

Carcinogenic Role and Clinical Significance of Histone H3-H4 Chaperone

Anti-silencing Function 1 B (ASF1B) in Lung Adenocarcinoma

Congkuan Song^{1#}, Yaolin Song^{2#}, Xiaoxia Wan^{2#}, Zhihong Zhao², Qing Geng¹

¹Department of Thoracic Surgery, Renmin Hospital of Wuhan University, Wuhan, China.

²Department of Thoracic Surgery, Ezhou Central Hospital, Ezhou, China.

#These authors contributed equally to this work.

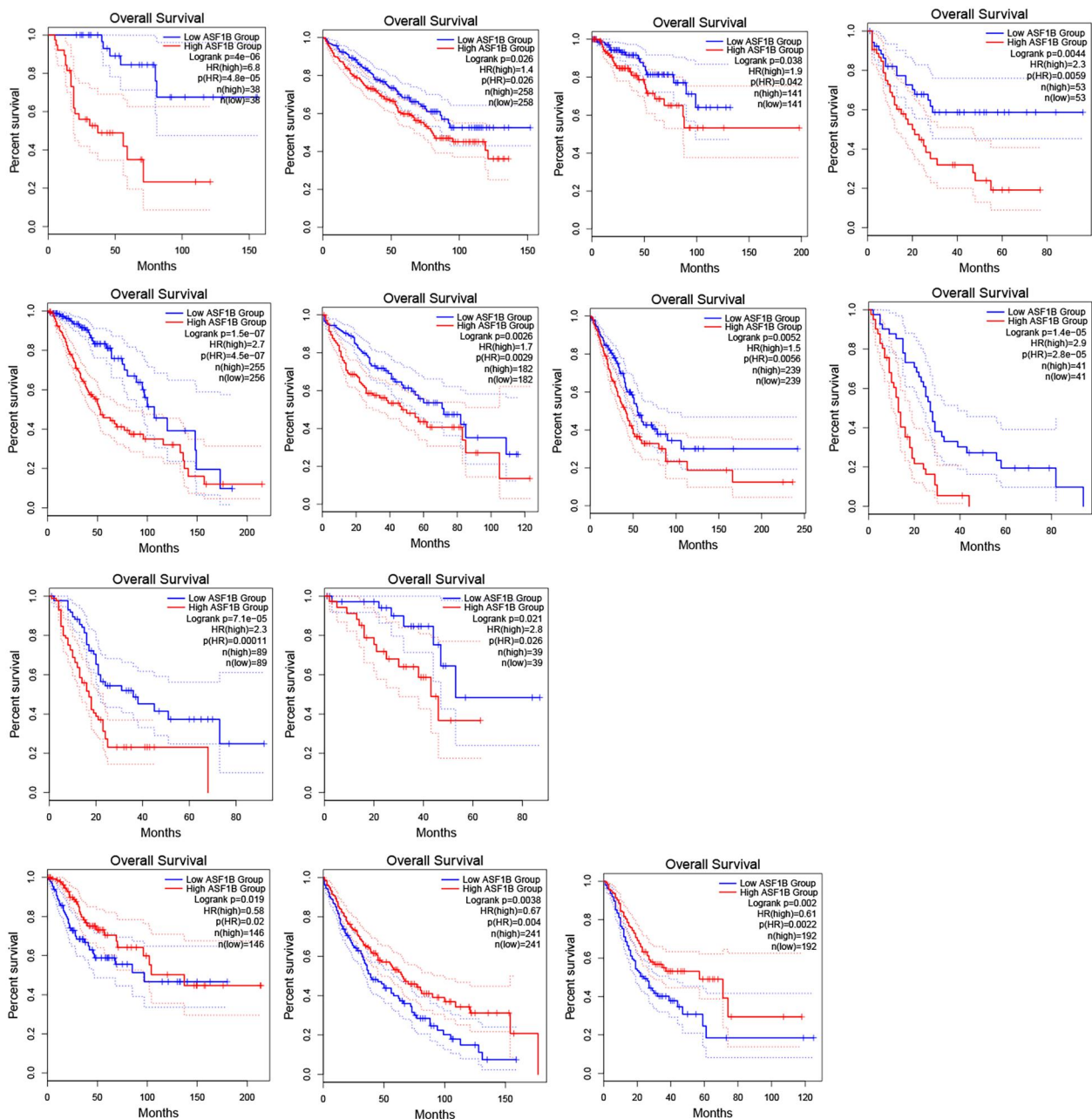


Figure S1: The relationship between the expression of ASF1B and the patient outcome from GEPIA2.

