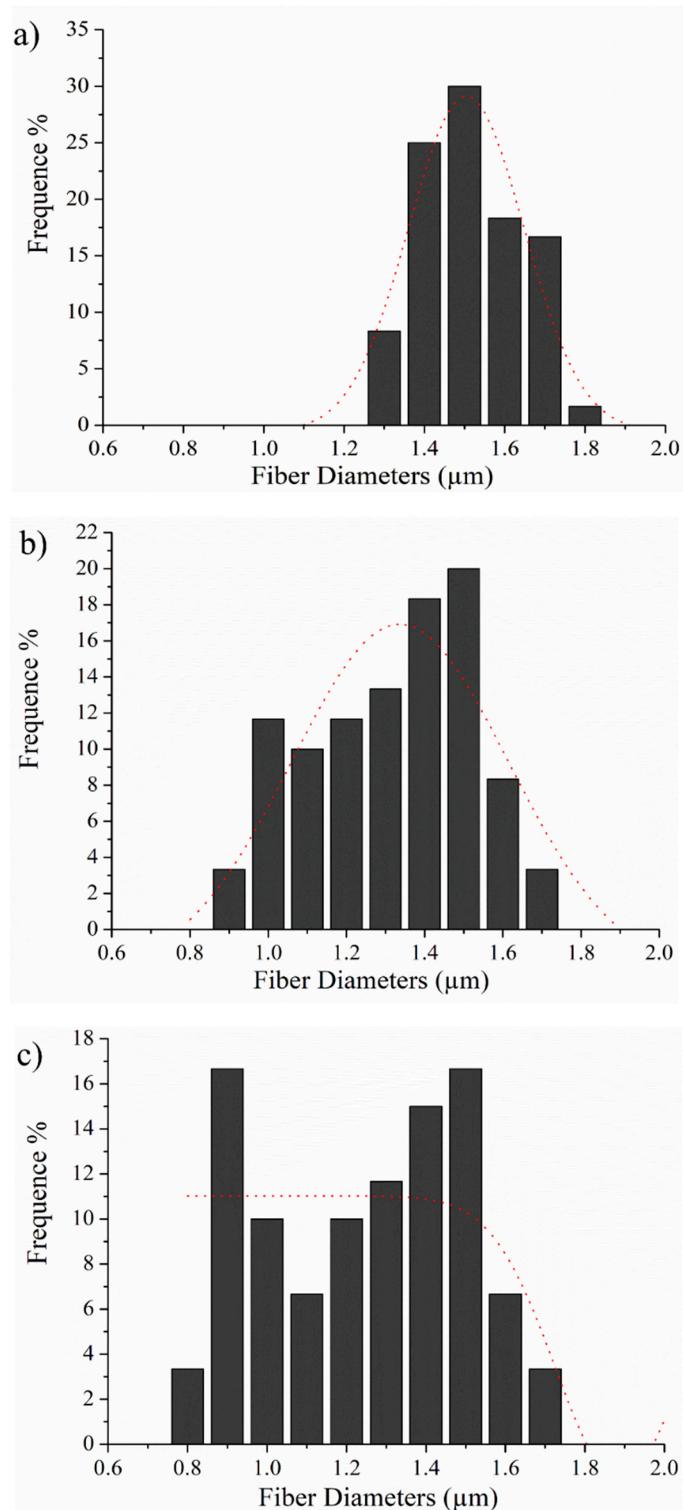


Figure S1: Histograms of fiber diameter distribution for: (a) PMMA, (b) PMMA SDS, and (c) PMMA SDS/HP β CD.

