

Supplementary Information

Controlled surface mineralization of metal oxides on nanofibers

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Figure S1. SEM micrographs of the nanoparticles (a) PS, (b) P($S_{0.99}$ -*co*-VPA_{0.01}), (c) P($S_{0.96}$ -*co*-VPA_{0.04}), (d) P($S_{0.90}$ -*co*-VPA_{0.10}).

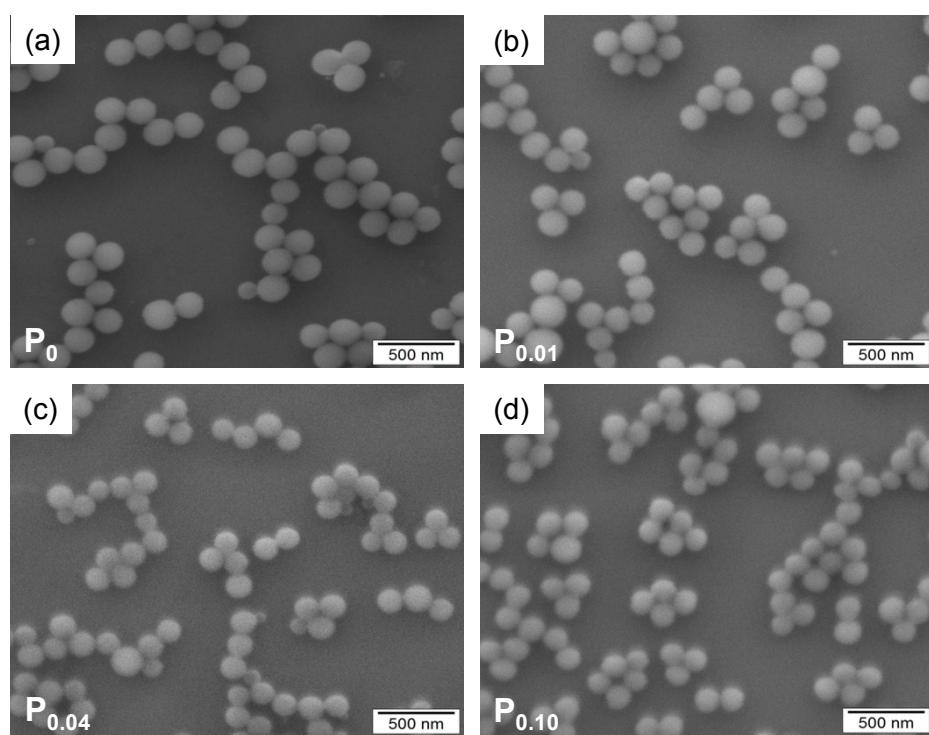


Figure S2. ^{31}P -NMR (202 MHz, DMF-d₆, 298K9 spectra of (a) P(S_{0.99}-*co*-VPA_{0.01}), (b) P(S_{0.96}-*co*-VPA_{0.04}), (c) P(S_{0.90}-*co*-VPA_{0.10}).

