

## Washington University School of Medicine Digital Commons@Becker

---

### Open Access Publications

---

2013

# Secondary anionic phospholipid binding site and gating mechanism in Kir2.1 inward rectifier channels

Sun-Joo Lee

*Washington University School of Medicine in St. Louis*

Shizhen Wang

*Washington University School of Medicine in St. Louis*

William Borschel

*Washington University School of Medicine in St. Louis*

Sarah Heyman

*Washington University School of Medicine in St. Louis*

Jacob Gyore

*Washington University School of Medicine in St. Louis*

*See next page for additional authors*

Follow this and additional works at: [http://digitalcommons.wustl.edu/open\\_access\\_pubs](http://digitalcommons.wustl.edu/open_access_pubs)

---

### Recommended Citation

Lee, Sun-Joo; Wang, Shizhen; Borschel, William; Heyman, Sarah; Gyore, Jacob; and Nichols, Colin G., "Secondary anionic phospholipid binding site and gating mechanism in Kir2.1 inward rectifier channels." *Nature Communications*.4. 2786. (2013). [http://digitalcommons.wustl.edu/open\\_access\\_pubs/4317](http://digitalcommons.wustl.edu/open_access_pubs/4317)

This Open Access Publication is brought to you for free and open access by Digital Commons@Becker. It has been accepted for inclusion in Open Access Publications by an authorized administrator of Digital Commons@Becker. For more information, please contact [engeszer@wustl.edu](mailto:engeszer@wustl.edu).

---

**Authors**

Sun-Joo Lee, Shizhen Wang, William Borschel, Sarah Heyman, Jacob Gyore, and Colin G. Nichols

ARTICLE

Received 11 Jun 2013 | Accepted 16 Oct 2013 | Published 25 Nov 2013

DOI: 10.1038/ncomms3786

OPEN

# Secondary anionic phospholipid binding site and gating mechanism in Kir2.1 inward rectifier channels

Sun-Joo Lee<sup>1</sup>, Shizhen Wang<sup>1</sup>, William Borschel<sup>1</sup>, Sarah Heyman<sup>1</sup>, Jacob Gyore<sup>1</sup> & Colin G. Nichols<sup>1</sup>

Inwardly rectifying potassium (Kir) channels regulate multiple tissues. All Kir channels require interaction of phosphatidyl-4,5-bisphosphate (PIP<sub>2</sub>) at a crystallographically identified binding site, but an additional nonspecific secondary anionic phospholipid (PL(−)) is required to generate high PIP<sub>2</sub> sensitivity of Kir2 channel gating. The PL(−)-binding site and mechanism are yet to be elucidated. Here we report docking simulations that identify a putative PL(−)-binding site, adjacent to the PIP<sub>2</sub>-binding site, generated by two lysine residues from neighbouring subunits. When either lysine is mutated to cysteine (K64C and K219C), channel activity is significantly decreased in cells and in reconstituted liposomes. Directly tethering K64C to the membrane by modification with decyl-MTS generates high PIP<sub>2</sub> sensitivity in liposomes, even in the complete absence of PL(−)s. The results provide a coherent molecular mechanism whereby PL(−) interaction with a discrete binding site results in a conformational change that stabilizes the high-affinity PIP<sub>2</sub> activatory site.

<sup>1</sup>Department of Cell Biology and Physiology and the Center for Investigation of Membrane Excitability Diseases, Washington University School of Medicine, St. Louis, Missouri 63110, USA. Correspondence and requests for materials should be addressed to C.G.N. (email: cnichols@wustl.edu).

The eukaryotic inwardly rectifying potassium (Kir) channel family includes seven subfamilies expressed in tissues throughout the body and implicated in diverse physiological functions, such that mutations are causally linked to a number of metabolic and genetic diseases including diabetes, heart diseases, Andersen–Tawil syndrome and Epilepsy, ataxia, sensorineural deafness, tubulopathy syndrome<sup>1–3</sup>. Kir channels are regulated by diverse modulators including phosphorylation by protein kinase A, auxiliary proteins (sulphonylurea receptors, G protein  $\beta\gamma$  subunits),  $\text{Na}^+$ , ATP, pore blockers ( $\text{Mg}^{2+}$ , polyamine)<sup>4,5</sup> and ethanol<sup>6</sup>. However, and in common with many other channels and membrane proteins, all Kir channels require the binding of phosphatidyl-4,5-bisphosphate ( $\text{PI}(4,5)\text{P}_2$ ;  $\text{PIP}_2$ ) for activation<sup>7–9</sup>.

The study of  $\text{PIP}_2$  effects on ion channels has been facilitated by its low natural abundance (0.4–1.5% of anionic phospholipids)<sup>10</sup> and the availability of methods to manipulate  $\text{PIP}_2$  concentrations in the membrane during electrophysiological experiments, either through enzymatic reactions or charge screening by specific antibodies or polycations<sup>8,11,12</sup>. Other anionic lipid species such as phosphatidylglycerol (PG), phosphatidic acid (PA), phosphatidylserine (PS), phosphatidylinositol (PI) and cardiolipin are also present at variable, and generally much higher, concentrations in native membranes<sup>13,14</sup>. Membrane lipid composition can vary under different physiological and pathological conditions<sup>15–17</sup>, and has been invoked causally in various diseases conditions<sup>18–20</sup>. However, progress in understanding the role of specific lipids in controlling of membrane protein function has been limited<sup>21</sup> because they cannot be readily manipulated in native membranes. Reconstitution of purified proteins in synthetic membranes makes it possible to study function in defined systems of known lipid and protein composition, and the availability of purified eukaryotic Kir2 channel protein, reconstituted into liposomes of defined composition has led us to the discovery that bulk anionic phospholipids are required in the membrane to generate the high  $\text{PIP}_2$  sensitivity that is necessary for physiological activation<sup>22</sup>. Multiple anionic phospholipids ( $\text{PL}(-)$ s) including PG, PA, PS, PI and cardiolipin can fulfill this requirement, nonspecifically shifting  $\text{PIP}_2$  sensitivity of Kir2.x channels by  $\sim 100$ -fold<sup>22</sup>. Increasing  $\text{PL}(-)$  in the membrane increases both channel open probability and unitary conductance<sup>22</sup>.

Recent crystal structures of Kir2.2 and Kir3.2 channels reveal a defined  $\text{PIP}_2$ -binding site and provide clues to how this binding may open the channel, but the nature of the  $\text{PL}(-)$ -binding site and the molecular mechanism of the synergistic effect on  $\text{PIP}_2$  activation are unknown. Hansen *et al.*<sup>23</sup> showed that the anionic pyrophosphatidic acid (PPA) head group ( $-3$  formal charge) can bind to essentially the same site as  $\text{PIP}_2$  (refs 23,24). This observation might suggest that activatory  $\text{PL}(-)$ s bind to the same site as  $\text{PIP}_2$ , although such a scenario provides no intuitive explanation for the synergistic effect of anionic lipids on  $\text{PIP}_2$  action. Therefore, we began this study by characterizing the functional effect of PPA on Kir2.1 channel activation. We first showed that PPA is clearly inhibitory, with no detectable synergistic effect on  $\text{PIP}_2$  activation. We then used extensive docking simulations of  $\text{PL}(-)$  head groups to identify a putative  $\text{PL}(-)$ -binding site that is formed by two lysine residues (K64 and K219) from two neighbouring subunits. The role of the two residues was examined experimentally through cysteine mutation and subsequent modification by a methanethiosulphonate (MTS) reagent containing a strongly hydrophobic moiety (a decyl chain). Neutralization of K64 and K219, by cysteine substitution reduces channel activity, but ‘lipid-tethering’ by modification of K64C with decyl-MTS strikingly induces high-affinity  $\text{PIP}_2$  activation, even in the absence of  $\text{PL}(-)$ . These functional assays implicate K64 as the primary residue

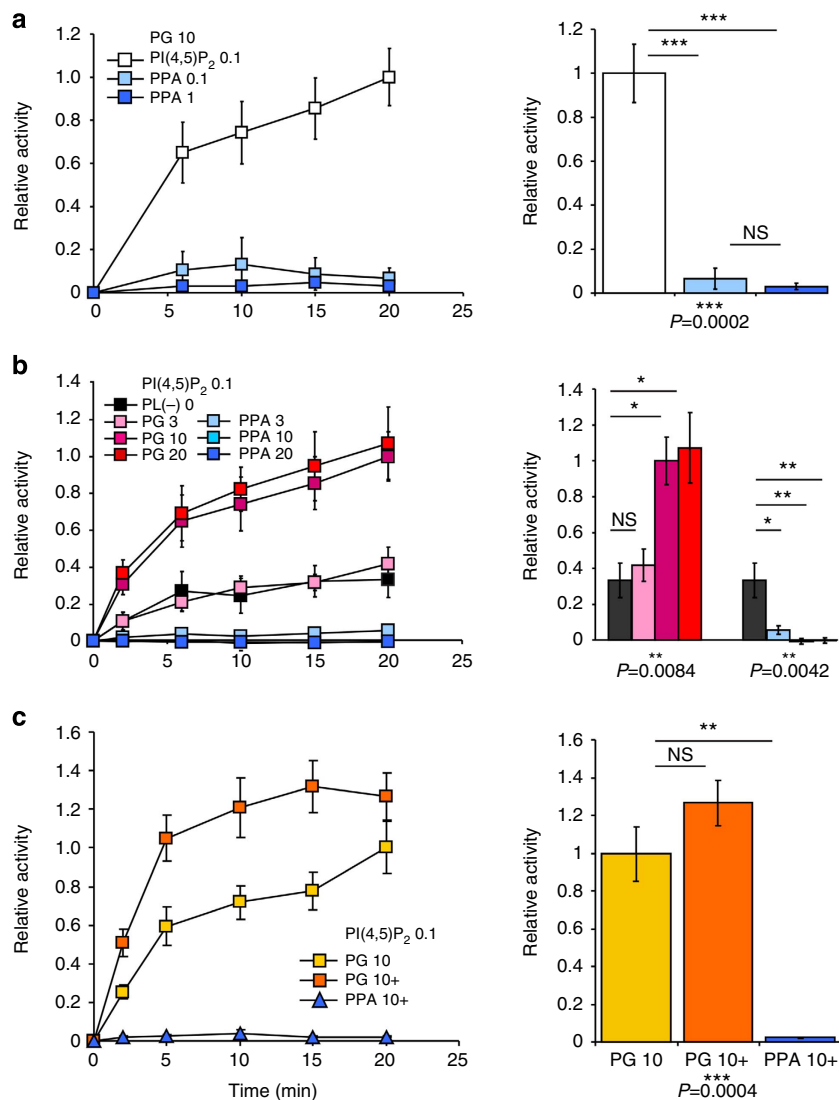
for  $\text{PL}(-)$  interaction and pulling of the cytoplasmic Kir domain towards the membrane as the molecular mechanism underlying the  $\text{PL}(-)$  synergistic effect on  $\text{PIP}_2$ -dependent channel activation.

