

Supplementary Information

Bacterial Cellulose-Polydopamine based Injectable Composite Hydrogel for
Enhanced Hemostasis in Acute Wounds

Kaushal R Shakya¹, Niranjana Chatterjee², Santosh K. Misra², Vivek Verma^{1,3,4,5*}

¹Department of Materials Science and Engineering, Indian Institute of Technology Kanpur, Kanpur-
208016, India.

²Department of Biological Sciences and Bioengineering, Indian Institute of Technology Kanpur,
Kanpur- 208016, India.

³Samtel Centre for Display Technologies, Indian Institute of Technology Kanpur, Kanpur- 208016,
India.

⁴National Centre for Flexible Electronics, Indian Institute of Technology Kanpur, Kanpur- 208016,
India.

⁵Centre of Environmental Science and Engineering, Indian Institute of Technology Kanpur, Kanpur-
208016, India.

corresponding author: vverma@iitk.ac.in

Table S1: Comparative description of previously reported hydrogels used for hemostatic outcomes in respect to preparation methods, advantages, and limitations.

| Method/Material | Advantages | Limitations | Ref. |
|---|---|---|--------------|
| Blue-light-activated double-network strategy AC/PDM Hydrogel: -Allyl cellulose (AC) -P (Dopamine methacrylamide) (PDM) Photo-initiator: lithiumphenyl-2,4,6-trimethylbenzoylphosphinate (LAP) | -Strong tissue adhesive strength -Fast gelation -Antibacterial properties -Tissue-like mechanical strength | - The use of AC and PDM prolongs and complicates the fabrication process. - The photo-initiator limited penetration depth may result in incomplete curing. material based on photo initiator can be delicate, limiting their robustness. | ¹ |
| One-step physical cross-linking method Quaternary ammonium chitosan /tannic acid (TA) Cross-linked by dynamic ionic bonds and hydrogen bonds between QCS and TA | -Superior reactive oxygen species scavenging activity -Broad-spectrum antibacterial ability -Rapid hemostatic capability | -The whole system is based on ionic and hydrogen bonds which may not provide full potential of material. | ² |
| Chitosan-based hydrogels Cross-linked by 3-(3,4-dihydroxyphenyl) propionic acid-modified chitosan (DCS) Sebacic acid-terminated polyethylene glycol modified by p-hydroxybenzaldehyde (PEGSH) | -Suitable stretchability (~780%) -High blood absorbability (1300% ± 50%) -Strong adhesion (~68.5 kPa) -Stretchable, self-adhesive, antibacterial | -Requires precise regulation of PEGSH proportion for favorable cytocompatibility -Involves extensive processing and complex modifications. | ³ |
| Incorporation of Keratin–catechin nanoparticles (KE-NPs) into cellulose hydrogel -Sodium carboxymethyl cellulose -Keratin–catechin nanoparticles (KE-NPs) | -Enhances mechanical properties -Provides antioxidant and antimicrobial properties | -Nanoparticles agglomeration poses Dispersion challenges. -Stirring the mixture in the dark under nitrogen protection at 37 °C makes the process complex. -Adding nanoparticles suspension with 0-2.5% w/w concentration complicates the weighing and solution preparation. | ⁴ |
| Current work: BC/PDA/CMC hydrogel -Bacterial cellulose (BC) -Polydopamine (PDA) -Carboxymethyl cellulose (CMC) | -Simple synthesis method -Easy storage conditions -No complex procedures used -Good hemostatic and antioxidant property | -less adhesive nature | - |

