Bestrophin-2 mediates bicarbonate transport by goblet cells in mouse colon

Kuai Yu, Emory University
Rafael Lujan, Universidad de Castilla-La Mancha
Alan Marmorstein, University of Arizona
Sherif Gabriel, University of North Carolina, Chapel Hill
Criss Hartzell Jr., Emory University

Journal Title: Journal of Clinical Investigation
Volume: Volume 120, Number 5
Publisher: American Society for Clinical Investigation | 2010-05-03, Pages 1722-1735
Type of Work: Article | Final Publisher PDF
Publisher DOI: 10.1172/JCI41129
Permanent URL: http://pid.emory.edu/ark:/25593/ffxfd

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Accessed August 3, 2019 9:51 AM EDT
Anion transport by the colonic mucosa maintains the hydration and pH of the colonic lumen, and its disruption causes a variety of diarrheal diseases. Cholinergic agonists raise cytosolic Ca2+ levels and stimulate anion secretion, but the mechanisms underlying this effect remain unclear. Cholinergic stimulation of anion secretion may occur via activation of Ca2+-activated Cl– channels (CaCCs) or an increase in the Cl– driving force through CFTR after activation of Ca2+-dependent K+ channels. Here we investigated the role of a candidate CaCC protein, bestrophin-2 (Best2), using Best2–/– mice. Cholinergic stimulation of anion current was greatly reduced in Best2–/– mice, consistent with our proposed role for Best2 as a CaCC. However, immunostaining revealed that Best2 localized to the basolateral membrane of mucin-secreting colonic goblet cells, not the apical membrane of Cl–-secreting enterocytes. In addition, in the absence of HCO3–, cholinergic-activated current was identical in control and Best2–/– tissue preparations, which suggests that most of the Best2 current was carried by HCO3–. These data delineate an alternative model of cholinergic regulation of colonic anion secretion in which goblet cells play a critical role in HCO3– homeostasis. We therefore propose that Best2 is a HCO3– channel that works in concert with a Cl:HCO3– exchanger in the apical membrane to affect transcellular HCO3– transport. Furthermore, previous models implicating CFTR in cholinergic Cl– secretion may be explained by substantial downregulation of Best2 in Cfr+/– mice.

The molecular identity of CaCCs has been elusive, but candidates include members of the bestrophin, CLCA, and TMEM16 families as well as CIC-3 (14–16). Among these candidates, bestrophins have received considerable attention for their roles in a variety of epithelia, including retinal pigment epithelium, airway, and gastrointestinal tract (14, 17). Best1 is expressed in proximal colon in mice and in human colonic epithelial cell lines (7, 8). Recently, we generated a mouse in which the first exons of the Best2 gene were replaced with LacZ and noted that murine Best2 was strongly expressed in colon (18). In the present study, our initial goal was to test the hypothesis that Best2 is an apical CaCC that mediates Cl– secretion. We expected that Best2 would be expressed on the apical surface of enterocytes in the colonic crypt, but were surprised to find that Best2 was expressed in the basolateral membrane of mucin-secreting goblet cells. We also found that Best2 was unlikely to participate in colonic Ca2+-activated Cl– secretion, but participated in colonic HCO3– secretion concomitant with mucin secretion. We therefore propose that HCO3– secretion plays a role in mucin expulsion from the crypt into the colonic lumen. Our findings also showed that colonic Ca2+-activated anion secretion was carried largely by HCO3–, and not by Cl–, as previously supposed.

Results

Lac-Z under control of the Best2 promoter is expressed in a proximal-distal gradient in the colon

Best2–/– mice were constructed with the Lac-Z gene replacing exon 2 and parts of exons 1 and 3 of the Best2 gene, so that the Lac-Z reporter (with a nuclear localization signal) was under control of the Best2 promoter (18). The expression of the mouse Best2 gene was evaluated by examining the expression of Lac-Z by X-gal staining. In a 2-month-old mouse, there was a gradient of expression.
from the cecum, which did not stain at all, to the rectum (Figure 1A). In younger mice, the level of expression was lower, and the staining of the distal colon was reduced compared with the older mice (Figure 1B). Staining localized in the nuclei of cells throughout the crypts (Figure 1C). The Best2−/− mouse was a true Best2 null, as shown by Western blot: extracts from WT colon or from mouse Best2-transfected HEK cells exhibit a 58-kDa band expected for Best2, whereas extracts from Best2−/− mice and untransfected HEK cells did not have the Best2 band (Figure 1D).

**Best2 is expressed in a subset of colonic cells in mice and humans**

To determine which cell types express Best2 in WT mice, the knockout-verified antibody was used to immunostain sections from a region of colon midway between anus and cecum. A subset of cells exhibited intense Best2 staining on the basal and lateral membranes (Figure 2A). These cells were located at all levels of the crypt as well as on the luminal brush border surface. At higher magnification (Figure 2, B and C), Best2-positive cells exhibited morphology typical of goblet cells (19). The Best2-positive cells at the tops of the crypts generally had a goblet-like appearance, with an expanded apical end presumably filled with mucin and a cell body that narrows as it approaches the basal lamina. Best2-positive cells deeper in the crypts had a more extensive basal surface and tapered apically. The Best2-positive cells did not exhibit apical actin staining with phalloidin (Figure 2, C and D) or with CFTR antibody (Figure 2E), whereas the apical membranes of the Best2-negative enterocytes, both in the crypt and at brush border of the luminal surface, stained intensely with phalloidin and with CFTR antibody. The relative absence of phalloidin staining of the apical membrane is consistent with the fact that goblet cells have fewer actin-containing microvilli than do enterocytes (20).

There are 3 bestrophin paralogs in mice and 4 in humans (Best1–Best3 and BEST1–BEST4, respectively; ref. 14). BEST2 and Best2 are 90% identical, and thus would be expected to function similarly. Our previous studies show that Best2 and BEST2 produce similar whole-cell Ca2+-activated anion currents (21). We immunostained human colon sections with specific anti-BEST2 antibodies and found that, like Best2, BEST2 was localized in the basolateral plasma membrane of a subset of colonic cells (Figure 2, G and H).

In human tissue, there was a considerable amount of intracellular staining at the basal end of the cells; this staining was punctate and appeared to be localized in vesicles (Figure 2H). In mouse colon, there was similar punctate intracellular staining in some cells (Figure 2D), but less than that observed in humans.