## Results

**PPA is a competitive inhibitor of Kir2.1 channels.** Recently resolved crystallographic structures of chicken Kir2.2 (ref. 23) reveal a specific  $\text{PIP}_2$ -binding site at the interface of the M2 helix and the cytoplasmic domain. These structures also show that an anionic lipid with a PPA head group (PPA) can bind at essentially the same site. This raises the possibility that the secondary requirement for anionic lipids to activate Kir2 channels<sup>22</sup> might be satisfied through binding to this site, perhaps if  $\text{PIP}_2$  was only required at a subset of the subunits. Alternatively, PPA may act as either a competitive analogue or competitive antagonist of  $\text{PIP}_2$  interaction and not as a secondary activator  $\text{PL}(-)$ . To resolve these possibilities, the functional role of PPA was first determined. Purified wild-type (WT) Kir2.1 protein was reconstituted into liposomes of different lipid compositions. First, to test whether PPA could act as a  $\text{PIP}_2$  analogue and activate channels in the absence of  $\text{PIP}_2$ , liposomes were formed with PPA (zero, 0.1 or 1%), plus 10% 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphoglycerol (POPG) (to supply the secondary nonspecific lipid requirement), and 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphatidylethanolamine (POPE) for the remaining fraction of the lipid (here and in all further liposome experiments). As shown in Fig. 1a, PPA did not activate the channel under these conditions, suggesting that PPA cannot act as a functional substitute for  $\text{PIP}_2$ . We then tested whether PPA could act as a  $\text{PL}(-)$  and suffice for the secondary anionic lipid effect in the presence of 0.1%  $\text{PIP}_2$ . Although increasing POPG augmented channel activity as shown before<sup>23</sup> (Fig. 1b), PPA significantly decreased channel activity at all concentrations and channel activity was completely inhibited at  $>10\%$  PPA. To further confirm the inhibitory effect of PPA, channel activity was measured in the presence of 0.1%  $\text{PIP}_2$ , 10% POPG with or without additional 10% of either POPG or PPA (Fig. 1c). Although additional 10% POPG (that is, 20% total POPG) further enhanced activity ( $P > 0.05$ , one-way analysis of variance, Turkey’s honestly significant difference *post hoc* test), 10% PPA again completely abolished the channel activity. These results are reminiscent of competitive inhibition of  $\text{PIP}_2$ -activated channel activity by other  $\text{PIP}_2$ s<sup>22,25</sup> and clearly show that PPA acts as a competitive inhibitor of  $\text{PIP}_2$  and not as an activatory  $\text{PL}(-)$ .

## Docking simulations identify a potential $\text{PL}(-)$ -binding site.

We previously used docking simulations to identify the crystallographic  $\text{PIP}_2$ -binding site (‘primary’ site)<sup>23</sup> as a main binding site for  $\text{PIP}_2$  as well as an additional (‘secondary’) site, located at the interface between two adjacent subunits near the end of the slide helix and the apex of the  $\beta\text{C}$ - $\beta\text{D}$  loop<sup>26</sup>. To determine potential docking sites for other anionic lipids, we have now performed extensive docking simulations of various lipids to Kir2.1, in both the absence (‘Apo-docking’) and presence (‘ $\text{PIP}_2$ -bound-docking’) of  $\text{PIP}_2$  at the crystallographic site. The searched space is shown in Fig. 2a,b; the  $\text{PIP}_2$  head group as located in the crystal structure (PDB 3SPI) was removed in the Apo-docking but present in  $\text{PIP}_2$ -bound-docking below. Tested lipids included negatively charged PPA, PA, PG, PS as well as neutral phosphatidylethanolamine (PE) and phosphatidylcholine (PC). Lipid head group structures and partial charges used for docking simulations are shown in Supplementary Fig. S1. Accepted poses, based on appropriate orientation relative to the bilayer normal and proximity to the membrane (see Methods),



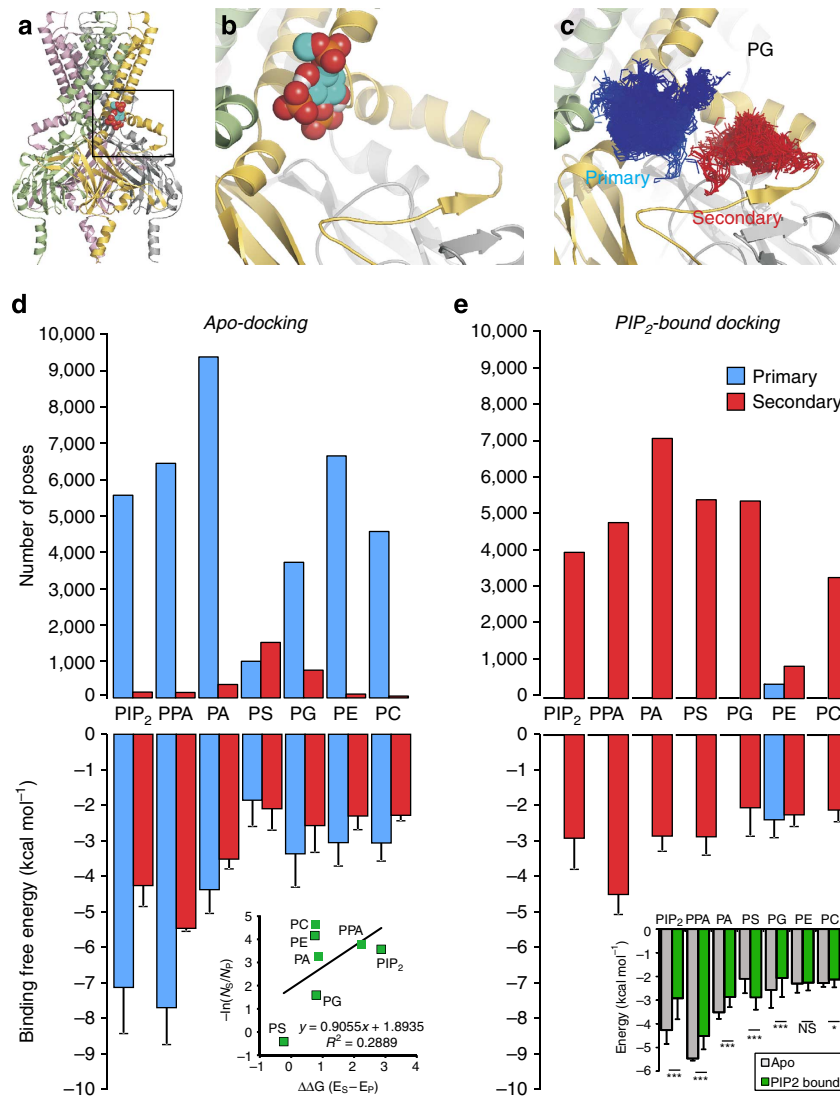
**Figure 1 | PPA is a competitive inhibitor of PIP<sub>2</sub> (left) <sup>86</sup>Rb<sup>+</sup> uptake (normalized to maximum activity of WT Kir2.1 in 0.1% PIP<sub>2</sub>, 10% POPG). (Right) Maximum activity for 20 min. (a) Ten percent POPG was present in each case, and PPA or PIP(4,5)P<sub>2</sub> as indicated. *n* = 3, mean ± s.e.m.; \*\*\*, *P* < 0.001 compared to the activity at PIP<sub>2</sub> 0.1 (*P* = 0.0002, one-way analysis of variance (ANOVA), Tukey's honestly significant difference (HSD) *post hoc* test). (b) PIP<sub>2</sub> (0.1%) was present in each case, and increasing POPG (pink) or PPA (blue) were added as indicated. Control experiment with no POPG or PPA is shown in black box. *n* = 3, mean ± s.e.m.; \*, *P* < 0.05 and \*\*, *P* < 0.001 compared to the activity at PL(-) 0 (*P* = 0.0084 and *P* = 0.0042, one-way ANOVA, Tukey's HSD *post hoc* test). (c) PIP<sub>2</sub> (0.1%) and 10% POPG were present in each case, with an additional 10% POPG (total 20%) or 10% PPA, as indicated. *n* = 3, mean ± s.e.m. \*\*, *P* < 0.01 compared to the activity at PG 10 (*P* = 0.0004, one-way ANOVA, Tukey's HSD *post hoc* test).**

were clustered using a K-clustering algorithm<sup>27</sup>. For each tested lipid, optimal clustering efficiency was achieved by division into two clusters (see Supplementary Fig. S2 and Methods), corresponding to the previously suggested primary and secondary phospholipid sites. As an illustrative example, every accepted (3,727 at primary and 758 at secondary out of 10,000) docked poses of PG is shown in Fig. 2c.

In Apo-docking, all tested lipids, except PS, docked to the primary site with higher frequency and greater binding affinity compared with the secondary site (Fig. 2d, top panel; Tables 1 and 2). The inconsistent behaviour of PS may originate from its unique zwitterionic head group structure with a positively charged amine and negatively charged carboxylic group. The relative distribution of docking events between the two sites was in reasonable agreement with the Boltzmann distribution, based on predicted binding free energies (Fig. 2d, inset). Predicted binding at the primary site was much stronger for PIP<sub>2</sub> and

PPA than for other lipids that at the primary site (Fig. 2d), but binding free energy at the secondary site was similar for all lipid species (maximum difference: 3.36 kcal mol<sup>-1</sup>, mean ± s.d.: -3.37 ± 1.24 kcal mol<sup>-1</sup>), consistent with the relatively non-specific and redundant character of the secondary PL(-)-activating effect<sup>22</sup>.