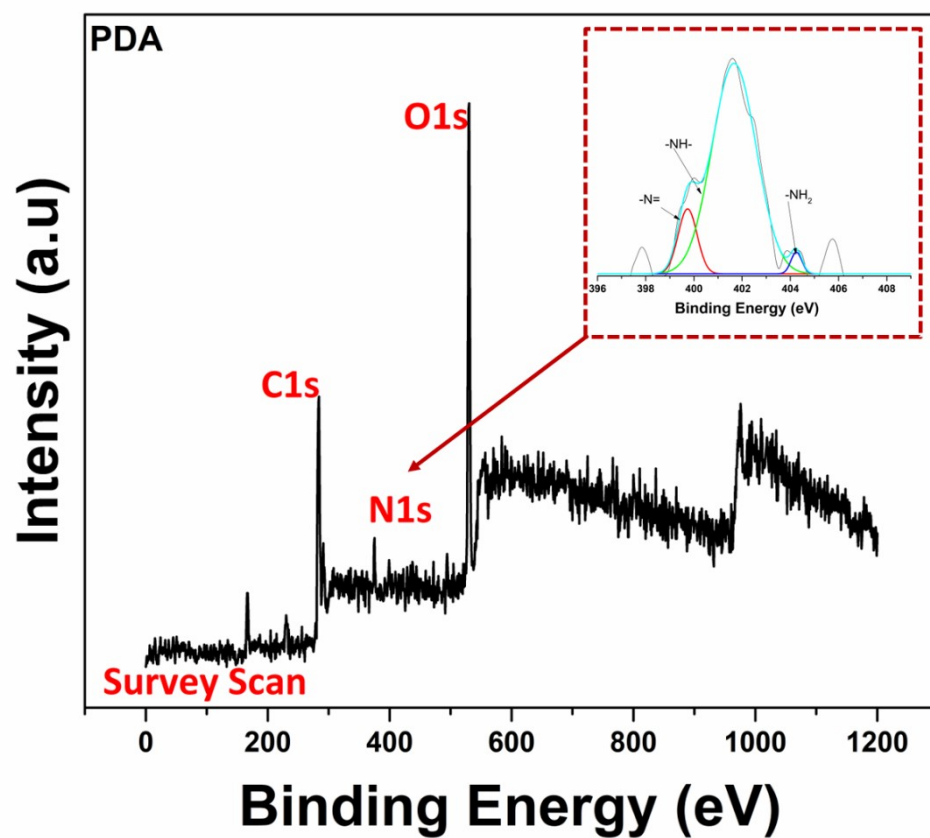


Figure S1: XPS survey spectra of polydopamine (PDA) with inset showing deconvoluted peak of nitrogen.

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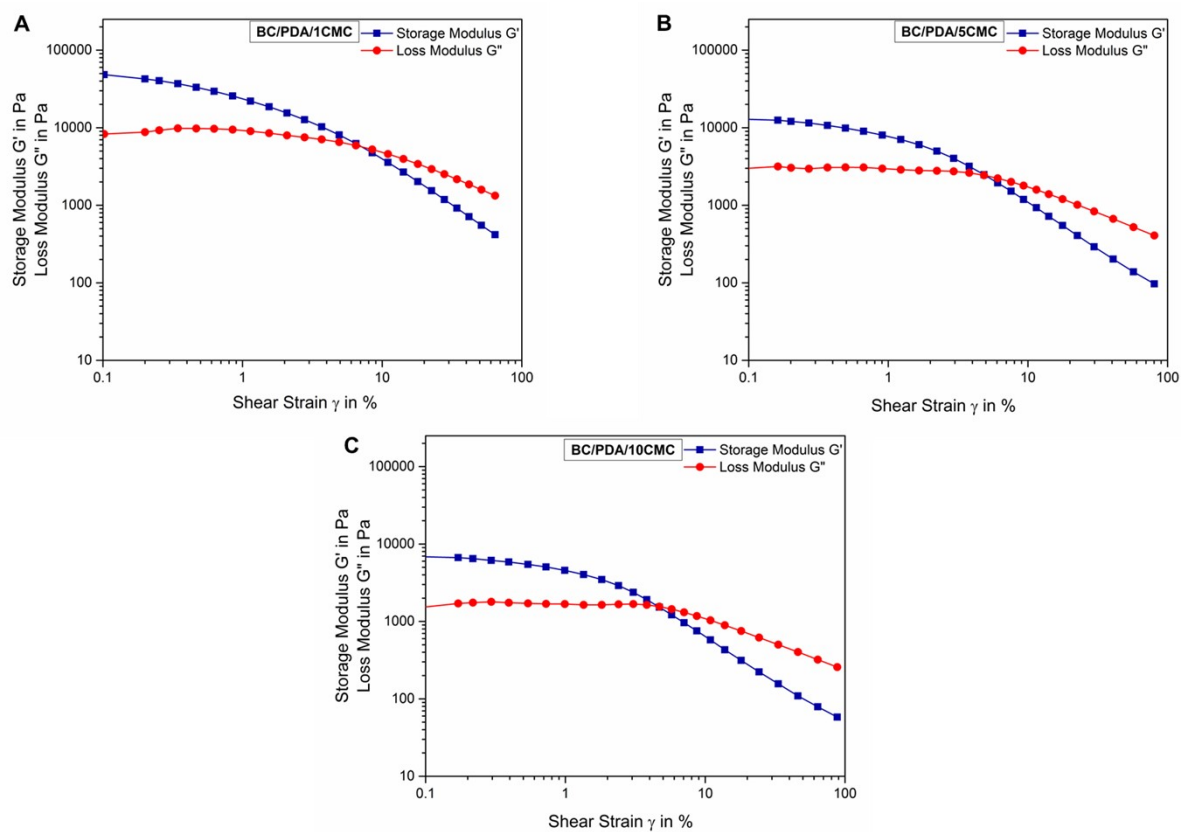


