Cellular/Molecular

Functional Coupling between Sulfonylurea Receptor Type 1 and a Nonselective Cation Channel in Reactive Astrocytes from Adult Rat Brain

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We previously identified a novel, nonselective cation channel in native reactive (type R1) astrocytes (NR1As) from injured rat brain that is regulated by cytoplasmic Ca\(^{2+}\) and ATP (NC\(_{Ca-ATP}\)) and exhibits sensitivity to block by adenine nucleotides similar to that of sulfonylurea receptor type 1 (SUR1). Here we show that SUR1 is involved in regulation of this channel. NR1As within the site of injury and after isolation exhibited specific binding of FITC-tagged glibenclamide and were immunolabeled with anti-SUR1 antibody, but not with anti-SUR2, anti-Kir6.1 or anti-Kir6.2 antibodies, indicating absence of ATP-sensitive K\(^{+}\) (K\(_{ATP}\)) channels. RT-PCR confirmed transcription of mRNA for SUR1 but not SUR2. Several properties previously associated exclusively with SUR1-regulated K\(_{ATP}\) channels were observed in patch-clamp experiments using Cs\(^{+}\) as the charge carrier: (1) the sulfonylureas, glibenclamide and tolbutamide, inhibited NC\(_{Ca-ATP}\) channels with EC\(_{50}\) values of 48 \(\mu\)M and 16.1 \(\mu\)M, respectively; (2) inhibition by sulfonylureas was lost after exposure of the intracellular face to trypsin or anti-SUR1 antibody; (3) channel inhibition was caused by a change in kinetics of channel closing, with no change in channel amplitude or open-channel dwell times; and (4) the SUR activator (“K\(_{ATP}\) channel opener”), diazoxide, activated the NC\(_{Ca-ATP}\) channel, whereas pinacidil and cromakalin did not. Also, glibenclamide prevented cell blebbing after ATP depletion, whereas blebbing was produced by exposure to diazoxide. Our data indicate that SUR1 is functionally coupled to the pore-forming portion of the NC\(_{Ca-ATP}\) channel, providing the first demonstration of promiscuity of SUR1 outside of the K\(^{+}\) inward rectifier family of channels.

Key words: sulfonylurea; SUR1; glibenclamide; diazoxide; cation channel; cell swelling; astrocyte; brain injury; patch clamp

Introduction

The antidiabetic sulfonylureas, which inhibit ATP-sensitive K\(^{+}\) (K\(_{ATP}\)) channels in pancreatic β cells and stimulate insulin release in diabetes mellitus, mediate their effect on K\(_{ATP}\) channels via a high-affinity sulfonylurea receptor (SUR) (Pantén et al., 1989; Aguilar-Bryan et al., 1995). Several isoforms of SUR, termed SUR1, SUR2A, SUR2B, and SUR2C, have been identified and cloned (Aguilar-Bryan et al., 1995; Inagaki et al., 1996; Isomoto et al., 1996; Lawson, 2000). These receptors belong to the ATP-binding cassette (ABC) transporter family, of which the cystic fibrosis transmembrane conductance regulator (CFTR), another ion channel modulator, is also a member (Higgins, 1992; Aguilar-Bryan et al., 1995).

K\(_{ATP}\) channels are heteromultimers, with Kir6.1 forming the conductive pores and SURx functioning as regulatory subunits (Shyng et al., 1997a; Tucker et al., 1997, 1998; Babenko et al., 1998; Drain et al., 1998). SUR imparts sensitivity to antidiabetic sulfonylureas such as glibenclamide and tolbutamide and is responsible for activation by a chemically diverse group of agents termed “K\(^{+}\) channel openers” (here termed “SUR activators”) such as diazoxide, pinacidil, and cromakalin (Aguilar-Bryan et al., 1995; Inagaki et al., 1996; Isomoto et al., 1996; Nichols et al., 1996; Shyng et al., 1997b). In various tissues, molecularly distinct SURs are coupled to distinct channel moieties to form different K\(_{ATP}\) channels with distinguishable physiological and pharmacological characteristics. The K\(_{ATP}\) channel in pancreatic β cells is formed from SUR1 linked with Kir6.2, whereas the cardiac and smooth muscle K\(_{ATP}\) channels are formed from SUR2A and SUR2B linked with Kir6.2 and Kir6.1, respectively (Fujita and Kurachi, 2000). In addition to these principal constructs, another inward rectifier channel, Kir1.1a, has been reported to link to another ABC ion channel regulator, CFTR (Ruknudin et al., 1998), or even to SUR itself (Ammala et al., 1996), thereby gaining sensitivity to sulfonylureas. To date, the only channels known to be regulated by SURx are inward rectifier K\(_{ATP}\) channels formed from Kir6.1, Kir6.2, or possibly Kir1.1a.

