

Figure S1. Characterization of CPM by UHPLC QE HF-X MS. (A) Total ion chromatogram of CPM. (B) Retention times, names and chemical structures of these 11 characterized compounds.

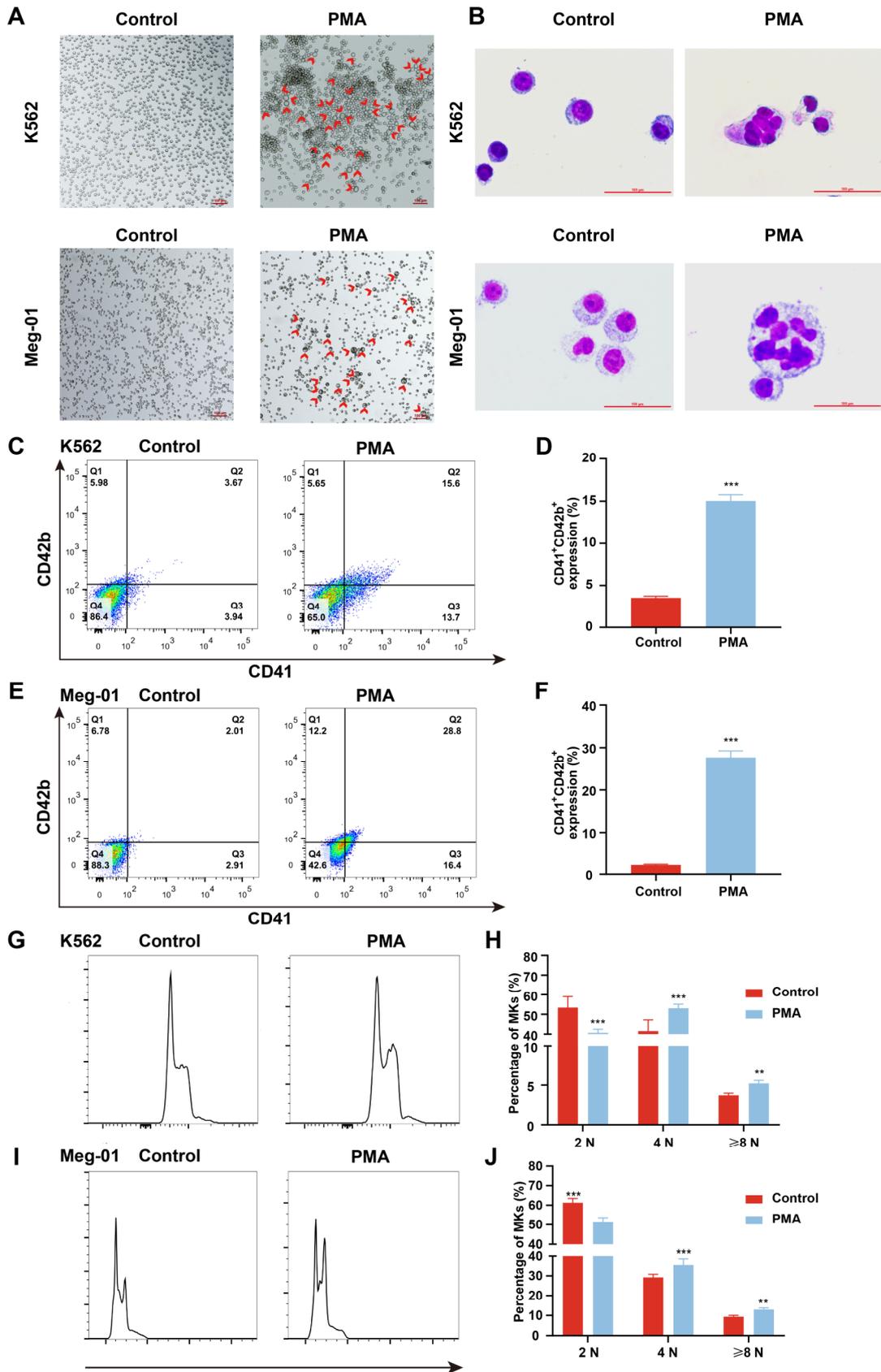


Figure S2. MK differentiation of K562 and Meg-01 cells induced by PMA. (A) Microscope photographs of K562 and Meg-01 cells with or without PMA treatment (1 nM) for 5 days were randomly captured at 10× resolution under the inverted light microscope. Scar bar: 100 μm. (B)

Giemsa staining of K562 and Meg-01 cells treated with or without PMA (1 nM) for 5 days. Scar bar: 100 μ m. (C, E) CD41 and CD42b expression of K562 and Meg-01 cells with or without PMA (1 nM) treatment for 5 days. (D, F) The proportion of CD41⁺CD42b⁺ cells in control and PMA-treated groups. Data are mean \pm SD (n=3). (G, I) DNA ploidy analysis of K562 and Meg-01 cells with or without PMA (1 nM) treatment for 5 days. (H, J) The percentage of 2N, 4N and \geq 8N cells in control and PMA-treated group. Data are mean \pm SD (n=3, ANOVA). * p < 0.05, ** p < 0.01, *** p < 0.001 vs. the control group.