Connexin43 Functions as a Novel Interacting Partner of Heat Shock Cognate Protein 70

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Regulation of connexin43 (Cx43) expression affects cell proliferation, differentiation and apoptosis in a gap junctional intercellular communication (GJIC)-independent manner. However, the underlying mechanisms of Cx43-mediated cell cycle suppression are still poorly understood. To elucidate the molecular mechanism of Cx43-mediated cell cycle suppression, we searched for Cx43 interacting proteins by using a proteomics approach. Here, we have identified a Cx43-interacting protein, heat shock cognate protein 70 (Hsc70). We confirmed that Hsc70 directly binds to the C-terminus of Cx43, whereas Hsc54, a splice variant of Hsc70, does not, that Cx43 competes with cyclin D1 for binding to Hsc70, and that the nuclear accumulation of cyclin D1 is reduced by overexpression of Cx43 in a GJIC-independent manner, which is restored by co-overexpression with Hsc70. As a result, the cell proliferation is regulated by Cx43. Our results suggest that Cx43-Hsc70 interaction probably plays a critical role during G1/S progression.

Connexins are integral membrane proteins that act as gap junction channel-forming proteins1,2. Gap junctions are specialized cell-cell junctions that directly link the plasma membrane between contacting cells. Low-molecular-weight (<1000 Da) metabolites and ions, including second messengers such as cyclic AMP, inositol triphosphate and Ca2+ can be selectively exchanged between adjacent cells through the gap junction. Gap junctional intercellular communication (GJIC) has fundamental roles, in regulating cell growth and differentiation, and in coordination of development, tissue function and cell homeostasis1,2.

Connexin43 (Cx43) is the most abundant gap junction protein in various tissues, and has long been viewed as a cell growth suppressor. Cx43 is ubiquitously expressed in human tissues and regulates cell growth and differentiation via multiple mechanisms. Cx43 plays various roles through not only its gap junction intercellular communication (GJIC)-dependent function, but also its GJIC-independent functions, such as hemichannels3–5 and protein-protein interaction6–12. It has been found that expression of Cx43 is inversely related to glioma cell proliferation and tumor grade13–15. The growth regulation of glioma cells is proposed to be more dependent on the behavior of connexins than the activity of GJIC. The ability of Cx43 to provide cell-cell communication via gap junctions was originally considered to be the only mechanism by which Cx43 regulates cell growth17–19. However, more recent studies have shown that Cx43 also controls cell growth independently of its ability to form gap junction channels between adjacent cells16–19. In HeLa cells transfected with mutated Cx43, which have point mutations in the second extracellular loop and do not display GJIC because of aberrant cytosolic localization of Cx43, the cell growth is still suppressed17. Moreover, the carboxy-tail of Cx43 localizes to the nucleus and inhibits cell growth in cardiomyocytes and HeLa cells19.

Therefore, it is possible that Cx43 may affect cell growth independently of gap junction formation. However, the exact mechanisms underlying GJIC-independent actions of Cx43-mediated cell cycle suppression and cell growth are still largely unknown. We hypothesized that Cx43 is involved in cell cycle suppression through a novel interacting protein, which regulates the cell cycle. We identified heat shock cognate protein 70 (Hsc70) as a novel interacting partner of Cx43 by co-immunoprecipitation. We confirmed that Cx43 directly interacts with Hsc70 via its C-terminus, and that this interaction plays a critical role in G1/S cell cycle progression through regulating nuclear accumulation of cyclin D1. Our findings are the first to suggest that the cell growth-suppressive properties of Cx43 probably are achieved by the interaction with Hsc70.
Results

Cx43 interacts with heat shock cognate protein 70. To identify a novel interacting protein of Cx43, we performed co-immunoprecipitation assays using anti-Cx43 antibodies to screen Cx43-interacting partners from HuH-7 cells, which abundantly express Cx43. As shown in Figure 1a, co-immunoprecipitation assays recovered several bands that were resolved by SDS-PAGE gels and visualized with Coomassie blue staining. MALDI/Q-TOF (matrix-assisted laser desorption ionization/quadrupole time of flight) mass spectrometry analysis of tryptic peptides identified a number of these proteins. One of them, heat shock cognate protein 70 (Hsc70), was reproducibly recovered in co-immunoprecipitation from the whole cell lysates of HuH-7 cells (Supplementary Fig. S1a). Another band obtained in the co-immunoprecipitation assay was identified by MALDI/Q-TOF analysis as β-tubulin (Fig. 1a), which has already been described as an interaction protein for Cx438.

