Tight junction CLDN2 gene is a direct target of the vitamin D receptor

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The breakdown of the intestinal barrier is a common manifestation of many diseases. Recent evidence suggests that vitamin D and its receptor VDR may regulate intestinal barrier function. Claudin-2 is a tight junction protein that mediates paracellular water transport in intestinal epithelia, rendering them “leaky”. Using whole body VDR−/− mice, intestinal epithelial VDR conditional knockout (VDRΔIEC) mice, and cultured human intestinal epithelial cells, we demonstrate here that the CLDN2 gene is a direct target of the transcription factor VDR. The Caudal-Related Homeobox (Cdx) protein family is a group of the transcription factor proteins which bind to DNA to regulate the expression of genes. Our data showed that VDR-enhances Claudin-2 promoter activity in a Cdx1 binding site-dependent manner. We further identify a functional vitamin D response element (VDRE) 5′-AGATAACAAAGGTCA-3′ in the Cdx1 site of the Claudin-2 promoter. It is a VDRE required for the regulation of Claudin-2 by vitamin D. Absence of VDR decreased Claudin-2 expression by abolishing VDR/promoter binding. In vivo, VDR deletion in intestinal epithelial cells led to significant decreased Claudin-2 in VDR−/− and VDRΔIEC mice. The current study reveals an important and novel mechanism for VDR by regulation of epithelial barriers.

Tight junction (TJ) structural components determine epithelial polarization and intestinal barrier functions1–4. Claudins, with approximately 24 members, are integral membrane proteins and components of tight junctions5. The complex expression pattern of claudins creates diversity in the barrier/channel property of TJs, which varies depending on the type of epithelium5. Claudin-2 and -10 tend to make tight monolayers leakier5,7–9. Claudin-2, a “leak” protein uniquely restricted to the proliferative zone of the crypt base10,11, forms a paracellular water channel that mediates paracellular water transport in epithelia and renders it more “leaky”11–15. Defective epithelial barrier function has been implicated in IBD16 and similarly, elevation of Claudin-2 is associated with active IBD12,17. In addition, Claudin-2 likely participates in cellular functions other than its known effects on TJ function. For example, we have demonstrated that Salmonella targets Claudin-2 to facilitate bacterial invasion18, and epithelial cells with Claudin-2 knockdown have significantly less internalized Salmonella than control cells with normal Claudin-2 expression. Claudin-2 has also been identified as a target of Wnt/β-catenin signaling19, which is essential for intestinal development, and a recent study reported that Claudin-2 and -12 contributed to vitamin D-dependent calcium homeostasis20. Because of this latter report, and because Vitamin D and its receptor (VDR) are implicated in the pathogenesis of various intestinal illness, including IBD21–31, we sought to determine how Claudin-2 is regulated by VDR signaling.

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In the current study, we hypothesize that Claudin-2 is a direct target of the VDR. Using whole body VDRΔ mice, intestinal epithelial VDR conditional knockout (VDRΔIEC) mice, and cultured human intestinal epithelial cells; we perform a series of molecular and biochemical experiments in vivo and in vitro to investigate VDR regulation of Claudin-2 expression in enterocytes.

**Results**

**Intestinal VDR deficiency in epithelial cells leads to reduction of Claudin-2 at the mRNA and protein levels.** In whole VDRΔ mice, we detected significantly decreased mRNA levels of Claudin-2 in intestine (Fig. 1A), whereas other Claudins, such as Claudin-1, -4, -7, -10, and -15, were not altered by the absence of VDR (data not shown). By immuno-blots, we further found that VDRΔ mice had the highest protein level of Claudin-2 in intestine. VDRΔIEC mice had intermediate levels of Claudin-2, whereas the VDRΔIEC had the lowest levels of Claudin-2 protein (Fig. 1B). Therefore, VDR expression correlates with protein levels of Claudin-2 in colonic epithelial cells in vivo (Fig. 1C).

