Phosphorylation regulates connexin43/ZO-1 binding and release, an important step in gap junction turnover

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ABSTRACT To investigate whether connexin phosphorylation regulates the known role of zonula occludens-1 protein (ZO-1) in gap junction (GJ) function, we generated and analyzed a series of phosphomimetic and phosphorylation-dead mutants by mutating known conserved regulatory serine (S) residues 255, 279/282, 365, 368, and 373 located in the C-terminal domain of connexin43 (Cx43) into glutamic acid (E) or alanine (A) residues. All connexin mutants were translated into stable, full-length proteins and assembled into GJs when expressed in HeLa or Madin-Darby canine kidney epithelial cells. However, mutants with S residues exchanged at positions 365, 368, and 373 exhibited a significantly altered ZO-1 interaction profile, while mutants with S residues exchanged at 255 and 279/282 did not. Unlike wild-type Cx43, in which ZO-1 binding is restricted to the periphery of GJ plaques, S365A, S365E, S368A, S368E, and S373A mutants bound ZO-1 throughout the GJ plaques, while the S373E mutant did not bind ZO-1 at all. Inability to disengage from ZO-1 correlated with increased GJ plaque size and increased connexin protein half-life, while maintaining GJ channels in an open, functional state. Quantitative clathrin-binding analyses revealed no significant alterations in clathrin-binding efficiency, suggesting that the inability to disengage from ZO-1 prevented maturation of functional into nonfunctional/endocytic channels, rather than ZO-1 interfering with GJ endocytosis directly. Collectively, our results indicate that ZO-1 binding regulates channel accrual, while disengagement from ZO-1 is critical for GJ channel closure and transition of GJ channels for endocytosis. Intriguingly, these transitional ZO-1 binding/release and channel-aging steps are mediated by a series of hierarchical phosphorylation/dephosphorylation events at S373, S365, and S368, well-known Cx43 Akt, protein kinase A, and protein kinase C phosphorylation sites located in the vicinity of the ZO-1 binding site.

INTRODUCTION

Direct cell-to-cell communication plays a pivotal role in many aspects of multicellular life, as well as in many disease states. Gap junction (GJ) channels consist of two hemichannels (also termed connexons) that dock head on in the extracellular space, bridging the plasma membranes and thus connecting the cytoplasm of apposed cells. These channels are assembled from four-pass trans-membrane...
proteins termed connexins (Cxs). Clusters of GJ channels in densely packed arrays, termed GJ plaques, typically consist of hundreds to thousands of individual channels (Bruzzone et al., 1996; Kumar and Gilula, 1996; Falk et al., 2009). Cx43-based GJs are of particular interest because of their ubiquitous expression in most tissues and the central role they play in the heart and vasculature (Lin et al., 2007; Remo et al., 2012; Lambiase and Tinker, 2015; Meens et al., 2015).

A number of independent studies using an array of imaging techniques convincingly demonstrate that newly synthesized GJ channels typically appear to accrue along the outer edge of GJ plaques while older, nonfunctional, and permanently closed GJ channels are removed from plaque centers (Gaietta et al., 2002; Lauf et al., 2002; Falk et al., 2009; Gilleron et al., 2011; Rhett et al., 2011; Nickel et al., 2013). This continuous and dynamic turnover of channels in GJ plaques correlates with the known and unusually short 1-5 h half-life of Cxs and GJs observed in vivo and in situ (with the notable exception of lens fiber cells, which do not maintain functional protein-degradation machinery in order to retain eye lens transparency; Fallon and Goodenough, 1981; Beardslee et al., 1998; Gaietta et al., 2002; Falk et al., 2009, 2014). Dynamic GJ channel renewal also correlates with more recent studies suggesting that only a small portion of channels in a GJ plaque appear to be functional at any given time (Bukauskas et al., 2000; Curti et al., 2012; Flores et al., 2012). Interestingly, continuous GJ channel endocytosis and renewal is important not only for regulating natural levels of GJIC (Kretz et al., 2003; Qiu et al., 2003; Niend et al., 2006; Wang et al., 2007; Nakano et al., 2008; Remo et al., 2011; Johnstone et al., 2012; reviewed in Laird, 2014; Kelly et al., 2015; Solan and Lampe, 2015; Falk et al., 2016), but also in circumstances where cells physiologically or pathologically separate from each other, such as development, wound healing, mitosis, apoptosis, leukocyte extravasation, ischemia, hemorrhage, edema, and cancer metastasis (Ginzberg and Gilula, 1979; Lo and Gilula, 1979; Leach and Oliphant, 1984; Baker et al., 2008; Gilleron et al., 2008; Boassa et al., 2010; Hesketh et al., 2010; Fong et al., 2014; Nimlaamoo et al., 2015). However, how GJ biosynthesis, maintenance, and specifically aging and turnover are regulated on a molecular level is not well understood and has been the focus of this investigation.

Cx43 is a well studied phosphoprotein whose function is regulated by a large number of serine (S) and tyrosine kinases and phosphatases that act on Cx43 at multiple sites during all stages of its life cycle. These stages include trafficking from the ER to the Golgi and plasma membrane and altering Cx oligomerization behavior, as well as GJ assembly, gating, internalization, degradation, and regulation of the cell cycle (Thevenin et al., 2013; Solan and Lampe, 2015; Falk et al., 2016). Several of these phosphorylation events have been linked to GJ internalization and turnover (Thevenin et al., 2013; Solan and Lampe, 2014, 2015; Falk et al., 2016). However, a comprehensive understanding of molecular events that regulate channel accrual, GJ plaque assembly, and turnover, or of events that transition functional/open channels into nonfunctional/closed channels that then may become endocytosed, has not been established.

Phosphorylation/dephosphorylation events on specific Cx43 C-terminal residues are also likely to control Cx43’s many known interactions with various binding partners, such as chaperones, scaffolding proteins, kinases, phosphatases, ubiquitin ligases, deubiquitinases, and endocytic and other regulatory proteins (Giepmans, 2006; Herve et al., 2012). Precise spatiotemporal control of their binding is crucial, as many of the binding proteins that regulate GJ function likely are not all bound at one time, but are recruited and interact with Cxs only at certain stages of their life cycles (Solan and Lampe, 2007; Thevenin et al., 2013).

