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## Correlation of transcription of MALAT-1, a novel noncoding RNA, with deregulated expression of tumor suppressor p53 in small DNA tumor virus models

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### Abstract

Although metastasis-associated lung adenocarcinoma transcript (MALAT)-1 is known to be consistently upregulated in several epithelial malignancies, little is known about its function or regulation. We therefore examined the relationship between MALAT-1 expression and candidate modulators such as DNA tumor virus oncoproteins human papillomavirus (HPV)-16 E6 and E7, BK virus T antigen (BKVTA<sub>g</sub>), mouse polyoma virus middle T antigen (MPVmTA<sub>g</sub>) and tumor suppressor genes p53 and pRb. Using suppressive subtractive hybridization (SSH) and real-time reverse transcriptase polymerase chain reaction (RT-PCR) assays, MALAT-1 was shown to be increased in viral onconeogene-expressing salivary gland biopsies from humans and mice. The results also indicated that MALAT-1 transcripts and promoter activity were increased in vitro when viral onconeogene-expressing plasmids were introduced into different cell types. These same viral oncogenes in addition to increasing MALAT-1 transcription have also been shown to inhibit p53 and/or pRb function. In p53 mutant or inactive cell lines MALAT-1 was also shown to be highly upregulated. We hypothesize that there is a correlation between MALAT-1 over-expression and p53 deregulation. In conclusion, we show that disruption of p53, by both polyoma and papilloma oncoproteins appear to play an important role in the up-regulation of MALAT-1. MALAT-1 might therefore represent a biomarker for p53 deregulation within malignancies.

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### Conflict of interest

Dr. Diatchenko is a cofounder and the Chief Scientific Officer of Algynomics, Inc., which has exclusive rights to develop and market proprietary genetic pain markers owned and patented by the University of North Carolina. The other authors have no potential conflicts to disclose.

## Keywords

p53; MALAT-1; transcription; BK virus; polyoma virus; papillomavirus; HPV

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## Introduction

Metastasis-associated lung adenocarcinoma transcript-1 (MALAT-1) is a novel large, noncoding RNA. The MALAT-1 gene, also known as the *gene*, is found on chromosome 11q13 and is well conserved among mammalian species [1]. The MALAT-1 transcript is widely expressed in normal human and mouse tissue, has been shown to localize to the nucleus[2] and its 3' end can be processed to yield a tRNA-like cytoplasmic RNA[3]. MALAT-1, has been shown to be a potentially generic marker for epithelial carcinomas[4–7] and is greatly up-regulated in lung adenocarcinoma metastasis[1] endometrial stromal sarcoma of the uterus[7], nonhepatic human carcinomas[5] and recently was reported to be overexpressed in placenta previa and to play a role in trophoblast invasion regulation [8]. Up-regulation of MALAT-1 also has been shown to predict unfavorable outcomes of drug therapy in patients with osteosarcoma[4] and its 3' end is an important biological motif in the invasion and metastasis of colorectal cancer cells [9]. These malignancies all have been associated with malfunction of p53[10; 11] a nuclear transcription factor that plays a role in cellular stress, including its accumulation during DNA damage and oncogene activation. P53 has been established as a key tumor suppressor, apoptosis inducer, and prognostic marker in cancer, with about 50% of human tumors encoding for a mutated p53 gene[12].

Several small DNA tumor viruses affect p53 function, including the polyomaviruses, the adenoviruses, and the papillomaviruses. These viruses play a role in the development of cancer in humans by encoding for proteins that interact with tumor suppressor genes p53 and pRb[13]. Polyomavirus large T antigen, for example, binds to and inhibits p53 and pRb[14–16]. The mouse polyoma virus middle T antigen (MPVmTAg) prevents p53-induced apoptosis through the phosphatidylinositol 3-kinase (PI3K) signal transduction pathway by way of PP2A interactions[17]. Adenovirus E1B[18] and high-risk human papillomavirus (HPV-16) E6[19] proteins both bind to and compromise p53 function, whereas HPV-16 E7 inhibits pRb function[20]

The genomes of small DNA tumor viruses have consistently been detected in certain malignancies. HPV is consistently detected in cervical cancer[21] and in a subset of oral cancers[22], whereas BK virus (BKV) DNA has been isolated in several human tumors, both integrated into the genome and episomally, including in bone, pancreatic islet cells, the kidney, the urinary tract, the prostate, and various brain tissues[23–29]. In addition, overexpression of polyoma virus mTAg has been shown to cause salivary gland enlargement in mice and pathology similar to that observed in human salivary gland disease[30; 31].

