

Induction of Δ Np63 by the Newly Identified Keratinocyte-Specific Transforming Growth Factor β Signaling Pathway with Smad2 and I κ B Kinase α in Squamous Cell Carcinoma^{1,2}

Nahoko Fukunishi^{*}, Iyoko Katoh[†],
Yoshiya Tomimori^{‡,3}, Keiichi Tsukinoki[§],
Ryu-ichiro Hata[§], Atsuhito Nakao[¶], Yoji Ikawa^{‡,4}
and Shun-ichi Kurata^{*}

^{*}Redox Response Cell Biology, Medical Research Institute, Tokyo Medical and Dental University, Tokyo, Japan;

[†]Department of Microbiology, Interdisciplinary Graduate School of Medicine and Engineering, University of Yamanashi, Yamanashi, Japan; [‡]Ikawa Laboratory RIKEN, Institute of Physical and Chemical Research, Saitama, Japan; [§]High-Tech Research Center, Kanagawa Dental College, Yokosuka, Japan; [¶]Department of Immunology, Interdisciplinary Graduate School of Medicine and Engineering, University of Yamanashi, Yamanashi, Japan

Abstract

The expression of *p63* (TP63/p51) occurs in the basal cells of stratified epithelia and is strongly enhanced at the early stages of squamous cell carcinomas (SCCs) of the head and neck, skin, cervix, and others. We analyzed a promoter/enhancer region (2k Δ N) that drives the predominant expression of Δ Np63 for sensitivity to Smad signaling pathways. Reporter assays in HepG2 cells showed a moderate activation of 2k Δ N by Smad2 and I κ B kinase α (IKK α), partners of the newly identified keratinocyte-specific transforming growth factor β (TGF- β) signaling, but not by other Smad molecules. In A431 cells, 2k Δ N was activated by Smad2 and IKK α , for which a Smad binding element (SMD2) at –204 was essential. Binding of Smad2 to the chromosomal SMD2 site was detectable. The association of Smad2 with IKK α was evident in the nucleus of A431, accounting for the enhancement of Δ Np63 expression by TGF- β . Moreover, both Δ Np63 and IKK α were necessary to maintain the noninvasive phenotype of this cell line. FaDu, an invasive, Smad4-deficient SCC, also allowed 2k Δ N transactivation by transfected Smad2 in the presence of endogenous IKK α . Reflecting the lack of chromosomal SMD2-Smad2 association and the absence of nuclear IKK α , however, endogenous Δ Np63 was not controlled by TGF- β or IKK α in FaDu. SCC tissue arrays showed nuclear accumulation of IKK α and *p63* intensification in well-differentiated noninvasive lesions. This study indicates that *p63* is a target gene of the proposed keratinocyte-specific TGF- β signal pathway for suppression of the malignant conversion of SCC.

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Address all correspondence to: Shun-ichi Kurata, Medical Research Institute, Tokyo Medical and Dental University, 1-5-45 Bunkyo-ku, Yushima, Tokyo 113-8510, Japan. E-mail: kushbgen@mri.tmd.ac.jp; or Iyoko Katoh, Department of Microbiology, Interdisciplinary Graduate School of Medicine and Engineering, University of Yamanashi, Yamanashi 409-3898, Japan. E-mail: iyoko@yamanashi.ac.jp

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³Present address: Nagahama Institute for Biochemical Science, Oriental Yeast Co., Ltd., 50 Kanoho, Nagahama-hi, Shiga 526-0804, Japan.

⁴Present address: Division of Cellular & Molecular Toxicology, Biological Safety Research Center, National Institute of Health Sciences, Tokyo, Japan.

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Introduction

Unlike *p53*, which is ubiquitously expressed to exert the tumor-suppressing function, *p63* (*TP63*, *p51*) [1,2] is required for the development of stratified epithelia including the skin and oral tissues [3,4]. A high level of expression of *p63* occurs not only in keratinocyte stem cells of normal stratified epithelia [5,6] but also in squamous cell carcinomas (SCCs) of head and neck, skin, and cervix as well as in carcinomas of urothelia and others [7–9]. After the intensification at the lower-grade carcinomas, however, *p63* expression diminishes during the malignant progression [10–13]. Although various genes induced by *p63* have been reported [14–19], it remains obscure how *p63* gene expression is enhanced at the limited stages of the specific lineages in tissue development and cancer progression.

The human *p63* locus has two separate transcriptional initiation sites to produce transactivator protein TAp63 and N-terminally truncated protein Δ Np63. Because Δ Np63 isoform expression is significantly more predominant than TAp63 in normal keratinocytes and SCCs [2,8,15], the Δ Np63 promoter immediately upstream of exon 3' is thought to control the level of Δ Np63 and overall functions of *p63*. An earlier report showed that Δ Np63 of zebrafish, the only *p63* transcript of this species, is induced by bone morphogenetic protein 2 (BMP2) through Smad binding elements (SBEs) in the promoter/enhancer region [20].

In addition to the canonical Co-Smad/R-Smad signaling pathways of TGF- β and BMP, varied modes of cross-talk between the Smad systems and other cellular signaling mechanisms have been studied [21,22]. Interestingly, a keratinocyte-specific TGF- β signaling pathway has been recently identified, in which I κ B kinase α (IKK α) instead of Smad4 acts like Co-Smad to interact with Smad2/3 (R-Smad) for transcriptional activation of the target genes [23,24]. Apart from the protein kinase activity required for the NF- κ B pathway activation in the cytosol, IKK α needs to translocate to the nucleus for this function. As earlier studies showed, IKK α -deficient mice manifest severe defects in the skin and limbs because of the blockage of keratinocyte differentiation [25,26].

The Smad2/3-IKK α pathway is activated in noninvasive well-differentiated (grade 1) SCCs but seems switched off on the malignant conversion into invasive (grade 3) SCCs [27]. These processes are observed in conjunction with nuclear translocation of IKK α in grade 1 SCCs and its cytosolic sequestration in grade 3 SCC. Intriguingly, a line of evidence indicated that both TAp63 and Δ Np63 activate transcription of the IKK α gene (*CHUK*, also termed *IKBKA*, *IKK-alpha*, *IKKI*, etc) in humans [18,28,29].

Because the *p63* expression pattern is well conserved over a wide range of species [20,30], we investigated how *p63* is influenced by the human Smad signaling systems. Our results indicate that the Δ Np63 promoter is activated by the keratinocyte-specific TGF- β signaling mechanism with Smad2 and IKK α . Functional interactions between IKK α and *p63* in SCC progression are discussed.

Materials and Methods

Cell Culture

HepG2 and A431 cells were obtained from the Japan Health Sciences Foundation. C2C12 and FaDu were from the American Type Culture Collection. HepG2, A431, C2C12 cells were propagated in Dulbecco modified minimum essential medium (D-MEM) with 10% fetal bovine serum (FBS). FaDu cells were maintained in MEM supple-

mented with sodium pyruvate, nonessential amino acids, glutamine, and 10% FBS.

Luciferase Reporter Assay

Chromosomal DNA obtained from a healthy donor (Y.T.) was amplified by polymerase chain reaction (PCR) to obtain 2k Δ N (-1848 to +152; Figure 1) and 2kTA (-1911 to +89, in which +1 was tentatively placed at the reported 5' end of TAp63 α complementary DNA [cDNA], NM_003722). The DNA segments were cloned into pGL3basic (Promega, Madison, WI) at the *Xho* I site and sequenced. *In vitro* mutagenesis was carried out with the GeneTailor Site-directed Mutagenesis System (Invitrogen, Carlsbad, CA). The Smad1 expression vector pcDNA3-Flag(N)-Smad1 and the Id1 promoter linked to the luciferase gene (BRE-luc) were from Dr Kohei Miyazono, University of Tokyo. Smad2 was described [31]. The Smad3, Smad4, and Smad5 cDNA were isolated from a HeLa cell cDNA library (RIKEN, Tsukuba, Japan), cloned into pRc/CMV (Invitrogen), and sequenced. An IKK α expression vector (CHUK, NM_001278.3), pCMV6-XL5-CHUK was purchased from OriGene Technologies, Rockville, MD. Plasmid DNA transfection was carried out with Attractene (Qiagen, Hilden, Germany) for HepG2, C2C12, and FaDu and with Superfect (Qiagen) for A431. Eighteen hours after transfection, cells were starved in 0.1% serum for 6 hours and were further incubated for 24 hours with or without TGF- β 1 (R&D Systems, Minneapolis, MN) at 10 ng/ml, a concentration relevant to experiments with epithelial cells [32]. We used the Luciferase Assay Kit (Promega) for FaDu and A431 and the Steady-Glo Luciferase Assay System (Promega) for HepG2 and C2C12 cells.