**Best2-positive cells are goblet cells**

To verify that the Best2-positive cells are goblet cells, we costained sections with antibodies against Best2 and the goblet cell mucin Muc-2. Muc-2 antibody intensely stained the apical portion of cells that were positive for Best2 (Figure 3A). The identity of these cells as goblet cells was confirmed using lectin staining. During biogenesis, mucins are heavily O-glycosylated with N-acetylgalactosamine and galactosyl(β-1,3)N-acetylgalactosamine at serine- and threonine-rich tandem repeat domains. These core sugars become heavily sialylated in mature mucin, which can be stained with *Sambucus nigra* agglutinin (SNA) and *Maackia amurensis* lectin–II (MAL-II), which recognize sialic acid. The apical region of Best2-positive cells stained with both SNA and MAL-II (Figure 3, B and C), confirming the identity of these cells as goblet cells.

Additional evidence that Best2 is expressed in the plasma membrane of goblet cells was provided by immunoelectron microscopy of goblet cells (Figure 3, D and E). There was no immunostaining in Best2−/− mice (data not shown), whereas in WT mice, immunogold particles were abundant on basal and lateral plasma membranes and in endoplasmic reticulum (Figure 3, F and G).

**Best2 mediates a Ca2+-activated and HCO3−-dependent current in colonic epithelium**

The finding that Best2 was localized in the basolateral membrane of goblet cells was surprising, because we expected that Best2 was a CaCC located in the apical membrane of enterocytes involved in Ca2+-stimulated Cl− secretion. To investigate the role of Best2 in colonic ion transport, we performed short-circuit current (Isc) analysis on pieces of distal colonic mucosa stripped of smooth muscle mounted in Ussing chambers. Transepithelial currents were recorded with the transepithelial voltage clamped to 0 mV. CFTR-mediated currents were activated by elevation of cAMP by forskolin application to the apical membrane of a subset of colonic cells (Figure 2, G and H).
cal (luminal) side, and Ca\(^{2+}\)-activated currents were activated by carbachol (CCh) application to activate G\(_{q}\)-coupled muscarinic AChRs on the basal (serosal) side.

Reduced CCh-activated \(I_\text{sc}\) in Best2\(^{-/-}\) mice. After equilibration of the mucosa, but before application of drugs, the baseline \(I_\text{sc}\) recorded in normal Krebs solution was the same in WT (9.8 ± 0.9 \(\mu\)A/cm\(^2\); \(n = 19\)) and Best2\(^{-/-}\) (12.0 ± 1.3 \(\mu\)A/cm\(^2\); \(n = 13\)) mucosa. Amplitudes of forskolin-activated \(\Delta I_\text{sc}\) were also the same in WT and Best2\(^{-/-}\) mice (40.6 ± 4.0 and 45.1 ± 7.1 \(\mu\)A/cm\(^2\), respectively; Figure 4A and C), whereas the CCh-activated currents were significantly smaller in Best2\(^{-/-}\) compared with WT mice (26.1 ± 0.7 versus 49.4 ± 4.5 \(\mu\)A/cm\(^2\); Figure 4B and D). The finding that forskolin-activated currents were not affected in the Best2\(^{-/-}\) mouse is consistent with our observation that CFTR immunostaining was the same in WT and Best2\(^{-/-}\) mice (data not shown) and showed that CFTR functions normally in Best2\(^{-/-}\) mice. The decrease in CCh-activated current in the Best2\(^{-/-}\) mouse colon is consistent with Best2 playing a role in this current.

CCh-activated current is composed of multiple components. Our previous studies have shown that Best2 has high permeability to HCO\(_3\)\(^{-}\) (21). For this reason, we investigated whether the CCh-activated current in the colon was carried by HCO\(_3\)\(^{-}\). HCO\(_3\)\(^{-}\) was removed from the serosal Krebs solution, and pH and ionic strength was maintained by addition of Na-HEPES and Na-glucuronate. As shown in Figure 4D, the recording was begun in HCO\(_3\)\(^{-}\)-containing solution that also contained Ba\(^{2+}\) and amiloride to block other cation channels. CCh stimulated a current that had a transient peak lasting approximately 1 minute and a prolonged plateau that typically lasted more than 15 minutes. When HCO\(_3\)\(^{-}\) was removed, the plateau current was completely abolished. Restoring HCO\(_3\)\(^{-}\) to the basal chamber restored the current. These results suggest that CCh stimulates 2 currents, a transient Cl\(^{-}\} current and a plateau HCO\(_3\)\(^{-}\} current.

The HCO\(_3\)\(^{-}\} component is mediated by Best2. To test this hypothesis more rigorously and to determine whether the CCh-activated currents were mediated by Best2, we examined the effect of HCO\(_3\)\(^{-}\} removal on preparations in which basolateral K\(^{+}\} channels were blocked by 5 mM Ba\(^{2+}\} and apical ENaC was blocked with amiloride. As above, under these conditions, CCh activated a current that rose to a peak before decaying to a steady plateau (Figure 4E). The CCh-activated current was smaller in Best2\(^{-/-}\) than in WT mucosa (Figure 4G). HCO\(_3\)\(^{-}\} removal greatly reduced the CCh-activated current in WT mucosa (Figure 4F). The greatest effect of HCO\(_3\)\(^{-}\} removal was seen on the plateau, with a smaller effect on the initial upstroke, which suggests that the plateau current was carried by HCO\(_3\)\(^{-}\} and that the initial component was carried by other ions, most likely Cl\(^{-}\}. Conversely, in Best2\(^{-/-}\} mucosa, HCO\(_3\)\(^{-}\} removal had a very small, statistically insignificant effect on the CCh-activated current (Figure 4G). These data support the conclusion that a component of the CCh-activated current is carried by HCO\(_3\)\(^{-}\} through Best2 channels. This conclusion is further supported by the observation that the CCh-activated currents recorded in the absence of HCO\(_3\)\(^{-}\} were identical in WT and Best2\(^{-/-}\} mucosa (Figure 4H).

The amplitudes of the Cl\(^{-}\} and HCO\(_3\)\(^{-}\} currents are quantified in Figure 4I. Cl\(^{-}\} current was defined as the current in HCO\(_3\)\(^{-}\} -free solution with Ba\(^{2+}\} and amiloride. The HCO\(_3\)\(^{-}\} current was defined in 2 different ways. (a) The amplitude of the current in HCO\(_3\)\(^{-}\} -containing Krebs solution at 10 minutes after CCh addition (i.e., the plateau) was carried largely by HCO\(_3\)\(^{-}\} because the current in HCO\(_3\)\(^{-}\} -free solution was transient and returned to baseline within 10 minutes of CCh application. (b) The difference between the amplitudes of the peak currents (approximately 3 minutes) in normal Krebs (carried by both Cl\(^{-}\} and HCO\(_3\)\(^{-}\}) and in HCO\(_3\)\(^{-}\} -free solution (carried largely by Cl\(^{-}\}) should represent...
the HCO₃⁻ current. The HCO₃⁻ current amplitude determined by these 2 methods agreed surprisingly well and showed that the HCO₃⁻ current in Best2⁺/⁻ mucosa was only 33% as large as WT.