We hypothesized that MALAT-1 expression is affected by dysregulation of tumor suppressor p53. We used viruses, viral gene products, and cell lines to test the modulation of a cloned MALAT-1 promoter and to determine the effect of p53 or p53 mutants on MALAT-1 expression. We also conducted in vivo testing of MALAT-1 expression in a transgenic mouse model of salivary gland disease (SGD) and in 4 subjects with HIV-associated SGD (HIVSGD).

## Materials and methods

### Subjects, animals and cell culture

Minor salivary glands (MSG) were dissected from the lower lips of 4 HIV positive patients with SGD and from 4 healthy control subjects (with or without ranula) in IRB approved protocol at either the UNC University Hospital dental clinic or UCSF Oral AIDS Center. Biopsy samples were snap-frozen and kept in liquid nitrogen or formalin-fixed and embedded in paraffin until the RNA extraction procedure. Four MMTV/PyV-mT transgenic mice and 4 wild type mice parotid samples were kindly donated from Dr. Lesley Ellies.

HSG cells are an epithelial cell line isolated by using tissue culture techniques from an irradiated human submandibular salivary gland which showed no neoplastic lesion[32]. HSG cells were obtained as a gift from Dr. B. Baum (NIH) and cultured in McCoy's 5A medium (Sigma). African monkey kidney cells or Vero cells (American Type Culture Collection [ATCC]) were cultured in DMEM (Sigma). All cell types were grown in medium supplemented with 10% fetal bovine serum (FBS) (Sigma), and 1% penicillin-streptomycin (pen/strep)(Gibco) unless otherwise stated and maintained in a humidified 37°C, 5% CO<sub>2</sub> chamber.

### Suppression Subtractive Hybridization (SSH)

One to two µg of pooled biopsied salivary glands from HIV-SGD patients and pooled control (healthy persons) RNA was sent to Evrogen (Russian Republic) for SSH. In brief, SMART technology was used to synthesize cDNA, which was subsequently digested with RsaI in preparation for subtraction in both directions (HIV-SGD as driver/control as tester, and vice versa). Adaptors were ligated to the two cDNA populations, followed by two rounds of hybridization and amplification. The secondary PCR products from the two subtracted populations were then ligated into the pAL9 vector.

### Sequencing of subtracted cDNA library

Two hundred and fifty positive clones from each of the subtractions were randomly selected and cultured in 1.5 mL LB-medium in duplicate 96-well plates at 37 °C overnight. Clones were selected by blue-white screening (Promega) and sequenced using a M13 forward promoter (-21) primer (5'-TGT AAA ACG ACG GCC AGT-3') and BigDye 3.0 sequencing mix (Applied Biosystems) before analysis by capillary electrophoresis on an ABI 3700 genetic analyser (Applied Biosystems). The sequences of the inserts of differentially expressed genes were identified using NCBI Blast search (blastn).

### RNA extraction and cDNA synthesis

Total RNA was extracted from pooled MSGs of HIV-SGD patients, pooled MSGs of healthy controls, mouse parotid frozen tissues and Vero cells using RNeasy MiniKit (Qiagen, USA). Total RNA isolation from formalin fixed tissue using the Optimum FFPE RNA Isolation Kit (Ambion Diagnostics, INC) according to manufacturer's instructions. The RNA was suspended in nuclease-free water and quantitated by UV spectrophotometry, aliquoted and stored at -80°C. One µg total RNA from the HIV-SGD and control tissues, mice parotid tissue and Vero cells were reverse transcribed to cDNA using random primers and the SuperScript™ II Reverse Transcriptase (RT) Kit (Invitrogen) as described by the manufacturer. Contaminating DNAs were removed by use of RQ1 DNase kit (Promega) as described by the manufacturer.