To confirm the interaction of Cx43 with Hsc70, lysates of HuH-7 cells were subjected to GST pulldown assays using the C-terminal tail of Cx43 as an affinity matrix (GST-Cx43CT228-382; see Supplementary Fig. S1b). The results showed that Hsc70 interacts with GST-Cx43CT228-382 but not with GST alone (Fig. 1b and Supplementary Fig. S1c). We also co-transfected Hsc70 with Cx43 to HuH-7 cells; a possible in vivo interaction between Hsc70 and Cx43 was investigated by co-immunoprecipitation (Fig. 1c, d and Supplementary Fig. S1d, e). Next, to confirm a subcellular interaction between Cx43 and Hsc70, we analyzed the distribution of endogenous Cx43 and Hsc70 by immunofluorescence with the corresponding antibodies. In HuH-7 cells, partial co-localization of both proteins was observed in cytoplasm (Fig. 1e).

To further verify the interaction between Cx43 and Hsc70, HeLa cells were also transfected with the expression vector of Cx43. In transfected HeLa cells, partial co-localization in the both proteins was observed in the cytoplasm (see Supplementary Fig. S2).

These results demonstrated that Hsc70 associates with Cx43 in the cytoplasm, and suggest that Hsc70 functions as a novel interaction partner of Cx43.

Identification of the Hsc70-binding domain in Cx43. To determine which domain of Cx43 is required for binding to Hsc70, three types of deletion GST fused Cx43-CT mutants were prepared (Fig. 2a and

![Figure 1](https://www.nature.com/scientificreports) | Hsc70 Interacts with Cx43 and Co-localizes with Cx43 in the Cytoplasm. (a) In vivo evidence for Hsc70-Cx43 interaction. SDS-PAGE and Coomassie staining of proteins obtained in co-immunoprecipitation assay with anti-Cx43 antibodies using lysates prepared from HuH-7 cells. Arrows indicate the positions of Hsc70 and β-tubulin. Both proteins were identified by MALDI/Q-TOF mass spectrometry analysis. (b) Binding of Hsc70 to GST-Cx43CT. Proteins extracted from HuH-7 cells were mixed either with GST or with GST-Cx43CT residues 228-382 resin. After washing, the bound proteins were analyzed on 12% SDS-PAGE followed by western blotting (WB) with anti-Hsc70 antibody. Full-length blots are presented in Supplementary Figure S1b. (c and d) Co-immunoprecipitation of Hsc70 with Cx43. Lysates were prepared from HuH-7 cells co-transfected with wild-type Cx43 and T7-tagged Hsc70 expression plasmids. Lysates were immunoprecipitated (IP) either by anti-Cx43 antibodies and control IgG (c) or by anti-T7 antibody and the IgG (d). The immunocomplexes were analyzed by western blotting (WB) using anti-T7 antibody (c) or anti-Cx43 antibodies (d). Full-length blots are presented in Supplementary Figure S1c and S1d. (e) Co-localization of endogenous Hsc70 and Cx43. HuH-7 cells were fixed and stained with anti-Cx43 antibodies and anti-Hsc70 antibody. The nuclei were stained with TO-PRO3. Co-localization of both proteins was observed in the cytoplasm.
Supplementary Fig. S1b). GST pulldown assays were carried out using the whole cell lysates extracted from HuH-7 cells. As shown in Figure 2b, one deletion mutant, GST-Cx43CT\textsubscript{D}\text{JM}228-263, bound Hsc70 to the same extent as did GST-Cx43CT\textsubscript{228-382}, while another deletion mutant, GST-Cx43JM228-263 protein, failed to interact with Hsc70, as did GST protein alone. Thus, the 263-382 region of Cx43 is required for binding to Hsc70.

Identification of the Cx43-binding domain in Hsc70. To determine the Cx43-binding domain in Hsc70, a series of T7-tagged Hsc70 expression vectors encoding various portions of Hsc70 molecular\textsuperscript{29} were transfected into HuH-7 cells (Fig. 2c and d, left panel). GST pulldown assays were carried out using the cell lysates and GST-Cx43CT\textsubscript{228-382}. As expected, GST-Cx43CT\textsubscript{228-382} bound to wild-type Hsc70 but not to D\text{385–543} mutant, which lacked the protein-binding domain in Hsc70 (Fig. 2d, right panel). Moreover, D\text{1-385} and D\text{543–646} mutant Hsc70 did not affect binding to GST-Cx43CT\textsubscript{228–382}. Taken together, these results demonstrated that the 385–543 region of Hsc70 is essential for binding to Cx43.

