

CORRECTION

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# Correction: Antimicrobial peptide moricin induces ROS mediated caspase-dependent apoptosis in human triple-negative breast cancer via suppression of notch pathway

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In this article [1], the figure legends for Figures 4 and 5 was interchanged. The correct Figs. 4 and 5 with legends are given in this erratum.

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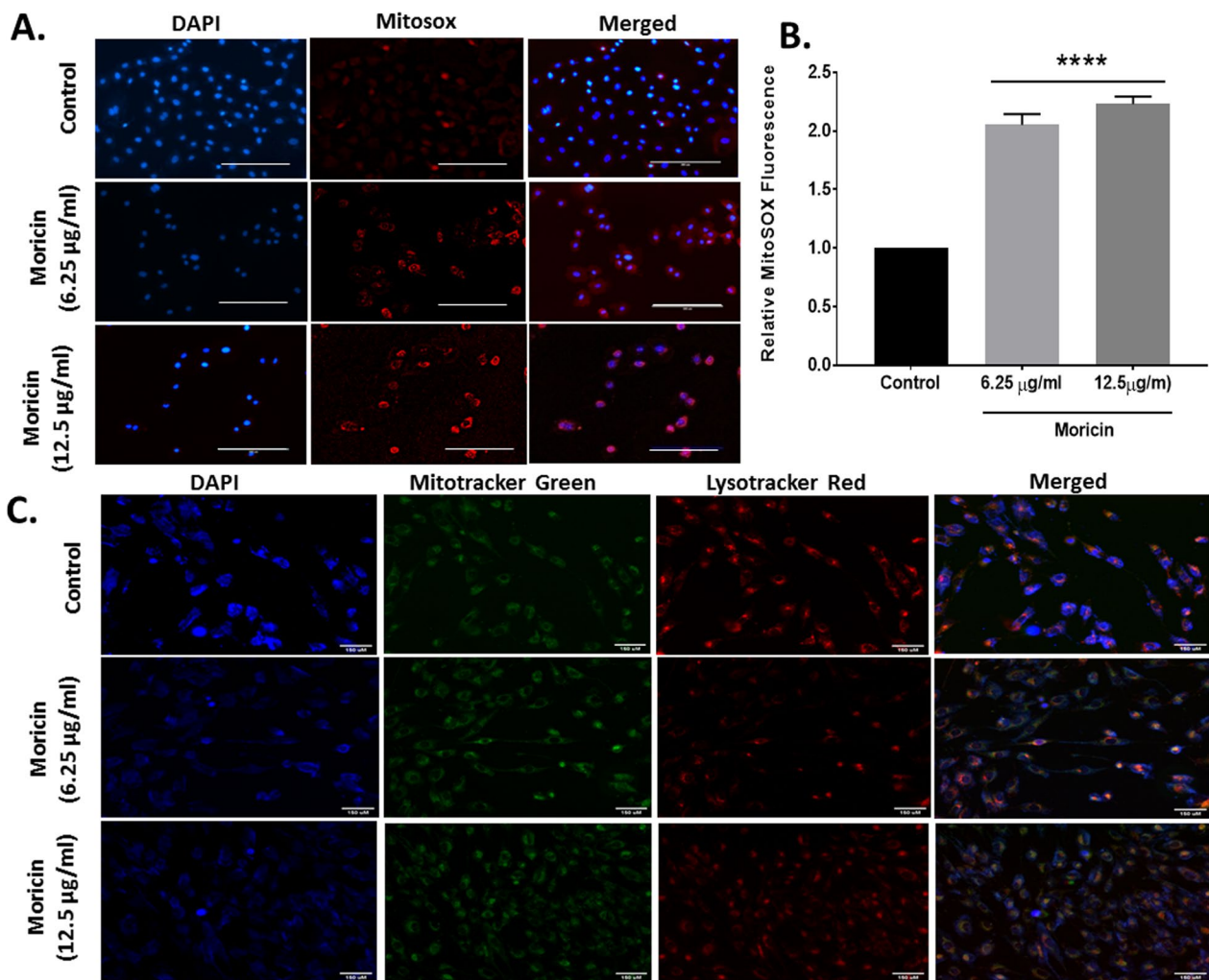
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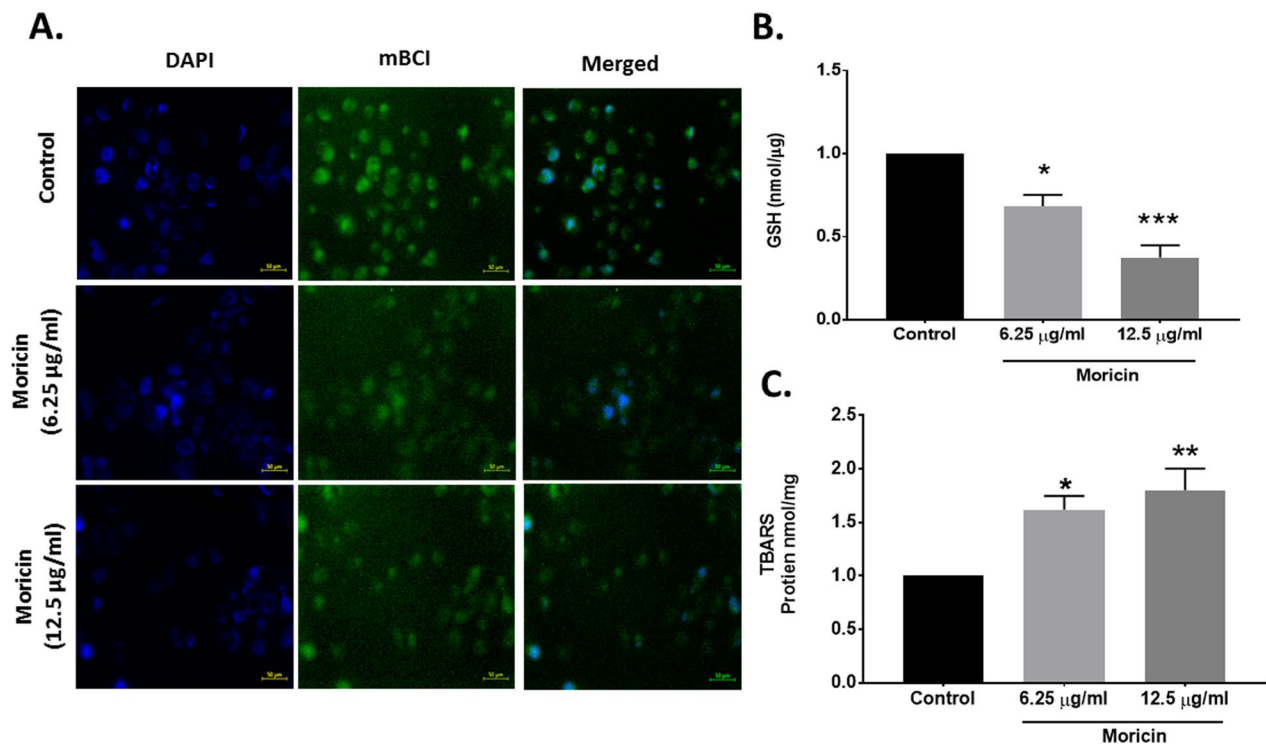
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**Fig. 4** Effect of moricin peptide treatment on structural damages of mitochondria and lysosomes and induced mitochondrial ROS generations in MDA-MB-231 cells. **A** Mitochondrial ROS was determined by microscopy with mitoSOX staining of moricin treated and untreated MDA-MB-231 cells treated with 6.25 µg/ml and 12.5 µg/ml moricin peptide. Images were taken with florescent microscopy (Zeiss Microsystems, GmbH, Germany) at  $\times 20$  magnification at scale bar 100 µm. **B** Represents level of mitochondrial ROS through relative mitoSOX florescence. **C** Florescent images of the MDA-MB-231 cells treated with 6.25 µg/ml and 12.5 µg/ml moricin peptide. Staining was done for the nucleus (blue), mitochondria (green) and lysosome (red) using Hoechst 33342 (10 µg/ml), Mitotracker Green FM (75 nM), and Lysotracker Red (100 