We recently identified a novel Ca\(^{2+}\)-activated, ATP-sensitive nonselective cation channel (NC\(_{Ca-ATP}\)) in native (not cultured) reactive (type R1) astrocytes (NR1As) from adult rat brain (Chen and Simard, 2001). This channel, which is nearly equally permeable to K\(^{+}\), Na\(^{+}\), Rb\(^{+}\), Li\(^{+}\), and Cs\(^{+}\), is readily distinguished from K\(_{ATP}\) channels, which are not permeable to Na\(^{+}\) and Cs\(^{+}\). The sensitivity of the NC\(_{Ca-ATP}\) channel to ATP suggested that it might belong to the ABC transporter family, prompting us to hypothesize that sulfonylureas might modulate it. Here we report that the NC\(_{Ca-ATP}\) channel exhibits physiological and pharmaco-
logical properties previously associated only with $K_{ATP}$ channels possessing the SUR1 regulatory subunit, that glibenclamide-mediated channel inhibition was prevented by anti-SUR1 antibody, and that blocking the channel with glibenclamide protected cells from blebbing secondary to Na azide-induced ATP depletion. Together, our data show that SUR1 is expressed, is coupled to, and regulates the pore-forming portion of the NC$_{CAS-ATP}$ channel in NR1A and that SUR-mediated regulation of this channel plays a role in blebbing and swelling of reactive astrocytes.

**Materials and Methods**

**Cell preparations.** All animal protocols were approved by the Institutional Animal Care and Use Committee of the University of Maryland. Native reactive astrocytes from adult brain were obtained from the glotic capsule that forms around gelatin sponges (Gelofom, Upjohn Co., Kalama-zoo, MI) implanted into a stab wound in the parietal lobe of 8-week-old Wistar rats as described previously (Perillan et al., 1999, 2000). Sponge pieces were harvested at 8 d and washed three times in PBS, pH 7.4. Washed pieces were placed in an Eppendorf tube containing artificial CSF (ACSF) composed of (mM): 124 NaCl, 5 KCl, 1.3 MgCl$_2$, 2 CaCl$_2$, 26 NaHCO$_3$, and 10 d-glucose, pH 7.4. $\approx 290$ mM O$_2$, with 20 U/ml papain, 10 mg/ml trypsin inhibitor, and 0.01% DNase (Worthington, Lakewood, NJ). The digestion system was transferred to an incubator (humidified 10% CO$_2$/95% air at 37°C). Culture medium was exchanged every 3 d. Culture medium was composed of (mM): 145 CsCl, 1.5 CaCl$_2$, 1 MgCl$_2$, 5 EGTA, 32.5 HEPES, 12.5 glucose, pH 7.4. All animal protocols were approved by the Institutional Animal Care and Use Committee of the University of Maryland.

**Histological staining.** Alternatively, for fluorescence imaging, cell nuclei were labeled using 4,6-diamidino-2-phenylindole (DAPI) (Sigma). Alternatively, when capturing low-power views of tissue sections, we used species-appropriate horseradish peroxidase-conjugated secondary antibodies. Specimens were examined using a Nikon Eclipse E1000 microscope. Images were captured using a SenSys digital camera (Roper Scientific) and processed using a personal computer running IPLab software (version 3.0, Scionographics). Post-processing of images was performed using Adobe Photoshop (version 5.0, Adobe Systems).

**Whole-cell patch clamp recording**

**Whole-cell patch clamp recordings.** For whole-cell macroscopic recordings, we used a nystatin perforated-patch technique with a bath solution containing (mM): 130 NaCl, 10 KCl, 1 CaCl$_2$, 1 MgCl$_2$, 32.5 HEPES, 12.5 glucose, pH 7.4. The pipette solution contained (in mM): 55 KCl, 75 K$_2$SO$_4$, 8 MgCl$_2$, and 10 HEPES, pH 7.2. Nystatin (50 mg) (Calbiochem) was dissolved in 1 ml of dimethylsulfoxide (DMSO). Working solutions were made before the experiment by adding 16.5 ml of nystatin stock solution to 5 ml of the base pipette solution to yield a final concentration of nystatin of 165 µg/ml and DMSO 3.3 µl/ml.

**For cell-attached patch recording,** we used a bath solution containing (in mM): 130 NaCl, 10 KCl, 1 CaCl$_2$, 1 MgCl$_2$, 32.5 HEPES, 12.5 glucose, pH 7.4. The pipette solution contained (in mM): 145 CsCl, 1 MgCl$_2$, 0.2 CaCl$_2$, 5 EGTA, 10 HEPES, pH 7.3. The measured osmolarity of the extracellular solution was $\approx 300$ mOsm (Precision Systems, Natick, MA).

**For inside-out patch recording,** we used a bath solution containing (in mM): 145 CsCl, 1.5 CaCl$_2$, 1 MgCl$_2$, 5 EGTA, 32.5 HEPES, 12.5 glucose, pH 7.4. The pipette solution contained (in mM): 145 CsCl, 1 MgCl$_2$, 0.2 CaCl$_2$, 5 EGTA, 10 HEPES, pH 7.3.

**For outside-out patch recording,** we used the pipette solution, which contained (in mM): 145 CsCl, 1 MgCl$_2$, 0.2 CaCl$_2$, 1 ATP, 1 EGTA, 10 HEPES, pH 7.3. The standard bath solution contained (in mM): 145 CsCl, 1.5 CaCl$_2$, 1 MgCl$_2$, 5 EGTA, 32.5 HEPES, 12.5 glucose, pH 7.4. SUR activators ($K_{ATP}$ channel openers) diazoxide, pinacidil, and cromakalin, and sulfonylureas, tolbutamide and glinclamide, were obtained from Sigma and dissolved in DMSO at concentrations of 10–100 mM for stock solutions.

