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Homeobox gene *Dlx3* is regulated by p63 during ectoderm development: relevance in the pathogenesis of ectodermal dysplasias

Nadezda Radoja¹, Luisa Guerrini², Nadia Lo Iacono^{2,3}, Giorgio R. Merlo³, Antonio Costanzo⁴, Wendy C. Weinberg⁵, Girolama La Mantia⁶, Viola Calabrò^{6,*} and Maria I. Morasso^{1,*†}

Ectodermal dysplasias (EDs) are a group of human pathological conditions characterized by anomalies in organs derived from epithelial-mesenchymal interactions during development. *Dlx3* and p63 act as part of the transcriptional regulatory pathways relevant in ectoderm derivatives, and autosomal mutations in either of these genes are associated with human EDs. However, the functional relationship between both proteins is unknown. Here, we demonstrate that *Dlx3* is a downstream target of p63. Moreover, we show that transcription of *Dlx3* is abrogated by mutations in the sterile α -motif (SAM) domain of p63 that are associated with ankyloblepharon-ectodermal dysplasia-clefting (AEC) dysplasias, but not by mutations found in ectrodactyly-ectodermal dysplasia-cleft lip/palate (EEC), Limb-mammary syndrome (LMS) and split hand-foot malformation (SHFM) dysplasias. Our results unravel aspects of the transcriptional cascade of events that contribute to ectoderm development and pathogenesis associated with p63 mutations.

KEY WORDS: *Dlx3*, p63, Transcription, Ectodermal dysplasias, Mouse development

INTRODUCTION

During embryonic development and organ formation, a series of signals between epithelial cells and the underlying mesenchymal cells are the basis for the formation of a variety of appendages and/or organs (Pispa and Thesleff, 2003). Anomalies in epithelial-mesenchymal-derived organs are characteristics of human pathological conditions defined as ectodermal dysplasias (EDs) (Priolo and Lagana, 2001).

Mutations in *DLX3* and *p63*, among other genes, have been directly linked with EDs. The *Dlx* and p63 families of transcriptional effectors are essential for the development of the epidermis and/or embryonic appendages (Panganiban and Rubenstein, 2002; Merlo et al., 2003; Morasso and Radoja, 2005; Koster and Roop, 2004). *Dlx3* expression has been detected in the hair follicle, tooth, limb bud, branchial arches, labyrinthine layer of the placenta, osteoblasts and epidermis (Morasso et al., 1995; Morasso et al., 1999; Hassan et al., 2004). Here, we present evidence that *Dlx3* is regulated by p63 as part of a transcriptional regulatory pathway relevant to specific EDs.

p63 regulates multiple signaling pathways, such as the bone morphogenetic protein (BMP) and fibroblast growth factor (FGF) pathways (Laurikkala et al., 2006; Barbieri and Pietenpol, 2006). Transgenic and knockout (KO) mouse models indicate that p63 has essential roles in the development and maintenance of the stratified

epidermis (Yang et al., 1999; Mills et al., 1999; Koster et al., 2004; Koster and Roop, 2004). The *p63* gene is transcribed from two distinct promoters, giving rise to proteins that either contain (TAp63) or lack (Δ Np63) the amino terminal transactivating domain. The TA and Δ N isoforms both possess the DNA-binding and oligomerization domains, and, by alternative splicing at the 3' end, produce isoforms with different C-termini, termed alpha (α), beta (β) and gamma (γ) (Yang et al., 1998). The α isoforms contain a sterile α motif (SAM) – a domain with reputed importance in protein-protein interactions (Qiao and Bowie, 2005). p63 isoforms act as transcriptional activators and/or repressors (Ghioni et al., 2002; King et al., 2003; Wu et al., 2005), and bind to two or more tandem repeats of RRRCWWGYYY, but preferentially activate the RRRCGTGYYY sequence (Osada et al., 2005).

Mutations in the *p63* gene have been associated with EDs that include ectrodactyly-ectodermal dysplasia-cleft lip/palate (EEC), limb-mammary syndrome (LMS), split hand-foot malformation (SHFM) and ankyloblepharon-ectodermal dysplasia-clefting (AEC) syndrome. There is a correlation between the position of the mutation and the observed abnormal phenotype (van Bokhoven et al., 2001; McGrath et al., 2001; van Bokhoven and Brunner, 2002). Mutations in the *DLX3* gene are linked to tricho-dento-osseous (TDO) syndrome, which, like AEC, is characterized by defects in the development of hair, teeth and bone, and by absence of overt limb malformations (Price et al., 1998).

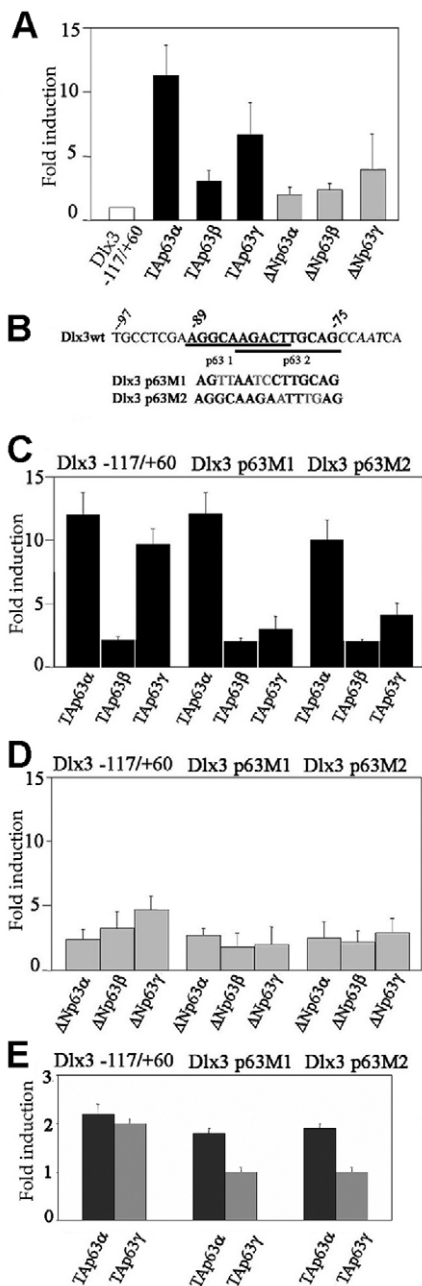
Partial-overlapping mRNA expression and phenotypes of specific human malformations caused by molecular lesions in either *p63* or *DLX3* suggest that these genes are components of common signaling pathways during embryonic development. Here, we show that p63 is able to bind and transactivate *Dlx3* both in vitro and in vivo. Mutant p63 proteins derived from AEC patients exhibit an impaired ability to transactivate *Dlx3*, indicating that the misregulation of the *DLX3* gene is involved in the pathogenesis of human syndromes associated to AEC.