Figure S2: Strain sweep graph of (A) BC/PDA/1CMC (B) BC/PDA/5CMC and (C) BC/PDA/10CMC samples.

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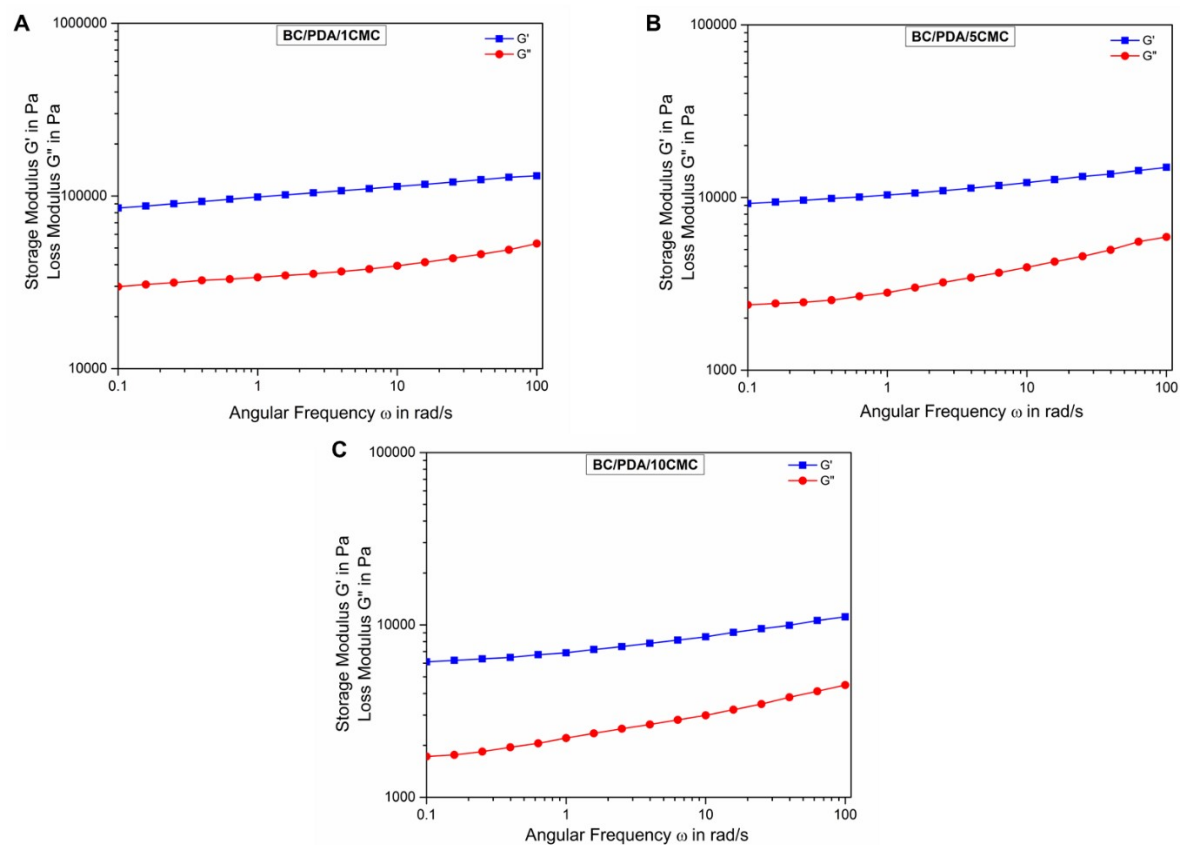


