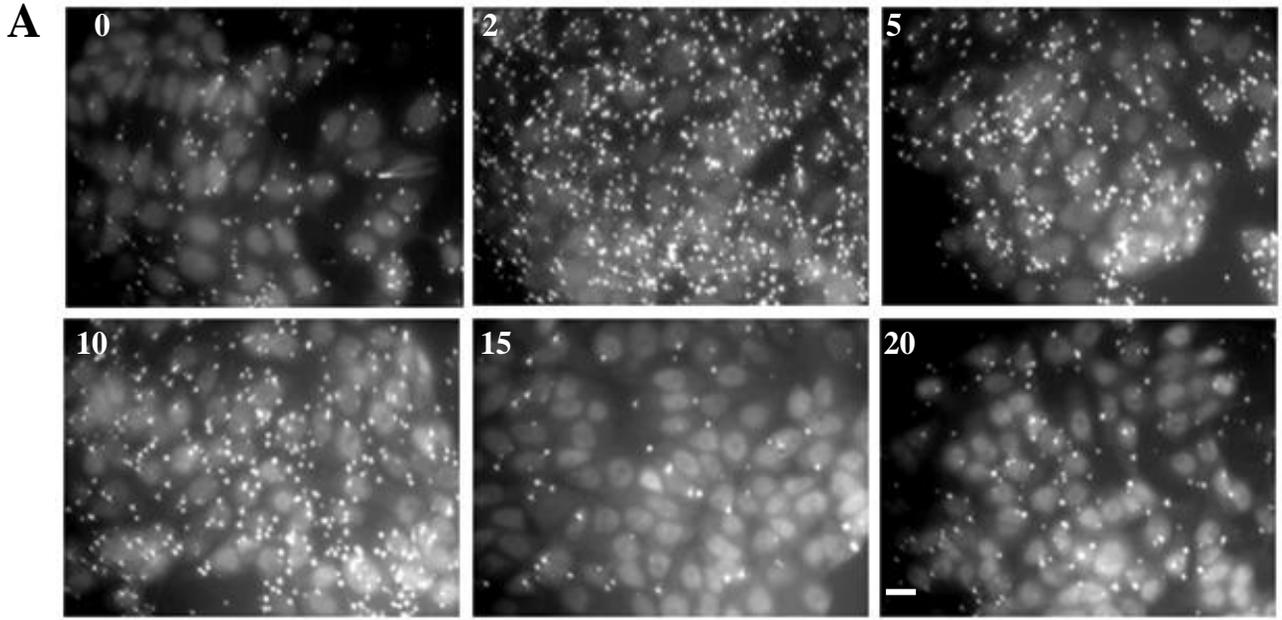
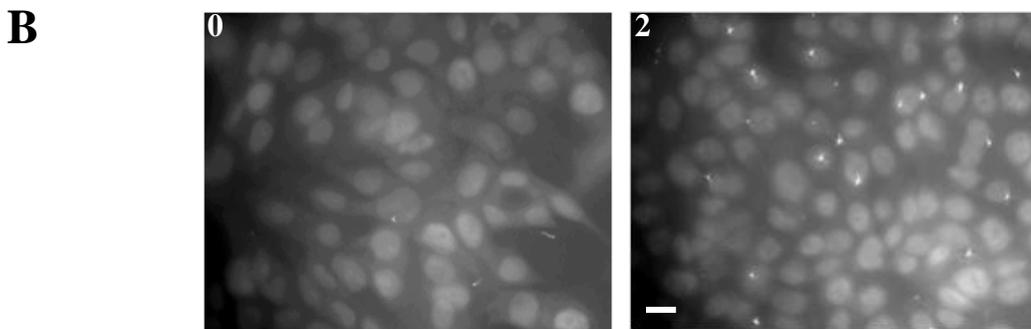


Supplementary Figures

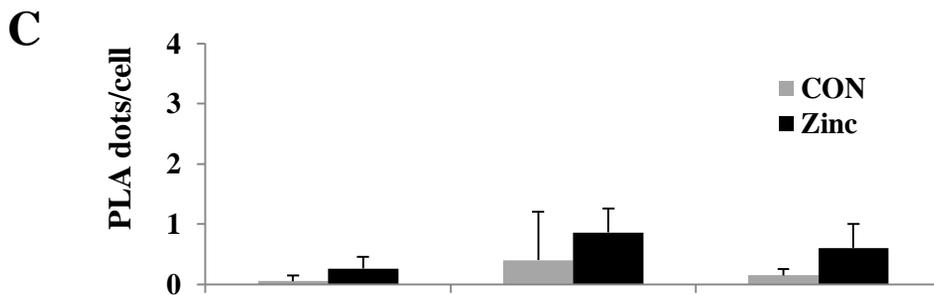
Figure S1



Treatment time (min)

