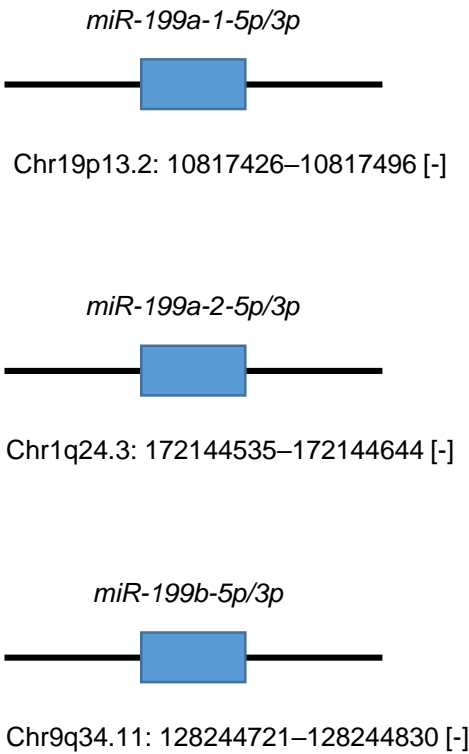


Figure S1

A



B

The guide strands

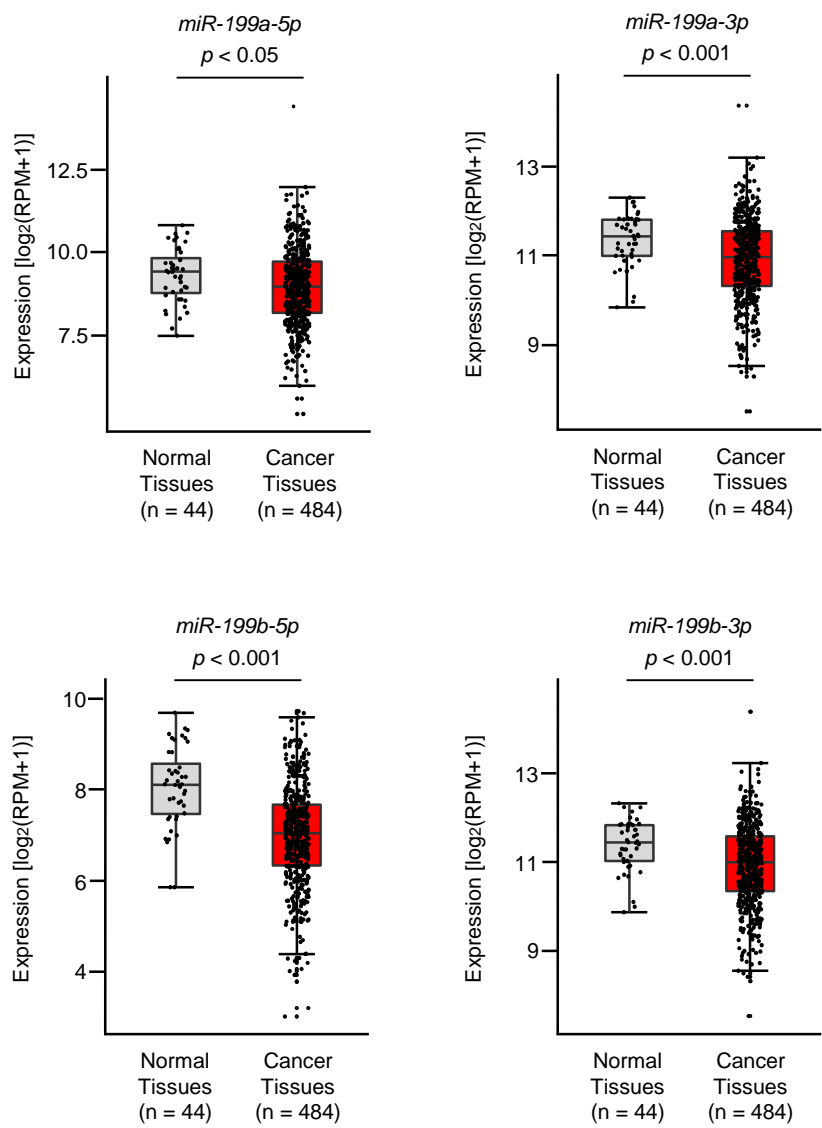
<i>miR-199a-1-5p</i>	CCCAGUGUUCAGACUACCUGUUC
<i>miR-199a-2-5p</i>	CCCAGUGUUCAGACUACCUGUUC
<i>miR-199b-5p</i>	CCCAGUGUUUAGACUAUCUGUUC

The passenger strands

<i>miR-199a-1-3p</i>	ACAGUAGUCUGCACAUUGGUUA
<i>miR-199a-2-3p</i>	ACAGUAGUCUGCACAUUGGUUA
<i>miR-199b-3p</i>	ACAGUAGUCUGCACAUUGGUUA