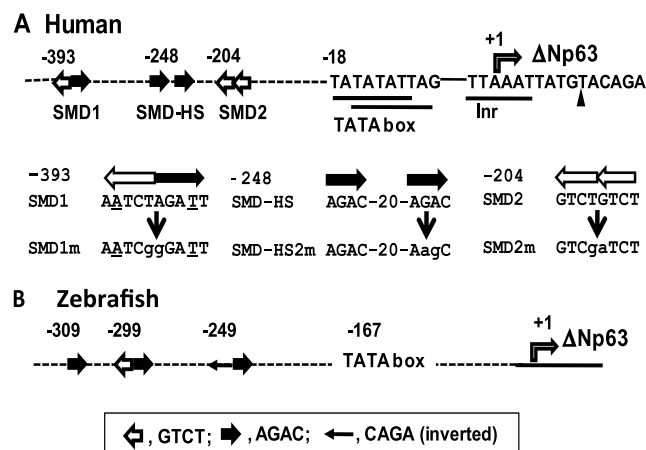


Figure 1. SBE-like sequences found in the promoter/enhancer region of Δ Np63. (A) The proximal promoter region of human Δ Np63 is schematically shown. The 5' end of the reported Δ Np63 α cDNA data (NM_001114980) is marked by a filled triangle. Nucleotide stretches matching the consensus sequences of initiators, 5'-(C/T)(C/T)AN(A/T)(C/T)(C/T)-3', and TATA box, 5'-TATA(T/A)A(T/A)(A/G)-3', are underlined. Provisional initiation site (position +1) of Δ Np63 is marked by a rectangular arrow. Filled and open arrows represent the SBE motif 5'-AGAC-3', and its complementary sequence 5'-GTCT-3', respectively. Relative positions are shown for SMD1 (a complete palindrome including 5'-TCT-AGA-3'), SMD2 (a tandem repeat of 5'-GTCT-3'), and SMD-HS (a repeat of 5'-AGAC-3' interrupted by 20 nucleotides). Mutated nucleotides in SMD1m, SMD2m and SMD-HS2m are in lower case. (B) The zebrafish (*Danio rerio*) Δ Np63 promoter/enhancer region with functional SBE sites are shown as described by Bakkers et al. [20].

Gene Silencing

SiLentfect (Bio-Rad, Hercules, CA) was used for small interfering RNA (siRNA) transfection. FaDu cells were transfected in MEM with 10% FBS. A431 cells were incubated with Opti-MEM serum-free medium (Invitrogen) during the siRNA transfection (18 hours) and refed with D-MEM with 10% FBS. A pair of siRNA duplex targeting *p63* (*p63*si) was designed by us [15] and synthesized by Takara Bio (Otsu, Japan). Control siRNA (Csi) and IKK α -targeting siRNA (IKK α si) were purchased from Ambion (Austin, TX; AM4611) and Dharmacon (Lafayette, CO; M-003473S), respectively.

Western Blot Analysis

Whole-cell lysates were subjected to Western blot analysis as described [14]. We purchased anti-*p63* (4A4, mouse), anti-*IKK α* (sc-7218, rabbit), anti-*Smad2* antibody (sc-101153, mouse), and anti-*Smad4* (sc-7966, mouse) antibodies from Santa Cruz Biotechnology (Santa Cruz, CA). A rabbit anti-Flag antibody (PM020; MBL International, Woburn, MA) was used for Flag-*Smad2* immunoprecipitation. Anti-HSP90 (ab1429, mouse) and anti-*p84* (ab487, mouse) antibodies were from Abcam (Cambridge, UK). An anti-phospho-*Smad2* (Ser465/467, #3101) antibody was from Cell Signaling Technology (Beverly, MA). Alkaline phosphatase (AP)-conjugated secondary antibodies (Cell Signaling Technology) were applied. Size markers (Biotinylated SDS-PAGE Standards; Bio-Rad) were detected with an AP-conjugated antibiotin immunoglobulin G (IgG; Cell Signaling Technology). Chemiluminescence (CDP-Star Reagent; New England Biolabs, Ipswich, MA) was captured with a camera luminometer (ECL Mini-camera; Amersham Pharmacia Biotech, Uppsala, Sweden), in which luminescence images appear white on the black background. Scanned digital images were processed by Photoshop (Adobe Systems, San Jose, CA) without a modification of the band intensity. Proteins were semiquantified with an imaging program, ImageJ (National Institutes of Health, Bethesda, MD).

Chromatin Immunoprecipitation

We used chromatin immunoprecipitation (ChIP) IT Expression Enzymatic, ChIP IT Control Kit (Active Motif, Carlsbad, CA), and an anti-*Smad2* antibody (51-1300; Zymed Laboratories, South San Francisco, CA). A431 and FaDu cells required 100 strokes of homogenization in Dounce homogenizer (Wheaton, DuPage County, IL) for efficient nuclear release. A SMD2-containing segment was amplified by PCR with primers 5'-ATCATTCTTGAAACCCCAAATCTA-3' (-411 to -388) and 5'-CCAGGAGACAGACAGGTAATAACTT-3' (-215 to -192). A segment upstream of *TAp63* was also amplified with 5'-TACATGTGCATGTGTTTGAGGTAG-3' and 5'-TAGAGCTGCT-TAGGGAAGTTTTGT-3'.

Reverse Transcription-Polymerase Chain Reaction

RNA isolated with RNawiz (Ambion) was further cleaned up using the RNeasy MinElute Cleanup Kit (Qiagen). Reverse transcription-polymerase chain reaction (RT-PCR; 30 μ l) contained the SuperScript One-Step RT-PCR reagent with Platinum Taq (Invitrogen), RNA (0.6 μ g), and the primers described previously [14,15].

Immunofluorescence Staining

Cells were cultured on the Lab-Tek chamber slides (Nunc, Roskilde, Denmark). Cancer tissues were obtained from patients with informed consent at Kanagawa Dental College Hospital (Yokosuka, Japan). SCC

tissue array slides, BC34011, and OR802 were from US Biomax (Rockville, MD). Formalin-fixed cells, tissue arrays, and sections were antigen-retrieved and probed with anti-*p63* (4A4) and anti-*IKK α* (sc-7218) antibodies in combination with secondary antibodies conjugated with Alexa Fluor 594 or 488 (Molecular Probes, Eugene, OR). Fluorescence image was acquired using a Radiance 2000 laser scanning confocal microscope (Bio-Rad) with LaserSharp 2000 imaging software (Carl Zeiss, Oberkochen, Germany) or fluorescence microscope (BZ-9000; Keyence, Osaka, Japan).

Matrigel Invasion Assay

After rehydration, the Matrigel culture inserts (BD Biosciences, San Jose, CA) were placed in the wells (24-well culture plates) containing the medium (0.5 ml) with 10% FBS. Cell suspension (5×10^4) in 0.5 ml of the medium with 0.1% bovine serum albumin was added to the chamber and incubated for 18 hours.

Results

Lack of BMP Sensitivity in the *TAp63* and Δ *Np63* Promoter/Enhancer Regions

The TA and Δ N promoter/enhancer regions of 2 kbp (2kTA and 2k Δ N, respectively) were linked 5' to a firefly luciferase gene (*luc*) (Figure 1A). We first tested 2kTA and 2k Δ N for sensitivity to BMP2 in C2C12 cells (Figure W1A). Activation of these promoters by the BMP-related Smad proteins was also assessed in HepG2 cells (Figure W1B). In comparison with the human *Id1* promoter [33], neither one showed a positive response in either experiment.

We next tested to see whether 2kTA and 2k Δ N can be activated by the canonical TGF- β signal by Smad2/3 paired with Smad4 in HepG2 cells (Figure W2A). Compared with the plasminogen activator inhibitor 1 (PAI-1) gene promoter [31], which was strongly activated by Smad3 transfection and the following TGF- β 1 stimulation (to nearly 40-fold), 2k Δ N showed a minimum detectable response to Smad2 (an approximately 2- to 3-fold activation), without an obvious response to TGF- β 1. Smad4 was not a good partner of Smad2 in the 2k Δ N activation.

Activation of the Δ *Np63* Promoter by Smad2 and IKK α

We considered the possibility of 2k Δ N activation by the newly identified TGF- β signaling mechanism. A431 cells derived from an epidermoid SCC persistently expressed endogenous *p63* at a high level [11]. In this cell line, the 2k Δ N-driven luciferase expression was significantly (20-fold) increased by Smad2 transfection (Figure 2A). TGF- β 1 (10 ng/ml) stimulation in the serum-starved (0.1% FBS) medium further enhanced the effect of Smad2 (to ~40-fold). Plasmid-mediated IKK α expression elicited a minor response of 2k Δ N (two-fold). Combined transfection of Smad2 and IKK α (2:2), followed by incubation with TGF- β 1, also caused 2k Δ N activation by a magnitude of ~20. In contrast, no enhancement was detected by Smad3-only and Smad4-only transfection or by the combination of Smad2-Smad4 or Smad3-Smad4. Luciferase expression by 2kTA remained unchanged (Figure 2C). The PAI-1 promoter also responded to TGF- β 1 (seven-fold activation) when Smad3 was transfected (Figure 2D). In contrast to the positive effects of Smad4 on PAI-1 in HepG2 cells (Figure W2A), Smad4 failed to activate this promoter in A431 cells, and even elicited a suppressive effect on Smad3.

