

CORRECTION

# Correction: Auranofin Inhibits Retinal Pigment Epithelium Cell Survival through Reactive Oxygen Species-Dependent Epidermal Growth Factor Receptor/ Mitogen-Activated Protein Kinase Signaling Pathway

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There are a number of errors in Table 1. The “Application” value for “P38MAPK” in line 4 should read “WB”. The “Dilution” value for “MAPKAPK2” in line 13 should read “1:1000 (WB)”. The “Dilution” value for “HSP27” in line 15 should read “1:1000”. Please view the correct [Table 1](#) here.



## OPEN ACCESS

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**Table 1. Primary antibodies used for immunodetection.**

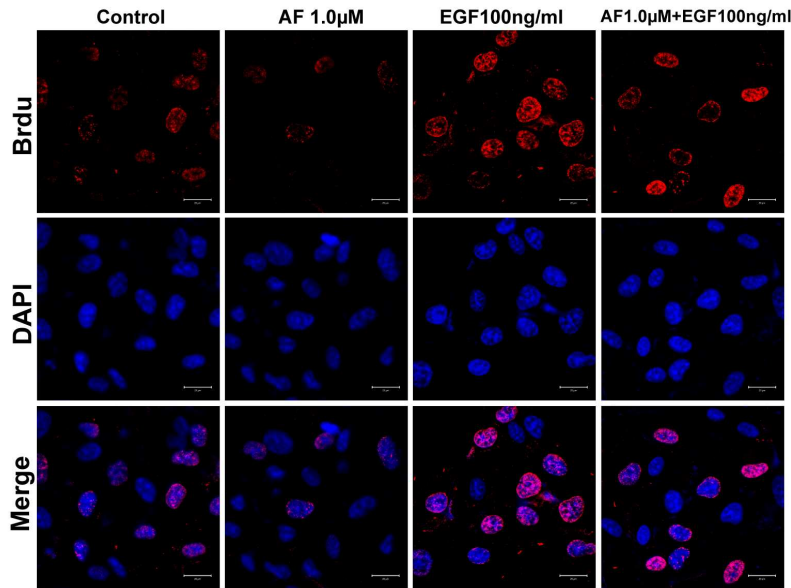
Name	Species	Manufacturer	Product number	Application	Dilution
pEGFR	Rabbit	Cell Signaling	3777	WB, IF	1:1000(WB),1:100(IF)
EGFR	Rabbit	Cell Signaling	4267	WB, IF	1:1000(WB),1:100(IF)
pP38MAPK	Rabbit	Cell Signaling	4511	WB, IF	1:1000(WB),1:100(IF)
P38MAPK	Rabbit	Cell Signaling	9212	WB	1:1000
pJNK	Rabbit	Cell Signaling	4668	WB	1:1000
pJNK	Mouse	Cell Signaling	9255	IF	1:100
JNK	Rabbit	Cell Signaling	9252	WB	1:1000
pERK	Rabbit	Cell Signaling	4370	WB, IF	1:1000(WB),1:100(IF)
ERK	Rabbit	Cell Signaling	9102	WB	1:1000
p-c-Jun	Rabbit	Cell Signaling	3270	WB, IF	1:1000(WB),1:100(IF)
c-Jun	Rabbit	Cell Signaling	9165	WB	1:1000
pMAPKAPK2	Rabbit	Cell Signaling	3007	WB, IF	1:1000(WB),1:100(IF)
MAPKAPK2	Rabbit	Cell Signaling	3042	WB	1:1000
pHSP27	Rabbit	Cell Signaling	9709	WB, IF	1:1000(WB),1:100(IF)
HSP27	Mouse	Cell Signaling	2402	WB	1:1000
BrdU	Mouse	Proteintech	66241	IF	1:500

WB, Western blot; IF, immunofluorescence.

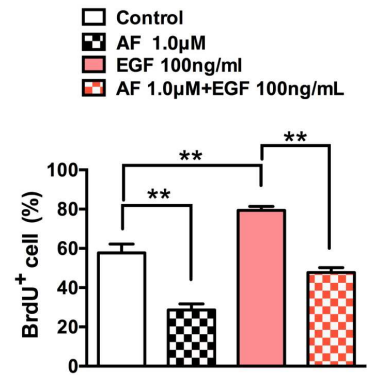
doi:10.1371/journal.pone.0172599.t001

In Fig 5, Fig 5C shows the incorrect image under the “AF 1.0  $\mu$ M” header for “0h”. Please view the correct [Fig 5](#) here.