## Northern Blot

Total RNA was isolated using TRizol (Invitrogen) as described by the manufacturer. Contaminating DNAs were removed by use of RQ1 DNase kit (Promega) as described by the manufacturer. 10 ug of RNA was subjected to electrophoresis and Northern blotting. MALAT-1 was hybridized with a <sup>32</sup>P-labeled oligonucleotide probe for MALAT-1

## Semi-quantitative and quantitative real-time RT-PCR

Semi-quantitative RT-PCR was performed using previously published primers for T Ag[33] and Taq Polymerase (Qiagen, USA). The following program was used for amplification: 95°C for 2 min (1x); 94°C for 45 sec, 56°C for 45 sec, 72°C for 30 sec with 2 sec increase per cycle (35x), 72°C for 10 min (1x). Amplified cDNA was electrophoresed on 2% agarose gel (Sigma). Real time RT-PCR was performed with LightCycler 480 Syber Green I Master Mix in the presence of transcribed cDNA and 0.25mM of gene specific primers. Experiments were performed in duplicate and  $\beta$ -actin was used as an internal control. Duplicate or triplicate Ct values were averaged, normalized to an average  $\beta$ -actin Ct value, and fold activation in healthy vs diseased tissue was calculated using the  $2^{-\Delta\Delta Ct}$  method. For Vero cells, gene expression values are presented as the changes (*n*-fold) in T Ag transcript levels, with the levels in non-transfected/mock samples arbitrarily set to 1.

## Immunofluorescence/Immunohistochemistry

Frozen sections were cut from minor salivary glands (HIV-SGD and control glands) and from mouse parotid glands (transgenic and wild type) for immunofluorescent analysis. Tissue sections were fixed, blocked, then stained with either PAb416 (Genetex) antibody specific for SV40 T antigen or IgG isotype control. PAb416 has been shown to cross react with BKV Tag and is commonly used for BKV Tag detection. Both antibodies were incubated with fixed cells for 1 hr at 37°C followed by a fluorescencein-conjugated anti-mouse (Sigma) antibody (1:20). Slides were overlaid with Vectashield (Vector Laboratories) then subjected to immunofluorescence microscopy.

Formalin fixed sections were deparaffinized, and washed. Slides were incubated in 3% hydrogen peroxide and blocked and incubated with PAb416 (Genetex). DAKO LSAB+ peroxidase kit (DAKO Corporation) was used according to manufacturer's specifications.

## Plasmid cloning, transfection and infection

A 5.5 kb region upstream of MALAT-1 transcription site was amplified using primers 5' GGGACGCGTAAAGAGGATTCTATCTAACAAGGA3' and 5' GGGCTCGAGGAAACGTGAAAACCCACTCT3' and inserted into pSeap2-Basic expression cloning vector (Clontech) using restriction sites MluI and XhoI. To determine MALAT-1 promoter activity, Vero cells were first infected with BKV as previously described[34] with 64 HAU of virus for 24 h. At 24 h post infection (hpi), virus was removed from the culture media, cells were washed with 1X PBS and replaced with fresh medium containing the MALAT-1 promoter plasmid and  $\beta$ -gal construct plus transfection reagent. At stated times post transfection the cell monolayers or supernatant were collected for immunoblot assays, beta-galactosidase enzyme assay (Promega) or secreted alkaline phosphatase enzyme assay (Promega) according to manufacturers instructions.

## Immunoblotting

Total cell protein was extracted using 1% SDS lysis buffer (1% (w/v) SDS, 0.05M Tris.Cl pH8, 1mM DTT). Protein concentrations were determined using the BioRad protein assay, and equal amounts of protein were electrophoresed on a 10% Bis-Tris polyacrylamide minigel (Invitrogen). PAb416 (1:200) (Genetex) in 5% non-fat dry milk in 0.1% Tween-20

PBS (PBS-T) was used to detect T Ag expression and Actin (C-11)-R sc-1615-R (1:1000) (Santa Cruz Biotechnology) in 1% BSA/TBS-T for actin expression. After washing in PBS-T/TBS-T, blots were probed with a horseradish peroxidase-conjugated secondary antibody (1:10,000) (Promega). Antibody complexes were detected using SuperSignal West Pico Chemiluminescent substrate (Thermo scientific) and exposed to film (Kodak).