¹Developmental Skin Biology Unit, NIAMS, NIH, Bethesda, MD 20892, USA.

²Department of Biomolecular and Biotechnological Sciences, University of Milan, Via Celoria 26. 20133, Milan, Italy. ³Dulbecco Telethon Institute c/o Istituto Tecnologie Biomediche CNR, 20100 Milan, Italy. ⁴Department of Dermatology, University of Rome 'Tor Vergata', Viale Oxford 81. 00133 Rome, Italy. ⁵Division of Monoclonal Antibodies, CDER/FDA, Bethesda, MD 20892, USA. ⁶Department of Structural and Functional Biology, University of Naples, Via Cinzia 26. 80126 Naples, Italy.

*Co-senior authors

†Author for correspondence (e-mail: morassom@mail.nih.gov)



MATERIALS AND METHODS

DNA constructs

The -117 to +60 DNA fragment of the *Dlx3* promoter (Park and Morasso, 1999) was inserted into the pGL3-Basic and pCAT-Basic vectors (Promega). Mutations in the p63-binding sites of the *Dlx3* promoter were obtained using the ExSite Mutagenesis kit (Stratagene). The coding sequences for Δ Np63 α , Δ Np63 β , and Δ Np63 γ were cloned into pBK-CMV (Stratagene). The TAp63 α , TAp63 β and TAp63 γ constructs were a gift from E. Candi (University of Rome 'Tor Vergata', Rome, Italy) and G. Melino (University of Rome 'Tor Vergata', Rome, Italy). The p63 mutants L518F, L518V and Q540L (AEC); E639X (SHFM); F5525 (EEC); and G76W and Δ AA (LMS) were kindly provided by H. van Bokhoven (Radboud University, Nijmegen, The Netherlands).

Cell culture

Primary mouse keratinocytes were grown according to Park and Morasso (Park and Morasso, 1999). Normal human epidermal keratinocytes [NHEK; a gift of M. Simon (SUNY, Stony Brook, NY, USA)] were derived from

Fig. 1. Transcriptional regulation of the *Dlx3* promoter by p63. (A) Fold-induction increase, compared with wild type, of *Dlx3*_{-117/+60} co-transfected with vectors expressing the different p63 isoforms. (B) Sequence of the mouse *Dlx3*-promoter region containing the two overlapping p63-binding sites (p63 site1 and p63 site2; underlined) and CCAAT box (italics). The mutated *Dlx3* sequences are shown in gray. (C-D) *Dlx3*_{-117/+60}, *Dlx3*p63M1 and *Dlx3*p63M2 constructs were co-transfected with vectors expressing the TAp63 (C) and Δ Np63 (D) isoforms, with activity shown relative to the basic activity of *Dlx3*_{-117/+60}. Transient transfection experiments were performed using mouse keratinocytes (C-D) and Saos-2 cells (E). Basal activity of the reporter was set to 1. Each histogram bar represents the mean of three independent transfection duplicates. Standard deviations are indicated. *Dlx3* p63M1, mutated at p63-binding-site 1; *Dlx3*p63M2, mutated at p63-binding-site 2.

newborn foreskin and cultured in K-SFM (Invitrogen). The human osteosarcoma Saos-2 cell line was maintained in RPMI 1640 and 10% FCS. The immortalized human keratinocyte HaCaT and H1299 non-small-lung-carcinoma cell lines were grown in DMEM and 10% FBS.

Transient transfection

Transient transfections of keratinocytes were performed with FuGENE6 (Roche) in a 1:3 ratio. PRL-SV40 vector was used as an internal control. Luciferase activity was measured 24-36 hours after transfection using the Dual-Luciferase Reporter Assay System (Promega). Transient transfections of Saos-2 and H1299 cells were performed according to Calabrò et al. (Calabrò et al., 2002). CAT reaction was performed 48 hours after transfection using 90 μ g of cell extract. β -Gal was used to normalize for transfection efficiency.

EMSA analysis

Nuclear extracts and EMSA analysis were carried out according to Park and Morasso (Park and Morasso, 1999) using the *Dlx3* p63site1+2 and non-specific competitor AP-2 binding site.

Chromatin immunoprecipitation assay

Chromatin immunoprecipitation (ChIP) was performed with chromatin from mouse keratinocytes and H1299 cells transfected with TAp63 α according to Caretti et al. (Caretti et al., 2004) using no antibody, IgG or p63 antibody (Santa Cruz, H137 and 4A4). Real-time PCR was performed using the Mx3000P System (Stratagene) and SyberGreen MasterMix (Applied Biosystems) with independent DNA samples and the following oligonucleotides: for mouse, *Dlx3*(F) 5'-GAGAAAGCGCGAGCGT-GTTTTGCC-3' and *Dlx3*(R) 5'-CCGGCTGTCGGTTCAGTCGCTGCGT-3'; for human, *DLX3*(F) 5'-AGAGAGGCGGAAGAGACGAG-3' and *DLX3*(R) 5'-GAGGAGGAGGAGAGAAGGA-3'; and for JAG2(F) 5'-CAAGTGGTGAACAAGGGAGACT-3' and JAG2(R) 5'-ACTG-CTGCCTTCTGGAAACTC-3'. Data are presented as fold differences relative to input and values are obtained by IgG with the formula $2^{[(Ct_{IgG}-Ct_{Input})-(Ct_{Ab}-Ct_{Input})]}$, where Ct is threshold cycles, IgG is normal rabbit IgG, Ab is specific antibody and Input is input genomic DNA. ACHR amplification was performed as a control for H1299 transfected with TAp63 α using ACHR(F) 5'-TGCCTCGGGTGAACCTAAGATG-3' and ACHR(R) 5'-GCCTCATTCGTCTTGGGAAC-3'.

