BMC Molecular Biology

CORRECTION

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Correction to: A protocol for custom CRISPR Cas9 donor vector construction to truncate genes in mammalian cells using pcDNA3 backbone

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Correction to: BMC Molecular Biology (2018) 19:3
https://doi.org/10.1186/s12867-018-0105-8

The original article [1] contains three erroneous mentions of usage of a restriction enzyme—*Bst*Z17I—in the Methods section as displayed in the following sentences:

1. […] Therefore, *FOXO3* gene fragments (PCR products) had pcDNA3 sequences on the ends that corresponded to upstream and downstream sequences of the utilized restriction sites (*Dra*III for Arm1 and *Bst*Z17I for Arm2) in pcDNA3.
2. […] The intermediate vector (with *FOXO3* Arm 1) was cut with *Bst*Z17I, which is on the other side of the neomycin resistance gene in the pcDNA3 plasmid compared to *FOXO3* Arm 1.
3. […] *FOXO3* Arm 2 was amplified with the primer pair specified in Table 1, producing a product that had sequences on each end that were identical to the sequences proximal to the *Bst*Z17I site in the intermediate *FOXO3* Arm 1 vector.

As such, the above three sentences should instead have stated the correct restriction enzyme—*Bsm*I—in place of where *Bst*Z17I was mentioned in each instance.

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The original article can be found online at https://doi.org/10.1186/s12867-018-0105-8.

Published online: 14 August 2019

Reference

1. Vazquez N, Sanchez L, Marks R, Martinez E, Fanniel V, Lopez A, Salinas A, Flores I, Hirschmann J, Gilkerson R, Schuenzel E, Dearth R, Halaby R, Innis-Whitehouse W, Keniry M. A protocol for custom CRISPR Cas9 donor vector construction to truncate genes in mammalian cells using pcDNA3 backbone. BMC Mol Biol. 2018;19:3. https://doi.org/10.1186/s12867-018-0105-8.

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