We then carried out a series of docking simulations of the same lipids with PIP<sub>2</sub> pre-bound at the primary site ('PIP<sub>2</sub>-bound-docking'; Fig. 2e). In this situation, PIP<sub>2</sub> precluded additional binding of all but one of the lipids at the primary site and shifted binding almost exclusively to the secondary site, without substantially altering binding free energies at that site (Fig. 2e). PE was one exception in that it still docked to the primary site, essentially next to the PIP<sub>2</sub> head group, presumably facilitated by the small head group size and high charge density in the amine group. These simulations, using a rigid backbone model, therefore indicate that when PIP<sub>2</sub> is pre-docked at the primary site, the



**Figure 2 | Docking simulations of anionic lipids to Kir2.1 without or with PIP<sub>2</sub> pre-bound at the crystallographic site.** (a) The tetrameric Kir2.1 homology model based on the PIP<sub>2</sub>-bound chicken Kir2.2 crystal structure (3SPI). Subunits are coloured in yellow, silver, pink and green in order. The PIP<sub>2</sub> ligand is shown in spheres at the crystallographic PIP<sub>2</sub> site. The space searched for docking simulations is marked by a black square, and in detail in **b**. (c) All accepted poses of PG in 'Apo-docking' are shown, subgrouped into two clusters: 3,727 poses are in the primary site (blue) and 758 poses in the secondary site (red). (d) Results from Apo-docking without crystallographic PIP<sub>2</sub>: the number and the mean binding free energy of accepted poses in primary (blue) or secondary (red) site are shown (mean  $\pm$  s.d.).  $n$  corresponds to the number of accepted poses at each site shown in Table 1. Inset shows the energy difference between primary ( $E_p$ ) and secondary ( $E_s$ ) site versus inverse natural log of docking frequency ratio ( $N_s/N_p$ ) between the two sites, which reasonably well follows the Boltzmann distribution. Data points refer to individual lipids from the Apo-docking simulation, and the line is the best fit to the data set. (e) Results (as in **d**) from PIP<sub>2</sub>-bound-docking simulated with PIP<sub>2</sub> pre-bound at the crystallographic-binding site (mean  $\pm$  s.d.). Inset shows the binding free energy at the secondary site from Apo- and PIP<sub>2</sub>-bound-docking, plotted for each lipid with their mean and s.d.  $n$  corresponds to the number of accepted poses at the secondary site in the Apo- and PIP<sub>2</sub>-bound docking respectively shown in Table 1. The decrease in binding free energy is substantially significant (\*,  $P < 0.05$  and \*\*\*,  $P < 0.001$  compared to the binding free energy in Apo-docking) for all anionic lipids determined by a two-tailed Student's  $t$ -test.

secondary site becomes the preferential site for PL(–) binding and no new sites become apparent.

#### Two lysines are the main determinants of PL(–) interaction.

To identify residues that directly interact with lipids at the secondary site, hydrogen bonding between lipids and neighbouring residues was assessed (Supplementary Table S1). Figure 3a shows residues that have more than 10% hydrogen bond (H-bond) frequency out of all H-bonds detected between the accepted poses combined from Apo-docking (left) and PIP<sub>2</sub>-bound-docking (right) experiments. The H-bond frequency results were

qualitatively similar for both cases, with two features of note. First, K64, at the N-terminal end of the slide helix of one subunit and K219 at the apex of the  $\beta$ C- $\beta$ D loop of the adjacent subunit are the two key residues that make direct contacts with the docked lipids (Fig. 3). In typical docked structures, these two positive side chains contributed to coordination of the negative phosphate group (Fig. 3b,c). Functional tests (below) demonstrate that these two residues are necessary for PL(–) activation and that chemical tethering of residue 64 to the membrane generates PL(–)-independent, high sensitivity to PIP<sub>2</sub> activation. Second, the residues found in the secondary site do not overlap with the residues forming the primary site (Fig. 3b). This observation

demonstrates that the secondary site is indeed a unique and separate site from the primary PIP<sub>2</sub>-binding site.

**Mutant channel activity is reduced in cell membranes.** To experimentally test the functional relevance of the secondary site

as a nonspecific PL(–)-binding site, we mutated the K64 and K219 residues to cysteines, either individually (K64C, K219C) or together (K64/219C) in a ‘cysteine-less’ Kir2.1 background construct (CLS\_Cont; see Methods). Both *in vitro* biochemical (Supplementary Fig. S3) and mammalian cell-based (Fig. 4) assays show that CLS\_Cont is functionally equivalent to WT Kir2.1 channels. Mutant channel activity was first tested using a cellular <sup>86</sup>Rb<sup>+</sup> efflux assay (Fig. 4a) after transient expression in mammalian COSm6 cells (Fig. 4a). Although efflux rates were not different between WT Kir2.1 and CLS\_Cont, each mutation substantially decreased channel activity, demonstrating that each residue is functionally important for maintenance of channel activity in mammalian cell membranes.

We carried out additional patch-clamp experiments on these transfected cells (Fig. 4b). In cell-attached patches, robust inwardly rectifying currents were measured through WT Kir2.1 as well as CLS\_Cont channels. Following excision, channel currents were maintained, and typically increased slightly, as rectification gradually decreased because of slow loss of polyamines from within the recessed patch<sup>28</sup>. In contrast, inward rectifier currents were either very small, or undetectable, in cell-attached membrane patches in which mutant channel proteins (K64C, K219C and K64/219C) were expressed. Although K64/219C activity remained undetectable, spermine-sensitive

**Table 1 | Number of accepted poses.**

	Apo-docking		PIP <sub>2</sub> -bound docking	
	Primary	Secondary	Primary	Secondary
PIP <sub>2</sub>	5,568	157	0	4,030
PPA	6,448	147	0	4,853
PA	9,373	364	0	7,169
PS	1,005	1,525	0	5,482
PG	3,727	758	0	5,448
PE	6,652	104	395	891
PC	4,574	45	0	3,335

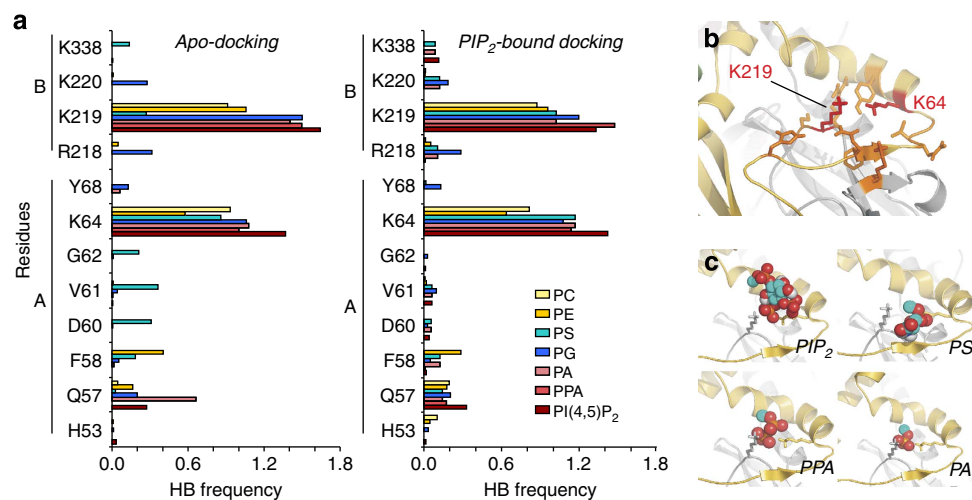
PA, phosphatidic acid; PC, phosphatidylcholine; PE, phosphatidylethanolamine; PG, phosphatidylglycerol; PIP<sub>2</sub>, phosphatidyl-4,5-bisphosphate; PPA, pyrophosphatidic acid; PS, phosphatidylserine.

Number of accepted poses of Apo-docking and PIP<sub>2</sub>-bound docking in the primary and secondary sites are presented for individual lipids.

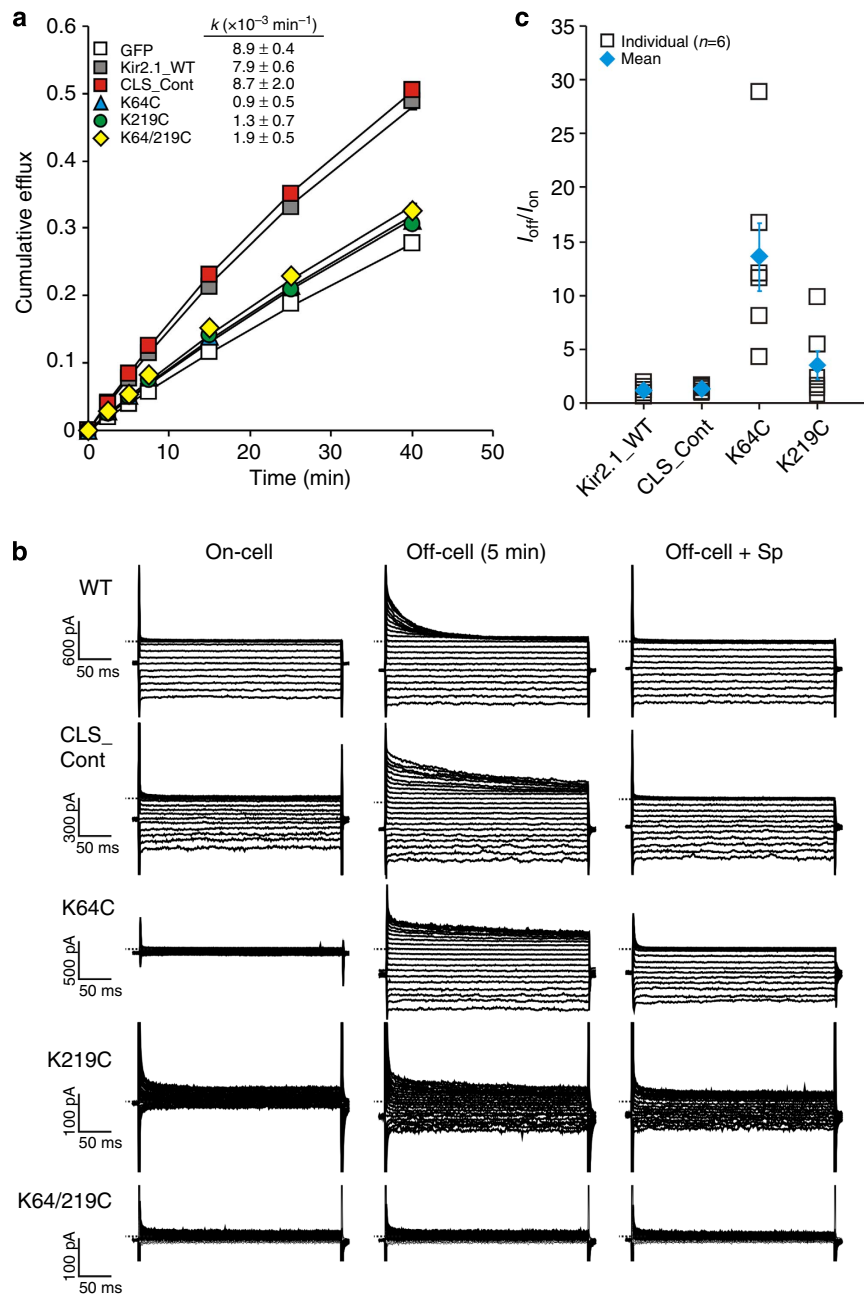
**Table 2 | Binding free energy of accepted poses.**

	Apo-docking		PIP <sub>2</sub> -bound docking		Secondary	P-value
	Primary	Secondary	Primary	Secondary		
PIP <sub>2</sub>	–7.133 ± 1.301	–4.266 ± 0.588		–2.922 ± 0.883	1.344 ± 1.061	<0.0001
PPA	–7.702 ± 1.042	–5.467 ± 0.090		–4.516 ± 0.564	0.950 ± 0.571	<0.0001
PA	–4.381 ± 0.665	–3.517 ± 0.278		–2.863 ± 0.432	0.654 ± 0.514	<0.0001
PS	–1.861 ± 0.736	–2.099 ± 0.607		–2.886 ± 0.519	–0.787 ± 0.799	<0.0001
PG	–3.370 ± 0.939	–2.572 ± 0.757		–2.064 ± 0.802	0.507 ± 1.103	<0.0001
PE	–3.054 ± 0.662	–2.302 ± 0.386	–2.405 ± 0.503	–2.262 ± 0.330	0.040 ± 0.508	0.251
PC	–3.063 ± 0.508	–2.281 ± 0.159		–2.131 ± 0.326	0.150 ± 0.363	0.002

PA, phosphatidic acid; PC, phosphatidylcholine; PE, phosphatidylethanolamine; PG, phosphatidylglycerol; PIP<sub>2</sub>, phosphatidyl-4,5-bisphosphate; PPA, pyrophosphatidic acid; PS, phosphatidylserine. Binding free energy differences at secondary site between Apo- and PIP<sub>2</sub>-bound docking (mean ± s.d.) of all poses of individual lipids clustered into each site are listed with P-value for statistical significance based on unpaired *t*-test. *n* corresponds to the number of accepted poses shown in Table 1 for each lipid and site respectively.



