**Abstract**

TLE1 is a Groucho-related transcriptional repressor protein that exerts survival and antiapoptotic function in several cellular systems and has been implicated in the pathogenesis of cancer. In the present study, we found that TLE1 is a regulator of anoikis in normal mammary epithelial and breast carcinoma cells. The induction of apoptosis following loss of cell attachment to the extracellular matrix (anoikis) in untransformed mammary epithelial MCF10A cells was associated with significant downregulation of TLE1 expression. Forced expression of exogenous TLE1 in these cells promoted resistance to anoikis. In breast cancer cells, TLE1 expression was significantly upregulated following detachment from the extracellular matrix. Genetic manipulation of TLE1 expression via overexpression and downregulation approaches indicated that TLE1 promotes the anoikis resistance and anchorage-independent growth of breast carcinoma cells. Mechanistically, we show that TLE1 inhibits the Bit1 anoikis pathway by reducing the formation of the proapoptotic Bit1-AES complex in part through sequestration of AES in the nucleus. The mitochondrial release of Bit1 during anoikis as well as exogenous expression of the cytoplasmic localized Bit1 or its cell death domain induced cytoplasmic translocation and degradation of nuclear TLE1 protein. These findings indicate a novel role for TLE1 in the maintenance of anoikis resistance in breast cancer cells. This conclusion is supported by an immunohistochemical analysis of a breast cancer tissue array illustrating that TLE1 is selectively upregulated in invasive breast tumors relative to noninvasive ductal carcinoma in situ and normal mammary epithelial tissues. *Mol Cancer Res; 10*(11); 1-14. ©2012 AACR.

**Introduction**

The Groucho (Gro)/TLE family of transcriptional corepressors represents a group of gene regulators that influence the transcriptional activity of a wide range of genes (1). As corepressors, Gro proteins do not bind to DNA directly, but instead bind to other DNA-binding transcription factors to form large multiprotein complexes (2). The binding of Gro to such a regulatory complex most often results in the decreased translation of the target gene. For instance, in Wnt gene regulation, binding of Gro to the transcription factor LEF results in the displacement of the Wnt-activator β-catenin and in decreased translation of the Wnt gene (3). This is accompanied by the recruitment of histone deacetylase to the gene and the subsequent removal of acetyl groups from the DNA-bound histones, which results in further gene silencing.

Two groups of the Gro/TLE protein family have been identified. The first group, TLE1–TLE4, shares homologous structural motifs including an N-terminal oligomerization domain (Q-domain), a CcN linker, and a C-terminal Histone deacetylase (HDAC)-binding WD domain (1). Normally, TLE1 is required to be in a homotetramer to be active. The second group, which includes the Amino Enhancer of Split (AES), retains the same N-terminal oligomerization domain as the other TLE family members, but importantly lacks the WD domain. Because of this, AES is considered a dominant-negative for TLE1–4 family members, as AES is able to oligomerize and sequester TLE proteins, preventing them from forming homotetramers and repressing gene transcription.

As transcriptional repressors, the Gro/TLE proteins play important roles in several processes including regulation of neurogenesis and a number of developmental processes (4, 5). Recently, data have begun to emerge indicating a prosurvival and/or antiapoptotic role of groucho proteins, TLE1 in particular. Overexpression of TLE1 in chicken embryo fibroblast led to significant growth stimulation and conferred anchorage-independent growth (6). In mature neurons, exogenous expression of TLE1 prevented cell death and apoptosis (7). Our previous studies have shown that TLE1 is antiapoptotic by blocking the apoptosis induced by the release of mitochondrial Bit1 (Bcl2-inhibitor of transcription 1) protein to the cytoplasm (8). Recently, Seo and
colleagues showed that TLE1 functions to inhibit apoptosis induced by doxorubicin in synovial sarcoma cells (9). Taken together, these observations provide a case for the role of TLE1 in promoting cell survival via its antiapoptotic function. It is conceivable that TLE1 may suppress an apoptosis gene transcription program or alternatively it may upregulate a survival-promoting gene transcription program. Consistent with this notion, TLE1 positively regulates Bcl2 expression (8) and ErbB1 and ErbB2 signaling (10), 2 survival pathways that influence tumorigenesis.

TLE1 seems to be a part of the novel integrin-dependent Bit1 apoptotic pathway (8). Following loss of cell attachment, mitochondrial-bound Bit1 protein is released to the cytoplasm and initiates a caspase-independent apoptosis, which is unresponsive to various antiapoptotic treatments including Bcl-2, Bcl-xl, and akt. Importantly, Bit1 is unique among cell death inducers in that it is negatively regulated by integrin engagement and therefore represents a critical pathway in understanding cell death caused by extracellular matrix (ECM) detachment (anoikis). Considering that anoikis resistance is a determinant of cancer metastasis, this also places Bit1 as a regulator of metastasis (11). However, the molecular mechanisms underlying Bit1 anoikis function and its regulation by integrin have not been fully elucidated. Bit1, once released from the mitochondria or ectopically expressed in the cytoplasm, forms a complex with AES, and the Bit1-AES complex represents the proapoptotic component that mediates the effect of Bit1 on apoptosis (8). One of the downstream targets of the Bit1 signaling pathway appears to be the extracellular signal-regulated kinase (Erk; ref. 12). Normal and transformed cells in which Bit1 expression is downregulated exhibit increased Erk activation and such elevated Erk activity contributes in part to the enhanced anoikis resistance of the Bit1 knockdown cells. Recently, we have also shown that the serine/threonine kinase protein kinase D (PKD) acts as an upstream activating kinase that promotes the apoptotic and anoikis function of Bit1 (13).

In light of previous reports documenting the antiapoptotic and growth promoting functions of TLE1, we examined in the present study the effect of TLE1 expression in the survival and anoikis resistance of breast carcinoma cells. We discovered a critical role for TLE1 in suppressing anoikis in breast cancer cells, and that its inhibitory effect on anoikis is in part through circumventing the Bit1-mediated anoikis pathway. Interestingly, the nuclear TLE1 protein is targeted and channeled to degradation in the cytoplasm by Bit1 during anoikis, indicating that turning off the TLE1 transcriptional machinery may represent an important mechanism in the induction of anoikis by Bit1.

