

Supplementary Information for

Macrophage-Mediated Delivery of Microcapsules for Enhanced Photodynamic Therapy of Colon Cancer

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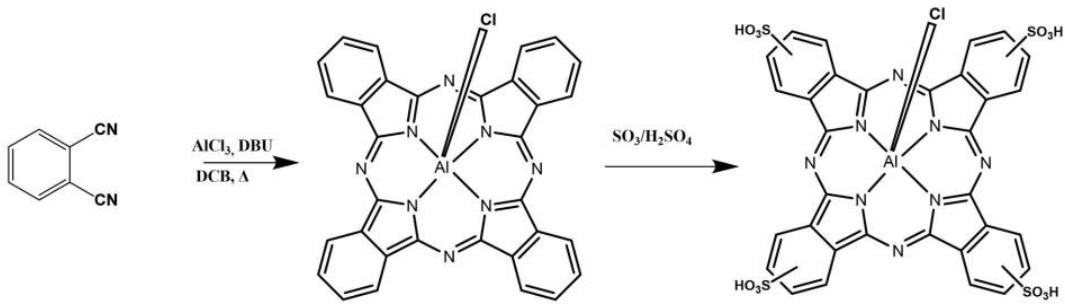


Fig. 1S. Synthesis scheme of PS.

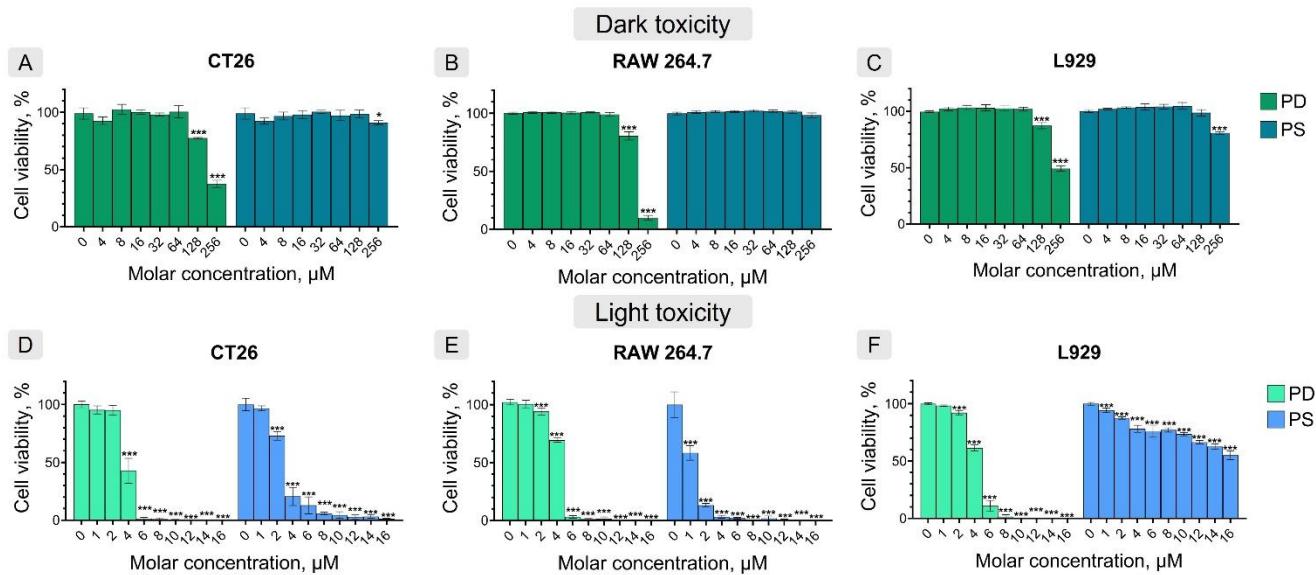


Fig. 2S. Dark toxicity (A, B, C) and light toxicity (D, E, F) assessments of photosensitizers PD and PS. Data are presented as mean \pm SD, n = 5, (*), (**), and (***)) indicate groups that are significantly different from control (0 μ M) with p < 0.05, p < 0.01, and p < 0.001, respectively.