S2: Histograms

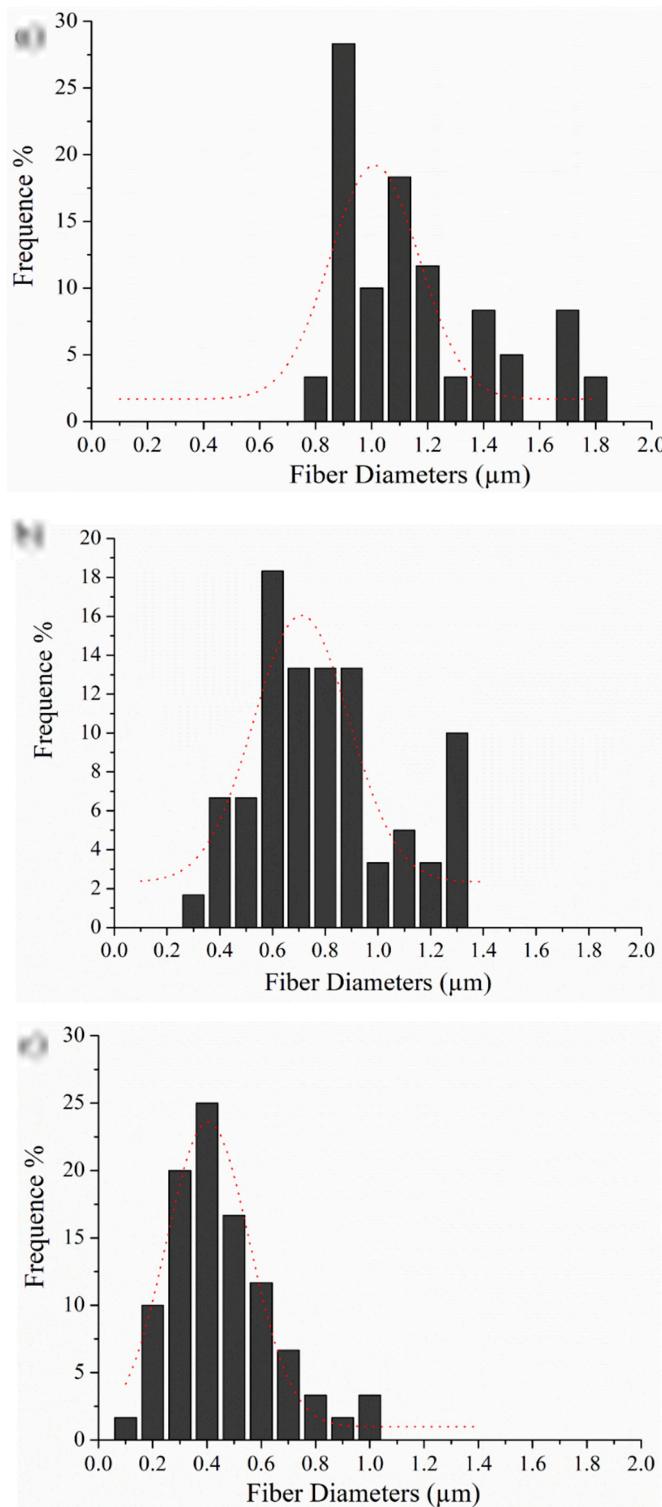


Figure S2: Histograms of fiber diameter distribution for: (a) PLGA, (b) PLGA SDS, and (c) PLGA SDS/HP β CD.