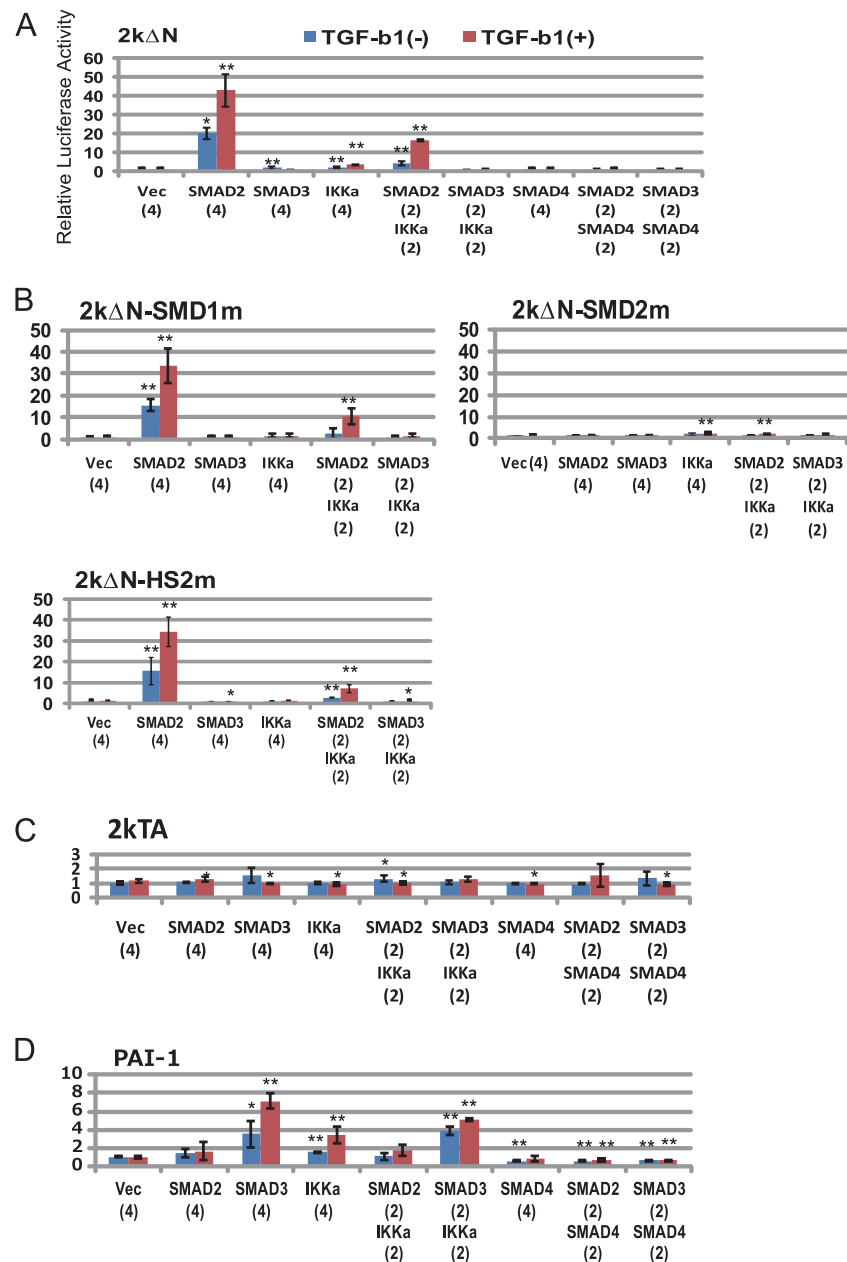


Figure 2. Activation of the Δ Np63 promoter by Smad2 and IKK α , and by stimulation with TGF- β in A431 cells. Results of the luciferase assay with 2k Δ N (A), 2k Δ N mutants (B), 2kTA (C) and the PAI-1 promoter (D) are shown. Cells were incubated with (+) or without (-) TGF- β 1 (10 ng/ml [32]) for 24 hours after serum starvation for 6 hours. Data (mean \pm SD; $n \geq 3$) are presented as relative luciferase activities in comparison with the control activity (1.0) produced by 2k Δ N-luc transfected with empty expression vector pRcCMV (Vec). Statistical significance: ** $P < .01$, * $P < .05$, versus control experiment with Vec. Indicated in parentheses are amounts of the cytomegalovirus promoter-driven gene expression vectors relative to 2k Δ N-luc (=1). IKK α indicates IKK α .

Within the 2k Δ N region, we detected multiple SBE-related sequences referred to as SMD1 (starting at position -393), SMDHS-1/2 (-248), and SMD2 (-204) (Figure 1A). A double-nucleotide mutation was introduced to each site. The mutation of SMD2 (5'-GTCTGTCT-3') to SMD2m (5'-GTCgaTCT-3') was sufficient to cancel the activation of 2k Δ N by Smad2 and IKK α , with or without stimulation with TGF- β 1 (Figure 2B). The SMD1m and SMDHS2m mutations did not affect the *luc* expression significantly. Thus, SMD2 was determined as the common target site for Smad2 and IKK α . Furthermore, this result ruled out the possibility that IKK α ac-

tivates 2k Δ N through NF- κ B because the Rel/NF- κ B complex binds to 5'-GGGRNNYYCC-3' (R, purine; Y, pyrimidine).

Requirement of IKK α for Smad2-Induced 2k Δ N Activation

We performed IKK α silencing followed by luciferase assay with 2k Δ N-luc (Figure 3A). In comparison with the control Csi-introduced A431 cells, luciferase activity produced by the IKK α si-transfected cells was decreased by approximately 50% on Smad2 cotransfection with or without induction of TGF- β 1. The levels of 2k Δ N enhancement by IKK α and the Smad2/IKK α combination declined likewise (~50%).

FaDu cells derived from a hypopharyngeal SCC display an invasive phenotype [34–36], still expressing a level of *p63* [8,15]. Without the entire *Smad4* gene [37,38], FaDu allowed activation of 2k Δ N by *Smad2* and *IKK α* . When 10% FBS was contained in the culture medium throughout the reporter assay, *Smad2* caused an approximately five-fold increase in the luciferase assay (Figure 3B, left), similar to the activation level in A431 (seven-fold) under the same conditions (right). We detected a blunted response of 2k Δ N to both *IKK α* and *Smad2* (~70% reduction in the luciferase activity) in *IKK α* si-transfected FaDu cells (Figure 3B, middle).

Western blot analysis indicated a significant (55%–65%) decrease in the *IKK α* protein 3 days after the *IKK α* si transfection in the two cell lines (Figure 3C). Thus, endogenously expressed *IKK α* seemed essential for 2k Δ N activation by transfected *Smad2* in the reporter assay.

Endogenous Δ Np63 Induction by TGF- β in A431

To assess whether *Smad2* is physically associated with the identified sequences upstream of the Δ Np63-encoding sequences on chromo-

some 3, we performed ChIP experiments (Figure 4A). DNA fragments coprecipitated with an anti-*Smad2* antibody, anti-RNA polymerase II antibody (anti-pol II), and a control nonimmune IgG were examined for the SMD2 target sequence and the glutaraldehyde phosphate dehydrogenase (*GAPDH*) gene promoter sequence by PCR. A corresponding segment in the TAp63 promoter/enhancer region was also tested as a control. In A431 cells, the SMD2 site was more enriched (four-fold or higher) in the anti-*Smad2* precipitate than in the control IgG precipitate, and the *GAPDH* promoter was more enriched in the anti-pol II precipitate than in the control. The TA segment was not selectively coprecipitated by any antibody examined. Thus, association of *Smad2* with the SMD2 site in A431 was strongly suggested.

In FaDu cells, however, the anti-*Smad2* antibody failed to precipitate the segment containing SMD2. Results with the anti-pol II and control IgG were comparable to those in A431. Binding of *Smad2* to the SMD2 site on the chromosomal DNA was less probable in FaDu.

