



Supplementary materials

IBtk α Activates the β -Catenin-Dependent Transcription of MYC through Ubiquitylation and Proteasomal Degradation of GSK3 β in Cancerous B Cells

Eleonora Vecchio ^{1,*}, Nancy Nisticò ¹, Gaetanina Golino ¹, Enrico Iaccino ¹, Domenico Maisano ¹, Selena Mimmi ¹, Annamaria Aloisio ¹, Maurizio Renna ², Angelica Avagliano ³, Alessandro Arcucci ³, Giuseppe Fiume ^{1,*},[†] and Ileana Quinto ^{1,†}

¹ Department of Experimental and Clinical Medicine, University of Catanzaro 'Magna Graecia', 88100 Catanzaro, Italy; nancynistico@unicz.it (N.N.); tania.golino@gmail.com (G.G.); iaccino@unicz.it (E.I.); maisano@unicz.it (D.M.); mimmi@unicz.it (S.M.); aloisio@unicz.it (A.A.); quinto@unicz.it (I.Q.)

² Department of Molecular Medicine and Medical Biotechnology, University of Naples Federico II, 80131 Naples, Italy; maurizio.renna@unina.it (M.R.)

³ Department of Public Health, University of Naples Federico II, 80131 Naples, Italy; angelica.avagliano@unina.it (A.A.); alessandro.arcucci2@unina.it (A.A.)

* Correspondence: eleonoravecchio@unicz.it (E.V.); fiume@unicz.it (G.F.)

† G.F. and I.Q. should be considered as co-last authors.

Table S1. Mass spectrometry-based identification of GSK3 β from the IBtk α co-immunoprecipitation experiment. HEK293T cells were transfected with pCMV6-IBtk α -FLAG plasmid or the corresponding empty vector (24 μ g of DNA/100-mm dish). Protein extracts (1,5 mg) were immunoprecipitated with anti-FLAG antibody (20 μ g), and the immunocomplexes were resolved by NuPAGE 4–12% SDS-PAGE. Gels were stained with colloidal Coomassie, and protein bands were excised for in-gel tryptic digestion. Chromatography of tryptic peptides and mass spectrometry were performed as previously described [8].

Accession no.	Description	Coverage	Unique Peptides
Q9P2D0	Inhibitor of Bruton tyrosine kinase	60.16	110
P49841	Glycogen synthase kinase-3 beta	7.88	2