Polarization behavior of bone marrow-derived macrophages on charged P(VDF-TrFE) coatings

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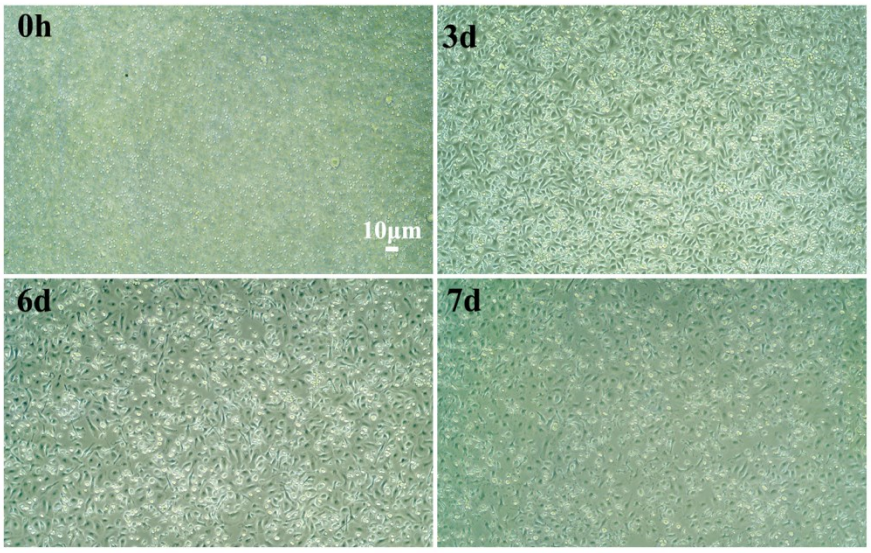

Figure S1. Induction of bone-marrow derived macrophages (BMDMs) by M-CSF.

Table S1. Primers used for real-time PCR.

Gene	Forward primer sequence (5'–3')	Reverse primer sequence (5'–3')
CD80	GCCTTGCCGTTACAACTC	TACTCGGGCCACACTTTT
TNF-α	CGCTGAGGTCAATCTGC	GGCTGGGTAGAGAATGGA
CD206	CAAGCGATGTGCCTACC	AATGCTGTGGATACTTGCC
IL-10	GCCCTTTGCTATGGTGTC	TCTCCCTGGTTTCTCTTCC
αM	CGCAAGAACACCAAGGA	TCAAAGAAGGCACGGATAC
β2	GGAGCATCGAGTATAGGCA	ACTGTATCCGTGAAACCCA
α5	ACTGCCACCTGCCAAGAT	GACTGTGCTGCTTGTCTGC
β1	AGCAACAATTCACCCACAG	TTCGAGACAGAGCAAGCA
Kv1.3	TGTGGATGGGAGGATGTT	CTTTGGGGAGTTTTCTGCT
actin	GCTCCGGCATGTGCAAAG	CCTTCTGACCCATTCCCAC

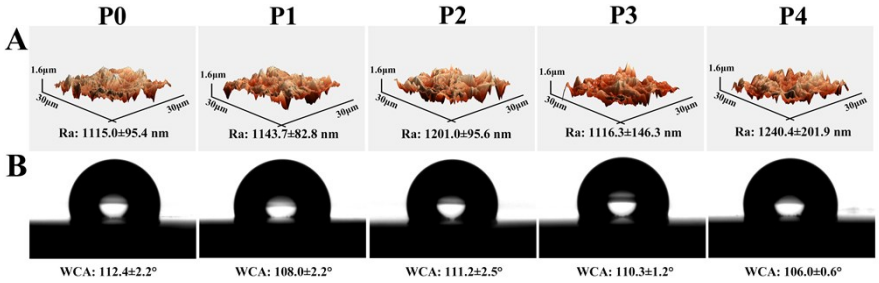

Figure S2. AFM images (A) and water contact angles (B) of the P(VDF-TrFE) coatings.

Table S2. Surface potential intensity of P(VDF-TrFE) coatings after electrically poled treatment.

Group	Surface potential (-mV)
P0	54
P1	65
P2	68
P3	71
P4	85

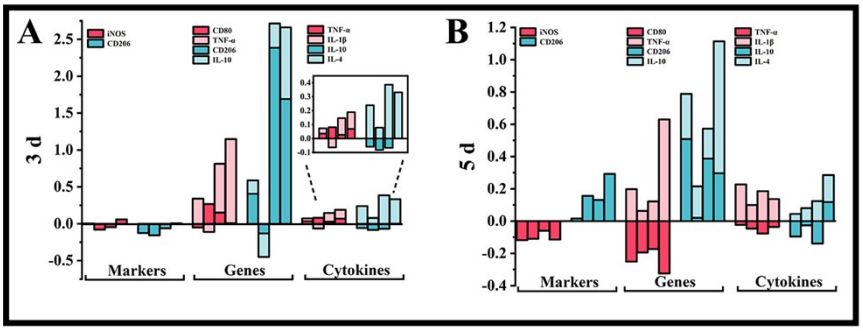

Figure S3. Semi-quantitative statistics of fluorescence intensity of macrophages surface markers and changes in gene expression and cytokines secretion of macrophages at 3 days (A) and 5 days (B) (each of the four bars represents the change range of P1, P2, P3, P4 compared with P0).

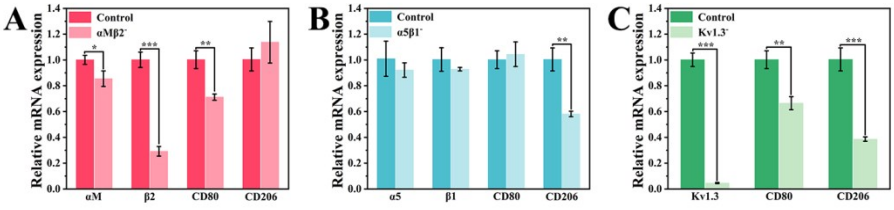

Figure S4. Effect of αMβ2 (A), α5β1 (B) and Kv1.3 (C) inhibition on mRNA expressions of corresponding integrin, potassium ion channel and phenotypic gene (CD80 and CD206) on P0 coatings.