

CORRECTION

# Correction: Functional implications of hexameric assembly of RraA proteins from *Vibrio vulnificus*

Saemee Song, Seokho Hong, Jinyang Jang, Ji-Hyun Yeom, Nohra Park, Jaejin Lee, Yeri Lim, Jun-Yeong Jeon, Hyung-Kyoon Choi, Minho Lee, Nam-Chul Ha, Kangseok Lee

The images for Figs 4 and 5 are incorrectly switched. The image that appears as Fig 4 should be Fig 5, and the image that appears as Fig 5 should be Fig 4. The figure captions appear in the correct order. Please see the corrected Fig 4 and Fig 5 below.

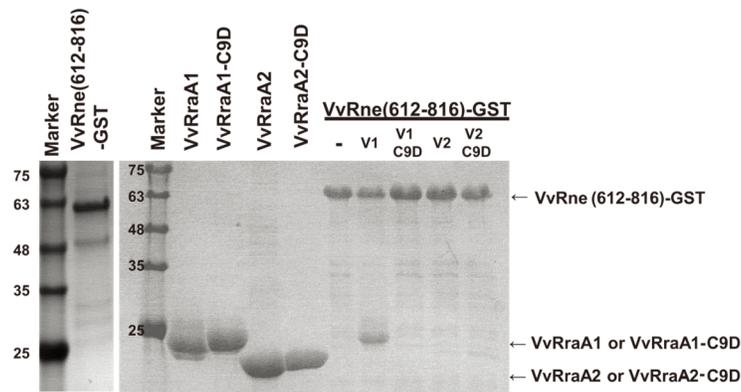


 OPEN ACCESS

**Citation:** Song S, Hong S, Jang J, Yeom J-H, Park N, Lee J, et al. (2018) Correction: Functional implications of hexameric assembly of RraA proteins from *Vibrio vulnificus*. PLoS ONE 13(1): e0191775. <https://doi.org/10.1371/journal.pone.0191775>

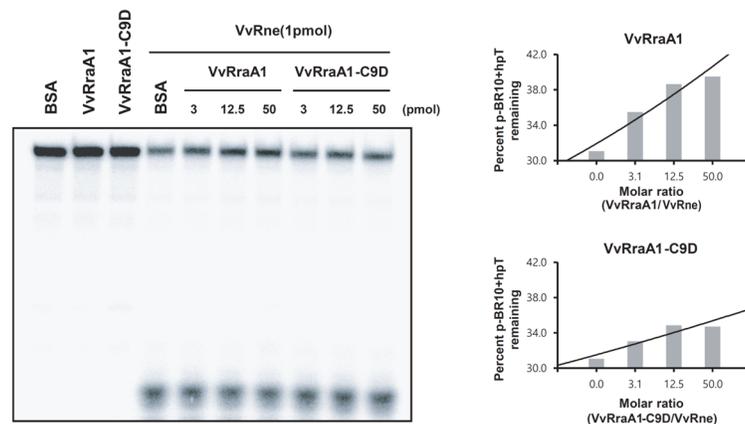
**Published:** January 19, 2018

**Copyright:** © 2018 Song et al. This is an open access article distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.



**Fig 4. Interactions of VvRNase E with VvRraA proteins.** Hexahistidine-tagged VvRraA1, VvRraA1-C9D, VvRraA2, VvRraA2-C9D, and the GST-fused VvRne (612–816 residues) were expressed and purified as described in the Methods section. The GST-fused VvRne protein was bound to GSH resin and incubated with VvRraA proteins and their C9D mutant proteins. Then, the proteins were eluted and the fractions were analyzed using SDS-PAGE. The protein bands were stained with Coomassie blue. Only VvRraA1 could tightly bind to VvRne.

<https://doi.org/10.1371/journal.pone.0191775.g001>



**Fig 5. Inhibition of VvRraA1 and VvRraA1-C9D on the cleavage of p-BR10+hpT by VvRNase E *in vitro*.** 0.5 pmol of 5'-end-labeled p-BR10+hpT RNA was incubated with 1 pmol of VvRne with varying concentrations of VvRraA1 and VvRraA1-C9D, 50 pmol of VvRraA1, or 50 pmol of BSA in 20  $\mu$ l of 1  $\times$  cleavage buffer at 37°C for 2 h for VvRne, VvRraA1 only, or BSA only controls. Samples were mixed with an equal volume of loading buffer, and then denatured at 65°C for 5 min and loaded onto a 12% polyacrylamide gel containing 8 M urea. The percentage of uncleaved p-BR10+hpT in the gel was quantitated using a phosphorimager and OptiQuant software.

<https://doi.org/10.1371/journal.pone.0191775.g002>

## Reference

1. Song S, Hong S, Jang J, Yeom J- H, Park N, Lee J, et al. (2017) Functional implications of hexameric assembly of RraA proteins from *Vibrio vulnificus*. PLoS ONE 12(12): e0190064. <https://doi.org/10.1371/journal.pone.0190064> PMID: 29261778