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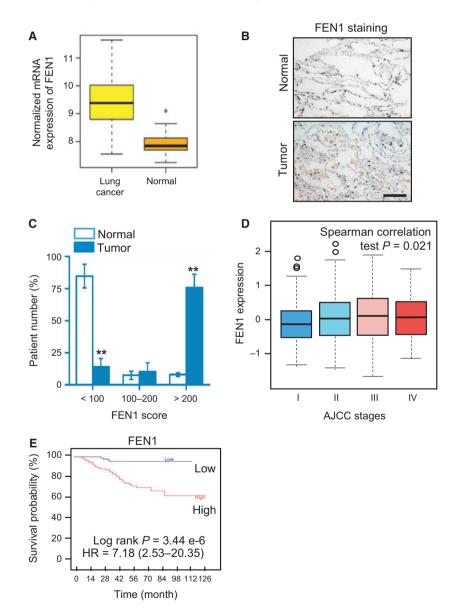
FEN1 promotes tumor progression and confers cisplatin resistance in non-small-cell lung cancer

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The authors of this article have supplied the following correction.

Figure 1E was mistakenly replaced by Fig. 1G of He *et al.* (2016) during the final assembly of the figures for review. The corrected figure is provided below. The authors apologize for the error.



Molecular Oncology **11** (2017) 1302–1303 © 2017 The Authors. Published by FEBS Press and John Wiley & Sons Ltd. This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited. **Fig. 1.** FEN1 overexpression was associated with lung cancer. (A) FEN1 expression analysis based on TCGA dataset showed that FEN1 mRNA levels were higher in lung cancer tissue than in normal tissue (*P < 0.01 vs control group). (B) FEN1 displayed significantly stronger staining (brown) in tumor specimens from clinical patients than from healthy counterparts. Immunohistochemistry was performed on formalin-fixed and paraffin-embedded tissues using antibodies against FEN1. Original magnification, ×400. Scale bars, 250 µm. (C). Number of patients' samples with FEN1 score > 200 was significantly higher in tumors than in normal tissues (*P < 0.01 vs control group). (D) FEN1 expression was correlated with the clinical stage of lung cancer. Spearman's correlation test, P = 0.021. (E) Kaplan–Meier analysis of survival of patients with lung cancer. Log rank P = 3.44 e-6.

References

He L, Zhang Y, Sun H, Jiang F, Yang H, Wu H, Zhou T, Hu S, Sekhar Kathera C, Wang X *et al.* (2016) Targeting DNA flap endonuclease 1 to impede breast cancer progression. *EBioMedicine* **14**, 32–43.