Supplementary information for

Loss of lysyl hydroxylase 2 activates CRP-mediated cancer invasion through MEK/ERK/MMP9 signaling in the bone-microenvironment

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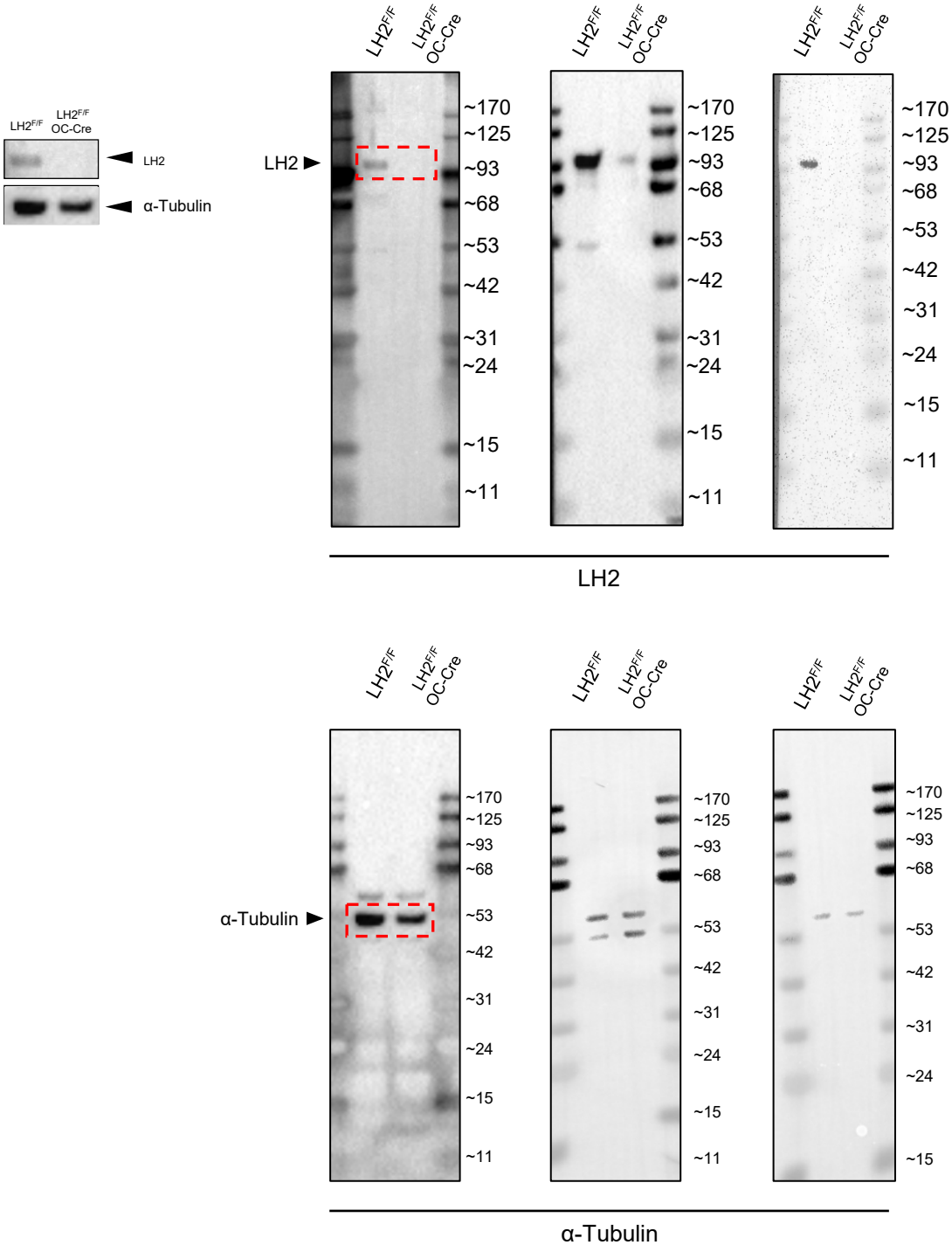
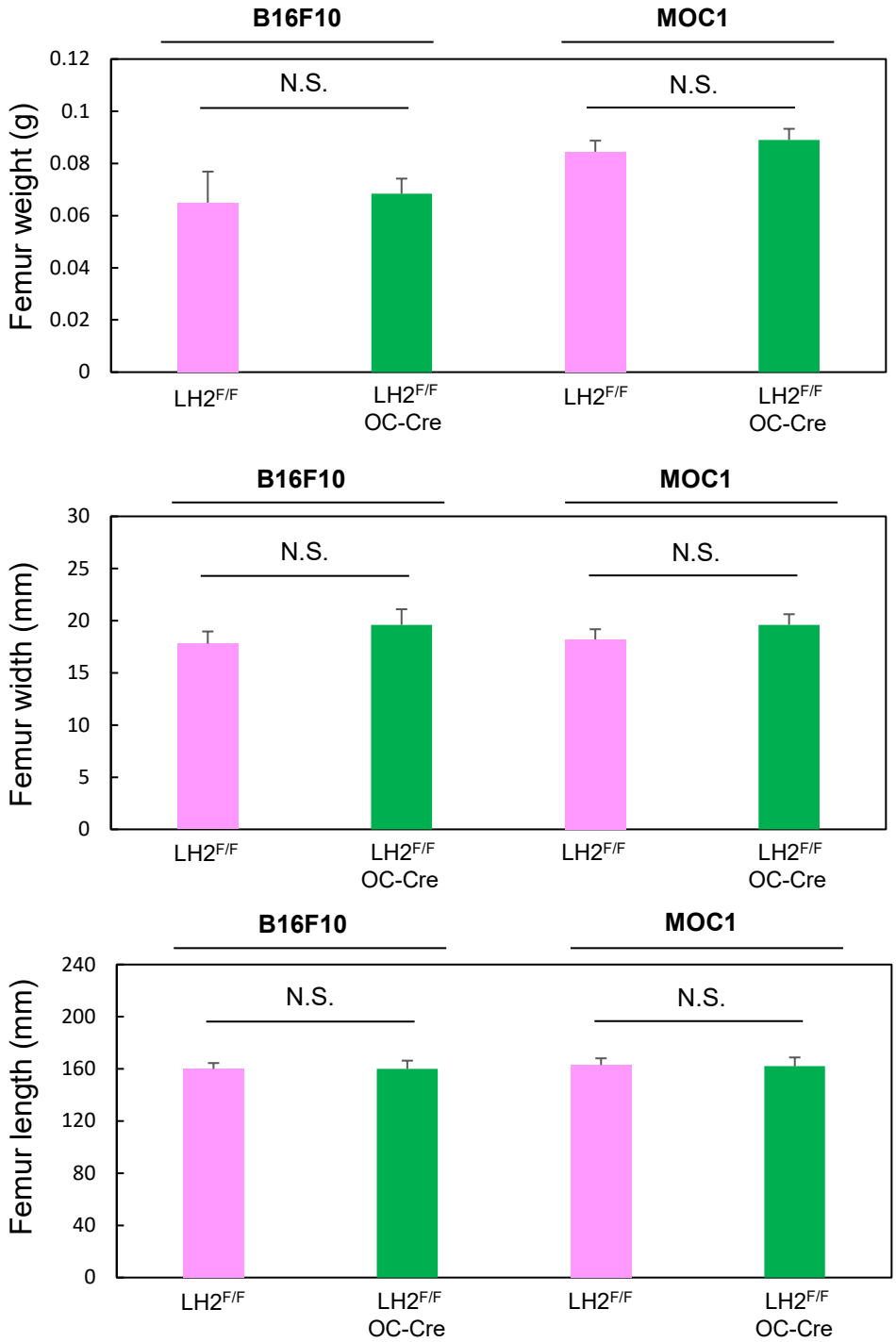
Supplemental Figure S1
(Full length blots of Figure 1B)



