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A Potassium Channel Protein Encoded by Chlorella Virus PBCV-1

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Abstract: The large chlorella virus PBCV-1, which contains double-stranded DNA (dsDNA), encodes a 94-codon open reading frame (ORF) that contains a motif resembling the signature sequence of the pore domain of potassium channel proteins. Phylogenetic analyses of the encoded protein, Kcv, indicate a previously unidentified type of potassium channel. The messenger RNA encoded by the ORF leads to functional expression of a potassium-selective conductance in *Xenopus laevis* oocytes. The channel blockers amantadine and barium, but not cesium, inhibit this conductance, in addition to virus plaque formation. Thus, PBCV-1 encodes the first known viral protein that functions as a potassium-selective channel and is essential in the virus life cycle.

Potassium channels function in eukaryotes and prokaryotes as selective transport proteins for passive K⁺ movement across membranes (1). Common to virtually all known K⁺ channels is a pore domain with eight highly conserved

amino acids, TXXTXG(Y/F)G (2, 3). In a functional channel, four subunits surround a pore in which these residues form the selectivity filter (4).

Sequence analysis of the 330-kb dsDNA genome of the plaque-forming chlorella

virus PBCV-1 (family *Phycodnaviridae*) (5) identified a short amino acid sequence in a small ORF (ORF A250R) which resembles the pore domain of K⁺ channel proteins. ORF A250R is predicted to encode a peptide of 94 amino acids (referred to as Kcv) with an isoelectric point of 8.7 and a molecular weight of 10.6 kD (Figure 1A). Hydropathy analysis of Kcv reveals two putative transmembrane domains (Figure 1B) separated by 44 amino acids that contain the K⁺ channel signature sequence TXXTXGFG (Figure 1A, amino acids 60 through 67) (2). The 26 amino acids surrounding this motif display, on average, 61% similarity and 38% identity to the pore domains of many K⁺ channel proteins (Figure 1C). Two structurally important aromatic amino acids are also conserved in the NH₂-terminal portion of the Kcv pore domain. In the bacterial channel KcsA, two W residues (amino acids 67 and 68 in Figure 1C) are part of a structure that acts as a cuff, keeping the pore open at the appropriate diameter for K⁺ passage (6). In Kcv as well as in several other K⁺ channel proteins, aromatic Y or F residues replace these residues (Figure 1C).

In contrast to the putative pore domain, the amino acid sequences of the two Kcv transmembrane domains differ markedly from other K⁺ channels. A phylogenetic comparison with eukaryotic Kir, Kv, and tandem K⁺ channels,

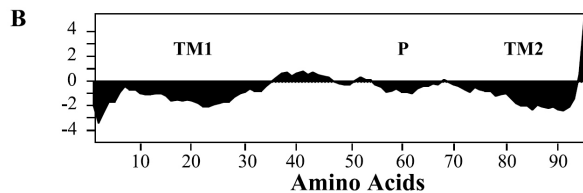
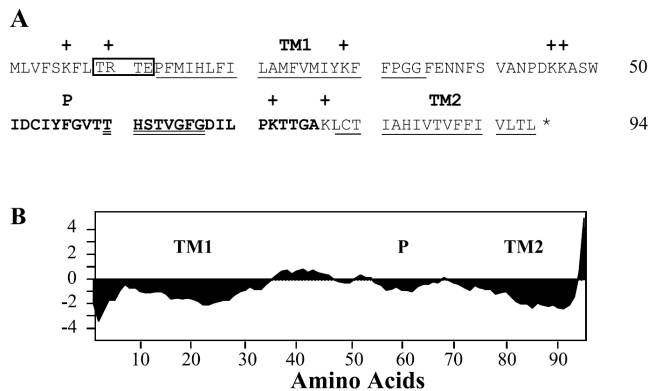
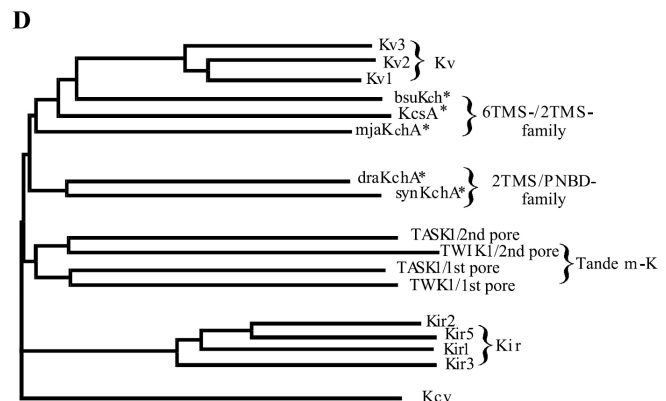