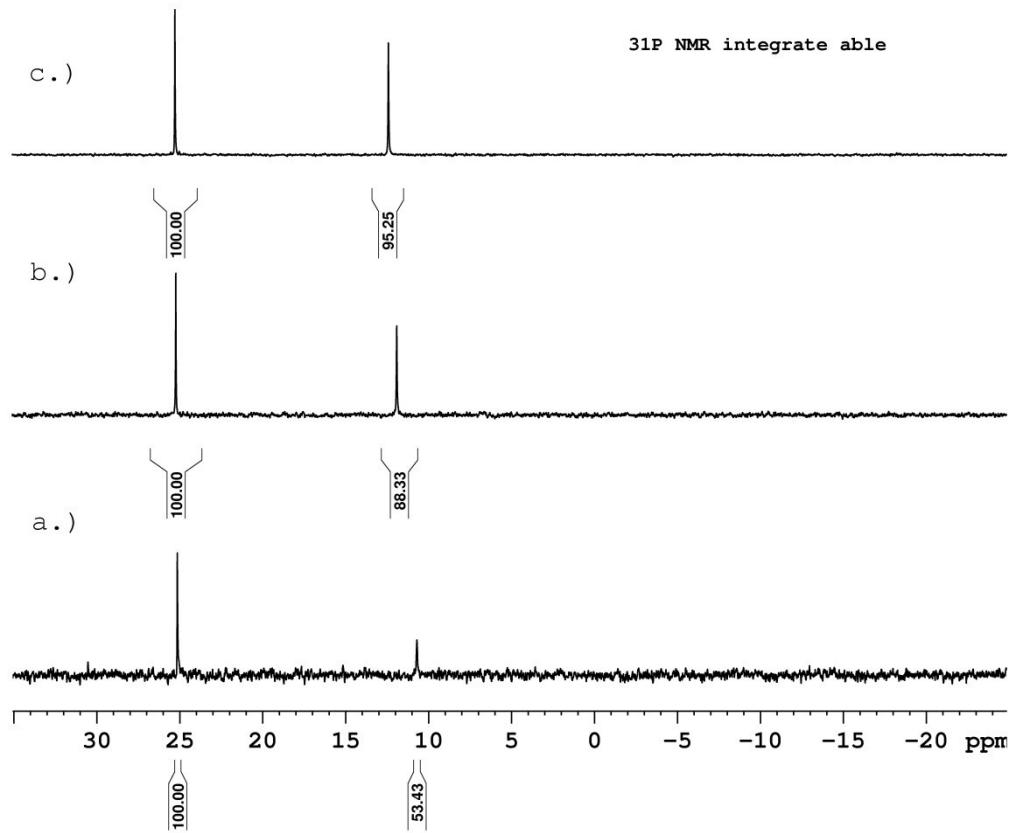


Figure S3. SEM micrographs of the electrospun fibers of (a) PS, (b) P(S_{0.99}-*co*-VPA_{0.01}), (c) P(S_{0.96}-*co*-VPA_{0.04}), (d) P(S_{0.90}-*co*-VPA_{0.10}).