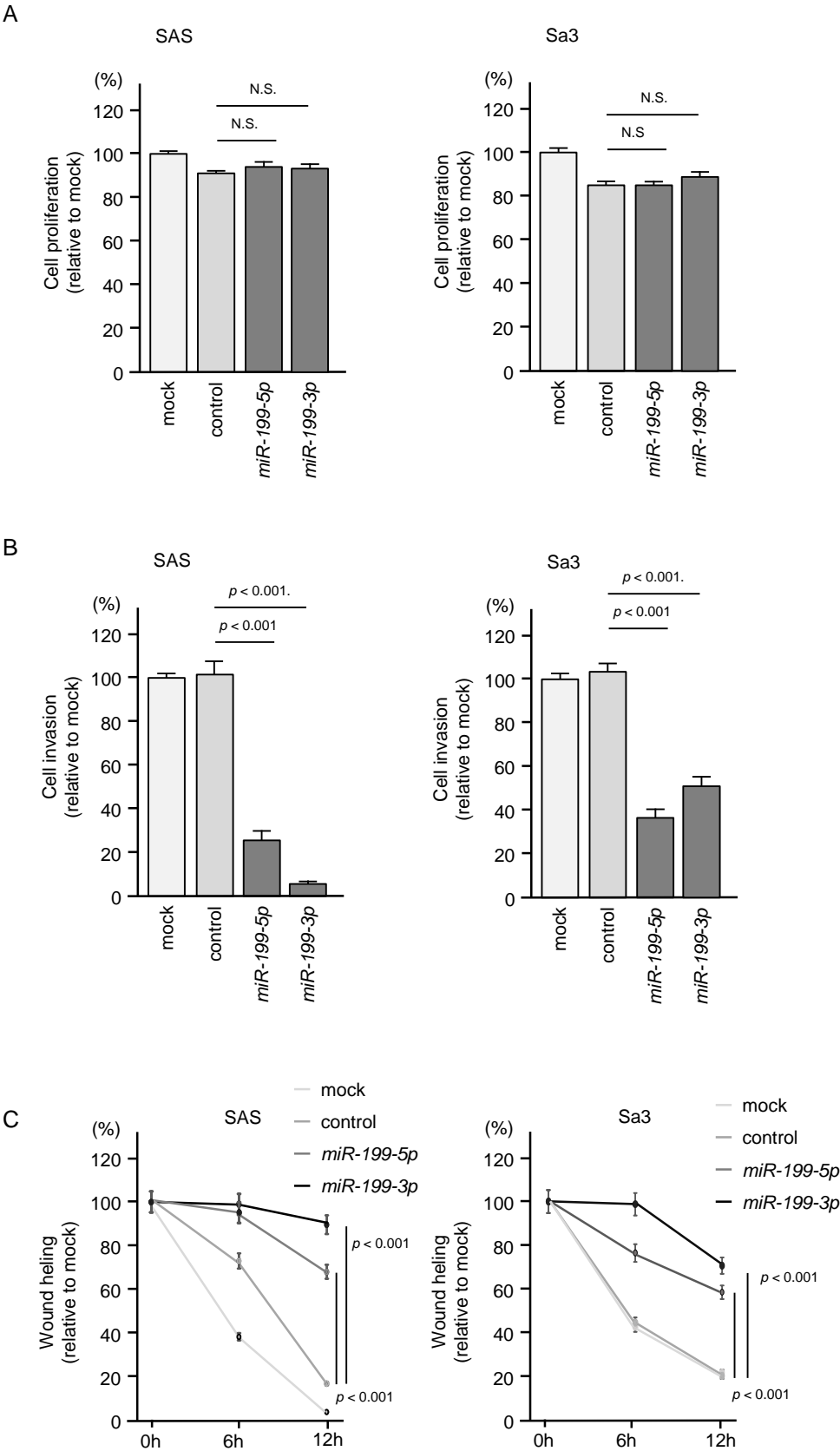
Supplementary Figure S1: Chromosomal locations and mature sequences of the *miR-199* family.
(A) The *miR-199* family is composed of 3 species: *miR-199a-1*, *miR-199a-2*, and *miR-199b* located on human chromosome 19p13.2, 1q24.3, and 9q34.11 respectively. (B) Mature sequences of the *miR-199* family are shown. miRBase shows that *miR-199-5p* annotates as the guide strand and *miR-199-3p* as the passenger strand. Mature sequences of *miR-199a-1-5p* and *miR-199a-2-5p* are identical. The mature sequence of *miR-199b-5p* differs in one base compares with *miR-199a-5p*. The seed sequences of the 3 miRNAs are identical (red letters). Mature sequences of *miR-199a-1-3p*, *miR-199a-2-3p* and *miR-199b-3p* are identical. The seed sequences of the 3 miRNAs are identical (blue letters).

Figure S2



Supplementary Figure S2: The expression levels of *miR-199* family.
The *miR-199* family (*miR-199a-5p*, *miR-199a-3p*, *miR-199-b-5p*, and *miR-199b-3p*) were evaluated by TCGA-HNSC dataset. All member of *miR-199* family were downregulated in cancer tissues.

