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AUTHOR CORRECTION

Correction for Tun-Yhong et al., "Tenofovir Disoproxil Fumarate Is a New Substrate of ATP-Binding Cassette Subfamily C Member 11"

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Volume 61, no. 4, e01725-16, 2017, https://doi.org/10.1128/AAC.01725-16. Page 5, Figure 4: the order of magnitude of TDF in Fig. 4C should read 1.75×10^4 instead of 1.75×10^3 and that of MTX in Fig. 4D should read 1.6×10^4 instead of 1.6×10^3. The correct figure is shown below.

Citation Tun-Yhong W, Chinpaisal C, Pamonsinlapatham P, Kaewkitichai S. 2017. Correction for Tun-Yhong et al., "Tenofovir disoproxil fumarate is a new substrate of ATP-binding cassette subfamily C member 11." Antimicrob Agents Chemother 61:e01753-17. https://doi.org/10.1128/AAC.01753-17.

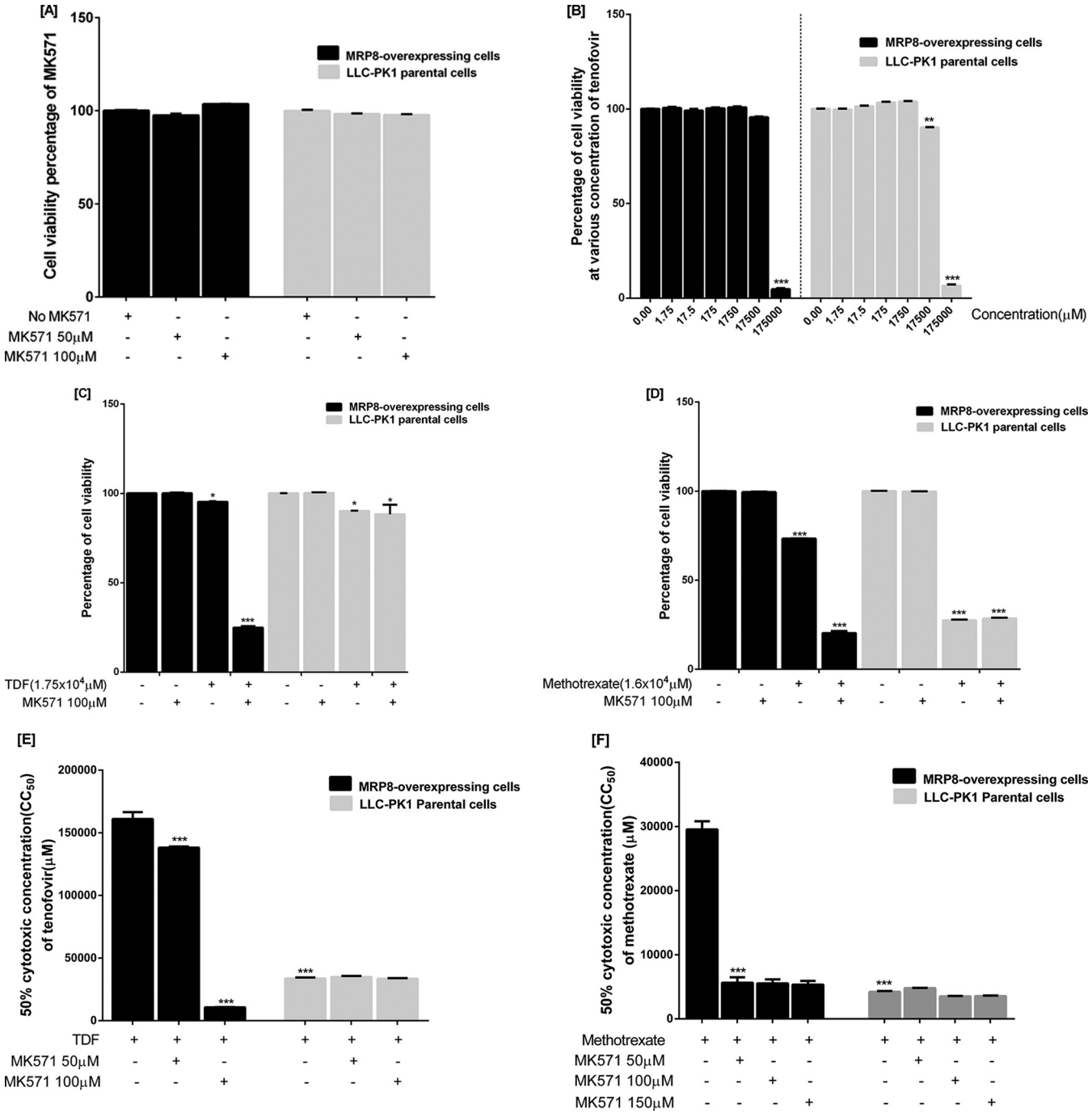

FIG 4 Cell viability assays with TDF and methotrexate in the presence and absence of the specific inhibitor MK-571. (A) Specific inhibitor MK-571 at various concentrations did not reduce MRP8-overexpressing and parental cell viability. (B) Cytotoxic effects of TDF on MRP8-overexpressing and parental cells. (C) MK-571 further reduced viability of the MRP8-overexpressing cells, but not parental cells, treated with TDF. (D) MK-571 also enhanced cytotoxicity of methotrexate only in MRP8-overexpressing cells. (E and F) Cytotoxicity assays showing methotrexate and TDF concentrations that reduced cell viability by 50% (CC_{50}) in MRP8-overexpressing LLC-PK1 or parental cells with or without the specific inhibitor MK-571. Statistical significance was analyzed by a two-way ANOVA multiple comparison assuming equal variance (*, $P < 0.01$; **, $P < 0.001$; ***, $P < 0.0001$). All values are the means ± standard deviations from five independent experiments.