One important binding partner of Cx43 in particular, the scaffolding protein zona occludens-1 (ZO-1), has been characterized in greater detail. ZO-1 is known to interact with GJs (Giepmans and Moolenaar, 1998; Toyofuku et al., 1998) and to bind to the periphery of GJ plaques, where newly synthesized channels accrue (Hunter et al., 2005; Baker et al., 2008). ZO-1 is a member of the membrane-associated guanylate kinase family (MAGUK) of proteins. Cx43 has been shown to interact with the PDZ-2 domain of ZO-1 through its extreme C-terminus, with the last four residues of Cx43 inserting into a binding pocket (Chen et al., 2008; Thevenin et al., 2013). Multiple independent investigations indicate that newly synthesized Cxs traffic to the plasma membrane, dock in the periphery of GJ plaques (the peri-nexus), and accrue to the outer edge of GJ plaques, while simultaneously older channels are internalized from plaque centers (Gaietta et al., 2002; Lauf et al., 2002; Falk et al., 2009; Cone et al., 2014). Rhett et al. showed that ZO-1 regulates the Cx43 Cx-to-GJ transition, suggesting that ZO-1-binding to Cxs slows down their accrual rate, and therefore balances GJ plaque size (Rhett et al., 2011). Studies by Chen and colleagues suggest that S373 phosphorylation on a peptide representing the most C-terminal portion of Cx43 protein decreases affinity for ZO-1 on eightfold (Chen et al., 2008). This finding was further corroborated by studies of Dunn and Lampe, who demonstrated that under conditions of induced Akt-activation (wounding, hypoxia), phosphorylation of S373 led to detachment of ZO-1 from Cx43 (Dunn and Lampe, 2014), suggesting a possible role of S373 phosphorylation/dephosphorylation in regulating Cx43/ZO-1 interaction. Additional phosphorylation events located in the vicinity of the ZO-1 binding site (e.g., S365 and S368, known respectively as protein kinase A [PKA] and protein kinase C [PKC] targets; Solan and Lampe, 2007); farther upstream in the Cx43 C-terminal domain (e.g., S255, S262, not conserved in humans; S279/S282, all known MAPK target sites); and located in the vicinity of the Cx43 internalization signals (S2, Y215,AVF218, S3, Y189[KLV219]) (Thomas et al., 2003; Fong et al., 2013) are also implied to contribute to GJ turnover (Johnson et al., 2013; Fong et al., 2014; Nimlaamoo et al., 2015; Spagnol et al., 2016). However, a sufficient understanding of how dynamic Cx43/ZO-1 binding and release is regulated on a molecular basis and how ZO-1 binding and disengagement relates to other Cx43 modifications that aid to transition functional/open into nonfunctional/closed GJ channels that lead to endocytosis is currently lacking.

To address this issue, we constructed and analyzed a series of phosphomimetic and phosphorylation-dead mutants in which multiple known S residues of the Cx43 C-terminal domain were mutated into glutamic acid (E) or alanine (A) residues. We expressed the mutants as native proteins (i.e., not fused to green fluorescent protein [GFP] or any other tag) in Cx-deficient HeLa and Madin–Darby canine kidney (MDCK) epithelial cells, and investigated whether 1) the mutants were translated into stable, full-length Cx43 polypeptides, 2) the mutants were able to form GJ plaques, 3) plaques still interacted with ZO-1, 4) Cx43/ZO-1 interaction was altered, 5) amounts of ZO-1 bound to the Cx43 mutants were modified, 6) GJ plaque size and number was different, 7) Cx43 mutant protein half-life was affected, 8) the endocytic rate of the mutated Cxs was altered, and 9) channel function (Lucifer yellow [LY] dye-transfer ability) was increased or decreased. Together with previous reports, our results presented here indicate that a series of hierarchical phosphorylation/dephosphorylation events at S373, S365, and S368 (known Akt, PKA, and PKC sites, respectively) regulate—through ZO-1 binding and its release—Cx docking, channel accrual, and transitioning of peripherally located functional (open) into centrally located nonfunctional (closed) GJ channels that can then be endocytosed.
RESULTS

Generation and expression of Cx43 constructs mutated at specific C-terminal serine residues

To determine whether attachment or detachment of ZO-1 to/from the Cx43 C-terminus may be regulated by phosphorylation of the Cx43 C-terminal domain, we generated a series of mutants in which several known regulatory S residues in the Cx43 C-terminus were exchanged (Figure 1A). Both phosphorylation-mimetic (serines, S, replaced with glutamic acid, E) and phosphorylation-dead (S replaced with alanine, A) mutants were generated. Sites known to be phosphorylated by Akt (S373), PKC (S368), PKA (S365), and MAPKs (S255, S279/S282) were targeted in this study (Figure 1B) (Solan and Lampe, 2007, 2014; Thevenin et al., 2013). All mutants were constructed untagged, as tagging Cx43 with a fluorescent probe, such as GFP on its C-terminus, is known to prevent Cx43 from interacting with ZO-1 (Hunter et al., 2005; Thevenin et al., 2013). All Cx43 constructs produced stable, correct–molecular mass protein products in addition to known Cx43-phosphorylation forms (P0, P1, P2) in Cx-deficient HeLa and MDCK cells (shown for MDCK cells and compared with endogenously Cx43-expressing pulmonary artery endothelial cell (PAEC) controls, Figure 1C) (Solan and Lampe, 2007, 2009; Thevenin et al., 2013). Under conditions applied, transfection efficiencies and resulting Cx43 protein levels were much higher in MDCK compared with HeLa cells (~90% and 25% transfection efficiencies, respectively) (compare Figure 2 and Supplementary Figure S1, as well as Figure 3, A and B, with Figure 3C). In addition, all mutants assembled efficiently into GJ plaques when expressed in HeLa or MDCK cells (Figure 3 and Supplementary Figure S1). In general, GJ plaques formed in HeLa cells were larger than plaques assembled in MDCK cells, consistent with a generally higher metabolic activity of these cancer cells. However, GJ plaque number per cell was increased in MDCK cells in a zipperlike manner more similar to the morphology of GJs formed by endogenously expressed Cx43 in situ, or in primary cells (compare Figure 2 and Supplementary Figure S1). We therefore performed morphological plaque analyses (size measurements and interaction with ZO-1, shown in Figure 2) primarily in HeLa cells, as larger plaques are more accessible to light microscopic analyses, while biochemical and functional analyses (shown in Figures 3–7) were performed primarily in MDCK cells. However, comparable results for all experiments were obtained in both cell lines.

Mutating Cx43 serines 365, 368, and 373 totally alters the distribution of ZO-1 on GJ plaques

To investigate Cx43/ZO-1 interactions, Cx43 S mutants were expressed in HeLa cells, Cx43 and ZO-1 proteins were coimmunostained, and GJ plaques were analyzed by high-resolution confocal fluorescence analyses (Figure 2). Cells expressing wild-type (WT) Cx43 exhibited the typical staining of ZO-1 along the outer periphery of GJ plaques, as reported previously by others and our own laboratory (Figure 2A, panels in row 1; Hunter et al., 2005; Baker et al., 2008). In contrast, both the A and E mutants of S365, S368, and S373 displayed a completely different distribution pattern of ZO-1 on Cx43 GJ plaques: In S365A/E, S368A/E and S373A mutants, ZO-1 was no longer restricted to the periphery, but was distributed over the entire plaque surface (Figure 2A, rows 2–6), while the S373E mutant no longer exhibited any detectable interaction with ZO-1 (Figure 2A, row 7). All described results were supported by linescan analyses (performed along the dotted white lines shown) for both green (Cx43), and red (ZO-1) fluorescence channels (Figure 2A, right panels). This result is in agreement with in vitro Cx43/ZO-1-PDZ2 peptide binding studies performed by Chen et al. (2008), as well as more recent work by Dunn and Lampe (2014). To clearly distinguish plaque peripheries from plaque centers and avoid potential image processing artifacts, only plaques viewed on their plane surfaces, were analyzed; plaques oriented edge-on were not considered. In addition, quantifying remaining levels of ZO-1 in the cytoplasm correlated well with amounts of ZO-1 bound to GJ plaques in each mutant. Mutants that showed ZO-1 binding throughout plaques (S365A/E; S368A/E; S373A) exhibited correspondingly lower cytoplasmic ZO-1 levels compared with WT and the S373E mutant (Figure 2B).

