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Original Citation

Mertins, Barbara, Psakis, Georgios, Grosse, Wolfgang, Back, Katrin Christiane, Salisowski, Anastasia, Reiss, Philipp, Koert, Ulrich and Essen, Lars-Oliver (2012) Flexibility of the N-Terminal mVDAC1 Segment Controls the Channel’s Gating Behavior. PLoS ONE, 7 (10). e47938. ISSN 1932-6203

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Flexibility of the N-Terminal mVDAC1 Segment Controls the Channel’s Gating Behavior

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Abstract
Since the solution of the molecular structures of members of the voltage dependent anion channels (VDACs), the N-terminal α-helix has been the main focus of attention, since its strategic location, in combination with its putative conformational flexibility, could define or control the channel’s gating characteristics. Through engineering of two double-cysteine mVDAC1 variants we achieved fixing of the N-terminal segment at the bottom and midpoint of the pore. Whilst cross-linking at the midpoint resulted in the channel remaining constitutively open, cross-linking at the base resulted in an “asymmetric” gating behavior, with closure only at one electric fields orientation depending on the channel’s orientation in the lipid bilayer. Additionally, and while the native channel adopts several well-defined closed states (S1 and S2), the cross-linked variants showed upon closure a clear preference for the S2 state. With native-channel characteristics restored following reduction of the cysteines, it is evident that the conformational flexibility of the N-terminal segment plays indeed a major part in the control of the channel’s gating behavior.

Introduction
Voltage dependent anion channels (VDACs) are the most abundant proteins in the mitochondrial membrane with about 10 000 copies per cell [1]. VDACs mediate the flow of energy carrying molecules like ATP, ADP, pyruvate or succinate [2]. With their N-terminus providing a surface for interactions with pro- and anti-apoptotic proteins, they also become key players in the control of apoptotic signaling [3,4,5]. In lipid bilayer recordings, VDACs show multiple conductance states: a high conductance, referred to as the open state, and several low conductance states referred to as closed or partly open [6]. The low conductance states are usually reported at a current of 40–60% of that of the high conductance. In the high conductance state the channel is weakly anion selective, but exhibits weak cation selectivity when partly or fully closed [7]. Despite their extensive characterization, first insights into the VDAC 3D-structure were only recently obtained. Independently solved X-ray structures revealed an atypical 19 stranded β-barrel with an α-helix spanning the middle of the pore [3,6,7]. Based on the current structural data, it is thought that alignment of the α-helix with the barrel wall yields the “open” state [8]. Despite the availability of structural information, the complexities of the VDAC conductance states and the molecular mechanism underlying their transitions are poorly understood. Current hypotheses favor control of the opening/closure of the channel through a complete or partial movement of the N-terminal helix with its flexible hinge region (amino acids 19–25) into [9] or out of the pore [10]. Although an inward α-helical movement could control the channel’s activity through blockage, it fails to account for the channel’s ion selectivity, which an outward-movement model addresses more adequately [10]. None of the proposed models, however, has received strong experimental support. Recent work questioned the previous hypotheses and claimed that the N-terminal helix is not responsible for the modulation of the channel’s gating at all [11].

Here, we report the engineering of two mVDAC1 channel variants, which under oxidative conditions deviated strongly from the typical native channel gating: one that has lost its symmetrical response to the applied potential (V3C-K119C) and one which was trapped in its conducting state (A14C-S193C). We confirm formation of the disulfide bonds between the aforementioned residues by SDS-PAGE and/or peptide mass fingerprint spectrometry as well as planar lipid bilayer recordings, thus attesting the current structural models. In addition, we demonstrate that the α-helix plays a major part in the voltage gating process by undergoing a combination of two structural changes: one allowing the far N-terminus to flip into the channel, and a second requiring a re-organization of the α-helical residues. Beside the observed S1 state we further identify the presence of two well-defined low-conductance sub-states (S2A and S2B), which reinforce the flexibility of VDACs, and use the above mechanism to address aspects of the channel’s ion selectivity.
Materials and Methods

1. Cloning and Site-directed Mutagenesis
   
   Plasmid vectors, harboring the murine VDAC1 (mVDAC1) gene, were kindly provided by Dr Jeff Abramson (University of California LA). The mVDAC1 gene was amplified by PCR, using the 5ʼ TTA ATA ATC GAG TTA TGC TTG AAA TTD CAC TGC TAG GC 3ʼ and 5ʼ-ATA AAT CAT ATG GCC GTG CCT CCC AC 3ʼ primers, introducing the XhoI and NdeI restriction sites respectively. PCR products were XhoI/NdeI (Fermentas) digested, PCR cleaned (Qiagen) and cloned into pre-digested with the same enzymes and dephosphorylated (alkaline phosphatase, NEB) pET20b plasmid vector (Neogen). Liguations were performed with T4-ratio, according to the manufacturer’s instructions. Transformants were selected on an LB-ampicillin (100 µg ml⁻¹) (Applichem) containing plates.

   For the construction of the Δ1–21 deletion variant, the XhoI/ NdeI-flanked mVDAC1 gene was PCR amplified by primers 5ʼ-TTA ATA ATC GAG TTA TGC TTG AAA TTD CAC TGC TAG GC-3ʼ and 5ʼ-ATA AAT CAT ATG GCC GTG CCT CCC AAC CTG ATT TAC CAG GAG-3ʼ and 5ʼ-GTC AGC CCC GAG GTT GAT GTG C-3ʼ were used for the introduction of the C127A mutation. Having confirmed the identity of the C127A construct by DNA sequencing (Qiagen), the latter was subsequently used in a second PCR round for the introduction of the final mutation (primers 5ʼ-GGG TAC TGC AGG-3ʼ, and 5ʼ-C GTA TGT GGG AGG GCA GGC CAT and 5ʼ-C CTT CTG GTA AAT GCA GCC ACC ACA ATG ATC CAG CAC CAG AT TC CG G-3ʼ).

   Further mutagenesis was performed on the cysteine-free mVDAC1 plasmid using the 5ʼ-CTT GCC AGG AAG TCC TGG AGG GAT ATG TTC-3ʼ and 5ʼ-GAA GAC ATC ACC CCT GCC GGA CTT GCC AGG-3ʼ primers in the first round (A14C mutation), and 5ʼ-GAA GAT TGG GTG GCC TTG ACC TGC CTG ACC CCC GAG GTT GTG GTG C-3ʼ were used for the introduction of the C127A mutation. Having confirmed the identity of the C127A construct by DNA sequencing (Qiagen), the latter was subsequently used in a second PCR round for the introduction of the final mutation (primers 5ʼ-CC AAC TAT CAG GTG CAT CCT GAT GCC GGC TCC TGG GCC GCA AAG G-3ʼ and 5ʼ-C TTT GGC CGA AAA GGC GGC ATC AGG ATG CAC CAG AT A TA TG G-3ʼ).