We further tested the specificity of intestinal VDR on expression levels of Claudin-2, in VDRΔIEC mice. We found that the mRNA levels of Claudin-2 were significantly lower in VDRΔIEC mice, compared to the VDR-lox mice (Fig. 1D). Claudin-2 protein was also significantly decreased in VDRΔIEC, where no VDR protein was detected in VDRΔIEC colon by Western blot (Fig. 1E). As expected, Claudin-3 and Claudin-7 were unchanged. These data indicate that intestinal VDR specifically regulates the expression levels of Claudin-2.

Colonic Claudin-2 expression is uniquely restricted to the proliferative zone10,11,18. In Fig. 1F, Claudin-2 staining (Green) was observed at the crypt base in VDRΔIEC mice. The density of Claudin-2 fluorescence staining was weak in the VDRΔ and VDRΔIEC intestinal epithelial cells. Claudin-7 was very stable in these mice and showed unchanged distribution and density (Fig. 1F).

**Vitamin D₃ treatment upregulates mRNA levels of Claudin-2.** For molecular mechanism studies in vitro, we used the human colonic epithelial SKCO15 cell line, which is widely used in studying TJs32,33. Vitamin D₃ is known to increase VDR expression and activate VDR signaling. Claudin-2 mRNA was significantly elevated in SKCO15 cells treated with 1, 25 vitamin D₃ (20 nM) for 24 hours, whereas Claudin-7 mRNA was not altered by vitamin D₃ treatment (Fig. 2A). Moreover, protein levels of Claudin-2 were increased by vitamin D₃ treatment in a dose-dependent manner (Fig. 2B). In contrast, the expression of Claudin-3 and 7 was unchanged in cells receiving vitamin D₃ treatment. These data suggest that the Claudin-2 gene could be a direct transcriptional target of the VDR.

VDR is generally present in the cytosol or bound to DNA in an inactive state and requires activation by binding ligand34. Upon binding to vitamin D, VDR translocates to the nucleus and binds to vitamin D response elements (VDREs) in target genes and induces gene expression. A previous study showed that ongoing protein synthesis is not required for this process to occur35. We treated human SKCO15 cells with vitamin D₃ (20 nM) in the presence or absence of cyclohexamide (CHX) to block protein synthesis. We chose to treat cells with vitamin D₃ at 20 nM because our dose-response data in Fig. 2B indicated that 20 nM is a suitable concentration to induce Claudin-2 expression. CHX is an inhibitor of eukaryotic protein biosynthesis and is commonly used to determine protein half-life. Therefore, in the cells treated with CHX only, the expression of Claudin-2 was significantly decreased (Fig. 2C SKCO15). CHX+vitamin D₃ treatment was able to stabilize the expression of Claudin-2 (Fig. 2C SKCO15+Vit.D3). Whereas vitamin D₃ induced Claudin-2 gene expression occurred in the absence of ongoing protein synthesis (presence of CHX), Vitamin D treatment did not induce Claudin-3 gene expression (Fig. 2C SKCO15+Vit.D3). These data further support the hypothesis that the Claudin-2 gene is a direct target of the VDR and not activated by secondary events, such as the synthesis of other transcription factors that are induced by VDR.

To study the effect of VDR overexpression on Claudin-2, we transiently transfected the human SKCO15 cells with a pCDNA-hVDR plasmid expressing human VDR. We found Claudin-2 expression increased after SKCO15 cells were transfected with pCMV-hVDR plasmids, whereas no change of Claudin-3 with VDR overexpression (Fig. 2D).

