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CORRECTION

Correction to: Evaluation of EpCAM-specific exosomal lncRNAs as potential diagnostic biomarkers for lung cancer using droplet digital PCR

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In Fig. 2c, the two pictures in the red boxes were pasted in the wrong place. Their positions should be swapped. Now, they were switched.

Xintong Shen, Yifeng Yang, and Yinfeng Chen contributed equally to this article.

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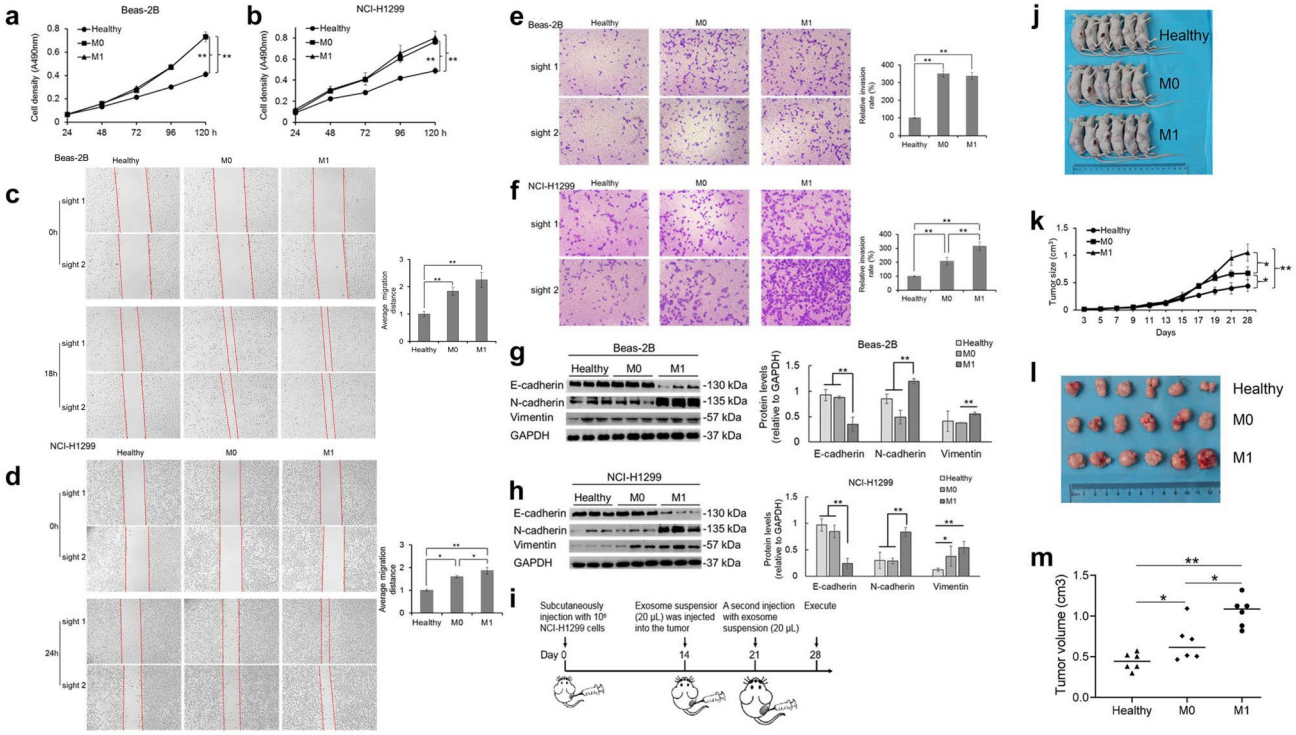

Fig. 2 The effect of EpCAM-specific exosomes on lung cancer. The total exosomes from healthy people; the EpCAM-specific exosomes from NSCLC patients without (M0)/with (M1) metastasis were isolated and resuspended in cell culture medium and further cocultured with Beas-2B and NCI-H1299 cells. The EpCAM-specific exosomes from M0 and M1 patients promote cell proliferation, cell migration, and invasion in Beas-2B (**a**, **c**, **e**) and NCI-H1299 cells (**b**, **d**, **f**). The western blot shows that the levels of E-cadherin are much lower, and the levels of N-cadherin and Vimentin proteins are higher in Beas-2B (**g**) and NCI-H1299 (**h**) cells cocultured with EpCAM-specific exosomes from M0 and M1 patients than those cocultured with total exosomes from healthy people. **i** NCI-H1299 cells were injected subcutaneously into the nude mouse to generate lung tumor, and then, total exosomes and EpCAM-specific exosomes were injected twice into lung tumors. (**j**, **k**, **l**, **m**) The tumor sizes were bigger in the mice injected with EpCAM-specific exosomes from M0 and M1 patients than those injected with total exosomes from healthy people

The original article has been corrected.