Channel Activity of a Viral Transmembrane Peptide in Micro-BLMs: Vpu_{1–32} from HIV-1

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Abstract: We report for the first time on pore-suspending lipid bilayers, which we call micro-black lipid membranes (micro-BLMs), based on a highly ordered macroporous silicon array. Micro-BLMs were established by first functionalizing the backside porous silicon surface with gold and then chemisorbing 1,2-dipalmitoyl-sn-glycero-3-phosphothioethanol followed by spreading 1,2-diphytanoyl-sn-glycero-3-phosphocholine dissolved in n-decane. Impedance spectroscopy revealed the formation of single lipid bilayers confirmed by a mean specific capacitance of 0.6 ± 0.2 μF/cm². Membrane resistances were in the GΩ-regime and beyond. The potential of the system for single channel recordings was demonstrated by inserting the transmembrane domain of the HIV-1 accessory peptide Vpu_{1–32}, which forms helix bundles with characteristic opening states. We elucidated different amilorides as potential drugs to inhibit channel activity of Vpu.

Introduction

Ion channels play key roles in the function of living cells, and as a consequence, there are many potential drug targets among them. Two main techniques are used to investigate ion channel activity: patch clamp experiments and recordings on planar lipid bilayers (black lipid membranes, BLMs), both allowing the study of ion channel behavior by recording the transmembrane current caused by opening and closing of the channels. From these recordings, single channel conductance as well as opening and closing probabilities can be deduced. While patch clamp experiments start from native membranes, planar bilayers are formed across apertures in thin films, often Teflon of 5 to 25 μm with hole diameters of 0.1–1 mm, with the advantage that the composition of the lipid membrane and the surrounding electrolyte solution on both sides of the membrane can be defined precisely. In particular, these planar lipid bilayers have been reassessed in recent years since ion channels have been identified as major drug targets. However classical BLMs, based on small apertures in thin Teflon films, are not suited for automation and miniaturization as required for biosensors and high-throughput screening assays for potential drugs. Thus, new strategies based on microfabricated apertures in silicon and glass have been recently developed, which allow for rather small apertures of 0.7 μm to a few millimeters to be generated. Lipid membranes suspended across 0.7–100 μm apertures have been prepared with various techniques ranging from depositing lipids in an organic solvent to spreading of a giant unilamellar vesicle. The long-term stability of these membranes is merely on the order of up to several hours. For biosensor applications, however, a long-term stability of several days is desirable. So far, this can only be achieved by solid support membranes, which enable the rational design

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of biosensor devices for high-throughput screening based on sophisticated micromachined chip technology. However, these membranes are not suited for single channel recordings, since first, the interface capacitance abolishes a continuous ion flow, and second, a GΩ seal between the bilayer and the support is hardly achieved. Moreover, solution exchange is not feasible since there is only a 1–5 nm water film that separates the bilayer from the support.

With the aim to combine the merits of both membrane systems, free-standing planar lipid bilayers and solid-supported membranes, we used macroporous silicon substrates with pore sizes of 1 μm that were covered with a lipid bilayer. In this hybrid membrane system, the membrane-suspended pores act as planar lipid bilayers, while the membranes attached to the pore rims provide mechanical stability as obtained for solidsupported membranes.

To elucidate the potential of this membrane type for investigating channel activity, a potential drug target for HIV treatment, Vpu, was investigated. Vpu is a small 81 amino acid integral membrane protein that forms cation-selective ion channels that is encoded by the human immunodeficiency virus type 1 (HIV-1). 18–20 Vpu comprises a highly hydrophobic N-terminal helical transmembrane (TM) domain encompassing residues 8–25 and a large hydrophilic phosphorylated cytoplasmatic C-terminal domain. 19,21–24 These two distinct structural domains of Vpu exhibit independent functionalities: while the latter domain is involved in the degradation of the virus receptor CD4 in the endoplasmic reticulum, the former domain enhances viral release/secretion from the cell surface. 21,22 Reconstitution of synthetic Vpu fragments in lipid bilayers identified ion channel activity for a sequence corresponding to the helical TM domain of Vpu, forming a water-filled bundle of homooligomers of an undetermined number of subunits in membranes. 22,25,26 Recent studies show that the cytoplasmatic domain encompassing two shorter amphipathic in-plane helices23,26 regulates the lifetime of the TM channel in the conductive state. 26 Amiloride derivatives have been shown to block channel activity and the enhancement of virus-like particle budding caused by Vpu and synthetic Vpu fragments in lipid bilayers identified ion channel activity, a potential drug target for HIV treatment.