Materials and Methods

Cell culture and transfection assays
MCF7, MDA-MB-231, MCF10A, and HEK 293T from American Type Culture Collection were cultured in Dulbecco’s modified Eagle’s medium (DMEM) with glutamine containing 10% FBS, penicillin, and streptomycin. Stable MDA-MD-231 control and DDK-TLE1 pool of cells were generated by transfection with DDK-TLE1 or empty vector construct (Origene) as described previously (11, 13). Briefly, transfections were carried out with Lipofectamine 2000 (Invitrogen) in OPTI-MEM (Invitrogen) according to the manufacturer’s protocol with cells plated 18 hours before transfection. 48 hours after transfection, transfected cells were cultured in the 1,000 μg/mL G418 to select for positive DDK-TLE1–expressing clones. Several G418 resistant control and DDK-TLE1 clones were harvested by ring-cloning, and the level of DDK-TLE1 expression was confirmed by immunoblotting against a specific DDK antibody. Two control clones and 3 positive DDK-TLE1 clones were pooled together to generate the control and DDK-TLE1 pools, respectively. In generating the MCF10A control and GFP-TLE1 pool of cells, the MCF01A cell line was transfected with GFP-TLE1 or empty GFP-tagged vector (Origene) using the Lipofectamine 2000 Plus reagent (Invitrogen), and transfected cells were then selected in the presence of 400 μg/mL G418 to generate the control and GFP-TLE1 pools as described above. In transient transfection assays, the total amount of plasmid used per transfection was normalized with the corresponding empty vector constructs.

Chemical reagents, antibodies, and plasmids
Poly(2-hydroxyethyl methacrylate (Polyhema) and the mouse monoclonal anti-B-actin antibody were obtained from Sigma Chemical Co. The mouse monoclonal anti-FLAG, anti-AES, anti-GFP, and anti-B-actin antibodies were purchased from Sigma. The polyclonal anti-TLE1 antibody was obtained from Abcam. The proteasomal inhibitor MG132 and the caspase inhibitor zVad-fmk were purchased from Calbiochem, whereas the chemical reagent bortezomib and was obtained from LC Laboratories. The anti-caspase-3, anti-cleaved caspase-3, anti-caspase-8, anti-cleaved caspase-8, and anti-PARP antibodies were obtained from Cell Signaling Technology. The anti-c-myc antibody was obtained from Santa Cruz Biotechnology. The mammalian expression vector encoding for mitochondrial Bit1 was generated as described previously (8, 13). The FLAG-TLE1 plasmids were obtained from Origene. The myc-DDK plasmids were provided by Dr. Renwei Chen (University of California, Santa Barbara, California).

siRNA and shRNA transfection
The TLE1-specific siRNAs (186345, 186346, and 110786) and the control, nontargeting siRNAs were purchased from Invitrogen. For transient transfection experiments, MDA-MB-231 cells (2 × 10⁵ cells) were transfected with 25 μmol/L of each siRNA by using the Lipofectamine RNAiMAX transfection reagent (Invitrogen). 48 hours after transfection, cells were harvested and subjected to immunoblotting, anoikis, or soft agar assays as described below.

To generate stable MDA-MB-231 TLE1 knockdown and control pools, MDA-MB-231 parental cells were transfected with pRS vector containing the short hairpin RNA (shRNA) against TLE1 (Origene) or the nontargeting scrambled shRNA (Origene). 48 hours after transfection, 1 μg/mL...
puromycin (Invitrogen) was added to the medium to select for puromycin-resistant clones. Individual puromycin-resistant clones were screened for TLE1 downregulation by immunoblotting using a specific antibody to TLE1. Three TLE1 knockdown-positive clones were pooled as well 2 control clones were also pooled for further characterization.

**Analysis of apoptosis, anoikis, and cell viability**

Apoptosis was assessed by determining the level of cytosolic nucleosomal fragments with the use of Cell Death Detection ELISA kit (Roche Molecular Biochemicals), according to the manufacturer’s instructions. To assess for anoikis cell death, cells were plated onto Polyhema-coated 96-well plates in complete growth medium containing 0.5% methylcellulose at a density of 1.0 × 10⁴ per well at various time points as previously described (8, 11). Detached cells were then collected and subjected to the Cell Death ELISA apoptosis assay (13). In some experiments, cell viability was quantified by alamar blue staining (Invitrogen) and subsequent fluorescence reading (485 nm excitation wavelength and 520 nm emission wavelength) using the microplate reader.

**Cell proliferation and soft agar assays**

To determine anchorage-dependent growth, cells were plated in a volume of 150 μL at a density of 2,000 cells per well in 96-well plates. At each indicated time, the number of metabolically active cells was measured with the use of MTT assay as described previously (11). Briefly, MTT reagent (Sigma) was dissolved in sterile PBS at 5 mg/mL. Ten microliters of this solution was added to each well in a 96-well plate (1:10 dilution). The plate was then incubated at 37°C for 4 hours. Later, the medium was gently aspirated away and the MTT precipitate was dissolved in 100 μL of a 50% MeOH-50% dimethyl sulfoxide (DMSO) solution. The precipitate was allowed to dissolve at room temperature for 10 minutes with gentle shaking. The resulting 550 nm absorbance was read on a microplate reader (BioTek Instruments). The anchorage-independent growth of cells was measured using the 96-well plate format (14). Briefly, 5,000 cells in 0.3% agar solution were plated onto wells precoated with 0.6% agar in culture medium. The anchorage-independent growth of cells was then quantified by alamar blue staining (Invitrogen) and fluorescence reading at 485 nm excitation wavelength and 520 nm emission wavelength with a microplate plate reader.

**Immunofluorescence microscopy**

Immunofluorescent staining was conducted as previously described (8). Briefly, cultured cells were transfected with FLAG-tagged TLE1 and myc-tagged Bit1 or CDD. Twenty-four hours after transfection, cells were fixed in 4% paraformaldehyde and permeabilized in 0.5% Triton X-100, followed by blocking with 20% FBS. The cells were then stained with anti-Flag (mouse, Sigma), and secondary antibodies (Alexa 488, Molecular Probes). The specimens were mounted with DAPI-containing media (Vectorshield) and analyzed on a confocal microscope (BioRad 1024 equipped with Lasersharp software OS2).