**VDR binds the Claudin-2 promoter in vitro and in vivo.** VDR is a nuclear receptor that acts as a transcription factor to regulate expression of its target genes21,36. We reasoned that VDR may bind to DNA promoters of Claudin-2, thus changing mRNA expression of the Claudin-2 genes. VDR’s effect on promoters of Claudin-2 was analyzed by CHIP assay. We designed primers to the nonrepetitive region near the transcriptional start site that specifically amplifies the Claudin-2 promoter. For negative controls, chromatin was immunoprecipitated with IgG or villin. The samples were amplified by conventional PCR. We found that VDR bound to the Claudin-2 promoter in vitro (SKCO15 cells, Fig. 3A) and in vivo (mouse colon, Fig. 3B). The expression of the other Claudin members, such as Claudin-1, was also tested. We found that VDR did not bind to the Claudin-1 promoter, either in vitro or in vivo (Figs. 3A,B). VDR is known to interact with nuclear receptor RXR in regulating gene expression. However there was no significant change in mRNA level of RXR in the VDRΔIEC intestine (Fig. 3C).

**Lacking VDR decreases Claudin-2 by abolishing VDR/promoter binding.** We reasoned that less Claudin-2 was generated in cells lacking VDR if VDR binds to the promoter of Claudin-2. We further
Figure 1. VDR status in intestinal epithelial cells leads to the change of Claudin-2 at both mRNA and protein levels in vivo. (A) Claudin-2 mRNA level and (B) Claudin-2 protein level in the intestinal epithelial cells of VDR$^{+/+}$, VDR$^{+/−}$, or VDR$^{−/−}$ mice. (C) Claudin-2/VDR protein relative in the intestinal epithelial cells of VDR$^{+/+}$, VDR$^{+/−}$, or VDR$^{−/−}$ mice. (D) Claudin-2 mRNA level and (E) Claudin-2 protein level in the intestinal epithelial cells of VDR KO (VDR$^{ΔIEC}$) mice. Data are expressed as mean ± SD. *P < 0.05. n = 3 mice/group. (F) Location and quantification of Claudin-2 protein in colons of mice in vivo. Images for each protein shown represent three separate experiments. n = 3 mice/group.
Figure 2. High levels VDR lead to increased Claudin-2 in human colonic epithelial SKCO15 cells in vitro. (A) Claudin-2 mRNA level increased post vitamin D₃ treatment. SKCO15 cells were treated with vitamin D₃ (20 nM) for 24 hours. Data are expressed as mean ± SD. *P < 0.05. n = 3 separate experiments. (B) Claudin-2 protein level and vitamin D₃ dose-dependent curve. SKCO15 cells were treated with indicated vitamin D₃ concentrations for 24 hours. Data are expressed as mean ± SD. *P < 0.05. n = 3 separate experiments. (C) Protein synthesis of Claudin-2 is high in vitamin D₃-treated SKCO15 cells. Data are expressed as mean ± SD. *P < 0.05. n = 3 separate experiments. (D) Claudin-2 expression after SKCO15 cell transfection with human VDR in a pCMV-hVDR plasmid. Data are expressed as mean ± SD. *P < 0.05. n = 3 separate experiments.
tested the effects of VDR in regulating mRNA and protein levels of Claudin-2 in human colonic epithelial SKCO15 cells. We found that Claudin-2 mRNA and protein expression were reduced when VDR was knocked down by siRNA (Fig. 4A,B). To examined the effect of one allele of VDR gene on the expression of Claudin-2, we chose VDR\(^{+/ -}\) and VDR\(^{-/-}\) mouse embryonic fibroblast (MEF) cells.\(^{37}\) We found that one allele of the VDR gene in the VDR\(^{+/ -}\) MEF cells was able to increase the expression of Claudin-2 protein (Fig. 4C). In contrast, Claudin-3 and -7 remained unchanged. At the transcriptional level, increased VDR mRNA was associated with elevated Claudin-2, but not Claudin-7 in VDR\(^{+/ -}\) MEFs. Claudin-2 mRNA was significantly decreased in VDR\(^{-/-}\) MEF cells (Fig. 4D). This result suggests that VDR deletion affects Claudin-2 mRNA. Additionally, if we knocked down Claudin-2 by siRNA, there was no reduction of VDR at either the protein or mRNA level (Fig. 4E,F). These data suggest that Claudin-2 is downstream of VDR signaling.