The HCO₃⁻ current is carried mainly by Best2
Although Ca²⁺ clearly activates anion secretion in colon, there is some question as to whether this is mediated by CaCCs or by CFTR. It has been proposed that CFTR channels are opened by elevation of cAMP produced by prostaglandins that are synthesized in response to tissue dissection and manipulation. Under these conditions of basal CFTR activation, Ca²⁺ may stimulate Cl⁻ flux through these open CFTR channels by activating Ca²⁺-activated K⁺ channels, hyperpolarizing the membrane, and increasing the driving force for Cl⁻ secretion (see Discussion). To determine whether the Ca²⁺-stimulated HCO₃⁻ current requires CFTR, we performed Iₑ analysis on colons from Cfr⁻/⁻ mice (homozygous Cfr⁻/⁻). In Cfr⁻/⁻ colons, both the total CCh-stimulated current and the HCO₃⁻-dependent component were significantly smaller than in WT colons (Figure 5A). These results are consistent with the suggestion that CFTR may be necessary for the CCh-stimulated current. However, we wondered whether the apparent dependence of the CCh-stimulated currents on CFTR could be explained by downregulation of Best2 in the Cfr⁻/⁻ mice. To test this idea, Western blots were performed on samples of the same colons studied in the Iₑ experiments (Figure 5B). Of the samples from Cfr⁻/⁻ mice, 2 gave little or no Best2 signal, whereas the other 2 samples gave weaker signals compared with the control mice. Immunofluorescence microscopy confirmed that the expression of Best2 was altered in Cfr⁻/⁻ mice. In Cfr⁻/⁻ mice, Best2 staining was weaker when viewed at identical settings and gains. However, more importantly, unlike WT colon, in which Best2 staining was intensely localized in the basolateral plasma membrane (Figure 5, C and D), in Cfr⁻/⁻ mice, Best2 staining was not sharply localized to the plasma membrane (Figure 5, E–H). In Cfr⁻/⁻ mice, Best2 staining was weakly and diffusely distributed in the cytoplasm (compare asterisks in Figure 5, D, F, and H). In addition, in Cfr⁻/⁻ mice, there was a considerable amount of Best2 accumulated in what appeared to be aggregates in the interstitium between crypts and the smooth muscle layer (arrowheads, Figure 5, E–G). These displaced accumulations of Best2 were never seen in WT tissue. These results suggest that the difference between WT and Cfr⁻/⁻ mice can be explained by an alteration in Best2 expression in the Cfr⁻/⁻ mice. These results raise the possibility that CFTR itself may not be required for Ca²⁺-stimulated Cl⁻ secretion.

We therefore reexamined other evidence supporting the idea that Ca²⁺ stimulates Cl⁻ secretion through CFTR. Evidence that the Ca²⁺-activated Cl⁻ current is carried by CFTR has been supported by the observation that the response to CCh is blocked when prostaglandin synthesis is inhibited by indomethacin (10, 11). Prostaglandins generated during tissue manipulation are believed to be responsible for a basal level of CFTR activation, but we considered an alternative possibility: namely, that indomethacin may directly block currents carried by Best2. In HEK-293 cells expressing Best2, we found that indomethacin at concentrations typically used by others in Ussing chamber experiments (10 μM) blocked Best2 currents by approximately 80% (Figure 6, B and E). We also measured the effect of indomethacin on currents generated by another candidate CaCC channel, anoctamin 1 (Ano1; see below). Indomethacin did not block Ano1 currents at the same concentration (Figure 6, D and E).

There is considerable evidence that CCh activates Ca²⁺-activated K⁺ channels (e.g., K₅Ca3.1, SK4, and IK1) reported to be necessary for CCh-stimulated Cl⁻ secretion (e.g., ref. 12). Activation of these channels could result in increased Cl⁻ efflux through CFTR (if CFTR channels were open). One line of evidence suggesting that K₅Ca3.1 channels are necessary for Ca²⁺-activated Cl⁻ secretion is the observation that clotrimazole, a blocker of K₅Ca3.1 channels, blocks the CCh-stimulated Cl⁻ current (12). We found that clotrimazole also blocked Best2 currents by approximately 80% and Ano1 channels expressed in HEK-293 cells by 70% (Figure 6, A and C). The block of Best2 channels is more slowly reversible than the block of Ano1 channels. Taken together, these data show that CFTR is unlikely to contribute significantly to the CCh-stimulated HCO₃⁻ current in mouse colon.

Ca²⁺-activated anion channels in isolated colonocytes
CCh activates a fast Cl⁻ current in addition to the slow HCO₃⁻ current carried by Best2. To gain additional insight into the channels that carry the Cl⁻ component, we performed whole-cell patch clamp recording on freshly isolated cells isolated from distal colon. These isolated colonocytes are expected to be a mixture of goblet cells and enterocytes. Cells patch-clamped with less than 20 nM free [Ca²⁺], in
the pipette had very small currents (Figure 7, A and D), but large Cl- currents were recorded with approximately 1 μM free \([Ca^{2+}]\) (Figure 7, B–D). We studied a total of 25 WT cells and 30 Best2–/– cells with high \([Ca^{2+}]\) solution, and observed 2 different current waveforms in different cells. One linear and time-independent waveform (Figure 7C) was observed in 36% of WT cells and 43% of Best2–/– cells, although the current was much smaller in Best2–/– cells. The percentage of cells with this current waveform roughly corresponded to the proportion

