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Supporting Information

Sonochemistry-assisted Reversible Addition-

Fragmentation Transfer Polymerization for Multicolor

Room-temperature Phosphorescent Polymers

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Figure S1. Conversion analysis by ¹H-NMR (DMSO-d₆) of the polymerization of PMAA-

CTA.



Figure S2. The GPC trace of PMAA-CTA.



Figure S3. The GPC trace of control group.



Figure S4. Conversion analysis by ¹H-NMR (DMSO-d₆) of the polymerization of MAA and

BA under ultrasound.



Figure S5. Conversion analysis by ¹H-NMR (DMSO-d₆) of the polymerization of MAA and

BA under control group.



Figure S6. Particle size testing at different ultrasound times.



Figure S7. Number-average molecular weight and size of ultrasound group.



Figure S8. Photos of RhB/GCP with different doping concentrations under sunlight, UV

irradiation, and UV off.



Figure S9. (a) CIE coordinates of afterglow luminescence of RTP materials with different

doping ratios, and (b) phosphorescence spectra.



Figure S10. (a) CIE coordinates of afterglow luminescence of RTP materials with different

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Figure S11. (a) RTP polymer color palette with different RhB/Fluc doping ratios. (b) RTP

polymer solutions with different RhB/Fluc doping ratios (top) and thin films (bottom).

Scheme S1. Synthesis of CTA



Scheme S2. Synthesis of Cz-OH



Scheme S3. Synthesis of RAFT agents



Scheme S4. Synthesis of GCP





Figure S12. ¹H-NMR spectrum of the Bis-(dodecyl thioalkyl thiocarbonyl) disulfide in CDCl₃.



Figure S13. ¹H-NMR spectrum of the CTA in CDCl₃.



Figure S14. ¹³C-NMR spectrum of the CTA in CDCl₃.



Figure S15. ¹H-NMR spectrum of Cz-OH in DMSO.



Figure S16. ¹³C-NMR spectrum of Cz-OH in CDCl₃.



Figure S17. ¹H-NMR spectrum of RAFT agents in DMSO.



Figure S18. ¹³C-NMR spectrum of RAFT agents in CDCl₃.



Figure S19. ¹H-NMR spectrum of PMAA-CTA agents in DMSO.