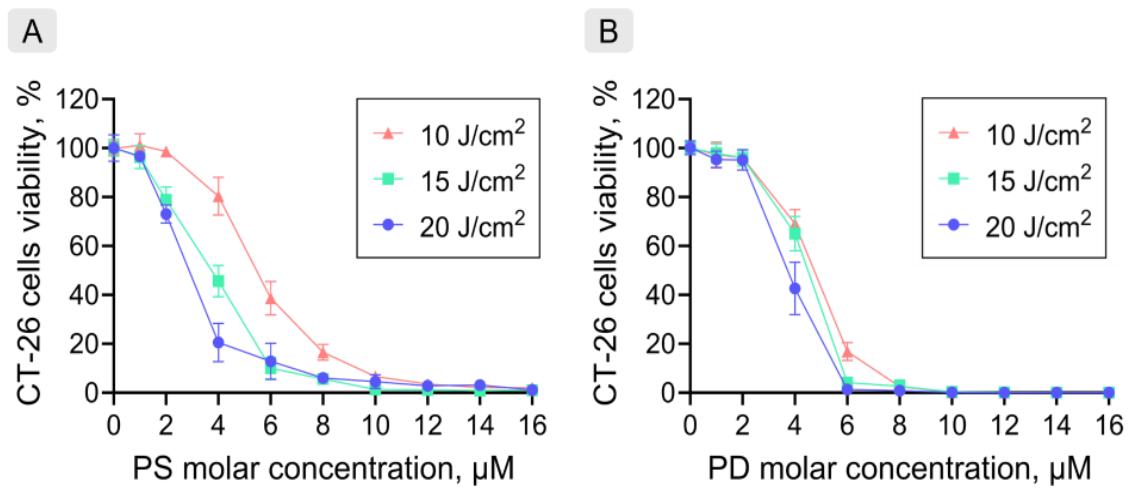


Fig. 3S. Assessment of light-induced toxicity of (A) PS and (B) PD at different energy densities (10, 15, and 20 J/cm²) on the CT-26 cell line. Results represent the mean \pm standard deviation of samples measured in five repetitions.

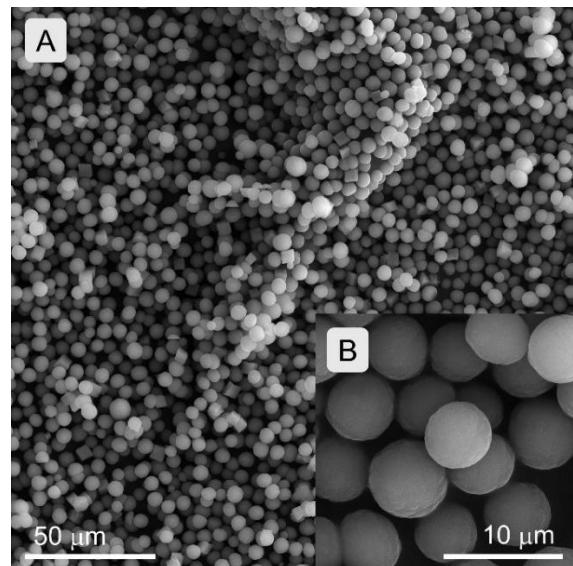


Fig. 4S. SEM images of vaterite cores at different magnifications: (A) 3.2 k \times ; (B) 10 k \times .

