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Mining of Ebola virus entry inhibitors identifies approved drugs as two-pore channel pore blockers[☆]

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ABSTRACT

Two-pore channels (TPCs) are Ca²⁺-permeable ion channels localised to the endo-lysosomal system where they regulate trafficking of various cargoes including viruses. As a result, TPCs are emerging as important drug targets. However, their pharmacology is ill-defined. There are no approved drugs to target them. And their mechanism of ligand activation is largely unknown. Here, we identify a number of FDA-approved drugs as TPC pore blockers. Using a model of the pore of human TPC2 based on recent structures of mammalian TPCs, we virtually screened a database of ~1500 approved drugs. Because TPCs have recently emerged as novel host factors for Ebola virus entry, we reasoned that Ebola virus entry inhibitors may exert their effects through inhibition of TPCs. Cross-referencing hits from the TPC virtual screen with two recent high throughput anti-Ebola screens yielded approved drugs targeting dopamine and estrogen receptors as common hits. These compounds inhibited endogenous NAADP-evoked Ca²⁺ release from sea urchin egg homogenates, NAADP-mediated channel activity of TPC2 re-routed to the plasma membrane, and PI(3,5)P₂-mediated channel activity of TPC2 expressed in enlarged lysosomes. Mechanistically, single channel analyses showed that the drugs reduced mean open time consistent with a direct action on the pore. Functionally, drug potency in blocking TPC2 activity correlated with inhibition of Ebola virus-like particle entry. Our results expand TPC pharmacology through the identification of approved drugs as novel blockers, support a role for TPCs in Ebola virus entry, and provide insight into the mechanisms underlying channel regulation. This article is part of a Special Issue entitled: ECS Meeting edited by Claus Heizmann, Joachim Krebs and Jacques Haiech.

1. Introduction

NAADP is a potent Ca²⁺ mobilizing messenger that triggers Ca²⁺ release from acidic Ca²⁺ stores such as lysosomes [1–5]. Critical to NAADP action are the endo-lysosomal two-pore channels (TPCs) [6–9]. Local Ca²⁺ fluxes through TPCs are increasingly implicated in membrane trafficking events [10,11]. These include roles for TPCs in regulating lysosome morphology [12], retrograde transport between late endosomes and the Golgi [13], and endocytic trafficking of receptors for LDL and growth factors [14,15], pigment granules [16] and integrins [17]. They also likely regulate non-vesicular trafficking by strengthening membrane contact sites between late endosomes and the

ER [18]. Consequently, TPCs are fast emerging as potential therapeutic targets in diverse disorders such as Parkinson's disease, fatty liver disease and cancer [12,14,17,19]. Moreover, TPCs have been identified as novel host factors for Ebola virus (EBOV) entry [20]. This virus enters cells through macropinocytosis, traffics through the endo-lysosomal system and, following binding to its intracellular receptor NPC1 [21], fuses with late endosomes/lysosomes to release its genome into the cytoplasm. Molecular or chemical inhibition of TPCs prevents EBOV infection likely at a late entry step, although the mechanism is unclear [20,22].

TPCs are dimeric ion channels, comprising a duplicated domain architecture, which are likely a bridge during the evolution of four-

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domain voltage-gated Ca^{2+} and Na^{+} channels [23–25]. Atomic structures of plant [26–28] and mammalian [29,30] channels are emerging providing detailed insight into the architecture of the pore. Two isoforms (TPC1 and TPC2) found in humans localise to distinct compartments within the endo-lysosomal system [9,31]. TPC2 is predominantly localised to late endosomes and lysosomes through an N-terminal targeting motif, which when mutated results in rerouting of the channel to the plasma membrane [32]. The activation mechanisms and ion selectivities of TPCs are areas of active investigation [33,34]. Although many studies have shown TPCs are required for NAADP-mediated Ca^{2+} signalling, NAADP appears to interact with TPCs indirectly through putative binding proteins [35–38]. In contrast, the endo-lysosomal lipid PI(3,5)P₂ has emerged as a direct channel activator that binds within the first domain of TPCs [29,30,39]. Interestingly, residues in the S4–S5 linker within this domain required for NAADP-mediated Ca^{2+} signals [40] fall within the PI(3,5)P₂ binding site [29] indicating a complex interplay between small molecule activators and accessory proteins.

Despite the growing pathophysiological importance of TPCs, there are currently few selective chemical tools to inhibit NAADP/TPC signalling. BZ194 [41], a N-alkylated nicotinic acid derivative, and Ned-19 [42], identified through a ligand-based virtual screen, are cell-permeable inhibitors of NAADP-evoked Ca^{2+} signals. But their indirect action, coupled with potentially promiscuous actions of their target NAADP-binding proteins on other channels [43], complicates the use of NAADP antagonists in a physiological setting. This underscores the need for direct channel inhibitors. We recently used a combination of homology modelling, molecular docking analyses and Ca^{2+} measurements to show that TPCs are likely to be direct targets for a number of voltage-gated Ca^{2+} and Na^{+} channel modifiers [24]. Tetrandrine [20], another Ca^{2+} channel antagonist has also recently emerged as a TPC blocker and potent inhibitor of EBOV entry. Finally, the flavonoid naringenin [44] has been shown to block TPCs, but with low affinity. Both tetrandrine and naringenin have multiple targets and are not approved for human use world-wide. There are therefore a limited number of drugs that directly inhibit TPCs.

Here, we apply a combination of structural modelling, virtual screening, Ca^{2+} measurements, electrophysiology and virus entry assays to identify approved drugs that both inhibit EBOV entry and block TPC activity.

2. Methods

2.1. Virtual screening

The structure of human TPC2 (Uniprot accession: Q8NHX9) in an open conformation was modelled on the cryoEM structure of PI(3,5)P₂-bound mouse TPC1 (pdb: 6C9A) [29] using SWISS-MODEL [45]. The first 40 residues were excluded due to lack of appropriate template. The final model was chosen using the default local model quality estimation method (QMEANDisCo) based on a novel version of QMEAN [45]. Additional model refinement was performed using GalaxyRefine [46].

A conformer library for the e-Drugs3D-2017 drug set [47] was prepared using Omega v3.0.1.2 (OpenEye Scientific Software, Santa Fe, NM). MakeReceptor (OEDOCKING 3.2.0.2; OpenEye Scientific Software, Santa Fe, NM) was used to generate the active site for virtual screening. This was centred on the pore vestibule and extended from above the selectivity filter to below the bundle crossing ($40 \times 40 \times 77 \mu\text{m}$). The starting pose was based on blind docking of verapamil using AutoDock Vina [48] with an exhaustiveness value of 24. Screening was carried out with FRED (OEDOCKING 3.2.0.2; OpenEye Scientific Software, Santa Fe, NM) using default parameters returning the top 10,000 conformers ranked by the Chemgauss4 score (Table S1). All but the top ranked conformer for each ligand were excluded. This analysis retrieved 1480 unique drugs and their metabolites. The top 200 are listed in Table S1.

Enrichment analysis of the top hits was performed by dividing the

proportion of a given drug class within the top 200 hits by the proportion of that class in the screen as a whole. Drugs were classified according to their annotated primary target in e-Drugs3D. This analysis was limited to classes where there were > 3 drugs. A class was considered enriched if values were > 1.5.

