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ERRATUM

Erratum to: RAGE-binding S100A8/A9 promotes the migration and invasion of human breast cancer cells through actin polymerization and epithelial–mesenchymal transition

Chonggao Yin[1] · Hongli Li[2] · Baogang Zhang[3,4] · Yuqing Liu[3,4] · Guohua Lu[1] · Shijun Lu[3,4] · Lei Sun[3,4] · Yueliang Qi[3,4] · Xiaolong Li[3,4] · Weiyi Chen[3,4]

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In the original publication, the images in Fig. 3 were mistakenly selected from other experiments in which similar procedures were performed. The corrected Fig. 3 is given in this erratum.

The online version of the original article can be found under doi:10.1007/s10549-013-2737-1.

✉ Baogang Zhang
zhangbg@wfmc.edu.cn; zbg0903@hotmail.com

[1] College of Nursing, Weifang Medical University, Weifang 261053, China

[2] Medicine Research Center, Weifang Medical University, Weifang 261053, China

[3] Department of Pathology, Weifang Medical University, Weifang 261053, China

[4] Department of Pathology, Key Clinical Specialty for Pathology of Shandong Province, Affiliated Hospital of Weifang Medical University, Weifang 261053, China

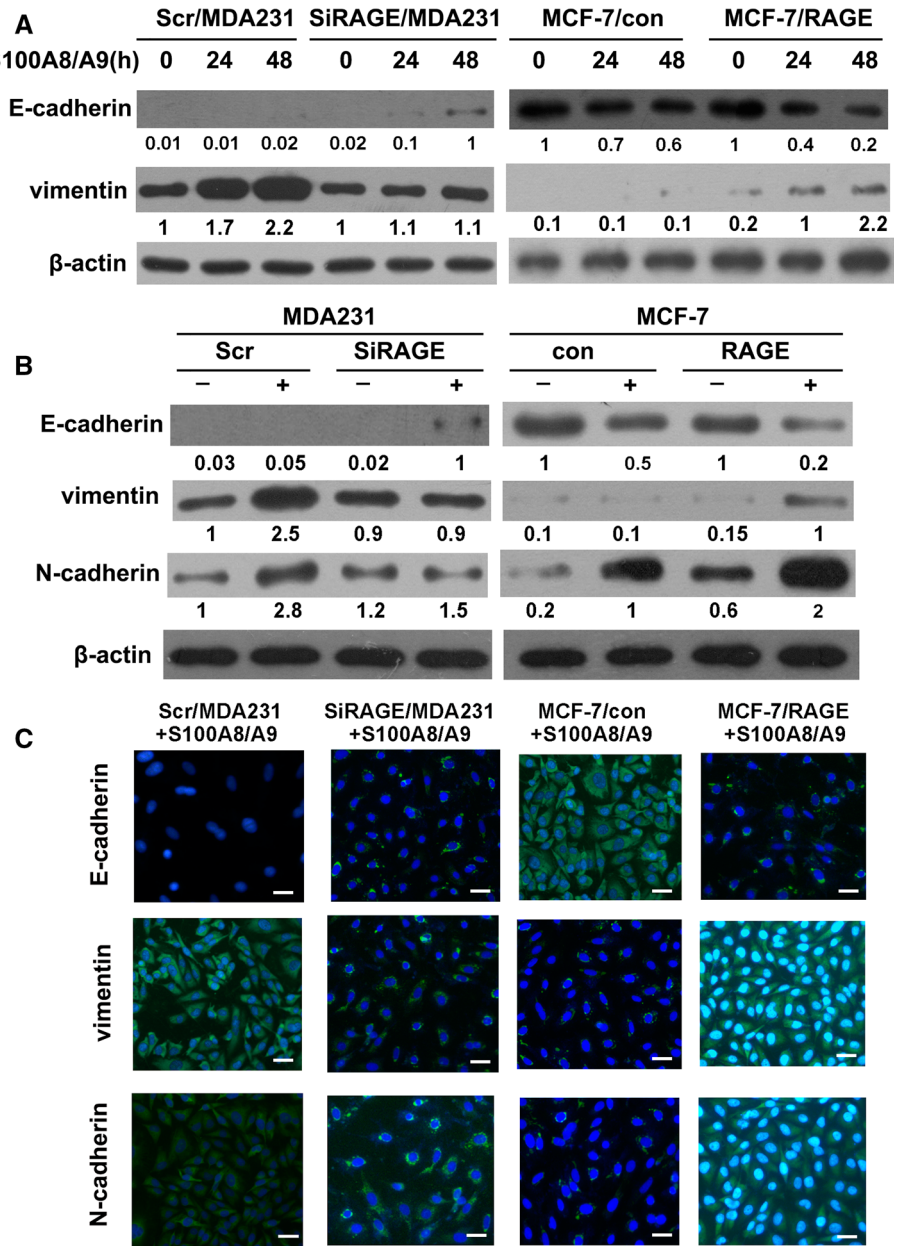

Fig. 3 RAGE binding to S100A8/A9-induced EMT through the NF-κB signaling pathway. **a** The expression of E-cadherin and vimentin in Scr/MDA231, SiRAGE/MDA231, MCF-7/con, and MCF-7/RAGE cells with 10 μg/mL rS100A8/A9 at different time points. β-actin was used as a loading control. Quantification of relative protein levels on three different western blots is shown below the *blots*. **b** Expression of epithelial markers, E-cadherin, as well as mesenchymal markers, N-cadherin and vimentin, was examined by western blot in Scr/MDA231, SiRAGE/MDA231, MCF-7/con, and MCF-7/RAGE cells with or without 10 μg/mL rS100A8/A9 for 48 h. β-actin was used as a loading control. Quantification of relative protein levels on three different western blots is shown below the *blots*. **c** Fluorescence microscopic staining of E-cadherin, N-cadherin, and vimentin (*green*) is indicated in the Scr/MDA231, SiRAGE/MDA231, MCF-7/Con, and MCF-7/RAGE cells with rS100A8/A9 for 48 h. Nuclear DNA was stained with DAPI (*blue*). *Scale bar* 20 μm. Data were collected in this set of figures from a representative of at least three independent experiments