Wnt5a controls Notch1 signaling through CaMKII mediated degradation of the SMRT corepressor protein.

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Running title: CaMKII regulates Notch1 signaling.

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**Keywords:** Notch1, Wnt5a, CaMKII, SMRT, RBP-Jk, Ubiquitination, Proteasomal degradation.

**Background:** Wnt and Notch signaling pathways belong to a highly conserved network that controls gene expression.

**Results:** Wnt5a enhances a Notch1 signaling through downregulation of SMRT by CaMKII.

**Conclusion:** CaMKII is crucial for the regulation between Notch and Wnt5a signaling.

**Significance:** The findings of this study may begin to shed some light onto what may be a signal cross-talk mechanism of Notch1 signaling and the CaMKII.

**SUMMARY**

Serine-threonine Ca²⁺/calmodulin-dependent protein kinase II (CaMKII) is the key component in non-canonical Wnt5a signaling and has been shown to regulate its signaling. In this study, we found that CaMKII induced by Wnt5a remarkably reduced the protein stability of the silencing mediator of retinoic acid and thyroid hormone receptor (SMRT), a co-repressor of Notch signaling, through proteasomal degradation. Wnt5a was found to enhance Notch1 intracellular domain (Notch1-IC) transcription activity, which could be inhibited by treatment with KN93, a CaMKII inhibitor. The kinase activity of CaMKII was essential for the activation of Notch signaling. We also determined that CaMKII could enhance the association between Notch1-IC and RBP-Jk. Furthermore, the physical association between RBP-Jk and SMRT was substantially suppressed by CaMKII. We demonstrated that CaMKII directly bound and phosphorylated SMRT at Ser1407, thereby facilitating SMRT translocation from the nucleus to the cytoplasm and proteasome dependent degradation. These results suggest that CaMKII down-regulated the protein stability of SMRT through proteasomal degradation.

Notch proteins participate in conserved pathways that regulate cell fate determination, cell differentiation, cell proliferation and cell death (1-4). Mammalian Notch receptors (Notch1–Notch4) are cell-surface type I transmembrane proteins that harbor a large extracellular domain involved in ligand binding and a cytoplasmic domain involved in signal transduction (5-7). The Notch1 receptor is processed in the trans-Golgi network by proteases of the furin (S1 cleavage) during transport to the cell surface, where it is expressed in heterodimeric form (8). At the cell surface, Notch can interact with ligands, Jagged and Delta. The transmembrane C-terminal fragment of Notch is generated via proteolytic cleavage (S2 cleavage), which subsequently cleaves the Notch fragment by gamma-secretase (S3 cleavage), an enzyme complex that contains presenilin (PS1), nicastrin (NCT), anterior pharynx defective (APH-1) and presenilin enhancer-2 (PEN-2). This process then results in the release of the Notch intracellular domain (Notch-IC) (9,10) (11-23). Notch1-IC translocates to the nucleus and interacts with various transcriptional activation complexes including CSL (CBF1/RBP-Jk, suppressor of hairless, Lag-1), coactivator protein MAML-1 (mastermind-like-1), and p300/CREB-binding protein, which possess histone acetyltransferase (HAT) activity, leading to transcriptional activation of downstream target genes such as those encoding Hes1, Hes5, Hes7, and...
Hey1, Hey2, HeyL, and Cyclin D (24-34). In the absence of Notch-IC, CSL recruits co-repressor factors, including SKIP (Ski-interacting protein), SMRT (silencing mediator for retinoid and thyroid hormone receptors), N-CoR (nuclear receptor co-repressor), and HDAC (histone deacetylases), resulting in the formation of a transcriptional repressor complex (6,35,36).

The SMRT corepressor participates in the repression of target gene expression through a variety of transcription factors, including the nuclear hormone receptors and promyelocytic leukemia zinc finger protein (37-39). The ability of SMRT to associate with these transcription factors and thereby to mediate repression is strongly inhibited by activation of tyrosine kinase signaling pathways, including pathways regulated by the epidermal growth factor receptor (40). Recent studies have emphasized the role of ubiquitination in regulating SMRT stability. Phosphorylation of SMRT by MEKK-1, Cdk2 and Pin1 induces conformational changes and reduces its binding affinity for transcription factors that promote its degradation (41,42). Phosphorylation of SMRT by IKKalpha has important meaning for the derepression of NF-kB and Notch target genes (43). Recent studies have emphasized the role of ubiquitination in regulating SMRT stability. The SMRT phosphorylation-dependent ubiquitination and degradation is an important negative feedback regulation that specifically terminates SMRT activation.

Wnt5a is a member of the Wnt family of secreted growth factor, which are 38-45 kDa secreted cysteine-rich proteins with hydrophobic signal peptides (44). Wnt5a is a highly conserved diffusible protein whose signal is transduced by Frizzled (Fz) receptors. In addition to Fzs, receptor tyrosine kinase-like orphan receptor 1/2 (Ror1/2), which is a single-pass transmembrane receptor with a tyrosine kinase domain, functions as a co-receptor (44,45). However, the effect of Fzs and Ror1/2 on Wnt5a signaling has not yet been examined. Wnt can be broadly divided into three signaling pathways: the canonical Wnt/β-catenin pathway, the non-canonical Wnt/Ca2+ pathway, and the non-canonical Wnt/planar cell polarity (PCP) pathway (46,47). Wnt5a is non-canonical pathway member of the Wnt family. Wnt5a is expressed in various adult tissues and are involved in a wide range of cellular processes. Signal transduction events through β-catenin-independent non-canonical Wnt5a elicits intracellular release of Ca2+, thereby activating the Ca2+/Calmodulin dependent protein kinase II (CaMKII), mitogen-activated protein kinase (MAPK), protein kinase C (PKC), and calpain to regulate cellular functions including cell migration (48-54). CaMKII is the key component in Wnt5a signaling and has been shown to regulate Wnt5a signaling. CaMKII is a multifunctional serine/threonine (Ser)/threonine (Thr) kinase that plays an important role in the regulation of cell growth, cytoskeletal organization, regulation of synaptic transmission, modulation of ion channel activity, learning and memory (55-58). CaMKII is important for prostate cancer cell survival and promotes their progression to an androgen-independent state (59). Furthermore, a recent study has shown that Calcium/calmodulin-dependent kinase II regulates notch-I signaling in prostate cancer cells (60). Wnt5a is transcribed based on multiple mechanisms, such as NF-kB, Hedgehog, TGF-beta, and Notch signaling cascades (61). Wnt5a expression in the human cells was induced by coculture and was blocked by gamma-secretase inhibition. Likewise, stimulation of Notch signaling by immobilized Jagged-1 promoted Wnt5a expression (62). Wnt5a acts as an essential downstream mediator of Notch–CSL signaling, which is dependent on expression in the keratinocyte compartment of FoxN1, a gene that plays a key role in hair follicle regulatory function (63). Despite these observations, the precise mechanisms underlying the relationship between the Notch1 and Wnt5a-CaMKII signaling pathways are still largely unknown.

In this study, we demonstrated that Wnt-5 induced CaMKII enhances the Notch1 signaling pathway through the downregulation of SMRT protein stability. CaMKII was shown to phosphorylate SMRT, which disrupts the SMRT-RBP-Jk transcription complex through SMRT protein degradation and polyubiquitination. In addition, the dissociation of SMRT from RBP-Jk is still largely unknown. CaMKII is important for prostate cancer cell survival and promotes their progression to an androgen-independent state (59). Furthermore, a recent study has shown that Calcium/calmodulin-dependent kinase II regulates notch-I signaling in prostate cancer cells (60). Wnt5a is transcribed based on multiple mechanisms, such as NF-kB, Hedgehog, TGF-beta, and Notch signaling cascades (61). Wnt5a expression in the human cells was induced by coculture and was blocked by gamma-secretase inhibition. Likewise, stimulation of Notch signaling by immobilized Jagged-1 promoted Wnt5a expression (62). Wnt5a acts as an essential downstream mediator of Notch–CSL signaling, which is dependent on expression in the keratinocyte compartment of FoxN1, a gene that plays a key role in hair follicle regulatory function (63). Despite these observations, the precise mechanisms underlying the relationship between the Notch1 and Wnt5a-CaMKII signaling pathways are still largely unknown.