**Figure 4**

I_{lc} in WT and Best2–/– distal colonic mucosa. Upward currents represent anion movement from serosa to lumen. (A and B) Representative I_{lc} in response to 10 μM forskolin (A) and 1 mM CCh (B) in WT and Best2–/– mice. (C) Peak I_{lc} amplitudes to forskolin or CCh in WT and Best2–/– mice (n = 6–10). *P < 0.05 versus WT, Student's t test. (D–I) I_{lc} mediated by Cl– and HCO3–. Solutions contained 5 mM Ba2+ and 0.1 mM amiloride. (D) Effect of HCO3– removal on CCh-stimulated current in WT mucosa. In normal Krebs solution, CCh induced a rapid peak followed by plateau. HCO3– removal inhibited the plateau. (E) Average I_{lc} in WT and Best2–/– mice. CCh was added at 2 minutes. (F and G) CCh-stimulated current in WT (F) and Best2–/– (G) mucosa in normal Krebs and HCO3–-free solutions. (H) Replot of data in F and G comparing Cl– currents in HCO3–-free Krebs in WT and Best2–/– mice. (I) Comparison of Cl– and HCO3– currents stimulated by CCh in WT and Best2–/– mucosa. Total current in normal Krebs; Cl, Cl– current in HCO3–-free solution; HCO3(P), amplitude of plateau current 8 minutes after CCh application in normal Krebs; HCO3(D), difference between currents at peak (about 1–2 minutes after CCh application) in normal Krebs and in HCO3–-free solution. n = 10 per data point.
of goblet cells existing in this region of the colon (e.g., Figure 2A). We believe that the cells with linear currents are goblet cells, because the linear current resembled that induced by Best2 expressed in HEK cells (Figure 7, compare C and H), and its amplitude was reduced 60% in Best2–/– cells (Figure 7F). The presence of a small linear current in the knockouts may be explained by an uncompensated leak current in some cells. The other waveform resembled a classical outwardly rectifying, time-dependent CaCC current (Figure 7B). The amplitudes of the outwardly rectifying currents were statistically the same in WT and Best2–/– mice (Figure 7E). The outwardly rectifying current was most likely mediated by the newly discovered Ano1 (TMEM16A) channel (22–24), because it resembled the currents induced by Ano1 expression in HEK cells (Figure 7, compare B and G). Transcripts for Ano1, Ano6, Ano7, and Ano10 were expressed in colonic mucosa (Figure 8A). Western blots showed that HEK293 cells transfected with Ano1 had a faint band at the expected mass of 110 kDa and a larger, diffuse band representing glycosylated protein (Figure 8B). A similar diffuse band was observed in extracts from distal colon. Staining frozen sections with the Ano1 antibody showed intense staining in interstitial cells of Cajal, as previously reported by Gomez-Pinilla et al. (25), as well as in cells of the brush border membrane (Figure 8C).

To determine whether Ano1 contributes to HCO3− permeability, we measured the relative HCO3− permeability and conductance of the Ano1 channel (Figure 7I). Ano1 had low permeability to HCO3− compared with Best2 (HCO3−/Cl− ratio, 0.3 versus 0.7). These data support the conclusion that Ano1 is responsible for a Ca2+-activated Cl− current in colon, whereas Best2 is responsible for a Ca2+-activated HCO3− current.

Figure 5
Best2 expression in colon from Ctr−/− mice. (A) Isc from Ctr−/− and WT mice, recorded as in Figure 4. Shown are results with normal Krebs and HCO3−-free solutions as well as the difference between them. n = 5 mucosal samples from 2 WT mice; n = 9 mucosal samples from 4 Ctr−/− mice. P < 0.01, all Ctr−/− versus all respective WT values. (B) Western blot of Best2 expression in WT (WT1 and WT2) and Ctr−/− (CF1–CF4) colon samples. Samples were the same ones used for Isc measurements in A. Blots were stained with Best2 (58 kDa) and GAPDH (37 kDa) antibodies. (C–H) Immunofluorescence of WT (C and D) and Ctr−/− (E–H) colon samples. Samples are adjacent tissue from the same colon used for Isc measurements in A. Arrowheads in E and F indicate interstitial accumulation of Best2 staining; asterisks in D, F, and H are shown for comparison of Best2 staining in the cytoplasm for comparison. Scale bars: 20 μm (C, E, and G); 10 μm (D, F, and H).
WT mice. \textit{Best2}⁻/⁻ mice lost an average of 10.2% ± 3.4% of their body weights after treatment with DSS, whereas WT mice lost only 3.7% ± 1.2% (Figure 9F). At the end of the 6-day DSS treatment period, MPO was substantially elevated in both groups, although the responses of the 2 groups were not significantly different (Figure 9G). \textit{Best2}⁻/⁻ mice recovered from DSS treatment less quickly than did WT mice: 5 days after DSS treatment, the \textit{Best2}⁻/⁻ mice had not regained the body weight they had lost, whereas WT mice were approximately 5% above their starting weight (Figure 9F). Furthermore, at 18 days after DSS treatment was ended, MPO levels in WT mice had returned to control levels, whereas MPO levels in \textit{Best2}⁻/⁻ mice remained 4.8-fold higher than the WT control level (Figure 9G).

Because there was such a large difference between groups in their recovery from DSS treatment, we examined the distribution of goblet cells 1 week after cessation of DSS treatment. In WT animals, there was tremendous hyperplasia of \textit{Best2}-positive goblet cells (Figure 9H). In some sections, it appeared that nearly every cell in the crypt was positive for \textit{Best2}. In colons prepared in parallel, control sections had less than 15 \textit{Best2}-positive cells per crypt (range, 8–15 cells), whereas DSS-recovery sections had greater than 25 positive cells (range, 25–40 cells). In addition, many of the cells at the brush border surface were also \textit{Best2}-positive (approximately 10 cells per 100-µm colon length), whereas in control colons, surface \textit{Best2}-positive cells were seen infrequently (less than 1 cell per 100-µm colon length). Because we cannot use \textit{Best2} staining to compare WT and \textit{Best2}⁻/⁻ goblet cell hyperplasia, we used peanut agglutinin (PNA), which recognizes the O-linked core galactosyl(β-1,3)N-acetylgalactosamine to label mucin. Virtually every \textit{Best2}-positive cell in the WT colon recovering from DSS treatment also stained with PNA (Figure 9H). In contrast, colon from \textit{Best2}⁻/⁻ mice recovering from DSS treatment exhibited little PNA staining (Figure 9I). These data clearly demonstrate that either mucin biogenesis or goblet cell differentiation is abnormal in \textit{Best2}⁻/⁻ colon when the animal is subjected to stress.
Figure 7
Anion currents recorded from freshly isolated colonocytes and transfected HEK293 cells. (A–C) Sample traces from isolated colonocytes with less than 20 nM free [Ca\(^{2+}\)] (A) or 1 \(\mu\)M [Ca\(^{2+}\)] (B and C), showing different waveforms observed in different cells. (D) Current-voltage relationships for WT colonocytes with less than 20 nM free [Ca\(^{2+}\)] as well as WT and Best2\(^{-/-}\) colonocytes with 1 \(\mu\)M free [Ca\(^{2+}\)]. (E) Comparison of current-voltage relationships of outwardly rectifying currents as in B from WT and Best2\(^{-/-}\) mice with 1 \(\mu\)M free [Ca\(^{2+}\)]. (F) Comparison of current-voltage relationships of linear currents as in C from WT and Best2\(^{-/-}\) mice with 1 \(\mu\)M free [Ca\(^{2+}\)]. (G) Representative currents in HEK cells transfected with Ano1 cDNA with 1 \(\mu\)M free [Ca\(^{2+}\)]. (H) Representative currents in HEK cells transfected with Best2 cDNA with 1 \(\mu\)M free [Ca\(^{2+}\)]. (I) Permeability and conductance, relative to Cl, of Ano1 and Best2 currents.
Discussion

These studies provide a number of insights. First, the function of bestrophins has been enigmatic since they were proposed to be Cl\(^{-}\) channels in 2002 (14, 16). For the first time to our knowledge, our results demonstrate a role of bestrophins in HCO\(_3\)- transport and open a window into the physiology of this family of channels. Second, our results provide insights into the poorly understood ion transport properties of colonic goblet cells, which are of fundamental importance in colonic physiology. Finally, we provide perspective on the regulation of colonic anion transport by proposing what we believe to be a novel mechanism for the role of cholinergic agonists.