Real-time PCR

For analysis of the expression of *p63* in mouse keratinocytes and embryonic tissues, the following oligonucleotides were used: TAp63(F) 5'-AGA-CAAGCGAGTTCCTCAGC-3', TAp63(R) 5'-TGCGGATACAATCC-ATGCTA-3', Δ Np63(F) 5'-ATGTTGTACCTGGAAAACAATG-3', Δ Np63(R) 5'-GATGGAGAGAGGGCATCAAA-3'. These oligonucleotides are designed in regions of the mRNA common to all isoforms (α , β and γ), and do not distinguish between these variants.

For *Dlx3* expression in mouse keratinocytes, the following oligonucleotides were used: *Dlx3*(F) 5'-ATTACAGCGCTCCTCAGCAT-3' and *Dlx3*(R) 5'-GCCTATAGGATCCCCCGTAG-3'. In embryonic

tissues: Dlx3(F) 5'-CGTTTCCAGAAAGCCCGTA-3' and Dlx3(R) 5'-CGTGGAAATGGGAAGATGTGT-3'. For normalization: GAPDH(F) 5'-TGTCAGCAATGCATCCTGCA-3' and GAPDH(R) 5'-TGTATGC-AGGGATGATGTTTC-3'.

For the analysis of *p63* and *Dlx3* mRNA levels in embryonic tissues, E10.5, E11.5 and E12.5 anterior or posterior limb buds from wild-type embryos were dissected, pooled in TRIZOL (Roche) and extracted, as indicated by the manufacturer. Real-time PCR was performed with a LightCycler (Roche) using FastStart DNA MasterPLUS SYBR-Green I (Roche). Standard curves were performed using wild-type cDNA with four calibration points: TQ; 1:3; 1:9 and 1:27. All samples were done in duplicates and the analysis was repeated twice. GAPDH was used for normalization, calculated using LightCycler Software 3.5.3. The results are expressed with the value relative to E10.5 (set at 1) for each mRNA.

Immunohistochemistry and whole-mount in situ hybridization

Immunohistochemistry was performed on 11 μ m cryostatic sections of E10.5 embryonic forelimbs. Sections were blocked with 10% goat serum in PBS for 40 minutes at room temperature. Antibodies used were: mouse monoclonal anti-p63 (4A4, 1:100, Santa Cruz) and rabbit anti-distal-less [pan-anti-Dlx, 1:100; kindly provided by G. Boekhoff-Falk (University of Wisconsin Medical School, WI, USA)]. As secondary antibody, anti-mouse-Cy2, anti-rabbit-Cy3 (1:100; Jackson Immuno-Research) and Envision anti-rabbit HRP (Dako) were used. Fluorescence micrographs were taken by confocal microscopy.

Whole-mount in situ hybridization was performed according to Acampora et al. (Acampora et al., 1999) on E10.5 *p63* KO embryos [Brdm2 line of *p63* KO kindly provided by D. Roop (Baylor College of Medicine, Houston, USA)] using a *Dlx3* probe (Morasso et al., 1995).

RT-PCR

Total RNA from human Saos-2 and HaCaT cells was prepared with TRIZOL (Roche). For reverse transcription (RT)-PCR, 3-4 μ g of total RNA were reverse-transcribed using SuperScript II (Invitrogen). The following oligonucleotides were used: Dlx3(F) 5'-ACCTACGGAGCCTCACC-3', Dlx3(R) 5'-ACTCAGGTTCTGTGCGTGAT-3', p63 α (F) 5'-GTCTCCATCTTCATATGGTAAC-3', p63 α (R) 5'-CACACTGACTGTAGAGCA-3', p63 β (F) 5'-GTCTCCATCTTCATATGGTAAC-3', p63 β (R) 5'-CTTGCCAAATCCTGACAATGCTGC-3', p63 γ (F) 5'-GAGGATAGCATCAGAAAACAGCAAG-3' and p63 γ (R) 5'-CTCCACAAGCTCATTCCTGAAGC-3'. For normalization: Cyclophilin(F) 5'-ATCACCA-TTGCTGACTGTGG-3', Cyclophilin(R) 5'-ACTCTGCAATCCAGC-TAGGC-3', GAPDH(F) 5'-GTCTCCATCTTCATATGGTAA-3' and GAPDH(R) 5'-CCACAGTCCATGCCATCACT-3' were used.

RESULTS AND DISCUSSION

The existence of malformations caused by either *p63* or *DLX3* gene mutations that translate to partially overlapping phenotypes suggests that these genes are transcriptional effectors in common signaling cascades regulating epidermal development. The severity of the phenotype in *p63*-null mice suggests that it is a crucial upstream regulator of these signaling pathways (Mills et al., 1999; Yang et al., 1999). Detailed analysis of the *Dlx3* proximal promoter region revealed a sequence with two *p63*-like overlapping binding sites immediately upstream of the CCAAT box, located from -89 to -80 bp (site 1) and from -84 to -75 bp (site 2) of the transcriptional start site (Park and Morasso, 1999). Because the expression patterns of *p63* and *Dlx3* overlap throughout embryonic development (Morasso and Radoja, 2005), we proceeded to test the ability of different *p63* isoforms to transactivate the *Dlx3* promoter. The Dlx3_{-117/+60} construct, which contains the two overlapping sites, was transiently transfected into primary mouse keratinocytes in either the absence or presence of expression vectors encoding TAp63 α , TAp63 β , TAp63 γ , Δ Np63 α , Δ Np63 β or Δ Np63 γ . The TA isoforms activated the Dlx3_{-117/+60} promoter at a magnitude of twelve-, three- and seven-