Working solutions were made before the experiment by adding stock solution to the bath solution to yield the desired concentration.
To calculate open-channel probability (amplitude histogram of records obtained at various potentials). To calculate open-channel probability, the all-points histogram was fit to a Gaussian function to an all-points histogram obtained during test pulse to concentration. (DMSO was tested in separate experiments and was found to be without effect at the concentrations used.) Antibodies used for electrophysiological experiments, which were the same as those used for immunolabeling, included anti-SUR1-antibody (catalog #sc-5789) and anti-Kir6.1 antibody (catalog #sc-11224), both from Santa Cruz Biotechnology.

Data analysis. Single-channel amplitudes used to calculate slope conductances were obtained by fitting a Gaussian function to an all-points amplitude histogram of records obtained at various potentials. To calculate open-channel probability (\(a \times p_o\)) at various potentials and with different test agents, the all-points histogram was fit to a Gaussian function, and the area under the fitted curve for the open channel was divided by the area under the fitted curve for the closed plus open channel.

For dwell time histograms, we used records with single-channel openings obtained during test pulse to \(E_m \approx -80\) mV, filtered at 1 kHz (3 dB; rise time, 330 \(\mu\) sec). As suggested previously (Sigworth and Sine, 1987), the distributions of open or closed times were compiled after conversion to the logarithm of the time interval, using 10 bins per decade for the abcissa of the histogram and a square root axis for the ordinate. Using this transformation, the probability density function (pdf) for a double exponential distribution is pdf = \(a_1 \times \exp[z_1 - \exp(z_2)] + a_2 \times \exp[z_2 - \exp(z_1)]\) \(t \leq t_1\), where \(z_1 = [\ln(t) - \ln(t_1)]\), \(z_2 = [\ln(t) - \ln(t_2)]\), \(t_1\) is time, and \(a_1\) and \(a_2\) relate to the number of open events having time constants of \(t_1\) and \(t_2\), respectively.

Junction potentials, which generally did not exceed 5 mV, were subtracted when appropriate. Holding currents were not subtracted from any of the recordings. Concentration–response data were fit to a standard logistic equation using the Levenberg-Marquardt algorithm as implemented in Origin 7.0 (Microcal Software, Northampton, MA). Data are given as mean ± SD.

**Results**

**Glibenclamide binding**

We used FITC-tagged glibenclamide to evaluate sulfonylurea binding sites, DAPI to label cell nuclei, and anti-GFAP antibodies as a marker of reactive astrocytes at the site of brain injury. Glibenclamide binding was found to be prominent in the layer of tissue corresponding to the gliotic capsule surrounding the gelatin sponge (Fig. 1A). Scattered areas of positive label were also found within the sponge itself (Fig. 1B). At higher magnification, glibenclamide-labeled cells were observed (Fig. 1C) that were invariably positive for GFAP (Fig. 1D), identifying them as reactive astrocytes. Overall, we estimated that >80% of the cells encapsulating the sponge and present within the sponge, as identified by DAPI, showed binding of glibenclamide. Specificity of binding was assessed by pretreating tissue sections with unlabeled glibenclamide before exposing them to FITC-tagged glibenclamide, a protocol that yielded sections with no fluorescent label (data not shown). In brain adjacent to the injury, no glibenclamide binding was apparent (Fig. 1A), although abundant label for GFAP was present (Perillan et al., 1999). These data indicated that sulfonylurea binding sites, which are not normally present in subcortical white matter, were expressed abundantly in GFAP-positive cells encapsulating and invading the foreign body after traumatic injury.

**Immunolabeling and RT-PCR for SUR1 and SUR2**

To further characterize the sulfonylurea binding sites, we assessed for expression of sulfonylurea receptors. Eight days after injury, strong labeling with anti-SUR1 antibody was found in the gliotic capsule surrounding the gelatin sponge (Fig. 2A). The gliotic capsule was harvested and, as reported previously, subjected to enzymatic digestion to release NRAs (Chen and Simard, 2001). Most of the cells released by this procedure are the NR1As, which are large, round, phase bright, and GFAP-positive and express the NCa,ATP channel (Chen and Simard, 2001). Examination at low magnification of fields of cells labeled with anti-SUR1 antibody

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**Figure 1.** Binding of glibenclamide. A, Fluorescence image of giotic capsule 8 d after injury labeled with FITC-tagged glibenclamide; parenchyma is shown in bottom right corner, gelatin sponge is shown in top left, and giotic capsule is arranged diagonally. B, Fluorescence image of gelatin sponge at higher power shows labeled cells lining cavities of the sponge. C, D, High-power fluorescence image of a single cell showing FITC-tagged glibenclamide (C) and anti-GFAP (D) immunolabeling of the same cell.

**Figure 2.** Immunolabeling and RT-PCR for SUR1 in giotic capsule and freshly isolated reactive astrocytes. A, Immunofluorescence image of giotic capsule labeled using anti-SUR1 antibody; Gelfoam sponge (gf) is shown at left, B, C, Low-power (B) and high-power (C) views of freshly isolated astrocytes immunolabeled using anti-SUR1 antibody and visualized using chromogenic (B) or fluorescent (C) secondary antibodies; D, RT-PCR for SUR1 and SUR2 in freshly isolated astrocytes and control tissues; SUR1-mRNA was present in insulinoma RIN-m5F cells (lane 2) and astrocytes (lane 3); SUR2 was present in cardiomyocytes (lane 4) but not in astrocytes (lane 5); lane 1 is the 100 bp DNA size marker, and lane 6 is the β-actin (540 bp) RT-PCR positive control reaction.
confirmed that nearly all cells expressed SUR1 (Fig. 2B). SUR1-positive cells examined at high magnification were invariably the large round cells typical of NR1A (Fig. 2C). Tissues and isolated NRAs were uniformly negative when labeled with anti-SUR2 antibody (data not shown).