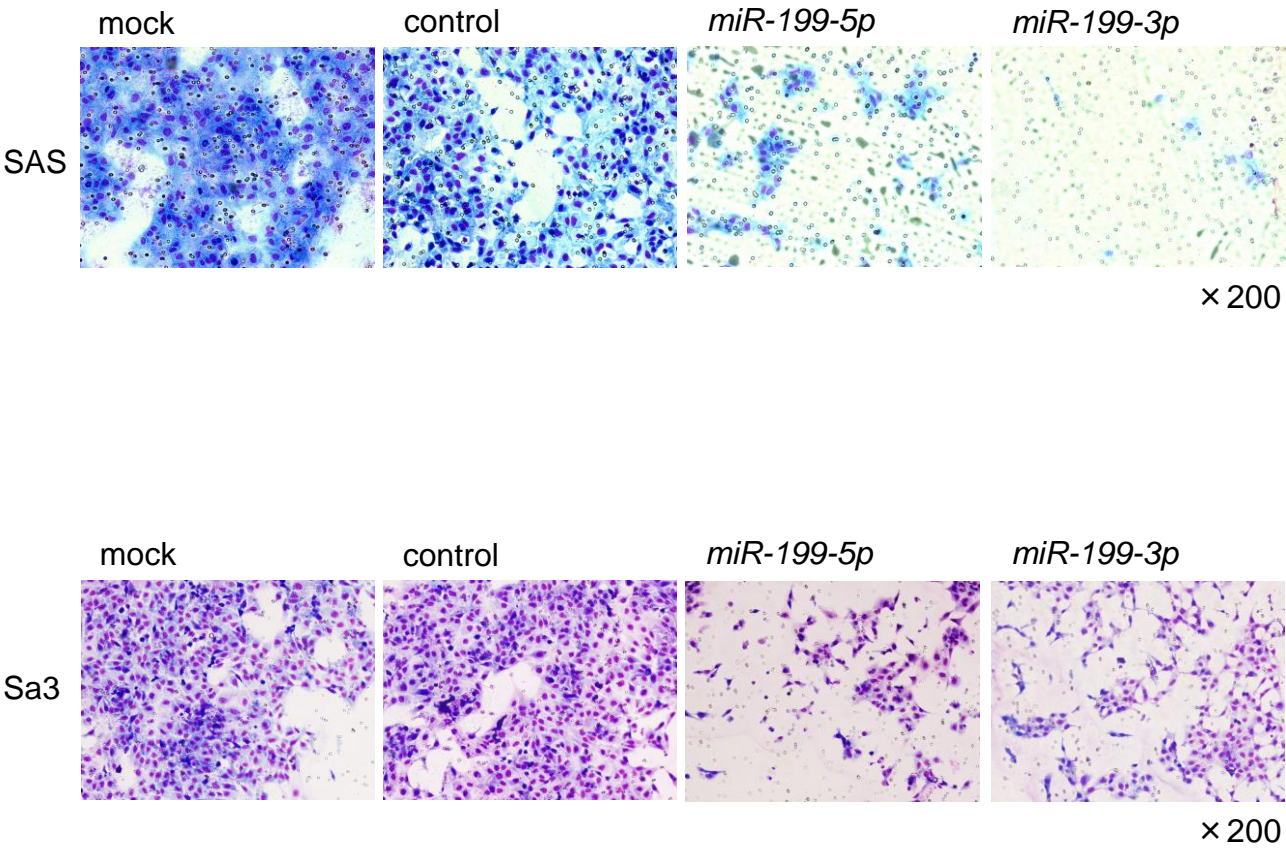
Figure S3



Supplementary Figure S3: Functional assays of cell proliferation, invasion and migration following ectopic expression of *miR-199-5p* and *miR-199-3p* in HNSCC cell lines (SAS and Sa3).

