frontiers in Oncology

CORRECTION
published: 27 January 2021
doi: 10.3389/fonc.2020.583698

Corrigendum: CDC27 Promotes Tumor Progression and Affects PD-L1 Expression in T-Cell Lymphoblastic Lymphoma

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Edited and reviewed by:
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Specialty section:
This article was submitted to Cancer Molecular Targets and Therapeutics, a section of the journal Frontiers in Oncology

Received: 15 July 2020
Accepted: 29 September 2020
Published: 27 January 2021

Citation:
Song Y, Song W, Li Z, Song W, Wen Y, Li J, Xia Q and Zhang M (2021) Corrigendum: CDC27 Promotes Tumor Progression and Affects PD-L1 Expression in T-Cell Lymphoblastic Lymphoma. Front. Oncol. 10:583698. doi: 10.3389/fonc.2020.583698

Keywords: CDC27, T-cell lymphoblastic lymphoma, PD-L1, cell cycle, APC/C

A Corrigendum on

CDC27 Promotes Tumor Progression and Affects PD-L1 Expression in T-Cell Lymphoblastic Lymphoma

Song Y, Song W, Li Z, Song W, Wen Y, Li J, Xia Q and Zhang M (2020). Front. Oncol. 10:488. doi: 10.3389/fonc.2020.00488

In the original article, there was a mistake in **Figure 3** and **Figure 4** as published. In **Figure 3**, we put the wrong picture of EdU result in shNC group. In **Figure 4B**, we put the wrong picture of Apoptosis result in Control group. The correct **Figure 3** and **Figure 4** appear below. And in the Materials and Methods parts, we give a supplementary statement that cell lines used for Immunofluorescence were transfected with plasmid without GFP tag.

The authors apologize for these errors and state that this does not change the scientific conclusions of the article in any way. The original article has been updated.