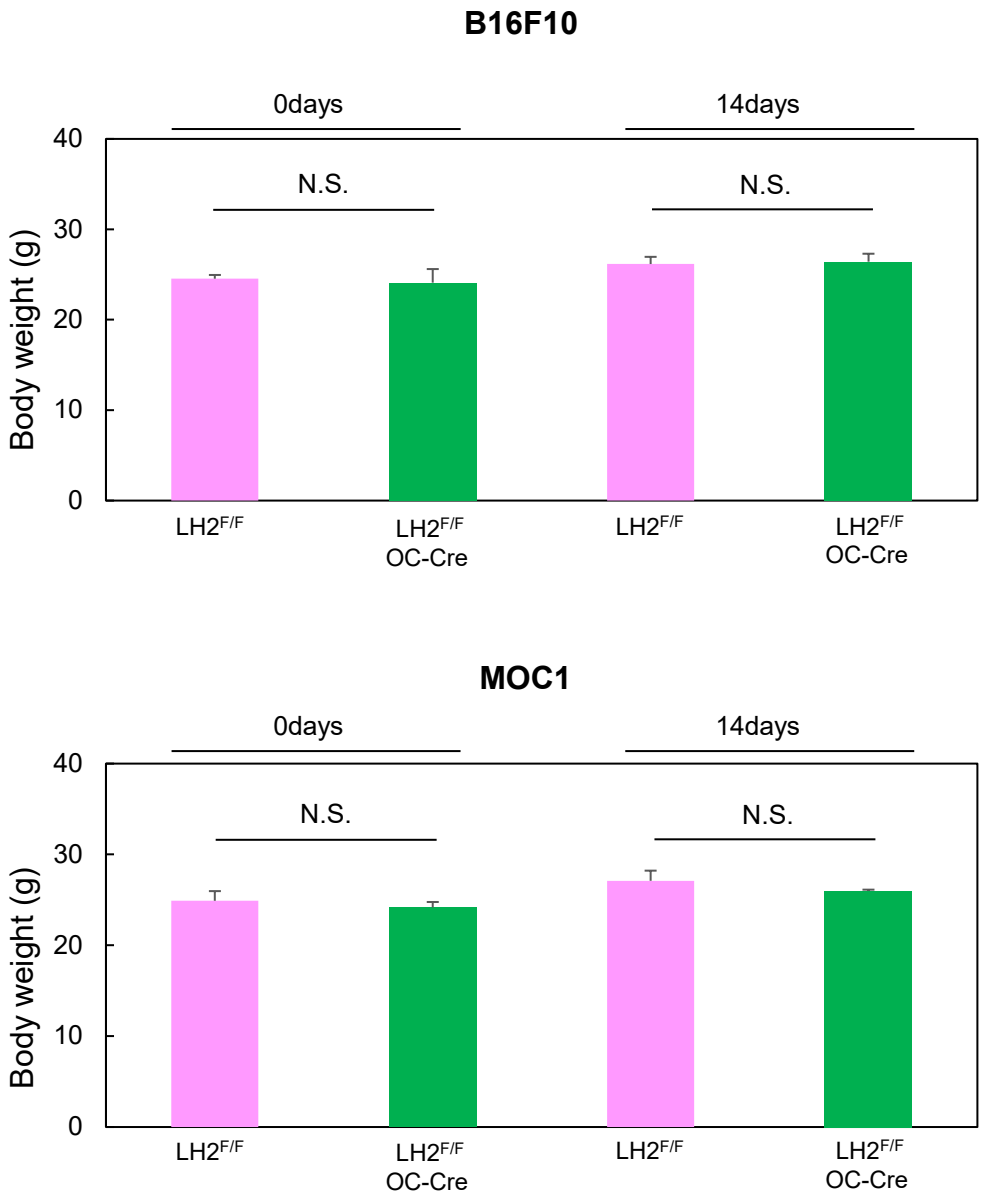
Supplementary Figure S1 | Full-length unprocessed Western blots of LH2 and α-Tubulin.
The samples derived from these experiments were used for immunoblotting shown in Figures 1A. LH2 and α-Tubulin were detected on separate membranes derived from the same samples. Three independent experiments are shown. All blots were detected under identical exposure and contrast settings.