**VDR-enhances Claudin-2 promoter activity in a Cdx1 binding site-dependent manner.** Cdx is a member of the caudal-related homebox gene family.\(^{38,39}\) Suzuki et al. reports that IL-6-induced Claudin-2 promoter activity requires Cdx binding sites.\(^{40}\) To assess whether vitamin D3 could enhance Claudin-2 promoter activity through Cdx binding sites, we used an in vitro reporter Luciferase assay. A schematic drawing of transcriptional binding sites in the wild-type (WT) Claudin-2 promoter and its
mutants is shown in Fig. 5A. Plasmids with WT or deletions of NFκB, STAT, or Cdx1 in the Claudin-2 promoter binding site (of ΔNFκB, ΔSTAT, or ΔCdx) were transfected into cells, respectively, and then treated with vitamin D3. Vitamin D3 enhanced WT-Claudin-2 promoter activity in both HCT116 and CaCO2 cells (Fig. 5B,C). Deletions of NFκB and STAT binding sites did not affect the Claudin-2 promoter activity. In contrast, deletions of Cdx1 binding sites clearly suppressed the promoter activity (Fig. 5B,C). These results demonstrate that vitamin D3-induced Claudin-2 expression requires Cdx1 binding sites in the Claudin-2 promoter sequence.

Identification of a functional VDRE sequence in the Claudin-2 promoter. As Claudin-2 promoter activity is strongly elevated by exposure to 1, 25 vitamin D3, we predicted the existence of a VDRE in the Claudin-2 promoter. Our results from the reporter assay suggested that Cdx binding sites are involved in vitamin D3-mediated increases in Claudin-2 expression. Therefore, studies were conducted...
**Figure 5.** Identification of a functional VDRE sequence in the Claudin-2 promoter. (A) A schematic representation of transcriptional binding sites in the WT Claudin-2 promoter and deletion mutants. Plasmids include wild-type (WT), binding site deletions of NFκB (ΔNFκB), STAT (ΔSTAT), or Cdx1 (ΔCdx) in the Claudin-2 promoter. (B) WT or mutant Claudin-2 reporter plasmids were transfected in HCT116 and (C) CaCO2 cells. Luciferase activity was measured in the cell monolayers incubated in the absence or presence of vitamin D₃ (20 nM) for 24 hours. Dual luciferase assays were performed and firefly luciferase activity was normalized to renilla luciferase activity. Data are expressed as mean ± SD. *P < 0.05. n = 3 separate experiments. (D) A schematic representation of VDRE deletion construct plasmids. Putative VDR-binding sites (containing AGATAACAAAGGTCA sequence) are designated as VDRE. Deletions of all VDRE binding sites (ΔVDRE), deletion of VDRE binding sites and adjacent bases (ΔD2), and non-VDRE deletion controls (ΔD3/ΔD4). (E) WT Claudin-2 reporter gene plasmids and the deletion mutant plasmids were transfected to HCT116 and (F) CaCO2 cells. Luciferase activity was measured in the cell monolayers incubated in the absence or presence of vitamin D₃ (20 nM) for 24 hours. Data are expressed as mean ± SD. *P < 0.05. n = 3 separate experiments.
to investigate the Cdx binding site region. VDRE sequence is AGATAACAAAGGTCA. A search of the
Cdx region revealed a DR3-type, which binds preferentially to directly arrangements of two hexameric
binding sites with three spacing nucleotides. PCR was used to construct deletions of all VDRE binding
sites (ΔVDRE), deletion of VDRE binding sites and adjacent bases (ΔD2) and non-VDRE deletion
controls (ΔD3/ΔD4). These fragments were separately cloned into the pGL3-basic firefly luciferase
reporter plasmid. A schematic drawing of the VDRE deletion and control mutants is shown in Fig. 5D.
The Claudin-2 promoter VDRE deletion constructs were transfected into cells and were subsequently
treated with vitamin D3 (20 nM). Deletions of VDRE (ΔVDRE and ΔD2) clearly lower the promoter
activity of vitamin D3 in HCT116 (Fig. 5E) and CaCO2 cells (Fig. 5F). In contrast, non-VDRE deletion
controls (ΔD3 and ΔD4) did not affect the Claudin-2 promoter activity induced by vitamin D3 (Fig.
5E&5F). Our results demonstrate that deletion of the VDRE sequence 5΄-AGATAACAAAGGTCA-3΄
in the Claudin-2 promoter region causes loss of its responsiveness to vitamin D3, and thus confirm that
Claudin-2 is a direct target of vitamin D receptor signaling in intestinal epithelial cells.