**Protein preparation and Western blotting assays**

Protein preparation and Western blotting were conducted as described previously (11, 13). Briefly, cells were harvested 24 to 48 hours after transfection with various constructs or siRNAs by adding ice-cold NP-40 lysis buffer [1% NP-40; 20 mmol/L Tris-HCL (pH7.4); 150 mmol/L NaCl; 10% glycerol, 2 mmol/L sodium vanadate; 1 mmol/L henylmethylsulfonfly fluoride; 10 μg/mL leupeptin, and 5 μg/mL aprotinin] and incubating at 4°C for 20 minutes. For immunoblot analysis, equal amounts of proteins were resolved on 4% to 20% gradient Tris-glycine gels (Invitrogen) and electrophoretically transferred to nitrocellulose membrane. The membranes were incubated with primary antibodies overnight at 4°C followed by secondary antibodies conjugated with horseradish peroxidase. Membranes were developed using the ECL detection system. Densitometric analysis was conducted using Image J software (13).

**Coimmunoprecipitation assay**

Coimmunoprecipitation assay was conducted as described previously (13). Briefly, transfected cells cultured in attached or detached conditions were harvested by washing once with PBS and resuspended in ice-cold Nonidet P-40 lysis buffer (1% Nonidet P-40, 20 mmol/L Tris-HCL, pH 7.4, 150 mmol/L NaCl, 10% glycerol, 2 mmol/L sodium vanadate, 1 mmol/L phenylmethylsulfonyl fluoride, 10 μg/mL leupeptin, and 5 μg/mL aprotinin) followed by a 20-minute incubation at 4°C. Cell debris were removed by centrifugation. Myc-tagged Bit1 was immunoprecipitated with anti-Myc-agarose conjugate (Abcam) and thoroughly washed with lysis buffer. Bound proteins were resolved by SDS-PAGE, and Western blotting was conducted using anti-AES antibody.

**Subcellular fractionation**

Subcellular fractionation was conducted as described previously (13). Briefly, transfected cells cultured in attached or detached conditions for 12 hours were harvested, washed once with PBS, resuspended in 1 mL of isotonie mitochondrial buffer [250 mmol/L mannitol, 70 mmol/L sucrose, 1 mmol/L EDTA, 10 mmol/L HEPES (pH 7.5)], and homogenized with 40 strokes in a Dounce homogenizer. The lysates were centrifuged at 500 × g for 5 minutes to eliminate nuclei and unbroken cells. The supernatant was further centrifuged at 10,000 × g for 30 minutes at 4°C to isolate the mitochondrial-enriched pellet, which was resuspended in isotonie mitochondrial buffer. Both the cytosolic supernatant and mitochondrial fraction were subjected to SDS-PAGE electrophoresis and immunoblotting. For separation of cytosol and the nuclear fraction, the NE-PER nuclear isolation kit (Pierce, No. 78833) was used and conducted as prescribed by the manufacturer. The protein concentration in the different fractions was quantified using the Bio-Rad protein assay kit with BSA as the standard.
Total RNA extraction and quantitative real-time PCR
Total RNA was extracted from 1.0 × 10^6 cultured cells using the RNeasy kit (Qiagen), quantified by spectrophotometry (NanoDrop 8000, Thermo Scientific). Total RNA was reverse transcribed using the Superscript First-Strand Synthesis Kit for RT-PCR (Invitrogen) as prescribed by the supplier. cDNA was quantified by real-time PCR on the ABI Prism 7900 Sequence Detection System (Applied Biosystems). Human TLE1 forward primer: CCCATATCCTGCTCCTTTTGTG and reverse primer: GGTGTAGGGTGTTGATCTTG were used. PCR was conducted using the SYBR Green PCR Core reagents (Applied Biosystems) based on manufacturer’s instructions. Amplification of the same cDNAs with human GAPDH primers was used for internal normalization.

Human breast tumor array analysis
Breast tumor tissue array slides containing intraductal carcinomas, invasive ductal carcinoma, and matched normal breast epithelial tissues were obtained from US Biomax, Inc. The immunohistochemistry procedure was conducted by Biomax Inc. on 2 tissue microarray slides. As described previously (11), tissue array slides were deparaffinized, hydrated, and subjected to antigen retrieval. The slides were then incubated in 2.5% normal horse serum for 30 minutes at room temperature followed by incubation with the primary antibody (1:100 dilution) for 1 hour at room temperature. The affinity-purified rabbit anti-TLE1 antibody (M-101) was purchased from Santa Cruz Biotechnology, whereas the affinity-purified rabbit anti-Bit1 antibody (HPA012897) was purchased from Sigma (11). Rabbit normal serum was used as negative control antibody to replace the primary antibody on control slide with 1-hour incubation. Tissue array slides were then washed and incubated with ImmPRESS reagent (Vector Laboratories) followed by treatment with peroxidise substrate DAB solution (DAKO Cytomation). Each experimental and control slide were scored for average staining intensity by 2 investigators with no knowledge of the pathologic status of the samples. These investigators scored staining intensity as 0, no staining; 1, low staining; 2, medium staining; or 3, high staining. The intensity of TLE1 or Bit1 staining was further measured as the average staining intensity of the majority of epithelial cells (75% to 100%) in each sample. The average staining was graded as 0, no staining; 1, slight staining; 2, moderate staining; and 3, strong staining.

Statistical analysis
Data are presented as mean (±SE). For Western blot and anoikis assays, experiments were carried out at least 3 times with duplicates. Statistical differences between groups were established at a P value < 0.05 using the 2-tailed Student t test. For breast tumor tissue array analysis, a one-way ANOVA with subsequent post hoc testing using the Tukey–Kramer multiple comparison test was used to compare the average staining intensity of each case type (11). All calculations were done using the NCSS statistical software.