S3: Contact angle

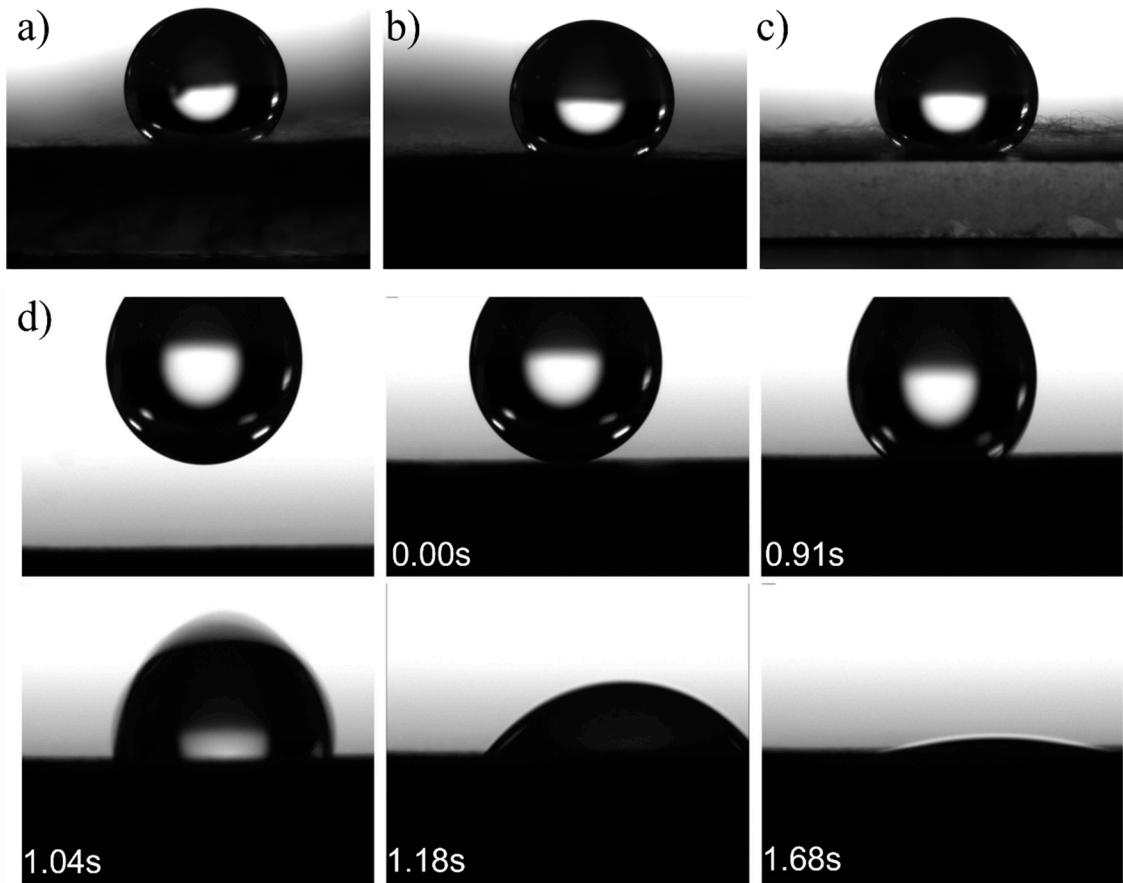


Figure S3: water contact angles images for the electrospun systems: (a) PMMA, (b) PMMA SDS, (c) PMMA SDS/HP β CD; (d) PLGA with the capture of video frames.

S4: calibration curve

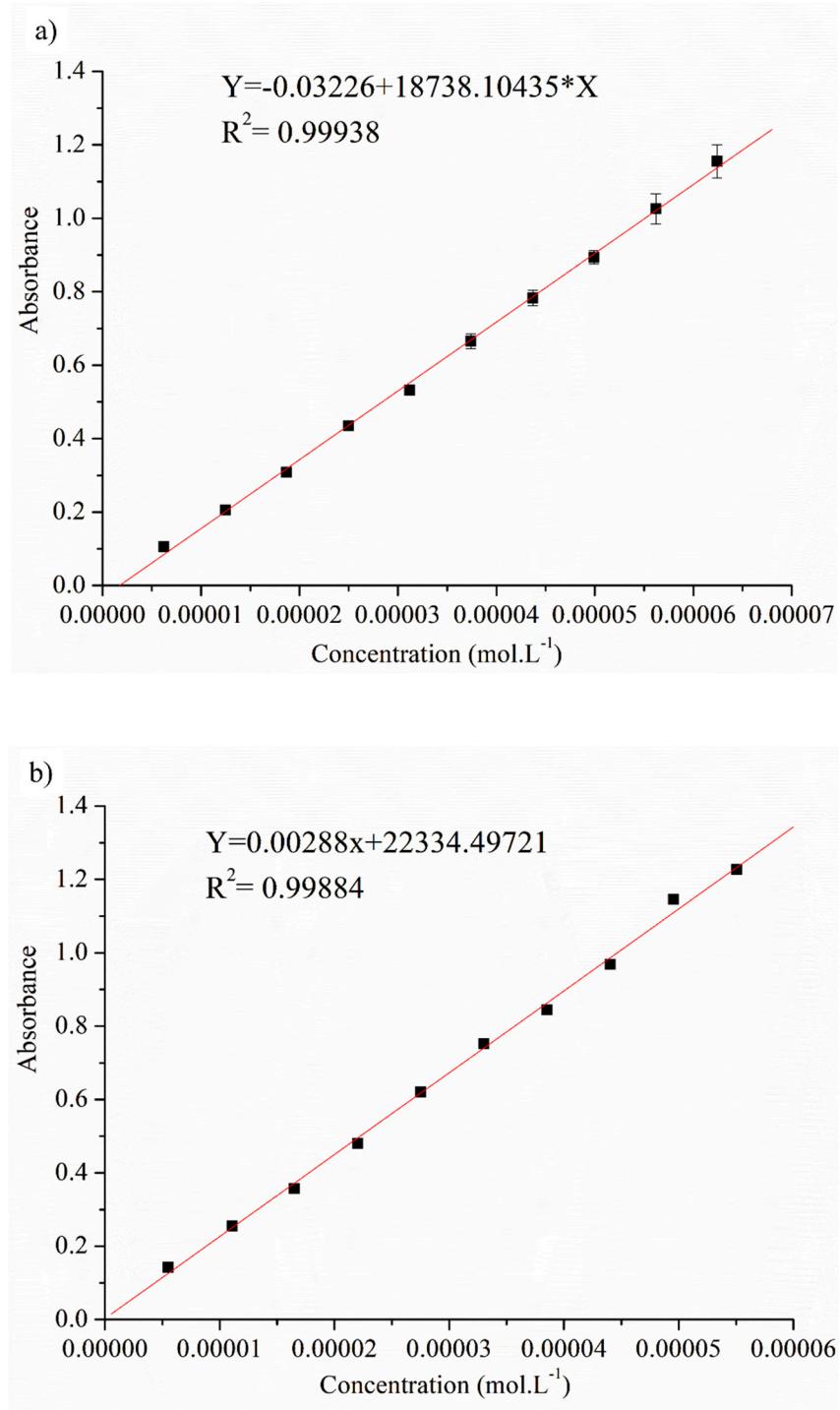


Figure S4: Calibration curve for SDS drug: (a) without HP β CD and (b) with HP β CD.