2.2. Ca^{2+} measurements

Ca^{2+} release assays using sea urchin egg homogenates were performed as described previously [49]. Briefly, egg homogenates were loaded with Ca^{2+} by sequential dilution in a cytosol-like medium supplemented with an ATP-regenerating system. Medium Ca^{2+} concentration was monitored fluorimetrically with Fluo-4.

2.3. Single channel current recording

Patch clamp recording in the excised inside-out configuration was performed as described in [32] using HEK cells expressing human TPC2 tagged with GFP at its C-terminus and where leucines 11 and 12 were replaced with alanine to promote cell surface targeting. The pipette (luminal) solution contained (in mM): 145 NaCl, 5 KCl, 2 CaCl_2 , 1 MgCl_2 , 10 HEPES and 10 MES (pH adjusted to 4.6 with NaOH). The bath (cytoplasmic) solution had the same composition except MES was excluded and pH was adjusted to 7.2 with NaOH. Osmolarity of the solutions was adjusted to ~300 mOsm with glucose.

The methods used for recording and analysing currents were reported previously [50]. Pipettes were coated with Sigmacote™, fire polished and minimally filled with the pipette solution. Currents were amplified using an Axopatch 200B amplifier, filtered at 1 kHz (–3 dB) with an internal 4-pole Bessel filter and digitized at 10 kHz with a Digidata 1322A interface and pClamp 10 suite. Unless otherwise stated, all recordings were at a holding potential of +40 mV.

For determining unitary current amplitude and unitary conductance, currents were analysed using the 50% threshold crossing criterion using ClampFit 10 and openings briefer than 700 μs (twice the filter rise time) were excluded from the analysis [51]. Due to uncertainty about the number of channels in most recordings, channel activity was expressed as NP_o .

For kinetic analyses of single channel records, a simple two state scheme ($C \rightleftharpoons O$) was initially used to idealize currents at full bandwidth (1 kHz) using the segmental k-means (SKM) hidden-Markov algorithm implemented in QuB [52]. Additional states were then systematically added to the initial scheme with various topologies using the MIL module of QuB. A dead time of 200 μs was retrospectively imposed to correct for missed events [51]. MIL fits a gating scheme to the single channel records and optimises the rate constants based on a maximum likelihood-based Markovian approach. The final gating schemes were derived when further changes in either the number of states or their connectivity did not produce significant (≥ 10) increment in the log likelihood (LL) ratio [50,52]. Closed time distributions were analysed when channel activity was reasonably high ($NP_o > 0.15$) and when only openings to a single current level were evident over a duration of ≥ 5 min [53].

Analyses were performed 10–15 s following NAADP/drug application in the bath solution to ensure adequate mixing.

2.4. Whole lysosomal current recording

Lysosomal currents were recorded using a modified patch clamp procedure as described in [54] and COS-7 cells expressing human TPC2 tagged with GFP at its C-terminus [7]. Briefly, cells were treated with 1 μM vacuolin-1, for 1–2 h and the resulting enlarged lysosomes accessed by breaking the plasma membrane with a glass pipette. The pipette (luminal) solution and bath (cytoplasmic) solution contained (in mM): 140 NaCl, 5 KCl, 10 HEPES (pH 7.4 with NaOH). After formation of a G Ω seal with a lysosome, break in was achieved by quick voltage

steps of +350 mV. Current was measured with an Axopatch 200B patch clamp amplifier that was controlled by pCLAMP9 software. The current protocol used 400-ms rapid alterations of membrane potential (RAMP) from -100 to $+100$ mV from a holding potential of 0 mV at 4 s intervals. The currents were filtered at 1 kHz with an internal four-pole Bessel filter, sampled at 5 kHz, and stored to a hard drive by Digidata 1440.

2.5. Ebola virus-like-particle entry assays

Virus-like particles (VLPs) for EBOV were generated as described in [55]. Briefly, HEK293T cells were co-transfected with constructs encoding the EBOV matrix protein, VP40, tagged with a β -lactamase reporter at its N-terminus [56] and the EBOV glycoprotein (GP; strain Zaire) [57], both a kind gift from Adolfo Garcia-Sastre. Cells were maintained in DMEM supplemented with 10% (v/v) FBS, 100 μ g/mL streptomycin and 100 units/mL penicillin at 37 °C in a humidified atmosphere containing 5% CO₂. Culture media were harvested after 48 h, cleared by centrifugation and VLP-containing supernatants stored at -80 °C until use.

The VLP entry assay [55] was adapted and optimised to a 96-well plate format. HeLa Kyoto cells were seeded at a density of 8×10^3 cells/well and maintained for 40 h prior to drug treatment in DMEM supplemented with 10% (v/v) FBS, 100 μ g/mL streptomycin and 100 units/mL penicillin at 37 °C in a humidified atmosphere containing 5% CO₂. Cells were incubated with drugs in DMEM + 2% (v/v) FBS for 1 h under culture conditions prior to the addition of VLPs (1:2 dilution). Cells were spinoculated at 450 \times g for 1 h at 4 °C, followed by incubation for 3 h under culture conditions to allow VLP entry in the continued presence of drugs. Cells were washed once with CO₂-independent medium supplemented with 10% (v/v) FBS and 1% (v/v) L-glutamine and loaded with the β -lactamase substrate, CCF2 (Live-Blazer FRET-B/G Loading Kit, Invitrogen) according to the manufacturer's instructions using the alternative substrate loading protocol. Cells were trypsinised, fixed with 1% paraformaldehyde for 20 min and analysed by flow cytometry using a LSRII flow cytometer (Becton Dickinson, USA) collecting at least 10,000 events. β -lactamase activity was assessed in live cells with the FlowJo software package by determining the proportion of blue fluorescent cells (cleaved substrate; excitation 405 nm) relative to green fluorescent cells (total substrate; excitation 488 nm) using filter sets for Pacific-blue and FITC, respectively.

2.6. Data analysis

Data are presented as mean values \pm standard error of the mean where indicated. Concentrations that caused 50% inhibition (IC₅₀) and Hill coefficients (n_H) were derived from fitting of inhibition curves using KaleidaGraph (Synergy software).

3. Results

3.1. Identification of putative TPC2 inhibitors through a structure-based virtual screen of approved drugs based on the TPC2 pore

Recent advances in the structural biology of TPCs [28,30] provide novel opportunities to gain insight into TPC pharmacology. We thus applied a structure-based virtual screening approach to identify novel lead compounds targeting the TPC pore. We took advantage of the recent cryo-EM structure of a mammalian TPC [29] to model the pore of human TPC2 (Fig. 1A) and used this model to perform an *in silico* screen of the e-Drugs3D database, which consists of \sim 1500 drugs approved by the FDA for use in humans. The results of this screen are depicted schematically in Fig. 1B where each compound is ranked according to its Chemguass4 score. This score is a measure of the predicted strength of interaction whereby more negative values indicate a stronger

predicted interaction.