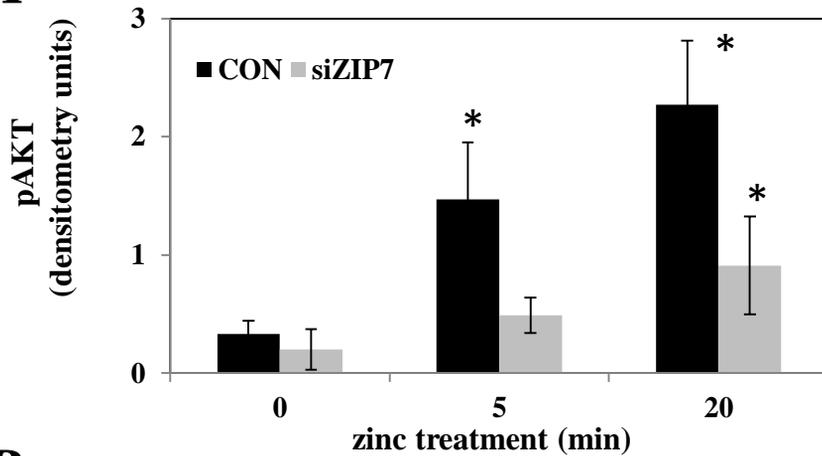
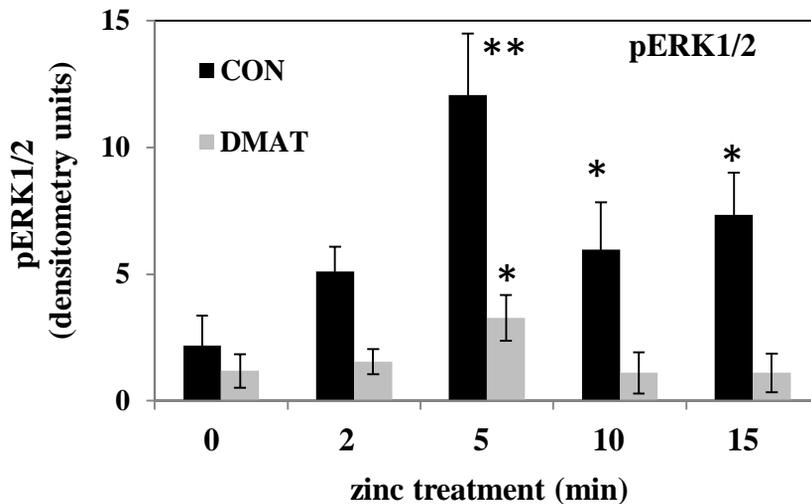
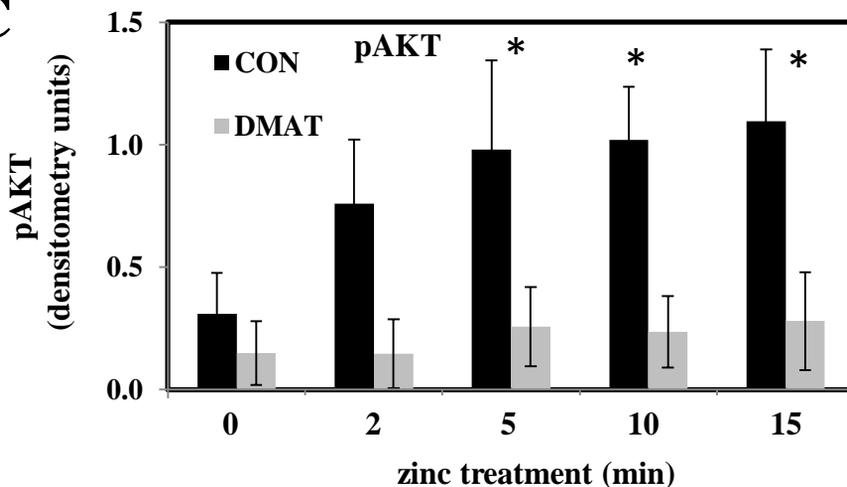


zinc treatment time (min)



Duolink Reagents	+	+	+
ZIP7 antibody	-	+	-
CK2 antibody	-	-	+

Proximity Ligation assay images and control results. (A) Representative figures of proximity ligation assay of ZIP7 and CK2 α stimulated with zinc in Fig 1E Scale bar = 10 μ m. (B) Representative figures of no antibody controls for proximity ligation assay in Figs 1E and 4B. Scale bar = 10 μ m. (C) Control results for proximity ligation assay lacking single and both antibodies and representative of n=3 experiments and expressed as mean \pm SD

