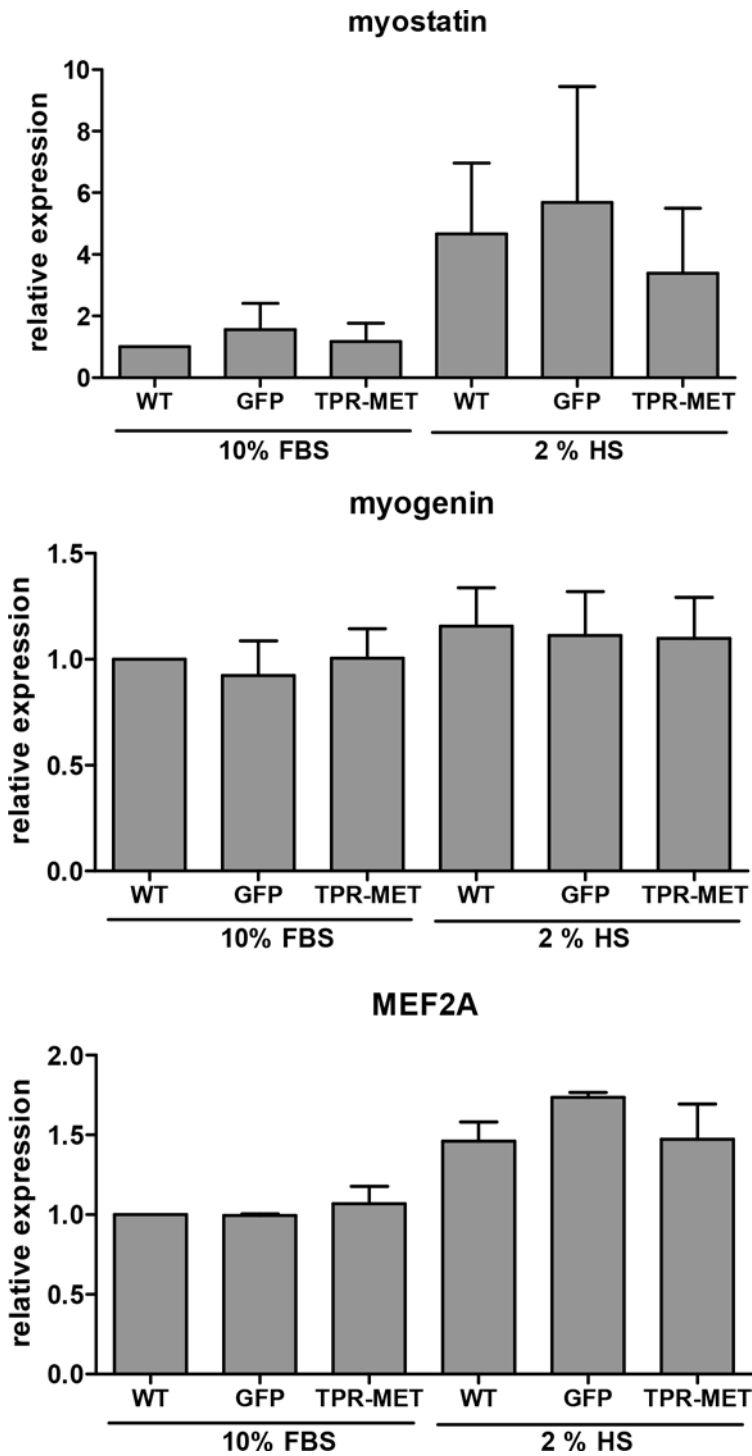


SUPPLEMENTARY TABLE AND FIGURES

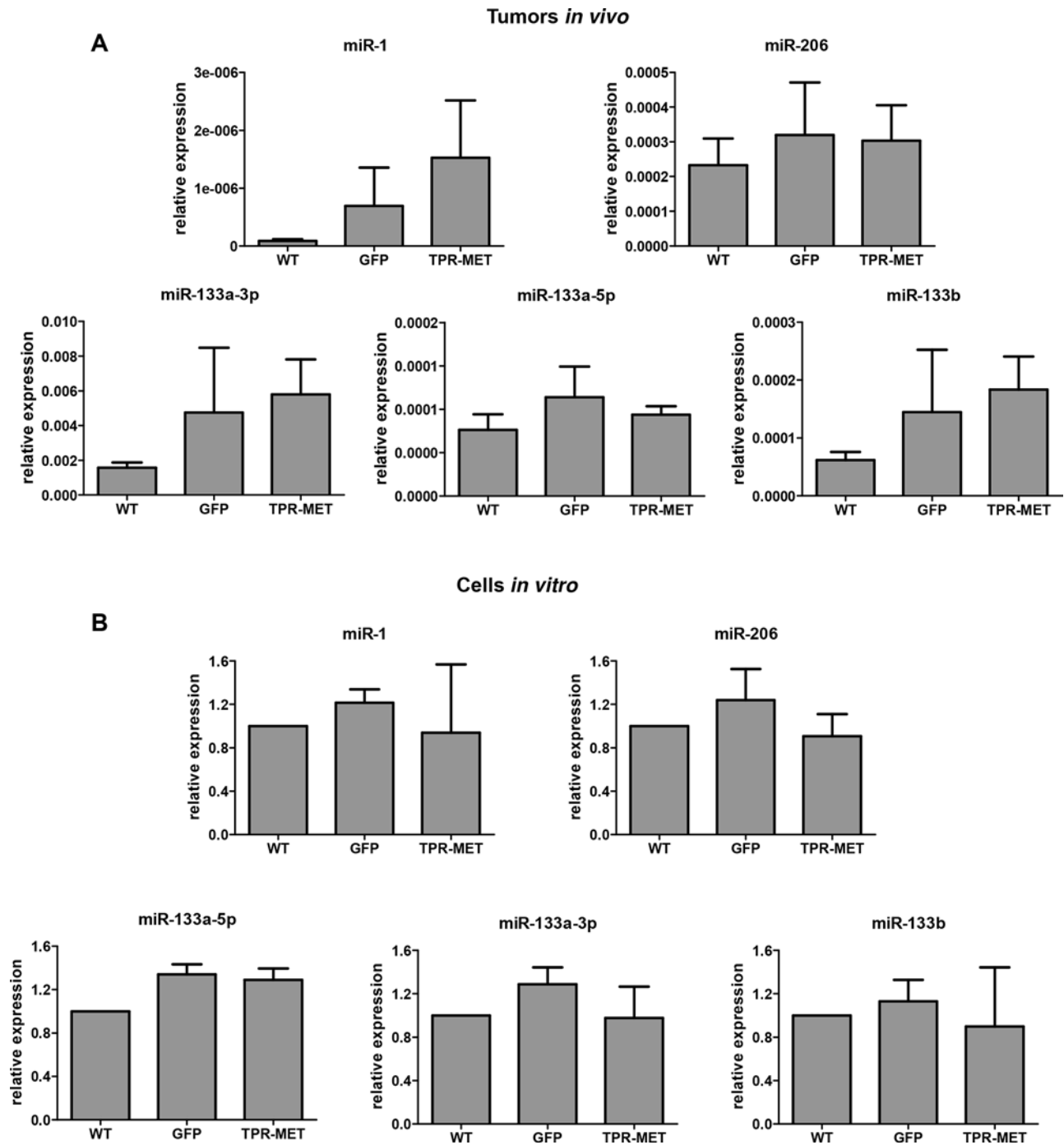
Supplementary Table 1: Values of the Pearson correlation between MET and myogenic differentiation markers in RMS tumors.

	MET	MYF5	MyoD	MEF2A	myogenin	myostatin	MYH2
MET		R = -0.68 <i>p</i> = 0.06	R = 0.87 <i>p</i> = 0.005	R = -0.33 <i>p</i> = 0.42	R = -0.43 <i>p</i> = 0.29	R = 0.26 <i>p</i> = 0.53	R = -0.55 <i>p</i> = 0.16
MYF5	R = -0.68 <i>p</i> = 0.06		R = -0.58 <i>p</i> = 0.14	R = -0.08 <i>p</i> = 0.86	R = 0.16 <i>p</i> = 0.71	R = 0.02 <i>p</i> = 0.96	R = 0.45 <i>p</i> = 0.26
MyoD	R = 0.87 <i>p</i> = 0.005	R = -0.58 <i>p</i> = 0.14		R = -0.37 <i>p</i> = 0.36	R = -0.42 <i>p</i> = 0.30	R = 0.27 <i>p</i> = 0.52	R = -0.39 <i>p</i> = 0.34
MEF2A	R = -0.33 <i>p</i> = 0.42	R = -0.08 <i>p</i> = 0.86	R = -0.37 <i>p</i> = 0.36		R = 0.92 <i>p</i> = 0.001	R = -0.49 <i>p</i> = 0.22	R = 0.69 <i>p</i> = 0.06
myogenin	R = -0.43 <i>p</i> = 0.29	R = 0.16 <i>p</i> = 0.71	R = -0.42 <i>p</i> = 0.30	R = 0.92 <i>p</i> = 0.001		R = -0.21 <i>p</i> = 0.62	R = 0.89 <i>p</i> = 0.003
myostatin	R = 0.26 <i>p</i> = 0.53	R = 0.02 <i>p</i> = 0.96	R = 0.27 <i>p</i> = 0.52	R = -0.49 <i>p</i> = 0.22	R = -0.21 <i>p</i> = 0.62		R = 0.0005 <i>p</i> = 0.99
MYH2	R = -0.55 <i>p</i> = 0.16	R = 0.45 <i>p</i> = 0.26	R = -0.39 <i>p</i> = 0.34	R = 0.69 <i>p</i> = 0.06	R = 0.89 <i>p</i> = 0.003	R = 0.0005 <i>p</i> = 0.99	