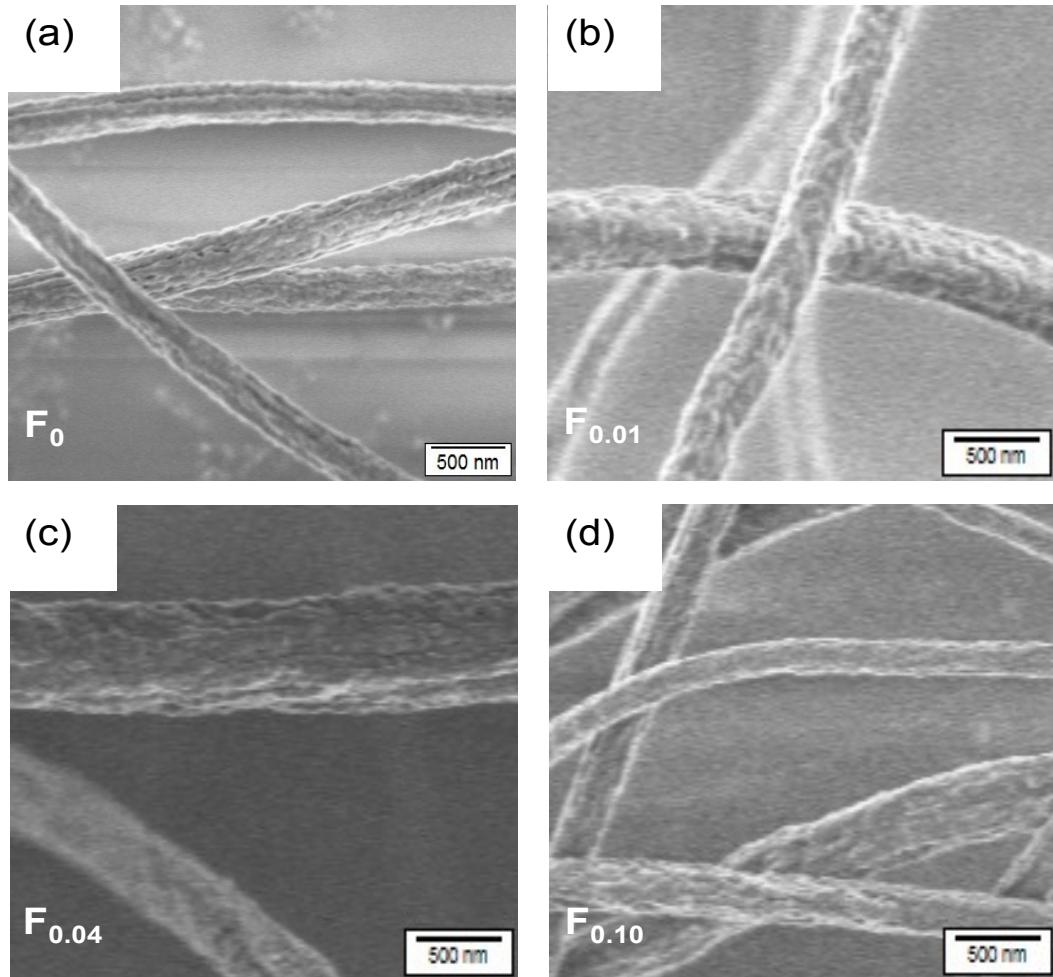


Figure S4. SEM micrographs of the electrospun fibers after ceria crystallization: (a) PS, (b) P(S_{0.99}-*co*-VPA_{0.01}), (c) P(S_{0.96}-*co*-VPA_{0.04}), (d) P(S_{0.90}-*co*-VPA_{0.10}).

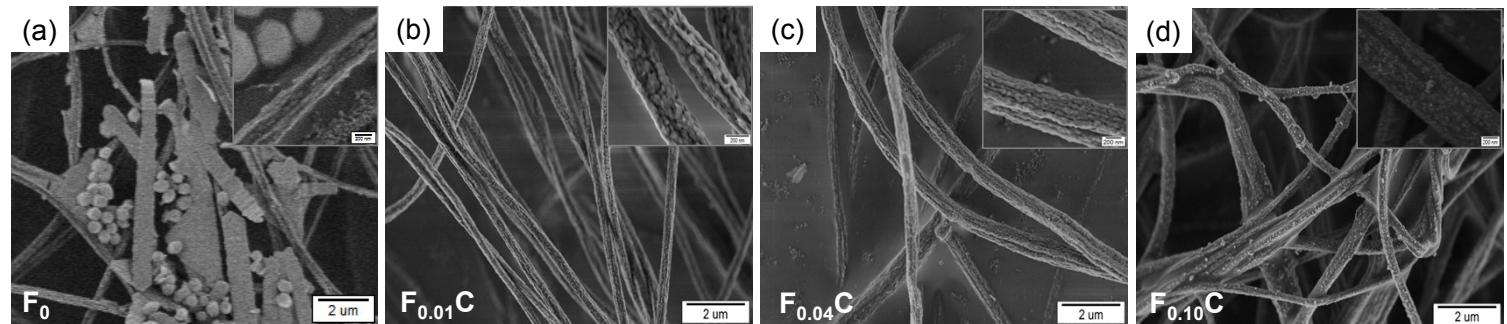


Figure S5. SEM micrograph showing the general view of the electrospun fibers after ceria crystallization of P(S_{0.99}-co-VPA_{0.01}).

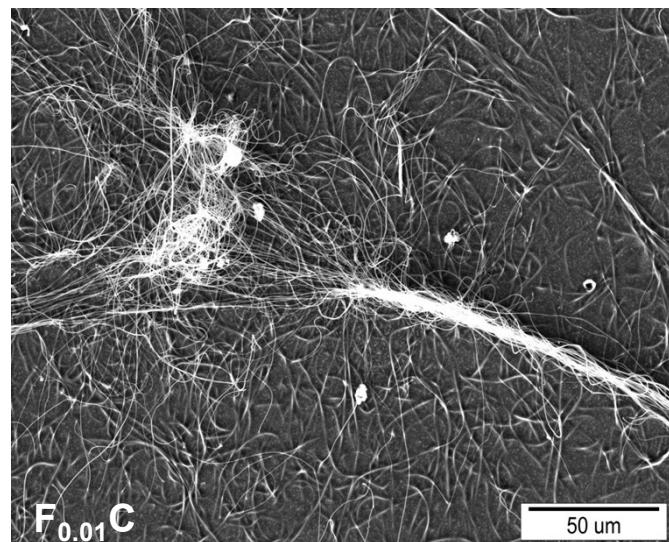


Figure S6. EDX spectrum and corresponding of the indicated area of a electrospun P(S_{0.96}-co-VPA_{0.04}) fibers after ceria crystallization.

