ENVIRONMENTAL SCIENCES


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The authors also note that an additional affiliation for Greg Deakin should be listed; the additional affiliation is Mushroom Research Group, Crops, Environment, and Land Use Programme, Teagasc, Ashtown, Dublin 15, Ireland. The corrected author and affiliation lines appear below. The online version has been corrected.


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The authors note the following: “We have recently become aware that errors were introduced into Figs. 1 and 2 during their composition. In Fig. 1, the control panels were reused in Fig. 1A and Fig. 1F, and the incorrect MTA1 panel was used in Fig. 1J. We have now replaced the panels with correct panels from the original experimental results. In Fig. 2, the ER panel in Fig. 2C was reused in Fig. 2H. We have now replaced the panel with the correct panel in Figure 2H. Additionally, the RT-PCR panels of Fig. 2E and 2F were originally spliced together. We have now replaced these RT-PCR panels with results from repeat experiments. These errors do not change the original scientific conclusions and validity of the result remains the same.” The corrected figures and their corresponding legends appear below.

**Fig. 1.** Recruitment to the regulatory region and modulation of BCAS3 by MTA1. (A) MTA1-associated chromatin from MCF-7 cells associates with regulatory region of BCAS3. (B) Levels of BCAS3 and MTA1 in MCF-7/MTA1 (T11 clone) and MCF-7/pcDNA cells. (C) Effect of MTA1 knockdown by RNAi on the levels of BCAS3. (D) Effect of T7-MTA1 or pcDNA on the BCAS3-luc activity in MCF-7 cells. Inset, expression of transfected T7-MTA1. (E) Status of BCAS3-luc activity in MCF-7/MTA1 and HC11/MTA1 clones with respective controls. (F) Recruitment of MTA1 onto BCAS3 regulatory region upon E2 stimulation. (G) BCAS3-luc reporter activity in MCF-7 cells transiently transfected with T7-MTA1 and treated with E2. (H and I) Effect of E2 treatment on BCAS3-luc activity and BCAS3 protein in MCF-7/pcDNA and MCF-7/MTA1 cells. Luciferase activity is represented as fold induction by E2 as compared to control (n = 3). (J) Effect of MTA1 depletion on BCAS3 expression in MCF-7 cells treated with or without E2 (n = 3).
**CELL BIOLOGY**


The authors note that, due to a printer’s error, the footnote “1L.Z., Y.M., and N.-y.F. contributed equally to this work” should instead appear as “1Y.M. and N.-y.F. contributed equally to this work.”

Additionally, within the author line, “Lei Zhang” should instead appear as “Lei Zhang.” The corrected author line appears below.

Lei Zhang, Yang Mei, Nai-yang Fu, Li Guan, Wei Xie, Hui-hui Liu, Chun-dong Yu, Zhenyu Yin, Victor C. Yu, and Han You.

**EDITORIAL**


The editors note that the author name Inder M. Verman should instead appear as Inder M. Verma. The corrected author line appears below. The online version has been corrected.

Inder M. Verma and David J. Harris.

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**Fig. 2.** Occupancy of BCAS3 enhancer module by ERα and regulation of gene expression. (A) E2 stimulation promotes ERα recruitment onto BCAS3 regulatory region in MCF-7 cells. (B) ERα occupancy of BCAS3 regulatory sequence upon E2 treatment. (C) Effect of ERα depletion upon E2 stimulation of BCAS3-luc activity in MCF-7 cells. (D) BCAS3-luc activity in HeLa cells transfected with ERα or vector and treated with E2. (E) Effect of E2 stimulation on BCAS3 mRNA in MCF-7 cells. (F) Inhibition of E2-induced BCAS3 expression by ICI-182780 in MCF-7 cells. (G) Effect of ERα knockdown on BCAS3 expression in MCF-7 cells treated with or without E2. (H) (Right) Recruitment of MTA1 onto BCAS3 regulatory region in response to E2 stimulation in ER-depleted MCF-7 cells. (Left) Status of ERα knockdown by RNAi (n = 3).

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With this issue, we relaunch the front matter of the journal in a magazine format, both in print and online. The new material will allow us to better explain science and to be part of the larger discussion that surrounds the core research published in the journal. We hope that the new content reaches an even broader audience than the scientific papers in the journal. Whether you are a scientist, a policy maker, or a fan of science, we will bring you stories that are engaging and accessible, requiring only an interest in science and an inquisitive mind to read.

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