To test whether mutating other S residues elsewhere in the Cx43 C-terminal domain would have a similar effect on Cx43/ZO-1 interactions, we mutated three additional known regulatory S residues: S255, S279, and S282 (279/282 together). Both A and E mutants at these sites displayed normal distributions of ZO-1 along the rim of GJ plaques that were similar to WT (Figure 2C). This result suggests...
that only S residues in the vicinity of the ZO-1 binding site regulate ZO-1 binding and release, while S residues located farther upstream (known MAPK sites located juxtaposed to the endocytic motifs S2 and S3; Fong et al., 2013) have no effect. Similar analyses performed with high-resolution wide-field fluorescence microscopy (150× primary magnification, Supplemental Figure S2), or AiryScan confocal superresolution microscopy (up to 140 nm resolution in X–Y, shown for representative mutants S255A and S365A, Supplemental Figure S3) showed comparable ZO-1/GJ plaque distribution patterns consistent with our high-resolution confocal analyses.

Cx43/ZO-1 coimmunoprecipitation correlates with altered ZO-1/GJ plaque distribution

To further investigate altered mutant Cx43/ZO-1 colocalization, we first performed coimmunoprecipitation analyses of Cx43 and ZO-1 from whole-cell lysates. Coimmunoprecipitation of ZO-1 from HeLa cells expressing WT or Cx43 mutants showed that indeed, significantly more ZO-1 interacted and coprecipitated with S365A and E, S368A and E, and S373A mutants, while significantly less ZO-1 coprecipitated with the S373E mutant when analyzed by one-way analysis of variance (ANOVA) with Dunnett comparison test (Figure 3, A and B).

To further refine our analyses and investigate whether increased/decreased ZO-1 coprecipitation indeed was specific for Cx43 localized in GJ plaques, we performed Triton X-100 solubility assays in Cx43 WT and mutant transfected MDCK cells. Based on the known resistance of GJs to dissolving in 1% Triton X-100 (Musil and Goodenough, 1991), plasma membrane GJs can be separated from the intracellular Cx43 pool (cCxs, Cx43 polypeptides). First, we found that much more ZO-1 was present in the insoluble than in the soluble fractions (Figure 3C). In addition, the level of ZO-1 that coprecipitated with S365A and E, S368A and E, and S373A mutants was increased, while the S373E mutant bound less ZO-1 than WT Cx43 GJs, consistent with our previously described fluorescence colocalization analyses (Figure 3C, top panel). Also note that Cx protein levels of mutant Cxs were increased (Figure 3C, bottom panel),

**FIGURE 2:** Mutating S365, S368, and S373, but not S255 and S279/S282, in Cx43 causes a dramatically altered ZO-1 distribution on GJs. HeLa cells expressing WT and respective Cx43 S mutants were processed for immunofluorescence confocal analyses at 24 h posttransfection and high-resolution confocal analyses. In the merged images, labeled Cx43 is shown in red and ZO-1 in green. Representative images of en face GJ plaques are shown in A and C. Boxed regions selected in the left panel are shown magnified in the panels to the right. Cx43 and ZO-1 fluorescence signals recorded along the dotted lines traversing GJ plaques are shown in the line scan graphs on the right. (A) In WT Cx43, ZO-1 is distributed only along the periphery of GJ plaques, in accordance with previous observations (Hunter et al., 2005). In contrast, in S365A and E, S368A and E, and S373A mutants, ZO-1 is distributed throughout plaques, while the S373E mutant does not bind ZO-1 at all. (B) Quantitative analyses of ZO-1 bound to GJ plaques and distributed in the cytoplasm (as a percentage) averaged from at least three independent experiments. Average percentages of ZO-1 levels were calculated from at least 20 images. Error bars are SEM. One-way ANOVA, followed by Dunnett multiple comparison analyses (compared with the WT), is shown. (C) Mutating S255, S279, and S282 in Cx43 into A or E did not alter ZO-1 distribution, resulting in the rim staining typical for WT Cx43.
consistent with increased protein half-life and impaired mutant protein turnover rates (see below). Similar results were also obtained when fractionation assays were performed in HeLa cells, albeit with a somewhat higher level of background due to the significantly lower transfection efficiency and thus overall lower Cx43 protein expression level in these cells (Supplemental Figure S4).

In addition, as a control, we separated whole cell lysates derived from primary PAECs that express Cx43 endogenously at high levels into soluble and insoluble fractions (Figure 3D). Also, in these cells, much more ZO-1 was present in insoluble than in soluble fractions. In addition, in agreement with the concept that Cx43 in GJs is highly phosphorylated compared with soluble (intracellular, secretory) Cx43 (Solan and Lampe, 2009), insoluble GJ Cx43 polypeptides migrated more slowly, compared with the soluble Cx43 protein pool. Taken together, these biochemical data fortify our confocal data, further supporting our conclusion that ZO-1 remains bound to Cx43 when critical Cx43 S residues are mutated.

Cx43 mutants that cannot disengage from ZO-1 form significantly larger GJ plaques

Because ZO-1 has been found to be involved in regulating GJ plaque size (Hunter et al., 2003; Rhett et al., 2011), we next assessed the effect of our S mutagenesis on mutant Cx43 polypeptide and Cx43 protein half-life. HeLa cells expressing WT or mutant Cx43 and immunostained with Cx43 antibodies were imaged and the Cx43 fluorescence signal at cell–cell contacts was quantified using ImageJ software (Figure 4, A and B; see Supplemental Figure S6 for details). Levels of GJ Cx43 at cell–cell contacts in S368E and S373E mutants were comparable to that in WT Cx43, while all other mutants (S365A and E, S368A, and S373A) displayed significantly more GJ plaque area (indicated by significantly increased fluorescence levels) than WT Cx43 (Figure 4, A and B). These results concur with the observed ZO-1 distribution differences presented in Figure 2, A and B, signifying that the S368E and S373E mutants accurately replicate steps that occur in aging GJ plaque channels—namely, that Cx43/ZO-1 interaction is naturally abolished, leading to reduced plaque size in these mutants. This is in contrast to the Cx43 mutants that prevent normal ZO-1 detachment, leading to increased overall GJ plaque area in these mutants. When any of our mutants were expressed with a C-terminal GFP tag, observed plaque size-differences were no longer detectable (unpublished data), further supporting the findings of Hunter et al. (2003, 2005) and Rhett et al. (2011) that Cx43/ZO-1 interaction regulates channel accrual/GJ plaque size.