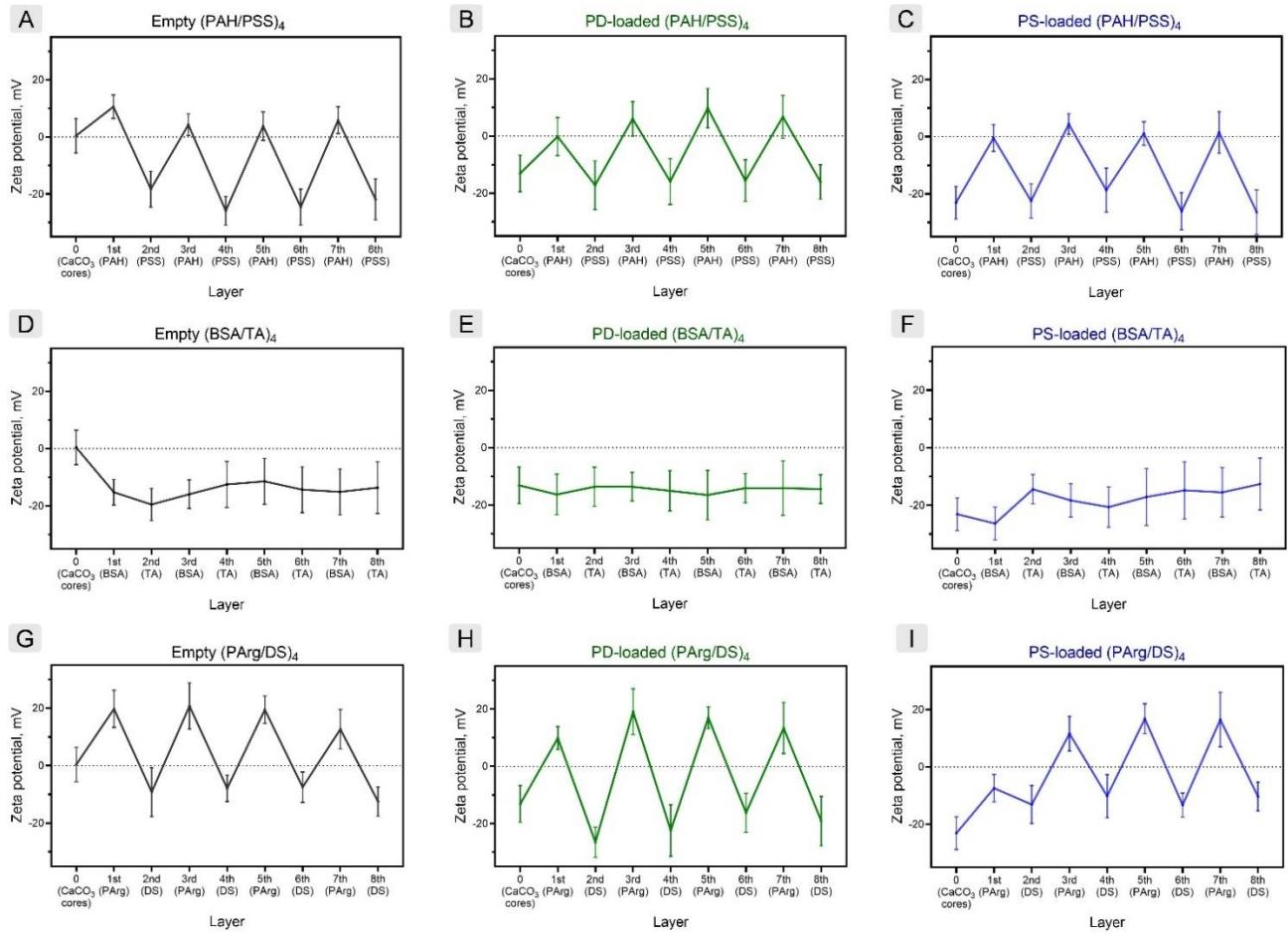


Fig. 5S. ζ -Potential profiles as a function of deposited layer number for different capsule formulations. Measurements were performed starting from either empty vaterite cores or dye-loaded cores, followed by sequential deposition up to the 8th polyelectrolyte layer (prior to core dissolution). (A) Empty (PAH/PSS)₄ system; (B) PD-loaded (PAH/PSS)₄; (C) PS-loaded (PAH/PSS)₄; (D) Empty (BSA/TA)₄; (E) PD-loaded (BSA/TA)₄; (F) PS-loaded (BSA/TA)₄; (G) Empty (PArg/DS)₄; (H) PD-loaded (PArg/DS)₄; (I) PS-loaded (PArg/DS)₄.