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**EXPERIMENTAL PROCEDURE**

**Cell Culture and Transfection** - Human embryonic kidney (HEK) 293 cells were grown in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% Fetal Bovine Serum (FBS), penicillin (100 U/ml), and streptomycin (100 μg/ml). Along with the HEK293 cells, PS1/2+ MEF cells and PS1/2− MEF cells were cultured in Dulbecco’s modified...
Eagle’s medium (DMEM) supplemented with 10% FBS, 100 U/ml penicillin, and 100 μg/ml streptomycin. For plasmid DNA transfection, cells were plated at a density of 2 × 10^6 cells/100-mm dish, grown overnight, and transfected with the appropriate expression vectors in the presence of the indicated combinations of plasmid DNAs via the calcium phosphate and liposome method (64).

**Luciferase reporter assay** - Human embryonic kidney 293 cells were co-transfected with 4XCSL-Luc (a 4-time repeating section of the RBP-Jk target sequence, CGTGGGAA, with the luciferase gene), Hes1-Luc, Hes5-Luc, mutating the RBP-Jk binding site of Hes5-Luc, and β-galactosidase coupled with the indicated vector constructs. 48 hours after transfection, the cells were lysed in chemiluminescence lysis buffer (18.3% of 1 M K_2HPO_4, 1.7% of 1 M KH_2PO_4, 1mM phenylmethylsulfonyl fluoride (PMSF), and 1mM dithiothreitol) and luciferase activity was assayed using a luciferase assay kit (Promega). The activity of the luciferase reporter protein in the transfected cells was normalized to the β-galactosidase activity in the same cells (65). These results represent the means ± average deviation of three independent experiments. R.L.U means relative luciferase units.

**Protein accumulation assay** - For the protein accumulation assay, the cells were seeded at a density of 50-60% confluence and incubated overnight. 48 hours after transfection, cells were treated with the proteasomal inhibitor MG-132 (Sigma-Aldrich). The MG-132 concentrations in the treated cells were 0, 1 and 5 μM and the treatment time was 6 hours. Protein levels were analyzed via immunoblotting (65).

**Protein stability assay** - For the protein stability assay, the cells were seeded at a density of 50-60% confluence and incubated overnight. 48 hours after transfection, the cells were treated with 0.1 mM cycloheximide for 0, 0.5, 1, 2, 4, or 6 hours to block the synthesis of new proteins. The cells were collected at each time point and the total lysates were then lysed in RIPA/ Laemmli buffer. Protein levels of endogenous Notch1 were determined by immunoblotting with the indicated antibody (65).

**In vitro binding assay** - The recombinant GST-SMRT proteins were expressed in *Escherichia coli* strain BL21, using the pGEX system as indicated (66). The GST-SMRT proteins were then purified using glutathione-agarose beads (Sigma), in accordance with the manufacturer’s instructions. Equal amounts of GST or GST-SMRT fusion proteins were incubated with the lysates of HEK293 cells, which had been transfected for 3 hours with combinations of expression vectors at 4°C, with rotation. After incubation, the beads were washed three times with ice-cold PBS, and boiled with 20 μl of Laemmli sample buffer. The precipitates were separated via SDS-PAGE, and the pull-down proteins were detected via immunoblotting with specific antibodies.

**Co-immunoprecipitation Assays** - The cells were lysed in 1 ml of RIPA buffer for 30 minutes at 4°C. After 20 minutes of centrifugation at 12,000 × g, the supernatants were subjected to immunoprecipitation with the appropriate antibodies coupled to protein A-agarose beads. The resulting immunoprecipitates were then washed three times in phosphate-buffered saline (PBS, pH 7.4). Laemmli sample buffer was then added to the immunoprecipitated pellets. The pellets were heated for 5 minutes at 95°C and analyzed via SDS-PAGE. Western blotting was conducted with the indicated antibodies.

**Western blot analysis** - The cultured cells were harvested and lysed in RIPA buffer (50mM Tris–HCl (pH 7.5)), 150mM NaCl, 1% Nonidet P-40, 0.5% Sodium deoxycholate, 0.1% SDS, 1mM PMSF, 1mM DTT, and 2μg/ml leupeptin and aprotinin for 30 min. The cell lysates were subjected to 20 min of centrifugation at 12,000g and 4°C. The resultant soluble fraction was then boiled in Laemmli buffer and subjected to SDS–PAGE. After gel electrophoresis, the separated proteins were transferred onto polyvinylidene difluoride (PVDF) membranes (Millipore) via electroblotting. The membranes were then blocked with Tris-buffered saline solution (pH 7.4) containing 0.1% Tween 20 and 5% nonfat milk. The blotted proteins were probed with the indicated antibodies.

**Antibody list** - The blotted proteins were probed using anti-Myc antibody (9E10), anti-HA (12CA5) antibody, anti-Flag M2 (Sigma Chemical Co.), anti-Notch1-IC (Cell Signaling), anti-RBP-Jk (Santa Cruz), anti-CaMKII (Santa Cruz), anti-SMRT (Santa Cruz), anti-Presenilin1 (D39D1) rabbit mAb, anti-Presenilin 2 (D30G3) rabbit mAb (Cell Signaling Technology) followed by incubation with anti-mouse horseradish peroxidase-conjugated secondary antibodies (Amersham Biosciences, Inc.). The blots were then developed using an enhanced chemiluminescence (ECL) system (Pierce).

**Site directed mutagenesis** - The site-directed mutagenesis of cDNA encoding for SMRT was performed using a QuikChange kit (Stratagene), and CaMKII regulates Notch1 signaling.
the mutagenic primer was S1407A (5'-GGGGACCCGGCCACCCgCTGTCCTCTCAGTGC AC-3') and S1434A (5'-GAGGACAGGCCCTGCgCGGAGGTTCACGC C-3') (mismatches with the SMRT cDNA template are indicated by lowercase letters). The mutations were verified by automatic DNA sequencing. **Immunocomplex kinase assay** - The cultured cells were harvested and lysed in buffer A, containing 50 mM Tris–HCl (pH 7.5), 150 mM NaCl, 1 mM phenylmethylsulfonyl fluoride (PMSF), 2 μg/ml of leupeptin, 2 μg/ml of aprotinin, 25 mM glyceraldehyde, 0.1 mM sodium orthovanadate, 1 mM sodium fluoride, 1% NP-40, 0.5% deoxycholate, and 0.1% SDS for 30 min at 4 °C. The cell lysates were then subjected to centrifugation at 12,000 × g and 4 °C for 20 min. The soluble fraction was incubated for 3 h with the appropriate antibodies against the indicated protein kinase at 4 °C. The immunocomplexes were then coupled to protein A-agarose during an additional hour of incubation at 4 °C, after which they were pelleted via centrifugation. The immunopellets were rinsed three times with lysis buffer and then twice with 20 μl of reaction buffer that contained 0.2 mM sodium orthovanadate, 10 mM MgCl2, 2 μCi [32P]ATP, 20 mM HEPES (pH 7.4). The phosphorylated substrates were separated by SDS-PAGE and quantified using a Fuji FLA7000 phosphorImager. The GST fusion proteins that were used as substrates were expressed in E. coli using pGEX-4T (Pharmacia) and purified using glutathione–Sepharose, as described previously (66,67). The protein concentrations were determined by the Bradford method (Shimadzu). **Immunofluorescence staining** - Assays were conducted as previously described with HEK293 cells plated at a density of 1x10⁵ cells per well onto cover slips (Fisher). Cells were transfected with a total of 0.5 μg of the appropriate DNA per well using the Lipofectamine-plus reagent (Invitrogen). The transfected cells were fixed with 4% paraformaldehyde in phosphate-buffered saline (PBS), and then permeabilized with 0.1% Triton-X 100 in PBS. Cells were blocked in 1% BSA in PBS. Anti-IA or anti-Flag M2 antibodies (Sigma Chemical Co.) were employed as the primary antibodies at a dilution of 1:100, washed three times in PBS, Alexa Fluor 488 (Invitrogen) or Alexa-555 (Invitrogen) conjugated anti-mouse secondary antibody (1:100) was added, and then the DNA dye ToPro-3 was used for nuclear localization. The stained cells were evaluated for localization via confocal microscopy (Leica TCS SPE). Each image consisted of a single Z section at the same cellular level. The final images were obtained using a confocal microscopy and analyzed with LAS AF software (Leica) and at least 300 cells were counted for each experimental condition. Scale bars represent 25 μm as indicated (68).