Our previous targeted docking analyses showed that voltage-gated Ca²⁺ and Na⁺ channel blockers likely interact with the TPC pore through a common 'ancestral' binding site [24]. Of the Ca²⁺ channel blockers analysed previously, nifedipine was predicted to interact most strongly with the pore and diltiazem was the most potent functionally [24]. Accordingly, nifedipine and diltiazem were recovered as top ranking blockers in the present screen (Fig. 1C, asterisk). Additionally, the mean rank and mean Chemguass4 scores of all Ca²⁺ channel blockers in the screen were lower than for Na⁺ channel blockers (Fig. 1C–D). This accords with our docking and functional analyses showing more avid interaction and more potent inhibition of TPCs by blockers of Ca²⁺ relative to Na⁺ channels [24]. Taken together, these analyses support our unbiased approach for identification of TPC inhibitors.

Subsequent efforts were focused on the top 200 hits recovered in the virtual screen. We performed enrichment analysis according to the primary target where annotated. As shown in Fig. 1D, this analysis revealed a number of drugs targeting different G-protein coupled receptors, HMG-CoA reductase, the estrogen receptor and the angiotensin II converting enzyme. This unbiased approach thus returns a number of drugs within distinct drug classes that potentially target the TPC2 pore.

3.2. Prioritisation of approved drugs that inhibit Ebola virus entry as putative TPC2 inhibitors

Because TPCs have emerged as novel host factors for EBOV infection [20], we reasoned that some of the FDA-approved drugs recently highlighted in high-throughput screens as EBOV entry inhibitors might function as TPC blockers. We thus compared our top 200 hits with two recent anti-EBOV screens [58,59] that together identified 73 unique EBOV entry inhibitors (Fig. 2A). Of these, 51 were present in e-Drugs3D and all but 5 were recovered in the TPC virtual screen (Fig. 2B). Notably, approximately, a third of these (14) ranked within the top 200 of the TPC screen (Fig. 2C). Inspection of the prioritised hit compounds (Fig. 2D) revealed the Ca²⁺ channel blocker bepridil, further underscoring this class of compounds as TPC blockers. Importantly, this analysis also returned 4 dopamine receptor antagonists and 5 selective estrogen receptor modulators (SERMs). The prevalence of drugs in these two classes is consistent with our enrichment analysis from the TPC screen (Fig. 1D). Thus, virtual screening against the TPC2 pore and physical screening for EBOV entry inhibitors converge on dopamine receptor antagonists and SERMs as candidate TPC inhibitors.

3.3. Dopamine antagonists and selective estrogen receptor modulators block endogenous NAADP-mediated Ca²⁺ signals

To examine the functionality of the drugs as TPC inhibitors, we first tested their effects on Ca²⁺ signals evoked by NAADP. We used sea urchin egg homogenates for this analysis as this preparation is considered the 'gold standard' for assessing NAADP-mediated Ca²⁺ signalling [60]. As shown in Fig. 3A, NAADP mediated a robust Ca²⁺ signal that was largely blocked by bepridil (100 μ M). Similar results were obtained with fluphenazine and pimozide, both dopamine receptor antagonists (Fig. 3A). We also tested the effects of the SERMs, raloxifene and clomiphene on NAADP responsiveness. Raloxifene almost completely blocked NAADP-evoked Ca²⁺ release whereas clomiphene effected a partial block (Fig. 3A).

The inhibitory effects of all these drugs were concentration dependent (Fig. 3B) with the following IC₅₀ and n_H values: bepridil (33 μ M, 1.5), fluphenazine (42 μ M, 1.3), pimozide (42 μ M, 0.96), raloxifene (28 μ M, 1.6) and clomiphene (94 μ M, 1.7). To determine selectivity, we examined the effects of the drugs on Ca²⁺ release by cyclic ADP-ribose, which mediates its effects through the ryanodine receptor. As summarized in Fig. 3C, all drugs were less effective at inhibiting responses to cyclic ADP-ribose than NAADP, attesting to specificity.

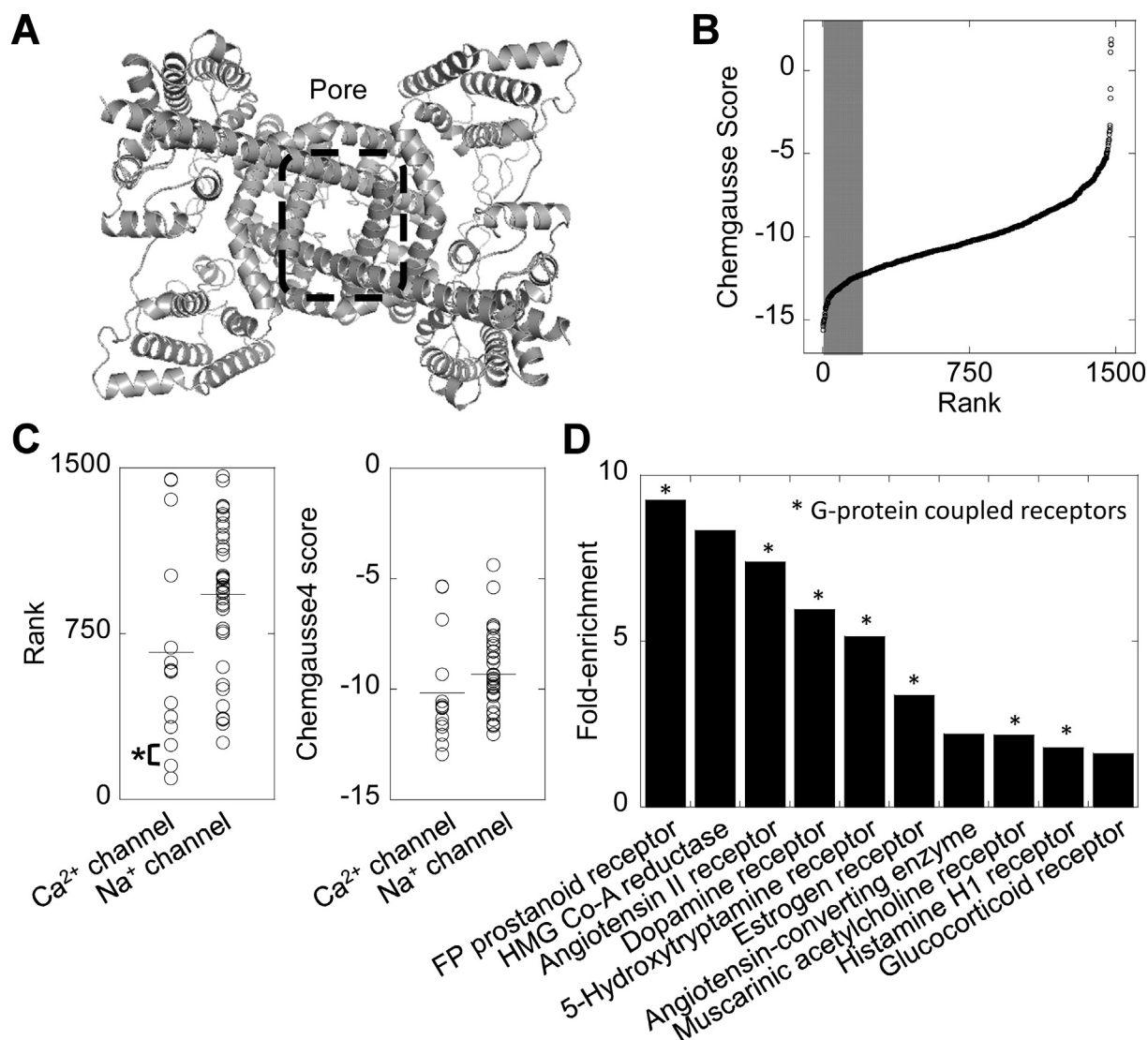


Fig. 1. A structure-based virtual screen of approved drugs based on the TPC2 pore.