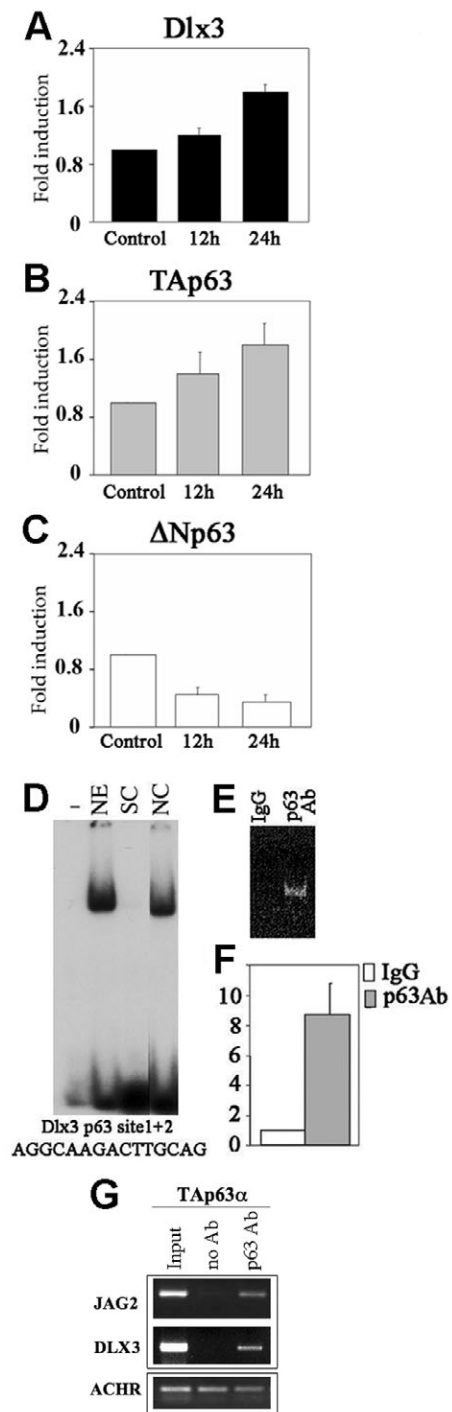


Fig. 2. *p63* and *Dlx3* expression in primary mouse keratinocytes cultured in vitro. *Dlx3* (A) and *TAp63* (B) mRNAs were induced, and Δ Np63 (C) mRNA was downregulated after 12- and 24-hours of high-[Ca²⁺] treatment. (D-F) *p63* binds to the *Dlx3* promoter region in vitro and in vivo. (D) EMSA assay performed with a DNA fragment that included the *Dlx3* *p63*-binding site 1 and 2 using nuclear extract from primary keratinocytes (NE), and in the presence of 100 M excess of specific (SC) and nonspecific (NC) competitors. (E-F) ChIP analysis on mouse keratinocytes with either control IgG or *p63* antibody (*p63* Ab) on the region of the *Dlx3* promoter containing the *p63*-binding sites by regular (E) and real-time (F) PCR. (G) ChIP analysis on TAp63 α -transfected H1299 cells with no antibody (no Ab) or *p63* antibody on DLX3 and JAG2 promoters. ACHR was used as a control.

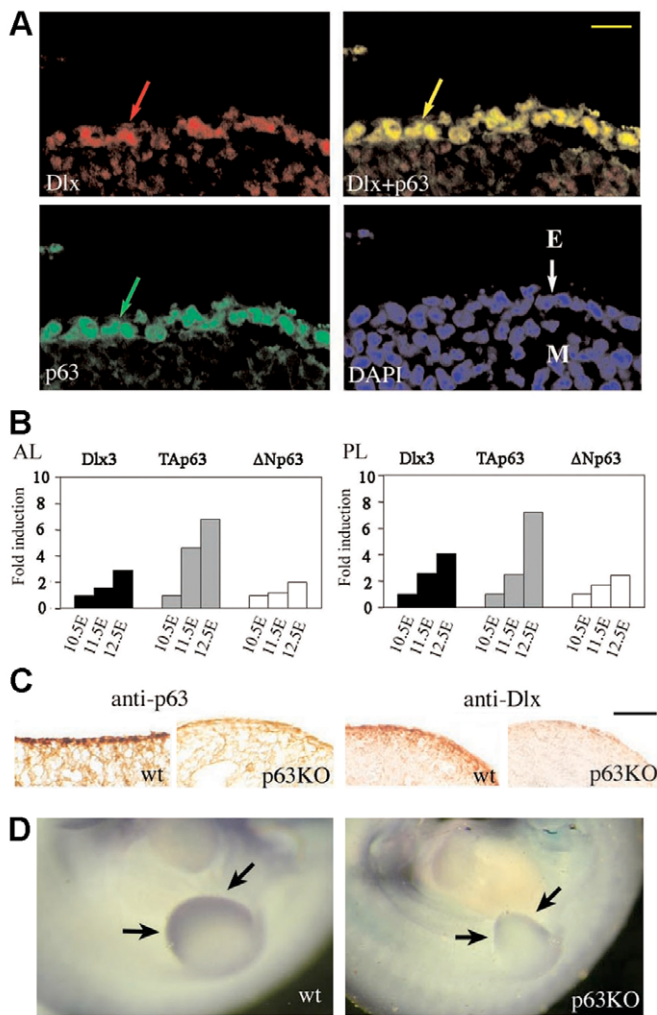


Fig. 3. p63 and Dlx3 colocalize in the embryonic ectoderm, and Dlx3 expression is downregulated in p63-KO embryos. (A) Histochemistry with anti-distal-less (pan-anti-Dlx; red) and p63 (green) antibodies on the dorsal forelimb ectoderm of E10.5 wild-type embryos (merge of Dlx and p63 expression is yellow). DAPI staining is also shown (blue). Arrows indicate the dlx-p63 double-positive nucleus. (B) Relative mRNA abundance, determined by real-time PCR, for *Dlx3*, *TAp63* and *ΔNp63* in the anterior limb (AL) and posterior limb (PL) of E10.5, E11.5 and E12.5 wild-type embryos. The relative abundance is expressed as fold-induction relative to the value at E10.5 (set at 1). (C) Histochemistry with anti-p63 and anti-Dlx on E10.5 wild-type and p63-KO limb-bud ectoderm. (D) Whole-mount in situ hybridization on wild-type and p63-KO E10.5 embryos with a *Dlx3* antisense probe. Arrows indicate limb buds. M, mesoderm; E, ectoderm. Scale bars: 10 μ M in A and 40 μ M in C. wt, wild type.

fold, respectively, compared to normal activation, whereas the exogenous expression of the Δ N isoforms resulted in a two- to four-fold greater transactivation compared with wild type (Fig. 1A). These effects are specific for p63 isoforms, because p53 did not transactivate the *Dlx3* promoter (data not shown).