**RESULTS**

**Wnt5a enhances the transcriptional activity of Notch1 target genes through CaMKII** - To determine whether Wnt5a is involved in regulating the transcriptional activation of Notch1 target genes, a reporter assay was performed with HEK293 cells using luciferase reporter genes. In this study, three types of luciferase reporter genes under the control of the Hes1 promoter, the Hes5 promoter, and the artificial four time repeat of RBP-Jk binding sequence were evaluated. We investigated the effect of Wnt5a on Notch1-IC transcriptional activity. HEK293 cells were transfected with 4XCSL-Luc and either the Notch1-IC or an empty vector. As expected, Notch1-IC mediated transcription activity increased in these samples (Figure. 1A). The Notch1-IC induced 4XCSL luciferase reporter activity was enhanced by co-transfection with Wnt5a in a dose-dependent manner (Figure. 1A). The bHLH proteins Hes1 and Hes5, both of which contain multiple CSL binding DNA sequences on their promoters, were identified as essential targets of Notch in epithelial cells. Therefore, we have
confirmed the effects of Wnt5a on the Notch1 signaling pathway, using the Hes1, Hes5 reporter and RMHes5 reporter systems, respectively. RMHes5-Luc was constructed by mutating the RBP-J binding site (-79 to -72: tgtgtaa) of Hes5-Luc to tgtgtga (69). Expression of Notch1-IC was determined to significantly induce activation of the Hes1 and Hes5 reporter systems (Figure. 1B-C). Wnt5a enhanced Notch1-IC induced natural Hes1 and Hes5 promoter transcriptional activity in a dose dependent manner, but not RMHes5 activity (Figure. 1B-C). Moreover, The Notch1-IC induced 4XCSL luciferase reporter activity was enhanced by purified Wnt5a in a dose-dependent manner (Figure. 1D). Based on these results, it was assumed that Wnt5a can activate the expression of the Notch1 target genes.

In order to delineate the possible role of CaMKII in the regulation of Notch1 signaling by Wnt5a in intact cells, we assessed the effects of the CaMKII inhibitor and CaMKII dominant negative on Notch1 signaling. We determined that the KN-93 or CaMKII dominant negative was able to block the effects of Wnt5a on Notch1-IC-induced transcriptional activity (Figure. 1E, F). From these results, it was assumed that Wnt5a can increase the expression of the Notch1 target gene through up-regulation of CaMKII. In order to demonstrate this assumption, Western bolt were performed for endogenous Hes1 and Hes5. The HEK293 cells were transfected with Myc-Notch1-IC, Wnt5a, or siCaMKII. The expression levels of Hes1 and Hes5 protein were increased by Wnt5a, as expected and the Wnt5a-induced expressions of Hes1 and Hes5 were decreased via coexpression of siCaMKII (Figure. 1G). Therefore, we hypothesized that CaMKII plays a role in the up-regulation of the Notch1 target gene transcriptional activity.

CaMKII activates Notch1 transcriptional activity - We attempted to determine whether or not CaMKII plays a role in the regulation of Notch1-IC mediated transcriptional activity. HEK293 cells were transfected with 4XCSL-Luc and either Notch1-IC or CaMKII. In these experiments, CaMKII was shown to activate Notch1-IC to stimulate transcription in a dose dependent manner (Figure. 2A). We have also confirmed the effects of CaMKII on the Notch1 signaling pathway, using the Hes1 and Hes5 reporter systems. Expression of the active form of Notch1 was determined to induce activation of the Hes1 and Hes5 reporter systems. CaMKII expression enhanced Notch1-IC-induced natural Hes1 and Hes5 promoter transcriptional activities, but not RMH5 activity (Figure. 2B, C).

To assess the role of CaMKII in the basal level of Notch1 signaling, we performed a transcription reporter assay. HEK293 cells were transfected with 4XCSL-Luc and CaMKII. The transcriptional activity of endogenous Notch1-IC in HEK293 cells was enhanced by CaMKII in a dose dependent manner (Figure. 2D). Similar results were also obtained using the Hes1-Luc and Hes5-Luc reporter systems (Figure. 2E-F). However, the absolute value of three reporter systems was higher in Notch1-IC and CaMKII cotransfected cells than in cells transfected with only CaMKII, suggesting that Notch1-IC and CaMKII produce synergic effects on reporter activity. Thus, taken together, our results indicate that CaMKII has a positive role in the regulation of Notch1 transcriptional activity.

CaMKII regulates Notch1 signaling activity through RBP-Jk dependent and Notch1-IC independent manner - To determine whether CAMKII regulates the transcriptional activation of Notch1 target genes through a Notch1-IC dependent mechanism, we performed a transcription reporter assay using PS1 and PS2 wild-type (PS1/2+/+) and PS1 and PS2-deficient (PS1/2-/-) MEF cells. We investigated the effect of CaMKII on the basal level of Notch1 transcriptional activity. PS1/2+/+ and PS1/2-/- MEF cells were transfected with 4XCSL-Luc and either the CaMKII. In these experiments, CaMKII was shown to activate the basal level of Notch1 reporter activity both in PS1/2+/+ and PS1/2-/- MEF cells (Figure. 3A, D). Similar results were also observed when using the Hes1-Luc, Hes5-Luc and RMHes5-Luc reporter systems in both PS1/2+/+ and PS1/2-/- MEF cells (Figure. 3B, C, E, F). These data demonstrate that CaMKII could activate the Notch1 reporter activity despite the absence of Notch1-IC generation, suggesting that CaMKII, at least in part, could regulate Notch1 target gene expression in a Notch1-IC independent manner. However, the absolute value of three reporter systems was higher in PS1/2+/+ MEF cells than PS1/2-/- MEF cells, suggesting that CaMKII could modulate Notch1 activity through a Notch1-IC dependent and independent manner.

To investigate whether the transcriptional activation of endogenous Notch1 target genes by CaMKII occurs through a RBP-Jk dependent manner, we performed a transcription reporter assay using the dominant negative form of RBP-Jk (RBP-Jk-DN). In the luciferase reporter gene assay with PS1/2 +/- MEF cells, RBP-Jk-DN was transfected and the effect of this transfection on the
transcriptional inhibition of Notch1-IC target genes was then assessed using 4XCSL-Luc. The transcriptional activity of endogenous Notch1-IC was activated by CaMKII but was inhibited by cotransfection with RBP-Jk-DN, suggesting that CaMKII activates Notch1 transcriptional activity through a RBP-Jk dependent mechanism (Figure. 3G). To investigate whether the kinase activity of CaMKII is necessary for the up-regulation of the transcriptional activity of Notch1 target genes, we used a dominant-negative, kinase deficient CaMKII mutant (CaMKII-K42M, called CaMKII-DN) to block the kinase activity of CaMKII. In the luciferase reporter gene assay with PS1/2−/− MEF cells, CaMKII-DN was transfected instead of CaMKII, and the effect of this transfection on the transcriptional activation of endogenous Notch1-IC target genes was assessed using 4XCSL-Luc. The transcriptional activity of endogenous Notch1-IC was enhanced by CaMKII but was not enhanced by transfection with the dominant negative CaMKII (Figure. 3H). We investigated the effect of CaMKII-DN on basal Notch1-IC transcriptional activity with 4CSL-Luc in PS1/2−/− MEF cells. Basal Notch1-IC transcriptional activity was decreased by CaMKII-DN in a dose dependent manner (data not shown). Similar results were also observed using the 4XCSL-Luc reporter system with KN-93, a CaMKII chemical inhibitor. The CaMKII induced 4XCSL luciferase reporter activity was inhibited by treating PS1/2−/− MEF cells with KN-93 (Figure. 3I). To determine the role of endogenous Notch1-IC in PS1/2+/+ and PS1/2−/− MEF cells was further evaluated by the knockdown of Notch1 using DAPT, which is a gamma-secretase inhibitor. These results confirmed that expression of Notch1-IC protein level was reduced by DAPT in the PS1/2−/− MEF cells (Figure. 3J). PS1 and PS2 expression levels in PS1/2+/+ and PS1/2−/− MEF cells were tested using specific antibodies (Figure. S1). Thus, the up-regulation of the Notch1-IC activity by CaMKII could be dependent on the intact kinase activity of CaMKII through RBP-Jk dependent and Notch1-IC independent mechanism.