A, structural model of human TPC2 viewed from the cytosolic face. The pore region is highlighted by the dashed box.

B, results of a virtual screen of the e-Drugs3D database against the pore region of TPC2. The top 200 drugs are marked by the shaded region.

C, plots showing the rank of individual drugs from the virtual screen targeting voltage-gated Ca^{2+} and Na^{+} channels. The horizontal lines represent the mean. Nicardipine (rank #154) and diltiazem (rank #247) are highlighted*.

D, enrichment analysis of drug classes within the top 200 hits. G-protein coupled receptors are highlighted*.

Taken together, these data identify dopamine receptor antagonists and SERMs as novel inhibitors of NAADP action consistent with their predicted interaction with TPCs.

3.4. Dopamine antagonists and selective estrogen receptor modulators inhibit single channel activity of TPC2 by reducing open time

To provide more direct evidence that these compounds are TPC blockers and to extend our analyses to mammalian systems, we analysed NAADP-mediated channel activity of human TPC2. We did this using cell surface targeted TPC2 to facilitate single channel recording in a null background. Na^{+} was used as the major cation to complement our previous analyses using Ca^{2+} as the permeant ion [32]. As shown in Fig. 4A, some spontaneous activity was noted in most of the excised patches. NAADP addition to the bath solution robustly enhanced channel activity in a concentration dependent manner. Current amplitude histograms are shown in Fig. S1A–C. Summary data quantifying the normalised open probability NP_o is shown in Fig. 4B. Analysis of current-voltage relationships indicated a unitary Na^{+} conductance

(γ_{Na}) of 86 ± 0.7 pS (Fig. 4C; $n = 3$). Single channel openings in the absence and presence of NAADP were best fit by a single exponential with a mean open time of ~ 5 ms (Fig. 4D). For a sub-set of recordings where we were confident that the openings represented a single active channel ($P_o \geq 0.15$, ≥ 5 min recording duration) [50], closed time distribution was also analysed (Fig. 4D). The distribution was best fit by a double exponential with time constants of 0.7 ± 0.4 ms and 47 ± 3.5 ms, constituting $49 \pm 1\%$ and $51 \pm 1\%$ of the distribution, respectively ($n = 4$). Kinetic modelling yielded two topologies with similar LL ratios that linearly connected the closed states (C_1 and C_2) with the open state (O) (Fig. 4F). Collectively, these data affirm the sensitivity of TPC2 to NAADP and its permeability to Na^{+} and provide a framework for assessing drug action at the single channel level.

We used this experimental setting to assess the effects of fluphenazine and raloxifene (representative of the two identified drug classes) on TPC2 channel activity. Fig. 5 shows continuous uninterrupted recordings of TPC2 channel activity in response to NAADP and subsequent addition of the drugs. Both drugs acutely inhibited NAADP-mediated channel activity (Fig. 5A–B). We also tested the effects of the

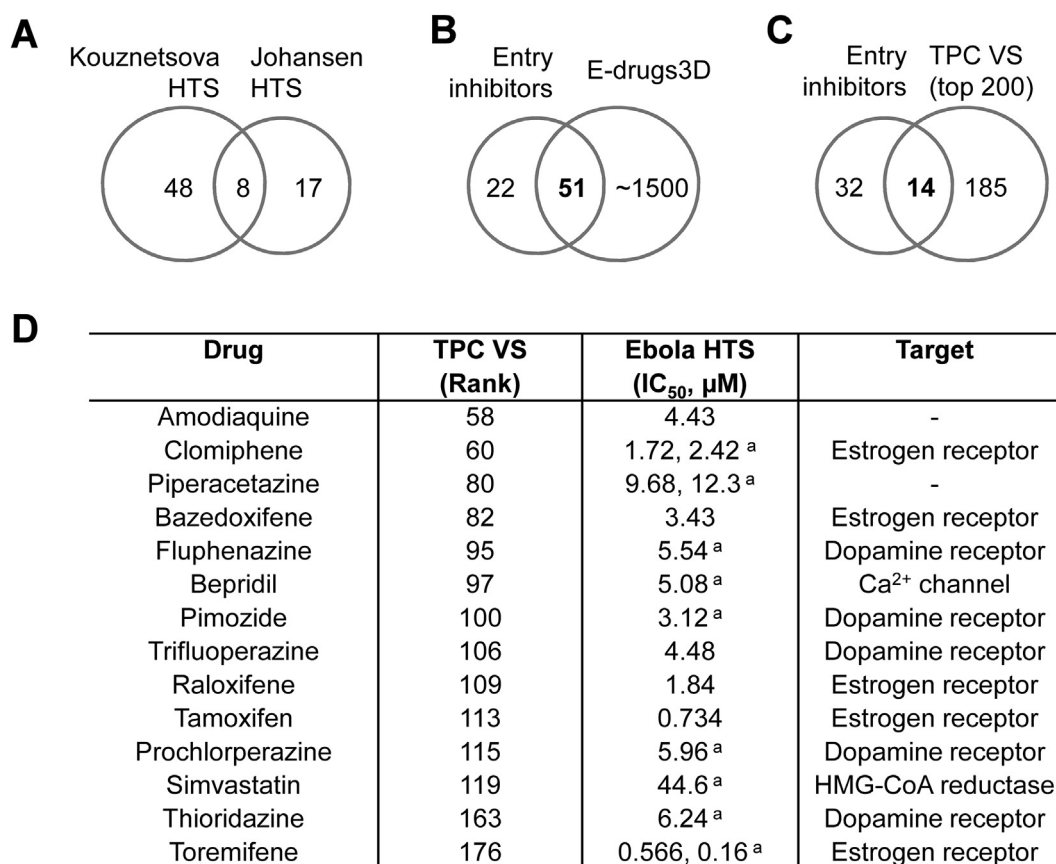


Fig. 2. Identification of drugs targeting dopamine and estrogen receptors as putative TPC2 inhibitors.

A, Venn diagram depicting the relationship between approved drugs identified as Ebola virus entry inhibitors in independent high through put screens by Kouznetsova [58] and Johansen et al. [59].

B, Venn diagram depicting the relationships between the entry inhibitors identified in A with the e-Drugs3D database used for the TPC virtual screen.

C, Venn diagram depicting the relationship between entry inhibitors present in the e-Drugs3-D database and the top 200 hits from the TPC virtual screen.

D, prioritised set of 14 drugs identified from C listing their rank in the TPC virtual screen, their reported IC₅₀ values for inhibiting virus entry (from Kouznetsova [58] and/or ^aJohansen et al. [59]) and their primary target (annotated in e-Drugs3D).

NAADP antagonist Ned-19. As shown in Fig. 5C, Ned-19 also similarly inhibited channel activity. To gain mechanistic insight into drug block, we analysed dwell time distributions. As shown in Fig. 5D–E, both fluphenazine and raloxifene reduced the mean open time from ~5 ms to ~3 ms without affecting the unitary current sizes. Current amplitude histograms are shown in Fig. S1D–F. These data are consistent with a direct interaction of the drugs with the pore. In contrast, Ned-19 affected neither the mean open time nor unitary current size (Fig. 5F) suggesting a distinct mechanism of action. Data summarizing the effects of the drugs on NP_{o} , open times and unitary conductance are shown in Fig. 5G. Taken together, these data suggest that fluphenazine and raloxifene effectively behave as pore blockers.

