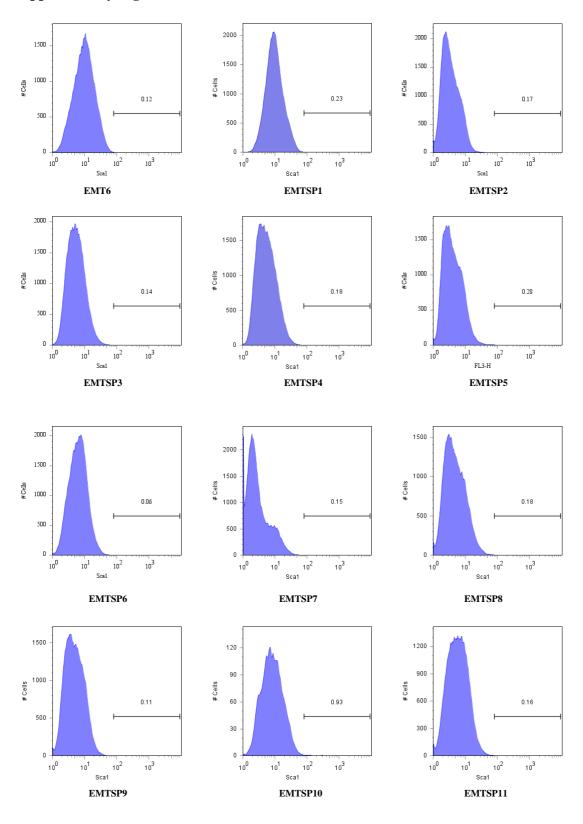
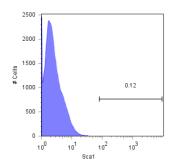
## SUPPORTING INFORMATION

## Figure S1. Sca 1<sup>+</sup> population among single cell EMT6 isolates and parental cells.

Cell populations that arose from single cells were seeded onto cell culture dishes (1×10<sup>6</sup> cells/100 mm dish) and harvested non-ezymatically in a cell dissociation solution (from Mediatech, Inc.) after 48 hours. Harvested cells were fixed overnight with 4% paraformaldehyde at 4C°, washed with PBS containing 1% fetal bovine serum that was free of Ca<sup>2+</sup> and Mg<sup>2+</sup> and the washed cells blocked with anti-mouse CD16/CD32 (mouse BD Fc Block<sup>TM</sup>) to avoid nonspecific binding of antibodies. FITC conjugated anti-mouse Sca1 antibody was used to stain each cell line. Control isotype antibodies which have no specificity for Sca1 were used as a negative control. Sca1 stained cell populations were analyzed by flow cytometry using a FACS Calibur 2 Sorter (Beckton Dickinson) and the number of cells in each phase of the cell cycle quantitated by FlowJo (Tree Star, Inc.) as indicated in each panel of the figure.

## **Supplementary Figure 1.**





EMTSP12