Discussion
The experimental focus of our current study was to investigate the molecular mechanisms whereby VDR
may act as a transcriptional factor to regulate the expression of Claudin-2. First, we provide molecular
biological evidence that the Claudin-2 gene is a direct target of the transcription factor VDR. A transcrip-
tional reporter study demonstrated Claudin-2 up-regulation by over-expressed VDR. CHIP-PCR
data demonstrated specific binding of VDR to the Claudin-2 promoter. VDR enhanced Claudin-2
promoter activity in a Cdx1 binding site-dependent manner. Next, we identified a functional VDRE
sequence within the Claudin-2 promoter. Knockout of VDR led to lower Claudin-2 at both mRNA and
protein levels. Increased VDR by vitamin D3 pretreatment was associated with elevated Claudin-2
mRNA and protein levels. This study highlights an important and novel mechanism for VDR regulation
of Claudin-2 critical to intestinal homeostasis.

Claudin-2 is a unique member of the Claudin family of transmembrane proteins as its expression is
restricted to leaky epithelium in vivo and correlates with epithelial leakiness in vitro. VDR is a nuclear
receptor that mediates most functions of vitamin D. Our data showed that activation of the CLDN2
gene occurred via a consensus VDRE in the promoter that is bound by VDR. VDR is expressed in a
wide range of tissues. Therefore, potentially, Claudin-2 can be induced in various tissues. We know that
multiple factors contribute to the upregulation of Claudin-2 at the transcriptional level. TNF-α, IL-1β and
IL-6 enhance claudin-2 promoter activity in a Cdx binding site-dependent manner. VDR has multiple critical functions in regulating innate and adaptive immunity, intestinal homeostasis, host response to invasive pathogens and commensal bacteria, and
tight junction structure. TJ structure plays a critical role in intestinal barrier and inflammation. Claudin-2 is enhanced in the inflamed gut of patients with IBD. The pathobiological importance of the VDR regulation of Claudin-2 could be complex. Hence, further insight into the mechanisms respon-
sible for VDR and barrier dysfunction in mucosal inflammation is needed, especially in in vivo systems
disease models.

In summary, for the first time, we identify CLDN2 gene is a direct target of VDR. Our findings reveal
a novel activity of VDR in regulation of TJs in primary cell structure and intestinal homeostasis. This
study fills an existing gap by characterizing the precise molecular mechanism of VDR in regulating
Claudin-2 and highlights the complex role of VDR in intestinal homeostasis. It also brings up the
possibility for restoration VDR-dependent functions and prevention of the intestinal barrier breakdown
in patients with intestinal disorders.

Materials and Methods
Animals. VDR+/+, VDR+/− and VDR−/− mice on a C57BL6 background were obtained by breeding
heterozygous VDR+/− mice. VDR flox mice were originally reported by Dr. Geert Carmeliet. VDRΔ IEC
mice were obtained by crossing the VDR flox mice with villin-cre mice (Jackson Laboratory, 004586,
Bar Harbor, Maine, USA), as we previously reported. Experiments were performed on 2–3 months old
mice. All animal work was approved by the Rush University Committee on Animal Resources. Euthanasia
method was sodium pentobarbital (100 mg per kg body weight) I.P. followed by cervical dislocation.

