

Correction Notice: Improved HTGTS for CRISPR/Cas9 Off-target Detection

Jianhang Yin^{1,2}, Mengzhu Liu¹, Yang Liu¹ and Jiazhi Hu^{1,2,*}

¹The MOE Key Laboratory of Cell Proliferation and Differentiation, Genome Editing Research Center, School of Life Sciences, Peking University, Beijing, 100871, China

²Peking-Tsinghua Center for Life Sciences, Peking University, Beijing, 100871, China

*For correspondence: hujz@pku.edu.cn

In page 6 of our Bio-protocol paper <https://bio-protocol.org/e3229>, we notice some mistakes and the correction details are listed below:

Step E3: 80 should be 42.4

Step F is added:

On-beads ligation for bridge adapter

	1x (μl)
10x T4 DNA ligase buffer	8
Bridge adapter (50 μM)	1.6
T4 DNA ligase (5 U/μl)	4
50% (wt/vol) PEG8000	24
DNA-beads complex	42.4
Total	80

Set the reaction in a 1.5 ml tube in a rotator and ligate overnight at RT

Step G1 is added:

Add 80 μl 2x B&W buffer and 160 μl 1x B&W buffer. Put the beads against the 1.5 ml tube magnet stand and remove the supernatant. Wash the beads using 400 μl 1x B&W buffer three times and 400 μl dH₂O once. Resuspend the beads in 80 μl dH₂O.

References

1. Yin, J., Liu, M., Liu, Y. and Hu, J. (2019). [Improved HTGTS for CRISPR/Cas9 Off-target Detection](https://bio-protocol.org/e3229). *Bio-protocol* 9(9): e3229.