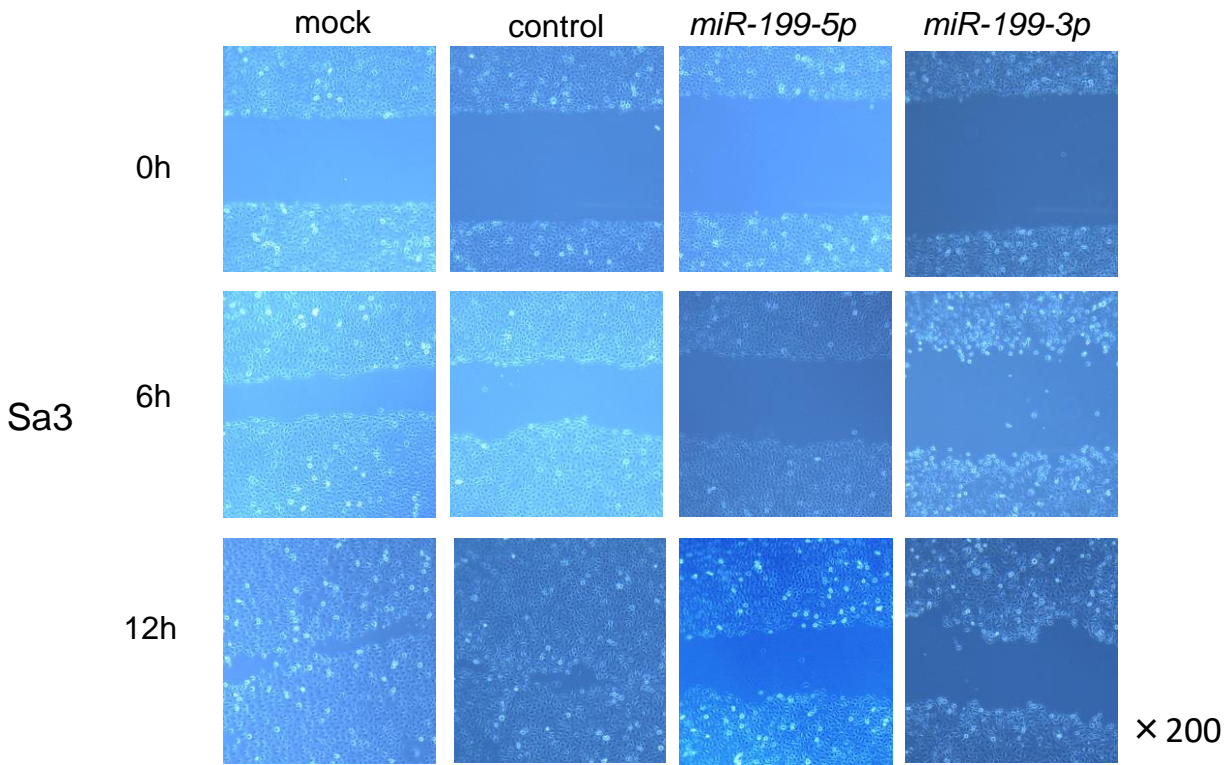
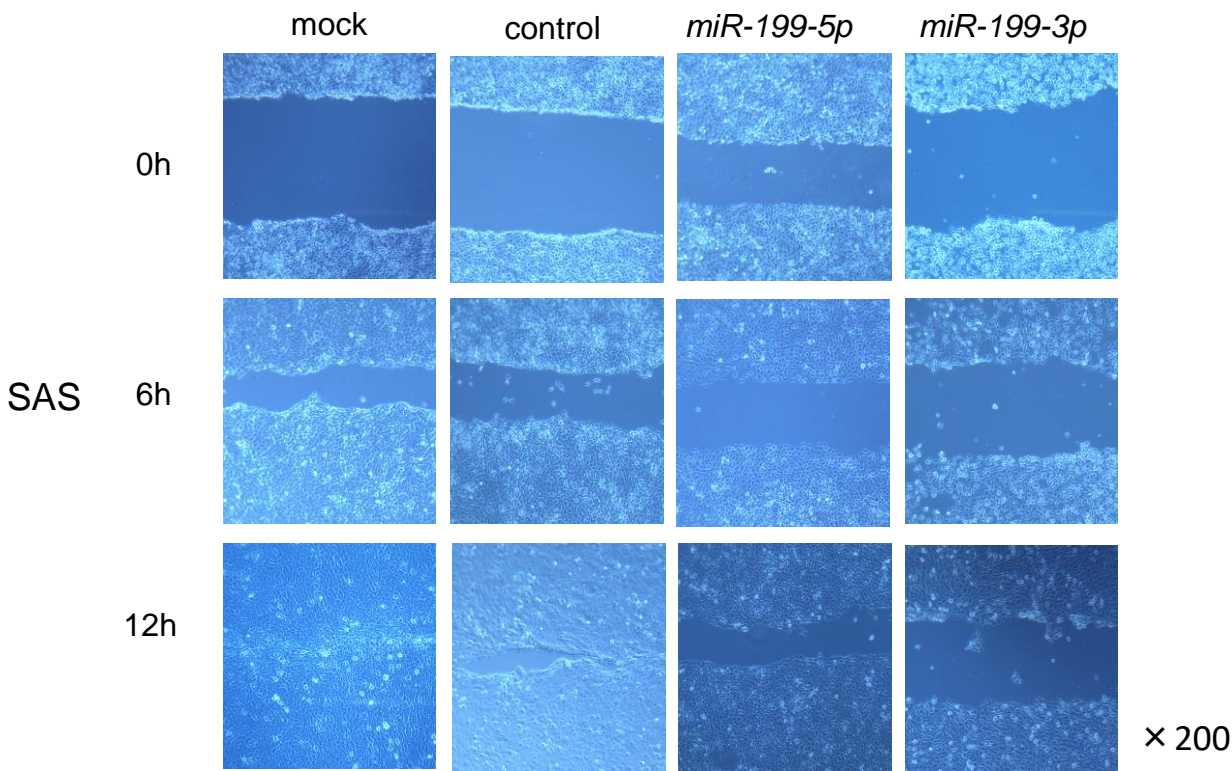
(A) Cell proliferation assessed by XTT assay at 72 h after *miR-199-5p* and *miR-199-3p* transfection (N.S., not significant). (B) Cell invasion assessed by Matrigel invasion assays at 48 h after seeding *miR-199-5p* and *miR-199-3p* transfected cells into chambers. (C) Cell migration assessed by wound healing assay at 0, 6, and 12 h after cell scratch formation.