In order to test each overlapping site (p63-binding-site 1 and p63-binding-site 2, Fig. 1B), we compared the activities of the *Dlx3*_{-117/+60}, *Dlx3*_{-117/+60} p63M1 (mutated p63 site 1) and *Dlx3*_{-117/+60} p63M2 (mutated p63 site 2) constructs (Fig. 1B) by co-transfection performed in the presence of the TA and Δ Np63 isoforms (Fig. 1C,D). Our results indicate that TAp63 α induction of

Dlx3 is mediated through either of the two p63 sites, whereas TAp63 γ required an intact regulatory region. These results are not cell-type specific, because a similar profile was obtained using Saos-2 cells (Fig. 1E).

Since *Dlx3* is induced in keratinocytes cultured in 0.12 mM Ca²⁺ (Park and Morasso, 1999), we compared the endogenous expression of the p63 isoforms with *Dlx3* after 12- and 24-hours of 0.12 mM Ca²⁺ treatment (Fig. 2A-C). The real-time PCR results showed a correlation between the upregulation of *Dlx3* and *TAp63* and the downregulation of Δ Np63 mRNAs associated by Ca²⁺-induced differentiation, and are consistent with the recent report by King et al. (King et al., 2006). The specificity of the PCR products was corroborated by sequencing (data not shown). TAp63 proteins are found in normal adult tissues (Nylander et al., 2002) and during mouse embryonic development (Koster et al., 2004) (also our own data). Findings of Koster et al. (Koster et al., 2004) support a role for TAp63 as a molecular switch for the initiation of epithelial stratification. Our findings support a working model in which, once transactivated by TAp63 α , *Dlx3* will in turn regulate the expression of terminal differentiation markers (Morasso et al., 1996).

In order to demonstrate direct binding to the p63 region in the *Dlx3* promoter, we performed EMSA with a fragment comprising -89 to -75 bp (*Dlx3* p63 sites 1 and 2) using nuclear extracts from primary keratinocytes (NE). A shift was detected (Fig. 2D, lane NE), and the complexes were competed with a specific competitor (Fig. 2D, lane SC), but not with a nonspecific DNA competitor (Fig. 2D, lane NC). We next evaluated whether p63 bound this region of the *Dlx3* promoter in vivo. In mouse keratinocytes, ChIP experiments were performed with a p63-specific antibody and analyzed by regular PCR (Fig. 2E) and real-time PCR (Fig. 2F). The data shows that p63 specifically binds to the *Dlx3* promoter in vivo (eightfold higher than with IgG control). Moreover, ChIP experiments on TAp63 α -transfected H1299 cells, which are devoid of p63, demonstrated the direct binding of TAp63 α to the *Dlx3* promoter (Fig. 2G).

To further explore the relationship of *Dlx3* and p63 in vivo, we analyzed their colocalization by immunofluorescence on E10.5 embryonic forelimb sections with anti-p63 and anti-distal-less antibodies. The latter reagent recognizes *Dlx3* in the limb ectoderm. p63 and *Dlx3* immunoreactivity were found to colocalize in the same nuclei (Fig. 3A). Comparison of the expression of *Dlx3*, *TAp63* and Δ Np63 in the limbs at embryonic stages E10.5, E11.5 and E12.5 was performed by real-time PCR. Between E11.5 and E12.5, the relative abundance of both *TAp63* and *Dlx3* mRNA increased from three- to eight-fold relative to their expression at E10.5 in the anterior (AL) and posterior limbs (PL) (Fig. 3B), whereas expression of Δ Np63 was only moderately increased. These results for p63 in the limb ectoderm of embryos at E10.5-E12.5 are in agreement with reported data (Koster et al., 2004) and show for the first time that a good correlation is observed between the expressions of *Dlx3* and *TAp63*.

To provide further evidence that p63 is an upstream regulator of *Dlx3*, we studied the effect of p63 ablation on *Dlx3* mRNA and protein expression in the *Brdm2* p63 KO mice (Mills et al., 1999) (Fig. 3C-D). As assessed by immunohistochemistry, the abundance of *Dlx3* protein was significantly reduced in p63-KO limb ectoderm (Fig. 3C). Analysis by whole-mount in situ hybridization with a *Dlx3* antisense probe demonstrated that the absence of p63 led to a downregulation of *Dlx3* (Fig. 3D).

We next studied the clinical relevance of p63-mediated regulation of *Dlx3* expression. We examined the transcriptional activity of the *Dlx3* promoter in the presence of p63 mutants causative of human AEC, EEC, SHFM or LMS (Fig. 4A) in the H1299 cell line. The p63 AEC mutants - L518V, L518F and Q540L - are all point