Preparations of NRA, enriched using gradient centrifugation to remove erythrocytes, were used in RT-PCR experiments to assess transcription of SURx mRNA. Insulinoma RIN-m5f cells and rat cardiomyocytes, which constitutively express KATP channels (Aguilar-Bryan et al., 1995), were used as positive controls for SUR1 and SUR2, respectively. Total RNA from NRA, RIN-m5f cells, and cardiomyocytes was reverse transcribed with oligo-dT primers to obtain cDNA, which was then amplified by PCR using primers unique for SUR1 and SUR2, respectively. Amplicons for SUR1 and SUR2 were found with control cardiac and NRA RNA, which are obtained only in very small numbers in our model of injury, exhibit a different electrophysiological profile than NR1As and not with NRA RNA (Fig. 2D, lanes 2 and 3, respectively). Amplicons for SUR2 were found with control cardiomyocytes but not with NRA RNA (Fig. 2D, lanes 4 and 5, respectively). Sequencing of the amplicon from NRA obtained from the agarose gel showed a 100% match with the corresponding 181 bp SUR1 sequence in the GenBank database. These data were consistent with SUR1, but not SUR2A, SUR2B, or SUR2C, mRNA transcription in NRA.

Immunolabeling for Kir6.1 and Kir6.2
The SURx are known to complex principally with Kir6.1 or Kir6.2 to form functional KATP channels (Aguilar-Bryan et al., 1995; Isomoto et al., 1996). We thus performed immunolabeling experiments on NRA using anti-Kir6.1 and anti-Kir6.2 antibodies to assess for the presence of KATP channels. Tissue sections were typically negative for both Kir6.1 (Fig. 3A) and Kir6.2 (data not shown). Similarly, for the majority of isolated cells, no labeling for either Kir6.1 (Fig. 3B) or Kir6.2 (data not shown) was evident, although rarely an occasional cell was labeled with anti-Kir6.1 antibody (Fig. 3B, arrow). Examination of positive cells at high magnification showed that they were invariably stellate-shaped astrocytes (Fig. 3C), which have previously been shown to express Kir6.1 KATP channels (Thomzig et al., 2001). By contrast, the more common large round cells, i.e., the NR1As, were not labeled with anti-Kir6.1 antibody (Fig. 3D). Notably, stellate astrocytes, which are obtained only in very small numbers in our model of injury, exhibit a different electrophysiological profile than NR1As and do not exhibit currents attributable to the NCa,ATP channel (Perillan et al., 2000). Lack of Kir6.1 and Kir6.2 labeling in NR1As indicated that KATP channels were not present in these cells, suggesting a different role for SUR1.

Sulfonylureas block the NCa,ATP channel
To this point, we had found that the NCa,ATP channel in NR1As and the KATP channel in pancreatic β cells bore important similarities: both channels exhibited similar sensitivity to the adenine nucleotides ATP, ADP, and AMP (Chen and Simard, 2001), and both channels were found in cells that expressed SUR1. Conversely, they bore an important difference: only the pancreatic β cells and not the NR1As expressed Kir6.x. We thus hypothesized that the NCa,ATP channel in NR1As might be formed from a non-Kir6.x pore-forming moiety that was regulated by SUR1. To test this, we performed a series of pharmacological and electrophysiological experiments designed to mimic experiments that...
have been used previously to characterize K\textsubscript{ATP} channels regulated by SUR1. For these experiments, care was taken to study only NR1As, which are readily identified by their large, round, phase-bright appearance, and to exclude stellate astrocytes as well as small phase-dark type R2 astrocytes (Dalton et al., 2003).

One hallmark of SUR1-regulated K\textsubscript{ATP} channels, when assessed either in pancreatic \( \beta \) cells or in expression systems expressing SUR1 plus Kir6.2, is channel block by the sulfonylureas, glibenclamide, and tolbutamide (Aguilar-Bryan et al., 1995; Inagaki et al., 1996). We used inside-out patches to assess block of the NC\textsubscript{Ca-ATP} channels by sulfonylureas. More than 90% of patches excised into solution with 1 \( \mu \)M Ca\textsuperscript{2+}, without ATP, exhibit 35 pS activity characteristic of this channel (Chen and Simard, 2001). We studied patches using Cs\textsuperscript{+} as the charge carrier to assure that no K\textsuperscript{+} channel, specifically K\textsubscript{ATP}, was contributing to patch activity. With no ATP and 1 \( \mu \)M Ca\textsuperscript{2+} in the bath, patches exhibited vigorous openings caused by multiple channels (Fig. 4A,Ba). Measurements of single-channel conductance at different potentials and calculation of slope conductance confirmed the presence of a 35 pS cation channel, consistent with the properties reported previously for this channel (Fig. 4C). Channel availability in this patch was profoundly diminished by subsequent addition of 10 \( \mu \)M glibenclamide (Fig. 4A,Bb), an effect that was partially reversed on washout of drug (Fig. 4Bc). Another sulfonylurea, tolbutamide, was also found to inhibit NC\textsubscript{Ca-ATP} channel activity. At low concentrations (\( \leq 1 \mu \)M), tolbutamide had no obvious effect on the channel, but as the concentration was increased from 1 to 300 \( \mu \)M, channel openings were increasingly diminished (Fig. 5A).

