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OPEN Author Correction: The lichen secondary metabolite atranorin suppresses lung cancer cell motility and tumorigenesis

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Correction to: Scientific Reports https://doi.org/10.1038/s41598-017-08225-1, published online 15 August 2017

The original version of this Article contained an error in Figure 2D, where the image for Atranorin 0 hr was inadvertently duplicated for the DMSO 0 hr panel.

The original Figure 2 appears below.

The original Article has been corrected.



Figure 2. Atranorin was identified as an active secondary metabolite from *E. vexans* with inhibitory activity against A549 cell motility. (a) TLC analysis performed using a Toluene: Dioxin: Acetic acid = 180: 45: 5 (v/v/v) solvent system showed that lichen extracts had inhibitory activity against A549 cell motility; 'a' denotes the location of the spot for atranorin. *L. cladonioides* was used as the standard control for atranorin; it contained atranorin (spot 'a') and norsticit acid (spot 'b'). (b) Chemical structure of atranorin. (c) MTT assay in A549 cells treated with atranorin and different doses. (d, e) Migration assays in A549 cells treated with 5 µg/mL atranorin, and quantitative analysis of wound length. (f, g) Invasion assays in A549 cells treated with 5 µg/mL atranorin and quantitative analysis of invaded cell numbers in each treatment. Quantitative data were obtained from three independent experiments (n=3). Data represent the mean \pm S.E.M. *p<0.05; **p<0.01; ***p<0.001 compared with DMSO-treated A549 cells.

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