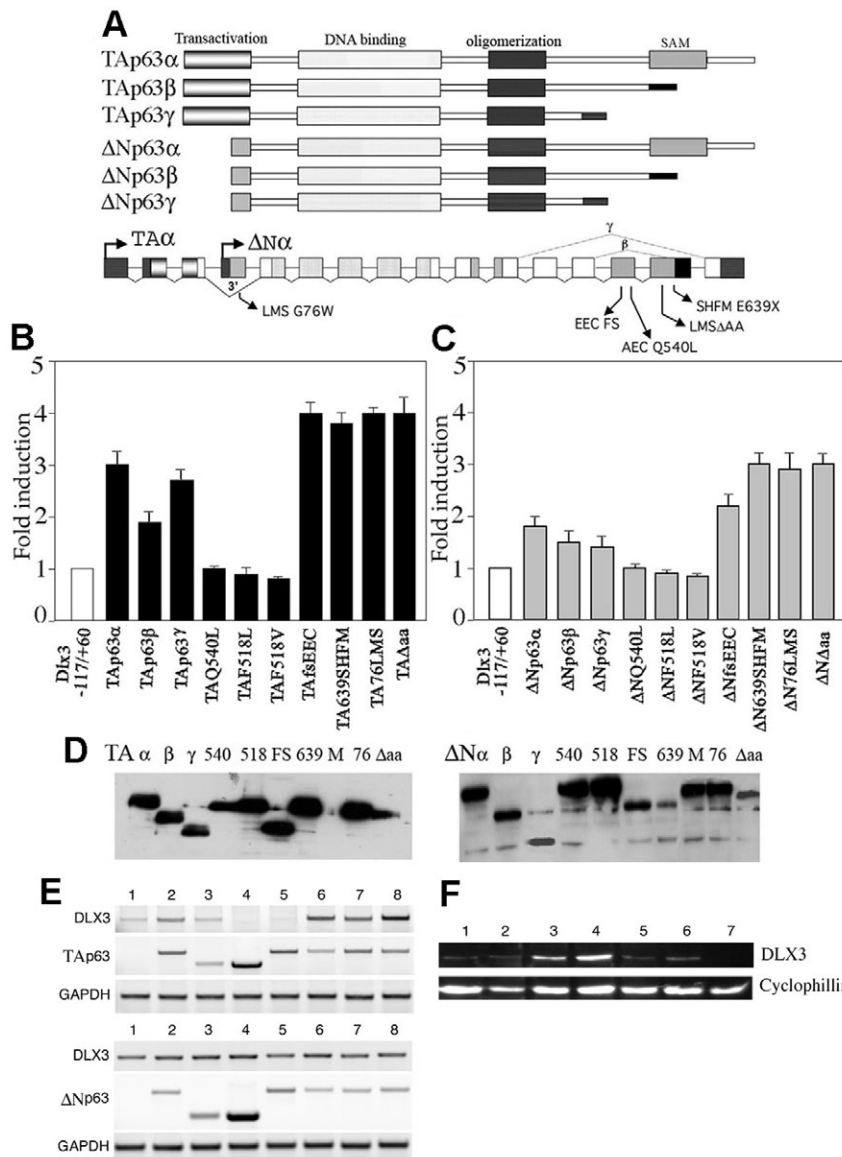


Fig. 4. Differential ability of p63-mutant proteins to transactivate Dlx3. (A) Schematic representation of p63 transcripts and mutations. (B-C) Transcriptional regulation of the *Dlx3* promoter by TAp63 (B) and Δ Np63 (C) wild-type and mutant proteins. H1299 cells were co-transfected with the *Dlx3* reporter plasmid and expression vectors for TAp63- and Δ Np63-mutant isoforms. The basal activity of the reporter was set to 1. Each histogram bar represents the mean of three independent transfections. Standard deviations are indicated. (D) Proteins were corroborated by western blot analysis with anti-p63 4A4 antibody (Santa Cruz). (E) The level of endogenous *Dlx3* mRNA upon transfection with p63 mutants in Saos-2 cells. Top panel, lanes: 1, mock; 2, TAp63 α ; 3, TAp63 β ; 4, TAp63 γ ; 5, TAp63F518V-AEC; 6, TAp63FS-EEC; 7, TAp63E639X-SHFM; and 8, TAp63 Δ AA-LMS. Bottom panel, lanes: 1, mock; 2, Δ Np63 α ; 3, Δ Np63 β ; 4, Δ Np63 γ ; 5, Δ Np63F518V-AEC; 6, Δ Np63FS-EEC; 7, Δ Np63E639X-SHFM; and 8, Δ Np63 Δ AA-LMS. GAPDH was used for normalization. (F) The level of endogenous *Dlx3* mRNA upon transfection with TAp63 α and TAp63 γ in HaCaT cells. Lanes: 1, mock; 2, TAp63 α 0.5 μ g; 3, TAp63 α 2 μ g; 4, TAp63 α 4 μ g; 5, TAp63 γ 0.5 μ g; 6, TAp63 γ 2 μ g; 7, TAp63 γ 4 μ g. Cyclophilin was used for normalization.

substitutions within the SAM domain, present only in TAp63 α and Δ Np63 α . The p63 α FS mutant contains a mutation found in EEC that generates a frameshift at amino acid 525, which leads to a premature stop. The E639X mutation, in exon 14, was isolated in a SHFM patient, whereas the 2-bp deletion, in exon 14 (Δ AA), was present in one family with LMS. The G76W mutation, in exon 3, was isolated from a LMS patient and affects all p63 isoforms. Co-transfection experiments were performed with expression vectors encoding TAp63 and Δ Np63 mutants. As shown in Fig. 4B and 4C, the AEC mutants failed to yield a significant level of reporter-gene expression despite having intact amino terminal and DNA-binding domains. These results point to a crucial role of the C-terminus of p63 α on the regulation of *Dlx3* transcription. The LMS-derived mutants G76W and Δ AA; as well as the FSEEC mutant, SHFM-derived E639X TA and Δ N proteins; showed a similar mode of regulation of *Dlx3*-promoter activity compared with the corresponding wild-type proteins (Fig. 4B,C). A similar profile, albeit with a lower amount of induction, was obtained upon transfection in HaCaT cells (data not shown). The level of expression of the mutant proteins was corroborated by immunoblot analysis with anti-p63 antibody (Fig. 4D).

The hampered ability of the AEC mutants, as well as the partial overlapping phenotypes of specific malformations caused by p63 (i.e. AEC) or *Dlx3* (i.e. TDO) gene mutations, suggest that *Dlx3* misregulation is involved in aspects of the pathogenesis of AEC. AEC is characterized by ectodermal dysplasia, ankyloblepharon and cleft lip with cleft palate, and by the lack of limb involvement. The absence of limb defects in AEC may reflect the possibility that a putative role of *Dlx3* in the limb is compensated for by other *Dlx* proteins (Panganiban and Rubenstein, 2002; Morasso and Radoja, 2005).

