ERRATUM



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Erratum to: Ascorbic acid improves pluripotency of human parthenogenetic embryonic stem cells through modifying imprinted gene expression in the Dlk1-Dio3 region

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Erratum

Following the publication of our article [1], we noticed that some incorrect images had been incorporated into figure twoB (included here as Fig. 1b) and threeF-H (included here as Fig. 2f-h) in error. The corrected figures are given below. This correction does not change the results or conclusion of the original study.

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References

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for alkaline phosphatase; (c) normal 46, XX karyotype at passage 20; (d) positive staining for OCT4; (e) positive staining for NANOG; (f) positive staining for TRA-1-60; (D1-F1) nuclear staining with Hoechst 33342; (D2-F2) merged images for OCT4, NANOG and TRA-1-60. Bar is 100 μm



Fig. 2 Differentiation abilities of human parthenogenetic embryonic stem cells. *In vitro* differentiated EBs displayed (**a**) positive AFP staining (endoderm), (**b**) positive SMA staining (mesoderm), (**c**) positive TUBULIN staining (ectoderm), and (**d**) expression of genes from endoderm (NF68KD), mesoderm (HBZ) and ectoderm (Albumin). Bar is 50 μm. (**e**) Efficiency of teratoma formation upon injection of human parthenogenetic embryonic stem cells into SCID mice; (**f**) neuro-ectoderm from ectoderm in teratoma; (**g**) cartilage from mesoderm in teratoma; (**h**) glandular tissue from endoderm in teratoma. Bar is 100 μm. EB, embryoid bodies; SCID, severe combined immunodeficiency