Figure S2**A****B****C**

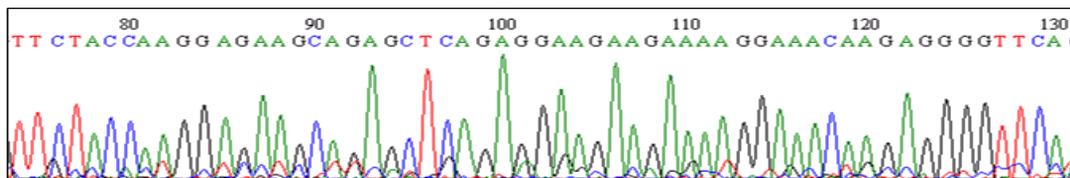
Densitometry analysis of pooled Western Blot results for Fig. 2. (A) Densitometry analysis of pooled Western Blot results from Fig 2D from n=3 experiments and expressed as mean \pm SD. * denotes significance from time zero ($P \leq 0.05$). (B) Densitometry analysis of pooled Western Blot results from Fig 2E from n=3 experiments and expressed as mean \pm SD. * denotes significance from time zero ($P \leq 0.05$) and ** denotes $P \leq 0.001$. (C) Densitometry analysis of pooled Western Blot results from Fig 2E from n=3 experiments and expressed as mean \pm SD. * denotes significance from time zero ($P \leq 0.05$).

Figure S3

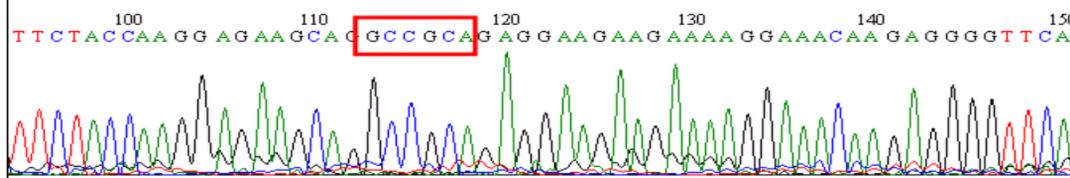
A

AGCTCA (Ser Ser) changed to GCCGCA (Ala Ala)