   For effective induction of disulfide cross-linking in mVDAC1, the cell pellet was washed three times in 50 mM Tris/HCl (pH 8.0), 100 mM NaCl and subsequently disrupted by emulsification (EmulsiFlex C5, Avestin). The crude cell extract was centrifuged (25402 g, 30 min, 4°C), the supernatant was discarded and the protein-containing pellet was washed with 100 mM NaCl, 1 mM EDTA, 2.22% (v/v) lauryldimethylamine-oxide (LDAO), allowing for a ten-fold dilution of the guanidinium chloride concentration, and the mixture was gently stirred overnight at 8°C. Following a centrifugation step (6400 g, 10 min, 4°C), for removal of insoluble aggregates, the supernatant was subsequently added drop-wise to 25 mM NaH₂PO₄/Na₂HPO₄ (pH 7.0), 10 mM NaCl, 1 mM EDTA, 1 mM DTT, 0.1% (v/v) LDAO (a further ten-fold dilution of the guanidinium chloride concentration), and the mixture was gently stirred overnight at 8°C for a further day. Finally the mixture was applied onto a pre-equilibrated Fractogel EMD-SE Hicap cation-exchange column (5 mL) (Merck) which was attached to an AKTA prime purification system (Amersham Biosciences). Elution was performed using a multi-segment gradient of 10–1000 mM NaCl in 25 mM NaH₂PO₄/Na₂HPO₄ (pH 7.0), 100 mM NaCl, 1 mM EDTA, 1 mM DTT, and 0.1% (v/v) LDAO. The mVDAC1-containing fractions were pooled and concentrated using a 10-kDa Amicon concentrator (Millipore) and applied onto a Superdex 200 (GE-Healthcare) size-exclusion column (5 mL) (Merck) which was attached to a Superdex 200 (GE-Healthcare) size-exclusion column (SEC) pre-equilibrated with 10 mM Tris/HCl (pH 8.0), 100 mM NaCl and 0.05% LDAO. SEC regularly yielded >95% pure protein as verified by SDS-PAGE. Size excluded mVDAC1 fractions were pooled and further concentrated. Protein concentrations were determined from the UV-Vis absorbance of tested samples at 280 nm (Nanodrop ND-1000, peqLab), using extinction coefficients as predicted by ProtParam [15]. Purified samples were stored at 4°C until further use.

2. Inclusion Body Production of mVDAC1 Variants

   E. coli BL21(DE3)-omp9 cells [F⁻, ompT ksdB (RKM - mB)] gal dcm (DE3) ΔlamB ompF:tn5 ΔompA ΔompC ΔompN::Ω] [14] were transformed with pET20b harboring the native or mutated mVDAC1 gene without its signal sequence. Transformants were grown in LB medium containing ampicillin (100 µg ml⁻¹) and 1.0% (w/v) glucose at 37°C. Induction was performed by addition of 1 mM isopropyl-β-D-thiogalactopyranoside (IPTG) (Gerbu Biotech.) at an optical density (λ = 595 nm) of ~0.6. Cells were allowed to grow for further 4 hours before harvesting (6040 g, 13 min, 4°C). Cell pellets were solubilized in 50 mM Tris/HCl (pH 8.0), 100 mM NaCl and were subsequently disrupted by emulsification (EmulsiFlex C5, Avestin). The crude cell extract was centrifuged (25402 g, 30 min, 4°C), the supernatant was discarded and the protein-containing pellet was washed three times in 50 mM Tris/HCl (pH 8.0), 100 mM NaCl, 10 mM EDTA, 2.5% (v/v) Triton-X-100 (Triton-TEN buffer) for removal of residual E. coli lipids, and three times in 50 mM Tris/HCl (pH 8.0), 100 mM NaCl (TEN buffer). The protein pellet was finally resuspended in TEN buffer, fast-frozen in liquid nitrogen and stored at −80°C until further use.

3. Protein Refolding and Purification

   Protein refolding was accomplished by rapid dilution in a two-step process. First, inclusion bodies (10 mg ml⁻¹) were denatured in 25 mM NaH₂PO₄/Na₂HPO₄ (pH 7.0), 100 mM NaCl, 1 mM EDTA, 10 mM DTT, 6 M guanidinium chloride. The denatured protein mixture was added drop-wise to 25 mM NaH₂PO₄/Na₂HPO₄ (pH 7.0), 100 mM NaCl, 1 mM EDTA, 1 mM DTT, 0.1% (v/v) LDAO, allowing for a ten-fold dilution of the guanidinium chloride concentration, and the mixture was gently stirred overnight at 8°C. Following a centrifugation step (6400 g, 10 min, 4°C), for removal of insoluble aggregates, the supernatant was subsequently added drop-wise to 25 mM NaH₂PO₄/Na₂HPO₄ (pH 7.0), 10 mM NaCl, 1 mM EDTA, 1 mM DTT, 0.1% (v/v) LDAO (a further ten-fold dilution of the guanidinium chloride concentration), and the mixture was gently stirred overnight at 8°C for a further day. Finally the mixture was applied onto a pre-equilibrated Fractogel EMD-SE Hicap cation-exchange column (5 mL) (Merck) which was attached to an AKTA prime purification system (Amersham Biosciences). Elution was performed using a multi-segment gradient of 10–1000 mM NaCl in 25 mM NaH₂PO₄/Na₂HPO₄ (pH 7.0), 100 mM NaCl and 0.05% LDAO. SEC regularly yielded >95% pure protein as verified by SDS-PAGE. Size excluded mVDAC1 fractions were pooled and further concentrated. Protein concentrations were determined from the UV-Vis absorbance of tested samples at 280 nm (Nanodrop ND-1000, peqLab), using extinction coefficients as predicted by ProtParam [15]. Purified samples were stored at 4°C until further use.