Cx43 competes with cyclin D1 for interaction with Hsc70. In previous studies, the interaction of cyclin D1 and Hsc70 has been reported\textsuperscript{30,31}. To confirm whether Hsc70 interacts with cyclin D1 or Cx43 \textit{in vivo}, co-immunoprecipitation assays were performed. Hsc70 was co-immunoprecipitated with cyclin D1 or Cx43 (Fig. 3a); however, no interaction between cyclin D1 and Cx43 was detected (Fig. 3b). We then reasoned that Cx43 could compete with cyclin D1 for interaction with Hsc70. To test this hypothesis, we performed \textit{in vivo} competition assays. HuH-7 cells were transfected with equivalent plasmids of Hsc70, cyclin D1, and increasing amount of Cx43 expression plasmids for competitive binding to Hsc70. Cx43 interfered with the binding of cyclin D1 to Hsc70 in a dose-dependent manner (Fig. 3c). Taken together, these results suggest that there are interactions between cyclin D1 and Cx43 for competitive binding at Hsc70.

Interaction of Cx43 with Hsc70 plays an important role during G1 to S phase transition. Hsc70 is involved in the progression of cell cycle during G1/S transition\textsuperscript{29–32}. To assess whether Cx43 exerts the cell growth-suppressive functions through interacting with Hsc70, resulting in an effect on cell proliferation, bromodeoxyuridine (BrdU) incorporation assays were performed. HuH-7 cells were transiently transfected with vectors encoding either EGFP, Cx43-EGFP, or Hsc70-EGFP, and or both Cx43- and Hsc70-EGFP. Forty-eight hours after transfection, the cells were labeled with BrdU for 2 hr and subsequently fixed and stained with anti-BrdU antibody (Fig. 4a and Supplementary Fig. S3a). The fraction of BrdU-positive cells over total number of transfected cells was determined. Approximately 40% BrdU-positive cells were observed in both control and Hsc70-transfected cells. Overexpression of Cx43 significantly decreased the BrdU incorporation, from 40% to 13%. Interestingly, co-transfection of Cx43 with Hsc70 was sufficient to restore the BrdU incorporation, from 13% to 33% (Fig. 4b and Supplementary Fig. S3b). To further investigate whether or not Cx43-mediated cell cycle suppression is independent of GJIC, BrdU incorporation assay in

Figure 2 | Interaction Domains between Cx43 and Hsc70. (a) A schematic representation of GST-Cx43CT fusion proteins used as a ligand. Consensus protein domains and regions with putative signaling functions are indicated. Abbreviations in Cx43: TM1-4, transmembrane domains 1–4; JM, juxtamembrane domain that binds microtubes; Y265, Src phosphorylation site; PP, proline-rich sequence that binds the SH3 domain of v-Src; DLEI, carboxy-terminal sequence involved in ZO-1 binding. (b) Identification of Hsc70-binding domain in Cx43. Whole cell lysates of HuH-7 cells and various forms of the GST-Cx43CT fusion proteins were used for GST pulldown assays. Binding experiments and western blotting (WB) were carried out as described in Figure 1b. (c) A schematic representation of Hsc70 and its deletion mutants used for overexpression. (d) Identification of Cx43 binding domain in Hsc70. Expression vectors encoding various Hsc70s were transfected into HuH-7 cells, and binding experiments were carried out as described in Figure 1B. The immunocomplexes were analyzed by western blotting (WB) using anti-T7 antibody.
HuH-7 cells was performed by using dominant negative form of Cx43 (DNCx43), which is mutant Cx43 with a deletion of amino acids 130–137 and does not form gap junction33,34 (see Supplementary Fig. S4a). Overexpression of DNCx43 significantly decreased the BrdU incorporation, from 39% to 11%. As expected, co-transfection of DNCx43 with Hsc70 was sufficient to restore the BrdU incorporation, from 11% to 30% (see Supplementary Fig. S4b and c).