Results

Characterization of the Macroporous Silicon Substrate.

Scanning electron microscopy (SEM) images of the front and backside as well as side views of macroporous silicon, prepared by photoelectrochemical etching,27 were taken. SEM images of the front side (Figure 1a) after the anodization procedure show square-shaped pores with an etch length of around 1.0–1.2 μm. The backside of the macroporous silicon substrate (Figure 1b) exhibits pores with diameters of 1.0 ± 0.1 μm. In contrast to the initial square-shaped photolithographically formed pore nuclei at the front side of the silicon wafer, the opened pores at the backside are circular and almost perfectly hexagonally ordered. Only very few defects are present. A surface porosity of 22 ± 3% was calculated by pixel analysis from the SEM images. A side view of a macroporous silicon substrate, where the backside of the pores has not yet been removed (Figure 1c), reveals that the pores grow perpendicular to the substrate surface. This almost perfect growth of each individual pore is a consequence not only of the lithographic pattern but also of the orientation of the silicon single crystal. Each pore grows with the same velocity, and thus, the thickness of the porous substrate is determined by the anodization time. Typically, a thickness of 50–200 μm is adjusted to ensure mechanical stability.

By means of impedance spectroscopy, the electrochemical characteristics of porous silicon substrates open to the electrolyte on both sides were investigated. Only an ohmic resistance of the electrolyte was detected with a small contribution from the platinum electrodes at frequencies below 1 Hz (see Supporting Information).
Impedance Analysis of Micro- and Porous Matrix-Supported BLMs. Two different preparation techniques were followed to form lipid bilayers suspending the highly ordered silicon macropore arrays with pore diameters of 1.0 \pm 0.1 \mu m:

(i) In a first approach, the backside of the porous silicon was first coated with a thin gold layer, which allows for the chemisorption of the phospholipid 1,2-dipalmitoyl-sn-glycero-3-phosphothioethanol (DPPTE) rendering the surface hydrophobic. Both, the gold coating and the chemisorption of DPPTE do not alter the impedance behavior corroborating the idea that the current flows only through the pores. Second, the phospholipid 1,2-diphytanoyl-sn-glycero-3-phosphocholine (DPhPC) dissolved in n-decane was spread across the functionalized porous matrix (see Figure 2b). The formation of a lipid bilayer was followed by means of impedance spectroscopy in a frequency regime of \(10^{-2} - 10^6\) Hz using the setup depicted in Figure 2a. For very large membrane resistances impedance measurements down to \(10^{-3}\) Hz were carried out (see Supporting Information). Electrical impedance spectroscopy enables one to obtain quantitative electrical parameters, which allow unambiguously distinguishing between single lipid bilayers and multilayers. Our measurements revealed the formation of single lipid bilayers consistent with a mean specific capacitance of \(0.6 \pm 0.2 \mu F/cm^2\). The bilayer formation process takes about 10–20 min. During this time period, the initially formed hydrophobic lipid droplet with a thickness of several micrometers thins out to form a planar lipid bilayer completing the self-assembled DPPTE submonolayer, which can be followed by the increase in capacitance leveling off at the characteristic specific capacitance of a single lipid bilayer (see Supporting Information). Due to the similarity of this hybrid system to classical black lipid membranes and the fact that the bilayer suspends pores with diameters of 1 \mu m, we termed these membranes micro-BLMs.