Figure 1. (A) Predicted amino acid sequence of chlorella virus PBCV-1 Kcv protein (GenBank accession AAC96618). Transmembrane regions (TM1, TM2) are underlined; pore region (P) is boldface; K⁺ channel signature sequence is double underlined; positively charged amino acids are marked by (+); a putative phosphorylation site (TRTE) is boxed. (B) Hydrophobicity profile (ordinate) was calculated according to Kyte and Doolittle (24) with a moving window of 19 amino acids and plotted against amino acid number (abscissa). (C) Alignment of the Kcv P domain with other K⁺ channel proteins (25). Amino acids similar or identical to Kcv are boldface. Following each sequence, the position of the last residue and the percent amino acid similarity and identity to Kcv (in brackets) are indicated. (D) Phylogenetic comparison of Kcv with eukaryotic and prokaryotic (denoted by asterisk) K⁺ channels calculated by CLUSTAL/CLUSTREE algorithms (26).

				% sim (id)
Kcv1	IDCIYFGVTT	HSTVGFQDIL	PKTTGA	76
Tandem-K	1st pore			
tok1/1	GNALYFCTVS	LLTVGLGDIL	PKSVGA	155 65 (50)
trek1/1	GSSFFFAQTV	ITTIGFGNIS	PRTEGG	155 65 (39)
Tandem-K	2nd pore			
C24A3/2	GTSLYFTLIS	FTTIGFGDIL	PSDYDF	279 58 (39)
T12C9/2	MDAFYYSFIS	LTTIGFGDIV	PENHDY	496 62 (35)
PAK family				
pak1	LHSLYWSIIT	MTTIGYGDIT	PQNLRE	354 58 (31)
pak2	FNSLYWITIT	SMTVGYGDIV	PVTTPE	340 62 (42)
EAG family				
eag	VTALYFTMTC	MTSVGFQDIL	AETDNE	466 50 (31)
herg	VTALYFTFSS	LTSVGFQDIL	PNTNSE	637 62 (31)
prokaryotic K⁺ channel genes				
KcsA	PRALWWSVET	ATTVGYGDLY	PVTLWG	88 62 (35)
ecoKch	MTAFYFSIET	MSTVGYGDIV	PVSESA	198 65 (46)
				61 (38)



and two prokaryotic K^+ channel families (7), places Kcv as an independent cluster, indicating significant sequence deviation (Figure 1D). Another unusual feature of Kcv is its short (12 amino acids) cytoplasmic NH_2 -terminus containing a consensus protein kinase ck2 phosphorylation site (Figure 1A, amino acids 9 through 12). The COOH-terminus of the Kcv protein is part of the second transmembrane region; Kcv thus appears to lack a COOH-terminal cytoplasmic tail. These structural data and the phylogenetic analysis suggest that Kcv represents a very primitive K^+ channel. This notion is supported by phylogenetic analyses of another viral protein that place the DNA polymerases from *Phycodnaviridae* near the root of all eukaryotic DNA polymerase delta proteins (5). This indicates that at least some PBCV-1-encoded proteins have long evolutionary histories.