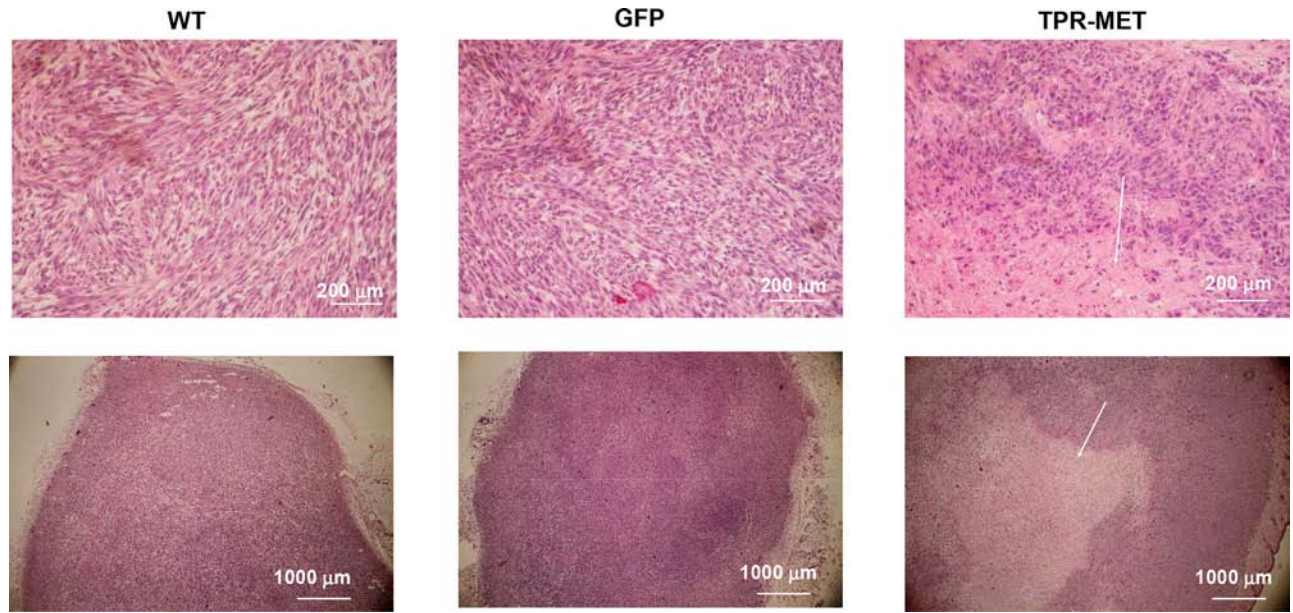
Expression of genes was evaluated by qPCR in paraffin-embedded RMS specimens, *n* = 8, R = correlation coefficient, *p* < 0.05 considered as significant.



Supplementary Figure 1: Myostatin and MEF2A are slightly upregulated in SMS-CTR cells cultured in DMEM with 2% HS. Expression of myostatin, myogenin and MEF2A was evaluated by qPCR in samples from SMS-CTR cells, $n = 2$. Data in graphs are represented as mean \pm SEM.



Supplementary Figure 2: The effects of TPR-MET are independent of myomiRs action. Expression of miR-1, miR-206, miR-133a-3p, miR-133a-5p and miR-133b was estimated by quantitative PCR in tumors formed by SMS-CTR cells after subcutaneous implantation to NOD-SCID mice, $n = 4-5$ (A) and in SMS-CTR cells *in vitro*, $n = 2$ (B) Data in graphs are represented as mean \pm SEM.



Supplementary Figure 3: Necrosis in tumors formed by SMS-CTR cells expressing TPR-MET. Representative images of tumors sections stained with hematoxylin-eosin show necrotic areas in TPR-MET tumors, which are indicated with arrows.