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CORRIGENDUM

Emodin-Induced Autophagy Against Cell Apoptosis Through the PI3K/AKT/mTOR Pathway in Human Hepatocytes [Corrigendum]

Zheng X, Yang S, Zhang R, Wang S, Li G, Zhou S. *Drug Des Devel Ther*. 2019;13:3171–3180.

The authors have advised due to an error that occurred inadvertently at the time of figure assembly, Figure 5J is incorrect. The correct Figure 5 is shown below.

The authors apologize for this error and advise since this image only displays the cell morphology and does not perform a semi-quantitative analysis, it does not affect the other results of Figure 5, nor does it affect the conclusion of the article.

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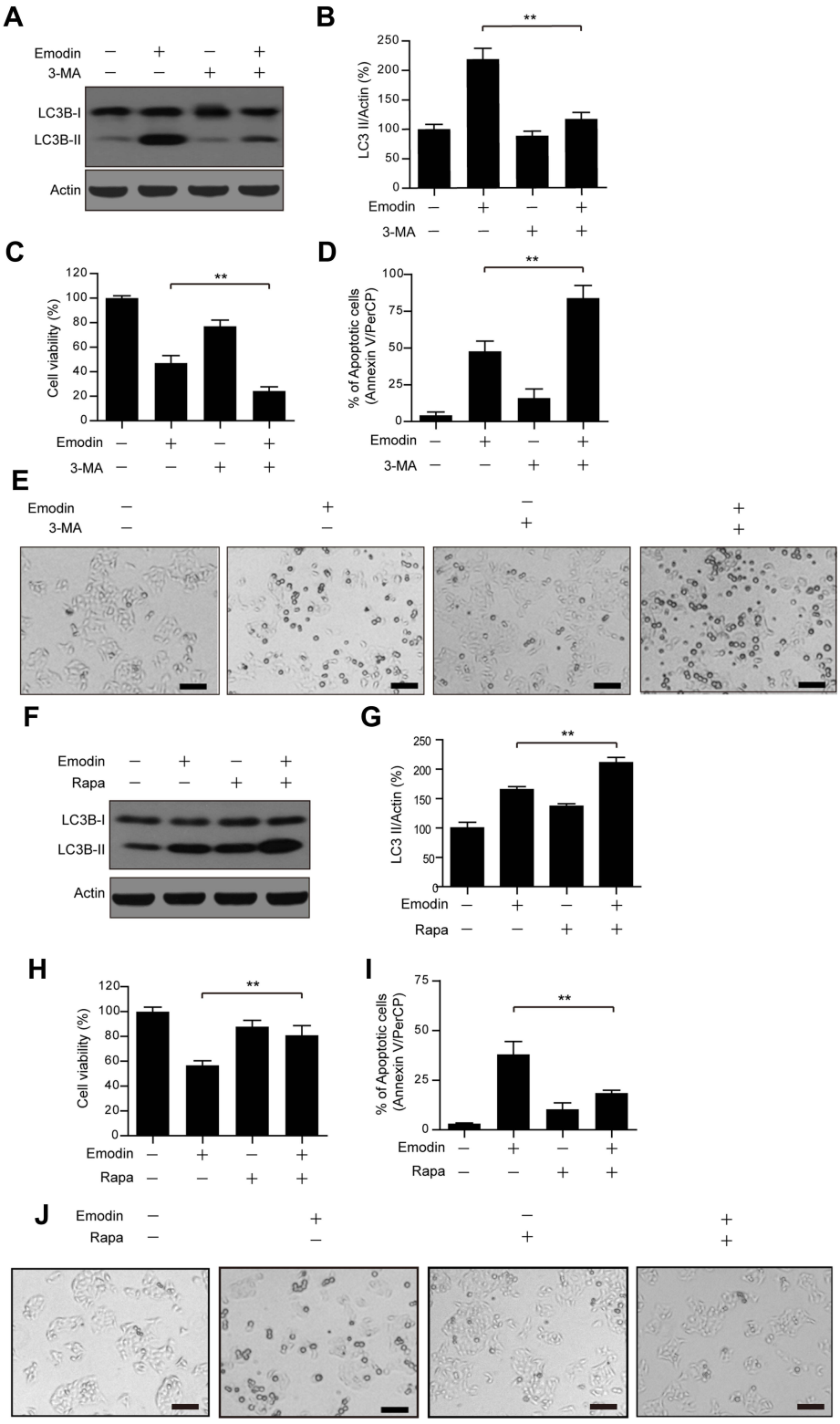

Figure 5 Emodin-induced autophagy played a protective role in L02 cells. (**A**) Cells were treated with or without emodin (40 μM) in the absence or presence of 3-MA (5 mM) for 24 h. The expression of LC3B-I/II was analysed by Western blot. (**B**) The intensities of LC3B-II in (**A**) normalized to actin were statistically analysed and represented as the mean ± SD for 3 independent experiments. ***P*<0.01. (**C**) Cell viability in (**A**) was determined by CCK-8 assay. Data are presented as the means ± SDs for 3 independent experiments. ***P*<0.01. (**D**) Cell apoptosis in (**A**) was detected by flow cytometry using annexin V/PerCP staining. Data are presented as the means ± SDs for 3 independent experiments. (**E**) Representative photos depicting the morphology of L02 cells treated in (**A**) Scale bars: 100 μm. (**F**) Cells were treated with or without emodin (40 μM) in the absence or presence of rapamycin (Rapa; 100 nM) for 24 h. The expression of LC3B-I/II was analysed by Western blot. (**G**) The intensities of LC3B-II in (**F**) normalized to actin were statistically analysed and are represented as the mean ± SD for 3 independent experiments. ***P*<0.01. (**H**) Cell viability in (**F**) was determined by CCK-8 assay. Data are presented as the means ± SDs for 3 independent experiments. ***P*<0.01. (**I**) Cell apoptosis in (**F**) was detected by flow cytometry using annexin V/PerCP staining. Data are presented as the means ± SDs for 3 independent experiments. (**J**) Representative photos depicting the morphology of L02 cells treated in (**F**). Scale bars: 100 μm.