Figure S4



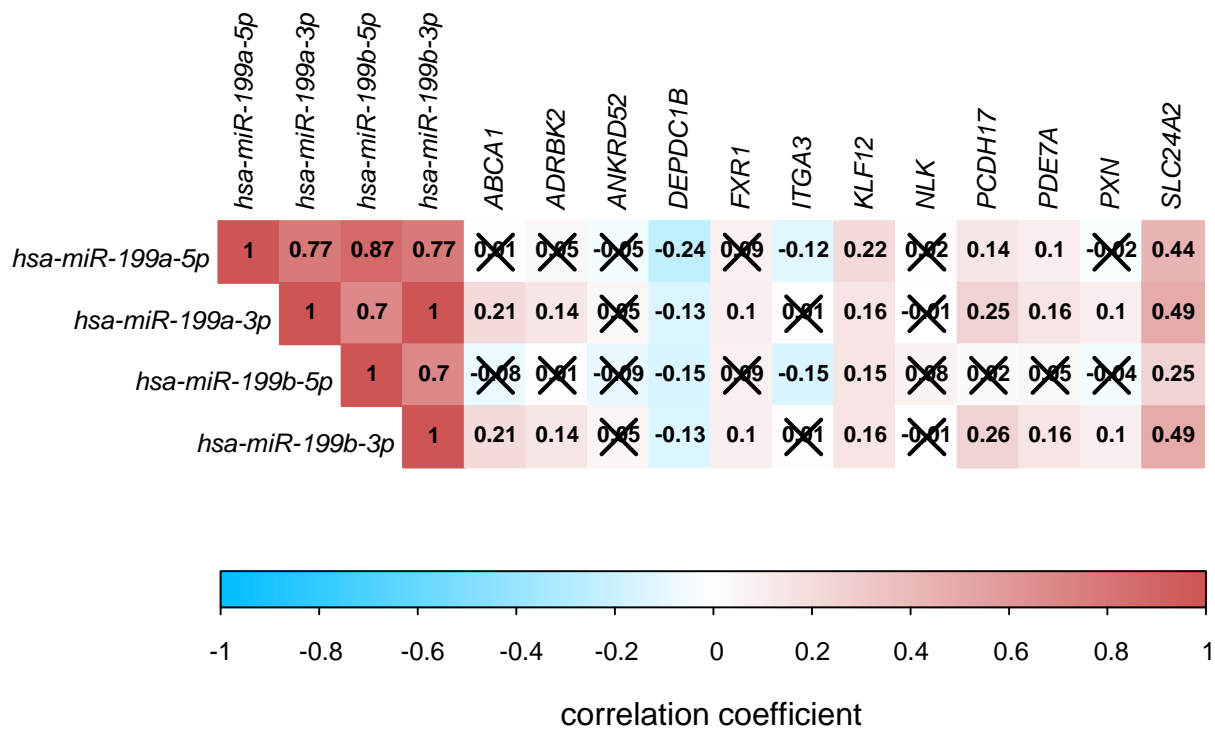
Supplementary Figure S4: Photomicrographs of cells subjected invasion assays.
Typical images of cells in invasion assays following *miR-199-5p* and *miR-199-3p* transfection into HNSCC cell lines, SAS and Sa3.

Figure S5



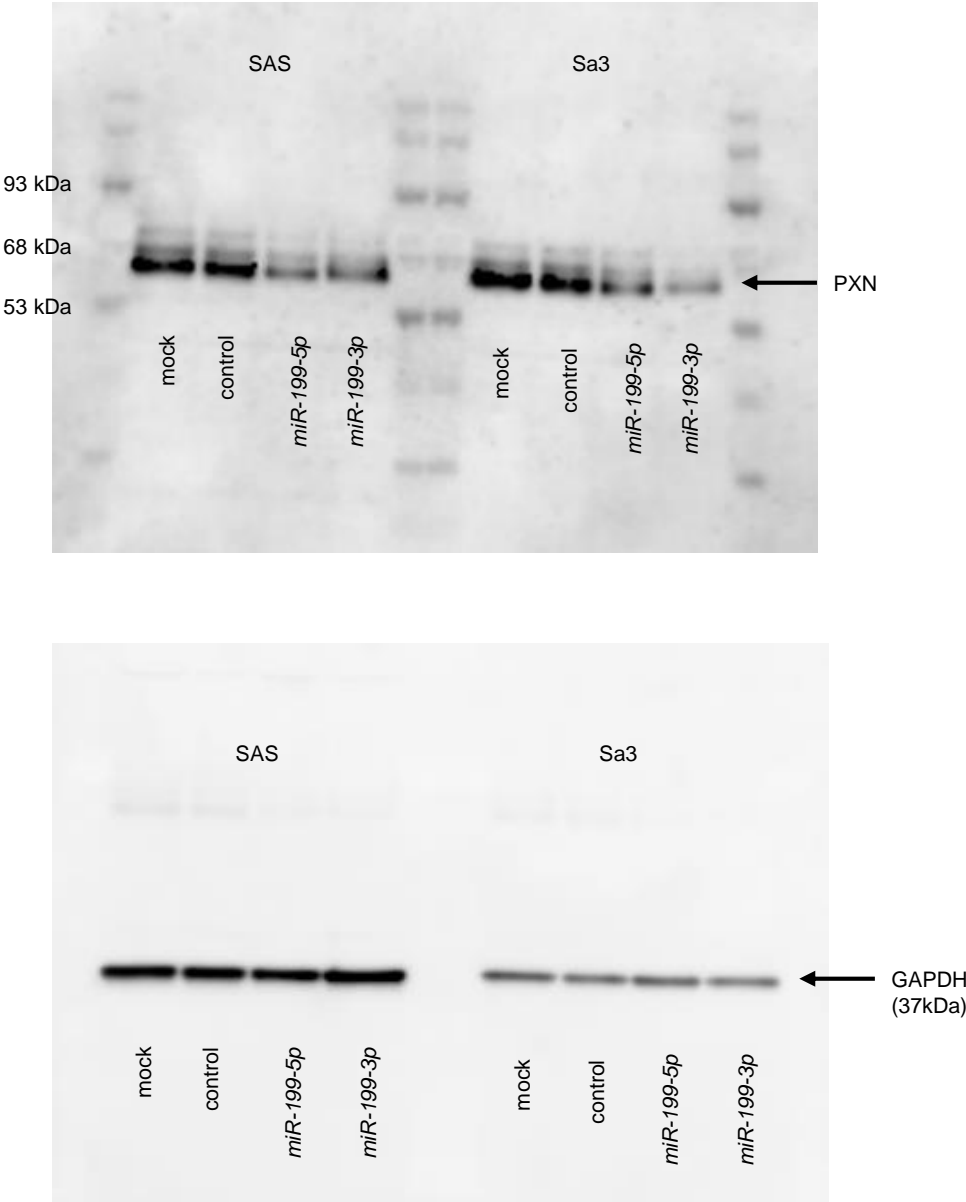
Supplementary Figure S5: Photomicrographs of cells subjected wound healing assays.
Typical images of cells in wound healing assays following *miR-199-5p* and *miR199-3p* transfection into HNSCC cell lines, SAS and Sa3.