Cx43 mutants with altered ZO-1 interaction profiles have drastically increased protein half-lives

To ascertain whether Cx43 protein half-life was affected in our mutants, MDCK cells expressing WT or mutant Cx43 were treated with cycloheximide to inhibit protein biosynthesis and remaining levels of Cx43 polypeptides were analyzed by Western blot analyses using Cx43-specific antibodies (Figure 5). All mutants, except S373E (5.3 h), were found to have significantly longer protein half-lives (∼12–21 h).
et al., 2008; Fong et al., 2014). None of the mutants revealed significantly altered clathrin-binding levels from those in WT (Figure 6, A and B). These results argue against a reduced endocytosis rate, and instead argue for an increased channel accrual rate caused by a deregulation in Cx43/ZO-1 binding and release characteristics in these 5 mutants.

GJ channels assembled from Cx43 mutants that cannot disengage from ZO-1 stay functional (open)
To test whether our Cx43 S mutants also exhibit altered channel activities, we performed scrape-loading Lucifer yellow (LY) dye-transfer assays in Cx43 WT or mutant-expressing MDCK cells (Figure 7). The extremely high transfection rate of MDCK cells (app. 90%; see Material and Methods) allowed us to perform these assays in transiently transfected cells. Distance of LY spreading away from the scrape 10 min past injury and fixation was determined by quantitatively measuring LY-fluorescence signals. LY in Cx43 WT-expressing MDCK cells exhibited robust dye transfer rates (∼100 μm/10 min; ∼4 cell width) compared with those in untransfected MDCK cells that were tested in parallel as negative controls. These cells took up the dye but were not able to spread it to neighboring cells (∼25 μm/10 min; ∼1 cell width) (Figure 7, A and B, panels 1 and 2).

The S365A mutant showed no difference in dye transfer from WT Cx43 (∼100 μm/10 min; ∼4 cell width), while in contrast, the S365E mutant (simulating permanent S365 phosphorylation) exhibited a significantly increased level of dye transfer and the highest dye transfer rates (∼135 μm/10 min; ∼5½ cell width) (Figure 7, A and B, panels 3 and 4). This result conforms to the proposed role of S365 and S368 in regulating transition from functional (open) into nonfunctional (permanently closed) channels that is controlled by S365 dephosphorylation/S368 phosphorylation (“gatekeeper event”; Solan and Lampe, 2007).

Increased mutant GJ plaque size results from increased channel accrual, rather than decreased endocytosis rates
To determine whether mutants exhibiting longer protein half-lives and/or increased GJ plaque size/numbers have decreased endocytosis rates, we conducted coimmunoprecipitation analyses of clathrin heavy chain (CHC) and Cx43 in MDCK cells (Figure 6). Previously, Cx43-based GJs have been shown to internalize using the clathrin endocytic machinery (Piehl et al., 2007; Gumpert et al., 2008; Fong et al., 2013); hence levels of clathrin bound to Cx43 can be used as a reliable measure to gauge the rate of GJ internalization (Gumpert et al., 2013). Hence levels of clathrin bound to Cx43 can be used as a reliable measure to gauge the rate of GJ internalization (Gumpert et al., 2013). Hence levels of clathrin bound to Cx43 can be used as a reliable measure to gauge the rate of GJ internalization (Gumpert et al., 2013). Hence levels of clathrin bound to Cx43 can be used as a reliable measure to gauge the rate of GJ internalization (Gumpert et al., 2013).
extrapolated and therefore are approximate.

prolonged half-lives of S365A and E, S368A and E, and S373A mutants, values were followed by Dunnett multiple comparison test results is shown for all mutants, as compared with WT Cx43. (C) Summarized average protein half-lives for all mutants. Owing to extremely prolonged half-lives of S365A and E, S368A and E, and S373A mutants, values were extrapolated and therefore are approximate.

with this mutant not forming a larger GJ plaque area than for WT; Figure 4) and ZO-1 redistribution results (with this mutant being unable to interact with ZO-1; Figure 2, A and B), supporting the concept that S373 is phosphorylated early after biosynthesis in Cx43 Cxs concomitant with transport to the plasma membrane. Owing to its inability to interact with ZO-1, in this mutant the ability to assemble GJ plaques appears to be perturbed. Indeed, in MDCK cells expressing the S373E mutant, many cell–cell borders did not exhibit detectable GJs (Supplemental Figure S1). Taken together, our results indicate that GJ assembly, channel function, and maturation into nonfunctional channels correlate with ZO-1 binding and release, and that these critical regulatory steps are controlled by a series of phosphorylation/dephosphorylation events that occur in a sequential order on S373, S365, and S368 in the Cx43C-terminal domain.

DISCUSSION

We found that mutating single amino acid residues in the Cx43 C-terminal domain can dramatically alter the distribution of a well-known Cx43 binding partner, ZO-1, from its distinct localization restricted to the peripheral rim of WT GJ plaques (Hunter et al., 2003, 2005; Baker et al., 2008; Rhett et al., 2011) into a pattern that suggests binding of ZO-1 throughout the entirety of GJ plaques, or into a complete loss of binding. These critical amino acid residues represent prominent S residues (S373, S368, S365) that are well known to regulate GJ function via their phosphorylation states. Mimicking phosphorylation/dephosphorylation of these residues by mutating them into phospho-mimetic and phospho-dead residues (glutamic acid [E] or alanine [A], respectively) lead to these dramatic changes in ZO-1 binding, suggesting that specific phosphorylation/ dephosphorylation events occurring on critical Cx43 S residues located in the vicinity of the C-terminal Cx43 ZO-1 binding site regulate Cx43/ZO-1 interaction, and if phosphorylation on these residues is permanently prevented (as in the S to A mutants) or simulated (as in the S to E mutants), ZO-1 no longer can bind, or can no longer disengage from Cx43 and remains bound throughout the mutant GJ plaques. Notably, similarly mutating regulatory S residues located farther upstream in the Cx43 C-terminal domain (S279, S282, S255 phosphorylated by MAPK) did not at all affect ZO-1 localization, and the pronounced rim association typical for WT Cx43 remained. Inability of mutant Cx43 to disengage from ZO-1 correlated with increased Cx43 mutant protein half-life, increased GJ plaque size and number, and reduced L& dye transfer ability, suggesting that disengagement of Cx43 from ZO-1 is necessary for normal GJ function, including transitioning GJ plaque channels from functional (open) into nonfunctional (permanently closed), which then may become endocytosed. Interestingly, phosphorylation/dephosphorylation on S373 and S365 appears to activate channel function (opens channels), while phosphorylation on S368 (and subsequent phosphorylations on upstream MAPK sites) appears to inactivate it (closes channels) and to transition channels for endocytosis. Obtained results correlate well with the described short half-lives of Cxs and GJs (Berthoud et al., 2004), and with previous findings that indicate that i) Cx43 phosphorylation on S373 regulates channel accrual (Dunn and Lampe, 2004), and with previous findings that indicate that i) Cx43 phosphorylation on S373 regulates channel accrual (Dunn and Lampe, 2004), ii) dephosphorylation on S365 followed by phosphorylation on S368 regulates cell–cell coupling efficiency (Solan et al., 2007), and iii) GJ channels that are phosphorylated on S368 remain permanently closed and are endocytosed from central plaque areas (Cone et al., 2014). However, a link between down-regulation of GJIC (which also can be achieved via channel closure) and GJ internalization, and the sequence and type of molecular events that regulate these transitional steps have never been established previously.