The membrane resistance \(R_m\) is determined in the very low-frequency regime. If \(R_m\) is larger than 10 G\(\Omega\), it is not detectable in the observed frequency regime and is classified as \(R_m > 10\) G\(\Omega\). Below 10 G\(\Omega\), the parameter \(R_m\) of the equivalent circuit shown in Figure S1a (Supporting Information) was fitted to the data with a relative fitting error of \(< 10\%\).

(ii) In a second approach suspended lipid bilayers were formed on macroporous silicon substrates without gold- and DPPTE-coating by simply spreading DPhPC in n-decane onto the porous matrix (see Figure 2c). Impedance spectra again clearly show the formation of an insulating lipid bilayer across the porous matrix characterized by a mean specific membrane capacitance of \((0.8 \pm 0.3) \mu F/cm^2\). Membrane resistances were again larger than 10 G\(\Omega\). This membrane type is termed matrix-supported BLM.

Long-Term Stability of Micro- and Matrix-Supported BLMs. The membrane capacitances and resistances achieved for micro-BLMs and matrix-supported BLMs based on silicon macropores resemble those obtained for classical BLMs, and the membrane resistances are sufficiently high to perform single channel recordings (see below). However, for ion channel biosensors, not only the electrical characteristics but also the long-term stability of these parameters are of major importance. Classical BLMs typically rupture in one single event, leading to a complete loss of membrane resistance and capacitance. To investigate the behavior of micro-BLMs, the time course of the membrane capacitance and resistance has been investigated by means of impedance spectroscopy (Figure 3). One hour after membrane formation, the micro-BLM exhibits a resistance of \(> 10\) G\(\Omega\). The value decreases over time, leading to a membrane resistance of 5 G\(\Omega\) after 24 h, 2 G\(\Omega\) after 50 h, and 1.5 G\(\Omega\) after 70 h. This indicates that only a limited area of the membrane suspending the pores ruptures, while other areas are still membrane-covered. This is corroborated by the capacitance measurements, which demonstrate that the capacitance of the micro-BLMs only slightly increases concomitant with the decrease in membrane resistance and the idea that the majority of the silicon substrate is then still covered by a membrane. In summary, a typical micro-BLM exhibits a membrane resistance of larger than 1 G\(\Omega\) for approximately 1 day. We simply defined 1 G\(\Omega\) as a threshold value, as this membrane resistance is sufficiently large to perform single channel recordings.
The time-dependent change in membrane resistance $R_m$ as a measure of long-term stability has also been investigated for matrix-supported BLMs. Although the achieved membrane resistances were in principle the same range as those obtained for micro-BLMs, a continuous time-dependent decrease in membrane resistance was not observed. Instead, after a certain time, a complete loss in membrane resistance was monitored, indicative of a complete rupture of the membrane. Such an event was observed during the recording of an impedance spectrum (Figure 4), indicated by the sudden decrease of impedance, leading to a spectrum displaying only the resistance of the buffer solution. To verify that the membrane was still intact prior to rupturing, part of the impedance spectrum that is characterized by the membrane capacitance $C_m$ and the electrolyte solution ($10^6-10^9$ Hz) was taken to extract the exact value of the specific membrane capacitance. A value of 1.05 $\mu$F/cm$^2$ was obtained, confirming the quality of the membrane before it ruptured. In general, membrane resistances for matrix-supported BLMs suited for single channel recordings were found within a broad time interval up to several hours.

In conclusion, only micro-BLMs exhibit the desired electrical and mechanical long-term stability for biosensor applications.

