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Reporting Summary

Nature Portfolio wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Portfolio policies, see our <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

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FOr	all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Confirmed
	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
	The statistical test(s) used AND whether they are one- or two-sided Only common tests should be described solely by name; describe more complex techniques in the Methods section.
	A description of all covariates tested
X	A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
	A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
	For null hypothesis testing, the test statistic (e.g. <i>F</i> , <i>t</i> , <i>r</i>) with confidence intervals, effect sizes, degrees of freedom and <i>P</i> value noted <i>Give P values as exact values whenever suitable.</i>
X	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
X	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
\boxtimes	Estimates of effect sizes (e.g. Cohen's <i>d</i> , Pearson's <i>r</i>), indicating how they were calculated
	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.
So	ftware and code
Poli	cy information about <u>availability of computer code</u>

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio guidelines for submitting code & software for further information.

Clampfit V10.1, Prizm V9, MS Excel, Coot V0.9.7, PHENIX V1.19.2-4158, UCSF ChimeraX V1.2, UCSF Chimera V1.15, Pymol V2.3.2, CrYOLO

Data

Data collection

Data analysis

Policy information about availability of data

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A description of any restrictions on data availability

Clampex 10.1

- For clinical datasets or third party data, please ensure that the statement adheres to our policy

V1.7.6, Hole V2.2.005, CTFFIND V4.0, cryoSPARC V3.3.1

Structural data that support the findings of this study have been deposited in the PDB with the deposition numbers D_1000261989, PDB-ID: 7TCI [https://doi.org/10.2210/pdb7TCI/pdb] (xKCNQ1-CaM) and D_1000262040 PDB-ID: 7TCP [https://doi.org/10.2210/pdb7TCP/pdb] (xKCNQ1-Cam In complex with ML277). Other previously published PDB files used in this manuscript were PDB-ID: 6v01 [http://doi.org/10.2210/pdb6V01/pdb] (KCNQ1-KCNE3-CaM complex with PIP2); PDB-ID: 5VMS, [http://doi.org/10.2210/pdb5VMS/pdb] (CryoEM structure of Xenopus KCNQ1 channel); PDB-ID: 7c73, [http://doi.org/10.2210/pdb7CR3/pdb] (human KCNQ2-CaM in apo state); PDB-ID: 7byl, [http://doi.org/10.2210/pdb7BYL/pdb] (Cryo-EM structure of human KCNQ4); PDB-ID: 6UZZ, [http://doi.org/10.2210/pdb6UZZ/pdb] (human KCNQ1-CaM complex); PDB-ID: 7CR0, [http://doi.org/10.2210/pdb7CR0/pdb] (human KCNQ2 in apo state); PDB-ID: 7VNQ,

[http://doi.org/10.2210/pdb7VNQ/pdb] (human KCNQ4-ML213 complex in nanodisc). The summary electrophysiological data that were used to generate the figures presented in the paper are attached as supplemental tables. All other data supporting the findings of this study are available within the paper, its Supplementary information files as well as a Source data file.							
Field-spe	ecific reporting						
Please select the o	ne below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.						
∠ Life sciences	☐ Life sciences ☐ Behavioural & social sciences ☐ Ecological, evolutionary & environmental sciences						
For a reference copy of the document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>							
Life scier	nces study design						
All studies must dis	sclose on these points even when the disclosure is negative.						
Sample size	We used sample sizes of 3 or more, to allow statistical comparison. No sample size calculation was performed as standard procedure in cellular electrophysiology experiments.						
Data exclusions	If electrophysiological data were too small or cells were below standard (leaky), or data were excessively noisy, they were omitted						
Replication	Data were replicated at least 3 times						

Reporting for specific materials, systems and methods

individuals. Randomization of allocation is not relevant to the present study

yes, blinded during experiments

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Control cells were studied along with test samples. The experimental studies and subsequent analysis were carried out by separate

Materials & experimental systems		Methods	
n/a	Involved in the study	n/a	Involved in the study
\boxtimes	Antibodies	\boxtimes	ChIP-seq
	Eukaryotic cell lines	\boxtimes	Flow cytometry
\boxtimes	Palaeontology and archaeology	\boxtimes	MRI-based neuroimaging
\times	Animals and other organisms		
\boxtimes	Human research participants		
\boxtimes	Clinical data		
\boxtimes	Dual use research of concern		

Eukaryotic cell lines

Randomization

Blinding

Policy information about cell lines	
Cell line source(s)	All cell lines were from ATCC: tsA201, and ltk- mouse fibroblasts.
Authentication	Authentication by ATCC, these were low passage number cells. No further authentification performed
Mycoplasma contamination	cell lines not tested recently
Commonly misidentified lines (See ICLAC register)	none used