With at least 15 different known phosphorylation sites being present in the Cx43 C-terminal domain and multiple protein kinases known to phosphorylate these sites (Solan and Lampe, 2005, 2007, 2009; Thévenin et al., 2013), it appears likely that many of the transitional steps that regulate GJ turnover, including Cx docking and channel accrual, opening and permanently closing GJ channels, and transitioning of functional (open) into nonfunctional (permanently closed) GJ channels, which then may be primed for clathrin-binding
and endocytosis, are regulated by distinct Cx phosphorylation events. As phosphomimetic and phospho-dead mutants bear the ability to arrest the Cx43 life-cycle at a relevant critical step, generating and analyzing these mutants allowed us to investigate the role each of these regulatory S residues may have in this process. Instead, activating/inactivating kinases/phosphatases, for example, via wounding or the generation of reactive oxygen species (ROS), may not have allowed us to do these analyses comparably well, as the latter approach is more likely to generate additional, unintentional, and potentially misleading phosphorylation events. In addition, it allowed us to investigate individual phosphorylation events under physiological, non-stress-induced conditions. Further evaluating Cx43S/E mutants together with previously published data finally permitted us to conclude i) that binding of ZO-1 to Cx43 and its subsequent release are regulated by Cx43 phosphorylation; ii) that disengagement from ZO-1 seems necessary for the transition of functional (open) channels into nonfunctional (closed) channels that then may be modified further to permit their endocytosis; and iii) that these transitional GJ channel activation and inactivation steps are regulated by a series of hierarchical phosphorylation/dephosphorylation events that occur on S373, S365, and S368, well-known Cx43 Akt, PKA, and PKC phosphorylation sites (schematically depicted in Figure 8).

Two S residues located in the Cx43 C-terminal domain, in particular, were found previously to play a critical role in regulating GJ channel function. S365 dephosphorylation (phosphorylated by PKA), termed “gatekeeper event” by the Lampe and Sorgen labs, was determined to be critical for successive phosphorylation of S368 (phosphorylated by PKC), an event that was linked to inhibition of GJIC, GJ channel closure, movement of channels to plaque centers, and the loss of GJ plaques from plasma membranes (Cone et al., 2014). S365 phosphorylation is constitutive in the heart, but is absent during hypoxia (Solan and Lampe, 2007). In addition, phosphorylation of S368 is increased in the heart undergoing hypoxia and Cx43 is lost from intercalated discs, suggesting that S368 phosphorylation might trigger GJ channel closure and potentially Cx43 internalization (Solan and Lampe, 2007; Cone et al., 2014), a hypothesis that concurs with our data.

Moreover, several labs previously gained evidence suggesting that phosphorylation/dephosphorylation of S373 in Cx43 (phosphorylated by Akt) may play an important role in regulating Cx43/ZO-1 interaction (Chen and Sudol, 1995; Park et al., 2006, 2007; Rhett et al., 2011; Dunn and Lampe, 2014). Our analyses indeed indicate that phosphorylation at S373 (mimicked in the S373E mutant) averted Cx43/ZO-1 interaction, while preventing S373 phosphorylation (mimicked in the S373A mutant) significantly increased Cx43/ZO-1 interaction (Figures 2 and 3). Some discrepancy, however, seems to remain as to whether S373 phosphorylation actually increases or decreases GJIC (Dunn and Lampe, 2014, and our study). Dunn and Lampe previously gained evidence that suggests that phosphorylation of S373 increases GJ size and cell–cell communication (Dunn and Lampe, 2014), while in our study S373 phosphorylation decreased communication (GJ plaque area remained unchanged in our study, Figures 4 and 7). These contradictory results may be explained by differing experimental conditions, as Dunn and Lampe’s experiments were performed under stress conditions (heart injury, hypoxia), which likely lead to strong activation of Akt, while our experiments were performed under normal, unstressed conditions where Akt kinase activity is expected to remain much lower. Importantly and independent of this difference, with respect to the S residue that regulates Cx43/ZO-1 interaction (S373), both our and the Dunn and Lampe studies concur.

Our comprehensive S mutagenesis further indicated that additional S residues are involved in the regulation of Cx43/ZO-1 interaction and GJ channel function. Indeed, dephosphorylation of S373 (which is thought to occur after Cx43 have arrived at the plasma membrane and docked to form GJ channels) may not be sufficient to render the newly accrued channels fully functional. Our LY dye transfer experiments indicate that the S373A mutant transfers dye comparably efficiently to WT and S365A mutant (Figure 7), while phosphorylation of S365 (mimicked in the S365E mutant, and reiterated in the S368A mutant, in which phosphorylation of S368 is prevented) resulted in significantly increased dye transfer abilities. Together with our evidence that the S366E mutant is unable to significantly transfer dye, and thus channels in this mutant apparently largely stay closed, our results are in good agreement with those of Solan and colleagues, who also found that phosphorylation of S365 keeps channels open, while phosphorylation of S368 closes channels (Solan et al., 2007).

So why after all may ZO-1’s release from GJ channels be required for their endocytosis? The work of Chen et al. (2008) indicates that the Cx43/ZO-1 interaction requires that two ZO-1 proteins undergo a structural dimerization known as “domain swap” as they bind two separate Cx43 polypeptides, a structural modification believed to make the PDZ-2 domain interaction of ZO-1 Cx43 specific. Thus, at least two ZO-1 proteins appear to bind to a Cx43 channel in a GJ plaque. As ZO-1 is a very large 220-kDa protein (almost the size of an entire Cx43-based Cx), its dimerization will generate a 440-kDa protein complex that in comparison dwarfs the size of the underlying bound connexon. Likely these huge ZO-1 protein complexes will efficiently shield the cytoplasmic surface of GJ plaques and will need to be removed before subsequent Cx43 modifiers (MAPKs, E3-Ub ligases,
clathrin/clathrin adaptors) can access, modify, and transition Cx43 GJ channels for endocytosis.

