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## Corrigendum

# Bcl-xL controls a switch between cell death modes during mitotic arrest

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Since the publication of this article the authors have noticed an error in Figure 7c. pS62DBcl-xL and pS62ABcl-xL are in the wrong order. The corrected figure is shown here.

The corrected article appears online together with this corrigendum. The authors would like to apologize for any inconvenience caused.

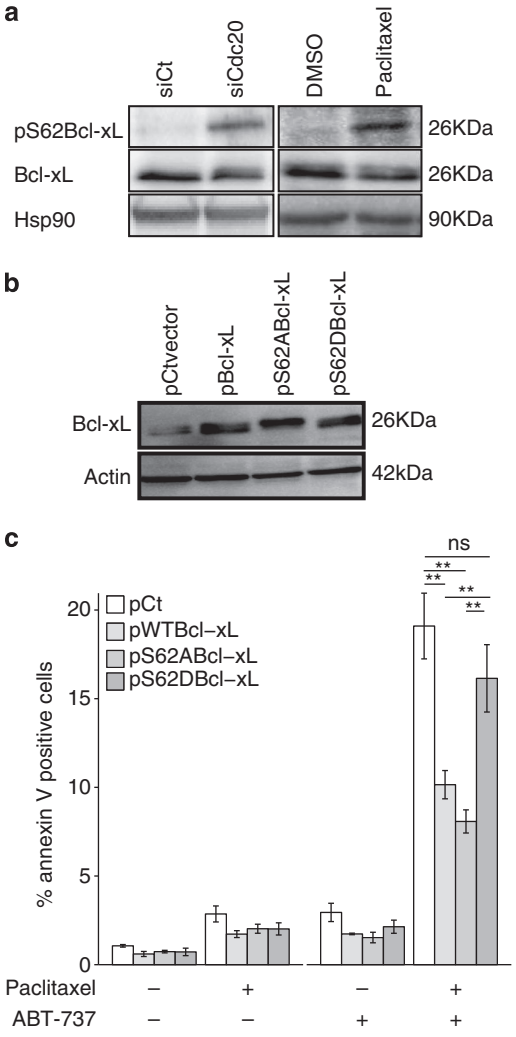


**Figure 7** Serine 62 phosphorylation of Bcl-xL occurring during prolonged mitotic arrest, promoted mitotic cell apoptosis. (**a**) Immunoblot analysis of S62 phosphorylation of Bcl-xL (pS62Bcl-xL) expression levels after Cdc20 depletion or paclitaxel treatment. MDA-MB-231 cells were transfected with either control or Cdc20 siRNA for 48 h or treated with paclitaxel for 12 h, harvested and the expression level of pS62Bcl-xL were finally evaluated by immunoblot analysis with S62-phospho-specific or total Bcl-xL antibodies. (**b**) Immunoblot analysis of Bcl-xL mutants. MDA-MB-231 cells were transfected with plasmids coding for wt Bcl-xL or S62(A or D)Bcl-xL mutants and analysed by immunoblotting 48 h later. (**c**) Apoptosis analysis of MDA-MB-231 cells after Bcl-xL or its S62 mutants (S62A- or S62D-Bcl-xL) overexpression, and treatment or not with paclitaxel and ABT-737, compared with control cells. MDA-MB-231 cells were transfected, respectively, with control plasmid (pCt), plasmids coding for Bcl-xL (pBcl-xL) or its S62(A or D) mutants (pS62(A or D)Bcl-xL). Cells were then treated with either DMSO or paclitaxel (70 nM), and finally stained with Annexin-V, and analysed by flow cytometry