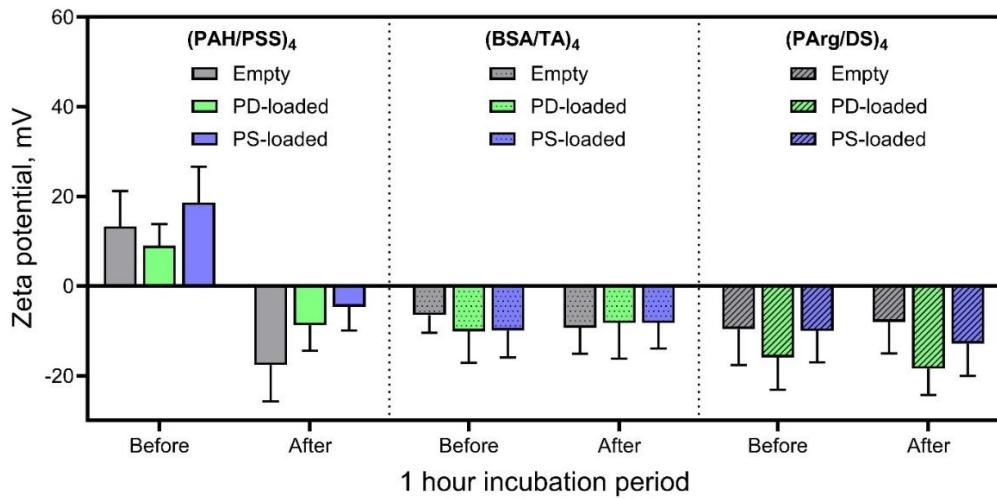
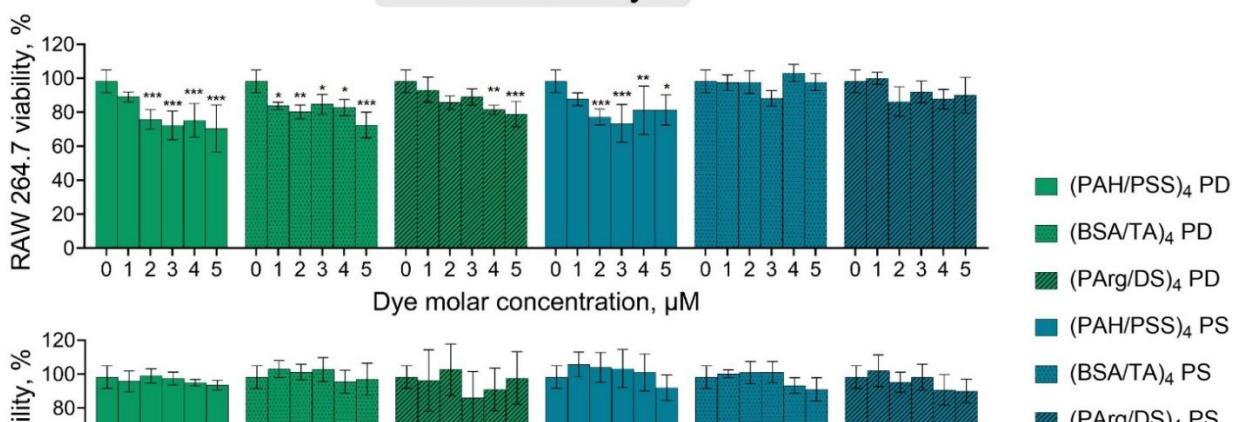


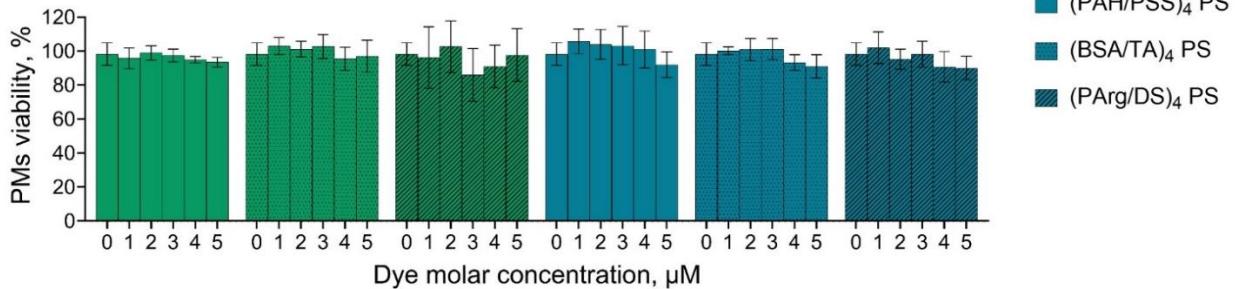
Fig. 6S. ζ -Potential values of microcapsules (after core dissolution) measured in deionized water before and after 1 h incubation in RPMI medium containing 10% FBS. Data are shown for (PAH/PSS)₄, (PArg/DS)₄, and (BSA/TA)₄ microcapsules, highlighting the effect of serum protein adsorption on surface charge.