In summary, the molecular machinery that drives accrual of newly synthesized channels to the periphery of GJ plaques and the subsequent transitioning of functional channels into nonfunctional channels (permanently closed) that then are endocytosed from central plaque areas appears complex, and is driven by a number of consecutive Cx43 phosphorylation and dephosphorylation events (Figure 8). Taking into account work by others, our results suggest that 1) S365 and S373 phosphorylation (by PKA and Akt, respectively)—probably occurring early in the secretory pathway—promote forward trafficking, potentially regulated via Akt phosphorylation of S373 and interaction with 14–3–3 protein (Park et al., 2006, 2007). In addition, phosphorylation on these residues appears to prevent premature binding of ZO-1 to plasma membrane-delivered Cxs, potentially allowing them to function as hemichannels before they can dock and accrue to GJ plaques. 2) S373 dephosphorylation then permits ZO-1 to bind to Cxs, and Cxs to dock and accrue to GJ plaques. These newly accrued channels are likely to be open and functional. 3) Subsequent dephosphorylation of S365 followed by concomitant phosphorylation of S368 (by PKC) then induces release of ZO-1 from Cx43 and GJ channels to close (permanently). Continuing accrual of newly synthesized channels to plaque edges then drives aged/closed channels inward that then, triggered by additional Cx43 posttranslational modifications, namely MAPK-mediated phosphorylation and ubiquitination (Fong et al., 2014; Nimlamool et al., 2015; Kells-Andrews and Falk, 2017), permits clathrin recruitment and GJ channel endocytosis. Interestingly, Solan et al. (2007) found that a Cx43 phosphomimetic mutant of S365 (S365D) underwent a large conformational change in its

FIGURE 7: Cx43 S mutants that cannot disengage from ZO-1 stay open. (A) MDCK cells were transfected with Cx43 WT and mutant constructs. Dishes with very high transfection rates (>90%) were scraped 24 h later with a razor blade in the presence of GJ-permeable LY dye. Untransfected cells were analyzed in parallel as negative controls. After a 10-min incubation period, cells were fixed and the distance that dye spread away from the scrapes (marked with dotted lines) was measured. Representative images are shown. (B) Quantitative analyses of three independent experiments in which more than 30 images for each construct were analyzed. Asterisks mark significant differences from WT Cx43 (one-way ANOVA followed by Dunnett multiple comparison test). (C) Cx43 protein levels in cells used in the dye transfer assays were analyzed by Western blot and were found to be comparable to Cx43 protein levels found in endogenously Cx43-expressing primary PAE cells. (D) Dye transfer rates in primary PAE cells analyzed in parallel were comparable to those in transfected Cx43 WT. Note that dye transfer rates in Cx43 S365E, S368A, and S373A were significantly higher compared with Cx43 WT, suggesting that channels assembled from these mutant Cxs stayed open. WCL = whole cell lysate.
MATERIALS AND METHODS

cDNA constructs and mutagenesis

Untagged Cx43 mutants were generated by the Quick Change mutagenesis method (Stratagene) using WT rat Cx43 cDNA cloned into pEGFP-N1 vector (Clontech) as previously described (Falk, 2000). A Cx43 authentic TAA stop codon was reintroduced to allow expression of full-length untagged Cx43. Forward mutagenesis primers were as follows (mutated codons are underlined): Cx43-Stop, 5′-CCT GAT GAC CTG GAG ATT TAA GAT CCA CCG GTC GCC ACC-3′; S365A, 5′-ATC GTG GAC CAA CGA CCT TCC GCA AGA GCC AGC CGC GCC ACC-3′; S365E, 5′-ATC GTG GAC CAA CGA CCT TCC GAA AGA GCC AGC AGC CGC GCC AGC-3′; S368A, 5′-CAA CGA CCT TCC AGC AGA GCC GCA AGG CCT CGG CCT GAT GAC-3′; S368E, 5′-CAA CGA CCT TCC AGC AGA GCC GCA AGG CCT CGG CCT GAT GAC-3′; S255, 5′-ATC GTG GAC CAA CGA CCT TCC GCA AGA GCC AGC CGC GCC ACC-3′; S255A, 5′-ATC GTG GAC CAA CGA CCT TCC GAA AGA GCC AGC AGC CGC GCC AGC-3′; S279A/S282A, 5′-CAA CGA CCT TCC AGC AGA GCC GCA AGG CCT CGG CCT GAT GAC-3′; and S279E/S282E, 5′-CAA CGA CCT TCC AGC AGA GCC GCA AGG CCT CGG CCT GAT GAC-3′. It thus is possible that in newly accrued (functional) channels in which S368 has not yet become phosphorylated, access of MAP kinases, ubiquitin ligases, and clathrin/clathrin-adaptors to the Cx43-C-terminus is simply sterically blocked, while in aged channels in which S368 is phosphorylated (nonfunctional, permanently closed), access—triggered by the conformational change—is permitted (Falk et al., 2016). Certainly, implementing a sequence of different regulatory modifications (binding/release of a scaffolding protein, early and late phosphorylation/dephosphorylation events, ubiquitination) instead of transitioning GJ channels from functional to nonfunctional by a single event increases specificity and flexibility, including the ability to abort and revert the chain of events in case changing cellular conditions call for a modification in GJIC and/or physical cell–cell adhesion. As the first reports now link altered, nonphysiological GJ turnover rates to disease (Berthoud et al., 2013; Gemel et al., 2014; reviewed in Laird, 2014; Kelly et al., 2015), and Cx43 polypeptides with altered or deleted C-terminal domains exhibit impaired turnover characteristics (Maas et al., 2007; Vreeburg et al., 2007; Remo et al., 2011), understanding the molecular machinery that drives the dynamic turnover of gap junctions appears particularly significant.

C-terminal domain that also included a distant, upstream-located domain that we now know harbors the Cx43 endocytic motifs (Thomas et al., 2003; Fong et al., 2013). It thus is possible that in newly accrued (functional) channels in which S368 has not yet become phosphorylated, access of MAP kinases, ubiquitin ligases, and clathrin/clathrin-adaptors to the Cx43-C-terminus is simply sterically blocked, while in aged channels in which S368 is phosphorylated (nonfunctional, permanently closed), access—triggered by the conformational change—is permitted (Falk et al., 2016). Certainly, implementing a sequence of different regulatory modifications (binding/release of a scaffolding protein, early and late phosphorylation/dephosphorylation events, ubiquitination) instead of transitioning GJ channels from functional to nonfunctional by a single event increases specificity and flexibility, including the ability to abort and revert the chain of events in case changing cellular conditions call for a modification in GJIC and/or physical cell–cell adhesion. As the first reports now link altered, nonphysiological GJ turnover rates to disease (Berthoud et al., 2013; Gemel et al., 2014; reviewed in Laird, 2014; Kelly et al., 2015), and Cx43 polypeptides with altered or deleted C-terminal domains exhibit impaired turnover characteristics (Maas et al., 2007; Vreeburg et al., 2007; Remo et al., 2011), understanding the molecular machinery that drives the dynamic turnover of gap junctions appears particularly significant.