Figure S6



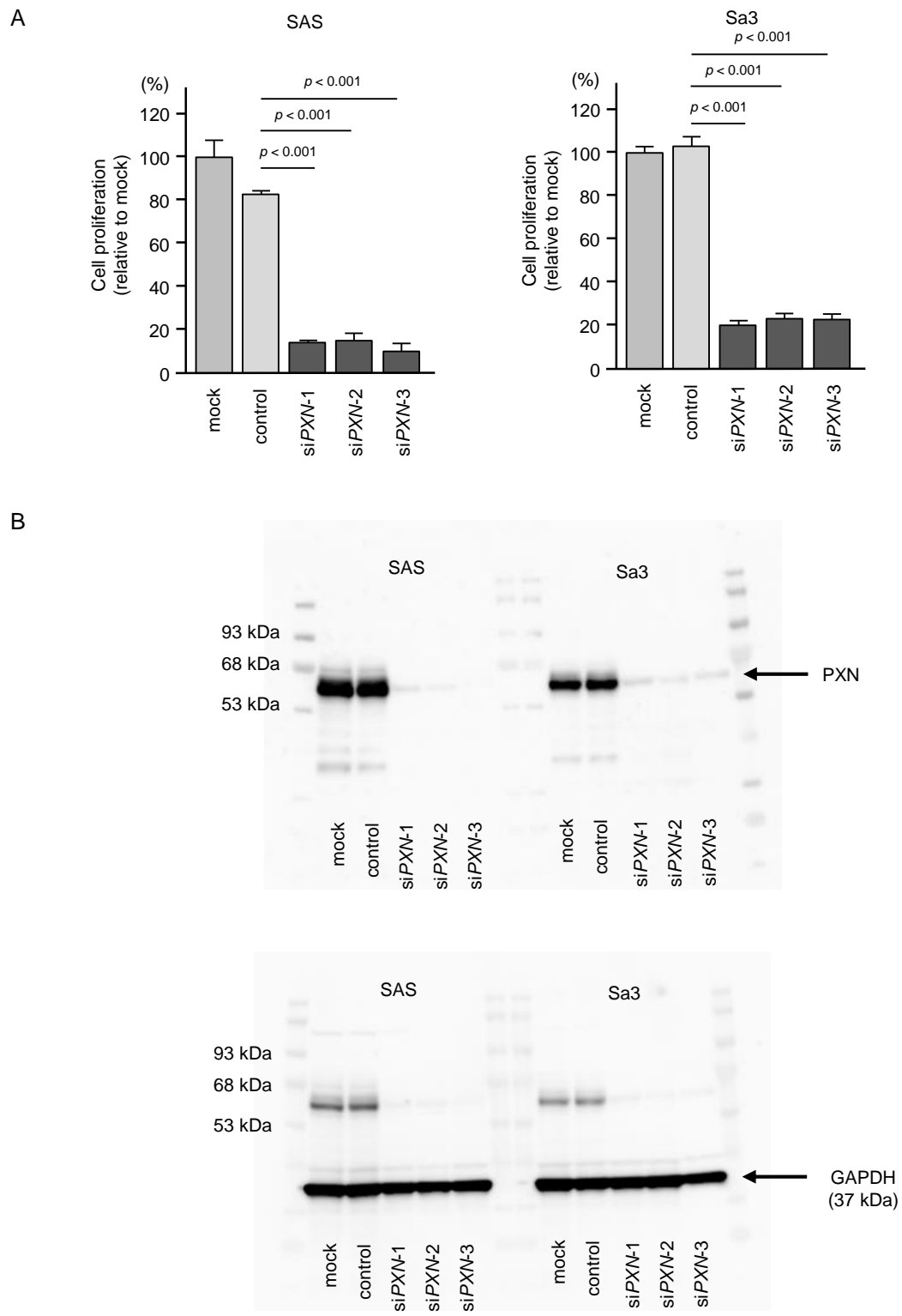
Supplementary Figure S6: Correlation between *miR-199* family and putative target genes in TCGA-HNSC. The plots showed the correlation among expression levels of *miR-199* family and putative target genes in cancer tissues from TCGA-HNSC according to Spearman's rank tests. The values in each cell showed correlation coefficients between miRNA/gene in each row and column. The color scale was created by correlation values. Red cells indicated positive , and blue cells indicated negative correlation. Cells without a cross represented that a relationship was significant (p -value < 0.05).