### **Beta galactosidase enzyme assay**

-Galactosidase Enzyme Assay System with Reporter Lysis Buffer (Promega) was performed according to manufacturer's instructions. Briefly, cells were collected, centrifuged and resuspended in 1X lysis buffer and transferred in triplicate to 96 well plates. Diluted sample was added to an equal volume of Assay 2X Buffer, which contained the substrate ONPG (o-nitrophenyl-beta-D-galactopyranoside). Samples were incubated for 30 minutes, terminated by addition of sodium carbonate, and the absorbance read at 420 nm with a spectrophotometer.

### **Secreted alkaline phosphatase reporter gene assay, chemiluminescent**

SEAP assay was performed according to manufacturer's instructions. Briefly, culture supernatant from transfected or mock (untreated) cells were collected at stated times post-transfection. Collected sample was diluted 1:4 in Dilution buffer, heated at 65°C for 30 min then centrifuged at maximum speed to 30 s. Heat-treated samples were then transferred to a microplate (black or white), inactivated with Inactivation Buffer then treated with Substrate Reagent. Chemiluminescence was measured using a luminometer.

### **Statistical Analysis**

One-way ANOVA tests were performed to determine statistically significant differences in MALAT-1 promoter activation and expression using Graph pad software.

### **miRNA Prediction**

The entire MALAT-1 sequence was imported into an online program mfold (<http://www.bioinfo.rpi.edu/applications/mfold>) to predict the presence of micro RNAs within the genome.

## **Results**

### **Differential gene expression in HIV-SGD by SSH**

Differential gene expression in minor salivary glands from patients with HIV-SGD and healthy control subjects were examined with the use of SSH. Pooled RNA from four HIV-SGD positive subjects and from four HIV negative subjects was utilized to diminish individual variation. Two SSH libraries were constructed, disease (plus) and healthy library (minus). The minus library was enriched for genes whose expression was suppressed in HIV-SGD and the plus library was enriched for genes whose expression was induced by disease development. 250 randomly selected clones were sequenced from the minus SSH library and 250 clones from the plus library; of those 232 and 208 contained cDNA inserts, respectively. The SSH library was automatically processed using a specifically created software tool. This processing included 3 basic steps: finding adaptor sequences on the 5 and 3 ends of a clone as markers and extracting a fragment from the clone, blasting the batch of fragments, clustering alignments and linking with unique gene identification. We were able to align 436 sequences with known or predicted human genes. Our total number of alignments reached 502 due to 57 chimeric clones which had 2 or three clones from different RNA's linked together during SSH. Of the 436 cDNAs 73 sequences were

identified that did not match to any known or predicted mRNAs but aligned with the human genomic DNA and with EST databases. These were considered novel.

A non redundant set of the genes was created which excluded highly expressed clones present in both disease and healthy genes in the plus library. There were 6 genes found more than once among sequenced SSH clones in each of the libraries (Table 1). It was shown previously that a number of cDNA fragments corresponding to a gene in the SSH library correlates with a degree of differential expression of the gene[35]. Considering all of the differentially expressed clones detected by SSH analysis, major functional classes of transcriptionally regulated genes in minus library included signal transduction (9%), cell cycle(8%), immune/defense(8%), synthesis/development/differentiation (8%) and transport (8%). Transcriptionally regulated genes in the plus expression library included those involved in signal transduction (15%), synthesis/development/differentiation(13%), apoptosis(8%), metabolism(8%) and transport. The largest category of de-regulated genes were unknown, among these was the metastasis associated lung adenocarcinoma transcript-1 (MALAT-1).

