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Supporting Information

Title

Enhanced Bone Regeneration via PHA Scaffolds Coated with Polydopamine-Captured BMP2

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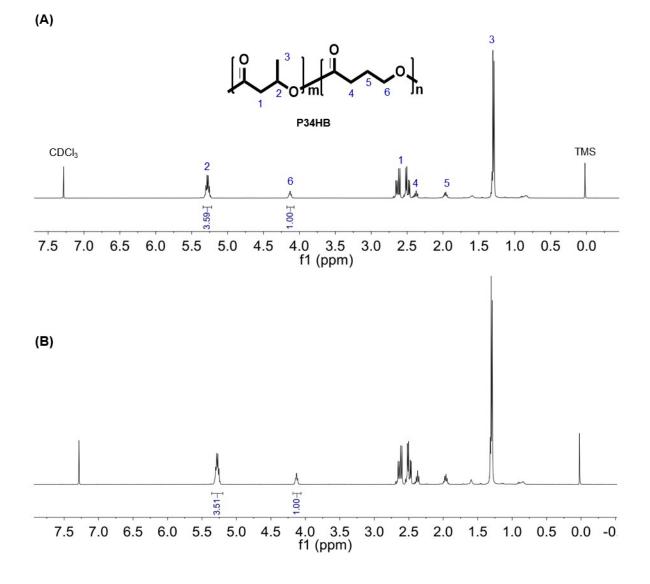


Figure S1. Structural analysis of PHA materials using ¹H Nuclear magnetic resonance (NMR). (A) P34HB raw materials were biosynthesized using engineered *H. bluephagenesis* TD; (B) P34HB materials were obtained after FDM.

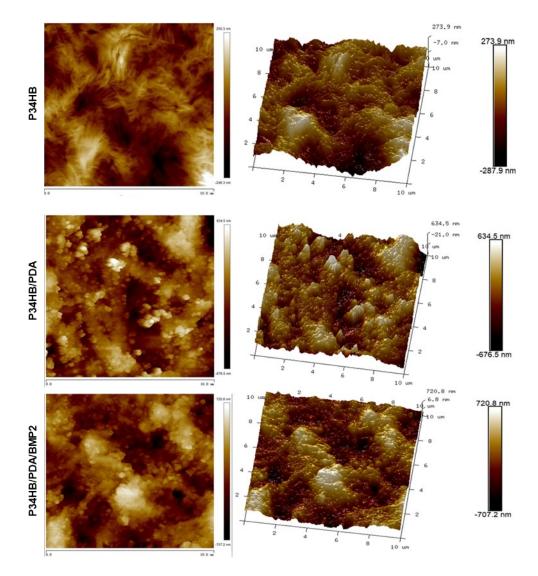


Figure S2. Surface morphology of different P34HB films determined by atom force microscope (AFM).



Figure S3. The hydrophility of P34HB films including P34HB, PDA-coated P34HB and BMP2-functionalized P34HB, as determined by a video optical contact angle measuring instrument.

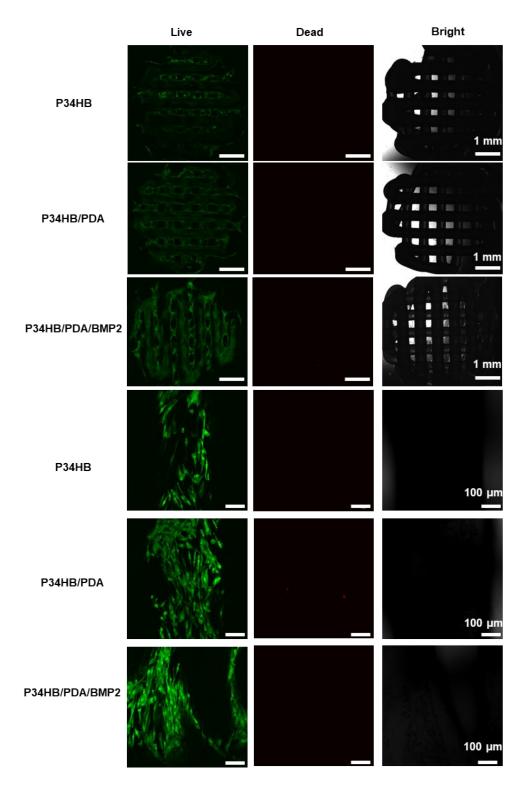


Figure S4. Representative confocal fluorescence images of ADSCs cocultured with scaffolds for day 7 and stained with LIVE/DEAD assay kit (live cells, green fluorescence; dead cells, red fluorescence).

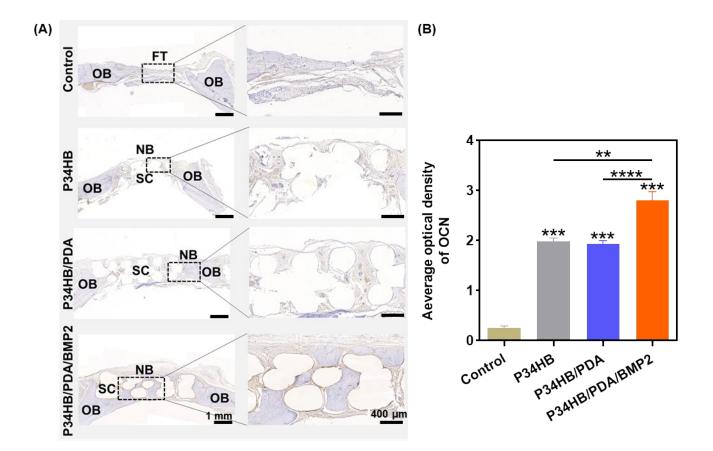


Figure S5. Histological staining and observation of bone tissue decalcified sections after 8 weeks of post-implantation. (A) Immunohistochemical staining of OCN; (B) Quantified average optical density at region of interest (ROI); OB, original bone; FT, fibrous tissue; NB, new bone; SC, scaffolds. The control group in the figure represented that there were no scaffolds. All data were showed as means \pm SD. **P < 0.01, ***P < 0.001, ****P < 0.0001, one-way analysis of variance (ANOVA) with Tukey's post hoc test.

 Table S1. Primer sequences of gene-related osteogenesis in RT-PCR.

Gene	forward	reverse
RUNX2	5'-CACTGGCGCTGCAACAAGA-3'	5'-CATTCCGGAGCTCAGCAGAAT-3'
Collal	5'-GCTTGGTCCACTTGCTTGA-3'	5'-GAGCATTGCCTTTGATTGCTG-3'
OCN	5'-TGTGGAGGGTTGTGGGTGT-3'	5'-TCAACTGGGGTGGGGTTTT-3'
OPN	5'-GATGAATCTGATGAACTGGT-3'	5'-GTGATGTCCTCGTCTGTAGCA-3'
GAPDH	5'-GCACCGTCAAGGCTGAGAAC-3'	5'-TGGTGAAGACGCCAGTGGA-3'