**CaMKII modulates the binding of RBP-Jk to Notch1-IC or SMRT** - To observe the effects of Wnt5a on the molecular interactions between Notch1-IC and RBP-Jk, co-immunoprecipitation was performed in HEK293 cells that had been cotransfected with Myc-tagged Notch1-IC, Flag-tagged RBP-Jk, and Wnt5a. Immunoprecipitation was performed on cell lysates using an anti-Flag antibody, and immunoblotting was performed using the anti-Myc antibody. Notch1-IC and RBP-Jk were co-immunoprecipitated, but when they were cotransfected with Wnt5a, the band intensity corresponding to the interaction between Notch1-IC and RBP-Jk was enhanced (Figure. 4A). We evaluated the physical association between Notch1-IC and RBP-Jk by Wnt5a in immunocomplexes that were collected using anti-Myc antibody. These results confirmed that the interaction between Notch1-IC and RBP-Jk was enhanced in the presence of Wnt5a. In the presence of KN-93, the association between Notch1-IC and RBP-Jk by Wnt5a was reduced, suggesting that CaMKII kinase activity might play an important role in the Notch1-IC–RBP-Jk interaction (Figure. 4B). We next evaluated the effects of KN-93 on the association of endogenous Notch1-IC and RBP-Jk by Wnt5a in intact cells. In these experiments, HEK293 cells were transiently transfected with Wnt5a. 42 hours after transfection, cells were treated with exogenous KN-93 for 6 hours, and then the affinity between these two proteins was measured. Wnt5a facilitated the physical association between Notch1-IC and RBP-Jk and this binding was substantially suppressed by KN-93 treatment, suggesting that Wnt5a induced CaMKII activation plays a critical role in the endogenous Notch1-IC-RBP-Jk binding (Figure. 4C). In order to further assess whether CaMKII plays a positive role in Notch1-IC–RBP-Jk binding, co-immunoprecipitation was performed in HEK293 cells with co-transfection of Myc-tagged Notch1-IC, Flag-tagged RBP-Jk, HA-tagged CaMKII, and HA-tagged CaMKII-DN. In the presence of CaMKII, the interaction between Notch1-IC and RBP-Jk was higher than in the absence of CaMKII transfection and had no effect on CaMKII-DN (Figure. 4D).

To evaluate the effects of Wnt5a on the molecular interactions between SMRT and RBP-Jk, co-immunoprecipitation was performed in HEK293 cells that had been co-transfected with Flag-tagged SMRT, Myc-tagged RBP-Jk, and Wnt5a. SMRT and RBP-Jk were co-immunoprecipitated, but when they were co-transfected with Wnt5a, the band corresponding to the SMRT-RBP-Jk complex disappeared (Figure. 4E). We also examined the effects of KN-93 on the association of endogenous RBP-Jk and SMRT by Wnt5a in intact cells. In these experiments, HEK293 cells were transiently transfected with Wnt5a. 42 hours after transfection, the cells were treated with exogenous KN-93 for 6 hours, and then affinity between these two proteins was measured. Wnt5a inhibited the physical
association between RBP-Jk and SMRT and this binding was substantially rescued by KN-93 treatment, suggesting that Wnt5a induced CaMKII activation plays a critical role in the endogenous RBP-Jk-SMRT binding (Figure. 4F). To confirm the role of CaMKII in the regulation of RBP-Jk-SMRT transcription repressor complex, co-immunoprecipitation was performed in HEK293 cells that had been co-transfected with Flag-tagged SMRT, Myc-tagged RBP-Jk, HA-tagged CaMKII and HA-tagged CaMKII-DN. SMRT and RBP-Jk were co-immunoprecipitated, but when they were co-transfected with CaMKII, the band corresponding to the SMRT-RBP-Jk complex disappeared (Figure. 4G). Furthermore, the dominant-negative mutant of CaMKII (CaMKII-DN) had no effect on the binding between RBP-Jk and SMRT, suggesting that the kinase activity of CaMKII is essential for the physical binding between RBP-Jk and SMRT (Figure. 4G). Interestingly, in the cell lysate immunoblot, the level of endogenous and ectopic expressed SMRT proteins, but not Notch1-IC and RBP-Jk, was significantly down-regulated upon coexpression with Wnt5a or CaMKII (Figure. 4E-G), which shows that CaMKII may regulate the steady state level of SMRT protein. These results indicate that Wnt5a mediated CaMKII activation facilitates not only the formation of the Notch1-IC-RBP-Jk transcription activator complex but also the dissociation of the RBP-Jk-SMRT transcription repressor complex due to the down regulation of SMRT protein level.

**CaMKII down-regulates the steady state level of SMRT protein** - Next, HEK293 cells were subjected to western blot analysis to determine whether CaMKII plays a role in the regulation of the SMRT protein level. Cells were co-transfected with Flag-tagged SMRT, HA-tagged CaMKII and HA-tagged CaMKII-DN. We found that the SMRT protein level was reduced upon co-transfection of CaMKII but was not reduced upon co-transfection of CaMKII-DN (Figure. 5A). This result showed that the kinase activity of CaMKII is essential for the regulation of the SMRT protein level. In an effort to determine whether the degradation of SMRT proteins by CaMKII is mediated by the proteasome pathway, the proteasome inhibitor MG132 was administered to SMRT and CaMKII-expressing cells. The cells were then subjected to 6 hours of proteasome inhibitor treatment, and the SMRT protein level was measured via an immunoblot assay. Our studies demonstrated that the SMRT protein level was reduced in the presence of CaMKII, but was significantly restored by treatment with MG132 (Figure. 5B). In an effort to evaluate the possible role of CaMKII in the regulation of SMRT protein stability, we transfected HEK293 cells with Flag-tagged SMRT, HA-tagged CaMKII and HA-tagged CaMKII-DN, and the quantity of remaining SMRT was analyzed after various periods of cycloheximide treatment. We determined the protein stability of SMRT in HEK293 cells by cycloheximide treatment with or without CaMKII. After cycloheximide treatment, the SMRT protein level gradually decreased, where approximately half of the protein was degraded after 1.5 hours without CaMKII (Figure. 5C). The SMRT protein also rapidly decreased in the presence of CaMKII, where approximately half of the protein was degraded after 0.8 hours (Figure. 5C). However, the SMRT protein level gradually increased in the presence of dominant negative CaMKII, where approximately half of the protein was degraded after 5 hours (Figure. 5C). No reduction in the level of beta-actin, which was used as a control, was observed (Figure. 5C). HEK293 cells were subjected to western blot analysis to determine whether purified Wnt5a plays a role in the regulation of the endogenous SMRT protein level. The expression level SMRT protein was reduced by purified Wnt5a in a dose-dependent manner (Figure. 5D). We then attempted to characterize the involvement of CaMKII activity in the poly-ubiquitination of SMRT. Cells were co-transfected with Flag-tagged SMRT, GFP-tagged CaMKII and HA-tagged Ub, and were then subjected to ubiquitination analysis. Immunoblot analysis of the Flag immunoprecipitates from the transfected cells with an anti-HA antibody showed that CaMKII facilitated the ubiquitination of SMRT and KN93 prevented the ubiquitination of SMRT (Figure. 5E). These results showed that the kinase activity of CaMKII is essential for regulating the SMRT protein level.

**Physical interaction of CaMKII with SMRT in intact cells** - In order to delineate more precisely the manner in which CaMKII facilitates Notch1-IC transcriptional activity, we conducted a series of in vitro binding and communoprecipitation experiments. In the in vitro binding experiments, purified GST or GST-SMRT proteins were immobilized onto GSH-agarose. HA-CaMKII expressing cell lysates were incubated with either GST or GST-SMRT, both of which were immobilized onto GSH-agarose. The interaction between GST-SMRT and CaMKII was detected on
CaMKII regulates Notch signaling.

bead complexes (Figure 6A). However, no interaction was observed between GST and CaMKII (Figure 6A). We next attempted to determine whether CaMKII could physically associate with SMRT. Here, HEK293 cells were cotransfected with vectors encoding Flag-tagged wild-type SMRT and HA-tagged CaMKII and were then subjected to coimmunoprecipitation analysis after treatment with MG132 (5 μM) for 6 hours. Immunoblot analysis of HA immunoprecipitates from cells transfected with an anti-Flag antibody revealed that Flag-SMRT physically associated with HA-CaMKII in the cells (Figure 6B). We also examined whether endogenous CaMKII and SMRT could interact in intact cells. Using HEK293 cells, immunoblot analysis of the SMRT immunoprecipitates using an anti-SMRT antibody indicated that endogenous CaMKII and SMRT were physically associated (Figure 6C). SMRT has the positions of two silencing domains (SD1 and SD2) and of two nuclear receptor interaction domains (RID1 and RID2) in its structure. We investigated which of these domains might be involved in the interaction between SMRT and CaMKII. We used purified GST and GST-SMRT deletion mutant proteins: SMRT-SD1, SMRT-SD2, SMRT-RID1 and SMRT-RID2. In the in vitro binding studies, purified GST and GST–SMRT deletion mutant proteins were immobilized on GSH agarose. Cell lysates expressing HA-CaMKII were incubated with either GST or with GST–SMRT deletion mutants immobilized on GSH-agarose. The interaction between GST–SMRT-RID1 and CaMKII was detected on bead complexes (Figure 6D). In these experiments, we demonstrated that SMRT physically interacts with CaMKII via the RID1 domain.