### **MALAT-1 transcript and promoter activity is deregulated in the absence of p53 expression, in vitro**

A reporter assay was developed to assess activation of MALAT-1's enhancer/promoter region. First, we examined the MALAT-1 enhancer/promoter region 5.5 kb upstream of the ATG start site by means of open-source software (PROMO 3.0; available at [http://algggen.lsi.upc.es/cgi-bin/promo\\_v3/promo/promoinit.cgi?dirDB=TF\\_8.3](http://algggen.lsi.upc.es/cgi-bin/promo_v3/promo/promoinit.cgi?dirDB=TF_8.3))[36]. This program identifies putative transcription factor binding sites in DNA sequences from the species (or group of species) of interest. The similarity between the sequence examined and the transcription binding site is > 98% (or dissimilarity is < 2%). Figure 1A shows the transcription sites found in the enhancer/promoter region of MALAT-1. We then quantified the MALAT-1–secreted embryonic alkaline phosphatase (SEAP) by chemiluminescent assay (Roche Diagnostic) and transiently cotransfected the reporter construct into p53-negative, sarcoma osteogenic (SAOS) cells with or without plasmids encoding wild-type p53 or p53 mutants R273H (a contact mutation), R175H (a structural mutation), H178Y, and L22QW23S[37–39]. The first two mutants are considered hot-spot mutations; that is, they abolish the wild-type tumor suppression function of p53 [40; 41]. The H178Y mutant has been shown to rescue the loss of function G245S phenotype when coexpressed in vitro. The L22QW23S mutant maintains its ability to bind to p53-specific DNA elements but lacks transactivation activity[37]. Cotransfections were performed using p53 expression plasmids and MALAT-1 promoter in pSEAP vector and supernatant collected at 72 hours post transfection. Higher MALAT-1 expression levels were detected in cells that did not express p53 or that contained p53 mutations (Figure 1B). We also examined MALAT-1 expression profiles in cell lines containing p53 mutations (C33A, HSG), those containing DNA tumor viruses that sequester p53 (CaSki, SiHa), p53-negative cells (SAOS), and cells containing wild-type p53 (OKF6-Tert). Only the cell types expressing wild-type p53 showed low MALAT-1 levels (Figure 1C). These differences were statistically significant ( $P < 0.001$ ).

To further investigate the role of p53 in deregulating MALAT-1 expression, we over-expressed HPV16 oncogenes E6 and E7 which inhibit p53 and pRb respectively. Our results showed that over expression of HPV-16 E6 in oral keratinocytes significantly enhanced MALAT-1 expression while HPV-16 E7 did not. Coexpression of both E6 and E7 resulted in a net increase in MALAT-1 transcription (Figure 2A).

To confirm that viral oncogenes that inhibit p53 may play a role in MALAT-1 deregulation, we infected or transfected BK virus or viral genome into HSG or Vero cells. BKV T ag binds to and inhibits p53 and pRb function[16]. Both human parotid salivary gland (HSG)

and African monkey kidney (Vero) cells were used for in vitro studies because BKV has been shown to replicate in both cell types[34]. MALAT-1 transcript levels were 1.6-fold higher in BKV-infected salivary gland cells versus uninfected/mock cells 4 days after infection (Figure 2B). MALAT-1 transcription was also consistently increased in Vero cells that overexpressed BKV whole genome compared to empty vector and nontransfected according to Northern blot testing (Figure 2B). Semi-quantitative real-time reverse transcriptase polymerase chain reaction (RT-PCR; Figure 2C), and quantitative real-time RT-PCR (Figure 2D) for MALAT-1 in T antigen transfected Vero cells also showed an increase in MALAT-1 transcripts. A 2.3 and 5.5 fold increase in MALAT-1 at 24 and 48 hours post transfection was detected by qRT-PCR in Vero cells.

To assess the correlation between p53 and T ag oncogene expression, we measured T ag and p21 transcript levels in BKV infected Vero cells. T antigen is known to bind to and inhibit p53 activity, while p21 is a downstream effector of p53 and would be expected to decrease in the absence of p53 expression. The results as expected indicated that p21 transcripts were low with high levels of BKV T ag expression, and high with low levels of T ag expression (Figure 2E). To determine whether BKV T ag expression regulates MALAT-1 promoter activity, we transfected Vero cells with the MALAT-1 promoter followed by BKV infection. MALAT-1 promoter activity was up-regulated 7-fold at 72 hours post BKV infection in comparison to uninfected. BKV T ag expression within these cells was confirmed in the immunoblot assay (Figure 2F). BKV T ag comprises of several transcription binding sites including p53 and pRb. To determine whether p53 or pRb plays a role in MALAT-1 regulation, we co-transfected both Vero (wild-type p53) and HSG (mutated p53) cells with either BKV T ag or BKV T ag pRb-binding mutant and the MALAT-1 promoter expression plasmid. The results showed that MALAT-1 promoter activity was increased compared to vector only in Vero cells regardless of T ag phenotype (Figure 2G). Whereas, in HSG cells containing mutant p53, neither wild-type TAg nor Rb further enhanced MALAT-1 transcription above vector alone (Figure 2H).