Supplemental Figure S2



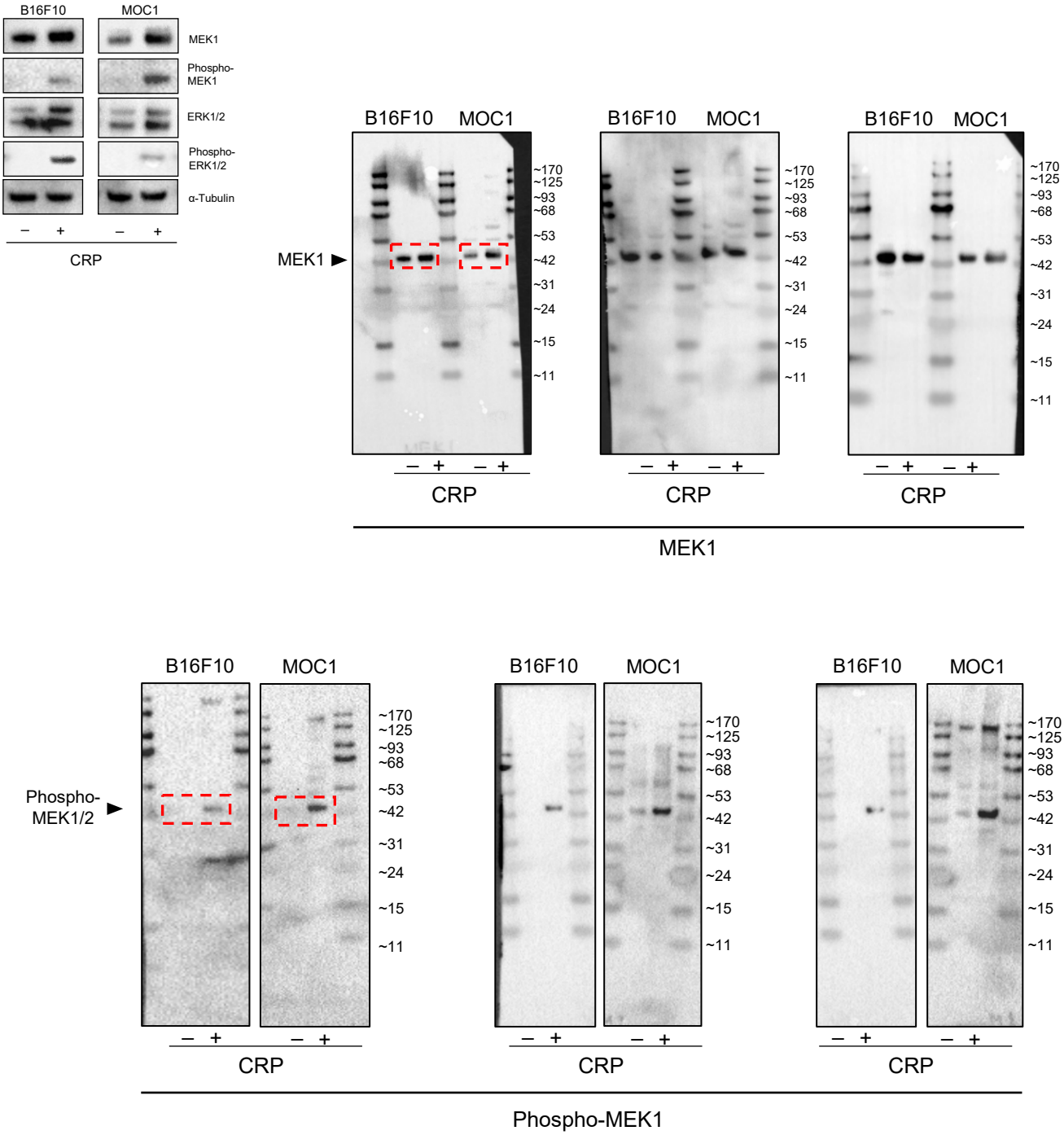
Supplementary Figure S2 | The morphology of femurs between LH2$^{F/F}$ and LH2$^{F/F}$ OC-Cre mice after cancer cell injection.
The weight, length, and width of femurs in LH2$^{F/F}$ and LH2$^{F/F}$ OC-Cre mice were collected and measured on day 14 after intra-femoral cancer cell injection. Data are presented as mean ± SD (n = 5 per group). No significant difference was observed between groups (Welch's t-test).

