	MDA468		TMD231	
	shControl	shMdm2	shControl	shMdm2
G0/G1	51.7	57.6	55.2	61.3
S	11.2	10.9	16.7	13.7
G2/M	34.0	30.3	14.5	21.5

Table S1 Knock-down of Mdm2 does not affect cell cycle.

Cell cycle analysis of MDA468 and TMD-231 cells that were serum starved for 24 h. followed by replacement of media with Complete DMEM (MDA468) or addition of 1% serum for 8 hours.

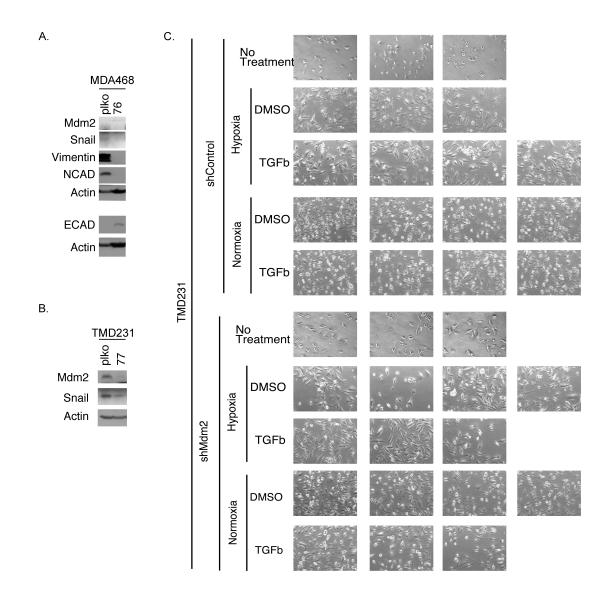


Fig. S1. Characterization of EMT in shMdm2 cell lines. A. MDA468 and B TMD231 cell lines generated from distinct shMdm2 constructs (TRCN0000003376 and TRCN0000003377 respectively) show similar changes in EMT marker levels as described in Figure 1by western blot. C. TMD231 cells do not undergo substantial morphological changes upon TGF β 1 or hypoxia treatment. shControl and shMdm2 cells were treated with DMSO or TGF β 1 under normoxia or hypoxia as described in the methods.

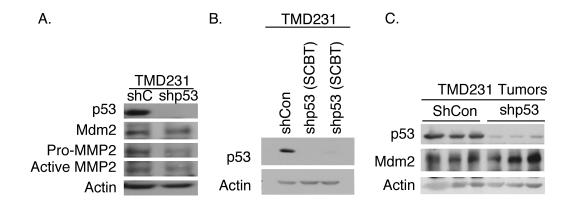


Fig. S2. Western blots of TMD231 cells with knocked-down p53 and shControl cells. A. Western blot of cells described in Figures 4 and 5 **B**. Western blot generated from cells transduced with different shp53 constructs. **C**. Western blot of *In vivo* tumors for Mdm2, p53 and Actin.