The phosphorylation of SMRT by CaMKII is critical for degradation of SMRT protein - We next conducted an in vitro kinase assay using CaMKII and purified GST–SMRT. The CaMKII immunocomplexes prepared from the HEK293 cells catalyzed the phosphorylation of purified recombinant GST–SMRT. Dominant negative CaMKII did not phosphorylate SMRT protein (Figure 7A). Next, we conducted an in vitro kinase assay with HA-CaMKII and purified GST–SMRT–deletion mutants (GST–SMRT-SD1, GST–SMRT-SD2, GST–SMRT-RID1 and GST–SMRT-RID2). Serial deletion mutants of SMRT were employed in the CaMKII phosphorylation reaction in order to determine the phosphorylation site of SMRT. CaMKII phosphorylated GST–SMRT-RID2 but did not phosphorylate the other three deletion mutants (Figure 7B). These results demonstrated that the phosphorylation sites were located between residues 1291-1495 of SMRT. This region harbors two conserved sequence for CaMKII phosphorylation (R-X-X-S/T): Ser1407 and Ser1434. Furthermore, using site-directed mutagenesis, we determined that the replacement of Ser1407 of SMRT with alanine reduced the in vitro phosphorylation of the recombinant protein by the CaMKII immunoprecipitates (Figure 7C). Moreover, we found that SMRT (S1407A) was resistant to CaMKII-induced degradation, which implies that the CaMKII-induced phosphorylation of SMRT is crucial for the degradation of the SMRT protein (Figure 7D). We then used coimmunoprecipitation to examine the role of phosphorylation in the physical association between RBP-Jk and SMRT. In these experiments, HEK293 cells were cotransfected with vectors coding for Myc-tagged RBP-Jk, HA-tagged CaMKII, Flag-tagged SMRT, and Flag-tagged SMRT (S1407A) and were then subjected to coimmunoprecipitation analysis. Immunoblot analysis of the Flag immunoprecipitates from the cells transfected showed that CaMKII does not prevent the physical association between RBP-Jk and SMRT (S1407A) in cells (Fig. 7E). Next, we examined the effect of CaMKII on the transcriptional repressive activity of SMRT and SMRT (S1407A) mutant in HEK293 cells. CaMKII was shown to activate the basal level of Notch1-induced transcriptional activity in the presence of wild type SMRT; however, CaMKII did not activate the basal level of Notch1-induced transcriptional activity in the presence of wild type SMRT(S1407A) (Figure. 7F). Moreover, we attempted to determine whether CaMKII could modulate SMRT localization within cells. To determine the effect of CaMKII on SMRT localization, HEK293 cells were cotransfected with vectors coding for HA-tagged CaMKII, Flag-tagged SMRT, and Flag-tagged SMRT (S1407A) and were then subjected immunofluorescence staining after treated with MG132 (5 μM) for 6 hours. As expected, SMRT was localized predominantly in the nucleus and CaMKII was distributed throughout the nucleus and cytoplasm (Fig. 7G, H). As shown in Figure 7G and 7H, coexpression of CaMKII facilitated the translocation of SMRT from the nucleus to cytoplasm. We also found that CaMKII was localized predominantly in the cytoplasm in the presence of SMRT. Furthermore, CaMKII did not...
affect the subcellular localization of SMRT (S1407A). We found that SMRT (S1407A) was localized predominantly in the nucleus, suggesting that CaMKII mediated phosphorylation of SMRT was critical to the regulation of cellular localization of the SMRT protein. These data suggest that CaMKII-mediated phosphorylation of SMRT is an important step in CaMKII mediated activation of Notch1 signaling in intact cells.

DISCUSSION
Wnt and Notch signaling pathways belong to a highly conserved network that controls gene expression during development, adult tissue maintenance and disease (70). Recent studies have implied that these pathways crosstalk in mammals (71,72). A recent study has suggested that CaMKII, a component of the non-canonical Wnt signaling pathway, regulates Notch1 signaling pathway (60). However the mechanism of this regulation has not been clearly established. In this study, we demonstrated that Wnt5a induced CaMKII enhances Notch transcriptional activity. CaMKII physically interacts with SMRT through the RID1 domain and disrupts the SMRT-RBP-Jk transcription complex through degradation of SMRT. Dissociation of SMRT from RBP-Jk provoked by CaMKII activates Hes1 and Hes5 transcriptional activity. CaMKII can phosphorylate SMRT and activate the Notch1 signaling pathway. The kinase activity of CaMKII was required for up-regulation of the transcription activity of Notch1 target genes. To the best of our knowledge, this is the first demonstration of CaMKII-mediated positive regulation of the Notch signaling through proteasomal degradation of SMRT.

Several recent studies have shown that ubiquitination mediated protein degradation plays a role in the regulation of Notch1 signaling through degradation of Notch1 ligands, Notch1 itself, and RBP-Jk (65,66,73-75). This study provides the first evidence that the CAMKII is an endogenous positive regulator of the Notch signaling pathway through degradation of the SMRT corepressor. When CAMKII activity was enhanced by Wnt5a, the Notch1 transcriptional activity was significantly increased in HEK293 cells. This was examined in two different experiments: treating the cells with a CaMKII inhibitor, and expressing the dominant-negative mutant CaMKII. In both experiments, the basal Notch1-IC transcriptional activity was substantially suppressed. On the other hand, when CAMKII was overexpressed, the Notch1 transcriptional activity was markedly increased in a dose-dependent manner. Our findings provide unequivocal evidence that CAMKII is an endogenous signaling agent that up-regulates Notch1-IC transcriptional activity. Previous studies have suggested that CaMKII regulates Notch1 signaling in C4-2B cells. Inhibition of CaMKII activity by KN93 leads to a decrease in the steady state level of Notch1-IC protein and down-regulates the expression of Notch1 target gene Hes-1 and its promoter activity (60). In contrast, the steady state level of Notch1-IC was not changed under these experimental conditions.

The transcriptional activation of downstream target genes by Notch1-IC depends on the association of Notch1-IC and RBP-Jk within the nucleus (6). Wnt5a and CaMKII facilitate the interaction between Notch1-IC and RBP-Jk and this interaction was inhibited by a CaMKII inhibitor or dominant negative CaMKII in intact cells. After binding to DNA, RBP-Jk recruits a co-repressor complex that includes SMRT and HDAC1 to exert its inhibitory effect on transcription (6). We found that Wnt5a and CaMKII inhibit the interaction between Notch1-IC and RBP-Jk and this interaction was rescued using a CaMKII inhibitor or dominant negative CaMKII in intact cells. Based on these results, we propose that endogenous CaMKII, which was induced by Wnt5a, may facilitate Notch1 transcriptional activity due to the stable association between Notch1-IC and RBP-Jk and dissociation between RBP-Jk and SMRT corepressor. Surprisingly, in the cell lysate immunoblot, the SMRT protein level was down-regulated upon co-transfection of Wnt5a or CaMKII and the SMRT protein level was restored using a CaMKII inhibitor or dominant negative CaMKII. These results show that Wnt5a may regulate the level of the SMRT protein.