4. Oxidation and Reduction of mVDAC1 Variants

   For effective induction of disulfide cross-linking in mVDAC1 variants, purified proteins were treated with 100 mM copper-phenanthroline (CuX₄Ph) as previously described [11,16]. CuX₄Ph reactions were quenched by addition of 0.5 mM EDTA. Disulfide bond reduction was mediated by addition of 10 mM DTT. To prevent potential interference of CuX₄Ph or DTT with the subsequently performed assays, treated protein samples were further purified using an Amicon Ultra-0.5 mL Filter Device (Millipore) according to the manufacturer’s instructions. The presence or absence of disulfide bonds was confirmed independently by non-reducing SDS-PAGE and peptide mass fingerprint spectrometry.
5. SDS-PAGE

Protein samples were resolved on 12% gels, using the Laemmli buffer compositions [17]. Homogeneity and purity were densitometrically determined. Gel-shift assays using CuX2Ph- and DTT-pre-treated samples indicated the presence and disruption of the disulfide cross-link, respectively.

6. Peptide Mass Fingerprinting

To cap free cysteines, 2-iodoacetamide (Applichem) was added in a final concentration of 1 mM to the mVDAC1 samples and stirred over night at 8°C. To remove free 2-iodo-acetamide the protein was precipitated using a four-fold excess of acetone and the samples were incubated for 2 hours at −20°C. Precipitated mVDAC1 was centrifuged (25402 g, 3 min, 4°C) and the pellets were washed with ice water and ice-cold ethanol. Pellets (341 μg each) were dried at 37°C, dissolved in 100 μL 10% acetonitrile and digested over night at 37°C with 10 μg sequencing grade trypsin (Promega) containing 25 μL 50 mM NH4HCO3 buffer at pH 8.0. For chromatographic separation of the tryptic peptides an 1100 Agilent HPLC system was used with a 150x3 mm Kinetex HPLC column (Phenomenex) at a temperature of 40°C and a flow rate of 0.2 mL/min. UV-detection of the peptides was carried out at 215 nm. The following gradient was applied by mixing water/0.1% TFA (solvent A) and acetonitrile/0.1% TFA (solvent B): 5% B for 10 minutes, linear increase to 60% B within 150 minutes, linear increase to 90% B in additional 5 minutes and then holding B at 90% for 10 minutes. Mass spectrometric detection of the peptides was done by online electrospray-ionisation-mass spectrometry (ESI-MS) with an LTQ-FIT mass spectrometer (ThermoScientific). An FTMS scan was carried out in the mass range of 400–2000 m/z, followed by three automated ion trap MS/MS scans. The disulfide-bridged peptides were identified from comparisons of predicted and actual masses.

7. Single Channel Conductance Recordings

Single channel conductance recordings were performed using the Black Lipid Membrane (BLM) technique [18,19,20,21,22,23]. A sample of n-decanol-solubilized diphytanoyl phosphatidylcholine membrane mixture (25 mg/mL) was painted over a 200 μm circular hole, separating the chamber compartments (polystyrene cuvette: CP2A, bilayer chamber: BCH-22A, Warner Instruments) filled equally with BLM buffer [10 mM Tris/HCl (pH 7.4), 1 M KCl, 5 mM CaCl2]. Protein (7.5–15 μg) was added to one compartment beside the planar lipid layer and a voltage-gradient was imposed across the membrane whilst waiting for insertion of single channels. Generally, ±40 mV voltages were applied, electric current was recorded using a Multipatch 700B patch-clamp amplifier connected to a Digidata 1440A A/D converter and traces were visualized through the pClamp 10.2 software (Axon Instruments). To avoid noise convolution of the recordings and to minimize the possibility of channels reconstituting in an antiparallel manner (see results and discussion), we analyzed only the recordings corresponding to the spontaneous insertion of up to five channels. Data were collected at 5 kHz and sampled at 200 Hz for further analysis. For each measurement, high and low conductance sub-states were defined relative to a clear baseline, and their corresponding differences to the baseline were evaluated by the software. Linear regressions were performed by Origin 7 (Origin Lab). For every determined conductance value, 15–84 separate measurements were averaged at different voltage steps. Statistical tests were performed using the GraphPad (GraphPad Software) online calculators. Note that values are given as mean ± SEM, unless otherwise stated.

Results

1. CuX2Ph Induced Cross-linking in mVDAC1 Variants Validates the 3EMN 3D-Structure

Basing our construct design (Figure 1A) on the murine VDAC1 structure (3EMN, [3]) we found that covalent fixing of the N-terminal segment by engineering disulfide bindings to the porin’s inner wall was favorable. Following the successful refolding and purification of mVDAC1 variants containing appropriate disulfide mutations at the positions A2-E121, A14-S193 and V3-K119, disulfide-bridge formation was confirmed by denaturing SDS-PAGE electrophoresis. Reduced double cysteine variants migrated generally slower and closer to their actual molecular mass than their non-linearized, cross-linked counterparts (Figure 1B). Of the investigated variants, only A2C-E121C partly resisted the CuX2Ph treatment (Figure 1B), as it failed to oxidize completely, implying that formation of the corresponding bridge here was structurally disfavored. In addition, peptide mass fingerprinting of the oxidized A14C-S193C variant confirmed quantitative disulfide formation, by detection of the S13-R15/T175-Y197 fragment (Figure S1). The corresponding fragment of the V3C-K119C variant could not be detected, most likely due to lack of ionization. As confirmed by SDS-PAGE, the fully reduced mVDAC1 variants had a limited half-life of ~22 hours (Figure S2) in the presence of air and hence showed a strong tendency towards oxidation. In essence, and in agreement with previously published works [11], our observations increased confidence that the 3EMN model [3] corresponds to that of the native and biologically most relevant open channel conformation.