**Figure 3 | Residue contributions to the secondary site.** (a) Hydrogen bond frequency between all accepted poses and surrounding residues at the secondary site are shown for Apo- (left) and PIP<sub>2</sub>-bound-docking (right). All residues having H-bonding frequency > 0.1 with any one of the tested lipids are shown. (b) Spatial location of the residues shown in a are depicted. The two residues, K64 and K219, of the highest contact frequency, are shown in red and others in orange. (c) Example structures of different lipids at the secondary site. K64, K219 and individual lipids (PIP<sub>2</sub>, PPA, PS and PA) are shown in stick and in atom colours.



**Figure 4 | Mutation of secondary site lysines inhibits channel activity in cell membranes.** (a) Cumulative  $^{86}\text{Rb}^+$  efflux from Cosm6 cells versus time for WT (grey) and mutant Kir2.1 channels (CLS\_Cont: red, K64C: blue, K219C: green, K64/219C: yellow). Nonspecific background flux was determined from GFP-transfected cells (white). Error bars are smaller than the symbols. Lines are the best fits of the single exponential equation described in the Methods, from which a rate constant ( $k$  in  $\text{min}^{-1}$ ) for each protein is obtained and shown in the figure. (b) Channel activity determined electrophysiologically on-cell (left), 5 min after excision (middle) and then in the presence of  $10 \mu\text{M}$  spermine (right). (c) Current increase on patch excitation (off-cell;  $I_{\text{off}}$ ) from cell-attached (on-cell;  $I_{\text{on}}$ ) configuration. The fold-increase in current is shown for individual patches (white square), and mean and s.e. for each construct ( $n = 6$ ).

inward rectifier currents slowly developed through K219C and especially K64C channels following patch excision, until current amplitudes were comparable to those measured from CLS\_Cont patches (Fig. 4b,c). This remarkable behaviour demonstrates that channels are indeed present in the cell membranes but substantially closed when exposed to the intact intracellular milieu. The normal cytoplasmic environment is strongly reducing<sup>29</sup>, but membrane excision exposes the cytoplasmic face of the membrane to an oxidizing environment. Given the proximity of the two residues within the protein, K64C/K219C double mutants are likely to be

disulphide bonded to each other (see Supplementary Fig. S4), but single mutant K64C and K219C will probably be reduced to free cysteines in the intact cell. They will be free to oxidize to surrounding proteins or lipids following patch excision, and we suggest that this oxidation then effectively tethers these residues to the membrane, thereby providing similar conformational energy to that which is normally provided by PL(−) interaction, to promote channel opening. This hypothesis is strongly supported by experiments below on recombinant channels in which the idea of ‘membrane tethering’ is directly assessed.



**Mutant channel activity is reduced in synthetic membranes.**

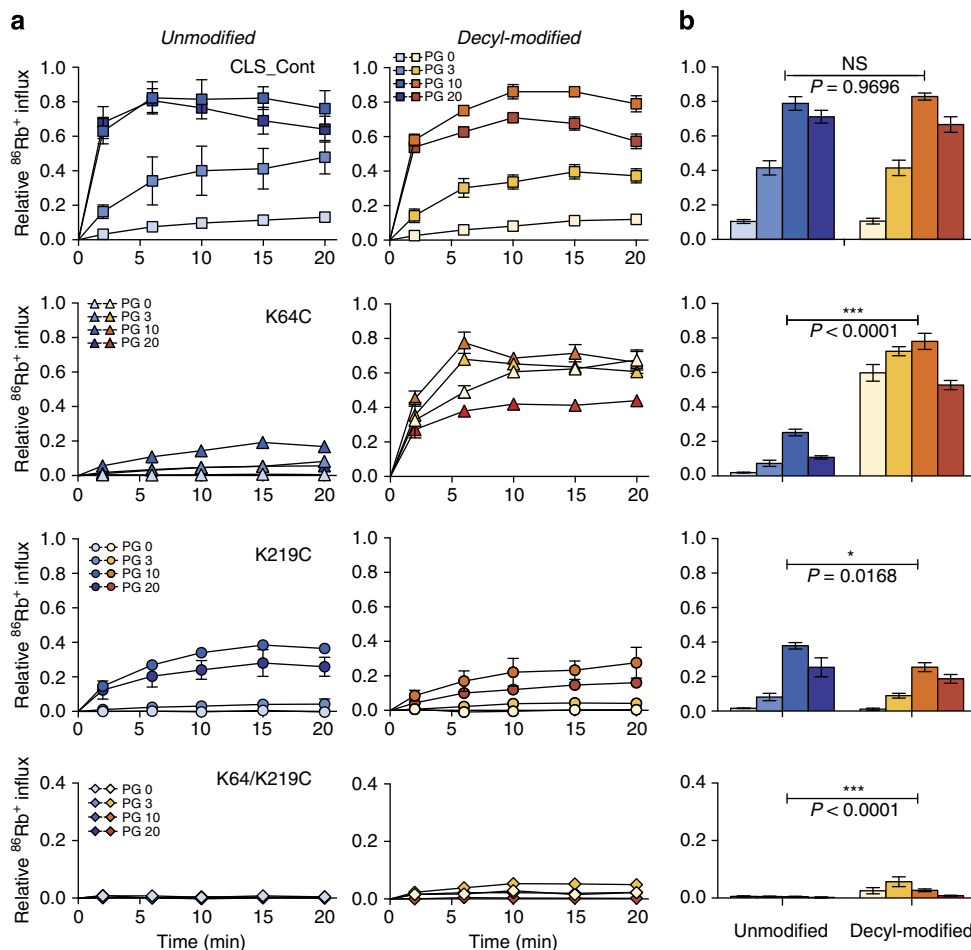
Channel activities were also assessed from recombinant proteins incorporated into liposomes of defined lipid composition and in which nonspecific influence from cellular components is excluded. Control and mutant proteins were expressed in *Saccharomyces cerevisiae*, and then purified and reconstituted into liposomes with increasing POPG concentrations in the presence of constant 0.1% PIP<sub>2</sub> (Fig. 5a, left). <sup>86</sup>Rb<sup>+</sup> influx into liposomes was measured as a function of time to assess channel activity. CLS\_Cont protein showed essentially identical PL(–)-dependent activation (Fig. 5a, left) to that of WT protein<sup>22</sup>; activity reached maximum at ~10% POPG and there was no significant further activation at 20% POPG. Similar PL(–)-dependent activation was still observed but maximal channel activity was reduced slightly for the K219C mutant, markedly so for K64C, and there was no detectable activity from the K64C/K219C double mutant (Fig. 5a, left).

**Membrane tethering at secondary site restores channel activity.**

Kir2 subfamily channels specifically require PI(4,5)P<sub>2</sub> for activation although all six other PIP species can interact with the PIP<sub>2</sub> site<sup>22,26,30,31</sup>. On the other hand, multiple different anionic phospholipids can satisfy the synergistic PL(–) effect with similar concentration dependence<sup>22</sup>. This suggests that the

molecular mechanism exerted by PL(–) binding might be through a simple conformational change. We hypothesize this to be a generic pulling of the Kir domain towards the membrane. To directly test this idea, purified cysteine mutant and control proteins were modified in the detergent micelle state by decyl-MTS. This places a very hydrophobic 10-carbon acyl chain on modified cysteines, which will strongly partition into the membrane phase if appropriately located, as with modification of K64C and K219C. PL(–)-dependent activation of modified proteins was then again assessed using the <sup>86</sup>Rb<sup>+</sup> influx assay. Decyl modification did not affect CLS\_Cont activation (Fig. 5a, right), indicating that the expected disulphide bond (cystine) generated by the remaining two cysteines on the extracellular site of the protein remains as a disulphide bond. In contrast, decyl modification substantially increased the activity of single mutant K64C channels (Fig. 5a, right) comparable to the activity of maximally active CLS\_Cont channels (that is, in 0.1% PIP<sub>2</sub> plus 10% PG). Even more strikingly, marked channel activity was now present at zero POPG, R64C + decyl becoming almost PL(–) independent (Fig. 5a, right). Decyl modification of K219C decreased maximum channel activity but had no significant effect on PL(–) dependence (Fig. 5a, right).

The double mutant (K64/K219C) was inactive before modification, but there was some low level of activity of decyl-MTS-



**Figure 5 | Channel activation by decyl modification of secondary site residues.** (a) Anionic lipid (PL(–))-dependent activation of CLS\_Cont, K64C, K219C and K64/219C was determined without (left, blue colours) or with (right, red colours) modification by decyl-MTS by measurement of <sup>86</sup>Rb<sup>+</sup> uptake into liposomes. PIP<sub>2</sub> (0.1%) was present in all cases, and POPG varied as indicated. <sup>86</sup>Rb<sup>+</sup> uptake was normalized to the valinomycin count of the same sample. (b) Maximum channel activities with or without modification are plotted. n = 6 (unmodified), 6 (modified) for CLS\_Cont, n = 6, 5 for K64C, n = 6, 5 for K219C and n = 6, 5 for K64/K219C. Error bars, s.e.m. \*, P < 0.05 and \*\*\*, P < 0.001 compared to unmodified protein (two-way analysis of variance).

modified channels (Fig. 5a, right). We suggest that inactivity of the unmodified channels may be a result of disulphide bond formation between the two introduced cysteines. The low level of activity following decyl-MTS treatment was probably attributable to subsequent modification of a low level of reduced cysteines in the presence of dithiothreitol ( $<50\ \mu\text{M}$ ). Disulphide bond formation between the two introduced cysteines is supported by the presence of dithiothreitol-sensitive dimers of the double-mutant protein, but not CLS\_Cont protein, in western blots (Supplementary Fig. S4).