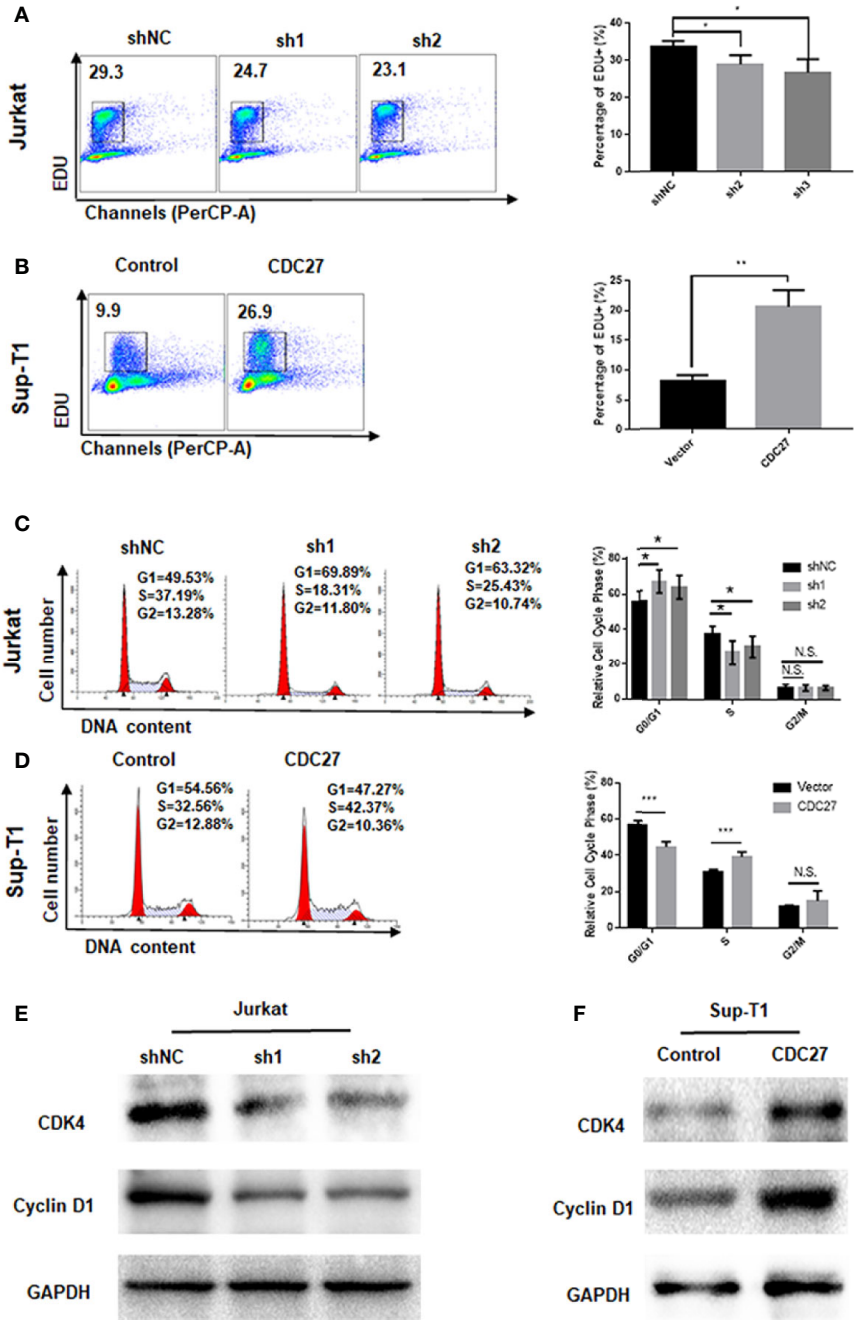

FIGURE 3 | CDC27 influence the G1/S phase transition. **(A, B)** Cell proliferation were assessed by EdU incorporation assay. Data are representative of at least three independent experiments. $^{**}P < 0.01$, $^{*}P < 0.05$. **(C, D)** Flow cytometry was used to examine the cell cycle by PI staining of both Jurkat and Sup-T1 cells. Images and qualification of the cell cycle distribution in three independent experiments are shown. $^{*}P < 0.05$, $^{***}P < 0.001$. N.S., not significant. **(E, F)** Western blot was performed to detect the expression levels of cell cycle related proteins in both Jurkat and Sup-T1 cells, respectively.

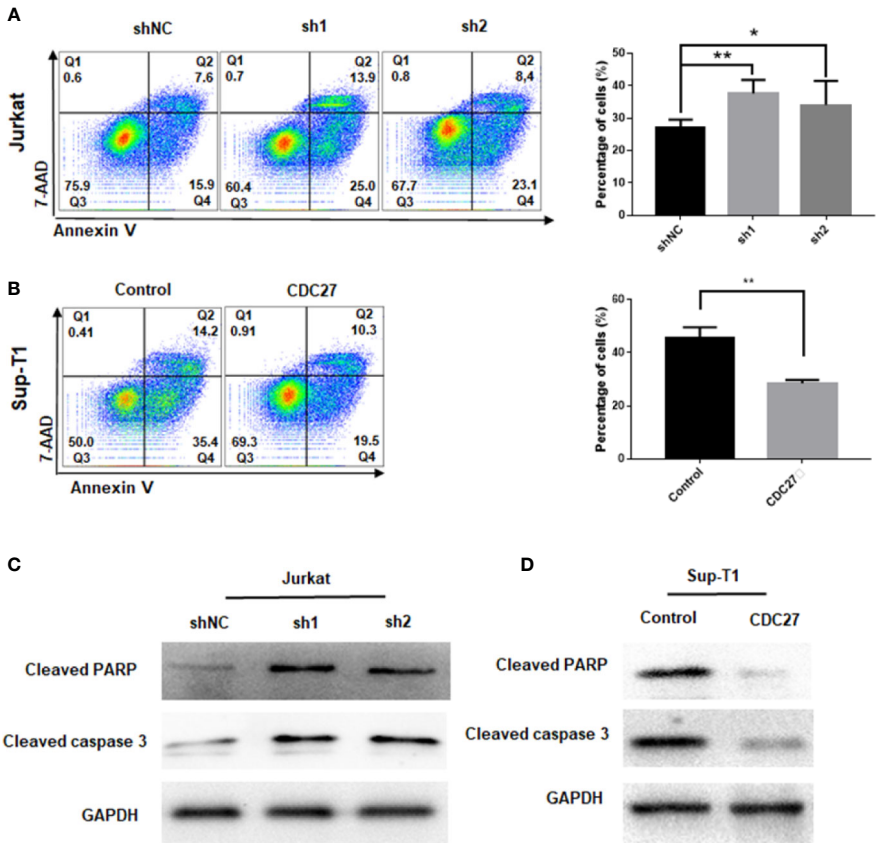

FIGURE 4 | CDC27 inhibits cell apoptosis in T-LBL cells. **(A, B)** Flow cytometry was used to examine the apoptosis as the sum of both Q2 and Q4 quadrants (early + late apoptosis) by Annexin V/7-AAD staining of both Jurkat and Sup-T1 cells. Apoptosis rates were expressed as the mean (Q2 + Q4) ± SD of values from experiments performed in triplicate by using Student's t-test. $^{*}P < 0.05$, $^{**}P < 0.01$. **(C, D)** Western blot was performed to detect the expression levels of apoptotic related proteins in both Jurkat and Sup-T1 cells, respectively.