

CORRECTION

Correction: A p53-independent role of Mdm2 in estrogen-mediated activation of breast cancer cell proliferation

Angelika Brekman, Kathryn E Singh, Alla Polotskaia, Nandini Kundu and Jill Bargonetti*

See related research by Brekman et al., http://breast-cancer-research.com/content/13/1/R3

Correction

Following publication of our article [1], the authors noticed that the data incorporated into Figure 4f was incorrect. The first column should be changed from 44, 13 and 46 to read 43, 13 and 44 instead. A corrected version of Figure 4f can be found overleaf.

Competing interests

The authors declare that they have no competing interests.

Published: 29 February 2012

Reference

1. Brekman A, Singh KE, Polotskaia A, Kundu N, Bargonetti J: A p53independent role of Mdm2 in estrogen-mediated activation of breast cancer cell proliferation. Breast Cancer Research 2011, 13:R3.

doi:10.1186/bcr3130

Cite this article as: Brekman A, et al.: Correction: A p53-independent role of Mdm2 in estrogen-mediated activation of breast cancer cell proliferation. Breast Cancer Research 2012, 14:302.

^{*} Correspondence: bargonetti@genectr.hunter.cuny.edu Department of Biological Sciences, Hunter College and The Graduate Center Biochemistry and Biology Programs, CUNY, 695 Park Ave, New York, NY 10065, USA



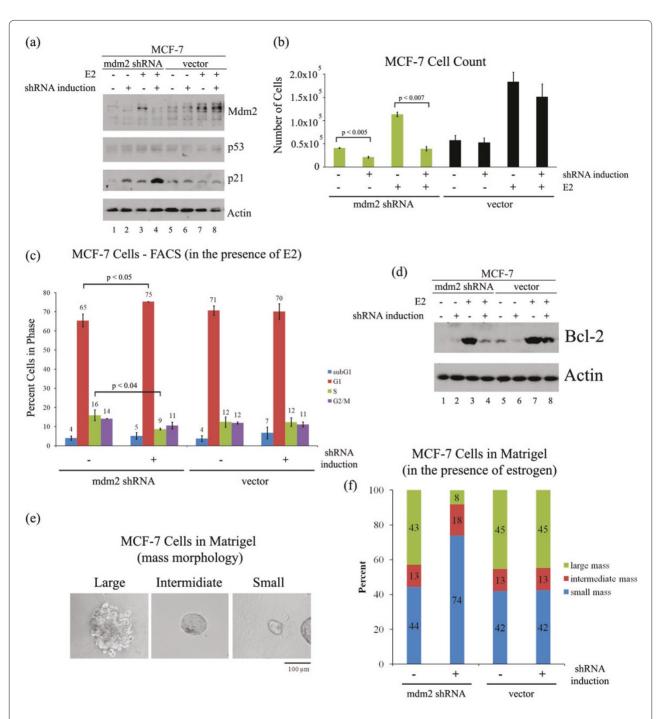


Figure 4. Mdm2 knockdown in estrogen-treated MCF-7 cells inhibits cell proliferation. Clonal MCF-7 cell lines with mdm2 shRNA or control vector were treated with 2 μ g/ml doxycycline (DOX) for three days to induce shRNA expression, followed by 10 nM estrogen (E2) for five days in the presence of DOX. (a) Western blot analysis of Mdm2, p53, p21 and Actin protein levels from whole cell lysates. (b) Number of cells was determined by Guava Viacount assay. 10,000 cells were seeded at beginning of treatments. (c) Fluorescence activated cell sorting (FACS). Cells were harvested, fixed in 30% ethanol, and cellular DNA was stained with propidium iodide. (d) Western blot analysis of Bcl-2 and Actin protein levels from whole cell lysates. (e) MCF-7 cells, grown in matrigel for three weeks, formed mass structures of three sizes: large, intermediate and small. (f) MCF-7 cells, grown in matrigel for three weeks in the presence of 10 nM estrogen and in the absence or presence of 2 μ g/ml doxycycline, were fixed and stained with propidium iodide. Masses of different sizes (large, intermediate and small) were counted and presented as percent of the total population. On average, a total of 300 masses were counted. Averages of three independent experiments are shown. The significance in the percent change of large and small masses was determined by the Student t-test (P < 0.05).