We next examined whether TGF- β could influence the endogenous *p63* expression. Two hours after the TGF- β 1 (10 ng/ml) stimulation of

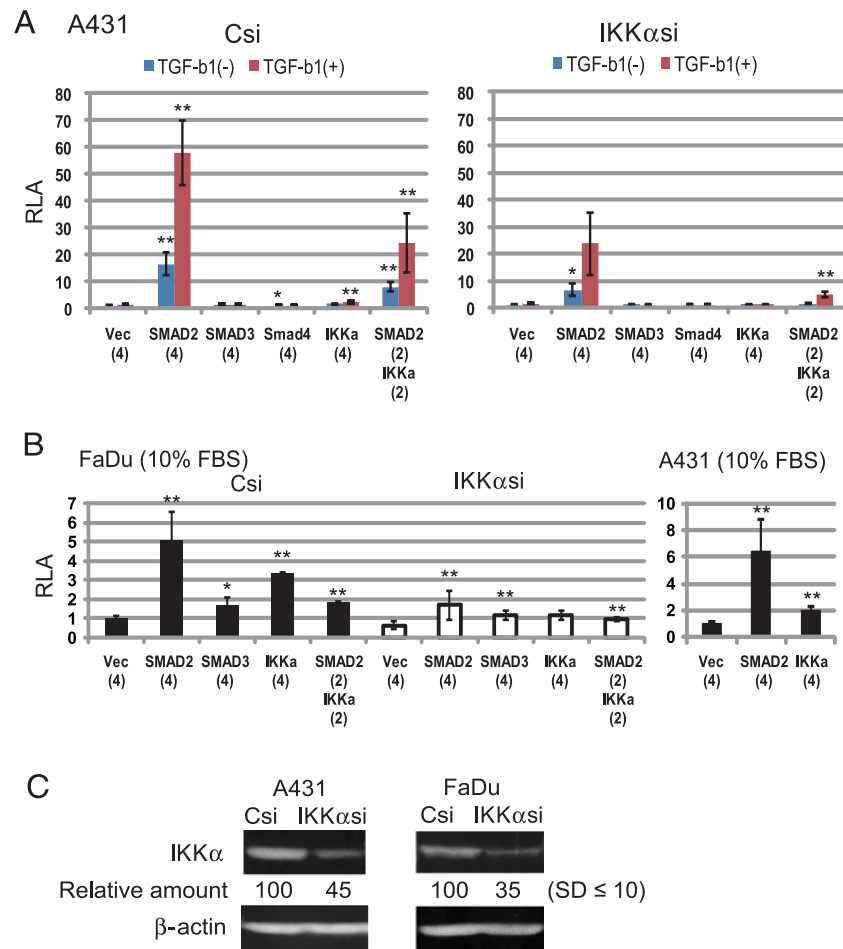


Figure 3. *Smad2* requires endogenously expressed *IKK α* for activation of the Δ Np63 promoter. (A) *Luc* reporter assay in A431 cells with *IKK α* silencing. Twenty-four hours after the siRNA (Csi, *IKK α* si) introduction, cells were seeded onto 24-well plates to perform the luciferase assay with 2k Δ N-luc and expression vectors of the *Smad* proteins and *IKK α* . Cells were incubated with (+) or without (–) TGF- β 1 (TGF-b1) as described in the legend to Figure 2. Luciferase activity 48 hours after the 2k Δ N-luc transfection was determined. Relative luciferase activity (RLA) is represented as mean \pm SD; $n \geq 3$. Statistic significance: ** $P < .01$, * $P < .05$, versus control experiment with Vec. (B) *IKK α* silencing in FaDu cells. We determined luciferase activities produced by the Csi- and *IKK α* si-transfected cells cultured with 10% FBS. Results with A431 under the same conditions are also shown. (C) Western blot analysis of A431 and FaDu cells with *IKK α* silencing. Day 3 cells after the transfection of Csi and *IKK α* si were analyzed for *IKK α* (85 kDa) and β -actin (43 kDa). Relative amounts of *IKK α* were obtained by standardization with β -actin for each cell line.

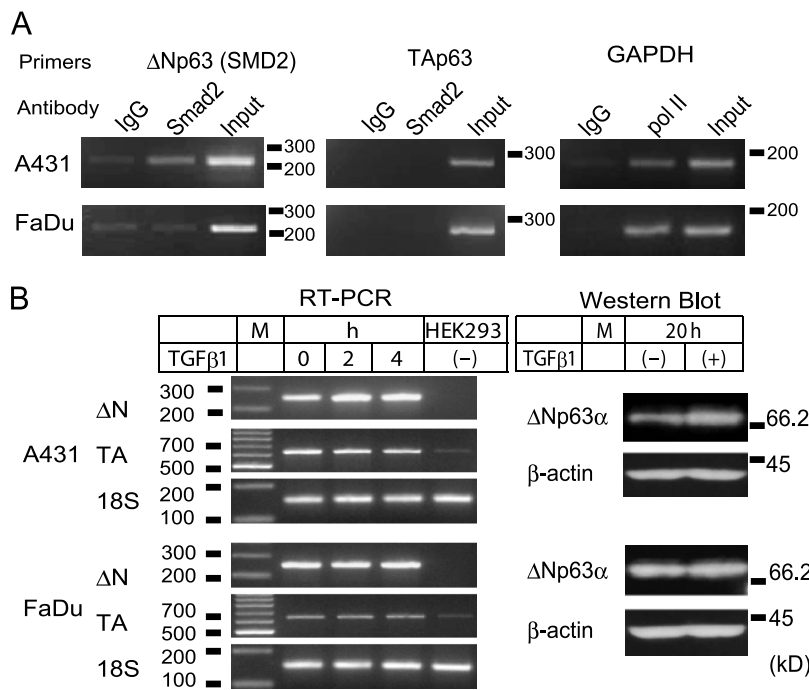


Figure 4. Endogenous Δ Np63 activation by TGF- β 1 in A431. (A) ChIP. A431 and FaDu nuclear extracts were examined for association of Smad2 with the SMD2 site. Antibodies used and amplified chromosomal segments, SMD2 (220 bp), TA (282 bp), and GAPDH (166 bp) are indicated. (B) Induction of endogenous Δ Np63 by TGF- β 1. Cells incubated with TGF- β 1 (10 ng/ml) for indicated periods (h) were analyzed for TAp63 and Δ Np63 mRNA by RT-PCR. Total RNA from HEK293 was tested as a negative control of Δ Np63 expression. Ribosomal RNA (18S) was amplified as a quantitative control. A 100-bp DNA ladder was loaded in lane M. Cells incubated with (+) or without (-) TGF- β 1 (for 20 hours) were also analyzed by Western blot analysis for Δ Np63 α . The blots were reprobbed with an anti- β -actin antibody. Sizes (kDa) and positions of the standard proteins are indicated.

serum-starved A431 cells, Δ Np63 messenger RNA (mRNA) was increased two-fold, which was retained to 4 hours (Figure 4B; RT-PCR). The TA-type mRNA was not altered. A 1.7-fold (1.7 ± 0.2) increase in Δ Np63 α , the most predominant p63 protein, was obvious 20 hours after the TGF- β 1 input (Figure 4B; Western blot). p63 expression in FaDu cells was not affected by TGF- β 1 as observed in RT-PCR and Western blot analysis. Thus, TGF- β 1 enhanced the chromosomal Δ Np63 transcription in A431 but not in FaDu.

Detection of Smad2- $IKK\alpha$ Association in the Nucleus of A431

The status of Smad2 and $IKK\alpha$ in the two SCC lines was analyzed by Western blot analysis (Figure 5A). Under the normal culture condition of A431, Smad2 existed in the cytosol and nucleus at a ratio of 8:2, showing nuclear localization of the phosphorylated form, P-Smad, reactive with antibodies recognizing phosphorylation at Ser465/467 of this protein (Figure 5A, right). In addition to a large amount of $IKK\alpha$ (85 kDa) in the cytosol, a small population of this protein (~14%) was detected in the nucleus. When Flag-tagged Smad2 was introduced into A431 by transfection, the cytoplasmic/nuclear distribution of the entire Smad2 molecules did not change significantly (Figure 5A, left). $IKK\alpha$ was efficiently coprecipitated with phosphorylated Flag-Smad2 from the nuclear extract by an anti-flag antibody. Although a large amount of $IKK\alpha$ was contained in the cytosol, it was poorly coprecipitated with Flag-Smad2 from the cytosol fraction.

FaDu cells contained a lesser amount of Smad2, ~45% in comparison with A431, showing its cytosolic and nuclear distributions at a ratio of

10:1 (Figure 5A, far right). Furthermore, $IKK\alpha$ was also decreased by ~50% in the total amount and hardly detectable in the nuclear fraction.

These differences between the two cell lines were confirmed by immunofluorescence analyses (Figure 5B). A431 contained Smad2 inside the nucleus as well as in the cytosol, whereas FaDu had a decreased amount of Smad2 showing a brighter signal outside the nucleus. A significant decrease in $IKK\alpha$ and the lack of the Smad4 protein in FaDu were also observed.