A

Dark toxicity

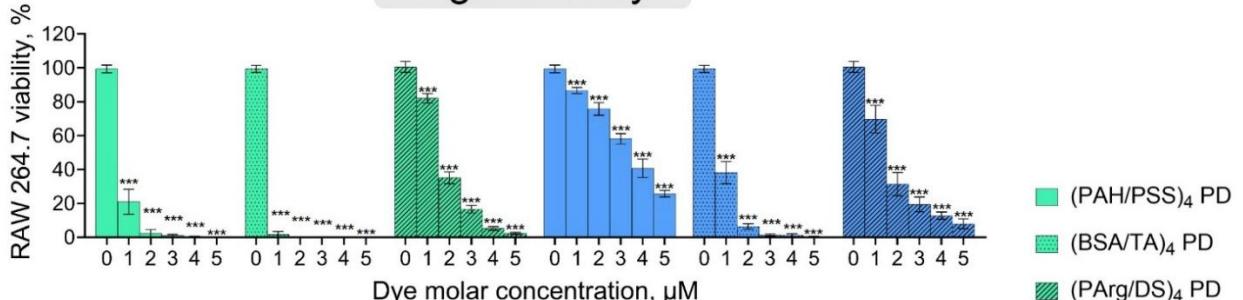


B



C

Light toxicity



D

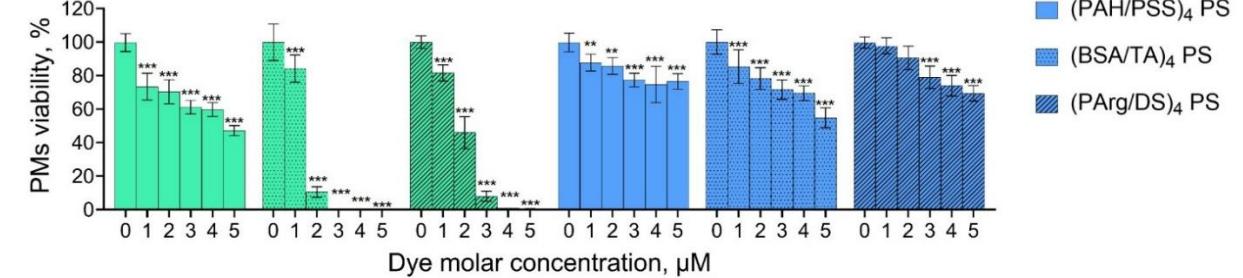


Fig. 7S. Evaluation of cytotoxicity for different microcapsule types loaded with photosensitizers (PS or PD) in macrophages. (A) Dark toxicity for RAW 264.7 cells. (B) Dark toxicity for PMs. (C) Light-induced toxicity for RAW 264.7 cells. (D) Light-induced toxicity for PMs. Data are presented as mean \pm SD, $n = 5$, (*), (**), and (***) indicate groups that are significantly different from control (0 μ M) with $p < 0.05$, $p < 0.01$, and $p < 0.001$, respectively.

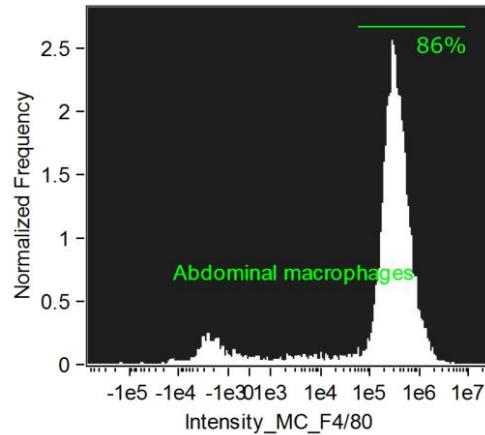


Fig. 8S. Intensity of F4/80 antibody staining in the primary culture of peritoneal macrophages obtained from the abdominal cavity of mice after cultivation and washing.

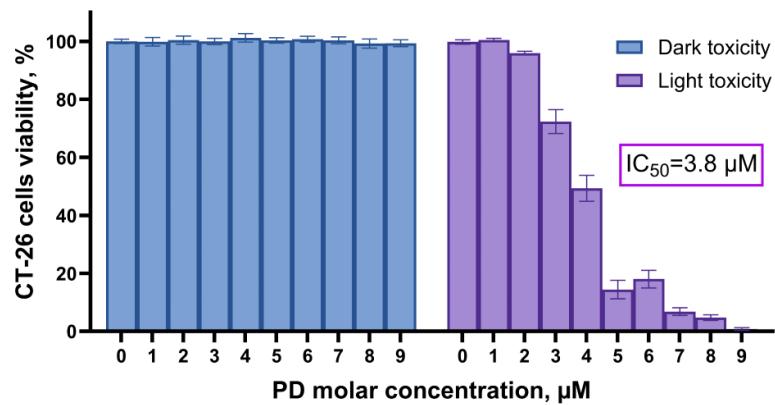


Fig. 9S. Dark and light cytotoxicity of (PArg/DS)₄ microcapsules loaded with PD against CT-26 cells. Results are presented as mean \pm SD (n = 5).