**Channel Activity of Synthetic Vpu$_{1-32}$ Peptide.** Micro-BLMs and matrix-supported BLMs, respectively, were subjected to an ion channel activity study of purified synthetic Vpu$_{1-32}$ comprising the transmembrane segment of Vpu. Micro-BLMs were utilized and the peptide dissolved in trifluoroethanol was added to the buffer solution of the cis-chamber after a stable baseline was monitored. Channel events were mostly observed shortly after peptide addition, indicating its insertion into the micro-BLM. Channel activity was observed for more than 15 individual bilayer preparations. The most frequently observed single channel openings of synthetic Vpu$_{1-32}$ displayed a conductance of $12 \pm 3$ pS in 5 mM Hepes, 0.5 M KCl, pH 7.4 and a less frequently observed one of $28 \pm 3$ pS (Figure 5a,b). Surprisingly, even 3 h after the first channel openings had been observed, distinct rectangular-shaped single channel events were discernible, with the most frequently observed at $30 \pm 2$ pS and the less frequently observed at $12 \pm 2$ pS conductance state. Figure 5c,d shows representative opening states monitored around 1 h after the first channel events were observed.

Besides micro-BLMs, matrix-supported lipid bilayers were also used for single channel recordings to investigate the influence of the chemisorbed DPPTE monolayer of the micro-BLMs on Vpu channel activity. Again, peptide was added to the buffer solution, and channel events were detected shortly after peptide addition. First, current traces were observed, which indicate a single channel undergoing transitions between a closed (C) and two open (O) states at a holding potential of 50 mV (Figure 6a,b). Two discrete channel openings with a mean conductance of $13 \pm 4$ pS and $25 \pm 2$ pS, respectively, were observed occurring initially with almost equal frequency, but were considerably less frequently observed as the closed state. In contrast to the single channel recordings obtained for micro-BLMs, in case of porous matrix-supported BLMs already after a short incubation time, single channel events could not be resolved unambiguously anymore, but rather noisy current traces were recorded (Figure 6c).

**Inhibition of Channel Activity by Amiloride Derivatives.** It has been shown that amiloride derivatives depress ion channel activity of full-length Vpu reconstituted in classical BLMs. The binding site for the inhibitors has been suggested to be located in the N-terminal transmembrane domain of Vpu. To prove if micro-BLMs are suited for the investigation of inhibitors of viral peptide channel activity, the influence of 5-(N,N-hexamethylene)amiloride (HMA), 5-(N,N-dimethyl)amiloride (DMA), and amiloride on the channel activity of Vpu reconstituted in micro-BLMs was investigated. The addition of an aqueous solution of HMA to the cis- and trans-chambers with a final concentration of 100 $\mu$M in each chamber resulted in

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complete inhibition of Vpu<sub>1-32</sub> single channel activity after approximately 10 min (see Supporting Information). Inhibition of single channel activity was also found for DMA within 10 min, while amiloride itself with a final concentration of 100 µM in 5 mM Hepes, 0.5 M NaCl, pH 7.4 and a holding potential of 50 mV did not inhibit the channel activity of Vpu.
Discussion

This work describes for the first time the development of micro-BLMs as a system well suited to the study of single ion channels formed by the N-terminal transmembrane domains of the viral protein Vpu from HIV-1. Impedance analysis of the membranes unambiguously proved the formation of single lipid bilayers consistent with the mean specific capacitance of 0.6 ± 0.2 μF/cm² that was also found for classical BLMs and nano-BLMs.2,29,30 To observe single channel events with low noise, high membrane resistances in the GΩ regime are required. Micro-BLMs exhibit membrane resistances larger than 1 GΩ, which are similar to those obtained for classical BLMs2 and much larger than those typically obtained for solid-supported membranes.2,31