Induction of an Invasion Activity Either by p63 or by $IKK\alpha$ Silencing

A431 cells displayed intense p63 immunofluorescence confined to the nucleus as observed by laser scanning microscopy (Figure 6A, left). Consistent with the Western blot analysis (Figure 5A), cells showing a stronger $IKK\alpha$ label in the cytosol than in the nucleus were seen predominantly. Nuclear accumulation of $IKK\alpha$ was rarely detected ($\leq 5\%$ cells). We carried out gene silencing by transfection of p63si and $IKK\alpha$ si. As reported other studies [18,28,29], p63 knockdown cells showed a substantial decrease in $IKK\alpha$. Conversely, $IKK\alpha$ knockdown cells also displayed an obvious decrease in p63. This result not only supported our results that $IKK\alpha$ facilitated p63 expression but also implied mutual activation between p63 and $IKK\alpha$ in A431.

FaDu cells also showed a reduction of $IKK\alpha$ on the elimination of p63 (Figure 6A, right). However, $IKK\alpha$ knockdown cells maintained the p63 level as in the control Csi-transfected cells. In Western blot analysis, all of the detectable p63 isoforms including Δ Np63 α and TAp63 γ were greatly decreased by p63si transfection (Figure 6C, left),

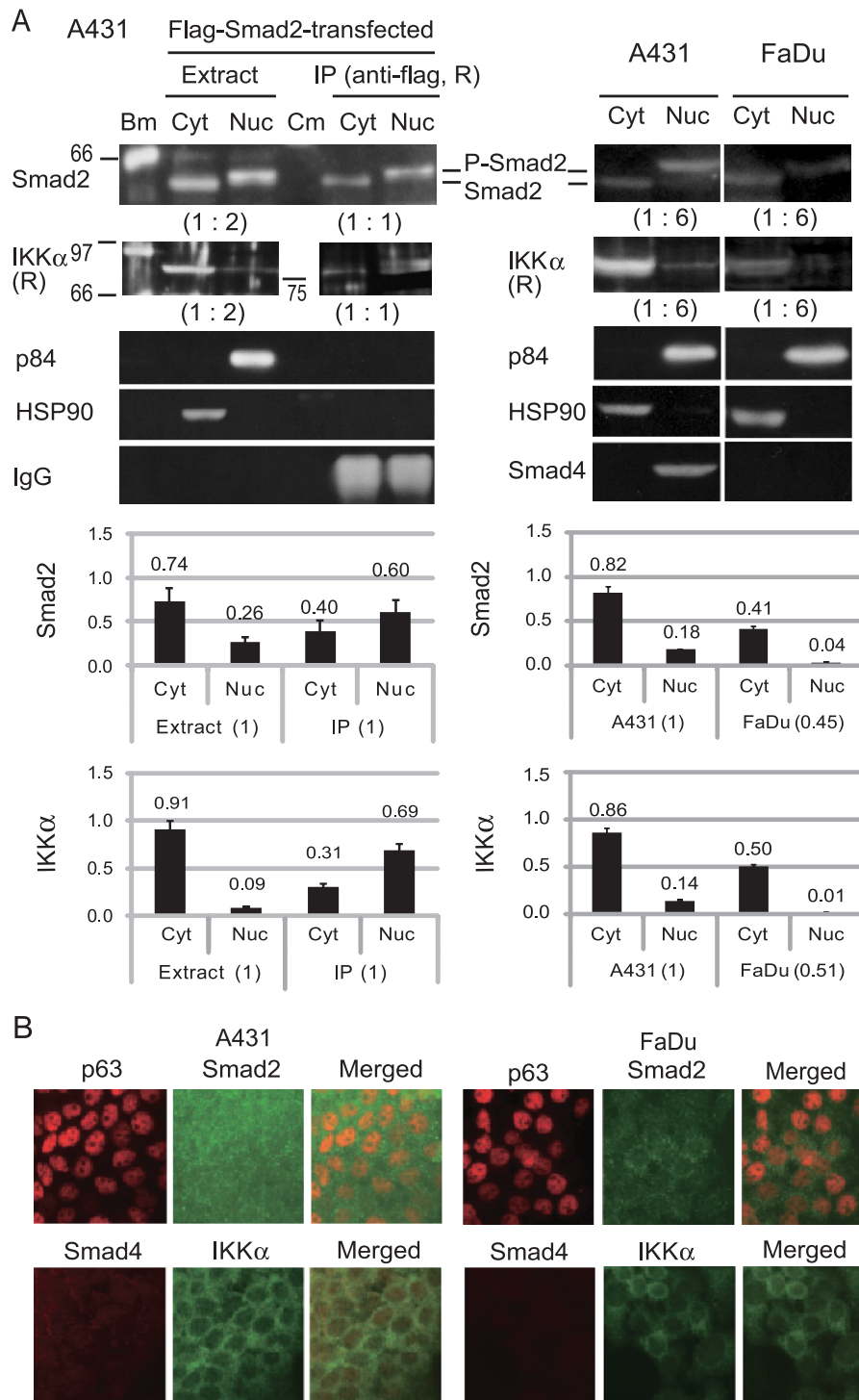


Figure 5. Status of the signaling molecules in A431 and FaDu. (A) Smad2 and IKKα were analyzed for the level of expression and subcellular localization by Western blot analysis. (Left panels) Cytosolic (Cyt) and nuclear (Nuc) fractions were obtained from A431 cells transfected with the Flag-Smad2 expression vector. Immunoprecipitation (IP) was carried out with a rabbit (R) anti-Flag antibody (PM020). Blot membrane was probed with anti-Smad2 (sc-101153), anti-IKKα (sc-7218), anti-p84 nuclear protein (ab487), and anti-HSP90 (ab1429) antibodies. Sizes (in kDa) and positions of the biotinylated standard proteins (Bm) and prestained color marker proteins (Cm) are indicated. Positions of Smad2 and phosphorylated Smad2 (P-Smad2) reactive with an anti-phospho-Smad2 (Ser465/467) antibody (#3101) are also indicated. Ratios of the sample amounts analyzed are indicated in parentheses. Graphs show relative amounts of Smad2 and IKKα in the fractions as determined by measurement of the band intensities ($n = 3$). (Right panels) Cytosol (Cyt) and nuclear (Nuc) fractions from untransfected A431 and FaDu cells were examined. Protein band intensities were measured and standardized with HSP90 and the ratio of sample amounts (1:6). Graphs show comparison of the total amount and subcellular distribution of Smad2 and IKKα between A431 and FaDu ($n = 3$). (B) Immunofluorescence analysis of formalin-fixed A431 and FaDu cells. Scale bars, 50 μm.