FIGURE 8: Predicted sequence of phosphorylation/dephosphorylation events indicated by our analyses and how they relate to ZO-1 binding and release, GJ channel accrual, opening and closing, and endocytosis. A scheme of a GJ plaque showing newly accrued functional channels (open) with bound ZO-1 located in the periphery (green), transitioning/aging channels (yellow), and aged, nonfunctional (closed) channels that no longer interact with ZO-1 and are primed for endocytosis in the plaque center (red) is shown. S373 is likely phosphorylated early on during the oligomerization of Cx43 into Cxs and forward trafficking towards the plasma membrane. Cx43 phosphorylated on S373 does not interact with ZO-1. Once S373 is dephosphorylated, ZO-1 can bind Cx43. S365 is phosphorylated concurrently with S373 dephosphorylation, or shortly before or after. GJ channels are open and functional. When S365 becomes dephosphorylated, S368 can be phosphorylated, causing channels to close. ZO-1 releases in response to these phosphorylation/dephosphorylation events. Earlier we reported that phosphorylation on S368, S255, S279, and S282 down-regulates GJIC (through channel closure) and causes Cx43 GJs to internalize (Fong et al., 2014; Nimlamool et al., 2015). This suggests that later MAPK phosphorylation events in the vicinity of the clathrin/clathrin-adaptor binding sites primes aged, nonfunctional (closed) GJ channels for endocytosis. Key works by others and us that correlate with our findings are referenced.
GAA CCT CCT GGG TAC AAG CTG G-3'. PCR mutagenesis reactions were performed with proofreading Pfu Ultra II polymerase (cat. no. 600670-51; Stratagene), per Quick Change mutagenesis protocol provided by Stratagene and DNA was digested with DpnI (cat. no. R0176-S; New England Biolabs) per Quick Change mutagenesis protocol provided by Stratagene and transformed into maximum- efficiency chemically competent DH5α Escherichia coli cells (cat. no. 18258-012; Invitrogen). All DNA constructs were sequenced to verify the presence of the intended mutations.

**Cell culture and transfections**

Cx-deficient HeLa cells (cat. no. CCL-2; American Type Culture Collection [ATCC]) were maintained at 37°C, 5% CO2 in high glucose DMEM. Media were supplemented for a final concentration of 10% with fetal bovine serum (FBS), 2 mM DMEM. Media were supplemented for a final concentration of 10% fetal bovine serum (FBS), 2 mM DMEM. Cells were grown on glass coverslips pretreated with poly-L-lysine. Cx-deficient MDCK cells (cat. no. CRL-2936; ATCC) (Dukes et al., 2011) and primary porcine PAECs, isolated as described previously (Baker et al., 2008), were maintained under the same conditions, except in low-glucose DMEM supplemented with the same additions as media used for HeLa cells. For transfection experiments, HeLa or MDCK cells were passaged 24 h prior to transfections and seeded to reach ~75% confluency. Cells were transfected with Lipofectamine2000 (cat. no. 11668027; Invitrogen) as recommended by the manufacturer. Transfection efficiencies of ~25% for HeLa and up to ~90% for MDCK cells were routinely achieved. PAECs endogenously expressing high levels of Cx43 served as a positive control.

**Immunofluorescence microscopy and quantitative image analyses**

Cells were grown on glass coverslips pretreated with poly-L-lysine (cat. no. P8920; Sigma-Aldrich), fixed with 3.7% formaldehyde at room temperature (RT) for 15 min, and permeabilized with 0.2% Triton X-100 in phosphate-buffered saline (PBS) at RT for 15 min. Cells were then blocked with 10% FBS in PBS at RT for 30 min and incubated with rabbit polyclonal anti-Cx43 antibodies (cat. no. 3512; Cell Signaling Technology) at 1:2000 dilution at 4°C overnight. Secondary antibodies (Alexa Fluor488-conjugated goat anti-rabbit, cat. no. A11008; Molecular Probes/Invitrogen) were used at 1:1000 dilution at RT for 1 h. Cells were rinsed in PBS and blocked with 10% FBS and then incubated with mouse monoclonal anti-ZO-1 antibodies (cat. no. 339100; Invitrogen) at 1:2000 dilution at 4°C overnight. Secondary antibodies (Alexa Fluor568-conjugated goat anti-mouse, cat. no. A11031; Molecular Probes) were used at 1:1000 dilution for 1 h at RT. Cell nuclei were stained with 1 μg/ml 4′,6-diamidino-2-phenylindole (DAPI). Cells were mounted using Fluoromount-G (cat. no. 0100-01; Southern Biotechnology) after coverslips were rinsed in distilled water. Slides were examined either with a Zeiss 880 confocal microscope, a Zeiss 880 confocal microscope equipped with AiryScan, or a Nikon Eclipse TE 2000E inverted fluorescence microscope, equipped with either 60×, 63×, or 100× NA 1.3 and NA 1.4 Plan-Apochromat oil-immersion objectives (Nikon Instruments; Carl Zeiss) and a Photronics CoolSnap HQ CCD camera (Roper Scientific). Images shown in Supplemental Figure S2 were acquired with a 100× objective with additional 1.5× auxiliary magnification to further boost CCD camera resolution. Wide-field images were acquired using MetaVue software version 6.1r5 (Molecular Devices). Quantitative image analyses comparing cytosolic and GJ bound ZO-1, line scan analyses, and GJ plaque area analyses were performed using ImageJ version 1.41o (National Institutes of Health; see legends of Supplemental Figures S5 and S6 for details). For line scan analyses, the “Line Scan Tool” in MetaVue software was used and data points were exported and analyzed in MS Excel. To calculate levels of cytosolic versus GJ-bound ZO-1, images were exported into ImageJ and areas of measurement were outlined and quantified. In each image, at least two areas of cytosolic ZO-1 in cells not expressing any Cx43 were compared with cytosolic levels of ZO-1 in cells that expressed Cx43 and GJ-bound ZO-1 calculated (Supplemental Figure S5). Data are averages of at least three independent experiments with at least 20 images per experiment and mutant. For GJ plaque size/number analyses, images were exported into ImageJ software and areas of cell–cell contacts containing visible Cx43 fluorescence signals were selected and quantified. Data are expressed as averages of performed experiments with at least 25–30 images per experiment and mutant.