Supplemental Figure S3



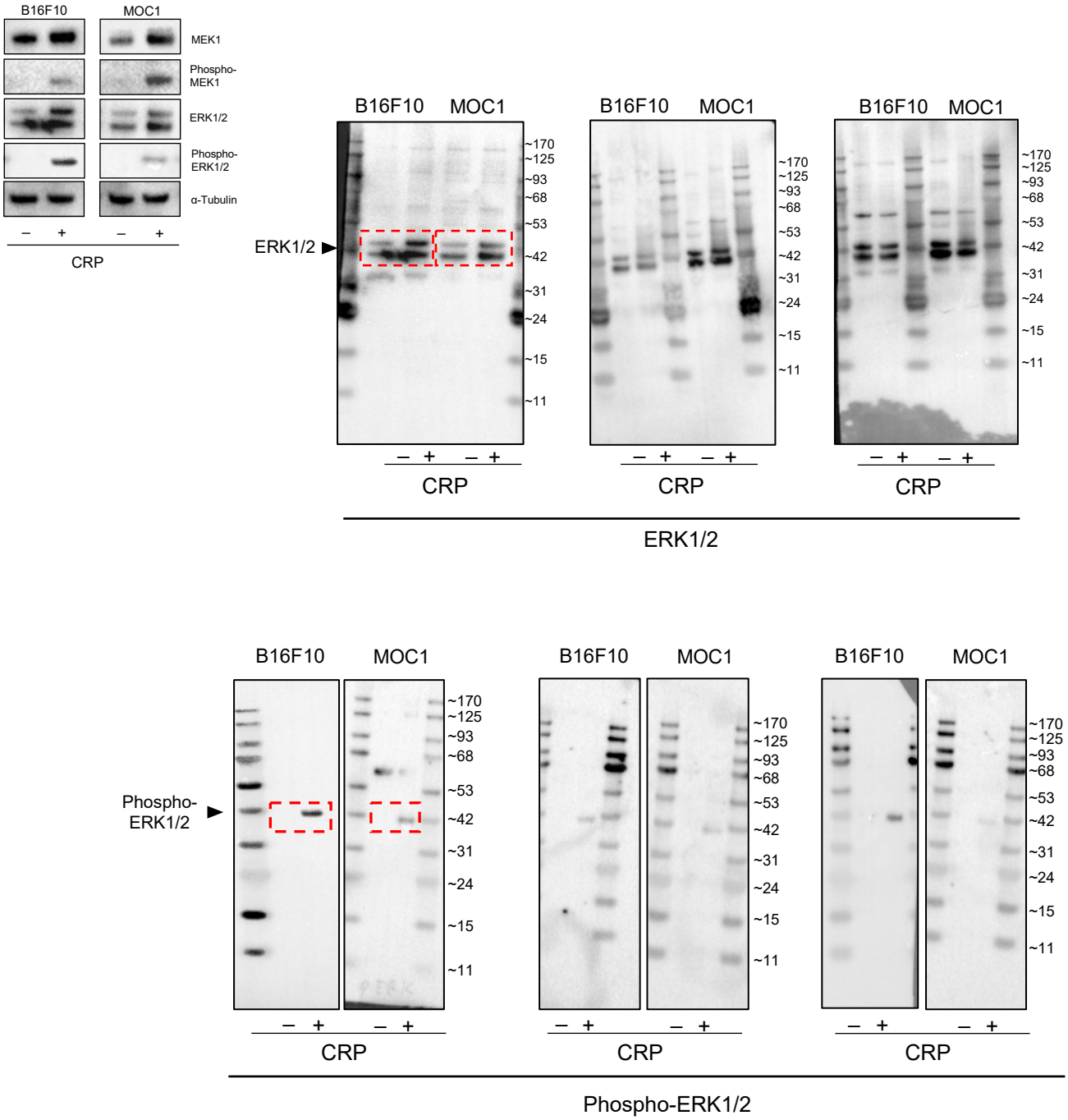
Supplementary Figure S3 | Body weight changes during bone invasion experiments.
Body weights of $LH2^{F/F}$ and $LH2^{F/F}$ OC-Cre mice were measured on day 0 and day 14 after intra-femoral cancer cell injection. Data are presented as mean ± SD (n = 5 per group). No significant difference was observed between groups (Welch's t-test).