2. Closure of the Native mVDAC1 Channel Occurs through at Least Two Low-conductance States

Native mVDAC1 exhibited the well observed [3,4,7,24,25,26,27] and expected gating behavior, responding linearly to the application of both positive and negative potentials, alternating between high and low conductance states (Figure 2A). In the open state (S0), the porin exhibited a conductance similar to those previously described (Table 1). When we calculated the mean of the low conductance state data (green and red squares, Figure 2B) we obtained a value of 2.34 ± 0.06 nS (n = 680) (Table S1). The latter conductance was ~1.4-fold different to that reported by Ujwal et al. [3] (Table 1). Thinking that this difference could not solely stem from random error, we attempted fitting the distribution of the low conductance data to the sum of two Gaussians (Figure 2C, Table S1). The sum of the two distributions worked better than the single Gaussian (Table S1) and enabled a clear identification of two closed states: a S1 (exhibiting 66% of the conductance of the S0 state) and a S2 (exhibiting 48% of the conductance of the S0 state) (Figures 2B, 2C). Interestingly, the determined S2 conductance value (1.90 ± 0.06 nS; n = 294); Figures 2B, 2C) in agreement with the values previously reported for channel closure (Table 1). Student t-tests were performed to statistically validate the differences between the observed conductances for 1) the S1 (n = 386) and S2 (n = 294) closed states and 2) for each closed state and the mean closed state (n = 680) derived from the single Gaussian fit (Table S1). S1 and S2 conductances were different not only with respect to each other (P = 10−4), but also with respect to the mean closed state conductance derived from the single Gaussian fit (Table S1). Furthermore, the S2 population (0.80–2.20 nS) indicated a double Gaussian distribution, revealing the presence of an additional low-conductance sub-state, S2B (Figure 2C). Hence, the conductance of the closed conformation is a composite of at least three separate, low conductance states, S1, S2A and S2B (Table S1). The average
dwell time of the channel in one state before switching to the next was 10.34 ± 1.65 s (n = 310) (Figure S3).

3. The Δ1-21-mVDAC1-Y22C (Δ21-mVDAC1) Deletion Construct Shows a Clear Preference for the Open State

To prove the dominant role of the N-terminal α-helix in any gating behavior of mVDAC1, we analyzed the N-terminal truncation variant Δ21-mVDAC1. Current recordings were obtained following the reconstitution of the Δ21-mVDAC1 construct in planar lipid membranes (Figure 3A). Upon application of +10 mV, single channels spontaneously reconstituted in the artificial bilayer in the open state and remained constitutively open (Figure 3A), exhibiting a conductance of 4.00 ± 0.04 nS (n = 20) in agreement with the reported observations of Abu-Hamad et al. [4] and De Pinto et al. [28]. T-test comparison of the S0 state conductances for native and Δ21-mVDAC1 proteins suggested that the compared values were similar (P = 0.80; sample size for native protein n = 680).

Figure 1. Cysteine mutagenesis on mVDAC1 and cross-linking of the N-terminal segment to the barrel wall. A) Cartoon representation of mVDAC1, based on the 3EMN structure [3], viewed perpendicular to the membrane plane. Location of the engineered cysteines and expected formation of disulfide bonds are shown as inlays. B) 12% Coomassie stained, non-reducing SDS-PAGE analysis of native and engineered mVDAC1 variants. Proteins were pre-treated with to copper-phenanthroline (oxidized) or DTT (reduced) as described in Materials and Methods. Increased electrophoretic mobility of protein bands indicated formation of disulfide cross-links. Following reduction by 10 mM DTT, “linearized” proteins migrated slower than their cross-linked counterparts.

doi:10.1371/journal.pone.0047938.g001

Figure 2. The gating profile of mVDAC1 shows switching events between an open (S0) and two closed (S1 and S2) states. A) Representative traces of the mVDAC1 gating activity at +10 and +40 mV. The observed S0, S1 and S2 states, in the +40 mV trace, are indicated by blue, red and green lines respectively. B) Ohm-plot used for the determination of the conductance values of the corresponding states. Reconstituted native mVDAC1 responded linearly to application of both positive and negative potentials, alternating between an open (S0; blue) and two major (S1; red and S2; green) closed states. Determined conductances were: S0; 3.94 ± 0.04 nS (n = 680), S1; 2.61 ± 0.01 nS (n = 386) and S2; 1.90 ± 0.06 nS (n = 294). Bars represent standard errors of a minimum of 15 replicates per measurement. C) Histogram of occurrences of low conductance values for native mVDAC1. The sum of two Gaussians indicated clearly the presence of two low conductance states for the native channel: S1 (red) and S2 (green). S1 and S2 conductances were different not only with respect to each other (P = 10^-4), but also with respect to the mean closed state conductance derived from the single Gaussian fit (Table S1). Applying the sum of two Gaussians to the S2 state (R^2:0.98) resulted in the identification of an additional population (S2B; dashed grey line), suggesting that S2 is the composite of two low conducting S2A (2.03 ± 0.02 nS, n = 224) and S2B (1.48 ± 0.02 nS, n = 70) sub-states (Table S1). Values are given as mean ± SEM.

doi:10.1371/journal.pone.0047938.g002
4. The Native and Cysteinless Channels Share the Same Gating Characteristics

The cysteinless variant (C127A-C232A) displayed native-like channel behavior, occupying the open S0 state with a conductance of $3.78 \pm 0.09 \text{nS}$ ($n = 28$, $P = 0.42$), the S1 state with a conductance of $2.55 \pm 0.12 \text{nS}$ ($n = 24$, $P = 0.24$) and the S2 closed state with a conductance of $1.78 \pm 0.09 \text{nS}$ ($n = 12$, $P = 0.69$) (Table 1, Figure 3B). Note that the statistics provided compare the observed conductance states of the cysteinless variant with the corresponding conductance states of the native protein. The sample sizes for the native protein were: S0 ($n = 680$), S1 ($n = 386$) and S2 ($n = 294$). Representative traces of the cysteinless channel are provided (Figure 3B). As expected, the behavior of the cysteinless variant was redox independent, since neither CuX$_2$Ph nor DTT treatment affected its gating characteristics (Figure 3B).

The average dwell times of both native ($10.34 \pm 1.65 \text{s}$, $n = 310$) and cysteinless ($10.45 \pm 5.06 \text{s}$, $n = 37$) constructs were similar ($P = 0.98$) (Figure 3S; Note that the average dwell times indicate the lingering in any state before transition to the next. Consequently, average dwell times correspond to switching events from and to all states). Furthermore both channels dwelled at the individual states for the same lengths of time (S0; $P = 0.66$, S1; $P = 0.87$ and S2; $P = 0.94$, Figure 4). Accordingly, any putative effects in the gating behavior of further modified mVDAC1 variants can only be attributed to the incorporation of the corresponding mutations and the disulfide bond-mediated immobilization of the N-terminal segment.