**Hypothesis.** The data presented here show that Best2 is expressed on the basolateral membrane of goblet cells and that disruption of Best2 causes a decrease in a CCh-activated HCO\(_3\)-dependent transepithelial current. Because Best2 is highly permeant to HCO\(_3\), when expressed heterologously in HEK cells (21), and HCO\(_3\)-dependent transport was reduced in the Best2\(^{-/-}\) mouse, we conclude that Best2 is a channel that mediates vectorial transepithelial secretion of HCO\(_3\). However, we should point out that we have not measured HCO\(_3\) secretion directly. Given the fact that Best2 may have other regulatory functions (14), it is possible that the HCO\(_3\) dependence of the transepithelial current might be caused by an effect of HCO\(_3\) on some other function of Best2. Also, it is important to note that Best2 is not only found on the plasma membrane, but also in intracellular organelles (Figure 2). Best2 may, therefore, have intracellular functions. Recently, it has been reported that Best1 is an ER protein that regulates ER Ca\(^{2+}\) transport (27).

How could passive movement of HCO\(_3\) across the basolateral membrane of the goblet cell mediate transepithelial HCO\(_3\)- transport? Our hypothesis is summarized in Figure 10B. In contrast to the classical view of ion transport in distal colon (Figure 10A), we propose that HCO\(_3\) moves passively across the basolateral membrane and is then transported into the colonic lumen by an apical Cl\(^{-}\)-HCO\(_3\) exchanger. Whether this is thermodynamically feasible depends upon some unknown variables, notably the potentials across the basolateral and apical membranes of goblet cells. However, we can calculate the range of membrane potentials that would permit such a mechanism to operate. We assume that serosal HCO\(_3\) is similar to plasma (approximately 25 mM). [HCO\(_3\)], levels have been experimentally determined to be 8–15 mM in colonocytes at normal arterial pCO\(_2\) (28). If these values also apply to goblet cells, the Nernst equilibrium potential for HCO\(_3\) across the basolateral membrane would range from –30 mV at 8 mM [HCO\(_3\)]\(_i\) to –13 mV at 15 mM [HCO\(_3\)]\(_i\). The transepithelial potential (TEP) of distal colon in our studies is typically –1 to –2 mV (apical negative relative to serosal; see also refs. 2, 12). This value is less than other epithelia, including the small intestine, partly because the colonic epithelium is more leaky; moreover, the high proportion of goblet cells in distal colon may contribute to the lower TEP. In any case, such a small TEP would dictate that the apical membrane potential must be close to the basolateral membrane potential. If secretion of HCO\(_3\) into the colonic lumen is powered by an apical Cl\(^{-}\)-HCO\(_3\) exchanger, the equilibrium luminal [HCO\(_3\)] concentration would be calculated as follows: ln([HCO\(_3\)]\(_i\)/[HCO\(_3\)]\(_o\))\(_m\) = ln([Cl\_]\(_i\)/[Cl\_]\(_o\))\(_m\) + \(\frac{\eta \times [Cl\_]\(_o\)}{[Cl\_]\(_i\)}\) – \(\frac{\eta \times [Cl\_]\(_o\)}{[HCO\(_3\)]\(_i\)}\) x \(\frac{F}{RT}\), in which n and m represent the stoichiometry of the Cl\(^{-}\):HCO\(_3\) exchanger, z is the valence (i.e., –1), V\(_m\) is the membrane potential, and F, R, and T have their usual thermodynamic meanings (29). With total [anion] of 160 mM, [HCO\(_3\)]\(_i\) of 10 mM, apical V\(_m\) of ~25 mV, and [Cl\_]\(_o\) of 40 mM, the luminal [HCO\(_3\)] would be predicted to be 29–37 mM, depending on the Cl\(^{-}\):HCO\(_3\) stoichiometry of the exchanger (29). Thus, a basolateral HCO\(_3\) channel and an apical Cl\(^{-}\):HCO\(_3\) exchanger could theoretically drive secretion of significant amounts of HCO\(_3\). HCO\(_3\) could be secreted even if the goblet cell basolateral membrane potential were more negative than –30 mV; however, a more negative membrane potential would result in an equilibrium [HCO\(_3\)] lower than 8 mM, which would likely produce an unphysiologically acidic intracellular pH. Whether these calculations reflect reality is limited by the little data available on the membrane properties of colonic goblet cells; we do not know what additional conductances are present in the apical and basolateral membranes.

HCO\(_3\) is thought to be important in secretion of mucins, which play a key role in the innate immune system of the gut. Mucins are stored in intracellular granules in a compacted form with high concentrations of Ca\(^{2+}\) and protons (30) that screen the negative charges in the highly polyanionic mucins. When mucins are exocytosed, they rapidly expand in volume by 1,000-fold (31). This
expansion is thought to be driven in part by electrostatic repulsion of negative charges that are unmasked during exocytosis. It has been suggested that HCO₃⁻ plays a critical role in unmasking these charges by raising the pH and sequestering Ca²⁺ (32).

Mechanisms of anion secretion in distal colon. Activation of CFTR, expressed in the apical membrane of enterocytes in the crypts (Figure 2E), is responsible for Cl⁻ secretion activated by agonists that elevate cAMP (5). Agonists that elevate [Ca²⁺], also stimulate anion secretion (e.g., refs. 6–8), but the mechanisms are unclear (3). Although it seems reasonable to suppose that Ca²⁺-stimulated Cl⁻ secretion is mediated by CaCCs, various evidence suggests that Ca²⁺ stimulates Cl⁻ secretion via CFTR.