Supplemental Figure S4
(Full length blots of Figure 5A)



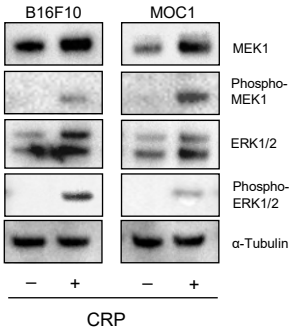
Supplementary Figure S4 | Full-length unprocessed Western blots of MEK1 and phospho-MEK1.
The samples derived from these experiments were used for immunoblotting shown in Figures 5A. MEK1 and phospho-MEK1 were detected on separate membranes derived from the same samples. Three independent experiments are shown. All blots were detected under identical exposure and contrast settings.

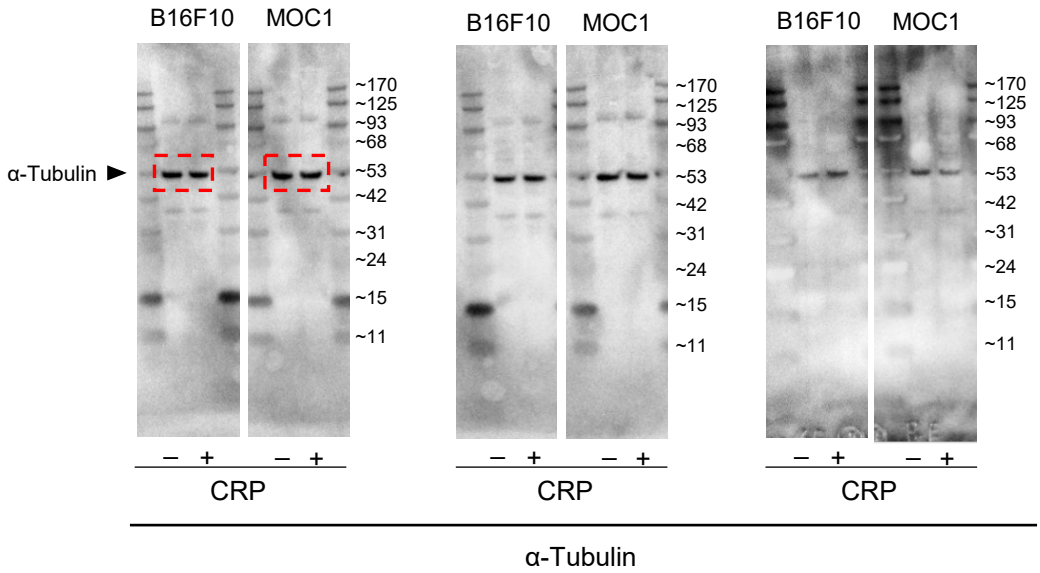
Supplemental Figure S5
(Full length blots of Figure 5A)



Supplementary Figure S5 | Full-length unprocessed Western blots of ERK1/2 and phospho-ERK1/2.
The samples derived from these experiments were used for immunoblotting shown in Figures 5A. ERK1/2 and phospho-ERK1/2 were detected on separate membranes derived from the same samples. Three independent experiments are shown. All blots were detected under identical exposure and contrast settings.