3.5. Dopamine antagonists and selective estrogen receptor modulators inhibit lysosomal TPC2 currents and entry of Ebola virus-like particles with similar potency

Because TPC2 in the plasma membrane may not faithfully reflect the properties of the channel in its native environment, we performed electrophysiological analysis of enlarged lysosomes expressing wildtype TPC2 (Fig. 6). For these experiments, TPC2 was stimulated with a saturating concentration of PI(3,5)P₂, which gives larger currents than NAADP in this experimental setting [54] most likely due to a direct interaction with the channel [61]. As shown in Fig. 6A, PI(3,5)P₂ stimulated robust currents in TPC2-expressing lysosomes. These responses were inhibited by fluphenazine in a concentration-dependent manner

with near full block achieved at 30 μM (Fig. 6A). Raloxifene also inhibited PI(3,5)P₂-mediated currents with maximal effects at 3 μM (Fig. 6A). The inhibitory effects of both drugs were reversed upon washout (Fig. 6A). Concentration-effect relationships showed that raloxifene was particularly potent at blocking TPC2 currents (IC₅₀ = 0.63 μM, n_H = 1.3) in this setting compared to fluphenazine (IC₅₀ = 8.2 μM, n_H = 1.3), respectively (Fig. 6B).

Finally, we examined the potency of dopamine receptor antagonists and SERMs on EBOV entry. As a surrogate for EBOV, we used Ebola VLPs carrying β-lactamase. Following fusion mediated by the EBOV glycoprotein, β-lactamase is delivered to the target cell cytoplasm where it mediates cleavage of a fluorescent substrate. Fig. 7A quantifies infection of HeLa cells by the VLPs. Infection was blocked by pre-treatment with EIPA (11.1 μM), an inhibitor of macropinocytosis, and by tetrandrine (1.2 μM; Fig. 7A). As shown in Fig. 7B, fluphenazine and pimozide inhibited Ebola VLP entry with IC₅₀ values of 1.3 μM (n_H = 1.7) and 3.6 μM (n_H = 2.8), respectively. Raloxifene, clomiphene and a third SERM, tamoxifen, also inhibited entry with IC₅₀ values of 0.61 μM (n_H = 1.1), 0.32 μM (n_H = 1.1) and 0.21 μM (n_H = 0.82), respectively. Enhanced potency of SERMs relative to dopamine receptor antagonists in blocking VLP entry matches that for inhibition of TPC2 currents (Fig. 6).

4. Discussion

TPCs are channels of physiological, and increasingly, patho-

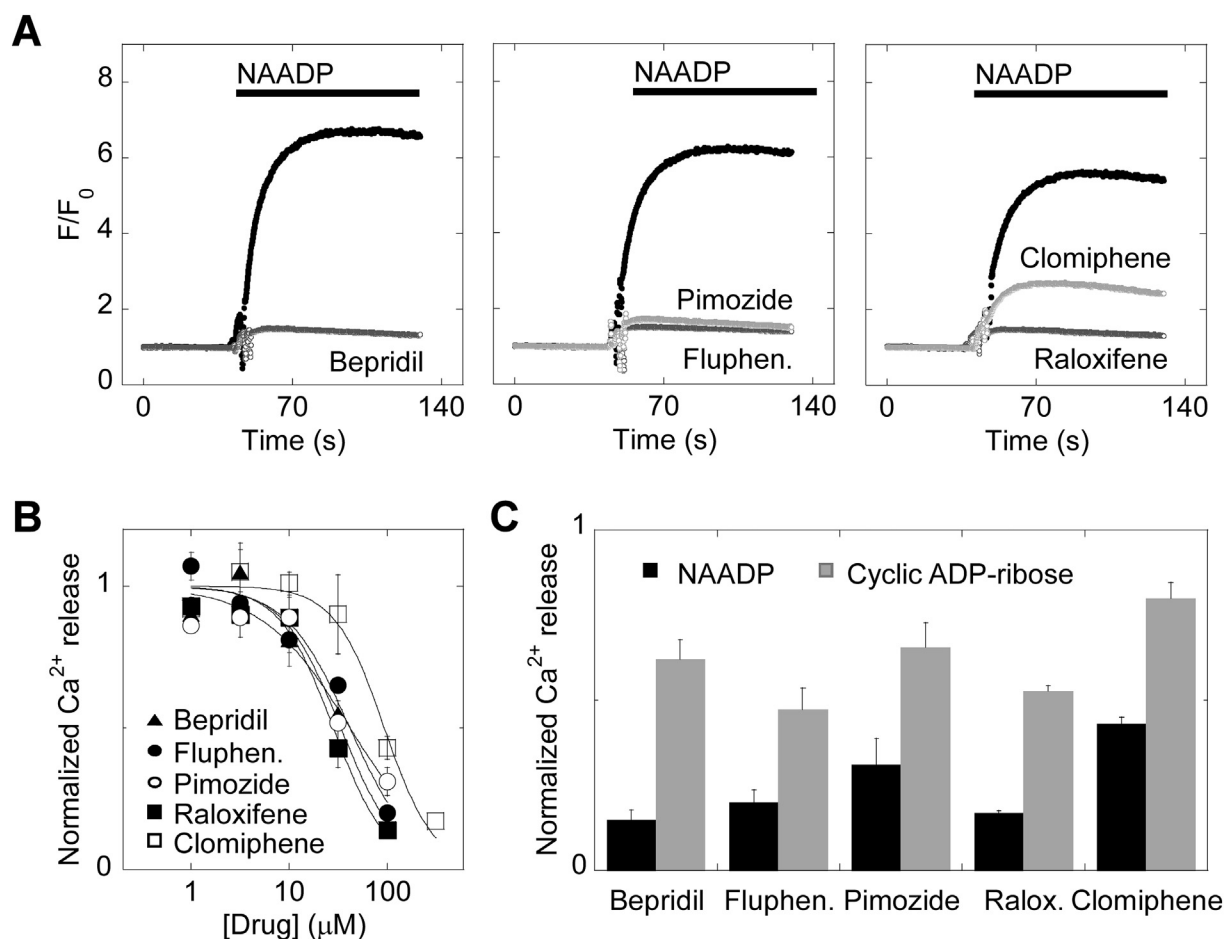


Fig. 3. Ca^{2+} channel blockers, dopamine antagonists and selective estrogen receptor modulators block endogenous NAADP-mediated Ca^{2+} signals. A, exemplar Ca^{2+} signals, recorded using Fluo-4 from sea urchin egg homogenates pre-incubated with the indicated drug (100 μM) or DMSO (black traces) for ~ 1 min prior to stimulation with NAADP (1 μM). B, concentration-effect relationships for blockade of NAADP responses by the various drugs ($n = 3$). Data are normalised to the NAADP response in the presence of DMSO. C, summary data quantifying the effect of the drugs (100 μM) on Ca^{2+} response to NAADP versus cyclic ADP-ribose ($n = 3$). Data are normalised to the NAADP and cyclic ADP-ribose responses in the presence of DMSO.

physiological importance [19]. However, their pharmacology is poorly defined and there are currently no approved drugs to target them. To begin to fill this gap, we performed a virtual structure-based screen using a database of approved drugs and the pore region of TPC2 (Fig. 1). Key to our strategy was i) the availability of structural information for mammalian TPCs, ii) the finding that TPCs are novel host factors for EBOV entry and iii) repurposing screens initiated in the wake of the 2014 Ebola epidemic, which yielded a number of approved drugs as novel EBOV entry inhibitors. To prioritise hits, cross-referencing the latter ‘wet’ data with our ‘dry’ modelling highlighted drugs targeting dopamine and estrogen receptors (Fig. 2). These drugs demonstrably inhibited TPCs in three independent assays (Figs. 3, 5, 6), and we correlated their potency with inhibiting Ebola VLP entry (Fig. 7). Single channel analyses confirmed an action of these drugs on the TPC2 pore (Fig. 5).

