## CORRECTION

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# Correction: Knockdown of THOC1 reduces the proliferation of hepatocellular carcinoma and increases the sensitivity to cisplatin



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Following the publication of the original article [1], the author identified errors in the images of Figs. 4G and 6D which were unintentionally caused during the figure assembly process. Specifically:

- Figure 4G: R-loop-Vector.
- Figure 6D: R-loop-shNC and Ki67-shNC.

The corrected figures are provided below:

The corrections do not affect the overall results, discussion, or conclusion of the article.

The online version of the original article can be found at https://doi. org/10.1186/s13046-020-01634-7.

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**Fig. 4** THOC1 enhances tumorigenesis in vivo. **a** Relative tumor volume, (**b**) images of tumor, and (**c**) tumor weight of PLC/PRF/5 stably transfected with shNC or shTHOC1 plasmids in BALB/c nu/nu mice (Student's *t* test; \*P < 0.05). **d** Relative tumor volume, (**e**) images of tumor, and (**f**) tumor weight of THOC1-expressing HepG2 cells in nude mice were compared with those of the control vector-transfected HepG2 cells (Student's *t* test; \*P < 0.05). **d** Relative tumor volume, (**e**) images of tumor, and (**f**) tumor weight of THOC1-expressing HepG2 cells in nude mice were compared with those of the control vector-transfected HepG2 cells (Student's *t* test; \*P < 0.05), \*\*P < 0.01). **g** THOC1 protein expression in subcutaneous xenografts was determined by immunohistochemistry. R-loop level was estimated by S9.6 staining, and cell proliferative activity was measured by PCNA and Ki67 staining (Student's *t* test; \*\*P < 0.001). Scale bar, 50 µm





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**Fig. 6** Luteolin reduces HCC proliferation by targeting THOC1 in vivo and enhances the anti-tumor effect of cisplatin. **a** Tumor growth curve, (**b**) representative images of tumor, and (**c**) tumor weight of PLC/PRF/5 cells stably transfected with shTHOC1 or shNC in BALB/c nu/nu mice treated with 50 mg/kg luteolin or saline as control, respectively (one-way ANOVA; \*\**P* < 0.01). **d** immunohistochemistry staining indicates the expressions of THOC1, R-loop, and proliferation markers (PCNA and Ki67) in tumors (one-way ANOVA; \*\**P* < 0.001). **e** Tumor growth curve, (**f**) representative images of tumor, and (**g**) tumor weight of PLC/PRF/5-bearing BALB/c nu/nu mice. Luteolin or cisplatin treatment significantly suppressed tumor growth. Furthermore, luteolin can enhance the antitumor effect of cisplatin (one-way ANOVA; \**P* < 0.05, \*\**P* < 0.001).





**Fig. 6** Luteolin reduces HCC proliferation by targeting THOC1 in vivo and enhances the anti-tumor effect of cisplatin. **a** Tumor growth curve, (**b**) representative images of tumor, and (**c**) tumor weight of PLC/PRF/5 cells stably transfected with shTHOC1 or shNC in BALB/c nu/nu mice treated with 50 mg/kg luteolin or saline as control, respectively (one-way ANOVA; \*\*P < 0.01). **d** immunohistochemistry staining indicates the expressions of THOC1, R-loop, and proliferation markers (PCNA and Ki67) in tumors (one-way ANOVA; \*\*P < 0.001). **e** Tumor growth curve, (**f**) representative images of tumor, and (**g**) tumor weight of PLC/PRF/5-bearing BALB/c nu/nu mice. Luteolin or cisplatin treatment significantly suppressed tumor growth. Furthermore, luteolin can enhance the antitumor effect of cisplatin (one-way ANOVA; \*P < 0.05, \*P < 0.01, \*\*P < 0.001)

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#### References

1. Cai, Bai S, Wang Y et al. H. Knockdown of THOC1 reduces the proliferation of hepatocellular carcinoma and increases the sensitivity to cisplatin. J Exp Clin Cancer Res. 2020;39:135. https://doi.org/10.1186/s13046-020-01634-7

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