We also demonstrated that the RID1 domain of SMRT physically interacts with CaMKII and the kinase activity of CaMKII was shown to be essential for the regulation of the SMRT protein level. Recent studies have suggested that SMRT, a member of Notch co-repressor complex, is phosphorylated and degraded in a proteasome-dependent manner (41-43,76). Degradation of SMRT is associated with the regulation of the Notch1 signaling pathway. In this study, we found that CaMKII stimulated the proteasomal degradation of ectopically expressed SMRT, and the reduction of SMRT levels by CaMKII was also observed through a proteasome-dependent mechanism. Our findings
show that the kinase activity of CaMKII plays a crucial role in the proteasomal degradation of SMRT. Furthermore, the kinase activity of CaMKII was critical to the binding and inhibition of the Notch1 signaling pathway. Amino acids Arg-X-X-Ser/Thr motif were preferentially phosphorylated by CaMKII and SMRT contains several putative serine and threonine phosphorylation sites (77,78). We found that SMRT contains a conserved sequence for phosphorylation by CaMKII. CaMKII phosphorylated the SMRT protein and the phosphorylation sites were shown to be located between residues 1291-1495 of SMRT, a region that harbors two conserved serine residues: Ser1407 and Ser1434. Furthermore, we demonstrated that CaMKII-mediated SMRT phosphorylation on Ser1407 results in a decrease in the degradation of SMRT protein. CaMKII was also shown to positively regulate the transcriptional activation of the Notch1-IC target genes and the stability of SMRT protein by phosphorylation of SMRT at the Ser1407 residue. Thus, enhanced SMRT phosphorylation may be a possible mechanism for the CaMKII-mediated ubiquitination and proteasomal degradation of SMRT (Figure. 8).

Previous reports have shown that MEKK1, Cdk2 and Pin1, CK2, and IKKalpha induced phosphorylation mediated SMRT translocation from the nucleus to the cytoplasm and proteasome-dependent degradation (41-43,76). In this report, we also found that CaMKII mediates a similar phosphorylation event on SMRT corepressors at S1407 and thereby facilitates translocation from the nucleus to the cytoplasm. These results explain why CaMKII-induced phosphorylation of SMRT was required for proteasomal degradation and translocation from the nucleus to the cytoplasm.

In summary, Wnt5a induced CaMKII enhances Notch transcriptional activity. CaMKII physically interacts with SMRT and disrupts the SMRT-RBP-Jk transcription complex through degradation of SMRT protein. Dissociation of SMRT from RBP-Jk induced by CaMKII activates Hes1 and Hes5 transcriptional activity. Furthermore, CaMKII can phosphorylate SMRT and activate the Notch1 signaling pathway. The kinase activity of CaMKII was required for the up-regulation of the transcription activity of Notch1 target genes. The results of this study suggest that CaMKII plays a pivotal role in the signal cross-talk between Wnt5a and Notch signaling through downregulation of the SMRT corepressor protein.
CaMKII regulates Notch1 signaling.

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CaMKII regulates Notch signaling.


Acknowledgements - We thank Raphael Kopan (Washington University Medical School, St. Louis, USA) for the providing Notch related constructs, Martin L. Privalsky (University of California, Davis, USA) for SMRT related constructs, S. Diane Hayward (Johns Hopkins University School of Medicine, USA) for SMRT, Kunihiro Matsumoto (Nagoya University, Japan) for CaMKII and Wnt5a, and Tetsuya Taga (Tokyo Medical and Dental University, Japan) for Hes-5-Luc and RMHes-5-Luc constructs. This study was supported by a grant of the Korea Healthcare technology R&D Project, Ministry of Health & Welfare, Republic of Korea. (A100455)

FOOTNOTES
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The abbreviations used are: Notch1-IC, Notch1 intracellular domain; CaMKII, Ca²⁺/calmodulin-dependent protein kinase II; SMRT, Silencing mediator of retinoic acid and thyroid hormone receptor; RBP-Jk, Recombining binding protein suppressor of hairless; Ub, Ubiquitin.

FIGURE LEGENDS

FIGURE 1. Wnt5a enhances the transcriptional activity of Notch1 target genes through CaMKII. (A-C) HEK293 cells were transfected with 50 ng of 4XCSL-Luc (A), 50 ng of Hes-1-Luc (B), 50 ng of Hes-5-Luc and RMHes5-Luc (C), and 50 ng of β-galactosidase, along with 100 ng of Notch1-IC and 50 ng (+), 100 ng (++) and 150 ng (+++) of Wnt5a, as indicated. (*) P < 0.05 versus Notch1-IC. (D) HEK293 cells were transfected with 50 ng of 4XCSL-Luc and 50 ng of β-galactosidase, along with 100 ng of Notch1-IC. Before harvest, HEK293 cells were treated with 25 ng/ml, 50 ng/ml and 100 ng/ml of Wnt5a protein for 8 hours, as indicated. (**) P < 0.05 (C) and P < 0.01 (D) versus Notch1-IC. (E) HEK293 cells were transfected for 48 hours with 50 ng of 4XCSL-Luc, 100 ng of Notch1-IC, 150 ng of Wnt5a and 50 ng of β-galactosidase. Before harvest, HEK293 cells were treated with KN-93 (5 μM) for 6 hours, the cells were lysed, and the luciferase activity was determined. (*) P < 0.05 versus Notch1-IC; (**) P < 0.05 versus Notch1-IC + Wnt5a. (F) HEK293 cells were transfected with 50 ng of 4XCSL-Luc and β-galactosidase, along with 100 ng of Notch1-IC, 150 ng of Wnt5a and 100 ng of CaMKII-DN, as indicated. 48 hours after transfection, the cells were lysed, and the luciferase activity was determined. DN, Dominant Negative. (*) P < 0.05 versus Notch1-IC; (**) P < 0.05 versus Notch1-IC + Wnt5a. (G) HEK293 cells were transfected with 0.5 μg of Myc-Notch1-IC, 0.5 μg of siCaMKII as indicated. 48 hours after transfection, the cell lysates were immunoblotted with anti-Hes1 and anti-Hes5 antibodies. Probing with an antibody to β-actin was used as a loading control.

FIGURE 2. CaMKII activates Notch1 transcriptional activity. (A-C) HEK293 cells were transfected with 50 ng of 4XCSL-Luc (A), 50 ng of Hes-1-Luc (B), 50 ng of Hes-5-Luc and RMHes5-Luc (C), and 50 ng of β-galactosidase, along with 100 ng of Notch1-IC and 50 ng (+), 100 ng (++) and 150 ng (+++) of CaMKII, as indicated. (D-F) HEK293 cells were transfected with 50 ng of 4XCSL-Luc and β-galactosidase, along with 100 ng of Notch1-IC, 150 ng of Wnt5a and 100 ng of CaMKII-DN, as indicated. 48 hours after transfection, the cells were lysed, and the luciferase activity was determined. (*) P < 0.01 (A, C) or P < 0.05 (B) versus Notch. (D-F) (*) P < 0.05 versus control.

FIGURE 3. CaMKII activates Notch1 transcriptional activity through RBP-Jk dependent and Notch1-IC independent manner. (A-C) PS1/2+/+ MEF cells were transfected with 50 ng of 4XCSL-Luc (A), 50 ng of Hes-1-Luc (B), 50 ng of Hes-5-Luc and RMHes5-Luc (C), and 50 ng of β-galactosidase, along with 100 ng of Notch1-IC and 50 ng (+), 100 ng (++) and 150 ng (+++) of CaMKII, as indicated. (D-F) PS1/2+ MEF cells were transfected with 50 ng of 4XCSL-Luc (D), 50 ng of Hes-1-Luc (E), 50 ng of Hes-5-Luc (F), and 50 ng of β-galactosidase, along with 50 ng (+), 100 ng (++) and 150 ng (+++) of CaMKII, as indicated. 48 hours after transfection, the cells were lysed, and the luciferase activity was determined. (*) P < 0.01 (A, C) or P < 0.05 (B) versus Notch. (D-F) (*) P < 0.05 versus control. (G) PS1/2+ MEF cells were transfected with 50 ng of 4XCSL-Luc and 50 ng of β-
galactosidase, along with 100 ng of CaMKII and 100 ng of RBP-Jk-DN as indicated. (*) P < 0.01 versus CaMKII. (H) PS1/2−/− MEF cells were transfected with 50 ng of 4XCSL-Luc and 50 ng of β-galactosidase, along with 100 ng of CaMKII and 100 ng of CaMKII-DN with 4XCSL-Luc. (*) P < 0.01 versus CaMKII. (I) PS1/2−/− cells were transfected with 50 ng of 4XCSL-Luc and 50 ng of β-galactosidase, along with 100 ng of CaMKII. 42 hours after transfection, HEK293 cells were treated with KN-93 (5μM) for 6 hours and the luciferase activity was determined. (*) P < 0.05 versus CaMKII. (A-I) 48 hours after transfection, the cells were lysed, and the luciferase activity was determined. DN, Dominant Negative. (J) Immunoblots of PS1/2+/+ and PS1/2−/− MEF cells treated with 1 μM of DAPT for 6 hours. The cell lysates were immunoblotted with anti-Notch1-IC antibodies. Probing with an antibody to β-actin was used as a loading control.