Results from the enrichment analysis of our virtual screen identified a number of drugs targeting distinct G-protein coupled receptors (Fig. 1D). Notably, a high throughput screen for drugs that block EBOV and Marburg virus (both filoviruses) infection also highlighted numerous drugs targeting G-protein coupled receptors [62]. Common drug classes in that screen and ours include those against 5HT, muscarinic and histamine receptors. This further links TPCs to EBOV infection. We focused our functional analyses on dopamine receptor antagonist and SERMs based on our cross-referencing strategy (Fig. 2). But

these compounds represent only a fraction of our top 200 hits. Thus, our screen offers considerable scope for identifying additional TPC blockers in the future.

The dopamine receptor drugs identified here as TPC blockers included several phenothiazines (fluphenazine, trifluoperazine, prochlorperazine and thioridazine) that are used as anti-psychotics (Fig. 2). Our data is consistent with a recent screening campaign published while this work was in progress, which also identified phenothiazines as inhibitors of NAADP-induced Ca^{2+} release from sea urchin egg homogenates [63]. Indeed, the IC_{50} values for fluphenazine, 42 μM (Fig. 3B) and $\sim 11 \mu\text{M}$ [63], are comparable. This congruence highlights the power of our *in silico* strategy for drug identification which, together with our electrophysiological analyses, points mechanistically to drug action on the TPC pore. We also identified the structurally distinct dopamine receptor drug, pimoziide, a diphenylbutylpiperidine. As well as targeting the dopamine receptor, pimoziide also targets voltage-gated Ca^{2+} channels [64]. The common action of pimoziide on TPCs and voltage-gated Ca^{2+} channels likely reflects an evolutionary kindred between these two members of the voltage-gated ion channel superfamily [24]. Although voltage-gated Ca^{2+} channels have been implicated in viral entry [65], their functional absence in non-excitable cells (such as HeLa cells used in this study) points to TPCs as the likely targets.

Our analyses identifying drugs targeting estrogen receptors as

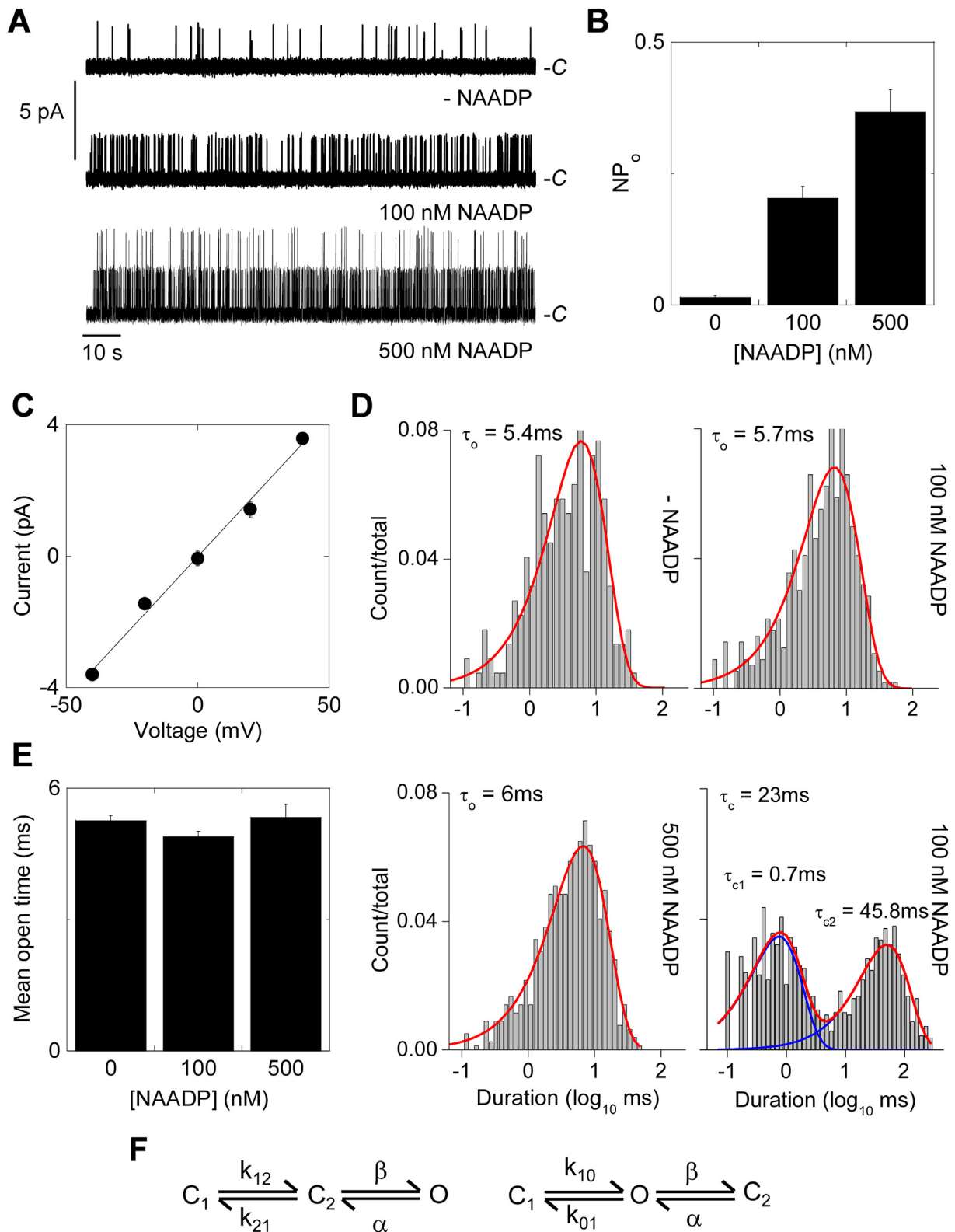


Fig. 4. NAADP stimulates single channel activity of TPC2 rerouted to the plasma membrane.

A, exemplar single channel recordings from inside-out patches excised from HEK cells expressing plasma membrane-targeted TPC2 in response to the indicated concentrations of NAADP. Na⁺ was the charge carrier and the holding potential was +40 mV. C denotes the closed state.