Evidence supporting a role for CaCCs in Ca²⁺-stimulated secretion. We note 5 lines of evidence in support for this role. (a) CaCCs have been previously described in acutely isolated colonocytes (33) as well as a variety of colonic cell lines, including T84, Caco2, and HT29 (34). (b) Although knockout of CFTR abolishes CAMP-induced volume decrease of colonic crypt cells mediated by CFTR, knockout of CFTR has no effect on Ca²⁺-induced volume decrease mediated by 4,4′-diisothiocyanatostilbene-2,2′-disulfonic acid–sensitive CaCC channels (35). (c) Although knockout of CFTR abolishes Ca²⁺-activated Cl⁻ currents in many mouse strains, several inbred strains of mice have been described in which Ca²⁺-activated Cl⁻ secretion is present even though the

Figure 9
Phenotype of Best2⁻/⁻ mice. (A and B) Body weights of male (A) and female (B) WT and Best2⁻/⁻ mice plotted versus age. (C–E) Hematoxylin and eosin–stained sections of WT (C) and Best2⁻/⁻ (D and E) colon. The circle in D indicates an area of inflammation characterized by many polymorphonuclear lymphocytes. Arrows in E denote lymphocytes (black) and an eosinophil (yellow). (F–I) Response to DSS treatment. (F) Body weights of WT and Best2⁻/⁻ mice given 3% DSS in their drinking water from day 1 to day 7 (n = 18 per group). (G) MPO activity in colon samples from WT and Best2⁻/⁻ mice before DSS treatment (control), at the end of 6 days of DSS treatment (DSS), and 18 days after DSS treatment was ended (recovery) (n = 6–10 per group). *P < 0.05. (H and I) Cells of WT and Best2⁻/⁻ distal colon, after 6 days of recovery from DSS, were stained with Best2 antibody (green) and PNA (red) to show unsialylated mucin. Scale bar: 20 μm. Images demonstrate hyperplasia of Best2-positive cells (H), whereas Best2⁻/⁻ mice had many fewer PNA-positive cells (I).
CFTR gene has been disrupted. This suggests that CaCC channel expression might be strain specific (36–38). (d) A subset of human cystic fibrosis patients exhibit residual colonic Cl– conductance that is likely mediated by CaCCs (39). (e) Rotavirus infection produces diarrhea by stimulating Ca2+-dependent Cl– secretion from crypts (40), and the rotavirus toxin NSP4 induces Ca2+-mediated Cl– secretion in both WT and CFTR-knockout crypts (41).

**Evidence against a role for CaCCs.** There are 3 lines of evidence that support the idea Ca2+ activates Kcnq3.1 channels that hyperpolarize the membrane and increase the driving force for anion secretion through CFTR (10–13). (a) It has been proposed that endogenous prostaglandin production stimulates adenylyl cyclase and that CFTR is partially activated under resting conditions. This idea is supported by the observation that suppression of prostaglandin production with indomethacin blocks CCh-stimulated Lm (10, 11). However, we found here that indomethacin also blocked Best2. (b) The observation that the CCh-activated Lm is potentiated by elevation of cAMP suggests a role for CFTR in Ca2+-stimulated Cl– secretion (10). However, the synergistic interaction between Ca2+ and cAMP could occur at a site other than CFTR. For example, because cAMP activates Kv7.1 channels, the resulting hyperpolarization would be expected to increase the driving force for Cl– efflux through CaCCs. (c) CCh-activated Lm is abolished or reduced in many mouse models of cystic fibrosis (42, 43) and in human biopsies from CF patients (11, 44, 45); however, as noted above, other studies have obtained different results. The differences may be explained by changes that occur as a secondary consequence of disruption of the gene encoding CFTR. CFTR is known to regulate a variety of ion channels and transporters, including CaCCs (46), and there may simply be compensatory changes in expression of other channels, including Best2. Finally, the idea that membrane hyperpolarization is required for a cholinergic response is not universal: cholinergic agonists can stimulate anion secretion when the membrane potential is clamped at a depolarized level (6) and when Kcnq3.1 channels are blocked pharmacologically (9). Although species differences may explain some discrepancies, the mechanisms of cholinergic action in colon are clearly murky.

**Figure 10**

Ion transport in distal colon. (A) Transporters and channels in colon. The distribution of channels and transporters was determined by immunofluorescence confocal microscopy. Secretion occurs in the crypt, and absorption occurs at the luminal brush border. Secretion is driven by transepithelial Cl– transport that occurs by active basolateral uptake of Cl– by the Na+/K+/2Cl– cotransporter NKCC1 (blue) and subsequent passive efflux via apical CFTR Cl– channels (green). Cl– transport is accompanied — paracellularly and possibly transcellularly — by H2O and Na+ (gray). Na+ and Cl– are then reabsorbed at the brush border surface by coupled Na+-H+ exchange (by NHE3; orange) and Cl–-HCO3– exchange (by SLC26A3; magenta) coupled to carbonic anhydrase (CA). Best2 (brown) is expressed basolaterally in goblet cells. Na+ is also reabsorbed by the ENaC (not shown). HCO3– is taken up by basolateral NBCe1 (pink), but apical mechanisms are not clear. (B) Interaction of goblet cells and enterocytes. Goblet cells (brown) are hypothesized to secrete HCO3– by transepacellular transport involving Best2 in the basolateral membrane and a Cl:HCO3– transporter in the apical membrane (see Discussion). Enterocytes (blue) secrete Cl– by transepacellular transport involving basolateral NKCC1 and apical CFTR.
We suggest that the colon has 2 kinds of CaCC currents, one likely mediated by Best2 and physiologically carried by HCO₃⁻, and another that may be mediated by an Ano1-encoded classical CaCC. Ca²⁺-stimulated Cl⁻ transport is likely to involve both CaCCs and CFTR, but the ratio of these pathways may be regulated.

**Mechanisms of HCO₃⁻ secretion in colon.** The transport proteins involved in HCO₃⁻ secretion from distal colon have not been clearly delineated (1, 2). In other tissues, in which HCO₃⁻ secretion has been studied more thoroughly (29), HCO₃⁻ transport is mediated by coordinated basolateral Na⁺HCO₃⁻ transporters and apical Cl⁻:H₂CO₃ exchangers. In the colon, the role of Na⁺HCO₃⁻ transporters is controversial (47, 48), and at least 3 different apical HCO₃⁻ efflux mechanisms have been identified (49). It remains unclear exactly which cells secrete HCO₃⁻ and how much of the HCO₃⁻ secretion occurs at the brush border and in the crypt. SLC26A3 is one of the Cl⁻:HCO₃⁻ exchangers in the apical membrane that mediates HCO₃⁻ secretion (50, 51). It has been proposed that CFTR-mediated HCO₃⁻ secretion is explained by direct effects of CFTR on SLC26A3 (52), but in the colon, SLC26A3 is located on the brush border membrane, whereas CFTR is expressed in the crypts (ref. 50 and H.C. Hartzell, unpublished observations). Thus, it seems unlikely that CFTR-mediated HCO₃⁻ secretion occurs by direct regulation of SLC26A3 in the colon. Another possibility is that CFTR stimulates HCO₃⁻ secretion indirectly by elevating extracellular Cl⁻ and thereby driving Cl⁻:HCO₃⁻ exchange in adjacent goblet cells.