To determine if Kcv functions as an ion channel, we expressed the protein in *Xenopus* oocytes (8). At 36 hours after Kcv mRNA injection, oocytes exhibited

distinct currents in voltage clamp assays (9) which differed quantitatively and qualitatively from control oocytes (Figure 2, A and B) (10). At voltages of +60 and -140 mV, average steady-state currents of Kcv mRNA-injected oocytes exceeded those of water-injected ones by factors of 8.6 and 8.2, respectively (11). The conductance introduced by Kcv mRNA injection consisted of an instantaneous and a time-dependent component (Figure 2A). The instantaneous I/V relation deviated from linearity by decreasing at extreme hyperpolarizing and depolarizing voltages. The time-dependent current activated and deactivated at negative and positive voltages, respectively. Figure 2B shows the instantaneous (I_i) and steady state-current (I_{ss}) from Kcv mRNA-injected oocytes and the steady-state component of water-injected oocytes as a function of voltage.

The selectivity of Kcv-mediated conductance was determined by obtaining I_i/V relationships at 2, 20, and 50 mM KCl (Figure 2C). Lowering the external

K^+ concentration (K^+_o) caused the current reversal voltage (V_r) to shift to more negative voltages and the inward current to decrease. Plotting V_r versus K^+_o in a Nernst plot yielded a slope of 60.0 mV per decade (Figure 2E). Figure 2D shows the time-dependent deactivation of the current following activation at a conditioning voltage of -160 mV. The current reversed at -20 mV, a value close to the reversal of I_i . This result was confirmed at all K^+_o values investigated (Figure 2E). Taken together, these analyses show that Kcv is a channel that conducts K^+ ions.

The cation selectivity of Kcv-mediated conductance was examined by replacing KCl with NaCl in the bath solution. Both the time-dependent and instantaneous currents were strongly depressed (Figure 3). Furthermore, V_r shifted negative by 68 mV, indicating that the Kcv-mediated conductance prefers K^+ over Na^+ ions. The P_K/P_{Na} permeability ratio from $n = 10$ I/V relations was 9.32 (12). Current reduction at positive voltages suggests an inhibitory effect of Na^+ on the K^+ outward current.

These results indicate that Kcv forms a K^+ -selective channel in oocytes (13). Kcv is the smallest K^+ channel protein known, primarily because of its short NH_2 - and COOH-terminal domains of other K^+