FIGURE 4. CaMKII modulates the binding of RBP-Jk to Notch1-IC or SMRT. (A) HEK293 cells were transfected for 48 hours with the indicated combinations of 0.5 μg of Myc-Notch1-IC, 0.5 μg of Flag-RBP-Jk, and 0.5 μg of Wnt5a. Cell lysates were subjected to immunoprecipitation with anti-Flag antibody and the immunoprecipitates were immunoblotted with anti-Myc antibody. The cell lysates were also subjected to immunoblot analysis with anti-Myc and anti-Flag antibodies, respectively. (B) HEK293 cells were transfected for 42 hours with the indicated combinations of 0.5 μg of Myc-Notch1-IC, 0.5 μg of Flag-RBP-Jk, and 0.5 μg of Wnt5a. Before harvest, cells were treated with KN-93 (4μM) for 6 hours. The cell lysates were subjected to immunoprecipitation with anti-Myc antibody and the immunoprecipitates were immunoblotted with anti-Flag antibody. The cell lysates were also subjected to immunoblot analysis with anti-Flag and anti-Myc antibodies, respectively. (C) HEK293 cells were transfected for 42 hours with 0.5 μg of Wnt5a. Before harvest, HEK293 cells were treated with KN-93 (5μM) for 6 hours. Cell lysates were subjected to immunoprecipitation with anti-RBP-Jk antibody and the immunoprecipitates were immunoblotted with anti-Notch1-IC antibody. The cell lysates were also subjected to immunoblot analysis with anti-Myc and anti-Flag antibodies, respectively. (D) HEK293 cells were transfected for 48 hours with the indicated combinations of 0.5 μg of Myc-Notch1-IC, 0.5 μg of Flag-RBP-Jk, 0.5 μg of HA-CaMKII and 0.5 μg of HA-CaMKII-DN. Cell lysates were subjected to immunoprecipitation with anti-Flag antibody and the immunoprecipitates were immunoblotted with anti-Myc antibody. The cell lysates were also subjected to immunoblot analysis with anti-Myc anti-HA and anti-Flag antibodies, respectively. (E) HEK293 cells were transfected for 48 hours with the indicated combinations of 0.5 μg of Flag-SMRT, 0.5 μg of Myc-RBP-Jk, and 0.5 μg of Wnt5a. Cell lysates were subjected to immunoprecipitation with anti-Flag antibody and the immunoprecipitates were immunoblotted with anti-Myc antibody. The cell lysates were also subjected to immunoblot analysis with anti-Myc-Notch1-IC antibody and the immunoprecipitates were immunoblotted with anti-SMRT antibody. The cell lysates were also subjected to immunoblot analysis with anti-SMRT and anti-RBP-Jk antibodies, respectively. (F) HEK293 cells were transfected for 42 hours with 0.5 μg of Wnt5a. Before harvest, HEK293 cells were treated with KN-93 (5μM) for 6 hours. Cell lysates were subjected to immunoprecipitation with anti-RBP-Jk antibody and the immunoprecipitates were immunoblotted with anti-SMRT antibody. The cell lysates were also subjected to immunoblot analysis with anti-SMRT and anti-RBP-Jk antibodies, respectively. (G) HEK293 cells were transfected for 48 hours with the indicated combinations of 0.5 μg of Flag-SMRT, 0.5 μg of Myc-RBP-Jk, 0.5 μg of HA-CaMKII and 0.5 μg of HA-CaMKII-DN. Cell lysates were subjected to immunoprecipitation with anti-Flag antibody and the immunoprecipitates were immunoblotted with anti-Myc antibody. The cell lysates were also subjected to immunoblot analysis with anti-Flag anti-HA and anti-Myc antibodies, respectively. (A, B, D, E, G) Cell lysates were subjected to immunoprecipitation with anti-Myc, anti-Flag and anti-HA antibody and the immunoprecipitates were immunoblotted with anti-Myc, anti-Flag and anti-HA antibody, respectively. DN, Dominant Negative. IgG16, immunoglobulin heavy chain. IgG1, immunoglobulin light chain. IB, immunoblot.

FIGURE 5. CaMKII down-regulates the steady state level of SMRT protein. (A) HEK293 cells were transfected with 0.5 μg of Flag–SMRT, 0.5 μg of HA-CaMKII and 0.5 μg of HA-CaMKII-DN as indicated. 48 hours after transfection, the cell lysates were immunoblotted with anti-Flag and anti-HA antibodies. (B) HEK293 cells were transfected with 0.5 μg of Flag–SMRT and 0.5 μg of HA-CaMKII as indicated. 42 hours after transfection, the cells were treated with various concentrations (0, 1, 5μM) of MG132 for 6 hours. 48 hours after transfection, the cell lysates were immunoblotted with anti-Flag and anti-HA antibodies. (C) HEK293 cells were transfected with 0.5 μg of Flag-SMRT, 0.5 μg of HA-CaMKII and 0.5 μg of HA-CaMKII-DN as indicated. 42 hours after transfection, the cells were then treated with 0.1 mM of cycloheximide for the indicated times. (D) HEK293 cell were treated with the indicated amounts of purified Wnt5a protein
CaMKII regulates Notch1 signaling.

for 8 hours. The cell lysates were also subjected to immunoblotting analysis with anti-SMRT. Probing with an antibody to β-actin was used as a loading control. (E) HEK293 cells were transfected with 0.5 μg of Flag-SMRT, 0.5 μg of HA-Ub and 0.5 μg of GFP-CaMKII as indicated. 42 hours after transfection, HEK293 cells were treated with KN-93 (5μM) for 6 hours. Cell lysates were subjected to immunoprecipitation with anti-Flag antibody and the immunoprecipitates were immunoblotted with anti-HA antibody. The cell lysates were also subjected to immunoblot analysis with anti-HA, anti-Flag and anti-GFP antibodies, respectively. Probing with an antibody to β-actin was used as a loading control. DN, Dominant Negative. IB, immunoblot.

FIGURE 6. Physical interaction of CaMKII with SMRT in intact cells. (A) Recombinant GST or GST–SMRT was immobilized onto GSH-agarose. HEK293 cells were transfected with 0.5 μg of HA-CaMKII. 48 hours after transfection, the cell lysates were subjected to GST pull-down experiments with immobilized GST or GST-SMRT. Proteins bound to GST or GST–Notch1-IC were analyzed via immunoblotting with an anti-HA antibody. (B) HEK293 cells were transfected with 0.5 μg of Flag–SMRT and 0.5 μg of HA-CaMKII as indicated. 42 hours after transfection, the cells were treated with MG132 (5μM) for 6 hours. The cell lysates were subjected to immunoprecipitation with an anti-HA antibody. The immunoprecipitates were then immunoblotted with an anti-Flag antibody. Cell lysates were also immunoblotted with anti-Flag and anti-HA antibodies. (C) HEK293 cells were lysed and subjected to immunoprecipitation with immunoglobulin G (IgG) and anti-SMRT antibodies as indicated. The immunoprecipitates were immunoblotted with an anti-CaMKII antibody. Cell lysates were also immunoblotted with anti-SMRT and anti-CaMKII. (D) Recombinant GST or GST–SMRT deletion proteins (GST-SMRT-SD1, GST-SMRT-SD2, GST-SMRT-RID1 and GST-SMRT-RID2) were immobilized onto GSH-agarose. HEK293 cells were transfected with 0.5 μg of HA-CaMKII. 48 hours after transfection, the cell lysates were subjected to GST pull-down experiments with immobilized GST or GST–SMRT. Proteins bound to GST or GST–SMRT were analyzed via immunoblotting with an anti-HA antibody. CBB, Coomassie Brilliant Blue staining. IgG, immunoglobulin heavy chain. IB, immunoblot.

