CORRECTION Open Access

Correction to: m6A mRNA methylation initiated by METTL3 directly promotes YAP translation and increases YAP activity by regulating the MALAT1-miR-1914-3p-YAP axis to induce NSCLC drug resistance and metastasis



Dan Jin^{1†}, Jiwei Guo^{2*†}, Yan Wu², Jing Du², Lijuan Yang², Xiaohong Wang³, Weihua Di⁴, Baoguang Hu⁵, Jiajia An⁶, Linggun Kong⁷, Lei Pan⁸ and Guoming Su⁹

Correction to: J Hematol Oncol 12:135 (2019) https://doi.org/10.1186/s13045-019-0830-6

The original article [1] contains errors in Fig. 2h, Fig. 2n and Fig. 6k:

- 1) In Fig. 2h, the protein band of YTHDF3 was mistakenly duplicated into the protein band of Cvr61.
- 2) In Fig. 2n, the image of Control+Vector treatment group was mistakenly duplicated into the siMETTL3 + YTHDF3 treatment group, and the image of METTL3 + siYTHDF3 treatment group was unintentionally duplicated into the METTL3 + YTHDF3 treatment group which were determined by transwell assay.
- 3) In Fig. 6k, the IHC image of ABCG2 was unintentionally duplicated onto the IHC image of ERCC1.

These errors were mistakenly introduced when organising the results; however, these errors have no bearing on the work's scientific conclusions as the statistical results are based on the correct pictures.

The authors would like to note the correct versions of each of the above-noted sub-figures ahead. The only changes are to the panels of Fig. 2h, Fig. 2n, and Fig. 6k, and the rest of the figures are identical to the published version; furthermore, no other errors were found after repeated checking of the published data.

The authors apologise to the Editor of Journal of Hematology & Oncology and to the readership for any inconvenience caused.

The original article can be found online at https://doi.org/10.1186/s13045-019-0830-6.

Full list of author information is available at the end of the article



© The Author(s). 2020 **Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/. The Creative Commons Public Domain Dedication waiver (http://creativecommons.org/publicdomain/zero/1.0/) applies to the data made available in this article, unless otherwise stated in a credit line to the data.

^{*} Correspondence: guojiwei0510@163.com

[†]Dan Jin and Jiwei Guo contributed equally to this work.

²Cancer research institute, Binzhou Medical University Hospital, Binzhou 256603. People's Republic of China

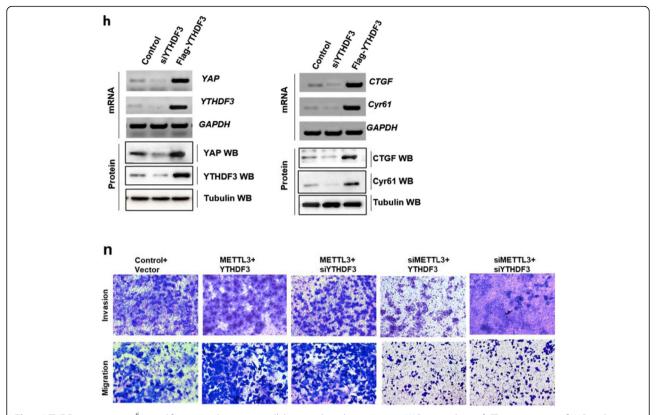


Fig. 2 YTHDF3 recognizes m^6A modification and promotes cellular growth and migration via YAP upregulation. **h** The expressions of YAP and its target genes, CTGF and Cyr61, were analyzed by RT-PCR and western blot. **n** The cellular invasion and migration growths were analyzed by transwell assay. Results were presented as mean \pm SD of three independent experiments. *P < 0.05 or **P < 0.01 indicates a significant difference between the indicated groups. NS, not significant

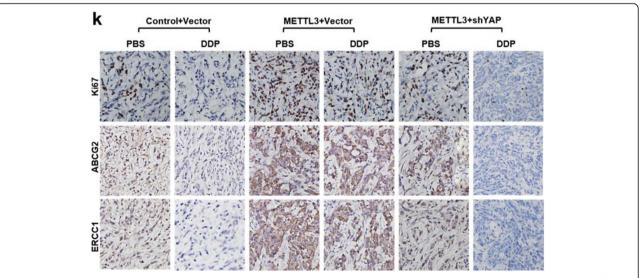


Fig. 6 The reduction of YAP m6A modification inhibits tumor growth and enhances DDP sensitivity in vivo. **k** The tumor nodules from control vector, METTL3 vector and METTL3 rough groups were treated with PBS or DDP every three days. The protein expression levels of Ki67, ABCG2 and ERCC1 were analyzed immunohistochemical staining assays (n = 5). Results were presented as mean \pm SD three independent experiments. **P < 0.01 indicates a significant difference between the indicated groups. NS, not significant

Author details

¹Clinical Medical Laboratory, Binzhou Medical University Hospital, Binzhou 256603, People's Republic of China. ²Cancer research institute, Binzhou Medical University Hospital, Binzhou 256603, People's Republic of China. ³Department of Thyroid and Breast Surgery, Binzhou Medical University Hospital, Binzhou 256603, People's Republic of China. ⁴Department of Pain, Binzhou Medical University Hospital, Binzhou 256603, People's Republic of China. ⁵Department of Gastrointestinal Surgery, Binzhou Medical University Hospital, Binzhou 256603, People's Republic of China. ⁶Department of Clinical Laboratory, Binzhou Medical University Hospital, Binzhou 256603, People's Republic of China. ⁷Department of Hepatobiliary Surgery, Binzhou Medical University Hospital, Binzhou 256603, People's Republic of China. ⁸Department of Respiratory and Critical Care Medicine, Binzhou Medical University Hospital, Binzhou 256603, People's Republic of China. ⁹Department of Nursing, Binzhou Polytechnic University, Binzhou 256603, People's Republic of China.

Published online: 03 August 2020

Reference

 Jin D, et al. m6A mRNA methylation initiated by METTL3 directly promotes YAP translation and increases YAP activity by regulating the MALAT1-miR-1914-3p-YAP axis to induce NSCLC drug resistance and metastasis. J Hematol Oncol. 2019;12:135 https://doi.org/10.1186/s13045-019-0830-6.