To establish a membrane system suited for biosensor devices and high-throughput screening systems based on single channel recordings, not only the membrane resistance itself but also a long-term stability of the membrane is required. The lifetime of a micro-BLM was defined here as the time period over which the membrane resistance exceeds 1 GΩ. On the basis of this definition, a mean lifetime of around 1 day was obtained. The most striking difference between classical BLMs, porous matrix-supported BLMs, and micro-BLMs was the finding that in the case of micro-BLMs the membrane resistance decreases with time. For conventional BLMs,2,31 spanning a small hole in a thin Teflon film, and for those bilayers more recently described and formed on individual apertures manufactured in silicon or glass supports,9,12 the membrane always ruptures in one single step, which can be deduced from the measured low resistance of several hundred Ω. Stabilities of conventional BLMs with pore diameters of 0.1–1 mm are reported to be typically in the range of several hours.2 Schmidt et al.9 reported a membrane suspending a 0.6–7 μm hole in silicon, which ruptures approximately after 1 h, whereas Cheng et al.12 obtained a stability of 5 h for a lipid bilayer suspended across an orifice of 128 μm in diameter before it completely ruptured. The notion that micro-BLMs do not rupture in one single step indicates that only membrane patches rupture that do not influence the stability of the entire membrane. This significantly different behavior of the membrane resistance of micro-BLMs and porous matrix-supported BLMs led us to conclude that the immobilization of the lipid bilayer on the porous silicon matrix via DPPTE chemisorbed on the pore rims decouples the free-standing membranes from each other.

Despite this difference in long-term stability and rupture mechanism, both membrane types, micro-BLMs as well as porous matrix-supported BLMs, allow in principle single channel recordings. Here, the suitability of micro-BLMs has been demonstrated by investigating the channel activity of the synthetic peptide Vpu. Over the past years it has been established that the transmembrane α-helix (residues 8–25) of the Vpu polypeptide exhibits ion channel activity. All that data were acquired from single channel recordings on classical planar lipid bilayers after reconstitution of full-length Vpu or truncated versions (Vpu2–27, Vpu2–30, Vpu2–37, Vpu2–51) of the protein.18,26,32,33 By means of the newly established micro-BLM system, single and multichannel openings of heterogeneous conductance in a highly resolved manner were recorded. The most frequent conductance state of 12 pS in 5 mM Hepes, 0.5 M KCl, pH 7.4 was largely observed as distinct single channel openings similar to the observation of Ma et al.33 They reported a single channel conductance of 12 ± 2 pS as the most frequently observed opening for Vpu2–37 under similar conditions using classical BLMs. The conductance level of 12 pS is also consistent with experiments carried out by Marassi et al.18 using full length Vpu and Vpu2–51 and Ewart et al.22 using Vpu constructs including the transmembrane domain. The observed 12 pS conductance state is also observed for full-length Vpu, with the most frequently occurring conductance of 12, 22,18,26 and 30 pS recorded in 0.5 M KCl. The finding that the second single channel conductance of 28 pS becomes more prominent after some time is consistent with the observations of Schubert et al. (Vpu2–27)31 and Marassi et al.18 using full length Vpu and Vpu2–51 and Kochendoerfer et al.20 (full length Vpu).

The results obtained for Vpu channel activity highlight that micro-BLMs are suited for single channel studies. They show advantages over classical BLMs, lipid bilayers suspending micromachined apertures, and porous matrix-supported BLMs. Using micro-BLMs, different single channel conductance states were resolved even 3 h after peptide had been added to the aqueous phase. In contrast, in the case of porous matrix-supported BLMs, single channel events could be observed only within the first 10–20 min. Already 10–20 min after peptide addition, large current bursts become visible, whose magnitudes increase over time presumably due to an increased number of inserted Vpu peptides. Such large conductance bursts of up to 430 pS were also reported by Ewart et al.22,24 using classical BLMs and were attributed to many simultaneously opened channels or the association of a large number of Vpu monomers to large channel aggregates, respectively. From our results we hypothesize that the chemisorbed DPPTE monolayer anchoring part of the micro-BLM to the support might reduce the lateral mobility of the lipids and thus influences the lateral mobility of incorporated Vpu monomers. This would result in a slower rate of assembly of large oligomeric bundles of Vpu monomers, and as a consequence, multichannel bursts due to oligomerization by large conducting channel aggregates as observed in matrix-supported BLMs become less probable within the observed time period. In addition, it is also conceivable that the incorporation probability of peptides is reduced due to the DPPTE-chemisorbed hybrid lipid bilayer, also resulting in less frequently observed multichannel bursts.