Figure S3: Frequency sweep graph of (A) BC/PDA/1CMC (B) BC/PDA/5CMC and (C) BC/PDA/10CMC samples.

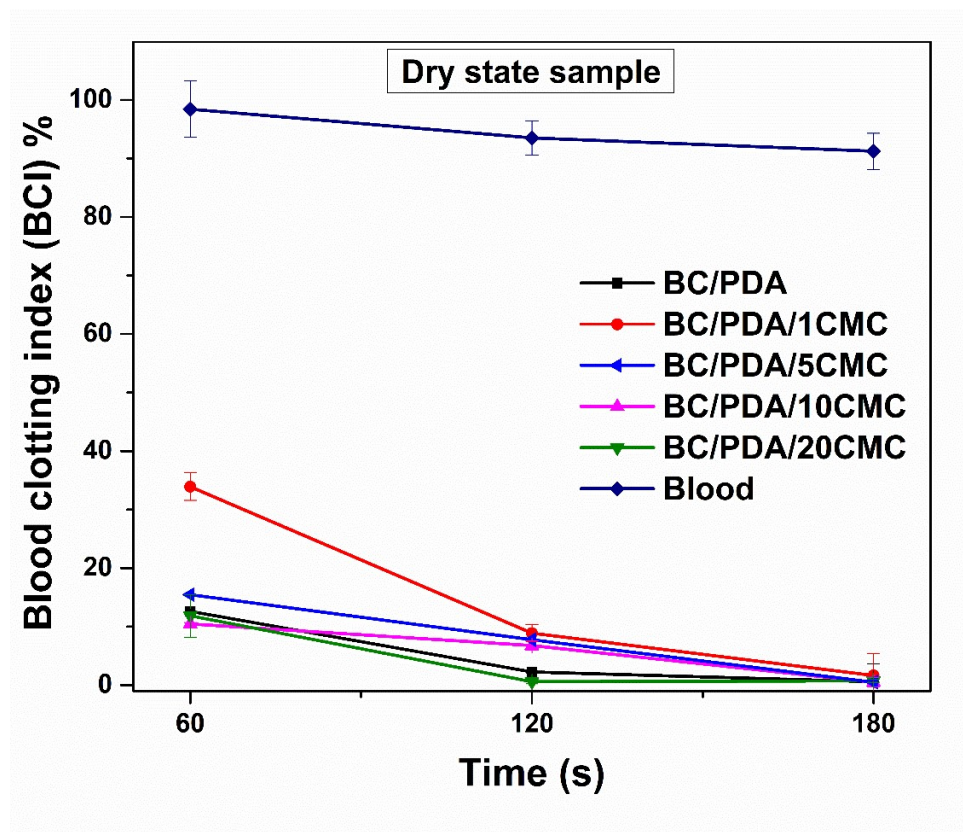


Figure S4: *In vitro* blood clotting study showing blood clotting index (BCI) at different time points on dry from of samples.