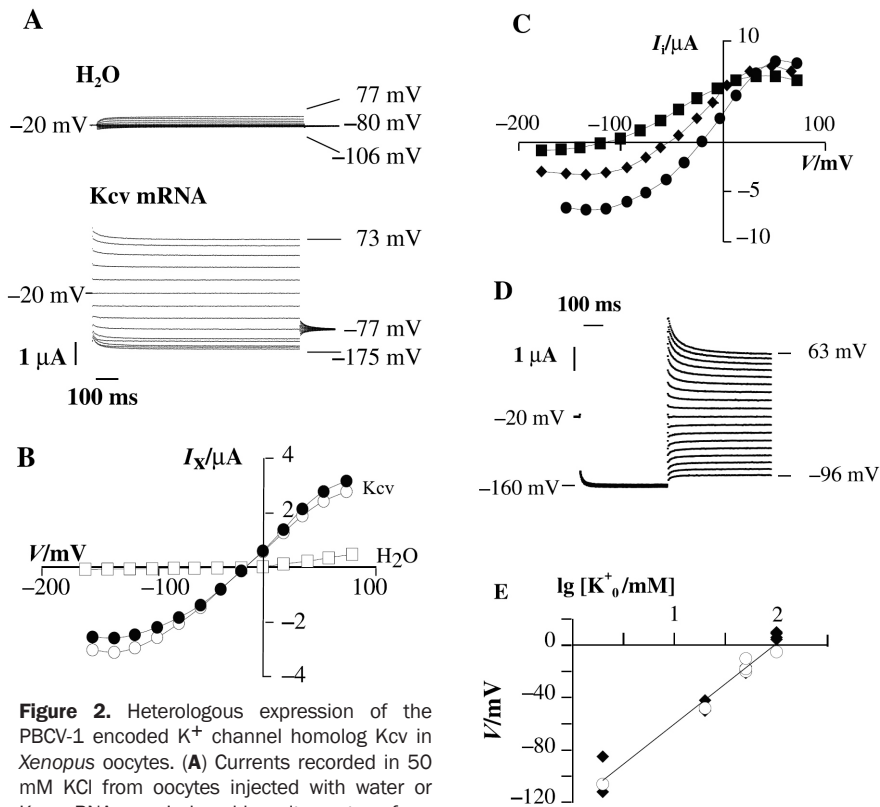


Figure 2. Heterologous expression of the PBCV-1 encoded K^+ channel homolog Kcv in *Xenopus* oocytes. (A) Currents recorded in 50 mM KCl from oocytes injected with water or Kcv mRNA were induced by voltage steps from holding voltage to test voltages as indicated. (B) I_i (solid symbols) and I_{ss} (open symbols) as a function of applied test voltages. Currents were measured after 3 ms (I_i) and at the end of the test pulse (I_{ss}). (C) I_i/V relations from a Kcv mRNA-injected oocyte in 2, 20, and 50 mM KCl measured as in (B). (D) Deactivation tail currents from a Kcv mRNA-injected oocyte in 50 mM KCl. Currents were elicited by clamping the oocyte to -160 mV for 1 s to activate time-dependent conductance. To follow current relaxation, voltage steps were applied in the ranges indicated. (E) Reversal voltages (V_r) obtained from I_i/V relations (solid symbols, $n = 9$ oocytes) and tail currents (open symbols, $n = 8$ oocytes) in Nernst plot against extracellular K^+ concentration. Linear regression to mean V_r from both types of analysis has a slope of 60.0 mV/decade.

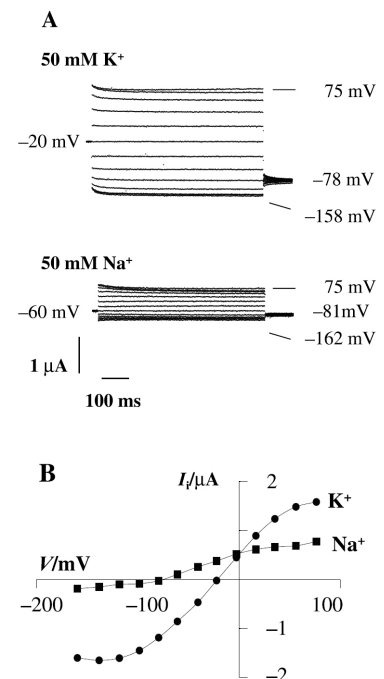


Figure 3. Selectivity of the Kcv-mediated conductance. Currents were elicited in response to voltage steps from resting voltage to test voltages in the ranges indicated. The bath solution in (A) contained 50 mM KCl or 50 mM NaCl. (B) I_i/V relation of data in (A).

channel proteins contribute to pore assembly and voltage sensitivity (14). Despite lacking these domains, Kcv displays several distinct properties with moderate voltage-sensitivity including (i) a decrease in I_i near both voltage extremes, (ii) a hyperpolarization-induced time-dependent activation, and (iii) a depolarization-induced time-dependent decrease of the conductance (Figure 2, A and B).

Oocytes expressing Kcv were exposed to amantadine, an antiviral drug which at concentrations $< 1 \mu\text{M}$ inhibits the influenza virus M2 protein (15). Amantadine inhibited the Kcv-mediated conductance (Figure 4A). Inhibition of I_i was voltage-independent (Figure 4A, lower panel). To quantify inhibition, we measured the effect of amantadine on the physiologically relevant steady-state current: At +60 and -140 mV, I_{ss} was half-inhibited by 2 and 0.8 mM amantadine, respectively (16). Thus, the effective concentration is about three orders of magnitude higher than required to inhibit M2. Amantadine inhibition of Kcv-mediated conductance reversed within minutes after removal of

the drug. In contrast, amantadine inhibition of M2 is essentially irreversible (15). These results, together with the fact that the two viral-encoded proteins have little or no structural similarity, suggest that amantadine inhibits the Kcv and M2 proteins by different mechanisms (17).