Ethics statement. The methods in animal models were carried out accordance with the approved
guidelines by the Rush University Committee on Animal Resources.

Mouse colonic epithelial cells. Mouse colonic epithelial cells were collected by scraping the tissue
from the colon of the mouse, including the proximal and distal regions. The cells were sonicated in
lysis buffer (1% Triton X-100, 150 mM NaCl, 10 mM Tris, pH 7.4, 1 mM EDTA, 1 mM EGTA, pH 8.0,
0.2 mM sodium ortho-vanadate, and protease inhibitor cocktail). The protein concentration was mea-
sured using the BioRad Reagent (BioRad, Hercules, CA, USA).

Cell culture. Human epithelial CaCO2 and SKCO15 cells were maintained on transwell inserts (0.33
or 4.67 cm², 0.4 mm pore. Costar, Cambridge, MA, USA) in DMEM supplemented with 10% fetal bovine
serum, penicillin-streptomycin (Penicillin, 100 I.U./ml/Streptomycin, 100 μg/ml), and L-glutamine
(4.5 g/L). Human colonic epithelial HCT116 cells, VDR+/− and VDR+/− MEF cells were cultured in DMEM medium supplemented with 10% (vol/vol) fetal bovine serum, as previously described.

**Immunofluorescence.** Colonic tissues were freshly isolated and embedded in paraffin wax after fixation with 10% neutral buffered formalin. Immunofluorescence was performed on paraffin-embedded sections (4 μm), after preparation of the slides as described previously followed by incubation for 1 hour in blocking solution (2% bovine serum albumin, 1% goat serum in HBSS) to reduce nonspecific background. The tissue samples were incubated overnight with primary antibodies at 4°C. The following antibodies were used: anti-Claudin-2, anti-Claudin-7 (Invitrogen, Grand Island, NY, USA). Samples were then incubated with secondary antibodies (goat anti-mouse Alexa Fluor 488 or goat anti-rabbit Alexa Fluor 488, Molecular Probes, CA; 1:200) for 1 hour at room temperature. Tissues were mounted with SlowFade Antifade Kit (Life technologies, s2828, Grand Island, NY, USA), followed by a coverslip, and the edges were sealed to prevent drying. Specimens were examined with a Zeiss laser scanning microscope (LSM) 710 (Carl Zeiss Inc., Oberkochen, Germany).

**Analysis of claudins distribution.** Fluorescence images were analyzed using image analysis software (LSM 710 META, version 4.2; Carl Zeiss Inc., Oberkochen, Germany). Each analysis was performed in triplicate from each tissue section on a total of 10 images per mouse sample (n = 5).

**Transient transfections.** Transient transfections were performed with Lipofectamine 2000 (Invitrogen, San Diego, CA, USA) in accordance with the manufacturer's instructions. Cells were seeded on 60 mm dishes overnight before transfection with DNA and were mixed with liposome reagent at a ratio of 1:1 before addition to cells. After a 24-hour transfection period, the proteins were extracted for western-blot analysis.

**Chromatin immunoprecipitation (ChIP) assays.** The ChIP assays were performed essentially as described by the manufacturer (Upstate Inc., Chalottesville, VA, USA). Briefly, SKC015 cells or scraped VDR+/−/VDR− colonic epithelial cells were treated with 1% formaldehyde for 10 min at 37°C. Cells were washed twice in ice-cold phosphate buffered saline containing protease inhibitor cocktail tablets (Roche, Nutley, NJ, USA). Cells were scraped into conical tubes, pelleted and lysed in SDS Lysis Buffer. The lysate was sonicated to shear DNA into fragments of 200–1000 bp (4 cycles of 10 s sonication, 10 s pausing, Branson Sonifier 250, Danbury, CT, USA). The chromatin samples were pre-cleared with salmon sperm DNA-bovine serum albumin-sepharose beads, then incubated overnight at 4°C with VDR antibody (Santa Cruz Biotechnology Inc., Dallas, Texas, USA). Immune complexes were precipitated with salmon sperm DNA-bovine serum albumin-sepharose beads. DNA was prepared by treatment with proteinase K, extraction with phenol and chloroform, and ethanol precipitation, and was subjected to PCR (Primers see supplement table 1).