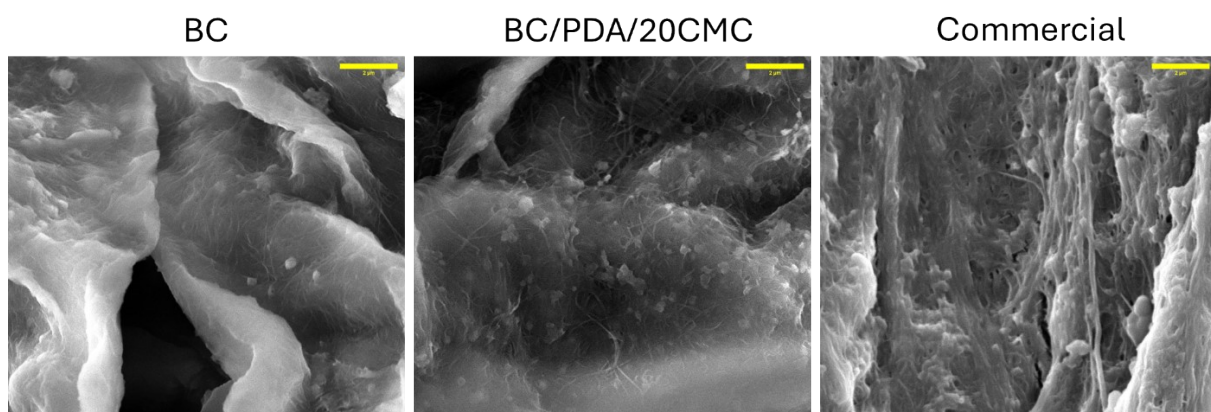


Figure S5: SEM images of platelet adhesion on samples at 7000 \times magnification. of BC, BC/PDA/20CMC, and commercial. Scale bar is equal to 2 μ m.

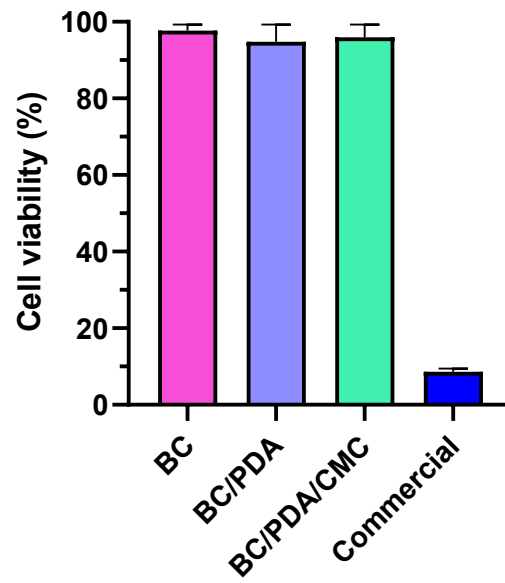


Figure S6: % Cell viability of NIH3T3 cells in presence of samples and controls using MTT assay.

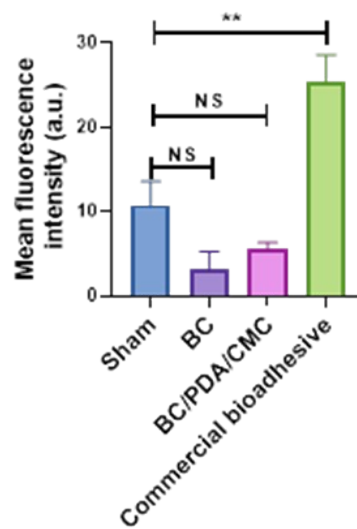


Figure S7: Semi-quantitative analysis of the expression of TNF- α (red).

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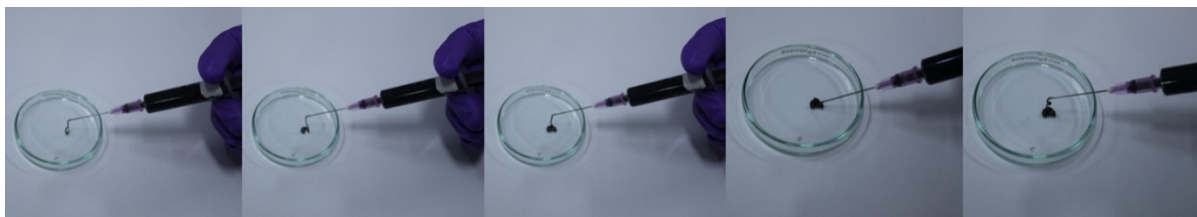


Figure S8: BC/PDA/20CMC hydrogel injecting through 2 mL syringe (23G).

References

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