S5: microscopies images

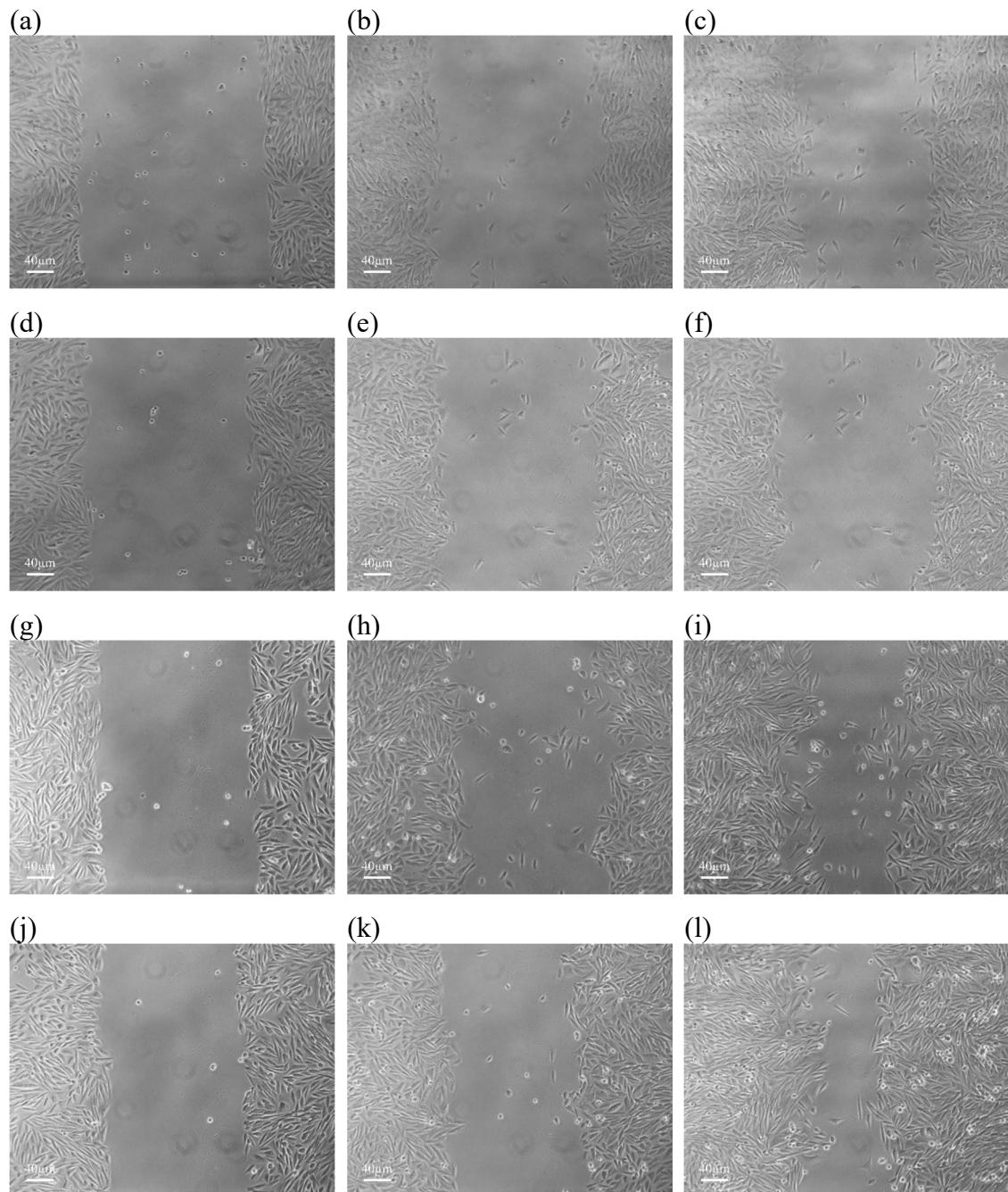


Figure S5: Microscopies images of a pseudo-wound created in a fibroblast cell L929 monolayer, at 0 hour (left column), at 10 hours (middle column), and at 24 hours (right column). The pseudo-wounds were treated with (a–c) PMMA, (d–f) PMMA SDS, (g–i) PMMA SDS/HP β CD, and (j–l) control group.

