

Figure S3

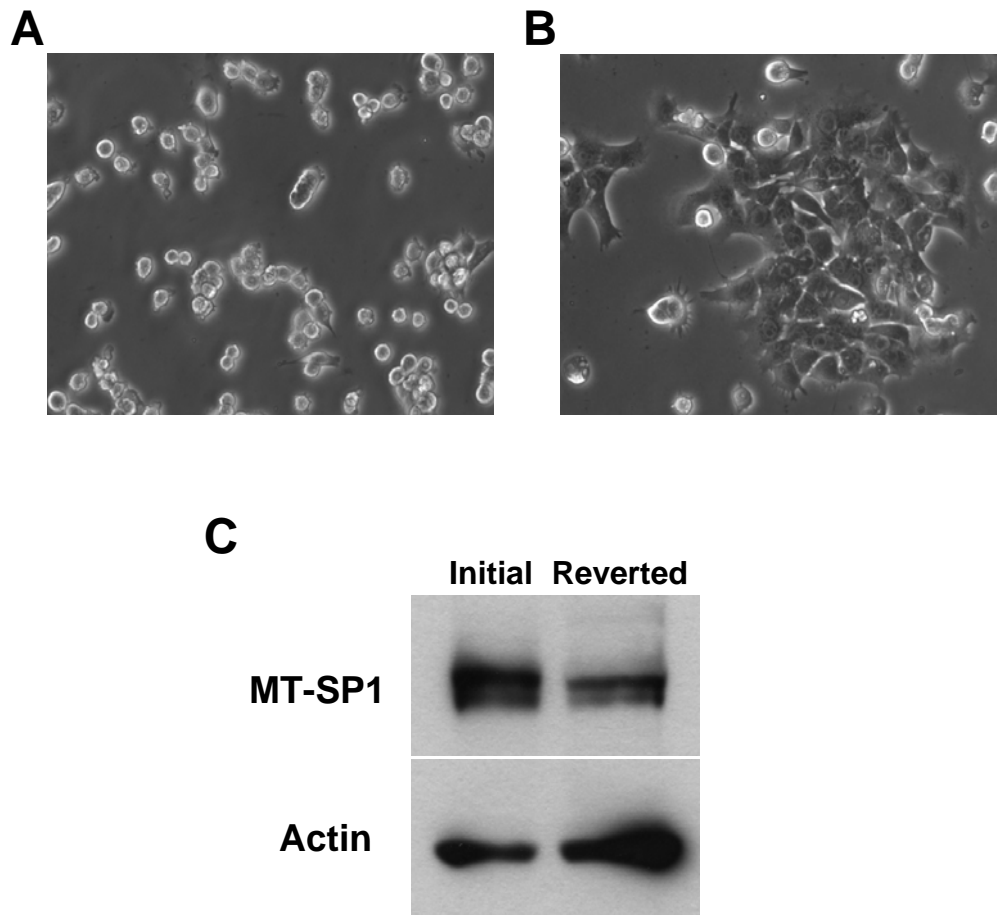


Figure S3: Matriptase (MT-SP1) protein levels in the 4T1 MT-SP1 A cells and in the subpopulation of these cells that spontaneously reverted to a “flat” morphology. **(A)** Bright field image of the initial 4T1 MT-SP1 A cells. **(B)** Bright field image of the “flat” morphology subpopulation established from 4T1 MT-SP1 A clone. **(C)** Western blot showing matriptase levels in the initial 4T1 MT-SP1 A cells (Initial) and in 4T1 MT-SP1 A cells that reverted to a “flat” phenotype (Reverted). Actin represents the loading control. Note that we purposefully overloaded the “Reverted” line to underline the decrease in matriptase level. The 4T1 MT-SP1 A cells that reverted to a “flat” phenotype were established after multiple (>20) passages. They were split using 0.05% trypsin solution in PBS after a short wash (~1min) with low concentration (0.025%) trypsin in PBS to remove less adherent cells. The cells were split 1:16 every three days.