Figure S7



Supplementary Figure S7: Full size image of Western blotting of Figure 5.

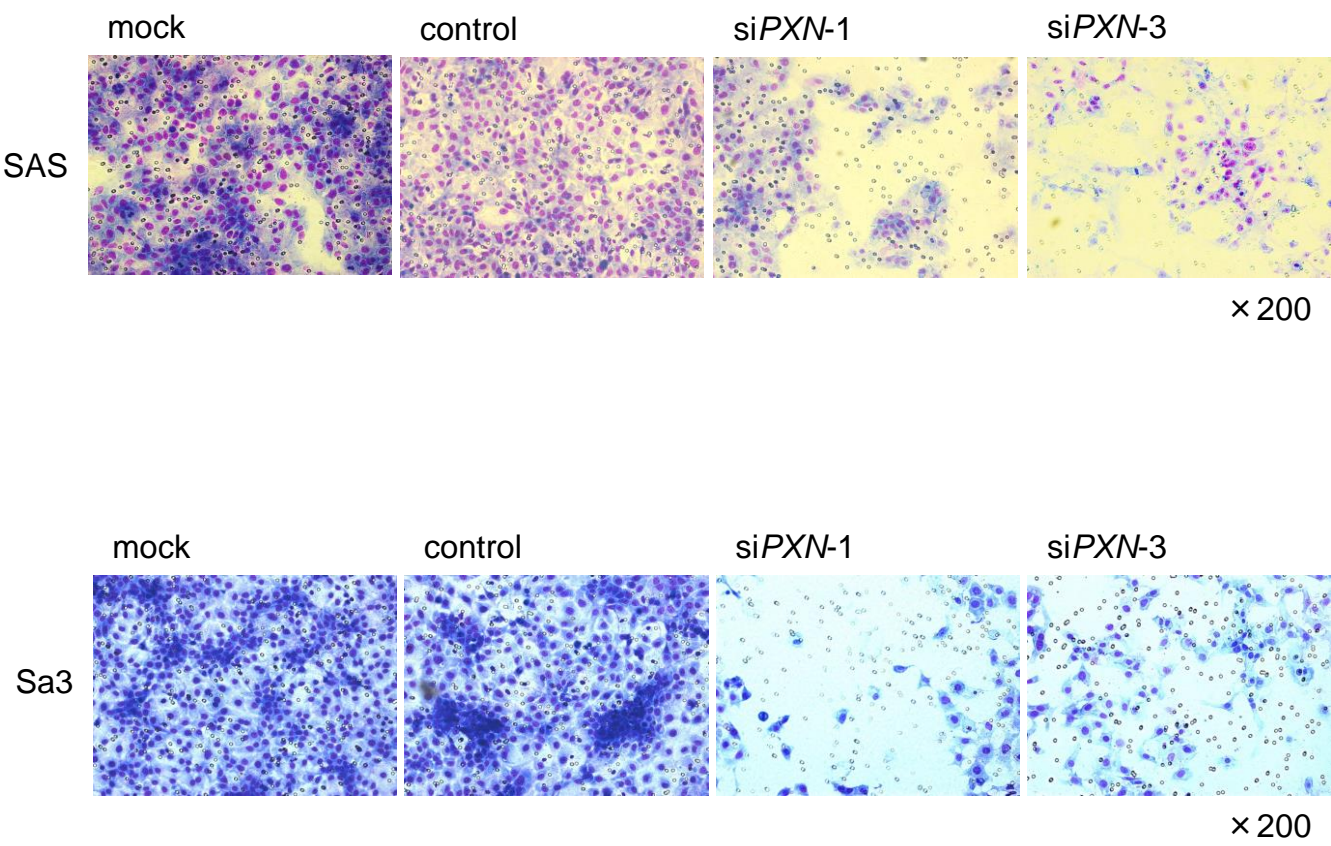
Figure S8



Supplementary Figure S8: siRNA knockdown efficiencies for *PXN* in HNSCC cell lines (SAS and Sa3 cells).