nM). Cell shrinkage and mitochondrial disruption are seen for cells treated with moricin but not in the control cells. Images were taken with florescent microscopy (Zeiss Microsystems, GmbH, Germany) at  $\times 20$  magnification at scale bar 150 µm. Results are the mean  $\pm$  S.E from three independent experiments and statistical analysis was determined one-way ANOVA test followed by Dunnett's post hoc comparison test \*\*\*\* $p < 0.0001$  vs untreated cells



**Fig. 5** Effect of moricin peptide treatment on Glutathione and TBARS level in MDA-MB-231 cells. **A** Microscopy analysis were performed in MDA-MB 231 cells to measure GSH in cells treated with 6.25 µg/ml and 12.5 µg/ml concentrations of moricin using Monochlorobimane (mBCI) staining. Images were taken with florescent microscopy (Zeiss Microsystems, GmbH, Germany) at  $\times 20$  magnification at scale bar 50 µm. **B** Represents the level of GSH (nmol/µg) in MDA-MB-231 cells treated with 6.25 µg/ml and 12.5 µg/ml concentrations of moricin. **C** Represents the level of TBARS (µM) in MDA-MB-231 cells treated with 6.25 µg/ml and 12.5 µg/ml concentrations of moricin. Results are the mean  $\pm$  S.E from three independent experiments and statistical analysis was determined one-way ANOVA test followed by Dunnett's post hoc comparison test \* $p < 0.05$ , and \*\*\* $p < 0.001$  vs untreated cells

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#### Reference

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