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# Author Correction: Design of a recombinant asparaginyl ligase for site-specific modification using efficient recognition and nucleophile motifs

Check for updates

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An earlier publication by Chua and colleagues reported a truncated construct of oaAEP1b, termed oaAEP1b-C247A-Δ351, which is similar to that reported in this study. In contrast to the previous study, oaAEP1b-C247A-aa55-351 is expressed in soluble form in *E. coli*, facilitated by an N-terminal fusion of a TrxA tag. This strategy eliminates the requirement for the denaturation and subsequent refolding of inclusion bodies using urea, thereby offering a more straightforward approach and ensuring the retention of protein activity.

While this prior publication was initially omitted from the reference list of this Article, the authors acknowledge that given the overlap between the two studies, citation of this earlier work is appropriate.

Chua N. et al. On the design of a constitutively active peptide asparaginyl ligase for facile protein conjugation. *FEBS Open Bio*, 13, 1195–1106 (2023).

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