

ERRATUM

The following was originally published in Volume 25, Number 1, pages 1–10 (DOI: <https://doi.org/10.3727/096504016X14685034103798>). Information in Figure 5C was incorrectly displayed. The corrected Figure 5 is provided here.

MicroRNA-200a Suppresses Cell Invasion and Migration by Directly Targeting GAB1 in Hepatocellular Carcinoma

Jianlin Wang,^{*1} Wenjie Song,^{*1} Weiwei Shen,^{†1} Xisheng Yang,^{*} Wei Sun,^{*} Sshibin Qu,^{*} Runze Shang,^{*} Ben Ma,^{*} Meng Pu,^{*} Kaishan Tao,^{*} Kefeng Dou,^{*} and Haimin Li^{*}

^{*}Department of Hepatobiliary Surgery, Xijing Hospital, Fourth Military Medical University, Xi'an, Shaanxi, P.R. China
[†]Department of Oncology, Tangdu Hospital, Fourth Military Medical University, Xi'an, Shaanxi, P.R. China

MicroRNA-200a (miR-200a) is frequently downregulated in most cancer types and plays an important role in carcinogenesis and cancer progression. In this study, we determined that miR-200a was downregulated in hepatocellular carcinoma (HCC) tissues and cell lines, consistent with the results of our previous study. Because a previous study suggested that downregulation of miR-200a is correlated with HCC metastasis, we aimed to elucidate the mechanism underlying the role of miR-200a in metastasis in HCC. Here we observed that overexpression of miR-200a resulted in suppression of HCC metastatic ability, including HCC cell migration, invasion, and metastasis, *in vitro* and *in vivo*. Furthermore, bioinformatics and luciferase reporter assays indicated that GAB1 is a direct target of miR-200a. Inhibition of GAB1 resulted in substantially decreased cell invasion and migration similar to that observed with overexpression of miR-200a in HCC cell lines, whereas restoration of GAB1 partially rescued the inhibitory effects of miR-200a. Taken together, these data provide novel information for comprehending the tumor-suppressive role of miR-200a in HCC pathogenesis through inhibition of GAB1 translation.

Key words: miR-200a; Hepatocellular carcinoma (HCC); Grb2-associated binding protein 1 (GAB1); Invasion and migration; Metastasis

¹These authors provided equal contribution to this work.

Address correspondence to Haimin Li, Department of Hepatobiliary Surgery, Xijing Hospital, Fourth Military Medical University, 127 West Changle Street, Xi'an, Shaanxi 710032, P.R. China. E-mail: lihaim@fmmu.edu.cn or Kefeng Dou, Department of Hepatobiliary Surgery, Xijing Hospital, Fourth Military Medical University, 127 West Changle Street, Xi'an, Shaanxi 710032, P.R. China. E-mail: gdwkgwx@fmmu.edu.cn

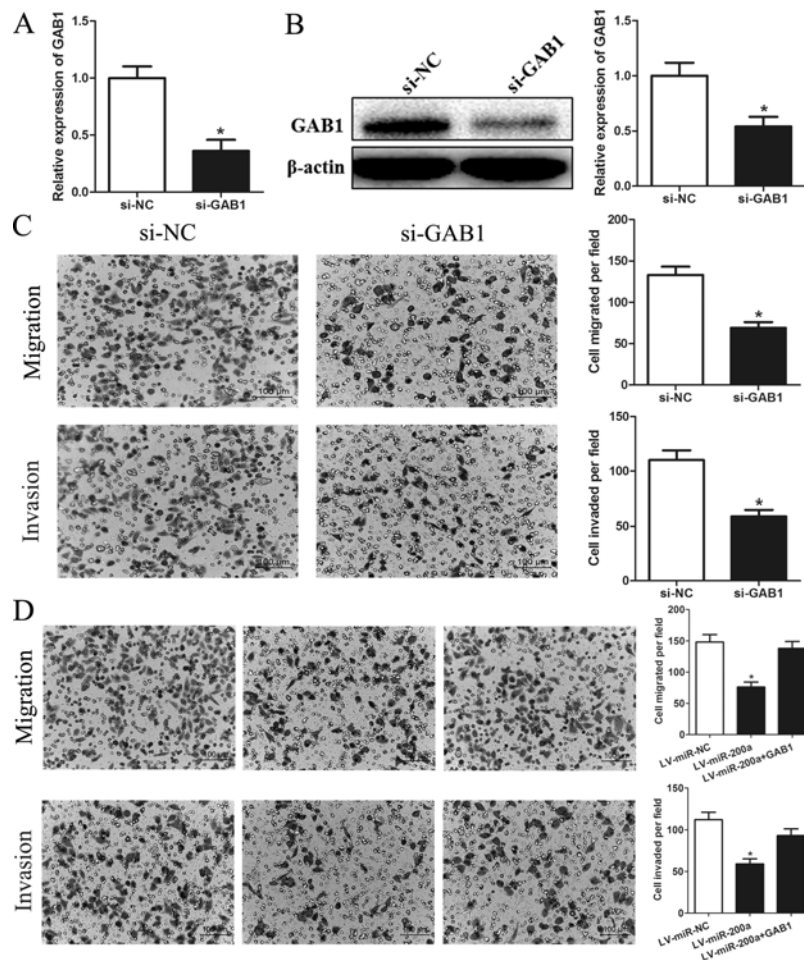


Figure 5. Alterations of GAB1 levels influence the effects of miR-200a on HCC cells. (A) Quantitation of GAB1 mRNA levels in SMMC-7721 cells by RT-PCR after transfection with siRNA targeting GAB1 (si-GAB1) or a negative control siRNA (si-NC). (B) Detection of GAB1 protein in SMMC-7721 cells by Western blot analysis after transfection with GAB1 siRNAs or si-NC. (C) GAB1 knockdown inhibited SMMC-7721 cell migration and invasion. (D) GAB1 reintroduction into SMMC-7721 cells partially rescued the miR-200a-mediated inhibition of cell migration and invasion. Migrated and invaded cells were counted in five randomly selected areas under a 200× microscope field. * $p < 0.05$ compared to the control.