To determine the modulation of endogenous *Dlx3* by p63, we used Saos-2 cells to express wild-type and mutant p63 (Fig. 4E). Interestingly, TAp63 α increased *Dlx3* expression (Fig. 4E, lane 2), and this effect was shared, although to different extents, with the EEC, SHFM and LMS mutants (Fig. 4E; lanes 6, 7 and 8). TAp63 β did not alter *Dlx3* levels (Fig. 4E, lane 3). Remarkably, an AEC mutant (Fig. 4E, lane 5) and the TAp63 γ isoform abolished *Dlx3* expression. Transfections of Δ Np63 isoforms, both wild-type and mutant, had no significant effect on *Dlx3* transcription (Fig. 4E, bottom panel). TAp63 α and TAp63 γ are potent transactivators (Barbieri and Pietenpol, 2006) and, in our in vitro studies, both isoforms were able to transactivate *Dlx3* at similar levels. However,

whereas TAp63 α induction was mediated through either of the two p63 sites in the *Dlx3* promoter, TAp63 γ required an intact regulatory region. Surprisingly, a slightly different outcome was obtained for the endogenous *Dlx3* regulation, where upregulation of *Dlx3* was detected with TAp63 α and a complete downregulation was found when expressing TAp63 γ in Soas-2 cells. To determine if these results could be attributed to cell context, the experiments were also performed in HaCaT cells. In these cells, overexpression of TAp63 α showed upregulation of endogenous *Dlx3*, whereas TAp63 γ once again caused a complete downregulation of *Dlx3* expression (Fig. 4F). These differences might be attributed to the specific p63RE-CCAAT-box chromatin architecture. An important feature of the *Dlx3* promoter is that the overlapping p63-binding sites are in close proximity to CCAAT box that binds NF-Y in keratinocytes (Park and Morasso, 1999). NF-Y is a general promoter organizer that pre-sets chromatin structure locally. A recent report shows that p63 α regulates the transcription of the *hsp70* gene through interactions with NF-Y (Wu et al., 2005). Although we have not determined the significance of NF-Y and p63 interactions on *Dlx3* transcriptional regulation, it might be proposed that there is a dual role for the overlapping p63-binding sites, and that *Dlx3* will be transcriptionally active or repressed depending on the specific p63 isoform bound to the promoter, on which of the p63 sites is occupied and on interactions with NF-Y CCAAT binding factor.

Dlx3 and p63 both function as part of a complex series of cascades that ultimately lead to the formation of ectoderm-derived organs. Unraveling the function of each protein at specific times of embryonic development will prove to be complex because of the differential expression of the p63 isoforms in distinct tissues (Nylander et al., 2002) and the cross-regulation with other developmentally relevant signaling pathways [i.e. FGF, BMP, and Notch (Laurikkala et al., 2006; Nguyen et al., 2006)]. The characterization of p63 target genes promises to improve our knowledge of the signaling cascades that are directly involved in normal ectodermal development. In summary, our study proves a functional relationship between p63 and *Dlx3*, with *Dlx3* demonstrated to be a direct target of p63. The findings also provide evidence that the misregulation of *Dlx3* is involved in the pathogenesis of p63 molecular lesions in AEC.