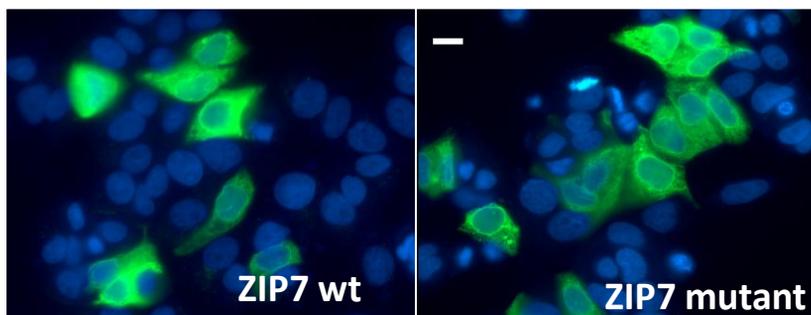
ZIP7
wild-type



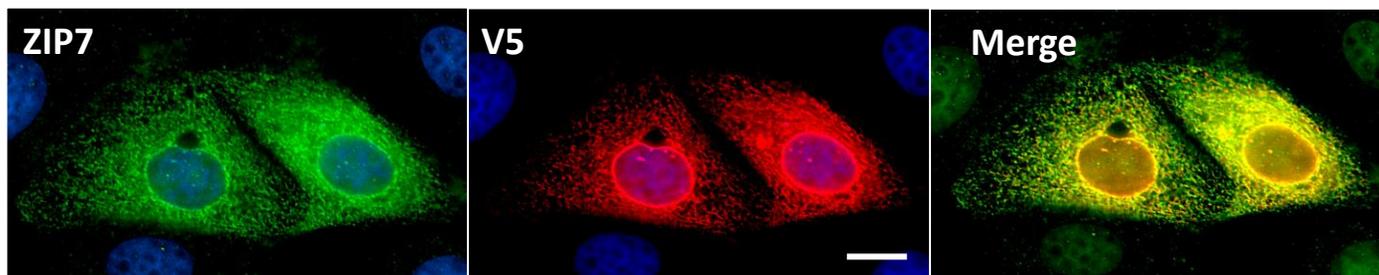
ZIP7
S275A
S276A



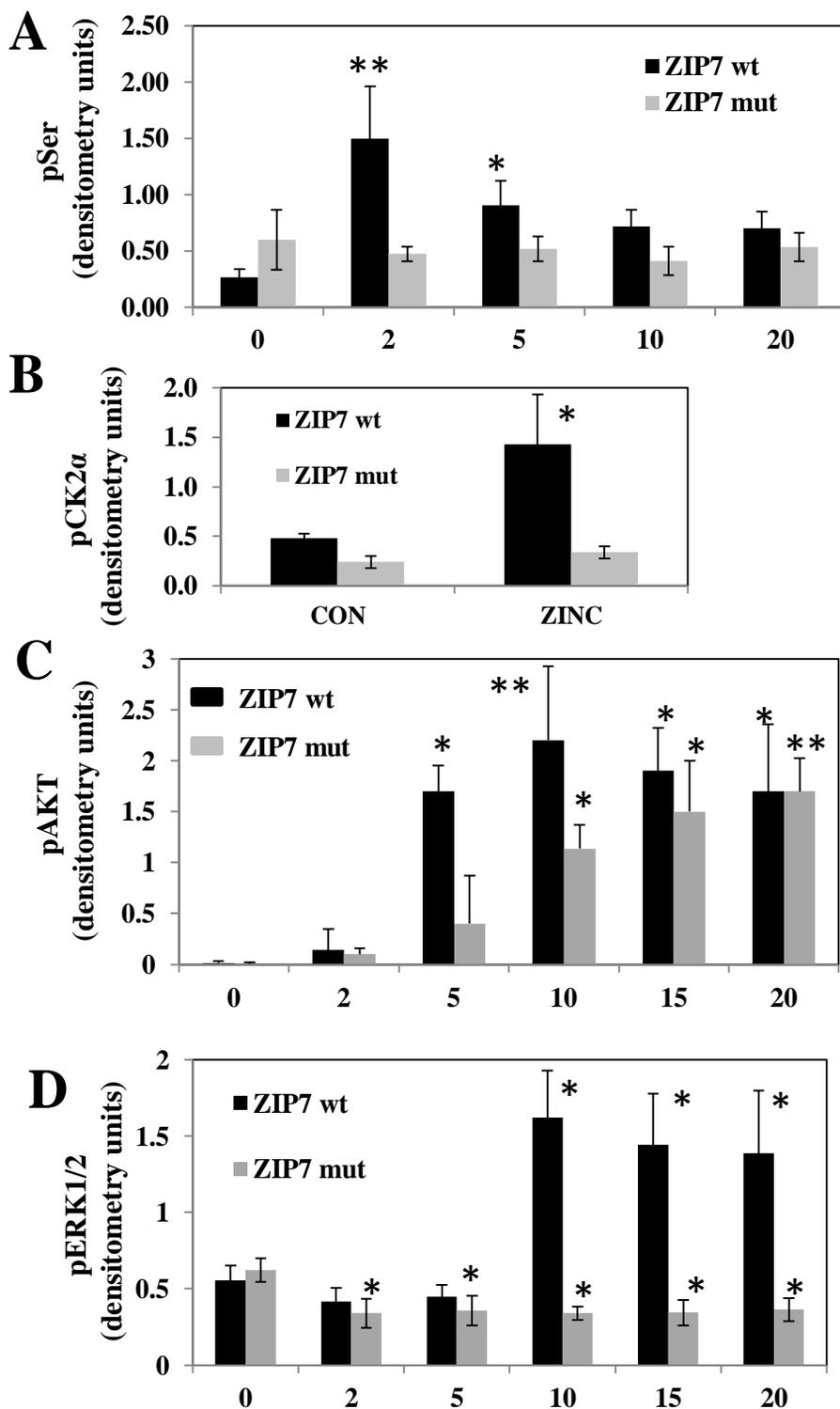
B



C



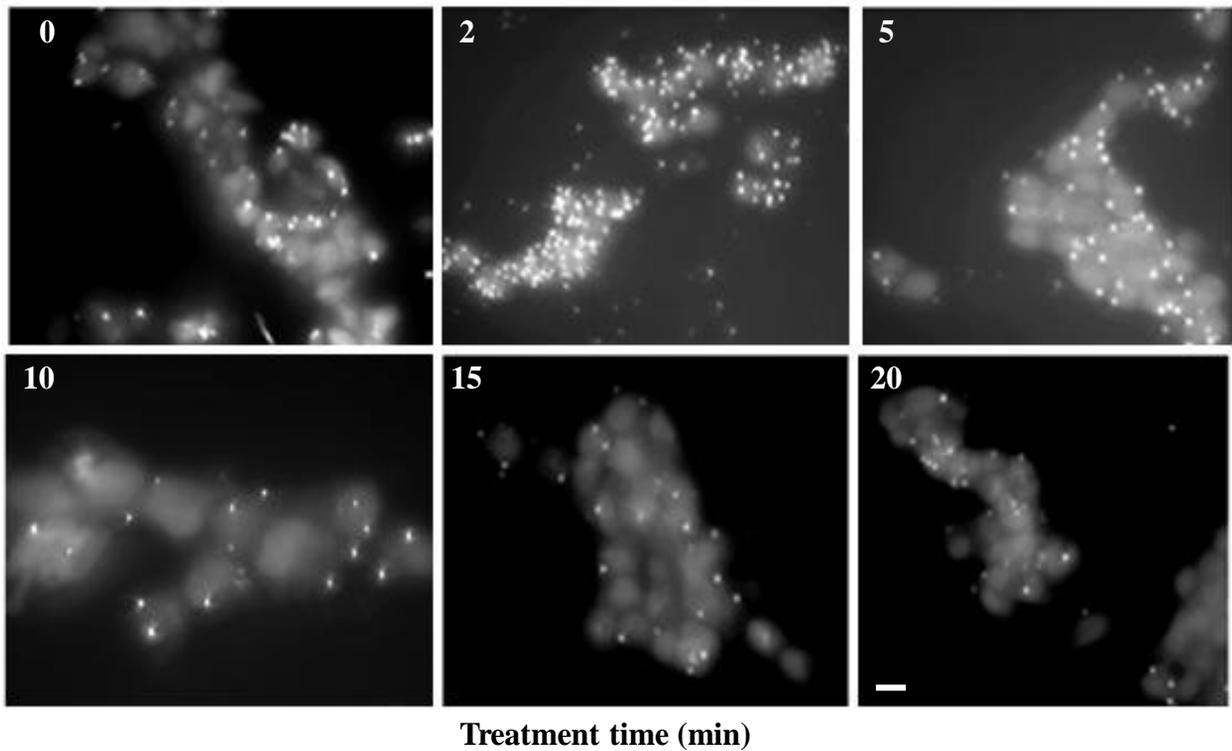
Characterisation of ZIP7 mutants and ZIP7 antibody. (A) Sequencing demonstration of S275 and S276 mutated to A275 and A276 in ZIP7 sequence. (B) MCF-7 cells transfected with either wild-type or mutant ZIP7 and probed with V5 antibody conjugated to Alexa-Fluor 488 (green). Nuclei stained blue with DAPI. Scale bar = 10µm. (C) MCF-7 cells transfected with wild-type ZIP7 and probed with ZIP7 antibody conjugated to Alexa-Fluor 488 (green), V5 antibody conjugated to Alexa-Fluor 594 (red) and nuclei stained blue with DAPI. Scale bar = 10µm.

Figure S4

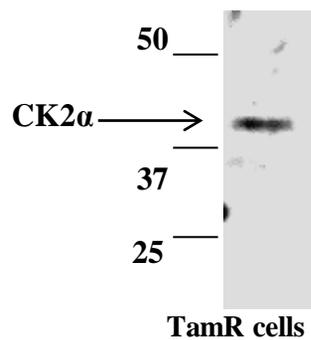
Densitometry analysis of pooled Western Blot results for Fig.3. (A) Densitometry analysis of pooled Western Blot results from Fig 3A from n=3 experiments and expressed as mean \pm SD. * denotes significance from time zero ($P \leq 0.05$) and ** denotes ($P \leq 0.001$). (B) Densitometry analysis of pooled Western Blot results from Fig 3C from n=3 experiments and expressed as mean \pm SD. * denotes significance from time zero ($P \leq 0.05$). (C) Densitometry analysis of pooled Western Blot results from Fig 3E from n=3 experiments and expressed as mean \pm SD. * denotes significance from time zero ($P \leq 0.05$) and ** denotes ($P \leq 0.001$). (D) Densitometry analysis of pooled Western Blot results from Fig 3E from n=3 experiments and expressed as mean \pm SD. * denotes significance from time zero ($P \leq 0.05$).

Figure S5

A



B



Proximity ligation assay images and CK2α antibody control. (A) Representative figures of proximity ligation assay of ZIP7 and CK2α stimulated with EGF and ionomycin in Fig 4B Scale bar = 10μm. (B) CK2α antibody recognises a single band on Western Blot of appropriate size in TamR cells