**Best2 is a colonic anion channel.** A considerable body of evidence has established that bestrophins are a type of Ca²⁺-activated Cl⁻ channel (14). However, we showed here that Best2 did not function as a classical Ca²⁺-activated Cl⁻ channel, as was initially anticipated. Best1 is expressed on the brush border membrane of colonocytes in proximal colon and has been suggested to participate in Cl⁻ secretion (7, 8), but it remains to be seen whether Best2 and Best1 differ in their localization and function.

Our findings here complement a growing realization that bestrophins are unlikely to be the Xenopus oocyte variety of Ca²⁺-activated Cl⁻ channels (CaCCs), such as the ones present in salivary gland and pancreatic acinar cells (14, 17, 53). Differences between bestrophins and CaCCs include differences in their Ca²⁺ sensitivity, voltage dependence and kinetics, methods of regulation, and bicarbonate permeability (54). The first definitive evidence that bestrophins were not classical CaCCs was the finding that knockout of Best1 in mice did not eliminate CaCC currents in retinal pigment epithelial cells in which Best1 is expressed (17, 55). These studies have led to the suggestion that bestrophins, in addition to functioning as Cl⁻ channels, may also have other regulatory functions, including regulation of voltage-gated Ca²⁺ channels (14).

**Speculation regarding disorders of ion transport in inflammatory bowel disease.** Ion transport is disrupted in many inflammatory diseases in airway, kidney, and gastrointestinal tract (2, 56–58). It is tantalizing that the Best2 gene is located within a susceptibility locus for inflammatory bowel disease (i.e., IBD6; refs. 59–61). Our observations that Best2⁻/⁻ mice exhibited enhanced inflammation, slow recovery from DSS-induced colitis, and altered mucin biogenesis raise the possibility that Best2 may play a role in inflammatory bowel diseases.

**Methods**

**Mice.** Generation of Best2⁻/⁻ mice has previously been described (18). In initial experiments, control mice were C57BL/6 mice, but later experiments used control mice bred with the same genetic background as that of Best2⁻/⁻ mice; results were the same. All animals were bred and maintained in accordance with NIH and institutional guidelines, and all procedures were approved by the Emory University Institutional Animal Care and Use Committee. Age-matched mice of either gender were randomly assigned to the experimental procedures. Animals were sacrificed by isoflurane inhalation anesthesia or CO₂ asphyxiation until breathing ceased, followed by decapitation.

**Human tissue.** Use of human tissue was approved by the Emory University Institutional Review Board as exempt from IRB review because the study does not meet Federal Regulations 45 CFR Section 46.102(f)(2) criteria for “research involving human subjects.” Tissue samples were deidentified and impossible for us to trace back to the patient’s identity.

**Immunofluorescence.** Colonic tissue was fixed in 4% paraformaldehyde buffered with 0.1M phosphate buffer (pH 7.4), frozen, and embedded in OCT compound. Sections were treated with 0.025% saponin in PBS at room temperature for 30 minutes, then rinsed in PBS and blocked with 3% BSA (or 1% cold water fish gelatin) and normal goat serum for 1 hour at room temperature. Sections were incubated with primary antibodies (diluted 1:500–1:1,000) overnight at 4°C in a humidified chamber. Antibodies included anti-Best2 (provided by A. Menini, Italian Institute of Technology, Trieste, Italy; or our own antibody raised against amino acids 386–485), anti-MUC2, anti-CFTR (3G11, provided by Cystic Fibrosis Foundation; or provided by N. Nameen, University of Pittsburgh), and anti-Ano1 (raised against amino acids 878–960). Ano1 cDNA was provided by U. Oh (Seoul National University, Seoul, South Korea). After washing 5 times with PBS, we incubated the sections with Alexa dye–conjugated (Invitrogen) or Daylite dye–conjugated (Jackson Immunochemicals) goat anti-rabbit secondary antibody for 2–4 hours at room temperature. The sections were washed again in PBS, mounted in Prolong Gold (Invitrogen), and examined using a Zeiss LSM-510 microscope. The anti-Best2 antibody stained HEK cells transfected with Best2, but not untransfected cells or cells transfected with mouse Best1 or Best3. The antibody did not stain tissue from Best2⁻/⁻ mice.

**Western blot analysis.** Colonos were flushed with PBS, after which the mucosa was scraped off a distal piece (0.5 cm) with a razor blade and homogenized in lysis buffer containing 1% Triton X-100, 1 mM EDTA, 50 mM Tris-HCl (pH 7.4), and protease inhibitor cocktail III (Calbiochem) plus 10 μM phenylmethylsulfonyl fluoride. Protein content was determined by BCA assay (Pierce), the samples were diluted in SDS Laemmli buffer, and aliquots were run on 7%–12% reducing SDS-PAGE, transferred to nitrocellulose, and detected with anti-Best2 or Ano1 antibody.

**Using chamber experiments.** We used 2 cm of the mouse distal colon. Muscle layers were removed by gentle dissection, and a piece of mucosa was mounted in an Ussing chamber (Physiologic Instruments Inc.) with an aperture of 0.3 cm² (or, in some experiments, 0.07 cm²). The standard Krebs-Henseleit solution contained 118 mM NaCl, 4.7 mM KCl, 23 mM NaHCO₃, 1.2 mM K₂HPO₄, 1.2 mM CaCl₂, 1.2 mM MgCl₂, 0.3 mM EDTA, and 10 mM glucose. The reservoirs were vigorously gassed with 5% CO₂ and 95% O₂ and maintained at 37°C by water jackets. Iₑ was measured using an automatic voltage clamping device (DVC-1000; Physiologic Instruments Inc.) that compensates for resistance of the solution between the potential measuring electrodes. TEP was recorded through 3M KCl-agar bridges connected to a pair of calomel half-cells. The transepithelial current was applied across the tissue via a pair of Ag/AgCl electrodes that were kept in contact with the mucosal and serosal bathing solution using 3M KCl-agar bridges. All experiments were done under short-circuit conditions. The Iₑ was negative when positive current flows from serosa to mucosa. The tissue was placed in the apparatus and equilibrated for 30–40 minutes to stabilize Iₑ before starting the experiment. The baseline value of electrical parameters was determined as the mean over the 2 minutes immediately prior to drug administration. A positive Iₑ corresponds to the net electrogenic secretion of anions or the net electrogenic absorption of cations. Carbamylcholine (CCh, 1 mM) and forskolin (10 μM) were used to activate the ion transport.
To reduce the contribution of cation currents to the $I_{n}$, the K+ channel blocker Ba2+ (5 mM) and the epithelial Na+ channel blocker amiloride (100 μM) were added at the start of most experiments.