### Membrane tethering at secondary site enhances PIP<sub>2</sub> sensitivity.

In the absence of POPG or other PL(-), WT Kir2.1 channels are very weakly activated by supra-physiological levels of PIP<sub>2</sub>, whereas 25% PL(-) left-shifts PIP<sub>2</sub> sensitivity of human Kir2.1 channels by  $\sim 100$ -fold, making the channel fully active at only  $\sim 0.1\%$  PIP<sub>2</sub> (ref. 22). If membrane tethering by decyl modification of K64C is functionally equivalent to the PL(-) interaction, then it should also cause left-shift in PIP<sub>2</sub> sensitivity of the modified channel in the absence of PL(-). To test this hypothesis, we assessed the POPG dependence of PIP<sub>2</sub> sensitivity in unmodified and decyl-modified K64C and K219C single mutants (Fig. 6). The maximal activity of unmodified K64C and K219C mutants was lower than CLS\_Cont but all showed a dramatic left-shift in PIP<sub>2</sub> sensitivity in the presence of POPG (Fig. 6, left). POPG dependence of PIP<sub>2</sub> sensitivity of CLS\_Cont and K219C was unaffected by decyl-MTS treatment (Fig. 6a–c, right). However, in striking agreement with the above prediction, the POPG-dependent shift was completely absent in decyl-

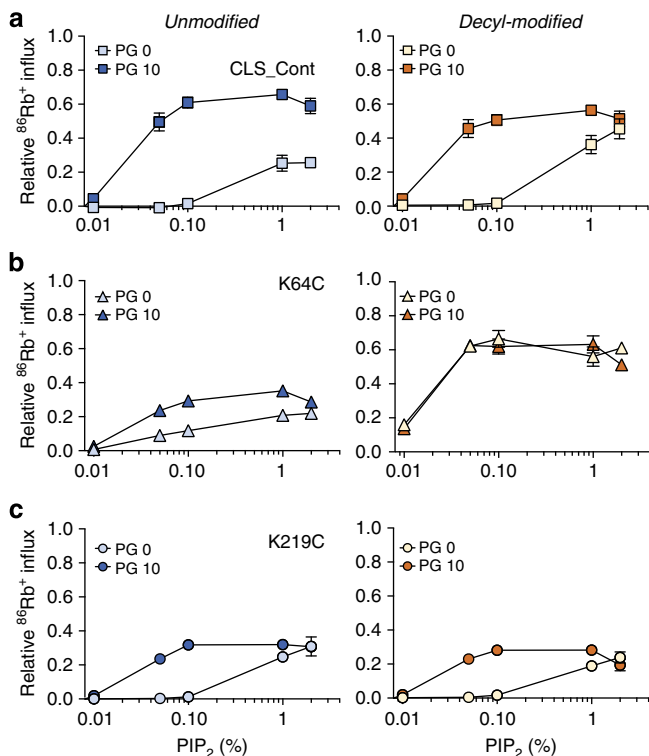
modified K64C (Fig. 6b, right), with the channels exhibiting identical high PIP<sub>2</sub> sensitivity in 0 or 10% POPG. This dramatic result not only suggests that K64 is the key residue for POPG interaction but also indicates that decyl modification of this residue can replace POPG interaction with this site to provide the membrane tethering that energetically sensitizes the channel to PIP<sub>2</sub> activation.

### Discussion

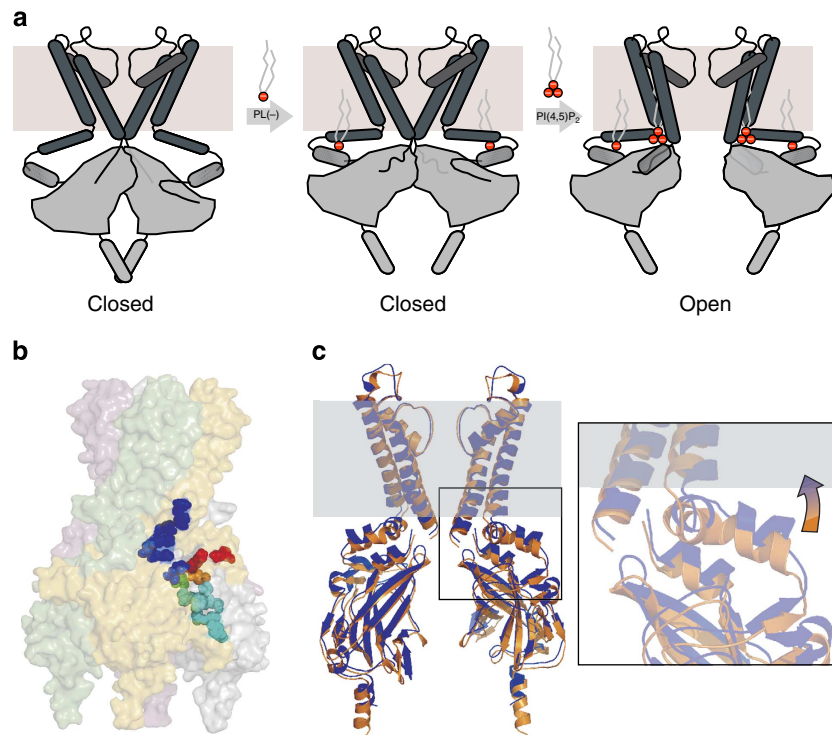
It is becoming increasingly clear that the lipid components of the membrane are major regulators of the activity of ion channels<sup>32</sup>. Interaction with PIP<sub>2</sub> or related PIPs is essential for activation of many channels, including all Kir subfamily members<sup>7–9</sup>, and recent crystal structures have demonstrated a discrete binding site in Kir2.2 channels that underlies this effect, and provides significant insight to the molecular mechanism<sup>23</sup>. More generally mechanistic details of channel–lipid interactions are far from clear. The realization of obligate synergistic PL(-) regulation of PIP<sub>2</sub>-mediated Kir2.1 and Kir2.2 channel activation<sup>22</sup> indicates a previously unrecognized complexity, requiring the simultaneous binding of both lipids, a mechanistic scenario that is currently without explanation. Moreover, the relatively low affinity of PL(-) for this action suggests that, although the interaction is key for physiological channel activity, crystallographic resolution of the interaction may not be forthcoming. To gain further insight to the molecular basis of the PL(-) action, we therefore turned to the powerful approaches of molecular simulation, followed by experimental analysis using biochemical manipulation of channel structure. Using these approaches, we have successfully defined a discrete binding site for PL(-) that is both necessary and sufficient for PL(-) action, and a mechanism by which the interaction leads to PIP<sub>2</sub> sensitivity at physiological PIP<sub>2</sub> levels (Fig. 7a).

Our docking simulations identify two sites to which anionic lipids could bind (Fig. 2). The first site (primary) at which the majority of docking events occurs corresponds to the PIP<sub>2</sub> site identified in X-ray crystal structures<sup>23,24</sup>. This agreement supports our Kir2.1 homology model and the parameters for ligand and docking simulations. When both this and an additional distinct PL(-)-binding (secondary) site are available, almost all lipids dock preferentially to the primary site (Fig. 2d), but if the primary site is occupied by PIP<sub>2</sub>, the secondary site becomes the most frequently docked site (Fig. 2e). Conversely, if the secondary site is pre-occupied by PIP<sub>2</sub>, 100% of PIP<sub>2</sub> docking is to the primary site, indicating that the secondary site is a spatially separate site to which anionic lipids can bind, such that even two PIP<sub>2</sub> molecules can bind to the two sites simultaneously (Fig. 2e).

Two important experimental features of the synergistic PL(-) effect<sup>22</sup> further support this argument. First, the calculated binding free energies at the secondary site vary little between test lipids (maximum difference:  $3.37\ \text{kcal mol}^{-1}$ , mean  $\pm$  s.d.:  $-3.21 \pm 1.24\ \text{kcal mol}^{-1}$ ), much less than the variability of binding free energies at the primary site (maximum difference:  $5.84\ \text{kcal mol}^{-1}$ , mean  $\pm$  s.d.:  $-4.37 \pm 2.22\ \text{kcal mol}^{-1}$ ; Table 2), consistent with the experimental finding that the secondary PL(-) effect is nonspecific and that the efficacy of different lipids are similar<sup>22</sup>. Second, high-affinity PIP<sub>2</sub> sensitivity is specific, no other phospholipids can substitute. In Apo-docking experiments the binding affinity of PIP<sub>2</sub> and of the competitive antagonist PPA were both much higher at the primary site than that of other anionic phospholipids (Fig. 2d). Interestingly, the pre-occupancy of PIP<sub>2</sub> at the primary site actually increases the efficacy of PL(-) binding at the secondary site. PIP<sub>2</sub> at the primary site may create a negative electric potential that prevents the binding of



**Figure 6** | Decyl modification abolishes PG-dependent shift of PIP<sub>2</sub> sensitivity activity-PIP<sub>2</sub> relationship for unmodified and decyl-modified CLS\_Cont (a), K64C (b) and K219C (c) obtained from the maximum <sup>86</sup>Rb<sup>+</sup> uptake over the course of 20 min from experiments (as in Fig. 5) with zero (light blue or yellow) or 10% (dark blue or orange) POPG. ( $n = 3$ –6 in each case, mean  $\pm$  s.e.m.).



**Figure 7 | Proposed model for secondary PL(–) regulation.** (a) Hypothetical model: (i) in the absence of PL(–)s in the membrane, or when PL(–) interaction is lost by neutralization of the two positive charges at the secondary site, the cytoplasmic Kir domain is displaced from the membrane surface. (ii) The presence of PL(–)s in the membrane—or decyl modification of one of these—residues pulls the Kir domain closer to the membrane, which in turn induces conformational change in the primary site to generate high-affinity PIP<sub>2</sub> interaction. (iii) PIP<sub>2</sub> can then fully activate channels at low concentrations (0.05%). (b) The surface of Kir2.1 is shown with the same colour codes in Fig. 2a. Residues (H53, R78, R80, K182, K185, K187 and K188) forming the primary site are shown in blue, and the secondary site (K64 and K219) in red. Residues implicated in interaction with other Kir channel modulators are also shown: L222 for cholesterol and Gβγ regulation are shown in green; the residues (F47, H42, L232, L245, P244, Y242, Y337 and L330) for ethanol and Gβγ are shown in cyan; N216 for Na<sup>+</sup> binding is shown in orange. (c) Structural models of Kir2.1 based on Kir2.2 bound to PIP<sub>2</sub> (3SPI, orange) and Kir3.2 additionally bound to Gβγ (4KFM, blue) to indicate potential motions associated with PL(–) interaction.

negative lipid to other locations of weaker binding free energy, thereby restricting binding to the secondary site, that is, the next available high-affinity site. It also seems likely that the strong negative charges of PIP<sub>2</sub> and PPA are most sensitive to the negative electric potential generated by PIP<sub>2</sub> at the primary site; the binding free energy of most lipids at the secondary site was diminished but the reduction was greatest for PIP<sub>2</sub> and PPA, which brought the binding free energies of all tested lipids at the secondary site to within 3.37 kcal mol<sup>−1</sup> of each other (Fig. 2e, inset). This may explain why PIP<sub>2</sub> is not more efficient than other PL(–) at this action; 10% PIP<sub>2</sub> is required to activate channels as much in zero POPG<sup>22</sup>, as 10% POPG activates channels in 0.1% PIP<sub>2</sub> (ref. 22) (Fig. 6a). Similar binding energies for both lipids at the secondary site provide a simple interpretation of their nonspecific and similar efficacy.