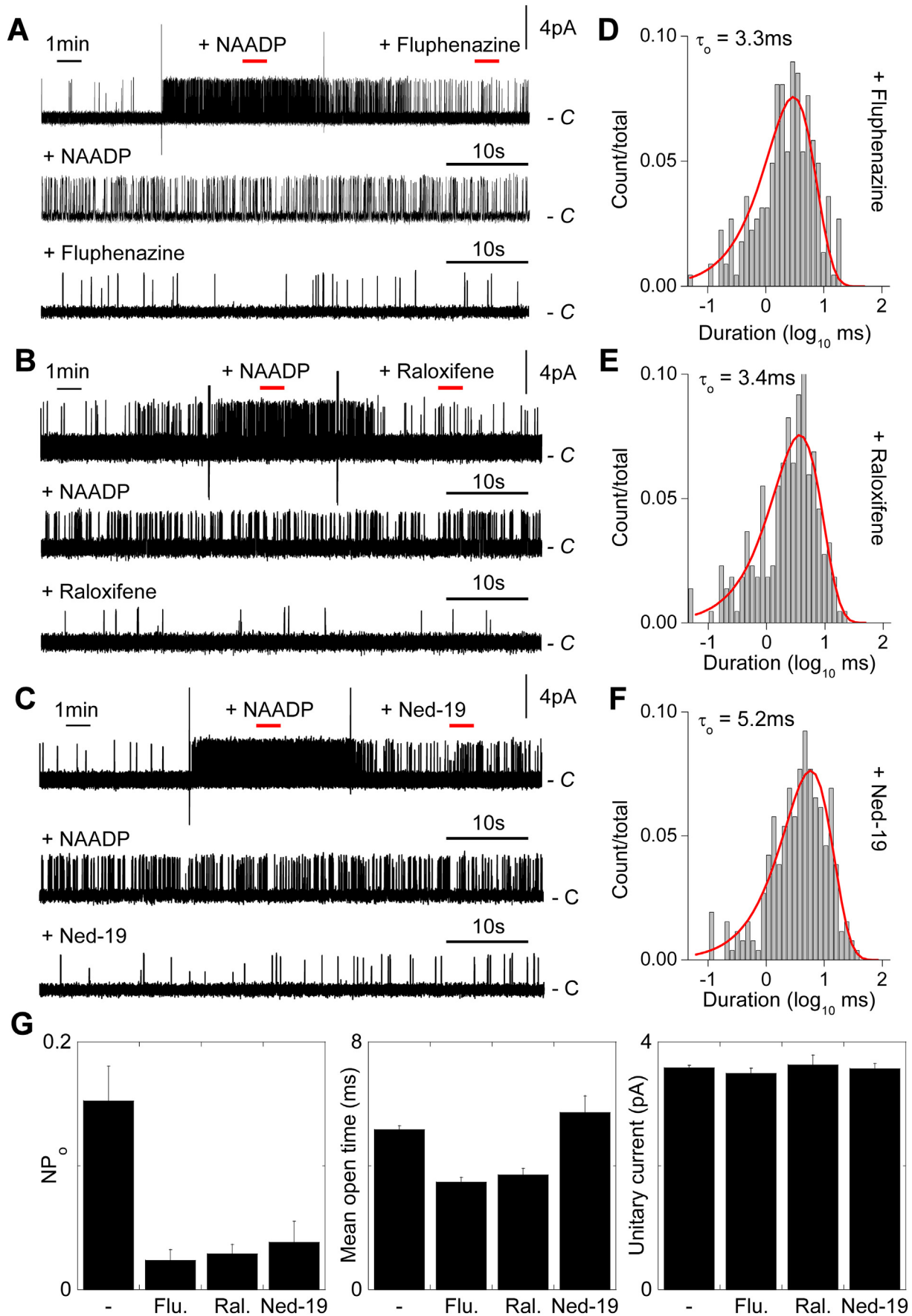
B, summary data quantifying open probability (NP_o; n = 3–28).

C, current-voltage relationships derived from records similar to those shown in A (n = 3).

D, exemplar open and closed time distributions at the indicated NAADP concentration.

E, summary data quantifying mean open time (n = 3–28).

F, likely gating schemes for TPC2.



(caption on next page)

Fig. 5. Dopamine antagonists and selective estrogen receptor modulators inhibit single channel activity of TPC2 by reducing open time. A–C, exemplar single channel records from inside-out patches expressing plasma membrane-targeted TPC2. Patches were sequentially challenged with 100 nM NAADP and either fluphenazine (100 μ M; A), raloxifene (100 μ M; B) or Ned-19 (1 μ M; C). Large vertical deflections are addition artefacts. Red bars indicate the sections of the upper trace in the presence of NAADP before and after drug addition that were expanded in the corresponding lower traces. D–F, exemplar open time distributions from records similar to A–C in the presence of the indicated drug. G, summary data quantifying open probability, mean closed time and unitary conductance for TPC2 activity in response to 100 nM NAADP in the absence ($n = 10$) and presence of the indicated drug ($n = 3$ –4). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

additional TPC blockers converged on first (clomiphene, tamoxifen, toremifene), second (raloxifene) and third (bazedoxifene) generation SERMs (Fig. 2). These drugs act as estrogen receptor agonists or antagonists, depending on the tissue, and are used for treating breast cancer and osteoporosis [66]. Off-target effects of these drugs are more than likely related to well-known rapid non-genomic actions of estrogen [67]. Indeed, inhibition of both viral entry and TPC activity by sub-micromolar concentrations of raloxifene is striking. Additional effects of SERMs such as destabilisation of either the EBOV glycoprotein [68] or lysosomal membranes due to their cationic amphiphilic nature [69] are possible. Nevertheless, both dopamine receptor antagonists and SERMs emerge as novel TPC blockers and provide independent scaffolds for small molecule development.

Estrogen receptor drugs appear more potent in blocking TPC activity (Fig. 6) and Ebola VLP entry (Fig. 7) than dopamine receptor drugs. This correlation points to a common mechanism of action (Figs. 6–7). The half maximal inhibitory concentrations by raloxifene were similar but fluphenazine was more effective at inhibiting VLP entry (Fig. 7) than blocking the channel (Fig. 6). This could reflect additional effects of fluphenazine underlying entry inhibition or the very different conditions under which the two assays were performed (acute drug treatment versus pre-incubation; broken versus intact cells). One other notable difference was the relative poor potency of the drugs in blocking Ca^{2+} signals in egg homogenates (Fig. 3) compared to lysosomal currents (Fig. 6). Again, this could relate to the way in which drugs were administered or the activating ligand. Alternatively, this could point to species differences in drug block. Nevertheless, our study suggests that TPCs may be the target for a number of EBOV entry inhibitors described previously for which mechanistic information was lacking. How TPCs promote EBOV entry is unclear at present. A recent study provided evidence that TPCs may also regulate entry of the MERS corona virus [70]. Pharmacological inhibition with tetrandrine analogues or molecular knockdown of TPCs was shown to reduce Furin

activity. This protease activity is required for priming of the envelope protein [71]. Thus, compromised activity upon TPC blockade could underlie reduced MERS corona virus entry. Such a mechanism is unlikely to regulate entry of EBOV because it is processed by endo-lysosomal cathepsins [72]. Moreover, entry of pre-cleaved EBOV is also blocked by tetrandrine [20] suggesting that TPCs act at a step distal to proteolytic processing. Clearly, further work is required to understand the mechanisms of TPC involvement in EBOV entry.