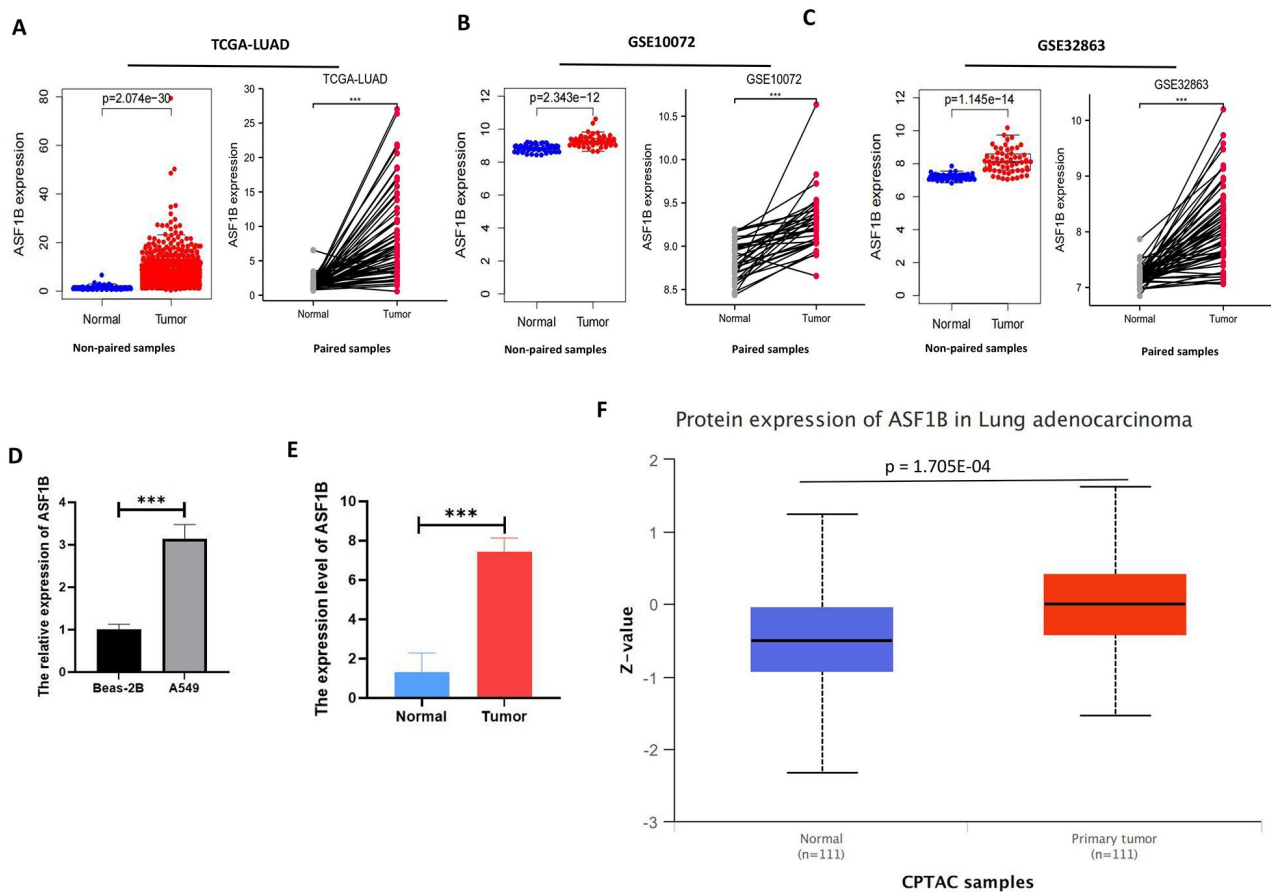
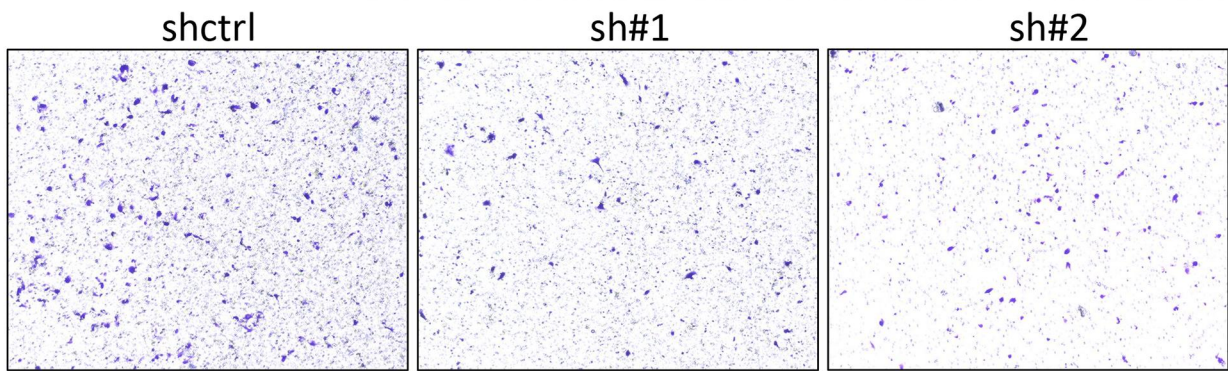


