

SOX2OT Long Noncoding RNA Is Regulated by the UPR in Oestrogen Receptor-Positive Breast Cancer

Carole Ferraro-Peyret ^{1,2,3,*}, Marjan E. Askarian-Amiri ^{1,4}, Debina Sarkar ^{1,4,5}, Wayne R. Joseph ¹, Herah Hansji ^{1,4,6}, Bruce C. Baguley ¹ and Euphemia Y. Leung ^{1,4,*}

¹ Auckland Cancer Society Research Centre, University of Auckland, 85 Park Rd, Grafton, Auckland 1023, New Zealand;

² Univ Lyon, ISPB Faculty of Pharmacy, INSERM 1052, CNRS5286, Cancer Research Centre of Lyon, F-69008 Lyon, France

³ Hospices Civils de Lyon, Biopathology of Tumours, GHE Hospital, F-69500 Bron, France

⁴ Molecular Medicine and Pathology Department, University of Auckland, 85 Park Rd, Grafton, Auckland 1023, New Zealand

⁵ Biochemistry Department, University of Otago, 710 Cumberland Street, Dunedin North, Dunedin 9016, New Zealand

⁶ Murdoch Children's Research Institute, The Royal Children's Hospital, Flemington Rd, Parkville VIC 3052, Australia

* Correspondence: carole.ferraro-peyret@univ-lyon1.fr (C.F.-P.); e.leung@auckland.ac.nz (E.Y.L.)

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Table 1. List of primers used in these experiments.

ATF4-F	5'- GTTCTCCAGCGACAAGGCTA-3'
ATF4-R	5'- ATCCTGCTTGCTGTTGTTGG-3'
BiP-F	5'- TGTTCACCAATTATCAGCAAACTC-3'
BiP-R	5'- TTCTGCTGTATCCTCTTCACCACT-3'
CHOP-F	5'- AGAACCAGGAAACGGAAACAGA-3'
CHOP-R	5'- TCTCCTTCATGCGCTGCTTT-3'
SOX2OT-F	5'- GCTCGTGGCTTAGGAGATTG -3'
SOX2OT-R	5'- CTGGCAAAGCATGAGGAACT -3'
SOX2-F	5'-AACCCCAGATGCACAA CTC-3'
SOX2-R	5'-GCTTAGCCTCGTCGATGAAC-3'
XBP1s-F	5'- CCTGGTTGCTGAAGAGGAGG-3'
XBP1s-R	5'- CCATGGGGAGATGTTCTGGAG -3'
HPRT-F	5'-TGAGGATTGGAAAGGGTGT-3'
HPRT-R	5'-GCACACAGAGGGCTACAATG-3'
GAPDH-F	5'-ACGGG AAGCTTGTCAAT-3'
GAPDH-R	5'-TGGACTCCACGACGTACTCA-3'
HPRT-F	5'-TGAGGATTGGAAAGGGTGT-3'
HPRT-R	5'-GCACACAGAGGGCTACAATG-3'

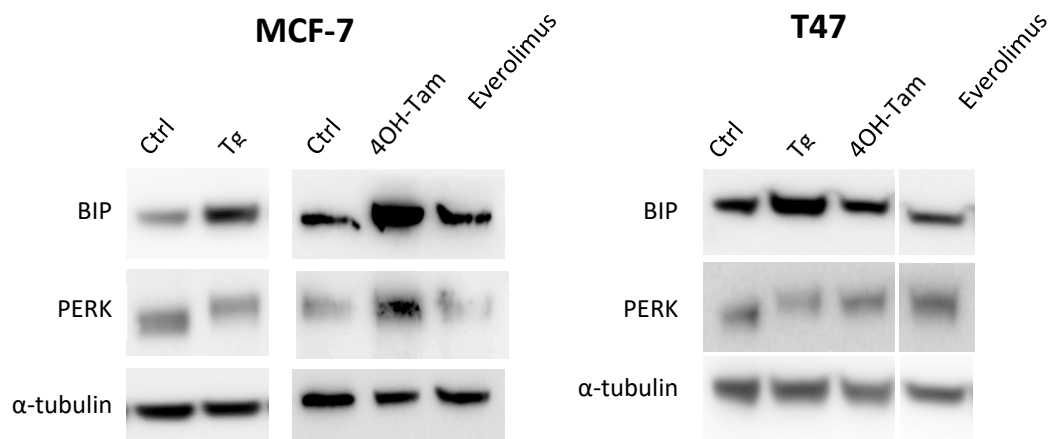


Figure S1. Expression of *SOX2OT* lncRNA and *SOX2* following UPR induction in ER+ breast cancer cell lines. MCF-7 (A) and T47D (B) cells were incubated in control medium (Ctrl) or in medium containing the ER stress-inducing agents thapsigargin (Tg, 300 nM), 4OH-Tam (0.1uM) or everolimus (10nM) for 16h. Protein expression levels of PERK and BIP protein expression were examined using Western Blot analysis. α -tubulin was used as internal control. Blots of Bip, PERK, and α -tubulin have been performed on the same electrophoresis gel, but cut and reconstituted.

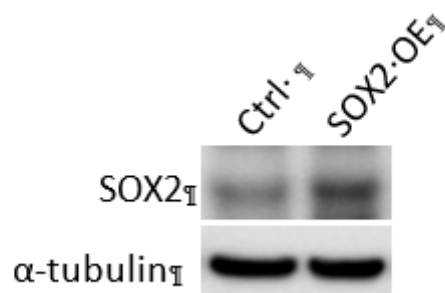


Figure S2. Comparison of SOX2 expression in control and SOX2 overexpressing MDA-MB-231. SOX2 protein expression was examined using Western Blot analysis. α -tubulin was used as internal control. Ctrl, control; SOX2 OE, SOX2 overexpression.



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