We measured the open-channel probability (\( n \times p_0 \)) at different concentrations of both glibenclamide and tolbutamide, normalized these values to those obtained at 1 \( \mu \)M and 0.1 \( \mu \)M for glibenclamide and tolbutamide, respectively, and fitted these values to a standard logistic equation. As shown in Figure 5B, the NC\textsubscript{Ca-ATP} channel was blocked by sulfonylureas in a dose-dependent manner. With tolbutamide, half-maximum inhibition (EC\textsubscript{50}) was observed at 16.1 \( \mu \)M with a Hill coefficient of 1.3, and channel activity was completely lost at concentrations >300 \( \mu \)M. With glibenclamide, the EC\textsubscript{50} was 48 nM with a Hill coefficient of 1.3 and in expression systems with SUR1, but not with SUR2 (Fujita and Kurachi, 2000).

Mechanism of channel inhibition by sulfonylureas

Characterizing the mechanism of sulfonylurea inhibition can be informative regarding the identity of the SUR isoform. With Kir6.2 as the channel moiety, inhibition does not occur directly by block of the channel pore but by an allosteric mechanism that affects channel kinetics. On the basis of analysis of channel kinetics, sulfonylurea-mediated inhibition of Kir6.2 via SUR1 is distinguishable from inhibition via SUR2 (Babenko et al., 1999).

We performed single-channel analysis to characterize the
mechanism of block of sulfonylureas on the NCa-ATP channel. First, it was evident from inspection of single-channel records that channel amplitude was not reduced with increasing concentration of drug (Fig. 5A). A reduction in channel amplitude is frequently observed when a blocking agent enters the pore of the channel and, with rapid off-time kinetics, produces a “flickery” block. This is the presumed mechanism for block of the NCa-ATP channel by intracellular Mg2+ (Chen and Simard, 2001), but it appeared not to be relevant to sulfonylurea-mediated inhibition of the NCa-ATP channel.

Open-channel dwell times were analyzed. As characterized previously, the NCa-ATP channel exhibits two open states, with dwell times (τo1 and τo2) of 2 and 8 msec, with 28% of openings being from the closed to the long open state (Chen and Simard, 2001). Similar observations were made here before drug application, with the example shown exhibiting an open probability (n × Po) of 0.63, with values of τo1 and τo2 of 1.9 and 8.2 msec, and with 31% of openings being from the closed to the long open state (Fig. 6A,D). After successive application of 3 μM (Fig. 6B,E) and 30 μM (Fig. 6C,F) tolbutamide, n × Po decreased to 0.44 and 0.09, respectively, and the number of openings from the closed to the long open state diminished to 23 and 10%, respectively (Fig. 6E,F); however, values of τo1 and τo2 were not appreciably affected by drug (τo1 = 1.9 and 1.9 msec and τo2 = 8.1 and 8.0 msec at 3 and 30 μM tolbutamide, respectively), indicating minimal effect on open-channel dwell times.

Closed-channel dwell times were also analyzed. Data from the same patch showed that both the duration and frequency of long closures increased with drug. Before drug application, the NCa-ATP channel exhibited two closed states, with dwell times (τc1 and τc2) of 1.7 and 6.3 msec, with the long closed state representing 18% of closures (Fig. 6G). With 3 μM (Fig. 6H) and 30 μM (Fig. 6I) tolbutamide, the time constant for the long closed state (τc2) increased to 14.1 and 30.3 msec, respectively, and the fraction of closings represented by the long closed state increased to 33 and 42%, respectively. Similar results were observed in four other patches. This form of channel inhibition, i.e., no effect on open-channel dwell times and a progressive increase in long closures, is typical of the inhibition produced by sulfonylureas acting on the KATP channel in pancreatic β cells (Gillis et al., 1989; Babenko et al., 1999).

SUR-mediated activation of NCa-ATP Channel

Another hallmark of SUR1-regulated KATP function, when assessed either in pancreatic β cells or in expression systems expressing SUR1 plus Kir6.2, is channel activation by the SUR activator diazoxide, with weak or no activation by pinacidil or cromakalin, which preferentially activate SUR2 (Fujita and Kurauchi, 2000). We found previously that cell-attached patches from healthy NR1As are usually quiet and that ATP depletion after exposure to Na azide or NaCN plus 2-deoxyglucose was required to activate NCa-ATP channels in this recording configuration (Chen and Simard, 2001). We used cell-attached patches to examine the effect of diazoxide on the NCa-ATP Channel, with Cs+ again being used in the pipette to prevent any contribution of K+ channels to the current. As found previously, typical cell-attached patches were mostly silent, but addition of 100 μM diazoxide to the bath elicited single-channel events soon after addition of drug (Fig. 7A). Single-channel events recorded at different membrane potentials after diazoxide are shown in Figure 7B, with the magnitude of the open-channel current plotted as a function of membrane potential in Figure 7C. Diazoxide activated a single-channel conductance of ~35 pS that exhibited weak inward rectification when measured in the cell-attached configuration, similar to previous observations on channel activation accompanying depletion of [ATP], (Chen and Simard, 2001). Similar findings were made in 8 of 18 cell-attached patches exposed to diazoxide, whereas both pinacidil (10 μM; n = 7) and cromakalin (100 μM; n = 4) were without effect in similar cell-attached patch experiments, consistent with SUR1- but not SUR2-mediated channel activation.