Isolation of colonic epithelial cells. A 4-cm piece of mouse distal colon was everted and rinsed with ice-cold Ca2+-free Ringer’s-type solution composed of 127 mM NaCl, 5 mM KCl, 5 mM Na-pyruvate, 5 mM glucose, 10 mM HEPES, 27 mM EDTA, and 1 mM MgCl2. This preparation was then incubated in normal Krebs solution for 10 minutes at 37°C. Isolated crypts were obtained by shaking and then treated with collagenase-P (20 μU/ml; Roche Applied Science) for 10 minutes. Isolated colonocytes from crypts were collected and replated in DMEM (Invitrogen) supplemented with 10% fetal bovine serum and penicillin-streptomycin (200 μg/ml and 200 U/ml, respectively). Studies were performed within 24 hours of the acute isolation.

X-gal staining of tissue. Distal colon (2 cm) was removed, opened lengthwise by a longitudinal incision, and pinned in a Sylgard-coated 35-mm dish. The tissue was fixed in 4% paraformaldehyde in 0.1M phosphate buffer (pH 7.4) for 1 hour; rinsed 3 times with rinse buffer containing 0.1M sodium phosphate buffer, 2 mM MgCl2, 0.01% sodium deoxycholate, and 0.02% NP-40; and incubated overnight in 20 mM potassium ferrocyanide, 20 mM potassium ferricyanide, and 1 mg/ml solution of X-gal in PBS. Tissue was examined with a microscope and photographed with a color CCD camera.

Immunoelectron microscopy. Immunoelectron microscopy was performed as described previously, using a preembedding immunogold method (62). Colons were fixed by immersion with 4% paraformaldehyde and 0.1% glutaraldehyde in 0.1M phosphate buffer, after which 60-μm sections were cut on a Vibratome (VT1000S; Leica) and blocked in Tris-buffered saline (TBS) containing 1% NGS. Sections were washed in PBS, and the sections were post-fixed with uranyl acetate followed by Reynolds’s lead citrate. Ultrastructural analyses were performed on 200-mesh nickel grids. Staining was performed on drops of 1% aqueous uranyl acetate, dehydrated in graded series of ethanol, and flat-embedded on glass slides in Durcupan (Fluka) resin. Regions of interest were cut at 70–90 nm on an ultramicrotome (Reichert Ultracut E; Leica) and collected on 200-mesh nickel grids. Staining was performed on drops of 1% aqueous uranyl acetate followed by Reynolds’s lead citrate. Ultrastructural analyses were performed in a Jeol-1010 electron microscope. To test specificity, the primary antibody was either omitted or replaced with 5% (v/v) normal rabbit serum. Under these conditions, no selective labeling was observed.

Patch clamp recording. Acutely isolated single colonocytes were recorded using conventional patch-clamp techniques. Fire-polished borosilicate glass patch pipettes were 3–5 MΩ. Experiments were conducted at room temperature (20°C–24°C). Because the liquid junction potentials were small (less than 2 mV), no correction was made. The standard pipette solution contained 146 mM CsCl, 2 mM MgCl2, 5 mM (Ca2+)-EGTA, 10 mM HEPES, and 10 mM sucrose (pH 7.3), adjusted with N-methyl-D-glucamine. The calculated Ca2+ concentrations were approximately 1 μM in high-Ca2+ solution. The standard extracellular solution contained 140 mM NaCl, 5 mM KCl, 2 mM CaCl2, 1 mM MgCl2, 15 mM glucose, and 10 mM HEPES (pH 7.4) with NaOH. This combination of intracellular and extracellular solutions set reversal potential ($E_{rev}$) for Cl- currents to 0, whereas cation currents carried by Na+ or Cs+ had very positive or negative $E_{rev}$, respectively. In HCO3−-containing solution, 140 mM NaHCO3 replaced NaCl on an equimolar basis, and the solution was equilibrated with 30% CO2 and 70% O2. Osmolarity was adjusted with sucrose to 303 mOsm for all solutions. Relative permeability was determined from the Goldman-Hodgkin-Katz equation, as previously described (21). Relative conductance was calculated from the slope of the current-voltage relationship ±25 mV of the reversal potential.

Induction of DSS colitis. Sex- and age-matched littermates, approximately 3 months old, received 3% DSS (MP Biomedicals) in drinking water for 6 days followed with normal water for recovery. Weight was recorded daily.

MPO activity. Neutrophil infiltration was quantified by measuring MPO activity. Colon samples were homogenized in 1:20 (w/v) of 50 mM phosphate buffer (pH 6.0) containing 0.5% hexadecyltrimethyl ammonium bromide (Sigma-Aldrich), on ice using a Polytron homogenizer. The homogenate was sonicated for 10 seconds, freeze-thawed 3 times, and centrifuged at 21,000 g for 15 minutes. The supernate (14 ml) was added to 1 ml mg/ml o-dianisidine hydrochloride (Sigma-Aldrich) and 0.0005% hydrogen peroxide (Sigma-Aldrich), and the change in absorbance at 460 nm over time was measured. One unit of MPO activity was defined as the amount that degraded 1 μmol peroxidase per minute at 25°C. Results are expressed as MPO units per microgram protein, determined by BioRad protein assay.

Statistics. Data are expressed as mean ± SEM. Statistical difference between means was evaluated by 2-tailed Student’s t test. The body weight difference between WT and Best2−/− mice was evaluated by factorial ANOVA of age (14 time points) and phenotype. Statistical significance was assumed for P values less than 0.05.

Acknowledgments

The authors thank Didier Merlin for the use of his Ussing Chamber setup and helpful discussion, Asma Nusrat for advice on histological and pathological analysis, Qinghuan Xiao for comments on the manuscript, Anna Menini for Best2 antibody, the Cystic Fibrosis Foundation for CFTR (3G11) antibody, Nadia Nameen for CFTR antibody, and Uthaek Oh for Ano1 cDNA. The authors’ work is supported by NIH grants GM60448 (to H.C. Hartzell), EY014852 (to H.C. Hartzell), and EY13160 (to A. Marmorstein); by an American Health Assistance Foundation grant (to A. Marmorstein); by an unrestricted grant to the Department of Ophthalmology and Vision Science at the University of Arizona (to A. Marmorstein); and by Emory University Core Grant for Vision Research P30 EY006360 (to H.C. Hartzell).

Received for publication September 10, 2009, and accepted in revised form February 17, 2010.

Address correspondence to: H. Criss Hartzell, Department of Cell Biology, Emory University School of Medicine, 615 Michael St., 535 Whitehead Building, Atlanta, Georgia 30322, USA. Phone: 404.242.5719; Fax: 404.727.6256; E-mail: Criss.Hartzell@emory.edu.


Hartnell HC. Ca\(^{2+}\)ing channels get the last laugh. Annu Rev Physiol. 2010;72:95–121.


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