Hsc54, a splice variant of Hsc70, encodes the protein lacking 153 amino acid residues of Hsc70 in a part of the protein-binding and variable domains (Fig. 5a)35,36. To test further whether Cx43 specifically interacts with Hsc70 via protein-binding domain, we performed pulldown assay using GST-Cx43CT228-382 to bind Hsc70- or Hsc54-overexpressed cell lysates. Predictably, Cx43 bound to Hsc70 but not to Hsc54 (Fig. 5b). In addition, we performed BrdU incorporation assay using Hsc54. Compared with Cx43-transfected groups, co-transfected Cx43 with Hsc54 recovered somewhat the level of BrdU incorporation; however, no significant difference was observed between these two groups (Fig. 5c and d). Similar results were also obtained in both HeLa (see Supplementary Fig. S5) and Caco-2 cells (data not shown).

Moreover, we examined whether the specific protein-protein interaction between Cx43 and Hsc70 is necessary for Cx43 to regulate G1/S transition. BrdU incorporation assays were carried out under treatment with 15-DSG, which has a peptidomimetic structure and is an immunosuppressive agent, binds specifically to Hsc70 and is also thought to preclude peptide binding to Hsc7029,30. We treated HuH-7 cells with 10 \( \mu \text{g/ml} \) 15-DSG. Compared with non-15-DSG-treated groups, in 15-DSG-treated groups, no significant differences in BrdU incorporation between control and Cx43-overexpressed cells were observed (Fig. 5e). Based on these results, it seems that the Cx43-mediated cell cycle suppression is probably achieved by the association with Hsc70.

Collectively, our data demonstrated that overexpression of Cx43 significantly suppresses G1/S cell cycle transition in a GJIC-independent manner, and that this suppression can be restored by co-overexpression with Hsc70 or treatment with 15-DSG, but not by

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**Figure 3** Association among Cx43, Cyclin D1, and Hsc70. (a) Hsc70 interacts with cyclin D1 or Cx43. Co-immunoprecipitation of Hsc70 with cyclin D1 or Cx43. Lysates were prepared from HuH-7 cells co-transfected with T7-tagged Hsc70 and cMyc-tagged cyclin D1 or wild-type Cx43 expression plasmids. Lysates were immunoprecipitated (IP) either by anti-cMyc antibody and control IgG (left) or by anti-Cx43 antibodies and the IgG (right). The immunocomplexes were analyzed by western blotting (WB) using anti-T7 antibody. (b) Absence of interaction between cyclin D1 and Cx43. Co-immunoprecipitation of Cx43 with cyclin D1. Lysates were prepared from HuH-7 cells co-transfected with wild-type Cx43 and cMyc-tagged cyclin D1 expression plasmids. Lysates were immunoprecipitated (IP) either by anti-cMyc antibody and control IgG (left) or by anti-Cx43 antibodies and the IgG (right). The immunocomplexes were analyzed by western blotting (WB) using anti-Cx43 antibodies (left) or anti-cMyc antibody (right). (c) Competition assay of Cx43 and cyclin D1 for binding with Hsc70. HuH-7 cells were transfected with equivalent plasmids of both T7-tagged Hsc70 and cMyc-tagged cyclin D1, and with increasing amounts of Cx43. Lysates were immunoprecipitated (IP) by anti-T7 antibody. The immunocomplexes were analyzed by western blotting (WB) using anti-T7-tag, anti-Cx43, and anti-cMyc-tag antibodies.
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Co-overexpression with Hsc54. These results suggest that the inter-

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Discussion

Cx43 has been reported to act as a tumor suppressor in various cell

lines, the Cx43-mediated tumor suppressive functions are GHIC-

independent in several cell lines, and the evidence has suggested a

possible involvement of binding to proteins that regulate carcino-
genesis12,23–25. These Cx43-binding partners seem to be related to

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co-overexpression together with Hsc70, but not with Hsc54, in HuH-7, HeLa, and Caco-2 cells.

Under heat-shock condition, Hsc70 translocated from cytoplasm to nucleus dramatically. We confirmed that under heat-shock condition, overexpression of Cx43 clearly prevents the nuclear translocation of Hsc70, and suppresses the nuclear accumulation of cyclin D1, and that these Cx43-mediated effects were abolished by co-overexpression with Hsc70. Competition assays demonstrated that Cx43 competes with cyclin D1 for the interaction with Hsc70. Our data suggest that the interaction of between Cx43 and Hsc70 may function as an important factor in the cell cycle transition via regulating nuclear accumulation of cyclin D1.