Kcv-mediated conductance was also exposed to the typical K^+ channel blockers Cs^+ and Ba^{2+} . Addition of 10 mM CsCl had only modest effects on the Kcv-mediated conductance. The inward current at -140 mV was inhibited by $9 \pm 4\%$, whereas the outward current at +60 mV was unaffected ($n = 4$ oocytes). In contrast, Ba^{2+} reduced the Kcv-mediated conductance in a voltage-dependent manner (Figure 4B) (18). This behavior supports the hypothesis that Ba^{2+} blocks inward current in K^+ channels after being drawn into the pore by negative voltage (19).

A plaque reduction assay (20) was employed to determine the importance of Kcv to virus replication (21). Amantadine inhibited PBCV-1 plaque formation by 50% at $2.8 \pm 0.2 \text{ mM}$ ($n = 4$) (16), that

is, at approximately the same millimolar concentration that affected the Kcv-mediated conductance (Figure 4C). Plaque formation was also inhibited 50% by $2.6 \pm 0.16 \text{ mM Ba}^{2+}$ ($n = 3$), a concentration sufficient to abolish Kcv inward current. Cesium (10 mM) had no apparent effect on PBCV-1 replication. These results, together with the finding that the Kcv gene is expressed early after virus infection (22) indicate that PBCV-1 replication depends, in a yet unknown manner, on a functional Kcv channel.

In conclusion, chlorella virus PBCV-1 encodes a functional K^+ channel protein, Kcv, which is important for virus replication. Prior to this study, only four virus-encoded proteins were thought to have ion channel activities: influenza virus A M2 protein, influenza virus B NB protein, and the human immunodeficiency virus proteins Vpu and Vpr (23). PBCV-1 is the first virus known to encode a K^+ -selective channel. Because of its exceptionally small size, Kcv may become a useful tool to study basic principles of channel assembly and function.