Results
Induction of anoikis in normal mammary epithelial cells is associated with downregulation of TLE1
To address the hypothesis that TLE1 plays a role in the anoikis resistance of breast carcinoma cells, we first examined the effects of TLE1 on anoikis of normal mammary epithelial cell line MCF10A. As previously described (15, 16), forced suspension of MCF10A cells resulted in a significant induction of apoptosis (Fig. 1A) which was associated with the downregulation of TLE1 expression both at the mRNA (Fig. 1B) and protein levels (Fig. 1C and D). To examine the significance of TLE1 suppression in MCF10A following loss of attachment, we stably expressed exogenous GFP-tagged TLE1 or the GFP alone in MCF10A cells. Exogenous TLE1 expression was confirmed by immunoblotting using anti-GFP antibody (Fig. 1E) and anti-TLE1 antibody (Supplementary Fig. S1). Immunofluorescence imaging also confirmed the nuclear localization of GFP-TLE1 (Fig. 1F) as previously described (1, 17). While the control and TLE1-expressing pools displayed similar levels of basal apoptosis in adherent conditions, the TLE1 expressing pool of cells showed significantly decreased apoptosis in suspension culture as compared with control cells (Fig. 1G). These findings indicate that TLE1 promotes survival of normal mammary epithelial cells in response to detachment from the ECM, and its downregulation following loss of cell attachment may contribute to induction of anoikis cell death program in breast epithelial cells.

Anoikis resistance of breast carcinoma cells is associated with increased expression of TLE1
We next examined the role of TLE1 in the anoikis resistance of human breast cancer cells. Consistent with the previous finding (18), the highly aggressive MDA-MB-231 cells showed significant resistance to anoikis as evidenced by the lack of apoptosis following a 24-hour culture in suspension (Fig. 2A). In marked contrast to that found in the highly anoikis sensitive MCF10A cells, TLE1 expression at both the mRNA (Fig. 2B) and protein (Fig. 2C and D) levels was significantly increased in the MDA-MB-231 cells following 24 hours in suspension culture as compared with the level in attached cells. Similar results were also obtained in another anoikis-resistant mammary carcinoma MCF7 cells (Fig. 2A), which exhibited significantly high levels of TLE1 protein following culture in suspension for 24 hours (Fig. 2C and D). These findings show that loss of cell-substrate attachment triggers induction of TLE1 expression in breast cancer cells.

TLE1 is a regulator of anoikis resistance in breast cancer cells
To examine the role of TLE1 upregulation in the anoikis resistance of breast carcinoma cells, we downregulated TLE1 expression in MDA-MB-231 cells via siRNAs. Two TLE1-specific siRNAs #1(186345) and #3(117086) showed a significant 50% to 70% downregulation of TLE1 expression (Fig. 3A). Suppression of TLE1 expression by these 2 specific TLE1-siRNAs did not induce a change in the morphology.
Supplementary Fig. S2A) or anchorage-dependent growth of MDA-MB-231 cells (Supplementary Fig. S2B). In contrast, culturing cells transfected with the TLE1 siRNAs in suspension showed reduced growth (Fig. 3B) and increased apoptosis (Fig. 3C) as compared with the parental and control siRNA–treated cells. It is noteworthy that no significant difference in basal apoptosis was observed between the TLE1-siRNA and control siRNA–treated cells in adherent conditions (Fig. 3C). The observed enhanced anoikis sensitivity is specific to TLE1 downregulation as forced overexpression of exogenous TLE1 attenuated the increased anoikis sensitivity of TLE1-siRNA–treated cells (Supplementary Fig. S3A and S3B). To complement the results from the transient transfection studies, we also generated stable TLE1 knockdown clones by infecting MDA-MB-231 cells with lentiviral expression vectors for TLE1-specific shRNAs.

Figure 1. TLE1 expression is downregulated during anoikis and expression of exogenous TLE1 inhibits anoikis in nontransformed breast epithelial cell line MCF10A. A to D, MCF10A cells were plated onto a polyhema-coated or uncoated tissue culture plates for 24 h in culture. In A, the cells were harvested and analyzed for apoptosis using the Cell Death ELISA. B, harvested cells were subjected to total RNA isolation and TLE1 mRNA expression levels were quantified by reverse transcription and real-time PCR analysis. Data represent relative values, normalized to the levels of GAPDH mRNA (see Materials and Methods). C, harvested cells were subjected to total cell lysate isolation, SDS-PAGE, and immunoblotting against specific antibodies to TLE1 and β-actin. D, the TLE1/β-actin ratios were determined by densitometry analysis of immunoblots in C from 3 independent experiments. E and F, expression of exogenous GFP-TLE1 was confirmed by immunoblotting (E) using the total cell lysates derived from control and TLE1-overexpressing pools against specific antibodies to GFP and β-actin and immunofluorescent imaging (F). G, control and TLE1-overexpressing cells were plated onto a polyhema-coated or uncoated tissue culture plates. Following 24 hours in culture, cells were harvested and analyzed for apoptosis by measuring the amount of DNA histone fragments (Cell Death Elisa). In A, B, D, and G, 3 independent experiments were conducted in triplicate, *, P < 0.05 by Student t-test.

(Continued)
or control shRNAs. As shown in Fig. 3D, TLE1 expression was reduced by 70% in a pool of TLE1 shRNA clones as compared with control shRNA pool of clones. The control shRNA and TLE1 shRNA pools exhibited similar growth kinetics (Supplementary Fig. S2C) and basal apoptosis (Fig. 3E) in monolayer culture. In contrast, when cultured without anchorage, the TLE1 shRNA pool displayed a significantly more apoptosis than the control shRNA pool (Fig. 3E). Consistent with the enhanced anoikis sensitivity induced by TLE1 downregulation, both transient and stable suppression of TLE1 resulted in decreased growth in soft agar (Supplementary Fig. S4A). TLE1 knockdown cells exhibited fewer and smaller colonies compared with the control cells (Supplementary Fig. S4B and S4C). Taken together, these results indicate that suppression of TLE1 induces detachment-induced apoptosis and attenuates the anchorage-independent growth of breast carcinoma cells.