We reported previously that Na azide activated the 35 pS NCa-ATP channel in the cell-attached configuration but not in outside-out patches with 1 mM ATP in the pipette, consistent with a mechanism involving ATP depletion rather than direct activation by Na azide (Chen and Simard, 2001). Having shown activation by diazoxide in a cell-attached configuration, we next...
sought to determine whether diazoxide would activate the channel in outside-out patches in the presence of ATP. When studied with 1 mM ATP and 1 mM Ca^{2+} in the pipette, and Cs^{+} on both inside and outside membrane surfaces, application of 1 mM Na azide to outside-out patches did not cause activation of any channel over the course of 2–3 min, as expected (Fig. 8A,B). After washout of Na azide, however, subsequent application of 100 μM diazoxide to the same patch resulted in brisk activation of channel activity (Fig. 8Ac,Bc), with partial loss of activity after washout of drug (Fig. 8Ad,Bd). Similar results were obtained in seven additional experiments. Thus, activation by diazoxide did not require a metabolically intact whole cell, consistent with diazoxide acting directly on the channel (Harvey et al., 1999), and in contrast to the action of Na azide, the effect of which is mediated via ATP depletion.

We also sought to show that the channel turned on in whole cells by ATP depletion was the same channel as that turned on by diazoxide. For these experiments, a nystatin perforated-patch technique was used to assure that metabolic disruption would come from drug application and not from cell dialysis. As reported previously (Chen and Simard, 2001), recordings obtained during step pulses showed significantly larger currents several minutes after Na azide (1 mM) compared with control (Fig. 9Ac,b). Subsequent application of 100 μM diazoxide (b), and after addition of 1 mM Na azide (c). Currents during step pulses under control conditions (a), after addition of 100 μM diazoxide (c), and after addition of 1 mM Na azide (b), and after addition of 1 μM glibenclamide (c).

We also performed the complementary experiment to show that current activation by Na azide-induced [ATP] depletion could be blocked by glibenclamide. As observed previously, when cells were studied using a whole-cell nystatin perforated-patch technique, exposure to Na azide (1 mM) caused activation of the NCCa-ATP channel (Fig. 9Ca,b). Subsequent addition of 1 μM glibenclamide caused full inhibition of the current that was turned on by Na azide (Fig. 9Cc). Together, these data suggest that sulfonylurea was acting on the same channel that was activated by Na azide-induced [ATP] depletion and that sulfonylurea could block the channel after it had been opened by [ATP] depletion.

**Tryptic digests**

Another characteristic feature of SUR-regulated K_{ATP} function is that tryptic digestion of the cytoplasmic but not extracellular face of the channel causes loss of inhibition by sulfonylureas without altering sensitivity to ATP and without changing the biophysical properties of the channel (Proks and Ashcroft, 1993; Lee et al., 1994). We thus assessed the effect of trypsin on the cytoplasmic side of the NCCa-ATP channel. Experiments were performed using an inside-out patch configuration with Cs^{+} on both sides of the membrane and 1 mM Ca^{2+} in the bath, which, as shown previously, produced robust channel activity (Fig. 10, first trace) that was strongly inhibited by 1 μM glibenclamide (Fig. 10, second trace). Subsequent washout of drug and exposure to 100 μg/ml trypsin for 3 min yielded a patch that still exhibited strong channel activity (Fig. 10, third trace) but was now completely unaffected by 1 μM glibenclamide (Fig. 10, fourth trace); however, the biophysical properties of the channel, including open-channel conductance, open-channel times, and Ca^{2+}-mediated activation, were unchanged, and the channel still maintained its typical sensitivity to ATP, showing reversible inhibition after addition of 1 mM ATP (Fig. 10, fifth and sixth traces). Similar observations were made in six patches. By contrast, exposure of the extracellular side to trypsin (four cells) had no effect on glibenclamide inhibition. These data on the NCCa-ATP channel paralleled exactly previous findings on K_{ATP} channels, giving additional supporting evidence that SUR1 was involved in regulation of the NCCa-ATP channel.
Anti-SUR1 antibody

To this point, all of our physiological and pharmacological data were consistent with the hypothesis that the pore-forming moiety of the NC$_{\text{Ca,ATP}}$ channel was functionally regulated by SUR1. To further test this hypothesis, we studied the effect of anti-SUR1 antibody on glibenclamide inhibition of the channel. Experiments were performed using an inside-out patch configuration with Cs$^+$ on both sides of the membrane and 1 mM Ca$^{2+}$ in the bath. As shown previously, channel activity was readily inhibited by addition of 1 mM glibenclamide to the bath (Fig. 11, trace 1). Subsequent washout of glibenclamide and application of anti-SUR1 antibody (final dilution, 1:1000; incubation time, 1 hr at room temperature) resulted in a patch with channel activity that was no longer inhibited by 1 mM glibenclamide (Fig. 11, trace 2). A similar finding was made in five patches from NR1A cells and in four patches from insulinoma cells (patches studied with KATP channel after [ATP]i depletion; data not shown). By contrast, application of anti-Kir6.1 antibody (final dilution, 1:1000; incubation time, 1 hr at room temperature) was without effect on glibenclamide inhibition in three patches from NR1As (Fig. 11, traces 3 and 4), showing that the effect of anti-SUR1 antibody was specific.