5. Disulfide Bond Formation in the A14C-S193C-mVDAC1 Double Mutant Favors Occupancy of the S0 and S2 States

As expected, fully reduced A14C-S193C-mVDAC1 displayed native-like channel characteristics alternating between a $3.75 \pm 0.12 \text{nS}$ high (S0) and a $1.70 \pm 0.1 \text{nS}$ low (S2) conductance states (Figure 5A, Table 1). The observed state conductances were similar to those of the native protein (S0; $P = 0.32$, S1; $P = 0.22$ and S2; $P = 0.30$. Sample sizes were $n = 30$ for every reduced-mutant state. Sample sizes for the native protein were as above). Disulfide cross-linking by CuX$_2$Ph after reconstitution of this as well as other variants in planar lipid bilayers failed, resulting in native-like channel recordings. However, pre-oxidation and subsequent voltage-driven lipid reconstitution resulted in current recordings supporting almost a lock in the S0 conductive state ($3.74 \pm 0.09 \text{nS}$; $n = 30$) (Figures 5A, 5B). The attained S0 conductance of the oxidized mutant was similar to that of the native protein ($P = 0.30$). In the minority of collected traces (30/358), and always following extended lingering in the S0 state, the channel finally relaxed to a low conductance state ($1.83 \pm 0.07 \text{nS}$), and a conductance of $2.55 \pm 0.12 \text{nS}$ (S1) ($P = 0.30$) (S2).

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<th>Salt composition</th>
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<tr>
<td>mVDAC1-A14C-C127A-S193C-C232A (oxidized)</td>
<td>This work</td>
<td>3.74±0.09</td>
<td>1.85±0.07</td>
<td>1 M KCl 5 mM CaCl$_2$</td>
</tr>
<tr>
<td>mVDAC1-A14C-C127A-S193C-C232A (reduced)</td>
<td>3.75±0.12</td>
<td>2.55±0.12</td>
<td>1.67±0.09</td>
<td></td>
</tr>
<tr>
<td>mVDAC1-V3C-K119C-C127A-C232A (oxidized)</td>
<td>3.76±0.10</td>
<td>1.77±0.09</td>
<td></td>
<td></td>
</tr>
<tr>
<td>mVDAC1-V3C-K119C-C127A-C232A (reduced)</td>
<td>3.98±0.04</td>
<td>~ 2.70</td>
<td>2.05±0.13</td>
<td></td>
</tr>
</tbody>
</table>

Table 1. Reported conductance values for VDACs following their reconstitution in planar lipids.

*Values given with their standard deviations; *b values given with their standard error; ND, not determined.

doi:10.1371/journal.pone.0047938.t001
n = 14), equivalent to S2 of native mVDAC1 (P = 0.86), omitting the S1 state and failing to reopen (Figure 5B). Furthermore, the average dwell time for the oxidized A14C-S193C variant in the S0 state (30.61 ± 6.39 s, n = 14) was 180 times longer (P < 10^-4) in comparison to that of the native mVDAC1 (0.17 ± 0.03 min = 10.34 ± 1.65 s, n = 310) (Figures 5C, S3).

### Figure 4. Dwell times for the individually attained conducting states by native and cysteinless mVDAC1 channels. S0, S1 and S2 conducting states were identified from collected traces, based on the values of Table 1 (±5%).

A) Native mVDAC1 occupied the S0 state (27.40 ± 6.76 s, n = 137) almost 10-fold longer (P = 0.0053) than the S1 (2.92 ± 1.34 s, n = 92) and S2 (2.45 ± 1.30 s, n = 81) states. Occupancy of both S1 and S2 was similar (P = 0.80). B) Cysteinless mVDAC1 occupied the S0 state (36.70 ± 4.41 s, n = 14) also ~10-fold longer (P < 10^-5) than the S1 (3.61 ± 1.34 s, n = 10) and S2 (3.10 ± 1.36 s, n = 13) states. Occupancy of both S1 and S2 was similar (P = 0.80). State-to-state dwell time comparisons between native and cysteinless channels showed no differences (S0; P = 0.66, S1; P = 0.87 and S2; P = 0.84). Values are given as mean ± SEM.

doi:10.1371/journal.pone.0047938.g004
A14C-S193C channel. The open state (blue squares) conducted the conductances of the observed states in the gating transitions of the (8% of the collected traces). B) Ohm-plot used for the determination of traces) with relaxation in the closed S2 state being strongly disfavored however, appeared to remain constitutively open (92% of the collected oxidized mutant exhibited an open state S0 with DTT and subsequently reconstituted A14C-S193C mVDAC1 ex- 
hibited the expected native-like channel gating. The oxidized mutant resulted in current recordings which showed channel opening under both positive and negative potentials (306 traces), but closure only under one or the other (300 traces) (Figures 6, 7A and 7B). Pre-reduction of the disulfide bond restored native gating characteristics (Figure 7A) with: S0 at 3.75±0.12 nS (n=30, P=0.32), S1 at 2.55±0.12 nS (n=20, P=0.24) and S2 at 1.67±0.09 nS (n=20, P=0.32), where P provides the strength of similarity to the corresponding conductance states of the native protein (native channel conductances - Table 1 - and sample sizes were as above).

The type of potential required for the cross-linked channel's closure was determined by the orientation of the protein within the BLM set-up. Thus, channels incorporating into the bilayer upon application of a positive potential conducted in the open state, and showed decreased voltage-dependent conductance (closed state) upon voltage reversal (Figure 7B). Likewise, channels incorporating into the bilayer at negative potentials conducted in the open state until the opposite voltage was applied (Figure 7B). The first disulfide-bridged V3C-K119C channel assumed a random orientation upon its reconstitution into the lipidic environment of the BLM set-up. Random channel orientation, following the reconstitution of membrane proteins into artificial bilayers, has been reported for the outer membrane porin G from E. coli [29], yeast VDAC [30], bacterial sodium channels [31] and pore forming peptides [32]. Though the biophysical details of these random insertions have not been fully resolved, it appears that depending on the lipid composition [30] and when protein is applied to the cis compartment (high salt concentration), a slightly more positive voltage on the cis side enables primary adsorption of the protein to the bilayer. Reconstitution is then completed when a full trans-negative voltage is applied. Subsequently reconstituted mVDAC1 channels (up to five), exhibited auto-directed insertion, in harmony with earlier observations [30,33,34,35], probably due to a re-duction of the energy barrier to subsequent insertions. This autodirected insertion ensured that the oxidized channels would not reconstitute in an anti-parallel manner. Had that been the case, one of the two would have been open in one voltage sign and the other in the opposite, a situation which would create the illusion of symmetrical voltage-dependent conductance changes. The oxidized A14C-S193C variant exhibited an open state S0 conductance of 3.76±0.10 nS (n=77), similar to that of the native channel (P=0.15; S0 conductance and sample size for the native protein were as above) (Figure 7B). The most-attained low-energy closed state (1.77±0.09 nS, n=30) was similar to the S2 state of the native channel (P=0.49; S2 conductance and sample size for the native protein were as above) (Figure 7B).