To confirm the results obtained with the downregulation approaches, we stably overexpressed TLE1 in MDA-MB-231 cells. Several TLE1-overexpressing and vector-transfected clones were selected and combined to generate the stable TLE1 and control pool, respectively (Fig. 3F). In monolayer culture, no significant differences in the growth rate (Supplementary Fig. S5A) and basal apoptosis (Fig. 3G) were observed between the control and TLE1-overexpressing pools. After a prolonged culture in suspension for 72 hours, the control pool underwent significant apoptosis, whereas the stable TLE1 pool exhibited significantly low degree of apoptosis (Fig. 3G). Consistent with this enhanced anoikis resistance, the stable TLE1 pool showed greater ability to grow in soft agar than the control pool (Supplementary Fig. S5B). These findings indicate that TLE1 provides protection from apoptosis under anchorage-independent conditions.

**TLE1 inhibits Bit1-induced anoikis**

On the basis of our previous finding that TLE1 counteracts Bit1 apoptosis function (8), we explored the notion that TLE1 is a target of mitochondrial Bit1 when this protein causes anoikis. To address this possibility, the highly aggressive and anoikis resistant MDA-MB-231 cells were transfected with the C-terminally myc-tagged Bit1, which associates with mitochondria or vector construct. The cells were then grown in suspension or left attached in culture dish. Consistent with previous findings (8, 11, 13), exogenous expression of mitochondrial Bit1 dose dependently enhanced apoptosis following culture in suspension for 24 hours (Fig. 4A). In contrast, the
Bit1- and control-transfected cells had the same level of spontaneous apoptosis when grown attached to a culture dish. As described previously (8), the Bit1-induced anoikis was independent of caspase activation as evidenced by the inability of the pan-caspase inhibitor z-Vad-fmk to block the Bit1 anoikis function (Fig. 4B). In addition, we also found no significant increase in the 85 kD PARP apoptotic fragment and in the active cleaved caspase-3 and cleaved caspase-8 fragments in the lysates of Bit1-transfected cells (Fig. 4C). Instead, we observed that the Bit1-induced anoikis was associated with significant down-regulation of the detachment-induced TLE1 expression (Fig. 4D and 4E). The effect of Bit1 on TLE1 expression occurred primarily at the level of protein accumulation, as Bit1 had no effect on the detachment-induced TLE1 mRNA (Fig. 4F).

Figure 3. TLE1 regulates anoikis sensitivity of breast cancer cells. MDA-MB-231 cells were transfected with control- or TLE1-specific siRNAs, and, 48 hours after transfection, cells were harvested and subjected to immunoblotting (A) with antibodies against TLE1 and β-actin. In B and C, control- and TLE1-siRNA–treated cells were plated onto a polyhema-coated or uncoated tissue culture plates and cultured for 24 hours. B, the growth of detached cells was measured by alamar blue staining and fluorescent reading. C, the level of apoptosis in attached and detached cells was quantified by measuring the amount of DNA histone fragments (Cell Death Elisa). B and C, results are representative of 3 independent experiments, *, P < 0.05 (Student t test) as compared with control siRNA–transfected cells. D, stable MDA-MB-231 control shRNA and TLE1 shRNA knockdown pools were generated as described in Materials and Methods, and the total cell lysates derived from control shRNA and Bit1 shRNA knockdown pools were subjected to immunoblotting using specific antibodies to TLE1 and β-actin. E, stable MDA-MB-231 control shRNA and TLE1 shRNA knockdown pools were plated onto a polyhema-coated or uncoated tissue culture plates. Following 24 hours in culture, cells were harvested and analyzed for apoptosis by measuring the amount of DNA histone fragments (Cell Death Elisa). F, stable MDA-MB-231–derived DDK-TLE1 and control-overexpressing pool of cells were generated as described in Materials and Methods. The expression of exogenous DDK-TLE1 was confirmed by immunoblotting using the total cell lysates derived from control and TLE1-overexpressing pools against a specific antibody to DDK and β-actin. G, the control and TLE1-overexpressing cells were plated onto a polyhema-coated or uncoated tissue culture plates. Following 72 hours in culture, cells were harvested and analyzed for apoptosis by measuring the amount of DNA histone fragments (Cell Death Elisa). In E and G, 3 independent experiments were conducted in triplicate, *, P < 0.05 by Student t test.
Considering that TLE1 confers significant protection from anoikis in breast cancer cells and has been previously shown to inhibit Bit1 apoptosis function (8), we examined the notion that loss of TLE1 may contribute to Bit1-induced anoikis in breast carcinoma cells. To address this possibility, MDA-MB-231 cells were cotransfected with C-terminally myc-tagged Bit1 and/or full-length DDK-tagged TLE1. The cells were then cultured on tissue culture plates, or on poly-HEMA–coated plates to induce anoikis. The ectopic expression of mitochondrial Bit1 and TLE1 was confirmed.

Figure 4. Induction of anoikis by Bit1 is associated with downregulation of TLE1 expression. A, MDA-MB-231 cells were transfected with the indicated amounts of C-terminally myc-tagged mitochondrial localized Bit1 construct. The total amount of DNA was normalized with the empty vector in each transfection. 24 hours after transfection, cells were plated onto polyhema-coated or uncoated tissue culture plates. Following 24 hours in culture, cells were harvested and analyzed for apoptosis by measuring the amount of DNA histone fragments (Cell Death Elisa). Three independent experiments were conducted in triplicate; *, P < 0.05 as compared with the corresponding attached condition (Student t test). B, mitochondrial Bit1–transfected cells were cultured in attached or detached conditions in the presence of z-Vad-fmk (50 μmol/L) or DMSO for 24 hours. Cells were then harvested and subjected to Cell Death ELISA. C and D, MDA-MB-231 cells transfected with C-terminally myc-tagged Bit1 or vector construct were plated onto polyhema-coated or uncoated tissue culture plates. Following 24 hours in culture, cells were harvested and subjected to total cell lysate isolation, SDS-PAGE, and immunoblotting against specific antibodies to caspase-3, cleaved caspase-3, caspase-8, cleaved caspase-8, PARP, TLE1, myc-tag, and β-actin. E, the TLE1/β-actin ratios were determined by densitometry analysis of immunoblots in D from 3 independent experiments; *, P < 0.05 by Student t test. F, MDA-MB-231 cells transfected with mitochondrial Bit1 or vector construct were plated onto polyhema-coated or uncoated tissue culture plates and cultured for 24 hours. Cells were then harvested and subjected to total RNA isolation, and TLE1 mRNA expression levels were quantified by reverse transcription and real-time PCR analysis.
via immunoblotting (Fig. 5A). Consistent with our previous findings (8, 11, 13), detachment induced a higher level of apoptosis in Bit1-transfected cells than in control cells (Fig. 5B). Importantly, ectopic TLE1 levels attenuated the high level of detachment-induced apoptosis in mitochondrial Bit1-transfected cells. Consistent with the inhibitory effect of TLE1 on Bit1 anoikis, the apoptotic activity of cytoplasmic localized Bit1 (cyto) was inhibited by TLE1 (Supplementary Fig. S6) as shown previously (8). It is noteworthy that similar attenuation of Bit1 anoikis was also observed in stable TLE1-expressing cells (Fig. 5C). To complement our results with ectopic TLE1 expression studies, we also examined the effect of decreased TLE1 expression on Bit1-mediated anoikis. Knockdown of TLE1 further increased the sensitivity of MDA-MB-231 cells to Bit1 anoikis (Fig. 5D).

