

Supplemental Information

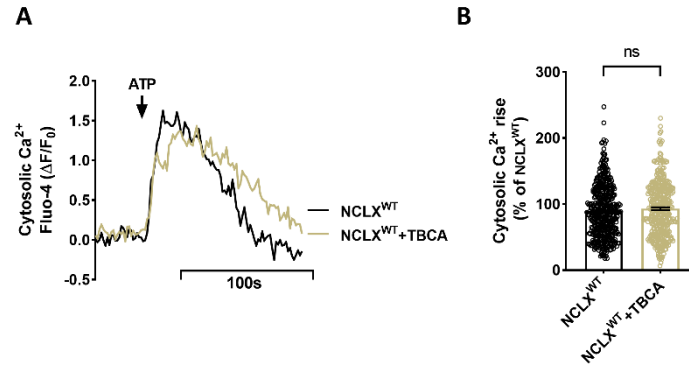


Figure S1. TBCA effect on Cytosolic Ca^{2+}

(A) Representative fluorescent traces of SH-SY5Y Cells co-expressing shNCLX and NCLX^{WT} (black), treated with TBCA (10 μM) for 2 hours (gold) and preloaded with Fluo-4 (2 μM). Cytosolic Ca^{2+} transient was triggered by application of extracellular ATP (100 μM) via Ca^{2+} -free Ringer's solution.

(D) Quantification of cytosolic Ca^{2+} influx amplitude in (A) (unpaired two-tailed t-test, $t=1.358$, $\text{df}=864$: ns; $n = \text{NCLX}^{\text{WT}}$ (452), NCLX^{WT}+TBCA (414)).

ns- Not significant. Error bars denote SEM.

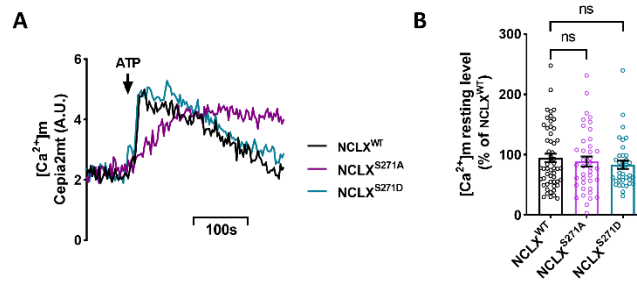


Figure S2

(A) Representative fluorescent (non-normalized) traces of ATP (100 μM) induced $[\text{Ca}^{2+}]_{\text{m}}$ transients in Cepia2mt expressing SH-SY5Y cells co-expressing shNCLX and NCLX^{WT} (black) or mutant NCLX^{S271A} (purple) \ NCLX^{S271D} (turquoise). A.U.- arbitrary units

(B) Quantification of $[\text{Ca}^{2+}]_{\text{m}}$ basal intensity in (A) (Welch and Brown-Forsythe ANOVA, $F(2,101.4)=0.8545$; Dunnett's T3 multiple comparisons test; NCLX^{WT} vs. NCLX^{S271A}: ns; NCLX^{WT} vs. NCLX^{S271D}: ns; $n = \text{NCLX}^{\text{WT}}$ (54), NCLX^{S271A} (40), NCLX^{S271D} (37)).

ns- Not significant. Error bars denote SEM.