Next, we investigated MALAT-1 levels in an in vivo model of BK T antigen-expressing salivary gland premalignant lesion (HIVSGD). Among all of the differentially expressed clones detected by SSH analysis, the most common category of deregulated genes was the “unknown” category, which included MALAT-1 (Table 1). Quantitative real-time RT-PCR detected a 2.6-fold increase in MALAT-1 expression in pooled DNA analysis from patients with HIVSGD versus healthy controls (Figure 3A). Semi-quantitative real-time RT-PCR likewise detected up-regulation of MALAT-1 in individual DNA samples from four patients with SGD (A–D) versus four healthy controls (A–D) (Figure 3A).

Immunohistochemical analysis of salivary gland biopsy samples from three patients with HIVSGD detected BKV T ag protein within the tissues as well as in a BKV positive kidney control with BKV nephropathy (BKVN) (Figure 3B, top). In addition, p53 was colocalized with BKV T ag within biopsied salivary glands from the patients with SGD (representative figure, Figure 3B, bottom). Downstream effectors of p53 (p21, PIGP-1, and WIG-1) were consistently down-regulated in semi-quantitative (left) and quantitative real-time (right) RT-PCR, whereas ING-3, negatively regulated by p53 expression and T ag was overexpressed, in both pooled cDNA (data not shown) and individual samples from patients with HIVSGD compared with healthy controls (Figure 3C).

In the murine model of polyomavirus-associated SGD, semi-quantitative real-time RT-PCR detected greater amplification of MALAT-1 cDNA from four transgenic mice with SGD, whereas transcripts were consistently reduced in four wild-type mice (Figure 3D, middle). Immunofluorescence studies detected T ag within biopsied salivary glands of transgenic mice but not in the wild-type mice (Figure 3D, top). Both ING-3 and MALAT-1 were

greatly upregulated in transgenic mice versus wild-type mice in quantitative real-time RT-PCR assay (Figure 3D, bottom).

## Discussion

MALAT-1 is known to be up-regulated in several epithelial malignancies, including non-small cell lung tumors, stromal sarcomas, and nonhepatic carcinomas[4–7; 42]. In this series of in vitro and in vivo experiments, we noted substantial MALAT-1 up-regulation in the presence of polyoma and papilloma oncoproteins that disrupt or negate p53 expression/function. In contrast, the pRb pathway did not appear to be critical to MALAT-1 regulation. Regardless of the mechanism of p53 deregulation—expression of frequently mutated p53 sites (Figure 1) or expression of tumorvirus oncoproteins targeting p53 (Figure 2 and 3)—MALAT-1 was consistently overexpressed. Thus MALAT-1 might represent a biomarker for p53 dysregulation within malignancies.

MALAT-1 expression in CaSki cells has been shown to be involved in cervical cancer cell growth, cell cycle progression, and invasion[43]. CaSki cells harbor HPV-16, a small DNA tumor virus that encodes for the E6 protein. This protein is known to form complexes with p53 and target it for degradation[43]. In contrast, the HPV-16 E7 protein is known to sequester pRb and inhibit its normal function in the cell cycle. The enhanced expression of MALAT-1 in the presence of HPV-16 E6, and its downregulation in the presence of HPV-16 E7, lend support to the hypothesis that p53, but not the pRb pathway, appears to be critical to MALAT-1 modulation (Figure 3).