Restraining the putative mobility of the first three amino acids, introduced upon potential reversal a “slow-motion” effect in the
channel’s gating behavior, with open-to-closed state transitions extending the channel’s average dwell time to 78.58±3.99 s (n = 8) with respect to the native protein (10.34±1.65 s, n = 310) (P<10⁻²⁴) (Figure S3). In 95% of the recorded events, the S1 state was avoided and S0-to-S2 transitions involved additional intermediates, with conductances exhibiting ≥5% difference to S0 and S1 conductances. S0-to-S1 transitions occurred only in ≥5% of the events and were independent of S2. The dwell-times of the

Figure 6. Cross-linking at the base of the pore abolishes symmetric voltage gating. CuX₂Ph pre-treated and subsequently reconstituted V3C-K119C-mVDAC1 conducted in the open state upon application of both negative and positive potentials (e.g. −40 mV) but closed only at one set of potentials (e.g. +40 mV), depending on its orientation in the lipid layer. As indicated in the corresponding traces (one step closing, two-step opening) and the adjacent cartoons, the gating behavior was dynamic in both directions, and was probably mediated by a series of short-lived intermediates (Note that the conductances of these intermediates exhibited ≥5% difference with respect to the conductances of the S0 and S1 states). The probable movement of the 5–14 residue part is sketched (dashed blue ellipticity). Cross-linking at the base of the channel is designated –S–S–. Polarity of the membrane is indicated.

doi:10.1371/journal.pone.0047938.g006
A

V3C-K119C ± 40 mV reduced

V3C-K119C - 40 mV oxidized

V3C-K119C + 40 mV oxidized

B

![Graph showing current (pA) vs. voltage (mV)]

C

![Bar graph showing dwell time (s) for different potentials]
channel in each one of the observed states are provided (Figure 7C). The cross-linked V3C-K119C variant, for a chosen initial potential (−40 mV), assumes the S0 state with a dwell time of 78.58±3.99 s (n=8) (Figure 7C). Note that for the transition to the next state the potential has to be reversed otherwise the cross-linked mutant channel would remain open. With the reversal (+40 mV), the channel lingers in the S0 state for 5.74±1.69 s (n=17) before switching to the S2 state (Figures 7A, 7C). The channel assumes the S2 state for 2.55±0.73 s (n=7), before another transition to S0 occurs (Figures 7A, 7C). To date, this is the first VDAC1 mutant to assume non-symmetrical voltage responses with both open-to-closed and closed-to-open transitions clearly occurring through a series of low-conducting sub states.

Discussion

VDACs have been the subject of extensive studies and lively discussions concerning their structural organization and gating mechanism [1,11]. The recently published mVDAC1 and hVDAC1 3D-structures fueled further debates [36,37] because they revealed a) a number of β-strands atypical for porins b) the formation of two parallel β-strands between β19 and β1 [3,6,7] and c) an alignment of the N-terminal α-helix with the inner pore wall [3] or at least a localization in the pore’s interior [7]. VDACs are capable of alternating between high and low conductance states, representing open and closed channel conformations respectively [2,3,27,38,39,40]. Based on past studies, it has been argued that crystal structures represent the open state of the pore while the NMR structure represented the closed state [8]. Brownian calculations and molecular dynamics simulations have revealed the dependence of the channel’s ion selectivity on salt concentration, with low salt concentrations favoring the permeability of anions in the open state [41,42,43,44]. The mechanistic details underlying the channel’s gating behavior and ion selectivity however, still remain largely unclear.

In this work, the 2EMN crystal structure [3] was used as a template for the design and construction of mVDAC1 double cysteine comprising variants. Formation of disulfide bridges in all our tested variants (Figure 1, Figure S1) attested the proposed 3D-model. Colombini disputed the organization of mVDAC1 in detergent-micelles, and consequently in the crystal structure, proposing instead a refolding mechanism involving a 19- to 13-strand transition [36]. This transition, however, should disfavor topologically the state alterations of our cross linked mVDAC1 variants. Nevertheless these variants exhibit gating, characteristic of S0-to-S2 transitions, albeit with lowered probability. In harmony with previous BLM and patch-clamp measurements [3,4,7,24,25,26,27,43,45] these results provide further support to the arguments of Hiller and co-workers that the 19-stranded 3D-structure represents indeed the biologically relevant open state [37].

To address the involvement of the N-terminal segment in the channel’s gating behavior we characterized the Δ21-mVDAC1 deletion variant. In agreement with previous observations [23,28], our mVDAC1 deletion variant produced current recordings characterized by some flickering. Popp et al. [25], reported a preference of the Δ3–20-mVDAC1 for a closed state and, since Δ2–12 exhibited native-channel characteristics, proposed a role for the N-terminal fragment in stabilization of the open state. Our data, however, indicated a clear arrest of the Δ21-mVDAC1 in the open state (Figure 3A), sharing similar characteristics with the Δ26-mVDAC1 deletion construct described by Abu-Hamam et al. [4]. An involvement thus of the N-terminal fragment in the maintenance of the overall integrity of the incorporated channel is highly plausible, as also demonstrated by NMR data [46], and cannot be restricted to stabilizing effects on the open conformation only. We suggest that flickering reflects rapid transitions between different sub-states of the channel, as we have observed noise reduction in bacterial porins engineered with a shorter-loop (unpublished data). Consequently, the observed flickering/noise may reflect complexities not solely attributed to the mobility of the N-terminal fragment, but also to other structural components of mVDAC1.

Under our BLM conditions, reconstituted mVDAC1 exhibited a 3.94±0.04 nS conductance in the open state (S0), consistent with previous observations, whereas the average conductance of the closed state was 2.34±0.06 nS, differing significantly from previously published values (Table 1) [3,27,39,41,47]. Instead the closed state of mVDAC1 shows a broad distribution between two already previously recognized but not defined major states (S1, S2) [48]. In addition we find that the S2 state is itself split into two further sub-states S2A and S2B in a 2:4:1 ratio. S2B exhibits a 1.48±0.02 nS conductance almost half of that of the S1 state (Figure 2C). Defining these main states and their transitions is essential not only for deciphering the channel’s functionality, but also in order to acquire a platform for the characterization of VDAC variants with potentially suble gating phenotypes. The latter may become pivotal when considering the VDAC-dependent mitochondrial metabolic transport.