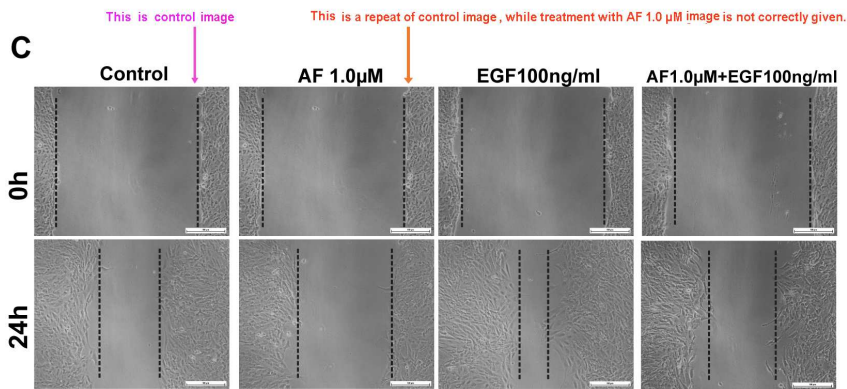
A



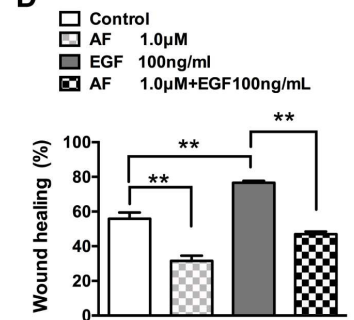
B



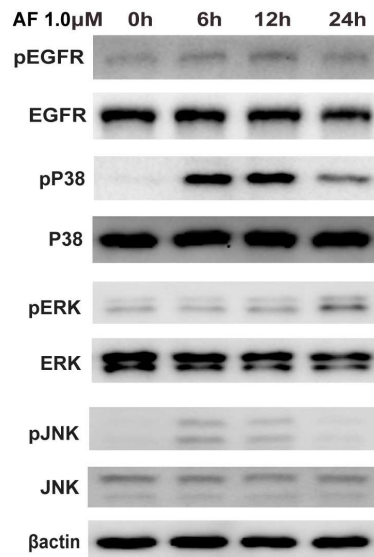
C



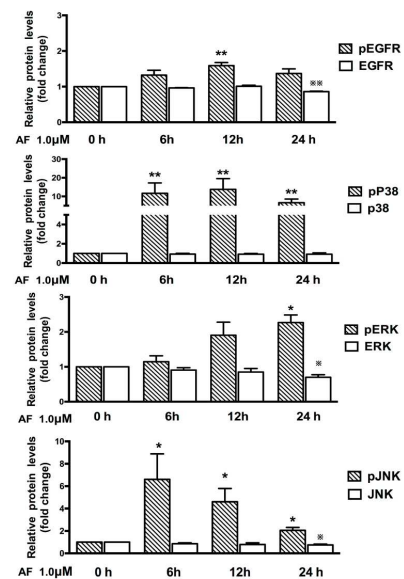
D



E



F



**Fig 5. Auranofin inhibits EGF-dependent proliferation and migration of ARPE-19 cells.** (A) Immunofluorescence microphotographs of proliferating ARPE-19 cell with BrdU (red) and DAPI (blue) staining after ARPE-19 cells were untreated or treated with AF (1.0  $\mu$ M) in the absence or presence of EGF (100ng/ml) for 24 hours and then subjected to BrdU labeling for 4 hours, followed by immunostaining with anti-BrdU antibody and DAPI. Scale bar = 20 $\mu$ m. (B) Quantitation data of the number of BrdU<sup>+</sup> cells shown in panel A. (C) ARPE-19 cells were subjected to wound healing assay, and then were left untreated or treated with AF (1.0  $\mu$ M) in the absence or presence of EGF (100 ng/ml) for 24 hours. Scale bar = 100 $\mu$ m. (D) Quantitation of the results shown in panel C. (E) ARPE-19 cells were treated with 1.0  $\mu$ M AF for 6, 12 and 24 hours. Cell lysates were subjected to Western blot for determination of total and phosphorylated EGFR, P38MAPK, ERK and JNK proteins.  $\beta$ -actin was used as a loading control. (F) Quantitative data of Western blot results shown in panel E from three experiments. The levels of the phosphorylated protein were compared with the control, \* P < 0.05, \*\* P < 0.01. The levels of the total protein were compared with the control, \*P < 0.05, \*\*P < 0.01. All data are mean  $\pm$  SEM.

doi:10.1371/journal.pone.0172599.g001

## Reference

1. Chen X, Tzekov R, Su M, Hong H, Min W, Han A, et al. (2016) Auranofin Inhibits Retinal Pigment Epithelium Cell Survival through Reactive Oxygen Species-Dependent Epidermal Growth Factor Receptor/Mitogen-Activated Protein Kinase Signaling Pathway. PLoS ONE 11(11): e0166386. doi:[10.1371/journal.pone.0166386](https://doi.org/10.1371/journal.pone.0166386) PMID: [27846303](https://pubmed.ncbi.nlm.nih.gov/27846303/)