Supplementary Figure 4.

SUMF1 is taken up into wild-type and Sumf2-/− MEFs. MEFs isolated from wild-type and Sumf2-/− mice were homogenized in RIPA buffer and the protein extracts analyzed by Western blotting using an anti-SUMF2 antibody (upper panel). For uptake, cellular extracts from recipient cells (WT and Sumf2-/− MEFs) cultured in SUMF1-Flag-conditioned medium were immunoblotted with anti-Flag and anti-beta-tubulin antibodies (lower panel).