The secondary PL(–)-binding site overlaps with likely sites for allosteric regulation by Na<sup>+</sup> (refs 24,33), cholesterol<sup>34,35</sup> and Gβγ<sup>8</sup> that have been identified in other Kir channels. Na<sup>+</sup> increases the PIP<sub>2</sub> sensitivity of Kir3.2 and Kir3.4 channels, binding in an analogous pocket generated by carbonyl oxygen atoms of the βC-βD loop and the D226 side chain (see Fig. 7b). The side chain of L222 in Kir2.1 (shown in green in Fig. 7b), located at the βC-βD loop, points towards the interior of the large β sheet of the Kir domain and is spatially close to where Na<sup>+</sup> binds in Kir3.2. Although cholesterol inhibits WT Kir2.1 channels<sup>34,36</sup>, mutation of this particular residue (L222I) abolishes cholesterol sensitivity<sup>35</sup>, and mutation of the

equivalent residue (I229) of Kir3.4 to leucine results in constitutively active channels in the absence of Gβγ (ref. 8). Other residues in the βC-βD loop are recognized to be involved in ligand regulation of other subfamily members. The same region (cyan Fig. 7b) is implicated in a common binding site for alcohol<sup>6</sup>, as well as Gβγ (ref. 37) and Gα<sub>i/o</sub> (ref. 38) for allosteric regulation of Kir3 channels. Convergence of allosteric sites is very consistent with the idea that they all regulate sensitization to PIP<sub>2</sub> activation.

The proposed secondary site was experimentally tested through single or double mutation of the two lysine residues. In intact cells, both single and double mutations of K64 and K219 significantly decreased channel activity, although channels are evidently in the membrane and can spontaneously activate following patch excision (Fig. 4). In purified proteins, the effect of the individual mutations is predominantly to reduce lipid sensitivity, in a manner consistent with loss of PL(–) binding and loss of allosteric stabilization of the PIP<sub>2</sub> interaction. Modification with decyl-MTS, which covalently attaches a 10-carbon acyl chain to the cysteine sulfhydryl group, dramatically enhances channel activity of the K64C mutant and effectively maximally enhances PIP<sub>2</sub> sensitivity even in the complete absence of PL(–), thus indicating that decyl modification of K64C is functionally equivalent to PL(–) interaction (Fig. 5). These equivalent effects of decyl-MTS modification and PL(–) interaction at the secondary site on PIP<sub>2</sub> sensitivity support the hypothesis that PL(–) interactions nonspecifically act to tether

the cytoplasmic Kir domain, specifically the N-terminal slide helix, to the membrane and that it is this conformational effect that serves to increase PIP<sub>2</sub> affinity and stabilize the open state, as other allosteric regulators do<sup>39,40</sup> (Fig. 7a).

Vertical motion or tethering of the cytoplasmic domain to the membrane seems to be a critical step for Kir channel opening<sup>40–42</sup>. Direct support for this idea comes from Kir2.2 crystal structures generated either in the absence<sup>43</sup> or presence<sup>23</sup> of anionic lipids (PIP<sub>2</sub> and PPA). These show marked differences in the location of the Kir domain relative to the potential membrane surface. In PIP<sub>2</sub>-bound structures, the C-linker adopts an  $\alpha$ -helical conformation and the Kir domain is pulled towards the potential membrane surface by  $\sim 6\text{ \AA}$ , relative to the Apo structure<sup>23</sup>. Comparison of recent Kir2.2 and Kir3.2 channel structures show that the additional binding of the allosteric regulator G $\beta\gamma$  to Kir3.2 further pulls the slide helix towards the membrane (4KFM)<sup>44</sup> relative to the PIP<sub>2</sub>-bound Kir2.2 structure (3SPI)<sup>23</sup>. Figure 7C shows Kir2.1 homology models based on these two structures to illustrate potential motions coupled to PL(–) interaction. Rotation of the cytoplasmic domain with respect to the membrane as a result of G $\beta\gamma$  binding is suggested to be part of the gating mechanism<sup>44,45</sup> and may well occur with PL(–) binding to Kir2 channels, although our experiments cannot address this possibility.

Pulling of the Kir domain of bacterial Kir channels (KirBac1.1) through membrane tethering also affects channel activity<sup>46</sup>. In this case, membrane tethering by decyl-MTS modification of introduced cysteines in the N-terminus or C-terminus ends of the slide helix resulted in opposite effects on KirBac1.1 activity, with tethering of the N terminus (R49C, that is, away from the pore axis) augmenting the channel activity, and tethering of the C terminus (L56C, close to the pore axis) reducing channel activity<sup>46</sup>. It is noteworthy that R49 of KirBac1.1 is structurally quite close to K64 of human Kir2.1, although these residues do not align exactly (Supplementary Fig. S5). Moreover, recent analysis of sequence elements affecting PL(–) activation of KcsA indicates residues in exactly the same N-terminal slide helix region<sup>47</sup>. These results suggest that cytoplasmic domain tethering to the membrane through PL(–) binding at the secondary site may be a broadly conserved gating mechanism shared between pro- and eukaryotic K<sup>+</sup> channels.

The K64 and K219 residues are loosely conserved among the seven eukaryotic Kir channel subfamilies (Supplementary Fig. S5). Kir2 and Kir4 channels contain both lysines, Kir1 and Kir6, and Kir7 subfamily members have one or other of the two basic residues and Kir3 and Kir5 channels contain neither. Interestingly, physiologically, Kir5 subunits only appear to form functional channels as heterotetramers with Kir4 subunits<sup>48</sup>, and PL(–) regulation might then be conferred on the channels by the Kir4 subunits. As discussed above, Kir3 channels exhibit allosteric regulation of PIP<sub>2</sub> activation by other modulators, including G $\beta\gamma$  and Na<sup>+</sup>. It seems possible that these other modulators may convey the same effect on GIRK channels that PL(–) has on Kir2 channels, consistent with the cytoplasmic domain, and in particular the N-terminal slide helix, being pulled towards the membrane in the G $\beta\gamma$ -bound crystal structure<sup>44</sup> (c.f. Fig. 7c).

## Methods

**Human Kir2.1 protein purification.** WT and mutant Kir2.1 proteins, tagged by Flag-His8 peptide at the C terminus, were expressed in and purified from the FGY217 strain of *S. cerevisiae*<sup>49</sup>. Expression levels and purification yields were similar for WT and mutant proteins. Mutagenesis was performed using QuickChange II site-directed mutagenesis kits (Stratagene Cloning Systems, CA) and verified by sequencing. Cysteine-less mutant Kir2.1 (CLS\_Cont) was generated by introduction of a series of mutations: C43S, C54V, C76V, C89I, C101L, C149F, C169V, C209S, C311A, C356S and C375S. The two remaining cysteines (C122 and

C154) are known to form a very stable intrasubunit disulphide bond<sup>50</sup> and were not substituted to maintain functional channels.

**<sup>86</sup>Rb<sup>+</sup> uptake assay.** Channel activity was assessed by <sup>86</sup>Rb<sup>+</sup> uptake into proteoliposomes<sup>25</sup>. POPE and POPG lipids were dissolved at 10 mg ml<sup>–1</sup> concentration in buffer A (450 mM KCl, 10 mM HEPES, 4 mM NMG, pH 7.4) with 35 mM CHAPS. PIP<sub>2</sub> was solubilized at 1 mg ml<sup>–1</sup> concentration in 10 mg ml<sup>–1</sup> POPE solution. One milligram of lipid mixture in 100  $\mu$ l was incubated at room temperature for 2 h, and 5  $\mu$ g of protein was added and incubated for another 20 min. A lipid–protein mixture was run through partly dehydrated G-50 beads pre-equilibrated by buffer A containing additional 0.5 mM EGTA, which led to proteoliposome formation. Then, the proteoliposomes were run through partly dehydrated G-50 beads pre-equilibrated by buffer B (400 mM sorbitol, 10 mM HEPES, 4 mM *N*-methyl-D-glucamine, 0.5 mM EGTA, pH 7.4) to remove external KCl. Four hundred microlitres of buffer C containing <sup>86</sup>Rb<sup>+</sup> (PerkinElmer, Waltham, MA) at  $\sim 4.5$  nCi was added at time zero. Equal amount of sample (60  $\mu$ l) was collected and run through cation exchange beads where the trace amount of <sup>86</sup>Rb<sup>+</sup> in the external solution was captured by the beads. Uptake was measured over the first 20 min after commencing the assay and normalized to the maximum uptake in valinomycin. In Fig. 1, the relative channel activity was obtained by renormalizing to the maximum activity of WT Kir2.1 in optimal lipid composition of 0.1% PI(4,5)P<sub>2</sub>, 90% POPE and 10% POPG. For cysteine modification, 100  $\mu$ M Decyl-MTS (Toronto Research Chemicals, Ontario, Canada) was added to protein–lipid mixtures and incubated for 15 min before proteoliposomes were formed by passage over a size-exclusion column<sup>46</sup>. Brain PI(4,5)P<sub>2</sub>, POPE, POPG, dioleoylglycerol pyrophosphate (PPA) were purchased from Avanti Polar Lipids.