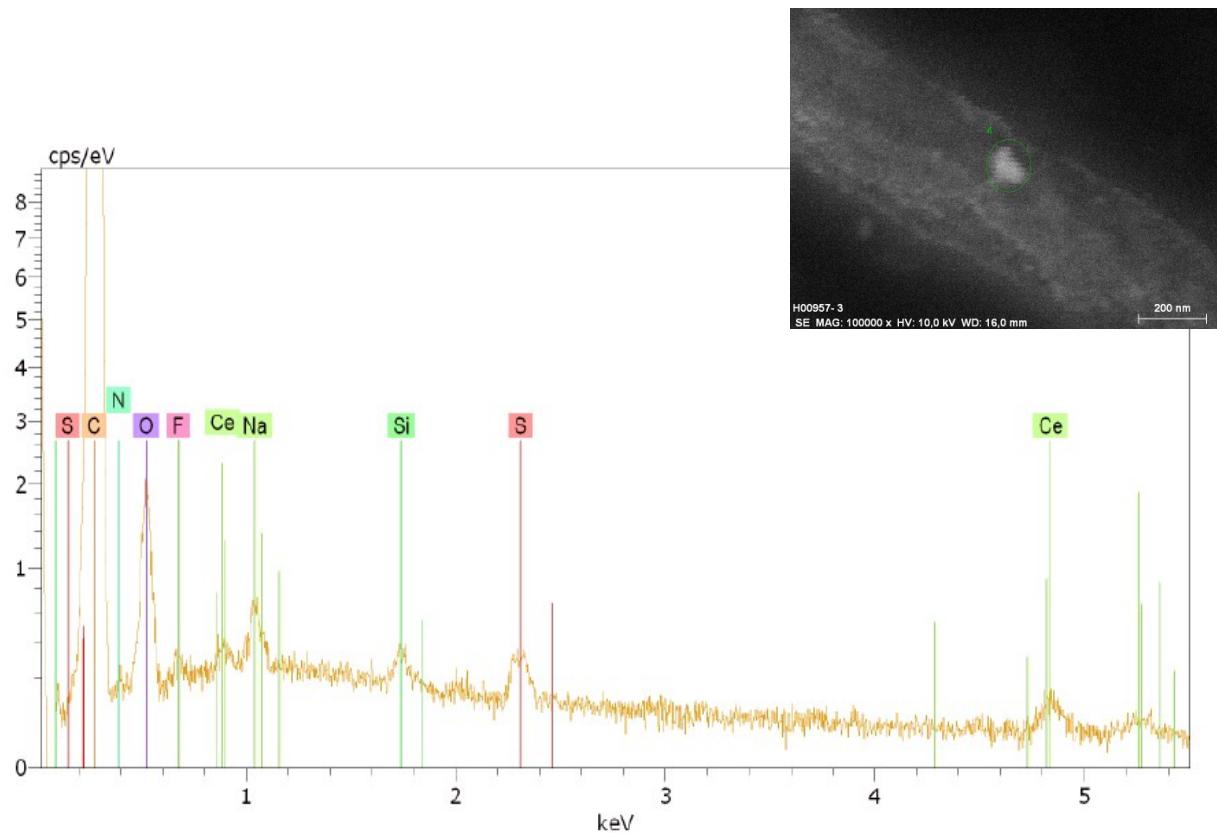


Figure S7. EDX spectrum and corresponding of the indicated area of a electrospun P(S_{0.96}-co-VPA_{0.04}) fibers after titania crystallization.