Although cellular studies repeatedly link TPCs to NAADP-mediated Ca^{2+} signalling, electrophysiological analyses of TPCs are less congruent with some studies failing to demonstrate appreciable Ca^{2+} permeability or NAADP activation [39,73]. Our previous whole cell recordings and single channel analyses demonstrated robust activation of Ca^{2+} and Cs^{+} currents in response to NAADP [32]. Here, we used Na^{+} as the major permeant ion. Comparing single channel conductances in this study (87 pS, Fig. 4) with those for Ca^{2+} (40 pS) and Cs^{+} (128 pS) reported previously [32] points strongly to TPC2 as a non-selective cation channel under these experimental conditions. This conclusion concurs with bilayer studies of TPC2 [74,75] and electrophysiological analyses of endogenous NAADP-activated TPC2 currents recorded from enlarged vacuoles (Na:Ca permeability ratio close to unity) [76]. TPCs interact with many other proteins [16] some of which likely act to effect and modify gating. Loss of accessory factors might well explain NAADP insensitivity in some studies. Indeed, such loss might also affect the ion selectivity of TPCs as accessory proteins are known to dictate the ion selectivity of other channels such as Orai [77] and MCU [78].

Our single channel analyses provide evidence that the drugs target the pore region of TPCs (Fig. 5). The reduction in open time (~ 2 -fold) was modest relative to the reduction in open probability (~ 10 -fold) pointing to an open channel block mechanism. This is compatible with effects produced by inhibitors known to interact directly with the pore of other channels [79–81]. Of note, phenothiazines have long been

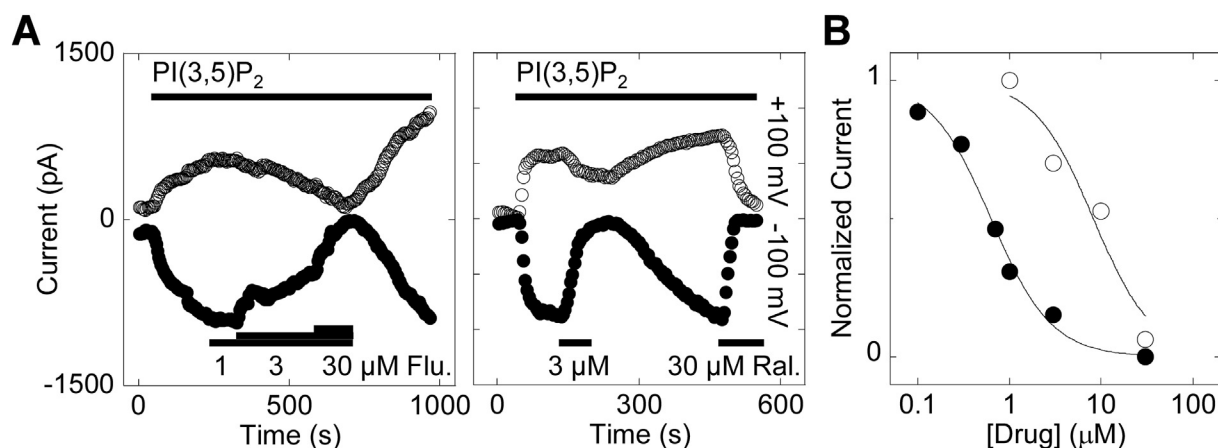


Fig. 6. Dopamine antagonists and selective estrogen receptor modulators inhibit lysosomal TPC2 currents. A, exemplar currents recorded from enlarged lysosomes isolated from HEK cells expressing lysosome-targeted TPC2 in response to PI(3,5)P₂ (1 μ M). The holding potentials were +100 mV (upper traces) and -100 mV (lower traces). Lysosomes were challenged with fluphenazine (Flu.) and raloxifene (Ral.) at the concentrations and for the times indicated. B, concentration-effect relationships for blockade of PI(3,5)P₂ currents by fluphenazine (open circles) and raloxifene (closed circles; $n = 2$). Data are normalised to the PI(3,5)P₂ response in the absence of the drugs.

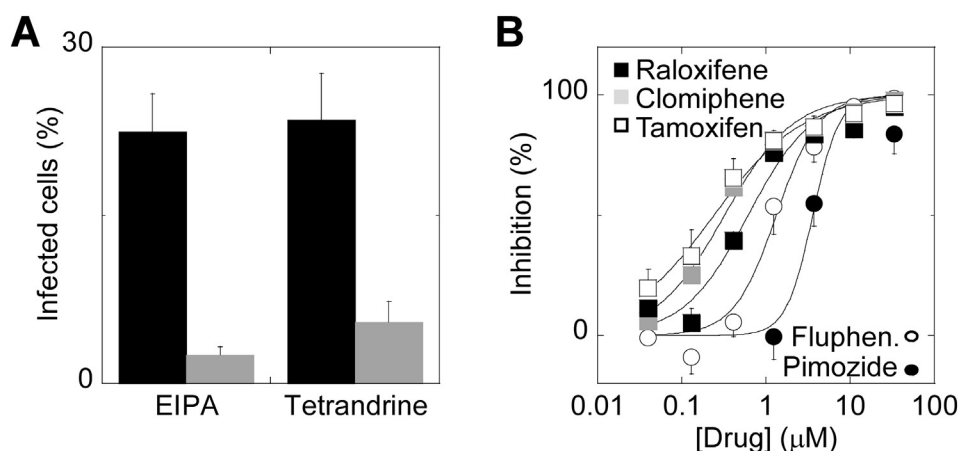


Fig. 7. Dopamine antagonists and selective estrogen receptor modulators inhibit entry of Ebola virus-like particles.

A, infection of HeLa cells with Ebola virus-like particles in the absence and presence of EIPA (11.1 μM; $n = 3$) or tetrandrine (1.2 μM; $n = 4$). B, concentration-effect relationships for blockade of virus-like particle entry by the indicated dopamine antagonist (squares) or SERMs (circles). $n = 3-4$.

known to antagonize calmodulin [82] and previous studies with trifluoperazine have suggested a role for calmodulin in NAADP-mediated Ca^{2+} signalling [83]. Our data, with fluphenazine at least, suggests an additional more direct action of phenothiazines on TPCs likely unrelated to calmodulin (Fig. 5). Clearly, further structure-function analyses are warranted.

In contrast to fluphenazine and raloxifene, Ned-19 did not affect open time (Fig. 5). It thus effectively acts as a gating modifier by reducing the overall frequency of channel openings through directly or indirectly influencing channel regions distal to the pore. Thus, our analyses uncover fundamentally distinct mechanisms for channel inhibition by the drugs and Ned-19. The lack of effect of Ned-19 (and also NAADP; Fig. 4) on open time support the notion that they act more remotely through associated NAADP-binding proteins. [38]. Uncertainty about the number of channels as well as the limited number of events in patches with low activity prevented us from performing close time distribution analysis upon channel inhibition. But further analysis is warranted not least due to the presence of a multiple closed times (Fig. 4F) which could reflect desensitized states.

In sum, we have identified approved drugs as TPC blockers. We used a novel strategy that affirmed the role of TPCs in EBOV entry and which should significantly aid in expanding the pharmacology of this ubiquitous class of endo-lysosomal ion channels.

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Transparency document

The Transparency document associated with this article can be found in the online version.

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