Recently, Cheneke, van den Berg and Movileanu [49] analyzed the gating transitions of the three major open states of the OepK channel from Pseudomonas aeruginosa and reported that the conformational fluctuations of the low amplitude current transitions stemmed from the highly flexible loop 7 occluding the pore. In VDACs however, the situation is less straightforward. Firstly, no major loop appears as a prominent candidate for channel occlusion, and secondly, although the N-terminal α-helix appears position-wise to exert control on the flux of metabolites [50], the latest NMR studies [51] revealed distinct mobility only for strands
β2-β7. Here, the N-terminal helix appeared relatively rigid, raising doubts as to its involvement in controlling gating. Interestingly, and following our observations, the observed low amplitude current transitions of OpdK [49] mimic the mVDAC1 S1→S2 transitions.

To further investigate the potential role of the N-terminal segment in voltage gating, we engineered and characterized double-cysteine mVDAC1 variants. Our cross-linked mVDAC1-A14C-S193C channel exhibited a clear preference for the open state (average dwell time 30.61±6.39 min = 1836.60±372.91 s), attaining hence the S2 state with a strongly reduced probability (Figures 5B, 5C and S3). Having achieved the closed state, this variant failed to re-open. The oxidized and subsequently reconstituted mVDAC1-V3C-K119C variant, moreover, exhibited asymmetric gating characteristics (Figure 6), conducting in the open state at both negative and positive potentials, and closing only upon application of the reverse potential, depending on its orientation in the lipid bilayer (Figures 7A, 7B). Surprisingly, transitions between open and closed states were mediated through a series of conducting conformers, enabling final relaxation to an S2 conductance state. Open-to-closed state transitions occurred more slowly than the wild type, extending the average dwell time of the cross-linked channel in the S0 state to 78.58±2.99 s (Figures 7C, S3). A complete switch to native-like channel behavior was achieved following reduction of the cysteines of both variants by DTT (Figures 5A, 7A).

The conductance behavior of the cross-linked mVDAC1 variants shows that the choice of residues for covalent linking is crucial for 1) observing unique gating phenomenotypes and 2) achieving different immobilization degrees of the N-terminal α-helix. This is clearly reflected in the intensity of the gating phenotype of the mVDAC1-A14C-S193C variant. Affixing residue 14 on the barrel wall immobilizes the N-terminal fragment more effectively than affixing residue 3, almost locking the channel in its open conformation. Yet, the remaining flexibility of the mVDAC1-A14C-S193C channel, still allowed for an unfavourable and permanent closure to the S2 state. The oxidized mVDAC1-V3C-K119C variant on the other hand, although having lost its symmetrical gating, exhibited reversible S0-to-S2 transitions. In consequence, these findings suggest that the full flexibility of the N-terminal fragment is required in achieving transitions to and/or from the S1 state, and that S0-to-S2 can happen independently of S0-to-S1 transitions, probably through meta-stable intermediates of an alternative route (Figure 6). Latest studies identified an intrinsic asymmetric effect on the closure of VDAC depending on the sign of applied voltage [45]. This finding points to the existence of alternate mechanisms for channel closure. Although further experimentation is required to decipher the role of these events, it seems that the S0-to-S1 and S0-to-S2 transitions may represent separate closing modes, as hypothesized by Hiller and Wagner, and that S1 is not just an intermediate before complete relaxation of the channel in the S2 state.

What information can be extracted from the closure of the cross-linked channels? With respect to the A14C-S193C-mVDAC1 channel, fixing residue 14 to the porin’s wall allows for mobility of residues 1–13 and the hinge region (KGFGYG). The arrest of the open state in this variant for ~30 min (Figure 5C) indicates that residue 14 and the stretches around it participate in native channel opening. Nevertheless, the hinge region remains free to undergo conformational changes, finally causing the channel’s closure. In doing so, it probably exerts mechanical strain on the relatively immobile α-helical part. We hypothesize that closure of this cross-linked channel is mediated by mechanical strain effects, stemming from the movement of the hinge region, which can further induce either a) the structural collapse of the barrel wall b) the irreversible unfolding of the N-terminal helix or c) a combination of the two. All scenarios could result in permanent channel closure as observed for the oxidized mutant.

In the V3C-K119C variant, where cross-linking at residue 3 allows for more N-terminal flexibility (extending from residue 4 to the hinge region), voltage gating lacks the symmetrical features of the native channel’s activity. This means that this cross-linked variant, although capable of conducting in the open state at both positive and negative potentials, closes only upon application of one. This indicates that only one direction of the applied electric field destabilizes the tethered N-terminal helix. Apparently, immobilization and/or reduced flexibility of the N-terminal segment make it less prone to unfolding by an applied electric field. Furthermore, the behavior of this channel upon potential reversal suggests that the residue 4–26 stretch still undergoes structural reorganizations allowing for slower S0-to-S2 transitions (Figures 6, 7C and S3). Interestingly, these structural changes can be reversed with the next reversal of applied potential. Thus, in contrast to the irreversible structural changes observed in the oxidized A14C-S193C variant, partial immobilization of the far N-terminal part allows for a voltage-driven reversible reorganization/refolding of the α-helix. Current studies [52] suggest that VDACs assume their open state upon N-terminal fragment detachment from the pore wall and subsequent translocation to the cytoplasmic side (“paddle” movement). The behavior of our cross-linked variants however indicated that, although a structural reorganization of the 1–14 residue segment is required for voltage gating, the so-called “paddle” movement is not required. Consequently, we propose that closure is mediated by the concurrence of 1) a flipping motion of the far N-terminal part into the middle of the channel and 2) a local unfolding/refolding event of the α-helix (Figure 8).