**<sup>86</sup>Rb<sup>+</sup> efflux assay.** COSm6 cells in 12-well plates were transfected with plasmids (400 ng) carrying Kir2.1 constructs cloned into pcDNA3.1(–), and pGreenLantern green fluorescent protein (GFP; 100 ng; Life Technologies, Grand Island, NY) mixed with Fugene 6 (Promega, Madison, WI) and cultured for an additional 24 h. New culture medium containing 1  $\mu$ Ci ml<sup>–1</sup> <sup>86</sup>RbCl was added, cells were incubated for a further 12 h and then washed twice with Ringer's solution composed of (in mM) 118 NaCl, 2.5 CaCl<sub>2</sub>, 1.2 KH<sub>2</sub>PO<sub>4</sub>, 4.7 KCl, 25 NaHCO<sub>3</sub>, 1.2 MgSO<sub>4</sub> and 10 HEPES (pH 7.4). Ringer's solution was repeatedly added and removed to collect released <sup>86</sup>Rb<sup>+</sup> from cells at 2.5, 5.0, 7.5, 15, 25 and 40 min time points (*t*). Finally, cells were lysed by 2% SDS solution to collect the remaining <sup>86</sup>Rb<sup>+</sup> within the cells. Cumulative efflux (*J*) relative to total <sup>86</sup>Rb<sup>+</sup> counts was computed to determine macroscopic channel activity. The channel-specific rate constant (*k*<sub>2</sub>) of transfected channel proteins was determined by fitting the data using MATLAB (Mathworks, Natick, MA) with the following equation, as described before<sup>51</sup>,  $J = 1 - e^{-(k_1 + k_2)t}$ , where rate constant *k*<sub>1</sub>, obtained from cells transfected with GFP alone, reflects endogenous background channel activity, and *t* is time.

**Electrophysiology.** COSm6 cells between passages 10 and 30 were grown on glass cover slips in 40-mm six-well plates and transiently transfected after reaching  $\sim 30$ –50% confluency. Each well of cells was transfected with 200 ng of GFP and 800 ng of the corresponding Kir2.1 plasmid using Fugene6 and used for electrophysiological experiments 24–48 h thereafter. Current responses were recorded with electrodes pulled from plain glass capillaries with a resistance of 1–4 M $\Omega$  and were filled with a buffer (K<sub>int</sub>) containing (in mM) 140 KCl, 1 EGTA, 1 K<sub>2</sub>-EDTA, 4 K<sub>2</sub>HPO<sub>4</sub> and adjusted to pH 7.3 (KOH). Symmetrical K<sub>int</sub> solutions, also containing 10  $\mu$ M spermine (Fluka, St. Louis, MO), were applied with a lightly pressurized perfusion system. Current responses were first elicited in the cell-attached patch-clamp configuration with a holding potential of  $-50$  mV and pulsed for 300 ms to voltages between  $-100$  and  $+100$  mV at 10 mV increments. The patch was then excised to the inside-out patch-clamp configuration and moved between various solutions; the voltage step protocol was performed at regular intervals. Currents were low-pass filtered at 1 kHz (Axopatch 1D), sampled at 10 kHz (Digidata, 1200) and converted to digital files in Clampex7 (Molecular Devices) and stored on an external hard drive for offline analysis using Clampfit 10.3 (Molecular Devices).

**Homology modelling of human Kir2.1 channels.** Homology models of human Kir2.1 were built based on the chicken Kir2.2 (PDB 3SPI) crystal structure<sup>23</sup>. Human Kir2.1 is 76% and  $\sim 90$ % conserved relative to chicken Kir2.2. Sequence alignment was performed using the ClustalW webserver<sup>52</sup> and is shown in Supplementary Fig. S5. One hundred homology models were generated through random seeding using the MODELLER 9.10 program<sup>53</sup>. Because of high sequence homology, modelled structures had identical backbone structures but with various rotamer orientations. PDBQT structure files, containing charge and atom type information, were generated from the PDB file of every homology model beginning with protonation, assigning Gasteiger charge<sup>54</sup>, merging of nonpolar hydrogen to the bonded carbon atom, and assigning atom type using Autodock Tool (ADT) 1.5.4 (ref. 55), and then the protein was aligned to the initial crystal structure using MacPyMOL programs (Schrödinger, New York, NY) to ensure that the docking search area was consistent among models.

**Docking simulations.** The head groups are critical determinants for lipid interactions with the Kir protein, and to facilitate simulations, essentially only the head group was modelled, including up to the first carbon atom of the glycerol moiety. Atom charges for PI(4,5)P<sub>2</sub> were obtained as published<sup>56</sup> and the charges for other anionic lipids were taken from the CHARMM force field<sup>57</sup> shown in Supplementary Fig. S1. The atom charge of pyrophosphatidic acid (PPA) was kindly provided by Wonpil Im (University of Kansas). ADT 1.5.4 was used to assign the atom type and torsion tree of each ligand<sup>55</sup>.

The searched region fully covered the slide helix, the lower end of the TM1 helix, the N-terminal  $\beta$ -strand of subunit B (yellow) and the adjacent part of neighbouring subunits A (green) and C (silver), which was necessary for identification of potential interactions at the subunit interface as shown in Fig. 2b. The parameters for the grid map generation and docking simulations were the same as used previously<sup>26</sup> using AutoDock-4.2.3 (ref. 58). Each docking simulation was run for ~8 h on a single processor of the local IBM x3650-m<sup>2</sup>. For each template, 10,000 poses were generated for each ligand. Poses were accepted based on their orientation and relative location with respect to the potential membrane. First, a pose was accepted if the angle between the molecular vector that is designated for each lipid (Supplementary Fig. S1) and the bilayer normal was less than 90°, that is, as will occur when the head group is either pointing away from the membrane or less than parallel to the membrane surface. Structural analysis was performed using in-house codes for MATLAB. The potential membrane surface was approximated by the centre of the slide helix based on studies of the binding of melittin (an amphipathic  $\alpha$ -helix) to membranes<sup>59,60</sup>. If the primary phosphate that is directly connected to the lipid tails was located deeper than 4.5 Å into the membrane, or more than 8 Å from the surface of the membrane (defined by the centre of the slide helix), the pose was rejected.

To elucidate representative binding sites, accepted poses were clustered into subgroups. A K-mean clustering algorithm<sup>27</sup> was adopted using the root mean squared deviation of the phosphorus atom position as a metric. The number of clusters for each ligand was determined to minimize the performance index (Q)<sup>61</sup>. The smaller the Q value, the greater the inter-subgroup variance and the smaller the intra-subgroup variance, which indicates better clustering performance (see Supplementary Fig. S2 and text). To determine the residues involved in ligand binding, hydrogen bonds were postulated based on the two following criteria<sup>62</sup>: first, the distance between the donor and acceptor atoms being shorter than or equal to 3.4 Å and second, the angular orientation being smaller than or equal to 30° between the two unit vectors. One joins hydrogen bond donor atom and hydrogen atom and the other joins hydrogen bond donor and acceptor atoms. Contact frequency was determined by counting the number of observed hydrogen bonds and dividing this by the total number of poses in each cluster.

**Statistical analysis.** Statistical significance was analysed using an unpaired *t*-test and either one-way or two-way analysis of variance as appropriate. Statistical significance of  $P < 0.05$ ,  $P < 0.01$  and  $P < 0.001$  is indicated by single, double and triple asterisks, respectively.