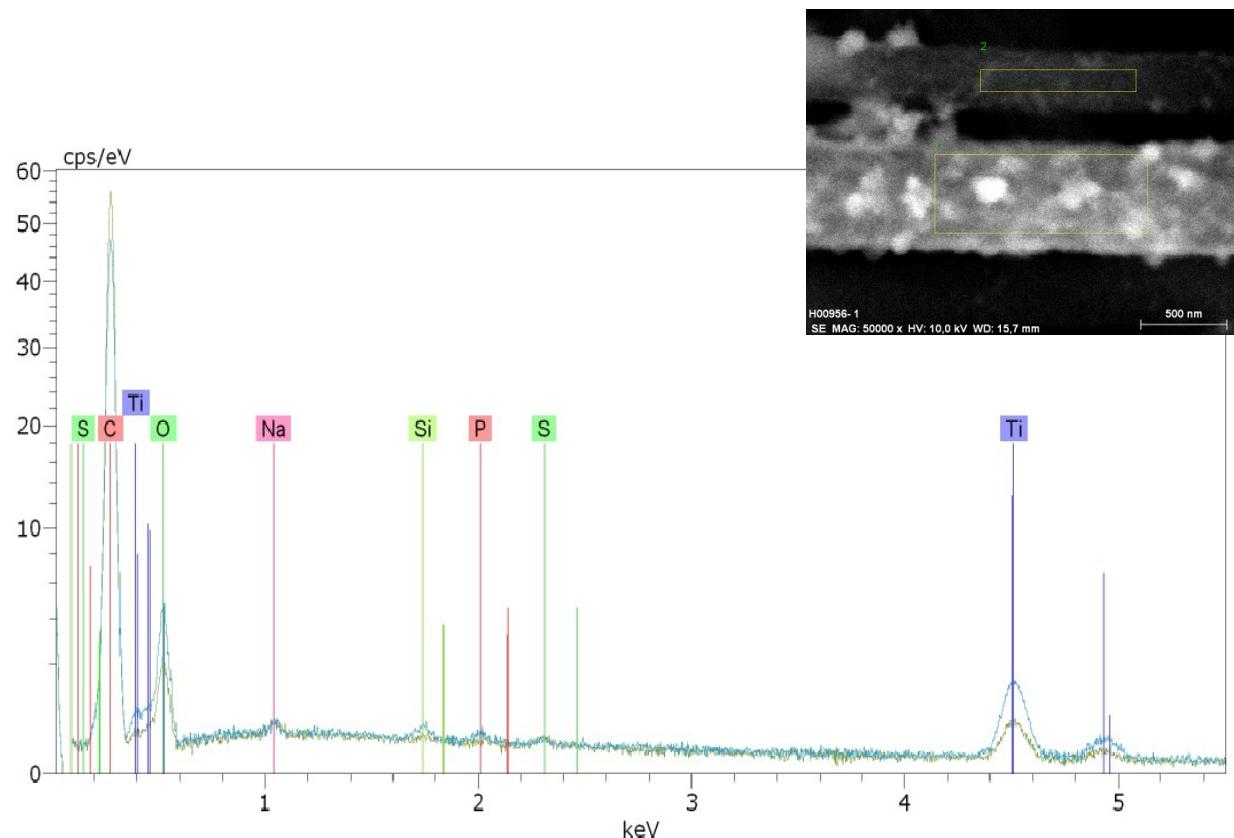


Table S1. Variation of average fiber diameter (AFD) of the electrospun fibers and molecular weight of the polymers as a function of comonomer concentration.

Sample Name	Mw (g/mol)	Sample Name	AFD (nm)
P₀	315,000	F₀	320 ±60
P_{0.01}	260,000	F_{0.01}	310 ±60
P_{0.04}	220,000	F_{0.04}	300 ±90
P_{0.10}		F_{0.10}	270 ±50

Table S2. Atomic concentrations of the fibers before and after ceria crystallization measured by XPS.

	C1s	O1s	P2p	Ce3d
F₀	100	0	0	NA
F₀C	91.7	7.2		1.1
F_{0.01}	96.1	3.9	0	0
F_{0.01}C	92.2	5.8	0.8	1.2
F_{0.05}	95.5	4.5	0	0
F_{0.05}C	84.0	11.1	3.4	1.5