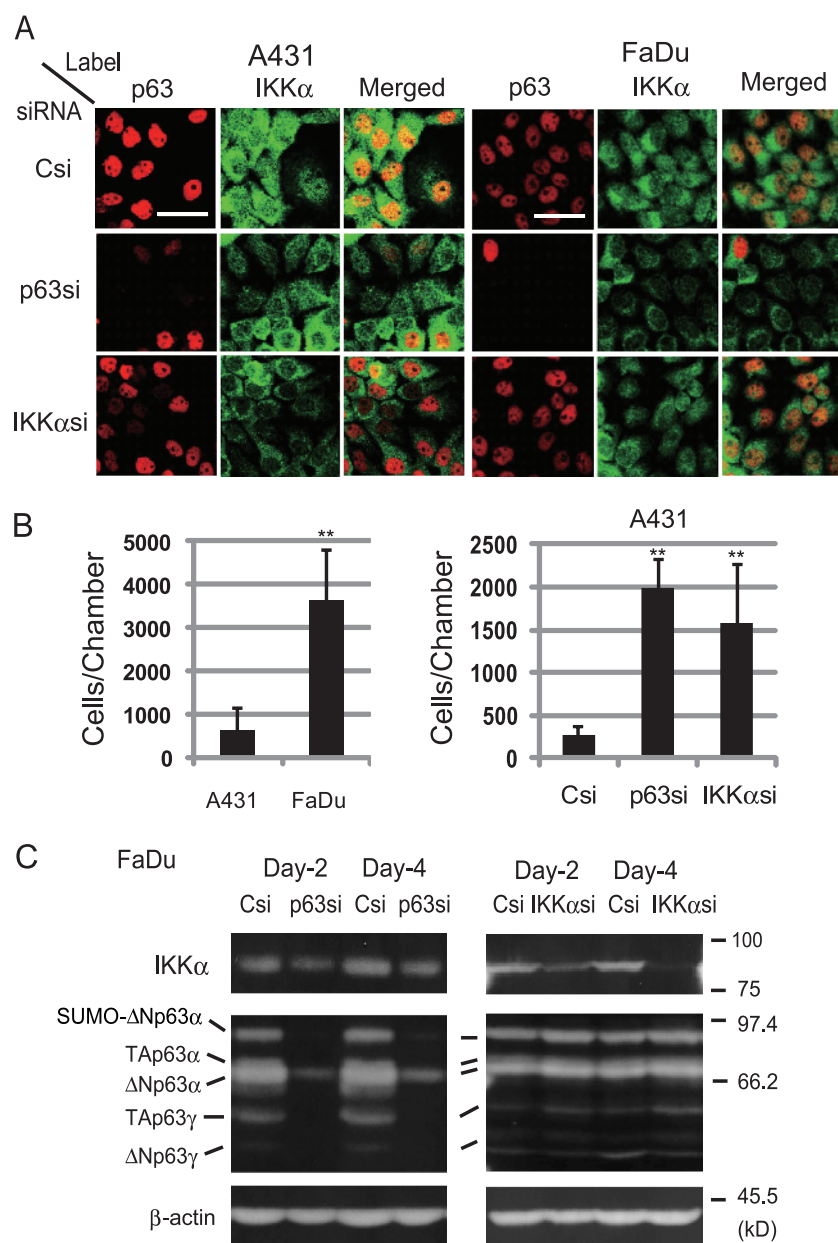


Figure 6. Induction of an invasion activity by p63 and IKK α knockdown in A431 cells. (A) Immunofluorescence analyses of siRNA-introduced A431 and FaDu cells. Two days after transfection with Csi, p63si, and IKK α si, cells were formalin-fixed for immunostaining. Scale bars, 50 μ m. (B) Invasion assay was carried out with Matrigel matrix cell culture inserts. A431 and FaDu cells without transfection (left) and A431 cells after siRNA (Csi, p63si, and IKK α si) transfection (right) were examined. Statistical significance: $^{***}P < .01$ versus A431 (left), $^{***}P < .01$ versus Csi (right). (C) Western blot analysis for p63 and IKK α in FaDu cells transfected with siRNA. Positions of the p63 isoforms were determined as described [15,47]. IKK α , β -actin, and sizes of the standard proteins are also shown.

which was accompanied by an obvious reduction of IKK α . However, IKK α silencing did not affect the p63 isoforms (Figure 6C, right). Given the lack of Smad2-SMD2 association (Figure 4A) and the absence of IKK α in the nucleus (Figure 5), the endogenous Δ Np63 expression in FaDu cells may not depend on the Smad2-IKK α pathway of TGF- β .

In an invasion assay with Matrigel-coated membrane, FaDu cells penetrated through the matrix protein complex, whereas A431 failed to do so (Figure 6B). Interestingly, either by p63 or by IKK α gene silencing, A431 gained an invasive activity, implying that both IKK α and p63 are involved in the maintenance of the noninvasive phenotype of A431 cells.

Changes in p63 and IKK α in the Progression of SCC

To assess alterations of p63 and IKK α during the SCC progression, we analyzed conventional oral carcinoma sections at different stages, hyperplasia (carcinoma *in situ*), well-differentiated (grade 1), and poorly differentiated (grade 3), for p63 and IKK α by immunostaining (Figure 7A). Four or six cases of each grade were examined. Normal gingival sections displayed nuclear p63 expression in the basal and suprabasal layers of the epithelium as described [3,4]. The IKK α label was spread throughout the cells, causing an enhanced signal in the cytoplasm. Hyperplasia lesions (4/4) displayed more enhanced p63 immunofluorescence than in the normal tissues. As observed in A431 (Figure 5), a

majority of the IKKα molecules existed in the cytoplasm. Intriguingly, well-differentiated SCC lesions (6/6) displayed a significant enhancement of p63 as well as nuclear accumulation of IKKα. Furthermore, in poorly differentiated, invasive SCC sections (5/6), loss of p63 positive cells, and relocalization of IKKα to the cytoplasm were profound.

To study more quantitatively the status of these proteins in SCCs, we analyzed SCC tissue arrays. The BC34011 slide contained 60 cores of different stages (grades 1-3) of SCC and 3 cores of normal tissues. Six cores of hyperplasia on OR802 were also examined. All of the tissue array immu-

nofluorescence images are presented in Figures W3 and W4. Cells (100 < n ≤ 350) in a typical field of view were sorted by 1) the presence (+) or absence (-) of p63 and 2) nuclear accumulation (N > C) or cytoplasmic/whole-cell distribution (N ≤ C) of IKKα. We obtained, for each core, composition (%) of the four categories: p63(+)/IKKα (N > C), p63(+)/IKKα (N ≤ C), p63(-)/IKKα (N > C), and p63(-)/IKKα (N ≤ C) (Figure 7B).

In normal gingiva and hyperplasia tissues, a large population (≥90%) showed the p63(+)/IKKα (N ≤ C) phenotype in six of eight cores. Interestingly, p63(+)/IKKα (N > C) cells were predominant (>50%)

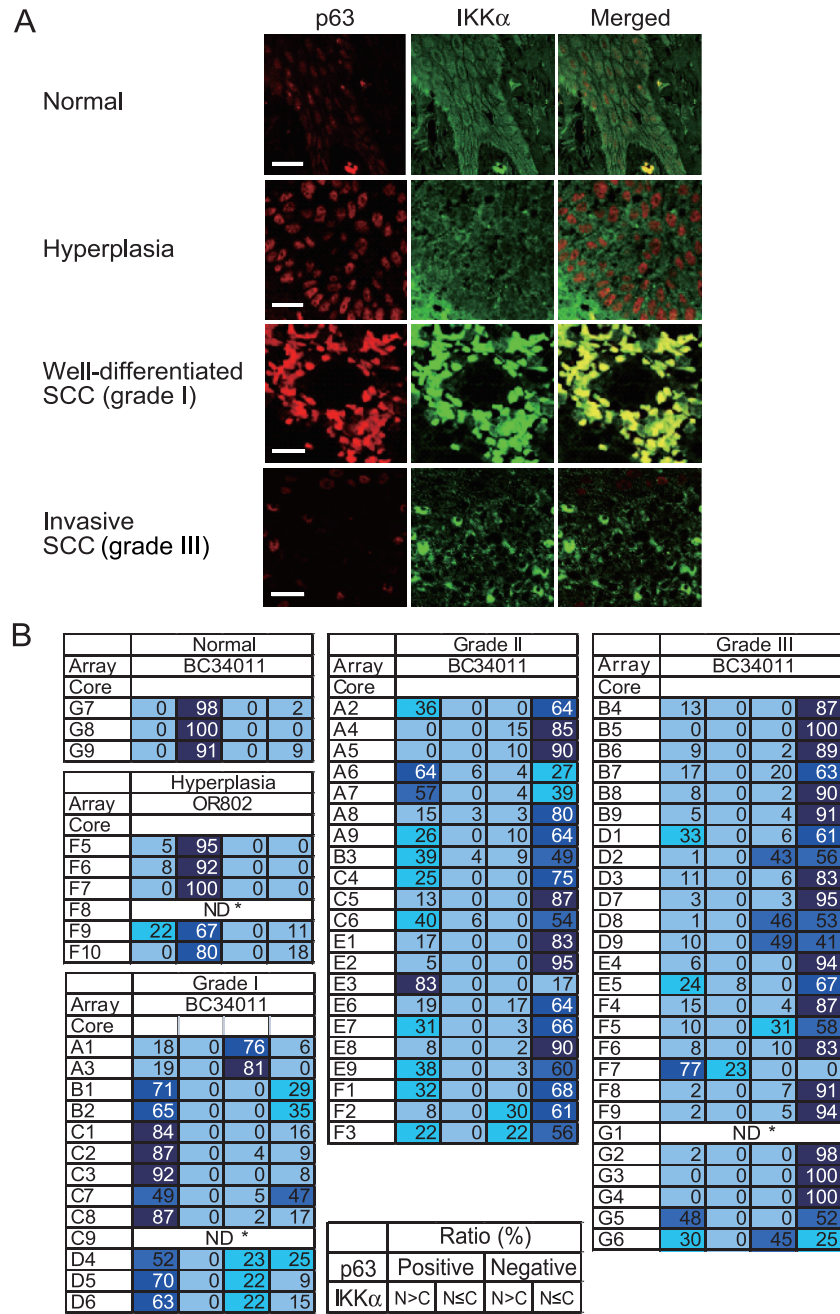


Figure 7. Amplification of p63 with IKKα in well-differentiated SCC. (A) Oral carcinoma sections at different pathologic stages were analyzed by immunofluorescence staining for p63 and IKKα. Typical images representing four or six individual samples of the indicated stages are presented. The well-differentiated SCC images contain nests of keratin pearls in the dark areas. Scale bars, 50 μm. (B) Summary of the SCC tissue array analysis. Composition (%) of the p63(+)/IKKα (N > C), p63(+)/IKKα (N ≤ C), p63(-)/IKKα (N > C), and p63(-)/IKKα (N ≤ C) populations was calculated for each core with predetermined cancer grade. Results are collectively shown for each category: normal, SCC grade 1, 2, 3 (indicated as I, II, III in array BC34011), and hyperplasia (OR802). ND* indicates not determined. (OR802-F8 and BC34011-G1 contained less than 10 labeled cells in each field of view. BC34011-C9 showed cells with cytoplasmically localized p63 and was therefore eliminated.)

in 9 of 12 cores of grade 1 SCC. Furthermore, the p63(+)/IKK α (N > C) population was diminished markedly in grade 2 (moderately well differentiated) tissues with a concomitant increase in the p63(-)/IKK α (N \leq C) population. In grade 3 SCCs, p63(-)/IKK α (N \leq C) cells were predominant (>50%) in 22 of 25 cores. Only a small number (\leq 10%) of p63(+)/IKK α (N > C) cells remained in 16 cores of grade 3.