SUR-mediated cell swelling

We reported previously that in freshly isolated NR1As, ATP depletion after exposure to Na azide caused cell blebbing and swelling that arose after activation of NC$_{\text{Ca,ATP}}$ channels (Chen and Simard, 2001). Here we sought to determine whether SUR activators and sulfonylureas could be implicated in these pathological responses. As found previously, cells exhibited a stable morphological appearance when observed for 30 min or more (Fig. 12, control). After addition of Na azide (1 mM), cell blebbing typically became apparent after 7–10 min (Fig. 12, Na Azide). Opening of this channel by diazoxide (100 mM) resulted in blebbing of the cells even without [ATP]i depletion (Fig. 12, Diazoxide); however, when cells were pretreated with glibenclamide (1 mM), addition of Na azide did not result in any appreciable blebbing, even after 30 min or more of observation (Fig. 12, Na Azide + Glibenclamide). This finding accorded with previous observations that glibenclamide protected from opening of the NC$_{\text{Ca,ATP}}$ channel after [ATP]i depletion (Fig. 9) and that opening of this channel preceded cell blebbing (Chen and Simard, 2001).

Discussion

The principal findings of this study are as follows: (1) the NC$_{\text{Ca,ATP}}$ channel in NR1As was regulated by pharmacological and physiological mechanisms that mimicked exactly those described for SUR1 regulation of the K$_{\text{ATP}}$ channel in pancreatic β cells; (2) glibenclamide-mediated block of the NC$_{\text{Ca,ATP}}$ channel was prevented by anti-SUR1 antibody; and (3) NR1As exhibited specific sulfonylurea binding sites attributable to SUR1 but were not immunolabeled with anti-Kir6.1 or -Kir6.2 antibodies. The latter observations, coupled with the fact that our electrophysiological data were obtained using Cs$^+$ as the charge carrier, indicate that these cells do not possess K$_{\text{ATP}}$ channels. Rather, our data are consistent with the hypothesis that SUR1 regulates the NC$_{\text{Ca,ATP}}$ channel in NR1As, and they suggest that this novel NC$_{\text{Ca,ATP}}$ channel, like the K$_{\text{ATP}}$ channel in pancreatic β cells, may be a heteromultimer composed of a channel moiety distinct from the SUR1 regulatory subunit.

Sulfonylurea agents act not only on various isoforms of SUR but also on other ABC family members, including CFTR and P-glycoprotein (Sheppard and Welsh, 1992; Schultz et al., 1996; Golstein et al., 1999); however, effects at these different sites are generally distinguishable by virtue of potency and mechanism of block. The sulfonylurea inhibitor glibenclamide is most potent at SUR1, with EC$_{50}$ values of ~2–30 nM (Yamazaki and Hume, 1997), compared with ~150 nM at SUR2A, and micromolar values at CFTR and P-glycoprotein. Another sulfonylurea, tolbutamide, is also highly effective at SUR1, with EC$_{50}$ values of ~5–30 μM (Yamazaki and Hume, 1997), compared with ~120 μM at SUR2A, and higher values at CFTR and P-glycoprotein. Thus, the values observed here with the NC$_{\text{Ca,ATP}}$ channel, 48 nM and 12 μM for glibenclamide and tolbutamide, respectively, accord well
with the hypothesized involvement of SUR1 in regulation of the NC\textsubscript{Ca-ATP} channel. Moreover, sulfonylurea inhibition of native pancreatic K\textsubscript{ATP} channels or of SUR1–Kir6.2 channels occurs via an allosteric mechanism that results in an increase in the closed dwell time of the channel, with no effect on the open-channel dwell time. Similarly, with the NC\textsubscript{Ca-ATP} channel, sulfonylurea inhibition was characterized by an increase in closed dwell times with no change in the open dwell times. These effects on channel kinetics contrast with the open-channel blocking mechanism reported with sulfonylurea block of CFTR (Schultz et al., 1996; Venglarik et al., 1996). Thus, in terms of both potency and mechanism, sulfonylurea block of NC\textsubscript{Ca-ATP} channels was indistinguishable from observations made with pancreatic K\textsubscript{ATP} and SUR1–Kir6.2 channels.

K\textsubscript{ATP} channels in pancreatic β cells are heteromultimers formed from Kir6.2 and the sulfonylurea receptor SUR1. Sensitivity to sulfonylureas, which is conferred by SUR1, is lost by exposure of the cytoplasmic but not the extracellular side to trypsin. Trypsin is believed to act on SUR1 itself, rather than on Kir6.2, because trypsin causes loss of sulfonylurea binding, whereas the biophysical properties of the channel itself are preserved. We thus reasoned that if similar findings could be made with the NC\textsubscript{Ca-ATP} channel, this would further advance the hypothesis of involvement of SUR1 in regulation of the NCCa-ATP channel. Indeed, we found that exposure of the cytoplasmic side of the NC\textsubscript{Ca-ATP} channel to trypsin resulted in complete loss of sensitivity to glibenclamide, without altering channel opening, conductance, Ca\textsuperscript{2+}-mediated activation, or sensitivity to block by ATP. Our findings with trypsin demonstrate that sulfonylurea sensitivity of the NC\textsubscript{Ca-ATP} channel is conferred by a regulatory peptide moiety susceptible to tryptic cleavage and that this moiety is distinct from the ATP binding site and the channel pore, with the pore itself not being functionally altered by exposure to trypsin. These findings, which again mimic observations made on pancreatic K\textsubscript{ATP} channels, provide evidence linking trypsin-sensitive SUR and the NC\textsubscript{Ca-ATP} channel.