Pulldown assays and immunofluorescence studies showed that Cx43CT228-382, especially 263–382, directly binds with Hsc70, and that Cx43 interacts with Hsc70 in the cytoplasm. The C-terminus of Cx43 is thought to be important in its interaction with key regulatory binding partners. Indeed, we showed that the recombinant C-terminal domain of Cx43 interacts directly with Hsc70, and that the deletion mutant Cx43JM228-263 does not. Moreover, we showed that Hsc54, a splice variant of Hsc70, is not able to bind to Cx43CT228-382. We extended these studies to show that overexpression of Cx43 in HuH-7, HeLa, and Caco-2 cells significantly suppressed the G1/S cell cycle transition, and that this effect was restored by co-overexpression with Hsc70 but not Hsc54. Considering the data together, we conclude that the C-terminus of Cx43 and the peptide binding domain of Hsc70 are important for their interaction with each other in regulating the cell cycle transition.

Figure 5 | Cx43 Binding Domain in Hsc70 Is Required for Cx43-Mediated Cell Cycle Suppression. (a) A schematic representation of Hsc70 and its splice variant, Hsc54, used for overexpression. (b) Pulldown assays with GST-Cx43CT fusion protein were performed using lysates prepared from Hsc70- or Hsc54-transfected HuH-7 cells. Expression vectors encoding Hsc70-EGFP and Hsc54-EGFP were transfected into HuH-7 cells, and binding experiments were carried out as described in Figure 1B. The bound proteins were analyzed by western blotting (WB) using anti-GFP antibody. (c) BrdU incorporation assays in HuH-7 cells. HuH-7 cells were transiently transfected with vectors encoding empty vector EGFP, Hsc54-EGFP, Cx43-EGFP alone, or both Hsc54- and Cx43-EGFP, respectively. BrdU incorporation and immunofluorescence staining were carried out as described in Figure 4a. (d) The fraction of BrdU-positive cells over total number of transfected cells. (e) Inhibition assay of protein–protein interaction. To inhibit the binding of proteins to Hsc70, HuH-7 cells were cultured with 10 μg/ml of 15-DSG for 24 hr. The fraction of BrdU-positive cells over total cells was determined. The data shown in (d) and (e) are means and SDs (error bars) of a representative experiment performed in triplicate at least. *p < 0.01. N.S.; not significant.
Several pieces of evidence have been accumulated suggesting a role for Hsc70 as a positive regulator of cell cycle transition and carcinogenesis\(^{29-32,38,45-49}\). Reports suggest that Hsc70 translocates from the cytoplasm into the nucleus with cyclin D1 during G1 to S phase transition\(^{30,31}\). In our experiments, Hsc70 translocated into the nucleus dramatically after heat shock; however, overexpression of Cx43 clearly prevented the nuclear translocation of Hsc70. These results suggest that Cx43 prevents Hsc70 from translocating into the nucleus through withholding Hsc70 in the cytoplasm. Interestingly, some proteins are known to bind to Hsc70. For example, previous reports showed that Hsc70 regulates the nuclear accumulation of cyclin D1, which is a key player in G1 to S phase cell cycle transition\(^{30,31}\). This evidence raises an interesting possibility that Cx43 may through directly binding to Hsc70 compete with

Figure 6 | Cx43 Prevents the Nuclear Translocation of Hsc70. (a) Hsc70 translocated into the nucleus under heat shock. HuH-7 cells were heat shocked at 42°C for 2 hr and stained with anti-Hsc70 antibodies. (b) Overexpression of Cx43 prevented the nuclear translocation of Hsc70 under heat shock. HuH-7 cells were transfected with control vector EGFP or Cx43-EGFP. Forty-eight hours after transfection followed by heat shocked at 42°C for 2 hr, cells were stained with anti-Hsc70 antibodies. (c) Ratio of nuclear/cytoplasmic (N/C) fluorescence intensities in Hsc70. After heat shock, at least 30 transfected HuH-7 cells of control EGFP or Cx43-EGFP were examined and quantified by confocal scanning microscopy. The data shown in (b) were plotted. The horizontal lines represent mean values. *p < 0.01.