Supplemental Figure S6
(Full length blots of Figure 5A)

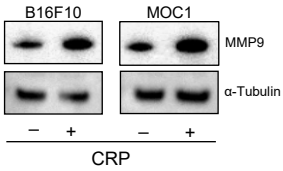


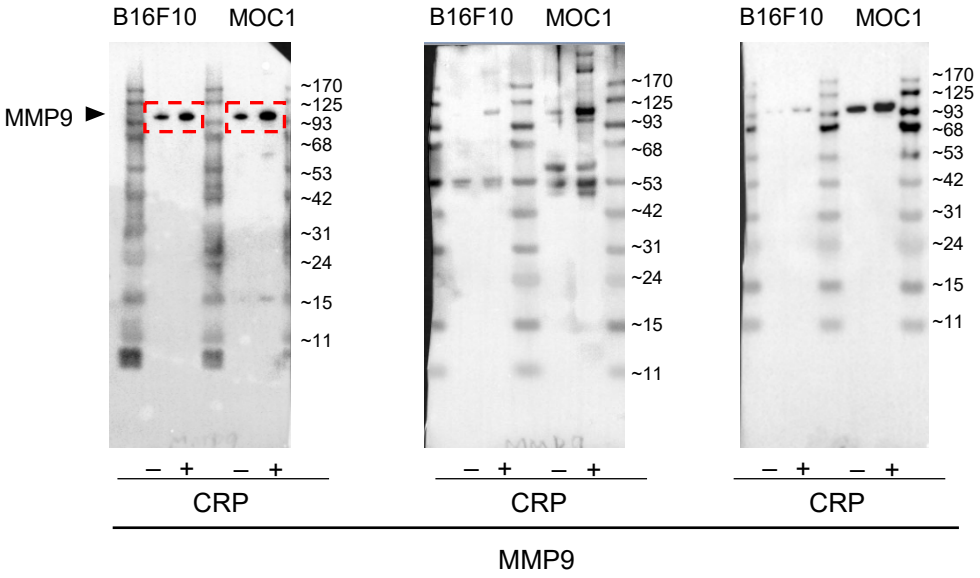


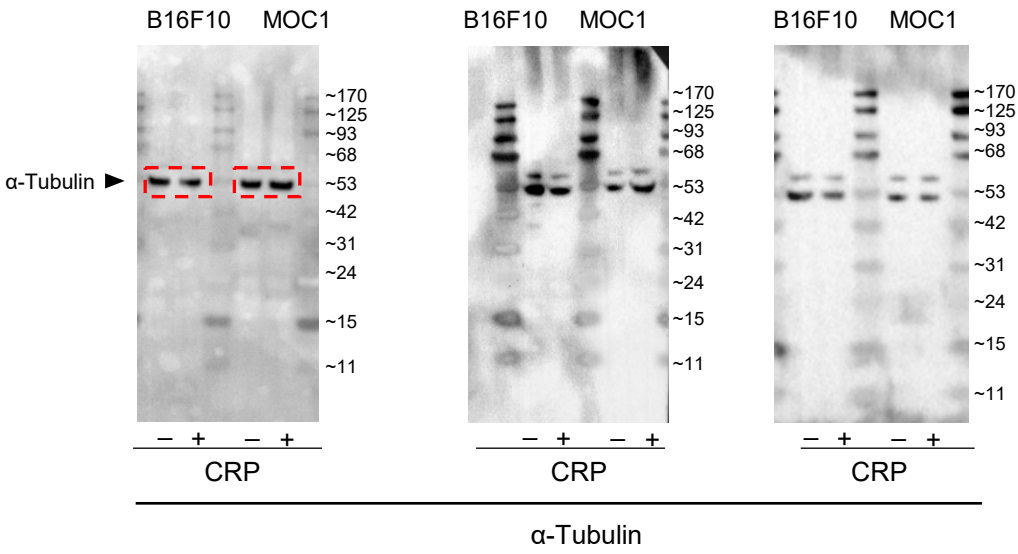
Supplementary Figure S6 | Full-length unprocessed Western blots of α-Tubulin.
The samples derived from these experiments were used for immunoblotting shown in Figures 5A. α-Tubulin was detected on separate membranes derived from the same samples. Three independent experiments are shown. All blots were detected under identical exposure and contrast settings.

Supplemental Figure S7

(Full length blots of Figure 5B)







Supplementary Figure S7 | Full-length unprocessed Western blots of MMP9 and α-Tubulin.

The samples derived from these experiments were used for immunoblotting shown in Figures 5B. MMP9 and α-Tubulin were detected on separate membranes derived from the same samples. Three independent experiments are shown. All blots were detected under identical exposure and contrast settings.