**Coimmunoprecipitation assays**

HeLa or MDCK cells transfected with WT or mutant Cx43 cDNAs were trypsinized and grown for 24 h, rinsed in ice-cold PBS, and lysed in lysis buffer (1× RIPA buffer, cat. no. 9806; Cell-Signaling Technology) in the presence of protease inhibitors (cat. no. P8340; Sigma-Aldrich) and 1 mM NaN3 and 1 mM β-glycerophosphate (to block phosphatase activity) for 20 min on ice. A quantity of 10 μl of Protein-G Dynabeads (cat. no. 10030D; Invitrogen)/sample was washed in 500 μl of PBS/0.02% Tween20. For ZO-1/Cx43 coimmunoprecipitations, the beads were incubated with 1 μl of mouse monoclonal ZO-1 antibody for 1 h at RT. For clathrin heavy-chain/Cx43 coimmunoprecipitation, the beads were incubated with 1 μl of mouse monoclonal clathrin heavy-chain antibody (cat. no. 610499; BD Transduction Laboratories) for 1 h at RT. Bead/antibody mixtures were washed twice in 500 μl PBS/0.02% Tween20 and added to cell lysates, which were first centrifuged at 14,000 rpm to remove cell debris and organelles. Cleared cell lysates (typically 800 μl prepared from a 6-cm-diameter culture dish of confluent cells) were incubated with antibody/beads conjugates for 2 h at RT, washed twice in 500 μl of lysis buffer, and eluted with 30 μl of 2 × SDS–PAGE sample buffer. Beads were boiled for 5 min, and eluted proteins were analyzed by SDS–PAGE and Western blot analyses.

**SDS–PAGE and Western blot analyses**

Proteins were resolved on 10% SDS–PAGE gels, transferred to nitrocellulose membranes, blocked with 5% nonfat dry milk in TBS/0.1% Tween20 (TBS-T), washed three times in TBS-T, and incubated overnight in primary antibodies in 5% BSA/TBS (BSA cat. no. A7906; Sigma Aldrich) at 4°C. Antibody dilutions were as follows: rabbit Cx43 (Cell Signaling), 1:2000; mouse ZO-1 (Invitrogen), 1:1000; mouse clathrin heavy chain (BD Transduction), 1:2000; α-tubulin (cat. no. T2096, Sigma-Aldrich), 1:5000. Blots were washed three times in TBS-T and incubated in the appropriate secondary antibodies (goat anti-mouse-HRP and goat anti-rabbit HRP, cat. nos. G21040 and G21234, respectively; Invitrogen) for 30 min at 1:3000 dilution at RT. Relevant proteins were detected with enhanced chemiluminescent reagent. ImageJ (National Institutes of Health) was used to quantify protein band intensities.

**Triton X-100 solubility assays**

Triton X-100 solubility assays were performed as described by Musil and Goodenough (1991) with the following modifications: 70–75% confluent 3.5-cm dishes of MDCK cells were transfected with WT or mutant Cx43 cDNAs using Lipofectamine2000. At 20 h posttransfection, cells were lysed for 20 min in 400 μl of prechilled lysis buffer containing 50 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1 mM EDTA, 0.5% sodium deoxycholate, and 1% Triton X-100. The buffer was supplemented with protease and phosphatase inhibitors as described above. Cells were scrapped off the dishes, and lysates were
transferred into 1.5 ml Eppendorf tubes and centrifuged at 10,000 × g for 5 min at RT to preclear cell lysates. A 180-μl sample of each resulting supernatant was centrifuged at 100,000 × g in a Beckman Coulter Airfigue ultracentrifuge for 8 min. Supernatant fractions were immediately removed and transferred into new tubes, while remaining pellets were resuspended in 20 μl of 2x SDS-PAGE sample buffer. For Western blot analyses, 40 μl of the soluble fraction (100,000 × g supernatants) were mixed with 13.3 μl of 4x SDS-PAGE sample buffer. Total fractions (10,000 × g supernatants) were prepared in the same manner in controls. For each construct, 15 μl of the soluble fraction and 15 μl of the total fraction were resolved by SDS-PAGE. The entire 20 μl of the insoluble fraction (100,000 × g pellet) was analyzed by SDS-PAGE using anti-ZO-1 antibodies. Probed membranes were then stripped and reprobed with Cx43 antibodies. Identical procedures were applied when HeLa or PAECs were analyzed.

Cx43 protein half-life analyses
A set of 70–75% confluent 3.5-cm dishes (seven per construct and experiment) were transfected either with WT or with respective mutant Cx43 cDNA using Lipofectamine2000. At 20 h posttransfection, six dishes each were treated with 50 μg/ml cycloheximide (cat. no. C6255; Sigma-Aldrich) and kept in an incubator at 37°C, 5% CO2. One dish was designated as an untreated control; it was rinsed at RT in 1x PBS and cells were lysed with 500 μl of 1x SDS-PAGE sample buffer. At various time points (30 min and 1, 2, 3, 4, and 6 h), remaining dishes were treated as described for the control. All samples were immediately boiled for 5 min and stored at –80°C for subsequent SDS-PAGE and Western blot analyses (15 μl/sample) using Cx43 antibodies. Probed membranes were stripped and reprobed with anti α-tubulin specific antibodies to control for equal loading. Intensities of Cx43 and α-tubulin bands were analyzed using ImageJ software and Cx43 signals were normalized to the intensity of the corresponding α-tubulin signals.

Scrape-loading dye-transfer assays
MDCK cells were seeded on poly-l-lysine-coated glass coverslips placed into 3.5-cm dishes and grown to 50-75% confluence. In parallel, 3.5-cm dishes not containing coverslips were also seeded with cells. Cells with and without coverslips were transfected with WT or mutant Cx43 cDNAs, leaving one set of dishes as untransfected controls. At 20 h posttransfection, cells (then close to confluency) in 1x PBS were rinsed with 1x SDS PAGE sample buffer, boiled for 5 min, and used for Western blot analyses to detect expression levels of Cx43. Cells in the dishes containing coverslips were rinsed in 1x PBS at RT, and 1 ml of 0.05 weight % LY (cat. no. L682; Invitrogen) in 1x PBS was added to each dish. Monolayers were scraped with a sharp razor blade and incubated at 37°C under 5% CO2 for 10 min. Cells then were rinsed quickly in 1x PBS and fixed in 3.7% formaldehyde for 10 min at RT. Cover slips were rinsed three times in 1x PBS and dye transfer was imaged with a 20x objective on a Nikon Eclipse TE 2000E inverted fluorescence microscope. Scrape-loading dye-transfer assays were also performed with PAECs as a positive control, as described previously (Baker et al., 2008). A comparable sample of whole-cell lysate derived from PAECs was analyzed in controls to compare Cx43 expression levels in MDCK cells with the expression level in endogenously Cx43-expressing cells.

Statistical analyses
All error bars were calculated as ±SEM in GraphPad Prism (version 4.0 for Macintosh) (GraphPad, La Jolla, CA). To calculate statistical significance, ordinary one-way ANOVAs were performed, followed with Dunnett multiple comparison tests (as compared with the WT), with 95% confidence (p ≤ 0.05). Asterisks represent statistically significant differences, where *p < 0.05, **p < 0.005, and ***p < 0.0005.

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