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References

- Acamora, D., Merlo, G. R., Paleari, L., Zerega, B., Postiglione, M. P., Mantero, S., Bober, E., Barbieri, O., Simeone, A. and Levi, G. (1999). Craniofacial, vestibular and bone defects in mice lacking the Distal-less-related gene *Dlx5*. *Development* **126**, 3795-3809.
- Barbieri, C. E. and Pietenpol, J. A. (2006). P63 and epithelial biology. *Exp. Cell Res.* **312**, 695-706.
- Calabrò, V., Mansueti, G., Parisi, T., Vivo, M., Calogero, R. A. and La Mantia, G. (2002). The human MDM2 oncoprotein increases the transcriptional activity and the protein level of the p53 homolog p63. *J. Biol. Chem.* **277**, 2674-2681.
- Caretti, G., Di Padova, M., Micales, B., Lyons, G. E. and Sartorelli, V. (2004). The Polycomb Ezh2 methyltransferase regulates muscle gene expression and skeletal muscle differentiation. *Genes Dev.* **18**, 2627-2638.
- Ghioni, P., Bolognese, F., Duijf, P., Van Bokhoven, H., Mantovani, R. and Guerrini, L. (2002). Complex transcriptional effects of p63 isoforms: identification of novel activation and repression domains. *Mol. Cell Biol.* **22**, 8659-8668.
- Hassan, M. Q., Javed, A., Morasso, M. I., Karlin, J., Montecino, M., van Wijnen, A. J., Stein, G. S., Stein, J. L. and Lian, J. B. (2004). Distal-less 3 (*Dlx3*) is expressed in bone: temporal recruitment of *Mx2*, *Dlx3* and *Dlx5* homeodomain proteins to chromatin for transcriptional regulation of osteoblast growth and differentiation. *Mol. Cell Biol.* **24**, 9248-9261.
- King, K. E., Ponnamperna, R. M., Yamashita, T., Tokino, T., Lee, L. A., Young, M. F. and Weinberg, W. (2003). DeltaNp63alpha functions as both a positive and a negative transcriptional regulator and blocks in vitro differentiation of murine keratinocytes. *Oncogene* **22**, 3635-3644.
- King, K. E., Ponnamperna, R. M., Gerdes, M. J., Tokino, T., Yamashita, T., Baker, C. C. and Weinberg, W. C. (2006). Unique domain functions of p63 isoforms that differentially regulate distinct aspects of epidermal homeostasis. *Carcinogenesis* **27**, 53-63.
- Koster, M. I. and Roop, D. R. (2004). Transgenic mouse models provide new insights into the role of p63 in epidermal development. *Cell Cycle* **3**, 411-413.
- Koster, M. I., Kim, S., Mills, A., DeMayo, F. J. and Roop, D. R. (2004). p63 is the molecular switch for initiation of an epithelial stratification program. *Genes Dev.* **18**, 126-131.
- Laurikkala, J., Mikkola, M. L., James, M., Tummers, M., Mills, A. A. and Thesleff, I. (2006). P63 regulates multiple signaling pathways required for ectodermal organogenesis and differentiation. *Development* **133**, 1553-1563.
- McGrath, J. A., Duijf, P. H. G., Doetsch, V., Irvine, A. D., de Wall, R., Vanmolkot, K. R. J., Wessagowit, V., Kelly, A., Atherton, D. J., Griffith, W. A. D. et al. (2001). Hay-Wells syndrome is caused by heterozygous missense mutations in the SAM domain of p63. *Hum. Mol. Genet.* **10**, 1-9.
- Merlo, G. R., Beverdam, A. and Levi, G. (2003). *Dlx* genes in craniofacial and limb morphogenesis. In: *Murine Homeobox Gene Control of Embryonic Patterning and Organogenesis* (ed. T. Lufkin). *Adv. Develop. Biol. Biochem.* **13**, 107-132.
- Mills, A. A., Zheng, B., Wang, X. J., Vogel, H., Roop, R. and Bradley, A. (1999). p63 is a p53 homologue required for limb and epidermal morphogenesis. *Nature* **398**, 708-713.
- Morasso, M. I. and Radoja, N. (2005). *Dlx* genes, p63, and ectodermal dysplasias. *Birth Defect Res.* **75**, 163-171.
- Morasso, M. I., Mahon, K. and Sargent, T. D. (1995). A *Xenopus* distal-less gene in transgenic mice: conserved regulation of distal limb epidermis and other sites of epithelial-mesenchymal interaction. *Proc. Natl. Acad. Sci. USA* **92**, 3968-3972.
- Morasso, M. I., Markova, N. G. and Sargent, T. D. (1996). Regulation of epidermal differentiation by a *Distal-less* homeodomain gene. *J. Cell Biol.* **135**, 1879-1887.
- Morasso, M. I., Grinberg, A., Robinson, G., Sargent, T. D. and Mahon, K. A. (1999). Placental failure in mice lacking the homeobox gene *Dlx3*. *Proc. Natl. Acad. Sci. USA* **96**, 162-167.
- Nguyen, B. C., Lefort, K., Mandinova, A., Antonini, D., Devgan, V., Della Gatta, G., Koster, M., Zhang, Z., Wang, J., Tommasi di Vignano, A. et al. (2006). Cross-regulation between Notch and p63 in keratinocyte commitment to differentiation. *Genes Dev.* **20**, 1028-1042.
- Nylander, K., Vojtesek, B., Nenutil, R., Lindgren, B., Roos, G., Zhanxiang, W., Sjöstrom, B., Dahlqvist, A. and Coates, P. J. (2002). Differential expression of p63 isoforms in normal tissues and neoplastic cells. *J. Pathol.* **198**, 417-427.
- Osada, M., Park, H. L., Nagakawa, Y., Yamashita, K., Fomenkov, A., Kim, M. S., Wu, G., Nomoto, S., Trink, B. and Sidransky, D. (2005). Differential recognition of response elements determines target gene specificity for p53 and p63. *Mol. Cell Biol.* **25**, 6077-6089.
- Panganiban, G. and Rubenstein, J. L. R. (2002). Developmental functions of the *Distal-less/Dlx* homeobox genes. *Development* **129**, 4371-4386.
- Park, G. T. and Morasso, M. I. (1999). Regulation of the *Dlx3* homeobox gene upon differentiation of mouse keratinocytes. *J. Biol. Chem.* **274**, 26599-26608.
- Pispa, J. and Thesleff, I. (2003). Mechanisms of ectodermal organogenesis. *Dev. Biol.* **262**, 195-205.
- Price, J. A., Bowden, D. W., Wright, J. T., Pettenati, M. J. and Hart, T. C. (1998). Identification of a mutation in *DLX3* associated with tricho-dento-osseous (TDO) syndrome. *Hum. Mol. Genet.* **7**, 563-569.
- Priolo, M. and Lagana, C. (2001). Ectodermal dysplasias: a new clinical-genetic classification. *J. Med. Genet.* **38**, 579-585.
- Qiao, F. and Bowie, J. U. (2005). The many faces of SAM. *Sci. STKE* **286**, re7: 1-10.
- van Bokhoven, H. and Brunner, H. G. (2002). Splitting p63. *Am. J. Hum. Genet.* **71**, 1-13.
- van Bokhoven, H., Hamel, B. C., Bamshad, M., Sangiorgi, E., Gurrieri, F., Duijf, P., Vanmolkot, K., van Beusekom, E., van Beersum, S., Celli, J. et al. (2001). P63 Gene mutations in EEC syndrome, limb-mammary syndrome, and isolated split hand-split foot malformation suggest a genotype-phenotype correlation. *Am. J. Hum. Genet.* **69**, 481-492.
- Wu, G., Osada, M., Guo, Z., Fomenkov, A., Begum, S., Zhao, M., Upadhyay, S., Xing, M., Wu, F., Moon, C. et al. (2005). DeltaNp63alpha up-regulates the *Hsp70* gene in human cancer. *Cancer Res.* **65**, 758-766.
- Yang, A., Kaghad, M., Wang, Y., Gillett, E., Fleming, M. D., Dotsch, V., Andrews, N. C., Caput, D. and McKeon, F. (1998). p63, a p53 homolog at 3q27-29, encodes multiple products with transactivating, death-inducing, and dominant-negative activities. *Mol. Cell.* **2**, 305-316.
- Yang, A., Schweitzer, R., Sun, D., Kaghad, M., Walker, R. T., Bronson, C., Tabin, A., Sharpe, D., Caput, D., Crum, C. and McKeon, F. (1999). p63 is essential for regenerative proliferation in limb, craniofacial and epithelial development. *Nature* **398**, 714-718.