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CORRECTION

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Correction: Photodynamic antitumor activity of aggregation-induced emission luminogens as chemosensitizers for paclitaxel by concurrent induction of apoptosis and autophagic cell death

Jia Wang,†[ab] Wenling Zhang,†[c] Ting Wu,†[b] Haisi Wu,[b] Yuan Zhang,[b] Siwan Wang,[b] You Ji,[b] Hui Jiang,[b] Ziting Zhang,[c] Chunming Tang,[b] Qiyun Tang,*[c] Xiaolin Li *[c] and Huae Xu *[ab]

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Correction for 'Photodynamic antitumor activity of aggregation-induced emission luminogens as chemosensitizers for paclitaxel by concurrent induction of apoptosis and autophagic cell death' by Jia Wang *et al.*, *Mater. Chem. Front.*, 2021, **5**, 3448–3457, https://doi.org/10.1039/D1QM00089F.

The authors regret that Fig. 2E was incorrect in the original article. The image showing the western blot data for p27 was duplicated in error. The corrected version of Fig. 2 is provided below.

The Royal Society of Chemistry apologises for these errors and any consequent inconvenience to authors and readers.

[a] *Jiangsu Key Laboratory of Molecular and Translational Cancer Research, Affiliated Cancer Hospital of Nanjing Medical University, Jiangsu Cancer Hospital, Jiangsu Institute of Cancer Research, Nanjing, 210009, China. E-mail: xuhuae@njmu.edu.cn*
[b] *Department of Pharmaceutics, School of Pharmacy, Nanjing Medical University, Nanjing 211166, China*
[c] *Department of Geriatric Gastroenterology, the First Affiliated Hospital of Nanjing Medical University, Nanjing, 210029, China. E-mail: tqy831@126.com, lxl@njmu.edu.cn*
† These authors contributed equally to this work.

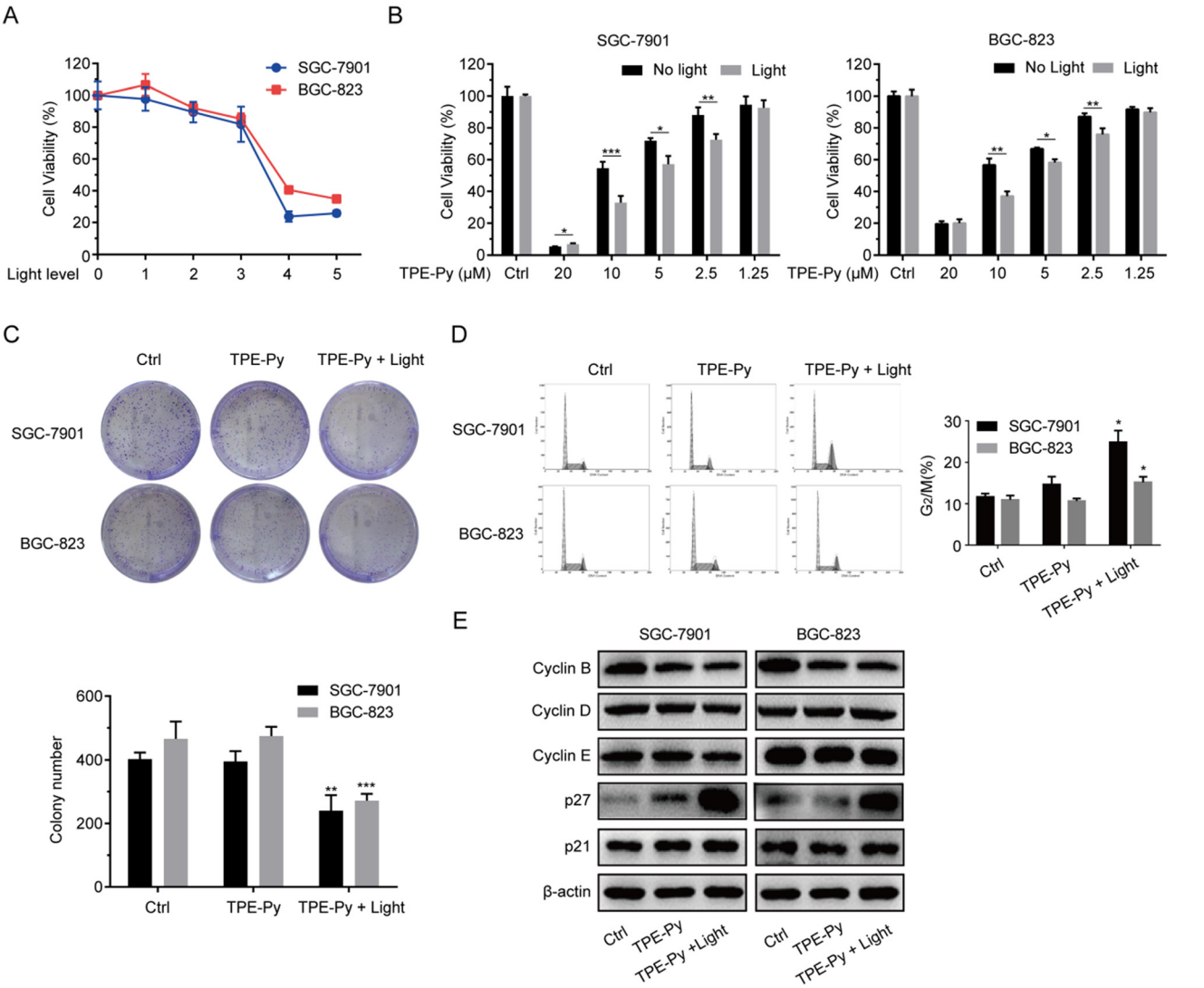

Fig. 2 **TPE-Py** photodynamically inhibits cell growth and induces cell cycle arrest in GC cells. (A) Cells were treated with different levels of white light (1, 5 lumens; 2, 50 lumens; 3, 150 lumens; 4, 350 lumens; 5, 1000 lumens) for 2 min. After 48 h, the cell viability was determined by MTS assay. (B) Cells were incubated with various concentrations of **TPE-Py** for 2 h, and then treated with or without 2 level white light. After 48 h, the cell viability was measured by MTS assay. (C) Representative images of colonies formed after treatment with 5 μM **TPE-Py** in the presence or absence of 2 level white light. (D) Cell cycle analysis of 5 μM **TPE-Py**-treated cells for 24 h in the presence or absence of 2 level white light by flow cytometry. (E) Representative immunoblots showed the effect of 5 μM **TPE-Py** in the presence or absence of 2 level white light on Cyclin B, Cyclin D, Cyclin E, p27 and p21. The results are expressed as mean ± SD of three independent experiments. $*p < 0.05$, $**p < 0.01$, $***p < 0.001$.