

Administration of BMP2/7 in utero partially reverses Rubinstein-Taybi syndrome–like skeletal defects induced by *Pdk1* or *Cbp* mutations in mice

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Corrigendum

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Corrigendum

Administration of BMP2/7 in utero partially reverses Rubinstein-Taybi syndrome-like skeletal defects induced by *Pdk1* or *Cbp* mutations in mice

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The authors regret the errors.

Corrigendum

Mutations in 2 distinct genetic pathways result in cerebral cavernous malformations in mice

Aubrey C. Chan, Stavros G. Drakos, Oscar E. Ruiz, Alexandra C.H. Smith, Christopher C. Gibson, Jing Ling, Samuel F. Passi, Amber N. Stratman, Anastasia Sacharidou, M. Patricia Revelo, Allie H. Grossmann, Nikolaos A. Diakos, George E. Davis, Mark M. Metzstein, Kevin J. Whitehead, and Dean Y. Li

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The control data provided in Figure 7, E, F, and G, were inadvertently provided from incorrectly matched samples. The correct images are below.

The authors regret the error.