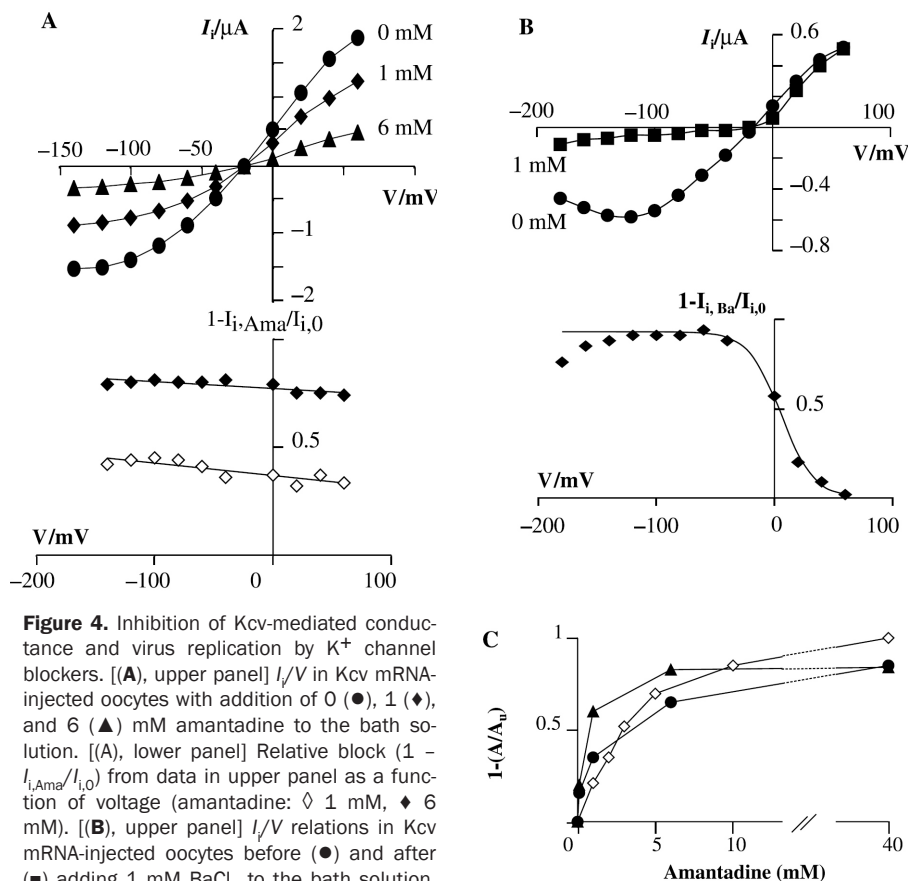


Figure 4. Inhibition of Kcv-mediated conductance and virus replication by K^+ channel blockers. [(A), upper panel] I_i/V in Kcv mRNA-injected oocytes with addition of 0 (\bullet), 1 (\blacklozenge), and 6 (\blacktriangle) mM amantadine to the bath solution. [(A), lower panel] Relative block ($1 - I_{i,Ama}/I_{i,0}$) from data in upper panel as a function of voltage (amantadine: \diamond 1 mM, \blacklozenge 6 mM). [(B), upper panel] I_i/V relations in Kcv mRNA-injected oocytes before (\bullet) and after (\blacksquare) adding 1 mM BaCl_2 to the bath solution. [(B), lower panel] Relative block ($1 - I_{i,Ba}/I_{i,0}$) from data in the upper panel as function of voltage. Data fitted (solid line) with Woodhull block model (18), yielding $b_{\text{max}} = 0.93$, $\delta = 0.91$, and $k^0 = 650 \mu\text{M}$ ($n = 3$ oocytes). (C) Relative inhibition ($1 - A/A_0$) of virus plaque formation (\diamond) and Kcv-mediated I_{ss} at +60 mV (\bullet) and -140 mV (\blacktriangle) as function of amantadine concentration in the external medium. Mean of four experiments; SE is smaller than symbols.

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3. Single-letter abbreviations for the amino acid residues are as follows: A, Ala; C, Cys; D, Asp; E, Glu; F, Phe; G, Gly; H, His; I, Ile; K, Lys; L, Leu; M, Met; N, Asn; P, Pro; Q, Gln; R, Arg; S, Ser; T, Thr; V, Val; W, Trp; and Y, Tyr. X indicates any residue.
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8. ORF A250R was amplified from PBCV-1 genomic DNA by polymerase chain reaction, using primers with restriction sites for Bam HI (5' end) and Xho I (3' end). The amplified fragment was cut with the two enzymes and ligated into predigested vector pSGEM (constructed from pGEM-HE, given to B. Plugge by courtesy of M. Hollmann, Max-Planck-Institut for Experimental Medicine, Göttingen, Germany). The cDNA templates were linearized with Nsi I and transcribed in vitro with T7 RNA polymerase. The mRNA was injected in *Xenopus laevis* oocytes. Membrane currents and voltages were recorded with conventional two-electrode voltage clamp method, using the Gene Clamp 500 amplifier under control of pCLAMP 5.5 software. Both electrodes were filled with 3 M KCl and had resistances between 0.4 and 1 megohms in 50 mM KCl. The oocytes were continually perfused (25°C) with experimental solution