Figure S2: Comparison of ASF1B expression levels between LUAD tumor and normal tissues. (A-C) Comparison of transcription levels between LUAD tumor and normal tissues, including paired and unpaired samples, in the TCGA (A), GSE10072 (B), and GSE32863 (C) cohorts. (D) The mRNA expression of ASF1B in the A549 and Beas-2B cells. (E) qPCR analysis revealed the mRNA expression of ASF1B in the LUAD tumor and normal tissues. (F) UALCAN portal analysis revealed the protein levels of ASF1B in tumor versus normal tissues.

Migration



Invasion

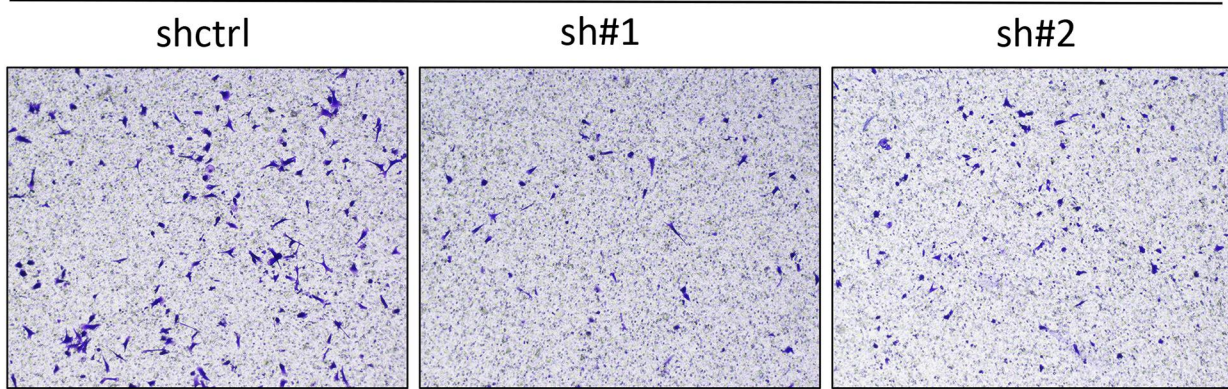


Figure S3: (A) Transwell assay revealed the migratory abilities of A549 cells. (B) Transwell assay revealed the invasive abilities of A549 cells.