To address the mechanism underlying the inhibitory effect of TLE1 on Bit1 anoikis function, we examined whether TLE1 can inhibit the formation of the Bit1-AES complex, which is the proapoptotic component that mediates the effect of Bit1 on apoptosis (8). As shown in Fig. 5E, AES coprecipitated with mitochondrial Bit1 in detached conditions, whereas no Bit1-AES complexes were detected in attached cells. Importantly, the ability of mitochondrial Bit1 to interact with AES during anoikis is inhibited by exogenous TLE1 (Fig. 5E). Because Bit1 complexes with AES in the cytoplasm (8), we speculate that TLE1 inhibits Bit1-AES complex formation via sequestration of AES in the

Figure 5. TLE1 antagonizes Bit1 anoikis function. A, MDA-MB-231 cells were transfected with C-terminally myc-tagged Bit1 together with DDK-tagged TLE1. 24 hours after transfection, the expression of exogenous Bit1 and TLE1 in transfected cells was confirmed by immunoblotting using specific antibodies to myc and DDK tags. The amount of plasmid transfected into cells was normalized with the vector construct. B, MDA-MB-231 cells transfected with the indicated construct were grown for 24 hours in suspension, and cells were then harvested and subjected to Cell Death ELISA assay to measure the amount of DNA histone fragments. C, stable MDA-MB-231–derived DDK-TLE1 and control-overexpressing pool of cells were transfected with mitochondrial Bit1 or vector construct, and, 24 hours after transfection, cells were plated onto polyhema-coated tissue culture plates. Following 48 hours in suspension culture, cells were harvested and analyzed for apoptosis by measuring the amount of DNA histone fragments (Cell Death Elisa). D, stable MDA-MB-231 control shRNA and TLE1 shRNA knockdown cells transfected with mitochondrial Bit1 or vector construct were subjected to anoikis and Cell Death Elisa assays as described in C. E, MDA-MB-231 cells were transfected with the indicated plasmids. 24 hours after transfection, cells were grown for another 24 hours in attached or detached conditions. Cell extracts were prepared, immunoprecipitated with agarose-immobilized anti-myc, and immunoblotted with anti-AES. F, MDA-MB-231 cells transfected with the indicated plasmids were grown in suspension for 24 hours, and cells were then harvested and subjected to nuclear isolation as indicated in the Materials and Methods. The resulting nuclear fraction was run on SDS-PAGE and immunoblotted with an anti-AES antibody. The Histone H2B was used as a nuclear marker. B to D, 3 independent experiments were carried out in triplicate, *, P < 0.05 by Student t test.
nucleus through hetero-oligomerization (1). Indeed, concomitant with the observed reduction of Bit1-AES complexes by exogenous TLE1 is the significant enrichment of AES in the nucleus (Fig. 5F). Collectively, these findings indicate that the induction of anoikis by Bit1 may involve suppression of the antiapoptotic TLE1 and that Bit1 anoikis activity can be inhibited by overexpression of TLE1 in breast cancer cells.

Bit1 induces cytoplasmic translocation and degradation of nuclear TLE1 protein during anoikis

Mitochondrial Bit1 markedly downregulates the level of endogenous TLE1 protein during anoikis (Fig. 4D and E). Consistent with this observation, we also found that the level of exogenous TLE1 protein in a stable TLE1-expressing pool of cells was significantly reduced by Bit1 in detached conditions, whereas the ectopic TLE1 expression was unaffected by Bit1 in attached conditions (Fig. 6A and B). These findings indicate that translocation of mitochondrial Bit1 to the cytoplasm during anoikis (8, 13, Fig. 6C) is required to decrease TLE1 protein levels. Consistent with the effect of mitochondrial release of Bit1 on TLE1 during anoikis, the cytoplasmic expression of proapoptotic N-terminally myc-tagged Bit1 (8, 13, Supplementary Fig. S6) and of the highly cytotoxic cell death domain (CDD) of Bit1 (localized to amino acids 1–62 of the molecule, Chen and colleagues; submitted for publication) resulted in a significant reduction in the TLE1 protein levels (Fig. 6D and E). We previously created negatively charged phosphomimetic mutation in the Ser5 amino acid within the mitochondrial localization sequence of the C-terminally GFP-tagged mitochondrial Bit1 (Bit1SSD-GFP and Bit1SSE-GFP), and such Bit1 phosphomimetic mutants exhibited enhanced cytoplasmic localization and, as a result, reduced mitochondrial import efficiency and displayed a more potent anoikis-sensitization activity (13). Indeed, these Bit1SSD-GFP and Bit1SSE-GFP mutants induced a higher reduction of TLE1 protein as compared with wild-type Bit1 (Bit1-GFP) and Bit1SSA-GFP mutant during anoikis (Fig. 6F). The observed reduction of TLE1 during anoikis is specific to Bit1 as the basal anoikis activity in HEK 293T cells, which we have previously shown to have a negligible amount of endogenous Bit1 expression (8), was not associated with alteration of TLE1 protein levels (Supplementary Fig. S7A).