These results suggest that the maximum expression of p63 at the stage of well-differentiated SCC is associated with IKK α nuclear accumulation. Furthermore, the loss of p63 during the malignant conversion seemed to occur concurrently with the decline of IKK α .

Discussion

We have shown that the Δ Np63 promoter is activated by the recently identified keratinocyte-specific TGF- β signal pathway with Smad2 and IKK α in SCC. This mechanism may underlie the suppression of malignant conversion by p63 and IKK α .

The SMD2 (5'-GTCTGTCT-3') site matching the SBE consensus was determined as the *cis*-acting sequence to respond to the Smad2/IKK α signal. The zebrafish Δ Np63 promoter also contains SBE-like sequences, with which epithelial cells responded to BMP2 *in situ* [20]. Our results do not necessarily dismiss the possibility that the human BMP signaling also contributes to p63 expression in other cell types. However, the functional SBE-like sequences in the Δ Np63 promoter are little conserved between human and zebrafish (Figure 1), implying that the disparity reflects diversification of the *cis*-regulatory sequences and the transcription factor usage in mammalian evolution.

By the ChIP, TGF- β 1 stimulation and gene silencing experiments, we concluded that transcription of chromosomal Δ Np63 is activated by the Smad2/IKK α signaling in A431 cells. In contrast, the pathway seemed dispensable for Δ Np63 in FaDu cells, despite the observation of 2k Δ N transactivation by Smad2 in an IKK α -dependent mode. This is not a contradiction between the assay results. Transfection of Smad2 in the reporter assay may have caused the pathway activation in FaDu. Furthermore, transcription of Δ Np63 on the chromosomal DNA can be influenced by the sequences upstream and downstream of the 2k Δ N region and by proteins interacting with them.

Conversely, recent studies showed transcriptional activation of the IKK α gene by p63 [18,28,29], the reverse reaction of our finding, hypothesizing mutual activation between p63 and IKK α . This hypothesis seems relevant to A431 cells in which IKK α knockdown decreased p63 and vice versa. The same level of invasion activity was induced by p63 and IKK α silencing in A431, implying that the mutual activation mechanism is important to maintain the non-invasive phenotype.

As many other SCC lines, A431 expresses both keratinocyte stem cell markers and an early differentiation marker, involucrin [39]. When the IKK α distribution pattern and the result of invasion assay are taken into account, we consider A431 to be an "advanced" hyperplasia. FaDu cells produce proinvasive molecules including furin, matrix metalloproteinase 3 (MMP-3, also termed *stromelysin-1*), and other MMPs, and characterized as an invasive SCC line [34,36], which was consistent with our invasion assay. Earlier studies showed that cell lines derived from SCCs all fail, to a greater or lesser extent, to complete the process of keratinization [40,41], indicating a limitation in well-differentiated SCC isolation as a cell line.

By an extensive histologic study in conjunction with biochemical analyses, nuclear translocation of IKK α was found essential for the keratinocyte-specific TGF- β pathway activation and for suppression of the malignant conversion of SCC [23,27]. Our immunofluores-

cence analyses with tissue arrays are consistent with the previous studies as well as our finding that IKK α acts in the p63-inducing mechanism. The down-regulation of p63 during the malignant conversion [10,11,13] may be associated with the significant decrease of IKK α as demonstrated by the gene silencing and invasion assay with A431 (Figure 6).

Invasion and metastasis of SCC involve "epithelial-to-mesenchymal transition" controlled by varied intracellular signaling pathways [42], production of proteases [43], and functional interactions with fibroblasts [44] and endothelial cells [45]. The p63 and Smad2/IKK α systems independently control many target genes in keratinocytes [14,16-19,23,27,46], which may collectively facilitate the proposed invasion-suppressing function of the keratinocyte-specific TGF- β signaling pathway. Mechanisms of the nuclear accumulation of IKK α and its elimination in the course of SCC progression await extensive studies.

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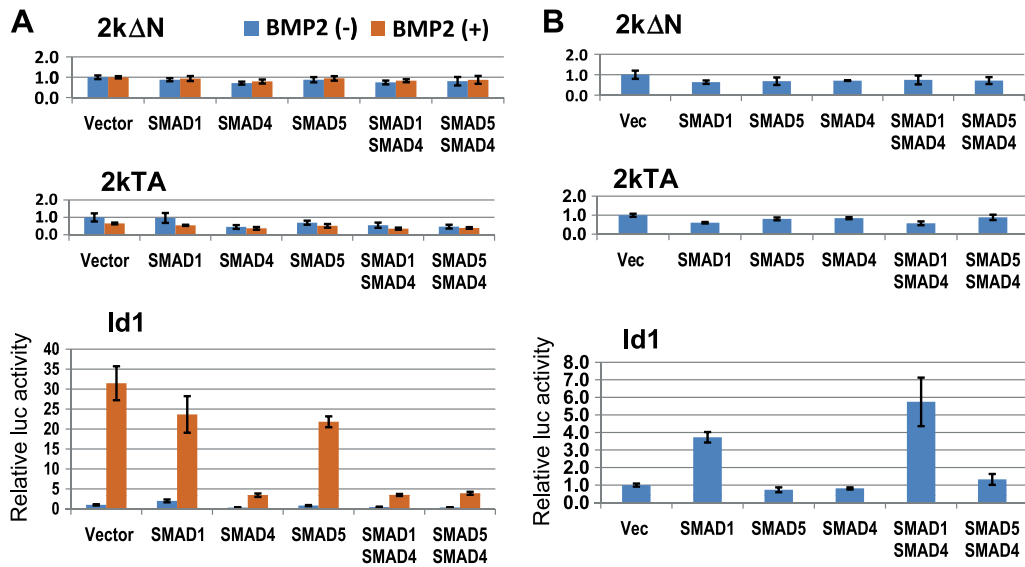


Figure W1. The p63 promoters do not respond to the BMP signal in C2C12 and HepG2 cells. (A) The TAp63 (2kTA), ΔNp63 (2kΔN), and control Id1 (*J Cell Physiol* 2002;193:299–318) promoters were tested for ability to respond to BMP2 by luciferase gene (*luc*) expression assay in C2C12 cells. After transfection of the Smad expression vector(s) with the *luc* plasmid at a ratio of 4:1, cells were incubated with (+) or without (-) BMP2. Luciferase activity is indicated in relation to the enzyme expression (1.0) driven by the *luc* plasmid with the empty vector (Vec) in the absence of BMP2. Data are represented as mean ± SEM. (B) The same promoters were examined for activation by Smad-1, -4, and -5 in HepG2 cells without BMP2 induction.