Figure 7 | Interaction of Cx43 and Hsc70 Decreases Nuclear Accumulation of Cyclin D1. (a) Effect of Cx43 on nuclear accumulation of cyclin D1. HuH-7 cells were transfected with control plasmid mRFP or Cx43-mRFP, or both Cx43-mRFP and Hsc70-DsRed2. Immunofluorescence staining with anti-cyclin D1 antibodies was carried out. Nuclear accumulation of cyclin D1 was examined and quantified by confocal laser scanning microscopy. (b) Ratio of nuclear/cytoplasmic fluorescence intensities in cyclin D1. At least 30 transfected HuH-7 cells of control mRFP or Cx43-mRFP, or both Cx43- mRFP and Hsc70-DsRed2 were examined and quantified. The data shown in (a) were plotted. The horizontal lines represent mean values. *p < 0.01.
Hsc70-associated proteins, such as cyclin D1 in the cytoplasm, and as a result, the accumulation of cyclin D1 in the nucleus is prevented. In fact, in our experiments, overexpression of Cx43 resulted in decreased nuclear accumulation of cyclin D1, and this effect was abolished by co-overexpression with Hsc70. Moreover, Cx43 probably could not directly interact with cyclin D1, but competes with cyclin D1 for interaction with Hsc70. As suggested above, Cx43-regulated nuclear accumulation of cyclin D1 could occur through competitive cyclin D1 binding to common partner Hsc70.

Previous reports showed that Hsc70 directly interacts with p27 during G1/S transition29,30 and that Cx43 increases the expression level of p27 via negative regulation of Skp2, which regulates p27 data provide the first evidence that the interaction of Cx43 withpendent mechanism that involves its COOH-terminal domain. Ourin the cell cycle suppression, whereas overexpression of Cx43 doesnot increase the expression of p27, but conversely decreases it. Ourresult suggests that the Cx43-mediated cell cycle suppression seemsnot to be dependent on Skp2-regulated p27 degradation pathway.

In summary, we showed that Cx43 possesses cell cycle suppressivefunctions in HuH-7, HeLa, and Caco-2 cells through a G1Jc-indepen-dent mechanism that involves its COOH-terminal domain. Ourour data provide the first evidence that the interaction of Cx43 with Hsc70 probably plays a critical role during G1 to S phase cell cycle transition in both Cx43-positive and -negative cell lines, and that Cx43 prevents nuclear translocation of Hsc70 resulting in the decreased nuclear accumulation of cyclin D1. Collectively, ourcurrent findings suggest that Cx43 is involved in the cell cycle progres-sion of G1/S transition through a novel interacting protein, Hsc70,which regulates nuclear accumulation of cyclin D1.

**Methods**

**Plasmid constructions.** The pEGFP.Hsc70 and pEGFP.Hsc54 were constructed as previously described31. The pBRed2.MSC70 and various forms of pCAT7.HHSC7032 were provided by Y. Imamura (Matsumoto Dental University). Various forms of GST-Cx4331 were provided by B. N. Giepmans (University of California San Diego [UCSD]). The pEGFP.Cx43, pmiRFP.Cx43, pDNA3.1.Cx43, and pEGFP.DNcX43 were constructed as described32-34. The pEY.cyclin D1 was provided by P. Zou (Keio University)35.

**Cell lines, cell culture, and transfection.** HuH-7 and HeLa cells were cultured in Dulbecco’s Modified Eagle’s Medium (DMEM) supplemented with 10% (vol/vol) fetal bovine serum (FBS), glutamine (2 mM/l), penicillin (100 U/ml), and streptomycin (100 U/ml). Caco-2 cells were grown in Minimum Essential Medium (MEM) containing 20% serum, 2 mM glutamine, 100 U/ml penicillin, and 100 U/ml streptomycin. For overexpression of targets, transient transfection of HuH-7, HeLa or Caco-2 cells was carried out using Lipofectamine 2000 reagent (Invitrogen, Carlsbad, CA) according to the manufacturer’s instruction. After 48 hr of transfection, cells were used for assays.

**RNA interference.** All siRNAs against human Skp2, and Silencer negative control number 1 siRNA (NC) were purchased from Santa Cruz Biotechnology (Santa Cruz, CA).

Lipofectamine 2000 (Invitrogen) was used to transfect siRNAs into the cells (final concentration, 10 nmol/l) according to the manual. After 48 hr of transfection, immunofluorescence analysis was carried out.