Despite the reduction of TLE1 protein levels in Bit1-transfected cells during anoikis, mitochondrial Bit1 expression has no significant effect on the detachment-induced upregulation of TLE1 mRNA (Fig. 4F), suggesting that Bit1 may regulate TLE1 at the level of protein accumulation. We first examined whether Bit1 may regulate the intracellular localization of TLE1. Translocation of mitochondrial Bit1 to the cytoplasm during anoikis (Fig. 6C) coincided with the cytoplasmic relocalization of the nuclear TLE1 protein (Fig. 6G and Supplementary Fig. S7C). Consistent with this finding, exogenous expression of the cytoplasmic localized Bit1 or its CDD domain also led to TLE1 accumulation in the cytoplasm (Fig. 6H). The observed cytoplasmic relocalization of TLE1 is likely specific to Bit1 as the basal anoikis in low Bit1-expressing HEK 293T cells did not show alteration of TLE1 nuclear localization (Supplementary Fig. S7B). The cytoplasmic enrichment of TLE1, however, became evident upon induction of Bit1-mediated anoikis (Supplementary Fig. S7C and S7D). Given that a number of critical anoikis regulators are subject to proteasomal degradation in response to detachment (19, 20), we then explored whether TLE1 is subsequently channeled to proteasomal degradation in the presence of Bit1 during anoikis. Cells transfected with mitochondrial Bit1 were plated in suspension culture in the presence or absence of proteasomal inhibitor, MG132. Figure 6I and J show that treatment of Bit1-transfected cells grown in suspension with MG132 exhibited increased levels of TLE1 protein compared with untreated Bit1-transfected cells, indicating that Bit1 may channel TLE1 to proteasomal degradation.

TLE1 is overexpressed in invasive breast tumors

On the basis of our in vitro evidence indicating that TLE1 promotes breast cancer cell anoikis resistance, which is a primary determinant of transformation and tumor aggressiveness, we explored the possibility that TLE1 may contribute to breast tumorigenesis in humans by screening a breast tumor tissue microarray for overexpression of TLE1. Both the normal breast epithelium tissue (Fig. 7A) and the noninvasive ductal carcinoma in situ (DCIS) lesions (Fig. 7A, iii) similarly showed low nuclear TLE1 staining. In contrast, the majority of tumor cells in a significant fraction of invasive breast carcinoma lesions consistently displayed moderate to strong TLE1 nuclear immunoreactivity (Fig. 7A, v). Quantification of the average TLE1 nuclear staining (Fig. 7B) confirmed the similar TLE1 expression between the normal and DCIS subgroups and the upregulation of TLE1 immunoreactivity in invasive breast carcinoma tissues as compared with normal/DCIS subgroups (ANOVA and subsequent Tukey post hoc analysis). We have previously shown that Bit1 expression is downregulated in a significant fraction of advanced invasive breast carcinoma tissues as compared with the normal breast epithelial tissue and DCIS lesions (11). To assess the relationship between TLE1 and Bit1 expression in breast carcinogenesis, immunohistochemical staining of Bit1 was conducted in the above studied breast tumor tissue arrays (Fig. 7A ii, iv, vi). Both the normal breast tissue and DCIS lesions, which exhibited low nuclear TLE1 staining, displayed moderate to strong cytoplasmic Bit1 immunoreactivity (Fig. 7A ii and iv), as previously observed (11). Importantly, a significant number of breast carcinoma tissues (43.2%) examined showed strong nuclear TLE1 and low cytoplasmic Bit1 staining (Fig. 7Avi; Table 1). Only a minor fraction (24.2%) of breast tumor tissue samples showed both strong nuclear TLE1 and cytoplasmic Bit1 staining (Table 1). The majority of the low TLE1-expressing breast carcinoma tissues displayed high cytoplasmic Bit1 immunoreactivity (21.2%; Table 1), and a small number of breast carcinoma tissues (10.6%; Table 1) display low levels of both TLE1 and Bit1 immunostains. Taken together, these data indicate that TLE1 is overexpressed in a fraction of human invasive breast carcinomas.
and that upregulation of TLE1 expression coupled with Bit1 downregulation may accompany the transition from DCIS to invasive carcinoma during breast cancer progression.

**Discussion**

In this report, we have provided evidence that the Groucho family member TLE1 is a regulator of mammary epithelial anoikis cell death program. Our findings also reveal an important link between cell–ECM interaction and the transcriptional regulation of TLE1 expression. In normal breast epithelial cells, loss of integrin engagement results in downregulation of TLE1 expression, and such TLE1 suppression may contribute to the anoikis sensitivity of untransformed epithelial cells. Conversely, in breast carcinoma cells, loss of matrix interaction results in the upregulation of TLE1 expression. The detachment-induced TLE1 expression functions to protect breast cancer cells from anoikis and promotes anchorage-independent growth, at least, in part, by inhibiting the Bit1 anoikis pathway. Importantly, we also show that the nuclear TLE1 is a target of the anoikis effector Bit1 and is channeled to the cytoplasm for proteasomal degradation by Bit1 during anoikis. The induction of TLE1 loss may represent a critical step in the anoikis function of Bit1. Taken together, these findings indicate a novel mechanism by which breast cancer cells evade anoikis via TLE1 expression which is regulated downstream of matrix attachment.

Gro proteins act as repressors for a wide range of gene targets (1, 2). Changes in Gro protein levels or repressor activity have a dramatic influence on the developmental fate of a cell. In neurons, TLE1 exerts an antidifferentiation or antiapneurogen function wherein ectopic expression of TLE in primary cultures of neural progenitor cells decreased neuronal differentiation and induced accumulation of proliferating progenitor cells (4, 5). TLE1 is also expressed in the proliferative stage of epithelial cells and is downregulated during differentiation (17). In addition to its role in cellular differentiation, our results indicate that TLE1 plays a critical role in protecting mammary epithelial and breast carcinoma cells from apoptosis induced by loss of matrix adhesion. Such findings are reminiscent of previous studies showing a potent prosurvival or antiapoptotic function of TLE1 in other cellular models. TLE1 has a strong growth promoting function in chicken embryo fibroblasts (6) while exogenous expression of TLE1 is sufficient to prevent apoptosis in neurons (7). Interestingly, neurons that are primed to die exhibit reduced expression of TLE1, which mimics our observation of loss of TLE1 expression in normal breast epithelial cells undergoing anoikis.