Recently, Teijido et al. [11] reported that the L10C-C127A-A170C-C232A-mVDAC1 mutant displayed native-like channel activity under oxidative conditions, and concluded that voltage gating did not involve and/or require rearrangements of the covalently linked N-terminal fragment. Contrary to that, both our mVDAC1 double cysteine variants showed under oxidative conditions distinct gating profiles, avoiding the S1 and relaxing only at the S2 state. The reasons for the observed deviant behavior of the L10C-A170C-mVDAC1 variant can be manifold. First, in situ oxidation as performed in [11] failed in our hands under the chosen BLM conditions and therefore required the usage of in vitro oxidized and purified double-cysteine mVDAC1 variants. Accordingly, proper disulfide bond formation in the used samples was verified by ESI-MS. Secondly, we only characterized mVDAC1 variants without a charged N-terminal affinity tag, thus avoiding possible gating interference. Finally, we restricted the number of single-channel incorporations to a minimum (≤5). Last but not least, and as previously indicated, the choice of disulfide-mediated fixing points is crucial for gating behavior and formation of arrested channels. For example, failure to completely immobilize the N-terminal helix results in native-channel behavior, whereas forcing the outward placement of the N-terminal region by generating cross-linked mVDAC1 homodimers via their N-termini [32] arrests gating like a deletion of the N-terminal helix itself.

As previously shown, channel closure provides a regulatory mechanism for control of substrate and/or ion efflux. In the SEMN crystal structure, salt bridges engaging residues K12-D16- K20 are evident. Recent studies have identified K12 and K20 to be important selectivity determinants. Toorroth-Horsefield and Neutze [53], proposed that even a modest rotation of the N-
Figure 8. Native channel voltage gating involves the concurrent movement of the very N-terminus into the pore and induction of channel narrowing by partial wall collapse. A) Cross section of a 3D-model of mVDAC1, based on the 3EMN structure [3], parallel to the membrane plane. Cross-linking at the base and midpoint of the pore is indicated (yellow circled asterisk). Possible dynamic movements of the α-helix and the 1–4 residue N-terminal part are described for the two cross-linked mutants relative to the native channel. In the native channel, the N-terminal fragment is capable of both a flip and a twist/turn movement. The concurrence of these movements together with a constriction of diameter of the pore, close the channel. Cross-linking at the bottom (V3C-K119C) or the middle (A14C-S193C) of the pore, prevent the flip and twist/turn motions of the N-terminal fragment respectively. Thin arrows indicate possible directions of movement. Thick arrows indicate potential tightening of the pore diameter. The asterisk denotes movement contributions from the hinge region. Note that with the proposed orientation both N- and C- termini are cytoplasmic (cis) in accordance with McDonald et al. [56]. IMS; intra-membrane space (trans). B) N-terminal fragment movements
in the context of the observed closed state conformations. In the open S0 state, the helix is located at the mid-point of the pore with the far N-terminus facing the cytoplasm (ci). Channel closure (S1 state) is initiated upon movement of the far N-terminus into the pore and is completed by the constricting movements of the pore’s wall. Oxidized mutations (yellow circled asterisk) become immunobound in an S2A state, exhibiting restricted N-terminal fragment flexibility and impede barrel wall movement. In the S2B state movements of both the barrel wall and the N-terminus complete channel closure. Effectors could control S1-to-S2 transitions enabling or inhibiting interactions with substrates thus regulating apoptotic responses. Footnote: While this manuscript was under review, Zachariae et al. [57] reported that the closure of voltage-dependent anion channels is largely regulated by the mobility of the β-barrel scaffold. These authors note however, that the mobility/flexibility of the N-terminal domain, albeit lower than that of the β-barrel, is also essential in shaping the channel’s gating characteristics. They also suggest that gating does not require the exit of the N-terminus from the inside of the channel, and show that minor structural reorganizations of the N-terminal α-helix, whilst still inside the pore, can have noticeable effects on the channel’s conducting capacity. Their observations are also in accordance with our proposed model of VDAC voltage-induced gating, where the conformational changes of the N-terminal fragment, in conjunction with dynamic movements of the barrel walls define transitions between specific conducting states. In essence, both works demonstrate the importance of the N-terminal fragment in a) the preservation of the overall integrity of the channel and b) the regulation of transitions/entry into states which, in combination with the dynamic movements of the barrel, eventually define the channel’s gating behavior.

doi:10.1371/journal.pone.0047938.g008

Supporting Information

Figure S1 Confirmation of disulphide formation in the A14C-S193C-mVDAC1 variant by mass spectrometry. MALDI-MS tryptic peptide mass fingerprint of A14C-S193C-mVDAC1. The peaks of the obtained chromatograms reveal the ionization profile of two cross linked fragments for the 993.0–995.4 and 1489–1493 mass range (m/z). Detected masses for the +3 and +2 ionization states are labelled (red and green) and are compared to the expected masses in the table below. Calculations were performed using the Peptide Mass Calculator v3.2 (Jef Rozenski, 1999).

(TIF)

Figure S2 Oxidation of reduced A14C-S193C-mVDAC1 over time. A) Non reducing SDS-PAGE profile of DTT (10 mM) treated A14C-S193C-mVDAC1. Following the removal of the reducing agent, protein aliquots (2 μg) were taken over a period of 8 days, mixed with SDS-PAGE sample buffer without β-mercaptoethanol and heated at 95°C (5 min) prior to 12% SDS-PAGE electrophoresis. Spontaneous oxidation of cysteines was visible (increased electrophoretic mobility-Ox.) after the first day. Red.; reduced. B) Percentage decrease of the fully-reduced A14C-S193C-mVDAC1 population with time. Percentages were determined from the densitometric analysis of the SDS-PAGE in (A). The half life of the reduced species was ~22 hours.

(TIF)

Figure S3 Average dwell times of native and engineered mVDAC1 variants. Native (n = 310) and cysteineless (n = 37) channels exhibited similar dwell times (P = 0.90) before making a transition to any state, whereas the dwell times of the oxidised cysteine-engineered variants (A14C-S193C; n = 14, V3C-K119C-initial potential; n = 8) were significantly extended with respect to the native protein (P<10−4 and P<10−4 respectively). Note that after potential reversal the capacity of the oxidized-V3C-K119C channel (n = 24) to alternate between individual states mimics that of the native channel (P = 0.60).

(TIF)

Table S1 Analysis of amplitude histograms fitted by Gaussian functions.

(DOC)

Acknowledgments

The authors wish to thank Dr Jeff Abramson for kindly providing us with the mVDAC1-harboring plasmid, Dr Uwe Linne for mass spectrometric analyses, Dr Manuel Maestre-Reyna for helpful discussions and Dr Sophia-Louisa Tsougarakis for the line editing of this manuscript.

Author Contributions

Conceived and designed the experiments: BM WG LOE. Performed the experiments: BM GP KCB AS PR. Analyzed the data: BM GP PR. Contributed reagents/materials/analysis tools: BM GP WG KCB AS PR. Wrote the paper: BM GP UK LOE.