

Retraction

Retracted: High Expression of PTGR1 Promotes NSCLC Cell Growth via Positive Regulation of Cyclin-Dependent Protein Kinase Complex

BioMed Research International

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BioMed Research International has retracted the article titled “High Expression of PTGR1 Promotes NSCLC Cell Growth via Positive Regulation of Cyclin-Dependent Protein Kinase Complex” [1]. A series of very similar articles on shRNA and cancer cell lines was identified by Byrne and Labbé [2], and the intertextual distance between this article and an article in the series [3] is lower than expected by chance. The following concerns were found:

- (i) The supposed nontargeting control shRNA sequence, 5' GCGGAGGGTTTCAAAGAAATATCTCGAG-ATATTCTTTCAAACCCCTCCGCTTTTTT-3', targets TPD52L2 (NM_199360). The same sequence was used as a nontargeting control in other articles identified by Byrne and Labbé.
- (ii) The article is very similar in methods and structure to two other studies that also use this incorrect sequence [4, 5].

- [4] X. Zhao, M. Chen, and J. Tan, “Knockdown of ZFR suppresses cell proliferation and invasion of human pancreatic cancer,” *Biological Research*, vol. 49, no. 1, 2016.
- [5] C. Zhang, J. Fu, F. Xue et al., “Knockdown of ribosomal protein S15A induces human glioblastoma cell apoptosis,” *World Journal of Surgical Oncology*, vol. 14, no. 1, article no. 129, 2016.

The authors did not respond to requests for comment.

References

- [1] X. Huang, W. Zhou, Y. Zhang, and Y. Liu, “High expression of ptgr1 promotes NSCLC cell growth via positive regulation of cyclin-dependent protein kinase complex,” *BioMed Research International*, vol. 2016, Article ID 5230642, 2016.
- [2] J. A. Byrne and C. Labbé, “Striking similarities between publications from China describing single gene knockdown experiments in human cancer cell lines,” *Scientometrics*, vol. 110, no. 3, pp. 1471–1493, 2017.
- [3] Y. Wang, T. Jin, X. Dai, and J. Xu, “Lentivirus-mediated knock-down of CEP55 suppresses cell proliferation of breast cancer cells,” *BioScience Trends*, vol. 10, no. 1, pp. 67–73, 2016.