We have previously reported the Bit1 anoikis effector whose apoptotic function is uniquely regulated by integrin-mediated cell adhesion and is independent from caspases (8). The apoptotic activity of mitochondrial Bit1 is dependent on its ability to form a complex with AES in the cytoplasm. Here, we show that the release of mitochondrial Bit1 channels the nuclear TLE1 protein to the cytoplasm for proteasomal degradation during anoikis. This finding is supported by our studies with the use of the apoptotic cytoplasmic localized Bit1 and its apoptotic fragment CDD. Importantly, the Bit1-induced TLE1 loss upon cellular detachment from ECM is critical for the mitochondrial Bit1 anoikis function, as enforced expression of TLE1 is sufficient to attenuate Bit1-mediated anoikis. Thus, our previous and current data raise a possibility that the specific role of Bit1 is to turn off a survival-promoting gene transcription program regulated by TLE1. In this regard, ECM-mediated cell survival may involve complex transcriptional regulation of target apoptotic and antiapoptotic gene(s) through the suppressive effect of Bit1 on TLE1.

The downstream target gene(s) of the TLE1-mediated survival promoting transcriptional program remains to be determined. In light of our findings, the TLE1 corepressor may function to suppress the expression of proapoptotic factors that are intricately involved in anoikis program. In
as evidenced in this paper (Fig. 1B, C, and D), may be
required for induction of anoikis (16). We are currently investigating the
downstream target gene(s) regulated by TLE1 through microarray
analysis and determining which of these gene(s) specifically
regulate the apoptotic program during anoikis.

The ability of TLE1 to impair anoikis in breast epithelial
cells may have a profound influence on breast tumorigenesis.
Anoikis resistance is a determinant of transformation and
tumor aggressiveness (22). Indeed, we found here that while
exogenous TLE1 expression confers increased anchorage-
independent growth, specific TLE1 downregulation attenuates
breast cancer cells’ anchorage-independent growth capacity.
Sonderegger and colleagues have previously shown that
TLE1 is required for Qin-induced transformation in chicken
embryo fibroblasts (6), but the mechanism for this TLE1
effect remains to be determined. Our results reveal a critical
role of TLE1 in anoikis suppression, which may be the basis
for its stimulatory effect on anchorage-independent growth
and transformation. It is therefore not surprising that TLE1
has been found to be overexpressed in various types of human
solid malignancies including cervical (17), lung (10), soft
tissue (23), brain (24), and in breast cancer in the present
study (Fig. 7A–B). In lymphoma, elevated TLE1 has been
identified as a marker for poor prognosis (25). Importantly,
Grg1 (the mouse homologue of TLE1) transgenic mice
developed lung tumors (10). Taken together, these findings
implicate TLE1 in the development of human malignancy
and raise the possibility of TLE1 as a putative oncogene.

As a summary, we found that found that TLE1 expression
is significantly downregulated in a normal mammary ep-
ithelial cell MCF10A line following loss of cell attachment,
and the loss of TLE1 expression contributes in part to the
anoikis sensitivity of these nonmalignant cells. In contrast,
TLE1 expression is induced in breast carcinoma cells during
anoikis. The upregulation of TLE1 protects malignant breast
cells from anoikis at least in part by inhibiting the casapse-
developmental anoikis function of Bit1. We further provide
evidence that Bit1 targets TLE1 to cytoplasm for proteaso-
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Table 1. Immunohistochemical analysis for nuclear TLE1 and cytoplasmic Bit1 expression in the breast tumor tissue population

<table>
<thead>
<tr>
<th>Nuclear TLE1</th>
<th>Cytoplasmic Bit1</th>
<th>Number of tissues (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Moderate to strong</td>
<td>Low</td>
<td>29 (43.2)</td>
</tr>
<tr>
<td>Moderate to strong</td>
<td>High</td>
<td>16 (24.2)</td>
</tr>
<tr>
<td>No or low</td>
<td>Low</td>
<td>7 (10.6)</td>
</tr>
<tr>
<td>No or low</td>
<td>High</td>
<td>14 (21.2)</td>
</tr>
</tbody>
</table>

Figure 7. TLE1 is overexpressed in invasive breast carcinoma tissues. A, breast tumor tissue array slides were stained with a rabbit polyclonal antibody to TLE1 (i, iii, v) or the affinity-purified rabbit anti-Bit1 antibody (ii, iv, vi). Images are representative of each respective case type: normal breast (i, ii × 10), DCIS (iii, iv × 10), and invasive breast carcinoma (v, vi × 10). B, the average staining intensity of each subgroup was determined as described in Materials and Methods. While no significant difference was found between normal and DCIS subgroups, the normal/DCIS were statistically significant from the invasive breast carcinoma tissues using the ANOVA and subsequent Tukey post hoc analysis (see Materials and Methods; *, P < 0.05).

mammary epithelial cells, BH3-only factors, Bim and Bmf,
are upregulated following loss of cell matrix attachment
and required for induction of anoikis (16). These proapoptotic
factors are potential candidate downstream target genes
that are suppressed by TLE1 during anoikis. Intriguingly, there
are data to suggest that transcriptional suppression of Bmf
requires histone deacetylase (HDAC) activity (21). Consid-
ering that HDAC function requires recruitment of TLE1, it
will be interesting to determine whether TLE1 plays a critical
role in the basal repression of Bmf. In this scenario, the loss
of TLE1 upon loss of matrix adhesion in breast epithelial cells,
as evidenced in this paper (Fig. 1B, C, and D), may be
sufficient to relieve the transcriptional repression of Bmf,
leading to its upregulation during anoikis as previously
observed (16). We are currently investigating the down-
stream target gene(s) regulated by TLE1 through microarray
analysis and determining which of these gene(s) specifically
regulate the apoptotic program during anoikis.
Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

Authors' Contributions
Conception and design: C. Brunquell, H. Biliran, R. Chen, E. Ruoslahti
Development of methodology: C. Brunquell, H. Biliran, R. Chen
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Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis): C. Brunquell, H. Biliran, E. Ruoslahti
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Study supervision: R. Chen, E. Ruoslahti

References

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