S6: microscopies images

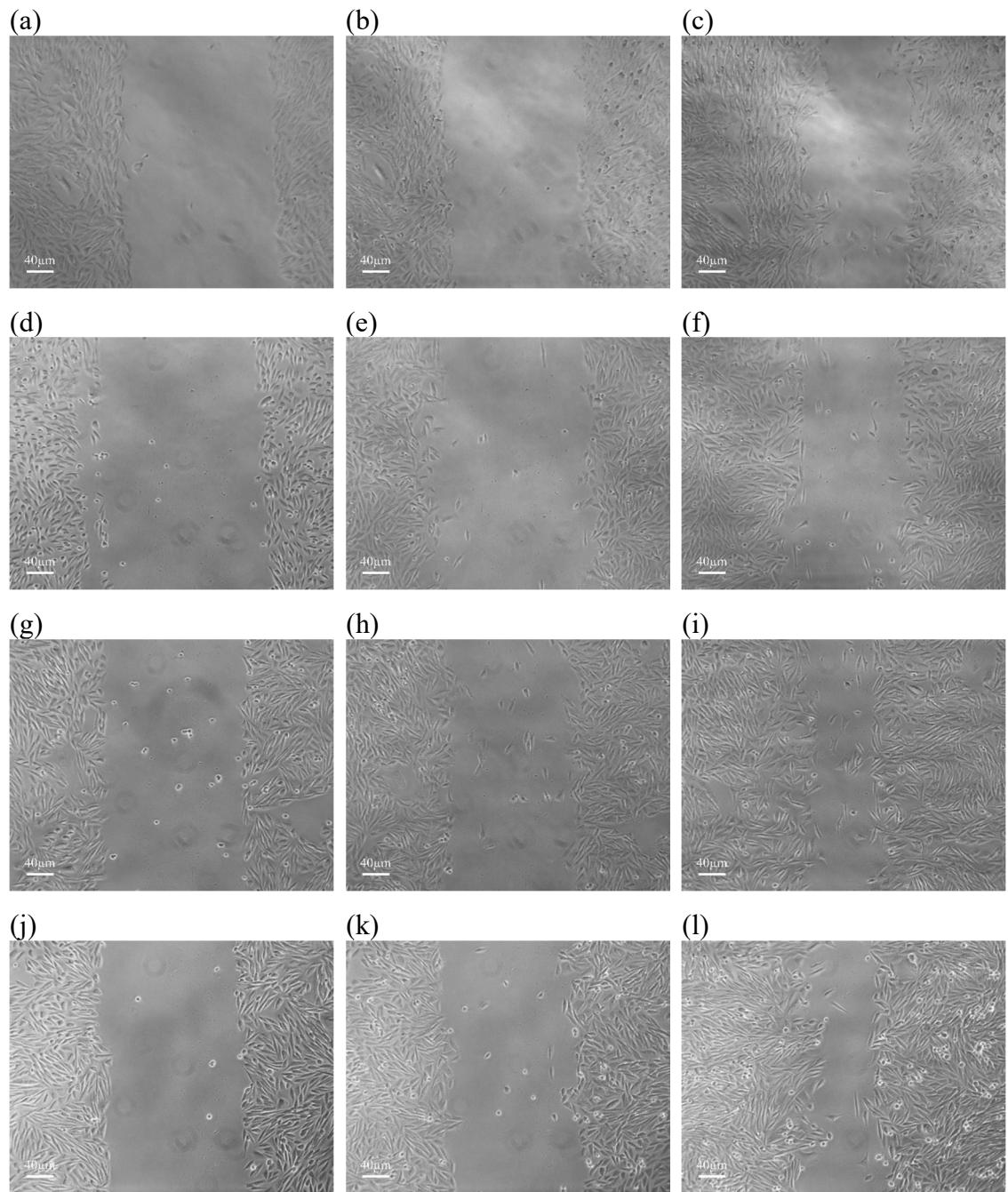


Figure S6: Microscopies images of a pseudo-wound created in a fibroblast cell L929 monolayer, at 0 hour (left column), at 10 hours (middle column), and at 24 hours (right column). The pseudo-wounds were treated with (a–c) PLGA, (d–f) PLGA SDS, (g–i) PLGA SDS/HP β CD, and (j–l) control group.