**Co-immunoprecipitation, GST pulldown assay, and western blotting.** The following antibodies were used. Anti-Cx43 (Sigma, St. Louis, MO), anti-T7-tag (Novagen, Madison, WI, USA), and anti-c-Myc-tag (Wako Pure Chemical, Osaka, Japan) were used in co-immunoprecipitation. Anti-Hsc70 (B-6), anti-cGFP (B-2), anti-cyclin D1 (M-20), anti-Skp2 (H-435), anti-p27 (C-19) and anti-GAPDH (4011) were from Santa Cruz Biotechnology (Santa Cruz). Co-immunoprecipitation, GST pulldown, and western blotting analyses were performed as previously described11-14. Co-immunoprecipitated proteins were separated on 12% SDS-PAGE gels and visualized by Coomassie brilliant blue. We used a service provider (GFOMINE, Inc., Korea) to analyze Cx43-co-immunoprecipitated proteins. Briefly, the protein bands were extracted and subjected to MALDI-TOF mass spectrometry analysis.

**Competition assay.** HuH-7 cells were transfected with 1 μg of both pCAT7.HHSC70 and pEY.cyclin D1.myc, and with increasing amounts of pCDNA3.1.Cx43 (0, 1, 2, and 4 μg, respectively). The cell lysates were co-immunoprecipitated with anti-T7-tag antibody. The immunoprecipitated proteins were immunoblotted with anti-T7-tag, anti-Cx43, and anti-cMyc antibody. The signals were measured by Image Quant LAS5000 (GE Healthcare UK Ltd., Buckinghamshire, England).

**BrDU incorporation assay.** For BrDU incorporation assay, cells were plated onto 35-mm glass covered dishes. After 20 hr, the cells were transfected with various plasmids of interest. After 48 hr of transfection, cells were labeled with 10 μM BrdU for an additional 2 hr. Then, the cells were fixed and detected by using 5-Bromo-2'-deoxyuridine Labeling and Detection Kit (Roche, Basel, Switzerland). For protein interaction inhibition, cells were cultured with 10 μg/ml of 15-deoxyspergualin (15-DSG) (Sigma-Aldrich, Tokyo, Japan) for 24 hr before BrdU incorporation. At least 100 transfected cells were counted in 40 fields for each sample, and each experiment was performed three times. The Student’s t test was performed for statistical evaluation of a difference; P < 0.01 was considered to be statistically significant.

**Immunofluorescence staining.** The cells were seeded on 35-mm glass-covered dishes for 20 hr prior to transfection with plasmids of interest. In heat-shock condition, transfected cells were heat-shocked at 42°C for 2 hr36. After 48 hr of transfection, the cells were rinsed twice with PBS and fixed with 2% PFA for 1 hr at room temperature. The cells were permeabilized with 0.3% Triton in PBS for 10 min and blocked with 3% skim milk in PBS. After washing five times with PBS, cells were incubated with secondary antibodies: Alexa Fluor 488 (594) goat anti-rabbit IgG (1:1000, Invitrogen-Molecular Probes); and Alexa Fluor 488 (594) goat anti-mouse IgG (1:1000, Invitrogen-Molecular Probes) for 1 hr at room temperature. Nuclei were detected by TO-PRO3 (1:1000, Invitrogen-Molecular Probes) staining. Images were analyzed by using confocal laser scanning microscopy (FV1000, Olympus, Tokyo, Japan).

**Nucleocytoplasmic localization analysis.** Nucleocytoplasmic localization of proteins was examined by confocal laser scanning microscopy (FV1000, Olympus). For evaluation of nucleocytoplasmic localization, the ratio of nuclear/cytoplasmic fluorescence intensities was determined and quantified in at least 30 cells per condition. The images were analyzed using Image J software (National Institutes of Health, Bethesda, MD, USA).

**Statistical analysis.** The data shown were evaluated for statistical significance with Student’s t test. P < 0.01 was considered to be statistically significant.
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**Author contributions**

P.D. and T.T. designed the experiments; T.H. conducted most of the experiments in this study; P.D. performed partial experiments; F.T. and Y.M. provided series plasmids of pEGFP.Hsc70 and pEGFP.Hsc54; H.H., Y.H. and E.O. gave technical support and conceptual advice; T.H., P.D. and T.T. wrote the manuscript.

**Additional information**

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