



ERRATUM

Erratum to: Structural and functional analyses of human DDX41 DEAD domain

Yan Jiang¹, Yanping Zhu¹, Weicheng Qiu¹, Yong-Jun Liu², Genhong Cheng³, Zhi-Jie Liu^{1,4}✉, Songying Ouyang^{1,3}✉

¹ National Laboratory of Biomacromolecules, Institute of Biophysics, Chinese Academy of Sciences, Beijing 100101, China

² Baylor Research Institute, Baylor Scott and White Health, Dallas, TX 75246, USA

³ Department of Microbiology, Immunology and Molecular Genetics, University of California Los Angeles, Los Angeles, CA 90095, USA

⁴ Institute of Molecular and Clinical Medicine, Kunming Medical University, Kunming 650500, China

✉ Correspondence: zjliu@ibp.ac.cn (Z.-J. Liu), ouyangsy@ibp.ac.cn (Songying Ouyang)

ERRATUM TO: PROTEIN CELL 2016

DOI 10.1007/S13238-016-0351-9

In the original publication of this article Fig. 2 has been incorrectly published. The correct Fig. 2 is provided in this erratum.

The online version of the original article can be found under doi:[10.1007/s13238-016-0351-9](https://doi.org/10.1007/s13238-016-0351-9).

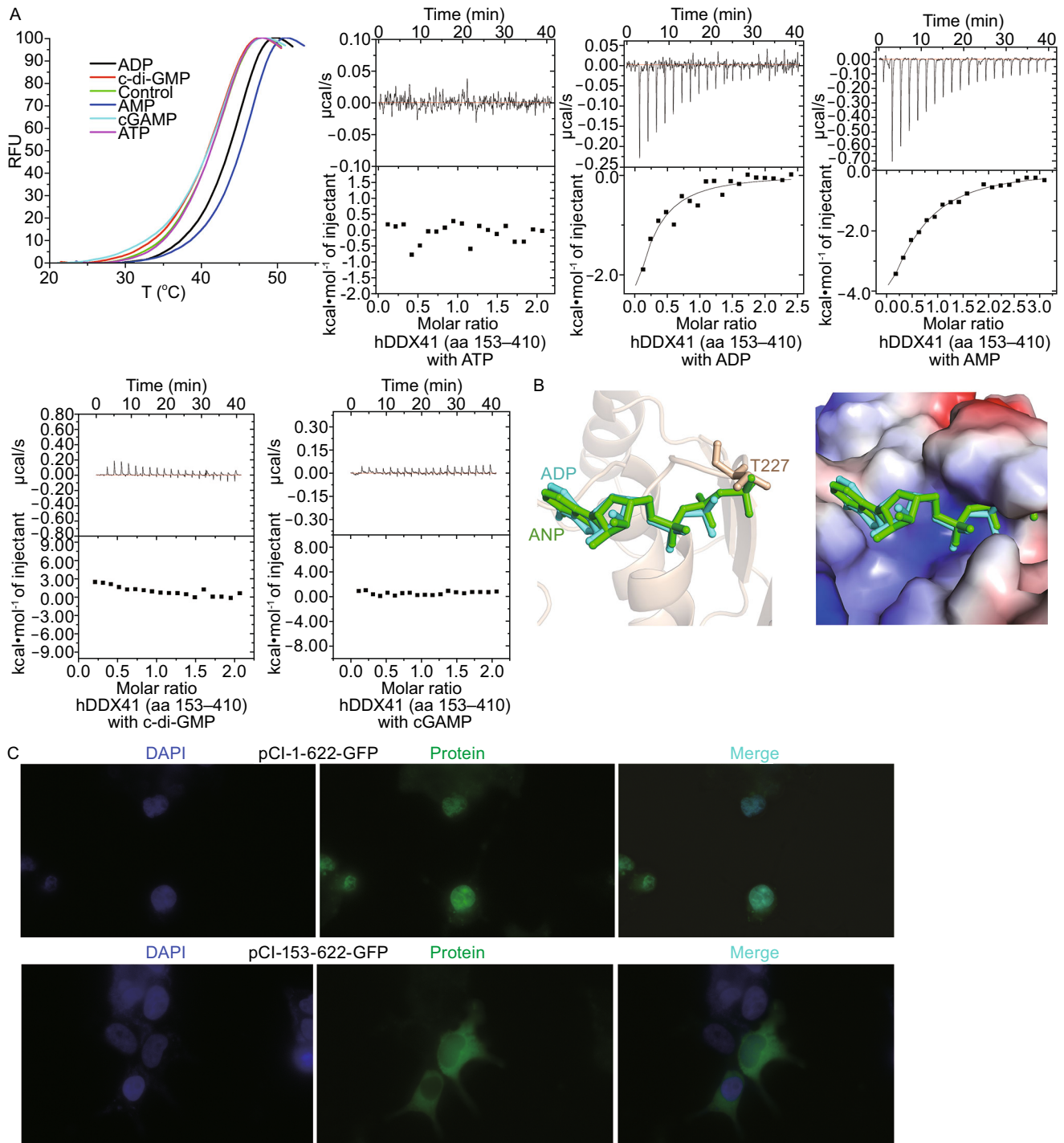


Figure 2. The binding of hDDX41 DEAD domain with different molecules and N-terminal region targets hDDX41 to the nucleus. (A) Thermal Shift Assay and Isothermal Titration Calorimetry of hDDX41 DEAD domain protein with ATP, ADP, AMP, c-di-GMP and cGAMP. **(B)** Left: the modeled ADP and ANP are colored in cyan and green. The γ -phosphate of ANP clashes with T227 of hDDX41. Right: surface electrostatic potential representation of the nucleotide binding pocket. Blue, positive potential; red, negative potential. The positively charged binding pocket is not big enough for ANP binding. **(C)** Fluorescence microscopy of HEK293T cells transfected with expression plasmids for GFP-tagged hDDX41 full length protein (1–622) and GFP-tagged hDDX41 N-terminal region deleted truncation (153–622). Nuclei are stained with DAPI.

OPEN ACCESS

This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and

reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.