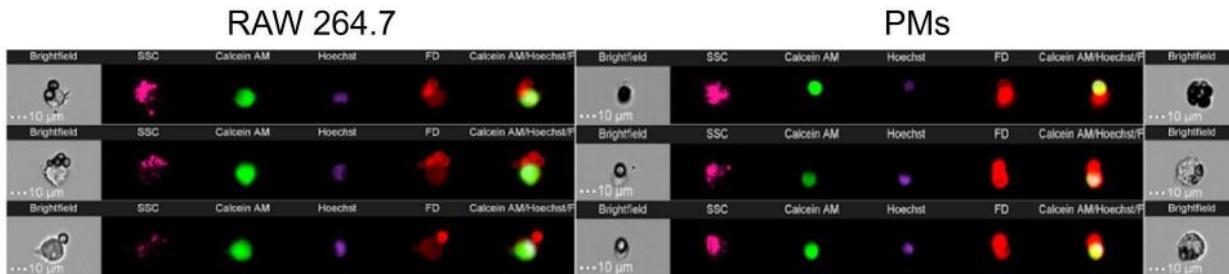


Fig. 10S. Visualization of RAW 264.7 cells and PMs with microcapsules using flow cytometry.

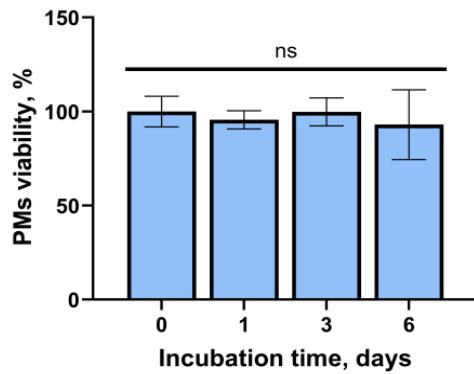


Fig. 11S. Long-term viability of peritoneal macrophages (PMs) exposed to (PArg/DS)₄ microcapsules loaded with Photoditazine (PD), assessed using the AlamarBlue assay on cultivation days 1, 3, and 6. Data are presented as mean \pm SD ($n = 5$), one-way ANOVA followed by Tukey's post hoc test.

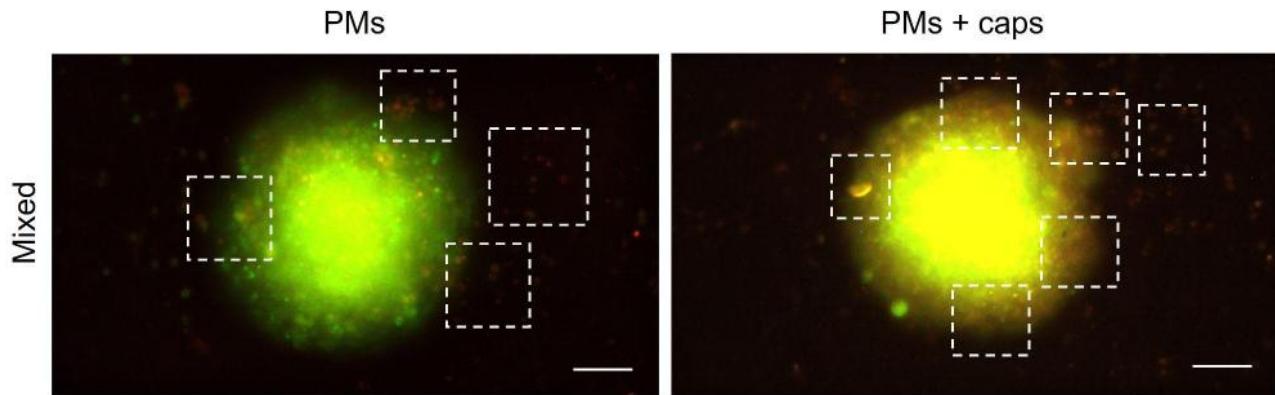


Fig. 12S. Detailed fluorescence images of PM migration to CT-26 spheroids. White dotted squares indicate visible clusters of macrophages. Scale bar: 100 μ m.