**Western-blot analysis.** Mouse epithelial cells were lysed in lysis buffer (1% Triton X-100, 150 mM NaCl, 10 mM Tris, pH 7.4, 1 mM EDTA, 1 mM EGTA, pH 8.0, 0.2 mM sodium ortho-ovanadate, and protease inhibitor cocktail) and the protein concentration was measured. SKC015 and MEF cells were rinsed three times in ice-cold HBSS, lysed in protein loading buffer (50 mM Tris, pH 6.8, 100 mM dithiothreitol, 2% SDS, 0.1% bromophenol blue, and 10% glycerol) and sonicated (Branson Sonifier, 250). Equal amounts of protein were separated by SDS-polyacrylamide gel electrophoresis, transferred to nitrocellulose (162–0112, Bio-rad, Hercules, CA, USA), and immunoblotted with primary antibodies. The following antibodies were used: anti-Claudin-2, anti-Claudin-3, anti-Claudin-7 (Invitrogen, Grand Island, NY, USA), anti-VDR, anti-Villin (Santa Cruz Biotechnology Inc., Dallas, Texas, USA), or anti-β-actin (Sigma-Aldrich, St. Louis, MO, USA) and were visualized by ECL (Thermo Scientific, Rockford, IL, USA). Membranes that were probed with more than one antibody were stripped before reprobing.

**Transcriptional activation.** After a 24-hour transfection period, the cells were lysed and luciferase activity was determined using the Dual Luciferase Reporter Assay System (Promega, Madison, WI, USA) with a TD-20/20 luminometer (Turner Designs, Sunnyvale, CA, USA). Firefly luciferase activity was normalized to Renilla luminescence activity and the activity expressed as relative units.

**Identification of functional VDRE.** PCR was used to construct deletion of entire VDRE binding sites (ΔVDRE), deletion of VDRE binding site with adjacent bases (ΔD2), and control (ΔD3/ΔD4). These fragments were separately subcloned into the firefly luciferase reporter plasmid pGL3-basic (Primers see supplement table 2). Deletions of different domains of the Claudin-2 promoter cloned into the in pGL3 vector, driving luciferase expression, were transfected into HCTC116/CaCO2 cells. Luciferase activity in cell lysates was assayed by the Dual Luciferase Reporter Assay System (Promega, Madison, WI, USA).

**Real-time quantitative PCR analysis.** Total RNA was extracted from mouse epithelial cells or cultured cells using TRIzol reagent (Invitrogen, Grand Island, NY, USA). The RNA integrity was verified by electrophoresis. RNA reverse transcription was performed using the iScript cDNA synthesis kit (Bio-Rad, Hercules, CA, USA) according to the manufacturer's protocol. The RT cDNA reaction products were
subjected to quantitative real-time PCR using CFX96 Real-time system (Bio-Rad, Hercules, CA, USA) and SYBR green supermix (Bio-Rad, Hercules, CA, USA) according to the manufacturer's protocol. All expression levels were normalized to β-actin levels of the same sample. Percent expression was calculated as the ratio of the normalized value of each sample to that of the corresponding untreated control cells. All real-time PCR reactions were performed in triplicate. Optimal primer sequences were designed using Primer-BLAST or were obtained from Primer Bank primer pairs listed in Supplement Table 3.

Statistical Analysis. All of the data are expressed as means ± SD. All of the statistical tests were two-sided and P values of less than 0.05 were considered to be statistically significant. Differences between two samples were analyzed using Student's t-test. The statistical analyses were performed using SAS version 9.2 (SAS Institute, Inc., Cary, NC).

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