- (about 2 ml min⁻¹) containing 50 mM KCl, 1.8 mM CaCl₂, 1 mM MgCl₂, and 5 mM HEPES/LiOH (pH 7.4). Mannitol was used to adjust the osmolarity to 215 mOsmol.
9. The amount of mRNA injected was limited to 1 ng/50 nl. Injection of higher amounts of mRNA or expression times longer than 48 hours caused oocytes to die.
 10. Two typical dominant endogenous currents were recorded in H₂O-injected oocytes: (i) an inactivating inward rectifying K⁺ current and (ii) a time-dependent outward rectifying Cl⁻ current [C. K. T. Bauer, J. R. Falk, W. Schwarz, *Pfluegers Arch.* **432**, 812 (1996); N. Dascal, *Crit. Rev. Biochem.* **22**, 317 (1987)].
 11. Steady-state currents at test voltage of +60 mV and -140 mV (voltage protocols as in Figure 2) recorded in oocytes injected with (i) H₂O: +0.2 ± 0.05 μA and -0.22 ± 0.08 μA (*n* = 9); (ii) Kcv mRNA: +1.63 ± 0.37 μA and -1.89 ± 0.44 μA (*n* = 17).
 12. The P_K/P_{Na} permeability ratio was calculated by least-squares fitting of reversal potentials in either external KCl or NaCl solutions by the GHK equation [B. Hille, *Ionic Channels of Excitable Membranes* (Sinauer, Sunderland, MA, 1992)], assuming permeation of K⁺ and Na⁺ ions only. Oocyte intracellular Na⁺ and K⁺ concentrations were assumed to be 20 mM [K. Kusano, R. Miledi, J. Stinnakre, *J. Physiol (London)* **328**, 143 (1982)] and 101 mM (based on the fitting of the Nernst plot in Figure 2E), respectively.
 13. However, the possibility exists that Kcv functions as a subunit, which up-regulates the activity of endogenous K⁺ channels in oocytes. To exclude this possibility, we constructed a site-specific mutation in the selectivity filter sequence of Kcv by replacing Phe⁶⁶ (F66) with Ala (A). If Kcv is a channel protein, the mutant protein should, by analogy to the Shaker channel, form a channel unable to conduct K⁺ currents (1). Using standard voltage-clamp assays, nine oocytes expressing KcvF66A had currents similar in kinetics to those of H₂O-injected oocytes with no additional K⁺ conductance compared to the H₂O-injected control cells. Hence, the absence of a prominent K⁺-selective current in KcvF66A-expressing oocytes confirms that Kcv functions as a channel protein in oocytes and that the observed currents are not due to activation of endogenous channel proteins.
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 16. Concentration for half-maximal inhibition estimated by fitting Michaelis-Menten type kinetics to data.
 17. The adamantyl group of amantadine is believed to interact with the hydrophobic lining of the M2 pore, whereas the ammonium group forms H bonds with the imidazole ring of His³⁷ [C. S. Gandhi et al., *J. Biol. Chem.* **274**, 5474 (1999)].
 18. Voltage-dependency of inhibition by Ba²⁺ was analyzed quantitatively based on a Woodhull block model [A. M. Woodhull, *J. Gen. Physiol.* **61**, 687 (1973)] by fitting data of relative block to

$$(1 - I/I_0) = b_{\max}/[1 + (k^0/B_a) e^{(z\delta F/RT)}]$$
 where I_0 is control and I blocked current, b_{\max} the maximal block, B_a the concentration of Ba²⁺, k^0 the dissociation constant of the blocking reaction at voltage = 0 mV, δ the fraction of the electrical field crossed by Ba²⁺, and $z = 2$ the valence of the blocking ion. R , T , and F have their usual thermodynamic meaning. Fitting yields for 1 mM Ba²⁺: $b_{\max} = 0.94$, $\delta = 0.9 \pm 0.03$, and $k^0 = 660 \pm 12 \mu\text{M}$ (*n* = 3 oocytes).
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 21. About 1200 plaque-forming units of virus PBCV-1, or a small plaque variant P1210 [D. Landstein, D. E. Burbank, J. W. Nietfeldt, J. L. Van Etten, *Virology* **214**, 413 (1995)], were mixed with 10⁸ host cells (*Chlorella* strain NC64A) and warm MBBM top agar containing various concentrations of ion-channel inhibitors. The mixture was layered onto MBBM nutrient agar (22). After incubating for 2 days at 25 °C, viral plaques were counted. Potassium concentration of MBBM growth medium was 1.3 mM.
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 27. We thank H. Terlau, D. Gradmann, and M. Blatt for helpful discussions. Supported in part by MURST in the framework of Cofin-99, NIH grant GM32441 to J.V.E., Small Business Innovative Research grant GM41333 to M.N., Telethon grants (#971 and #296.bi) to D.D. and A.M., and a traveling grant by the SmithKline Beecham foundation to B.P.