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CORRECTION

Wnt controls the transcriptional activity of Kaiso through CK1ε-dependent phosphorylation of p120-catenin

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There was an error published in *J. Cell Sci.* **124**, 2298–2309.

Some of the blots presented in Fig. 4A were inadvertently duplicated in Fig. 6D. The blots in Fig. 4A are correct as presented. The correct Fig. 6 is presented below. There are no changes to the figure legend, which is accurate. This error does not affect the conclusions of the study.

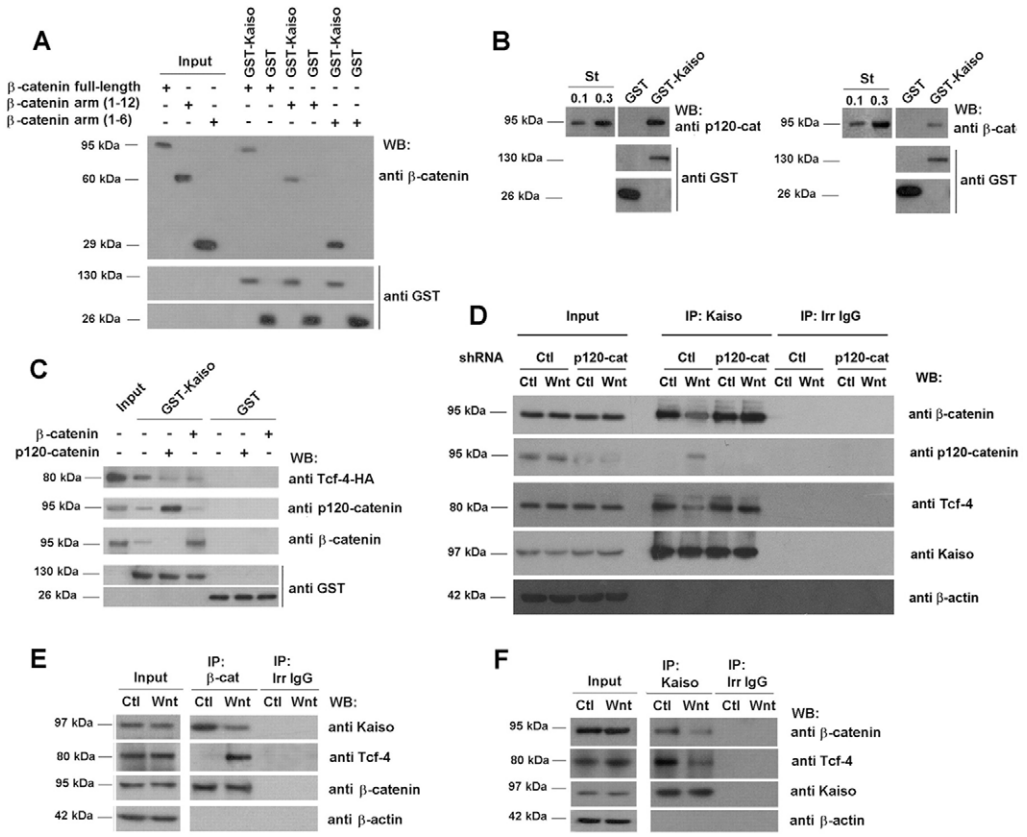

Fig. 6. Kaiso also interacts with β-catenin and its binding is regulated by p120-catenin. (A) 2 pmol of recombinant GST–Kaiso or GST as a control was incubated with 1 pmol of full-length β-catenin or two deletion mutants comprising armadillo repeats 1–12 (amino acids 120–683) or 1–6 (amino acids 120–420). Protein complexes were affinity purified with glutathione–Sepharose and analyzed by western blotting (WB) with anti-β-catenin antibody. Blots were re-analyzed with anti-GST antibody to ensure that a similar amount of fusion protein was present. (B) 2 pmol of recombinant GST–Kaiso, or GST as a control, was incubated with 2 pmol of p120-catenin (amino acids 102–911) (left-hand panel) or 2 pmol of full-length β-catenin (right-hand panel). Protein complexes were affinity purified with glutathione–Sepharose and analyzed by western blotting with the indicated antibodies. 'St' indicates the signal obtained with known amounts of recombinant proteins used as control. (C) Pull-down assays were performed by incubating 7 pmol of GST–Kaiso with 400 µg of whole-cell extracts from SW-480 cells transfected with pcDNA3-Tcf-4-HA. When indicated, 35 pmols of p120-catenin or β-catenin was added to the incubation medium. Protein complexes were affinity purified and analyzed by western blotting with the indicated antibodies. (D) SW-480 cells were infected with scrambled shRNA (Ctl) or shRNA specific for p120-catenin, and were treated with control or Wnt3a-conditioned medium for 6 hours. Kaiso was immunoprecipitated (IP) and immunocomplexes were analyzed by western blotting. (E) SW480 cells were treated with control or Wnt3a-conditioned medium for 6 hours. β-catenin was immunoprecipitated from SW-480 total cell extracts and the immunocomplex was analyzed by western blotting. In the input lane, 5% of each total cell extract used is shown. (F) Kaiso was immunoprecipitated with a specific antibody from total cell extracts of HT29-M6 cells treated with control or Wnt3a-conditioned medium for 6 hours. Associated proteins were analyzed by western blotting. All the data presented in this figure are representative of at least three independent experiments. Irr IgG, an irrelevant IgG used as control in the immunoprecipitation.

The authors apologise to the readers for any confusion that this error might have caused.

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