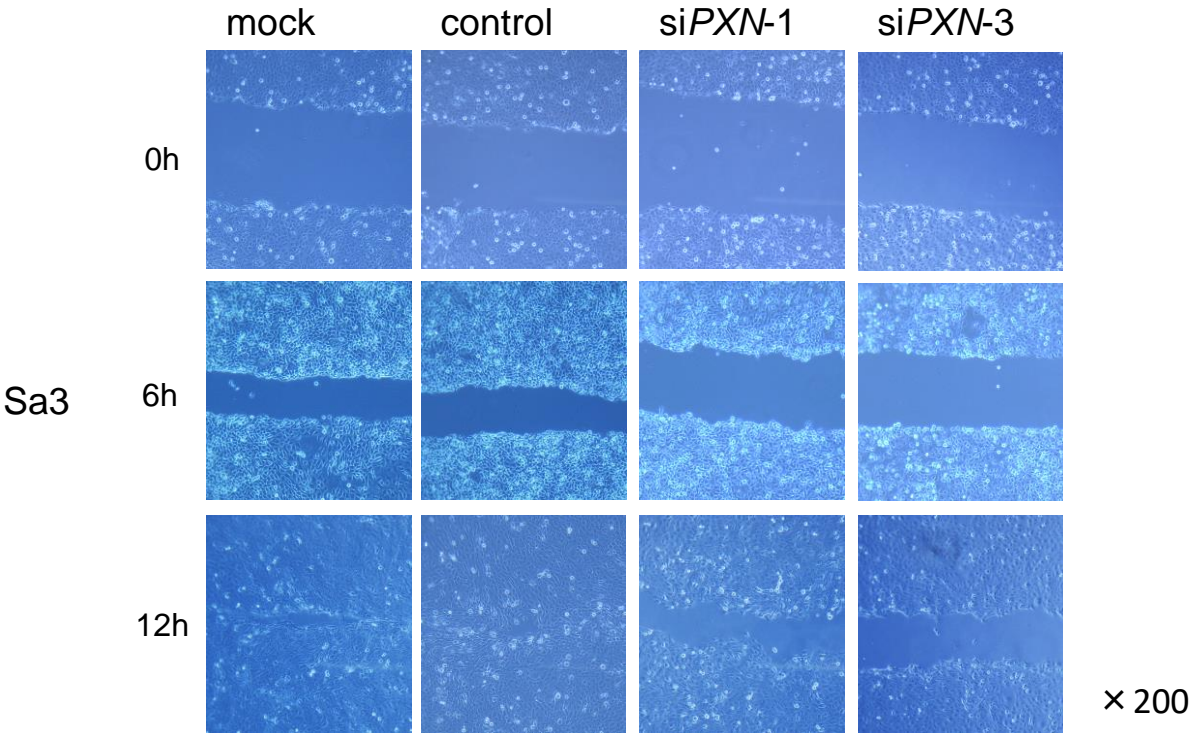
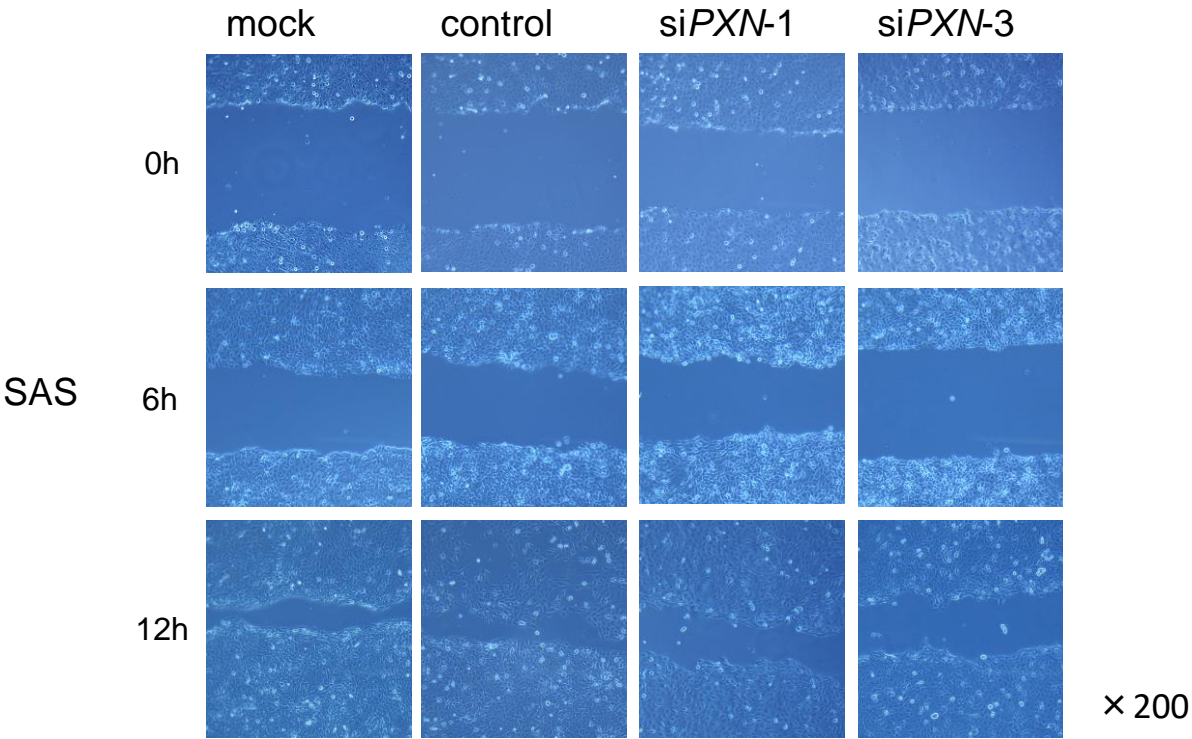
The efficiencies of *PXN* knockdown by siPXN-1, siPXN-2 and siPXN-3 were evaluated by qRT-PCR (A) and Western blotting (B). The *PXN* mRNA and protein levels were measured 72 h after siRNA transfection. GAPDH mRNA and protein levels were used as internal controls for qRT-PCR and Western blotting, respectively.

Figure S9



Supplementary Figure S9: Photomicrographs of cells subjected invasion assays.
Typical images of cells in invasion assays following si*PXN*-1 and si*PXN*-3 transfection into HNSCC cell lines, SAS and Sa3.

Figure S10



Supplementary Figure S10: Photomicrographs of cells subjected wound healing assays.
Typical images of cells in wound healing assays following si*PXN*-1 and si*PXN*-3 transfection into HNSCC cell lines, SAS and Sa3.