FIGURE 7. The phosphorylation of SMRT by CaMKII is critical for degradation of SMRT protein. (A) HEK293 cells were transfected with 0.5 μg of HA-CaMKII or 0.5 μg of HA-CaMKII-DN as indicated. 48 hours after transfection, the cell lysates were subjected to immunoprecipitation with an anti-HA antibody, and the resultant precipitates were evaluated for phosphorylation of SMRT by an immune complex kinase assay using GST–SMRT. (B) HEK293 cells were transfected with 0.5 μg of HA-CaMKII or an empty vector as indicated. 48 hours after transfection, the cell lysates were subjected to immunoprecipitation with an anti-HA antibody and the resulting precipitates were examined for phosphorylation of SMRT using an immune complex kinase assay with purified recombinant GST–SMRT-SD1, GST–SMRT-SD2, GST–SMRT-RID1 and GST–SMRT-RID2 proteins. (C) HEK293 cells were transfected with 0.5 μg of HA-CaMKII. 48 hours after transfection, the cell lysates were subjected to immunoprecipitation with an anti-HA antibody, and the resultant precipitates were evaluated for phosphorylation of SMRT using an immune complex kinase assay with GST–SMRT, GST–SMRT (S1407A) and GST–SMRT (S1434A). (D) HEK293 cells were transfected with 0.5 μg of Flag–SMRT, 0.5 μg of Flag-SMRT (S1407A), 0.5 μg of Flag-SMRT (S1434A) and 0.5 μg of HA-CaMKII as indicated. 48 hours after transfection, the cell lysates were immunoblotted with an anti-Flag antibody. Cell lysates were also immunoblotted with anti-Flag and anti-HA antibodies. (E) HEK293 cells were transfected with 0.5 μg of Flag–SMRT, 0.5 μg of Flag-SMRT (S1407A), 0.5 μg of Myc-RBP-Jk and 0.5 μg of HA-CaMKII as indicated. 48 hours after transfection, the cell lysates were subjected to immunoprecipitation using an anti-Flag antibody. The immunoprecipitates were then immunoblotted with an anti-Myc antibody. Cell lysates were also immunoblotted with anti-Flag, anti-Myc and anti-HA antibodies. (F) HEK293 cells were transfected with 50ng of 4XCSL-Luc, 50ng of SMRT, 50ng of SMRT (S1407A), and 50ng of β-galactosidase, along with 100ng of CaMKII, as indicated. 48 hours after transfection, the cells were lysed, and the luciferase activity was determined. The data was normalized with β-galactosidase. These results represent the means ± average deviation of three independent experiments. R.L.U means relative luciferase units. (G) HEK293 cells were transfected with empty vector, 0.5 μg of HA-CaMKII, 0.5 μg of Flag-SMRT, and 0.5 μg of Flag-SMRT (S1407A) as indicated. 42 hours after transfection, the cells were treated with MG132 (5μM) for 6 hours. 48 hours after transfection, the cells were fixed and CaMKII and SMRT were stained with Alexa Flou 488 (green) and Alexa Flou 555 (red) and examined by confocal microscopy. The DNA dye ToPro3.
was used to visualize nuclei. For each experiment, at least 300 cells were examined, and the images shown here represent the typical staining pattern for a majority of the cells. The mean nuclear localization intensity per cell was analyzed by imaging cytometric analysis. White Scale bar, 25 \( \mu \)m. (H) HEK293 cells were transfected with empty vector, 0.5 \( \mu \)g of HA-CaMKII, 0.5 \( \mu \)g of Flag-SMRT, and 0.5 \( \mu \)g of Flag-SMRT (S1407A) as indicated. 48 hours after transfection, the cells were fractionated into cytosolic (Cyto), and nuclear (Nuc) fractions. Cell lysates were also immunoblotted with anti-Flag and anti-HA antibodies. LaminB and \( \beta \)-actin were used as nuclear and cytoplasmic fraction marker. DN, Dominant Negative. IgG\textsubscript{H}, immunoglobulin heavy chain. IB, immunoblot.

**FIGURE 8.** Schematic diagram of CaMKII regulates Notch1 signaling through proteasomal degradation of SMRT. Frizzled (Fz), and receptor tyrosine kinase-like orphan receptor1/2 (Ror1/2) mediates Wnt5a/Ca\textsuperscript{2+} signal has not yet been elucidated, however, the formation of a trimeric complex with Fz and Ror1/2 and PDZ domain mediated binding with dishevelled (Dvl) could be important. In this study, we evaluated that the mechanism by which Wnt up-regulates Notch1 target genes. We demonstrated that Wnt5a induced CaMKII enhances Notch transcriptional activity. CaMKII physically interacts with SMRT and disrupts the SMRT-RBP-Jk transcription complex through degradation of SMRT. Dissociation of SMRT from RBP-Jk induced by CaMKII activates Hes1 and Hes5 transcriptional activity. Furthermore, CaMKII can phosphorylate SMRT and activate the Notch1 signaling pathway. The kinase activity of CaMKII is necessary for the up-regulation of the transcription activity of Notch1 target genes. The results of this study suggest that CaMKII is a crucial regulator for signal cross-talk between Wnt5a and Notch signaling via downregulation of the SMRT protein level.
Figure 1

A 4XCSL-Luc

B Hes1-Luc

C Hes5-Luc

D 4XCSL-Luc

E 4XCSL-Luc

F 4XCSL-Luc

G Myc-Notch1-IC : - * * * * Wnt5a : - - - - siCaMKII : - - - - KN-93 : - - - - CaMKII-DN : - - - - IB: Hes1 25 IB: Hes5 20 IB: β-actin 43

β-actin

Hes5

Hes1
Figure 2

A. 4XCSL-Luc

B. Hes1-Luc

C. Hes5-Luc

D. 4XCSL-Luc

E. Hes1-Luc

F. Hes5-Luc
Figure 3

A  

B  

C  

D  

E  

F  

G  

H  

I  

J
Figure 5

A

Flag-SMRT : - + + +
HA-CaMKII : - - + -
HA-CaMKII-DN : - - - +
IB: Flag 180
IB: HA 55

B

Flag-SMRT : - + + + +
HA-CaMKII : - - 0 1 5 (µM)
MG132 : - - + + +
IB: Flag 180
IB: HA 55

C

Cycloheximide : 0 0.5 1 2 4 6 (h)
Flag-SMRT : 180
Flag-SMRT + HA-CaMKII : 180
Flag-SMRT + HA-CaMKII-DN : 180
β-actin : 43

D

Wnt5a : 0 50 100 150 ng/ml
IB: SMRT 180
IB: β-actin 43

E

Flag-SMRT : + + +
HA-ub : + + +
GFP-CaMKII : - + +
KN93 : - - -
IB: HA 180
IB: HA 55
IP: Flag
**Figure 6**

**A**
HA-CaMKII:

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**B**

MG132:

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| IB: Flag   | 180        |
| IP: HA     | Flag-SMRT  |
| Cell Lysates |          |
| HA-CaMKII  |           |

**C**

IP: IgG SMRT

| IB: CaMKII | 55 |
| Cell Lysates | 55 |

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**D**

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Figure 7

A. Kinase assay

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<th>HA-CaMKII</th>
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B. GST-SMRT

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C. HA-CaMKII

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D. HA-CaMKII

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E. Myc-RBP-Jk

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F. 4XCSL-Luc

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G. CaMKII, SMRT, TOPRO3, Merge

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Figure 8

- **CaMKII**
- **Wnt5a**
- **Fz receptor**
- **Ror1/2**
- **CaMKII**
- **SMRT**
- **T**
- **RBP-Jk**
- **Proteasomal degradation**
- **CaMKII**
- **CaMKII**
- **CaMKII**
- **SMRT**
- **SMRT**
- **SMRT**
- **SMRT**
- **SMRT**
- **SMRT**
- **SMRT**
- **MAML**
- **MAML**
- **RBP-Jk**
- **RBP-Jk**
- **RBP-Jk**
- **Repressor complex**
- **Activator complex**
- **Notch1**
- **Notch1-IC**
- **S3 cleavage**
- **Cytoplasm**
- **Nucleus**
- **Target Gene Transcription ON (e.g., Hes1, Hes5, Hey1... etc)**

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Wnt5a controls Notch1 signaling through CaMKII mediated degradation of the SMRT corepressor protein

Eun-Jung Ann, Hwa-Young Kim, Mi-Sun Seo, Jung-Soon Mo, Mi-Yeon Kim, Ji-Hye Yoon and Hee-Sae Park

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