**Supplementary File 4**. Comparison of Ca2+-activated Cl--tail currents measured from wild type and *Ribeye-*korods filled with 0.5 mM EGTA.

|  |  |  |  |
| --- | --- | --- | --- |
| genotype | ICl(Ca) @ −10 mV (pA) | Modified Boltzmann-IV fits | Boltzmann fits |
| V1/2(mV) | dx (mV-e−1) | Vrev(mV) | Gmax (pA-mV−1) | span(pA) | V0.5 (mV) | dx(mV-e−1) |
| wt 0.5 EGTAn: 6 | −68 ± 13 | −20.4 ± 0.5 | −5.55 ± 0.95 | 35.8 ± 3.8 | 1.41 ± 0.44 | 67 ± 14 | −23.9 ± 0.8 | 3.20 ± 0.19 |
| ko 0.5 EGTAn: 5 | −27 ± 5**p: 0.036** | −22.8 ± 0.6**p: 0.016** | −5.28 ± 0.25p: 0.79 | 34.1 ± 3.0p: 0.73 | 0.70 ± 0.13*p: 0.17* |  27 ± 5**p: 0.042** | −27.4 ± 0.8**p: 0.013** | 3.52 ± 0.24p: 0.33 |

Notes: Ca2+-activated Cl--tail currents were generated with the same voltage step protocol used to measure peak-ICa amplitude (see the legend to Supplementary File 3), and Boltzmann fits were made as described in Materials and methods. Tail-current amplitudes were measured 3 ms after repolarizing the cell to −70 mV. Liquid junction potentials were *not* subtracted from the voltage values presented above (i.e., V1/2, Vrev and V0.5); see Supplementary File 2 for details.