Apart from conferring sensitivity to sulfonylurea inhibitors, SUR1 also confers sensitivity to K\textsuperscript{+} channel openers (SUR activators). Notably, this diverse group of agents activates SUR-linked channels but inhibits CFTR- and P-glycoprotein-linked channels (Sheppard and Welsh, 1992), making the SUR activators potentially more useful than sulfonylurea inhibitors as indicators of involvement of SUR. Moreover, these activators are essentially isoform specific in their effects, permitting provisional identification of SUR isoform (Babenko et al., 2000; Lawson, 2000). Diazoxide is highly effective at SUR1, with an EC\textsubscript{50} value of ~50 μM, and is essentially ineffective at SUR2A, whereas pinacidil is highly effective at SUR2A, with an EC\textsubscript{50} value of ~10 μM, and is essentially ineffective at SUR1 (Fujita and Kurachi, 2000). Thus, our finding that 100 μM diazoxide readily activated the NC\textsubscript{Ca-ATP} channel, whereas pinacidil did not, strongly supports the hypothesis of involvement of SUR1 in regulation of the NC\textsubscript{Ca-ATP} channel.

Overall, the pharmacological characteristics of the NC\textsubscript{Ca-ATP} channel closely recapitulate the pharmacological profile of native pancreatic β-cell K\textsubscript{ATP} channels and of heterologously expressed Kir6.2–SUR1 channels. In addition, glibenclamide block of the channel was found to be prevented by application of anti-SUR1 antibody to the cytoplasmic side. The most parsimonious explanation for the constellation of findings reported here is that SUR1, identified at the protein and mRNA levels, functionally regulates the NC\textsubscript{Ca-ATP} channel. This implies that the NC\textsubscript{Ca-ATP} channel, like the K\textsubscript{ATP} channel, is a heteromultimer formed from SUR1 regulatory subunits and molecularly distinct pore-forming subunits.

The identity of the putative pore-forming subunit of the NC\textsubscript{Ca-ATP} channel remains to be determined; however, several features distinguish it from Kir6.x pore-forming subunits: (1) it is nearly equally permeable to K\textsuperscript{+}, Na\textsuperscript{+}, Rb\textsuperscript{+}, Li\textsuperscript{+}, and Cs\textsuperscript{+}; (2) its conductance of 35 pS is significantly less than the ~80 pS conductance of the K\textsubscript{ATP} channel; (3) it is regulated by intracellular Ca\textsuperscript{2+}; and (4) it does not bind Kir6.1 or Kir6.2 antibodies. It seems unlikely that the pore-forming subunit of the NC\textsubscript{Ca-ATP} channel would simply represent a post-translational modification of the Kir6.x subunit of a K\textsubscript{ATP} channel, because this would imply a major modification of the selectivity filter and conductive pore and would require addition of a Ca\textsuperscript{2+} regulatory site. Full molecular characterization will require cloning of the pore-forming subunit of the NC\textsubscript{Ca-ATP} channel. Nevertheless, the data presented here favor the hypothesis that SUR1 regulates a novel pore-forming subunit. Thus, the NC\textsubscript{Ca-ATP} channel becomes the first example of promiscuity of SUR1 outside of the Kir family of inward-rectifier K\textsuperscript{+}-selective channels.

The normal physiological role of the NC\textsubscript{Ca-ATP} channel remains to be determined, but some observations related to its function may be noted. Our imaging experiments showed that cells exhibiting glibenclamide binding were GFP positive, indicating that they are reactive astrocytes, but that they comprised only a subpopulation of reactive astrocytes, being found most prominently in the gliotic capsule surrounding the foreign body and to a lesser extent within the sponge itself, but not within the brain parenchyma, even in regions adjacent to the injury. Reactive astrocytes are found in abundance in adjacent brain (Perillan et al., 1999), but they do not exhibit glibenclamide binding. We reported previously that NRAs harvested from the gelatin sponge implant lose the NC\textsubscript{Ca-ATP} channel when they are maintained in vitro under standard culture conditions (Chen and Simard, 2001), and we have since found that these cells retain expression of this channel when cultured under hypoxic conditions (our unpublished observation). Thus, we speculate that NR1A may be specifically adapted for hypoxic conditions within the gliotic capsule that are not found in adjacent areas of brain. The function of these cells encapsulating the sponge is unclear, but nonetheless it is evident that they have migrated to the sponge and that most (>80%) of the cells present within the sponge, as identified by DAPI labeling, exhibit glibenclamide binding. Thus, there is the intriguing observation that migrating reactive astrocytes appear to express the NC\textsubscript{Ca-ATP} channel only as they cross the border out of properly oxygenated parenchyma into the hypoxic region surrounding the foreign body. Because migrating glial cells are believed to have to shrink, to reduce their cell volume, to permit migration within the brain (Ransom et al., 2001), presumably the converse, i.e., swelling, would halt migration. Thus, migrating reactive astrocytes may use the NC\textsubscript{Ca-ATP} channel to sense arrival in a hypoxic environment, opening the channel to initiate swelling and reverse the shrinkage imposed previously that allowed migration. Notably, in this process, these cells may well serve as a significant reservoir of water taken up from the extracellular space and stored intracellularly as “cytotoxic edema,” a well known response to injury that can have deleterious effects on CNS function.

References