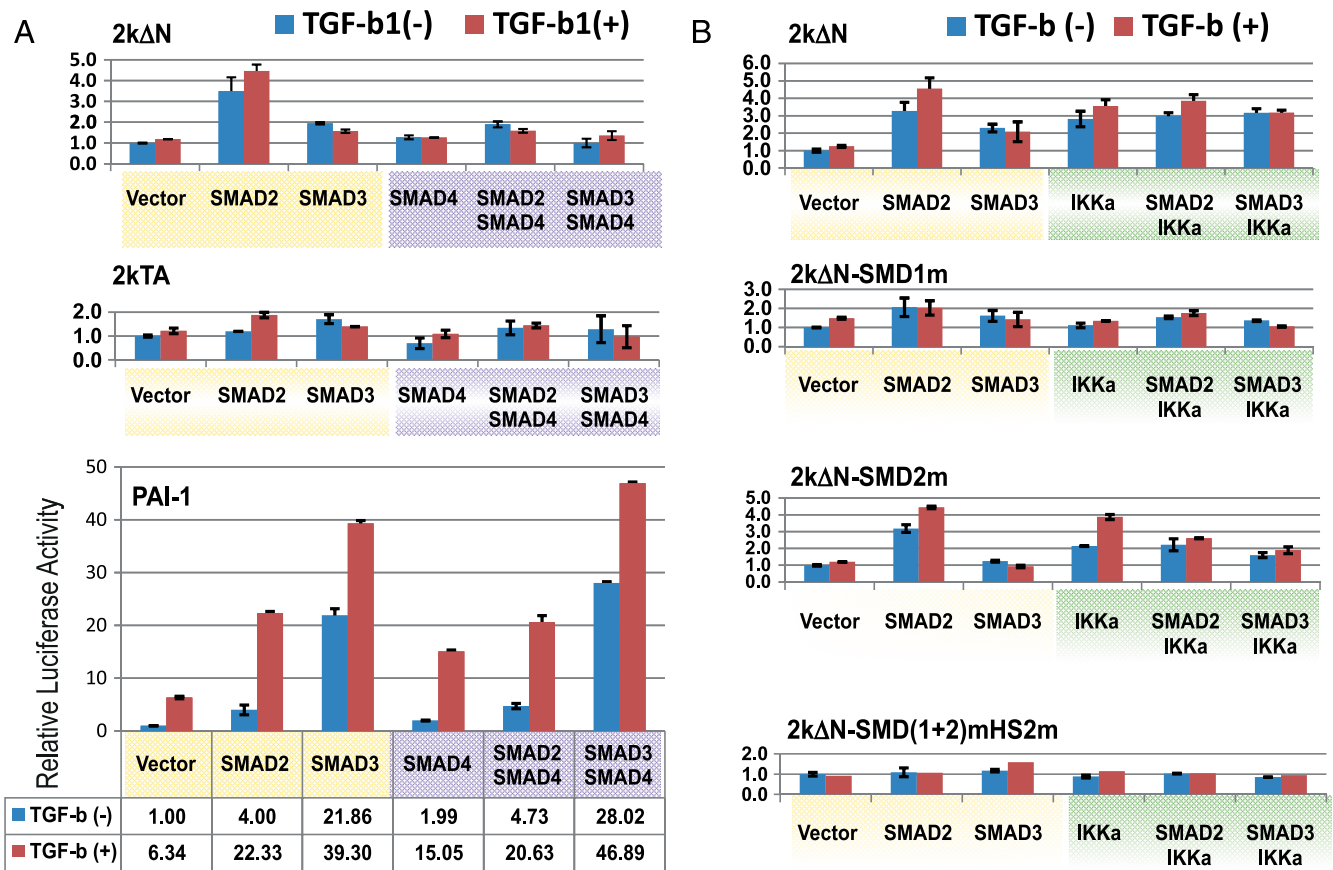


Figure W2. Moderate activation of the ΔNp63 promoter by Smad2 and IKKα in HepG2 cells. (A) The 2kTA, 2kΔN and PAI-1 promoter/enhancer regions were tested for sensitivity to Smad-2, -3, and -4 in HepG2 cells. After transfection of the Smad expression vector(s) and the *luc* plasmid at a ratio of 4:1, cells were serum-starved, and incubated with (+) or without (-) TGF-β1 (TGF-β1) for 24 hours. Relative luciferase activity is shown. Data are represented as mean ± SEM. (B) 2kΔN and its mutants were analyzed for response to Smad2, Smad3, and IKKα (IKKα).

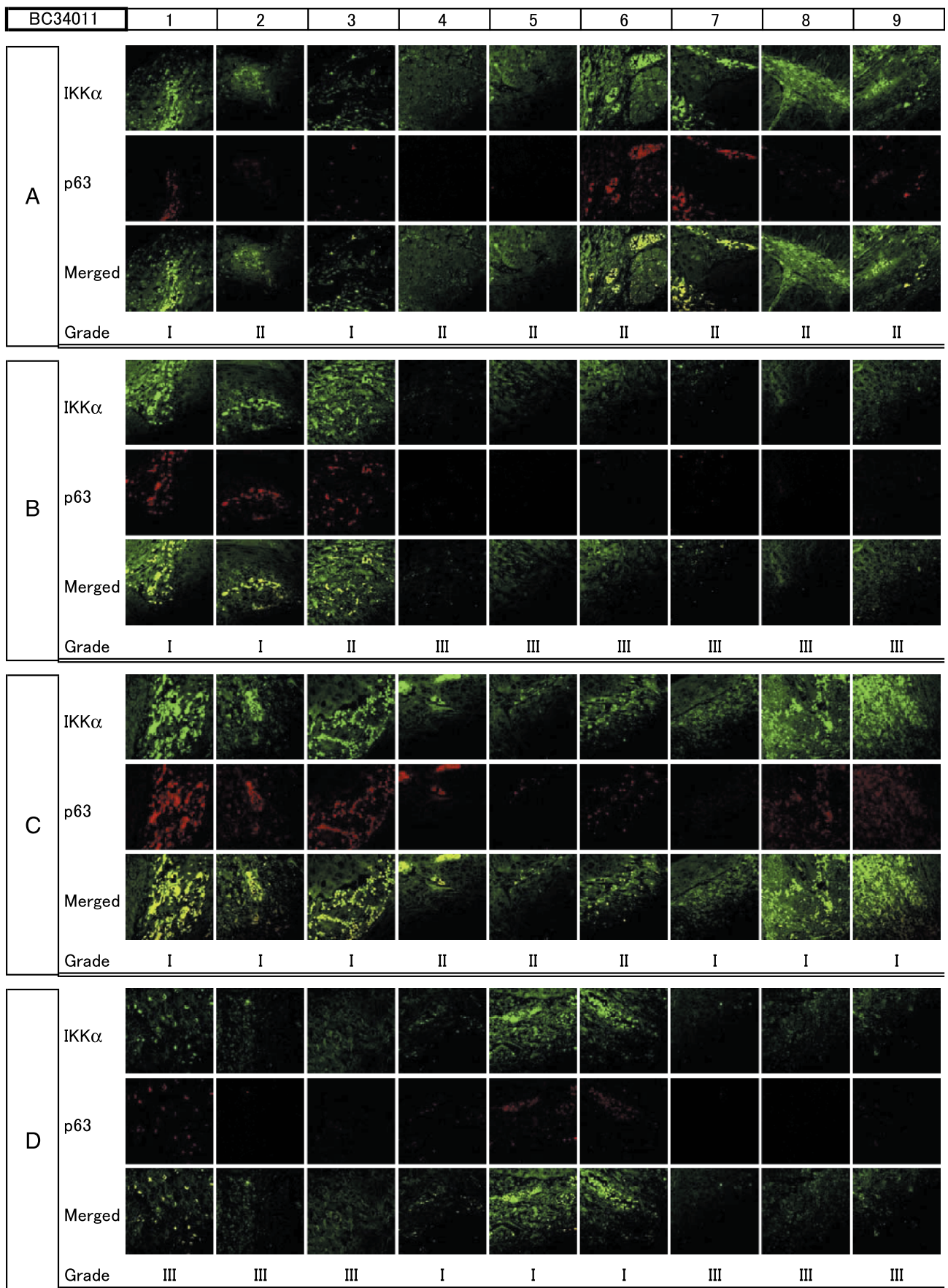


Figure W3. Immunofluorescence analysis of SCC tissue array BC34011. Immunofluorescence images of p63 (red) and IKK α (green) were obtained for every core placed at rows A to G and columns 1 to 9. Predetermined cancer grades (1, 2, and 3) are denoted by I, II, and III.

BC34011		1	2	3	4	5	6	7	8	9
E	IKK α									
	p63									
	Merged									
	Grade	II	II	II	III	III	II	II	II	II
F	IKK α									
	p63									
	Merged									
	Grade	II	II	II	III	III	III	III	III	III
G	IKK α									
	p63									
	Merged									
	Grade	III	III	III	III	III	III	Normal	Normal	Normal

Figure W3. (continued).

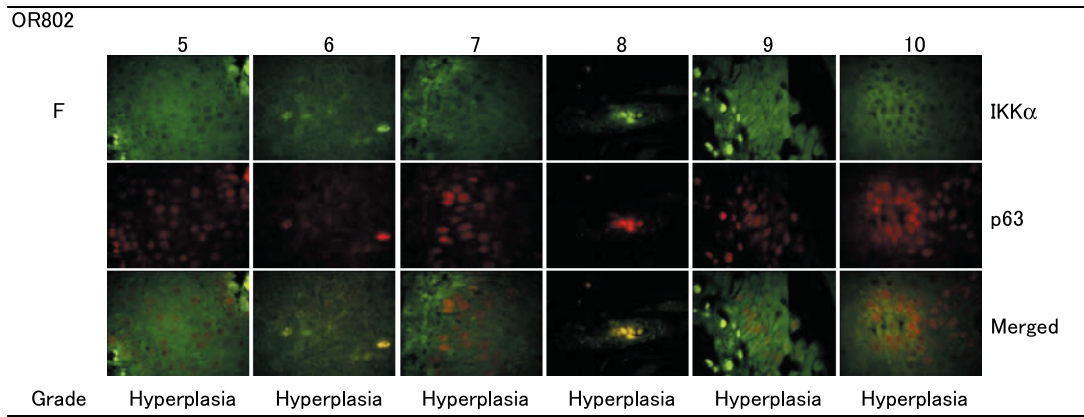


Figure W4. Immunofluorescence analysis of SCC tissue array OR802. Cores of hyperplasia (in row F, columns 5-10) are shown. Because F8 contained less than 10 countable cells with a ductal cell morphology, this core was omitted from the analysis shown in Figure 7B.