## References

- Pattnaik, B., Asuma, M., Spott, R. & Pillers, D.-A. Genetic defects in the hotspot of inwardly rectifying K<sup>+</sup> (Kir) channels and their metabolic consequences: a review. *Mol. Genet. Metab.* **105**, 64–72 (2012).
- Nichols, C. KATP channels as molecular sensors of cellular metabolism. *Nature* **440**, 470–476 (2006).
- Lopatin, A. N. & Nichols, C. G. Inward rectifiers in the heart: an update on I(K1). *J. Mol. Cell Cardiol.* **33**, 625–638 (2001).
- Hibino, H. et al. Inwardly rectifying potassium channels: their structure, function, and physiological roles. *Physiol. Rev.* **90**, 291–366 (2009).
- Xie, L.-H., John, S., Ribalet, B. & Weiss, J. Activation of inwardly rectifying potassium (Kir) channels by phosphatidylinositol-4,5-bisphosphate (PIP<sub>2</sub>): Interaction with other regulatory ligands. *Prog. Biophys. Mol. Biol.* **94**, 320–335 (2007).
- Aryal, P., Dvir, H., Choe, S. & Slesinger, P. A. A discrete alcohol pocket involved in GIRK channel activation. *Nat. Neurosci.* **12**, 988–995 (2009).
- Huang, C. L., Feng, S. & Hilgemann, D. W. Direct activation of inward rectifier potassium channels by PIP<sub>2</sub> and its stabilization by G $\beta\gamma$ . *Nature* **391**, 803–806 (1998).
- Zhang, H., He, C., Yan, X., Mirshahi, T. & Logothetis, D. E. Activation of inwardly rectifying K<sup>+</sup> channels by distinct PtdIns(4,5)P<sub>2</sub> interactions. *Nat. Cell Biol.* **1**, 183–188 (1999).
- Suh, B.-C. & Hille, B. Regulation of ion channels by phosphatidylinositol 4,5-bisphosphate. *Curr. Opin. Neurobiol.* **15**, 370–378 (2005).
- Hilgemann, D. Local PIP<sub>2</sub> signals: when, where, and how? *Pflügers Archiv.* **455**, 55–67 (2007).
- Li, Y. et al. KCNE1 enhances phosphatidylinositol 4,5-bisphosphate (PIP<sub>2</sub>) sensitivity of IKs to modulate channel activity. *Proc. Natl Acad. Sci. USA* **108**, 9095–9100 (2011).
- Rapedius, M. et al. H bonding at the helix-bundle crossing controls gating in Kir potassium channels. *Neuron* **55**, 602–614 (2007).
- Dowhan, W. Molecular basis for membrane lipid diversity: why are there so many lipids? *Annu. Rev. Biochem.* **66**, 199–232 (1997).
- van Meer, G., Voelker, D. R. & Feigenson, G. W. Membrane lipids: where they are and how they behave. *Nat. Rev. Mol. Cell Biol.* **9**, 112–124 (2008).
- Rosenberger, T. A., Hovda, J. T. & Peters, J. M. Targeted disruption of peroxisomal proliferator-activated receptor beta ( $\delta$ ) results in distinct gender differences in mouse brain phospholipid and esterified FA levels. *Lipids* **37**, 495–500 (2002).
- Fuhrman, B. J. et al. Erythrocyte membrane phospholipid composition as a biomarker of dietary fat. *Ann. Nutr. Metab.* **50**, 95–102 (2006).
- Sanders, C. R. & Mittendorf, K. F. Tolerance to changes in membrane lipid composition as a selected trait of membrane proteins. *Biochemistry* **50**, 7858–7867 (2011).
- Farooqui, A. A., Wells, K. & Horrocks, L. A. Breakdown of membrane phospholipids in Alzheimer disease. Involvement of excitatory amino acid receptors. *Mol. Chem. Neurobiol.* **25**, 155–173 (1995).
- Kuliszkiewicz-Janus, M. et al. 31P MRS analysis of the phospholipid composition of the peripheral blood mononuclear cells (PBMC) and bone marrow mononuclear cells (BMMC) of patients with acute leukemia (AL). *Cell Mol. Biol. Lett.* **14**, 35–45 (2009).
- Nouri-Sorkhabi, M. H., Gruca, M. A., Kuchel, P. W. & Gaskin, K. J. Phospholipid changes in children with pancreatic sufficiency and insufficiency. *Clin. Chim. Acta* **281**, 89–100 (1999).
- Fan, Z. & Makielski, J. Anionic phospholipids activate ATP-sensitive potassium channels. *J. Biol. Chem.* **272**, 5388–5395 (1997).
- Cheng, W., D'Avanzo, N., Doyle, D. & Nichols, C. Dual-mode phospholipid regulation of human inward rectifying potassium channels. *Biophys. J.* **100**, 620–628 (2011).
- Hansen, S., Tao, X. & MacKinnon, R. Structural basis of PIP<sub>2</sub> activation of the classical inward rectifier K<sup>+</sup> channel Kir2.2. *Nature* **477**, 495–498 (2011).
- Whorton, M. R. & MacKinnon, R. Crystal structure of the mammalian GIRK2 K<sup>+</sup> channel and gating regulation by G proteins, PIP<sub>2</sub>, and sodium. *Cell* **147**, 199–208 (2011).
- D'Avanzo, N., Cheng, W., Doyle, D. & Nichols, C. Direct and specific activation of human inward rectifier K<sup>+</sup> channels by membrane phosphatidylinositol 4,5-bisphosphate. *J. Biol. Chem.* **285**, 37129–37132 (2011).
- D'Avanzo, N., Lee, S. J., Cheng, W. W. & Nichols, C. G. Energetics and location of phosphoinositide binding in human kir2.1 channels. *J. Biol. Chem.* **288**, 16726–16737 (2013).
- Bagga, S. & Singh, G. N. Clustering method for categorical and numeric data sets. *GJCT* **6**, 49–52 (2011).
- Lopatin, A. N., Makhina, E. N. & Nichols, C. G. Potassium channel block by cytoplasmic polyamines as the mechanism of intrinsic rectification. *Nature* **372**, 366–369 (1994).
- Sies, H., Wahlländer, A., Waydhas, C., Soboll, S. & Häberle, D. Functions of intracellular glutathione in hepatic hydroperoxide and drug metabolism and the role of extracellular glutathione. *Adv. Enzyme Regul.* **18**, 303–320 (1980).
- Lopes, C. M. B. et al. Alterations in conserved Kir channel-PIP<sub>2</sub> interactions underlie channelopathies. *Neuron* **34**, 933–944 (2002).
- Rohács, T., Chen, J., Prestwich, G. D. & Logothetis, D. E. Distinct specificities of inwardly rectifying K<sup>+</sup> channels for phosphoinositides. *J. Biol. Chem.* **274**, 36065–36072 (1999).
- Hardie, R. C. & Muallem, S. Lipids in Ca<sup>2+</sup> signalling—an introduction. *Cell Calcium* **45**, 517–520 (2009).
- Ho, I. H. M. & Murrell-Lagnado, R. D. Molecular determinants for sodium-dependent activation of G protein-gated K<sup>+</sup> channels. *J. Biol. Chem.* **274**, 8639–8648 (1999).
- Romanenko, V. G. et al. Cholesterol sensitivity and lipid raft targeting of Kir2.1 channels. *Biophys. J.* **87**, 3850–3861 (2004).
- Epshtein, Y. et al. Identification of a C-terminus domain critical for the sensitivity of Kir2.1 to cholesterol. *Proc. Natl Acad. Sci. USA* **106**, 8055–8060 (2009).
- D'Avanzo, N., Hyrc, K., Enkvetchakul, D., Covey, D. F. & Nichols, C. G. Enantioselective protein-sterol interactions mediate regulation of both prokaryotic and eukaryotic inward rectifier K<sup>+</sup> channels by cholesterol. *PLoS One* **6**, e19393 (2011).
- Yokogawa, M., Osawa, M., Takeuchi, K., Mase, Y. & Shimada, I. NMR analyses of the G $\beta\gamma$  binding and conformational rearrangements of the cytoplasmic pore of G protein-activated inwardly rectifying potassium channel 1 (GIRK1). *J. Biol. Chem.* **286**, 9 (2011).
- Mase, Y., Yokogawa, M., Osawa, M. & Shimada, I. Structural basis for modulation of gating property of G protein-gated inwardly rectifying potassium ion channel (GIRK) by *i/o*-family G protein  $\alpha$  subunit (G $\alpha$ /o). *J. Biol. Chem.* **287**, 13 (2012).
- Logothetis, D. E., Lupyan, D. & Rosenhouse-Dantsker, A. Diverse Kir modulators act in close proximity to residues implicated in phosphoinositide binding. *J. Physiol.* **582**, 953–965 (2007).

40. Xie, L.-H., John, S., Ribalet, B. & Weiss, J. Activation of inwardly rectifying potassium (Kir) channels by phosphatidylinositol-4,5-bisphosphate (PIP<sub>2</sub>): interaction with other regulatory ligands. *Prog. Biophys. Mol. Biol.* **94**, 320–335 (2007).
41. D'Avanzo, N., Cheng, W. W. L., Wang, S., Enkvetchakul, D. & Nichols, C. G. Lipids driving protein structure? Evolutionary adaptations in Kir channels. *Channels* **4**, 139–141 (2010).
42. Enkvetchakul, D. & Nichols, C. G. Gating mechanism of KATP channels: function fits form. *J. Gen. Physiol.* **122**, 471–480 (2003).
43. Tao, X., Avalos, J. L., Chen, J. & MacKinnon, R. Crystal structure of the eukaryotic strong inward-rectifier K<sup>+</sup> channel Kir2.2 at 3.1 Å resolution. *Science* **326**, 1668–1674 (2009).
44. Whorton, M. R. & MacKinnon, R. X-ray structure of the mammalian GIRK2-beta-gamma G-protein complex. *Nature* **498**, 190–197 (2013).
45. Wang, S., Lee, S.-J., Heyman, S., Enkvetchakul, D. & Nichols, C. Structural rearrangements underlying ligand-gating in Kir channels. *Nat. Comm.* **3**, 617 (2012).
46. Enkvetchakul, D., Jeliakova, I., Bhattacharyya, J. & Nichols, C. Control of inward rectifier K channel activity by lipid tethering of cytoplasmic domains. *J. Gen. Physiol.* **130**, 329–334 (2007).
47. Iwamoto, M. & Oiki, S. Amphipathic antenna of an inward rectifier K<sup>+</sup> channel responds to changes in the inner membrane leaflet. *Proc. Natl Acad. Sci. USA* **110**, 6 (2013).
48. Pessia, M., Tucker, S. J., Lee, K., Bond, C. T. & Adelman, J. P. Subunit positional effects revealed by novel heteromeric inwardly rectifying K<sup>+</sup> channels. *EMBO J.* **15**, 2980–2987 (1996).
49. D'Avanzo, N. *et al.* Expression and purification of recombinant human inward rectifier K<sup>+</sup> (KCNJ) channels in *Saccharomyces cerevisiae*. *Prot. Expr. Purif.* **71**, 115–121 (2010).
50. Cho, H. C., Tsushima, R. G., Nguyen, T. T., Guy, H. R. & Backx, P. H. Two critical cysteine residues implicated in disulfide bond formation and proper folding of Kir2.1. *Biochemistry* **39**, 4649–4657 (2000).
51. Sala-Rabanal, M., Kucheryavykh, L., Skatchkov, S., Eaton, M. & Nichols, C. Molecular mechanisms of EAST/SeSAME syndrome mutations in Kir4.1 (KCNJ10). *J. Biol. Chem.* **285**, 36040–36048 (2010).
52. Larkin, M. A. *et al.* Clustal W and Clustal X version 2.0. *Bioinformatics* **23**, 2947–2948 (2007).
53. Sali, A. & Blundell, T. L. *Comparative Protein Modelling by Satisfaction of Spatial Restraints*. C86 (IOS Press, Amsterdam, 1994).
54. Huey, R., Morris, G. M., Olson, A. J. & Goodsell, D. S. A semiempirical free energy force field with charge-based desolvation. *J. Comput. Chem.* **28**, 1145–1152 (2007).
55. Morris, G. *et al.* AutoDock4 and AutoDockTools4: automated docking with selective receptor flexibility. *J. Comput. Chem.* **30**, 2785–2791 (2009).
56. Lupyan, D., Mezei, M., Logothetis, D. E. & Osman, R. A molecular dynamics investigation of lipid bilayer perturbation by PIP<sub>2</sub>. *Biophys. J.* **98**, 240–247 (2010).
57. Feller, S. & MacKerell, A. An improved empirical potential energy function for molecular simulations of phospholipids. *J. Phys. Chem. B* **104**, 7510–7515 (2000).
58. Morris, G. *et al.* AutoDock4 and AutoDockTools4: automated docking with selective receptor flexibility. *J. Comput. Chem.* **30**, 2785–2791 (2009).
59. Hristova, K., Dempsey, C. E. & White, S. H. Structure, location, and lipid perturbations of melittin at the membrane interface. *Biophys. J.* **80**, 801–811 (2001).
60. Lee, S.-J., Schlesinger, P. H., Wickline, S. A., Lanza, G. M. & Baker, N. A. Interaction of melittin peptides with perfluorocarbon nanoemulsion particles. *J. Phys. Chem. B* **115**, 15271–15279 (2011).
61. Savaresi, S. M., Boley, D. L., Bittanti, S. & Gazzaniga, G. Cluster selection in divisive clustering algorithms. *Proc. 2nd SIAM* **1401**, 299–314 (2002).
62. Bradley, M. J., Chivers, P. T. & Baker, N. A. Molecular dynamics simulation of the *Escherichia coli* NikR protein: equilibrium conformational fluctuations reveal interdomain allosteric communication pathways. *J. Mol. Biol.* **378**, 1155–1173 (2008).

### Acknowledgements

We thank Dr Wonpil Im for PPA charge information developed by his group to be compatible with the CHARMM force field parameters, and Dr Malcolm Tobias and the Center for High Performance Computing in Washington University, School of Medicine for computing resources. Financial support was provided by NIH Grant HL54171 (to C.G.N.), and AHA Fellowship 13POST14660069 (to S.-J.L.).

### Author contributions

S.-J.L. and C.G.N. conceived the project. S.-J.L. carried out computational modelling. S.-J.L., S.W. and W.B. carried out experiments and data analysis. C.G.N. and S.-J.L. wrote the paper.

### Additional information

**Supplementary Information** accompanies this paper at <http://www.nature.com/naturecommunications>

**Competing financial interests:** The authors declare no competing financial interests.

**Reprints and permission** information is available online at <http://npg.nature.com/reprintsandpermissions/>

**How to cite this article:** Lee, S.-J. *et al.* Secondary anionic phospholipid binding site and gating mechanism in Kir2.1 inward rectifier channels. *Nat. Commun.* **4**:2786 doi: 10.1038/ncomms3786 (2013).



This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivs 3.0 Unported License. To view a copy of this license, visit <http://creativecommons.org/licenses/by-nc-nd/3.0/>