Although the hindrance of large channels is particularly advantageous for the investigation of Vpu, this phenomenon may be disadvantageous for other diffusion-controlled channel-forming compounds. If the insertion probability is less and the lateral mobility of the lipids and channel-forming compounds is reduced, this would disfavor the channel formation process, and hence, the observation of single channel events becomes less likely in micro-BLMs, while it might still be observable in classical or matrix-supported BLMs.

References

Experimental Section

Materials. (100) n-type silicon wafers with a resistivity of 0.5 Ωcm were supplied from Wacker (Germany). Hydrofluoric acid p.a. (40%) was purchased from Merck (Darmstadt, Germany). 1,2-Di(2-hexyloxy-alkyl)-3,4-epoxycyclohexane (DPhPC) and 1,2-dilinoyl-sn-glycero-3-phosphatidylethanolamine (DPPE) were obtained from Avanti Polar Lipids (Alabaster, AL). The peptide Vpu1 were supplied from Wacker (Germany). Hydrofluoric acid p.a. (40%) (Alabaster, AL). The peptide Vpu1 and phosphothioethanol (DPPTE) were obtained from Avanti Polar Lipids (Alabaster, AL). The peptide Vpu1 was synthesized on a Pioneer Synthesizer from Applied Biosystems (Foster City, CA) in capacitive feedback configuration via Ag/AgCl electrodes. The electrode of the cis compartment is connected to ground. Single channel currents were all recorded at an applied voltage of 50 mV. Data were filtered at 1 kHz with an 8-pole Bessel filter, and the analogue output signal was digitized at a sampling rate of 5 kHz by using an A/D converter (Digidata 1322A, Axon Instruments). Data processing is performed using pClamp 8.0 software (Axon Instruments). To avoid electrostatic interference during measurements, the electrochemical cell is placed in a Faraday cage set on a mechanically isolated support.30

Single Channel Recordings. Channel recordings are conducted in the Teflon cell (Figure 2a). Electrical currents are recorded with an Axopatch 200B patch clamp amplifier (Axon Instruments, Foster City, CA) in capacitive feedback configuration via Ag/AgCl electrodes. The electrode of the cis compartment is connected to ground. Single channel currents were all recorded at an applied voltage of 50 mV. Data were filtered at 1 kHz with an 8-pole Bessel filter, and the analogue output signal was digitized at a sampling rate of 5 kHz by using an A/D converter (Digidata 1322A, Axon Instruments). Data processing is performed using pClamp 8.0 software (Axon Instruments). To avoid electrostatic interference during measurements, the electrochemical cell is placed in a Faraday cage set on a mechanically isolated support.30

Single channel experiments are carried out in symmetrically buffered solutions composed of 5 mM Hepes and 0.5 M KCl, pH 7.4 at room temperature. After formation of stable lipid bilayers, Vpu1 was added to the cis compartment from a 10 nM stock solution in triluoroethanol, leading to a final concentration of approximately 10 nM. The illustrated channel current recordings are representative of the most frequently observed conductance under the applied experimental conditions. Single channel conductance was calculated from the corresponding Gaussian fits to current histograms by using data from segments of continuous recordings lasting longer than 10 s. Openings shorter than 0.5 ms were ignored.

Conclusions

A planar chip-based arrangement of so-called micro-BLMs with resistances larger than 1 GΩ based on highly ordered macroporous silicon arrays, which are on average stable for 1 day, has been designed, which is the crucial step toward an automated and paralleled array for ion channel recordings that will pave the way for ion channel screening assays.

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Supporting Information Available: Impedance spectra, time-resolved change in capacitance during micro-BLM formation, and current traces before and after addition of HMA. This material is available free of charge via the Internet at http://pubs.acs.org.

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