In normal tissues, MALAT-1 is expressed in numerous cell types, with the highest relative expression found in the normal pancreas and lung (~2.0- and 1.6-fold relative increases in expression, respectively)[1]. Its expression is also associated with epithelial cell malignancies such as non-small cell lung cancer[1] endometrial stromal sarcoma[7] and nonhepatic carcinomas[5]. Of interest, all of these cancers are associated with p53 mutation (Table 2). However, normal salivary gland cells appear to express negligible MALAT-1[1]. Our study shows, for the first time, that the presence of a polyoma virus (BKV) is linked to up-regulation of MALAT-1 in patients with HIVSGD. Interestingly, the perinuclear BKV Tag staining pattern in HIVSGD is very similar to that detected by the Imperiale group in prostate dysplasia[2; 3]. Moreover, oncoprotein expression from three distinct DNA tumor viruses - HPV-16, BKV, and mouse polyoma virus -are capable of upregulating MALAT-1 potentially via p53 deregulation. Interestingly, using an mRNA folding prediction program it appears that the 3' end of the MALAT-1 gene may encode a microRNA (miRNA) (Figure 4). miRNAs are single-stranded RNA molecules of 21–23 nucleotides in length, which regulate gene expression. miRNAs are encoded by genes from whose DNA they are transcribed but miRNA are not translated into protein, that is they are non-coding RNA; instead each primary transcript (a pri-miRNA) is processed into a short stem-loop structure called a pre-miRNA and finally into a functional miRNA. Mature miRNA molecules are partially complementary to one or more messenger RNA (mRNA) molecules, and their main function is to down-regulate gene expression. Like known miRNAs, MALAT-1 is a non-coding transcript, folds into a short stem loop structure at the 3' end and is conserved across mammalian species. Modulation of MALAT-1 expression by p53, perhaps by miRNA regulation, is intriguing and opens the door toward understanding the regulation of this important malignancy-associated gene. Investigation of MALAT-1 as a miRNA is currently being explored and potential targets in SGD will be determined. In the meantime, MALAT-1 might represent a biomarker for p53 deregulation within malignancies.



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Figure 1A.

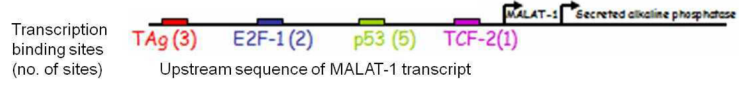


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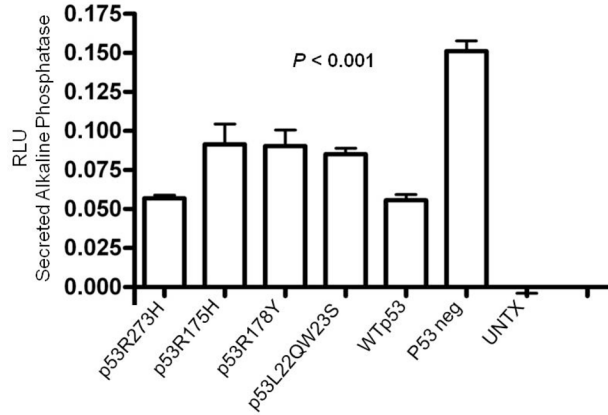


Figure 1C.

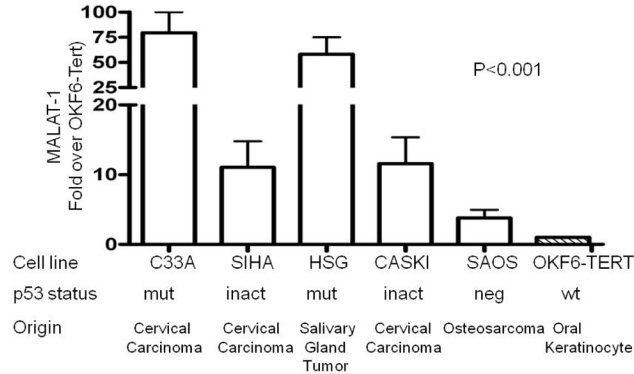
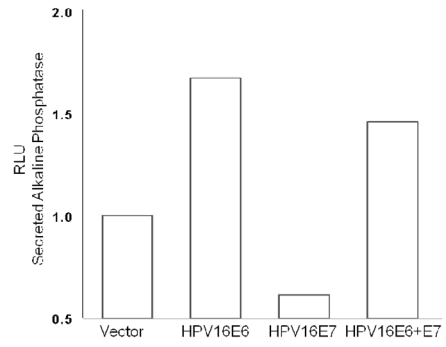


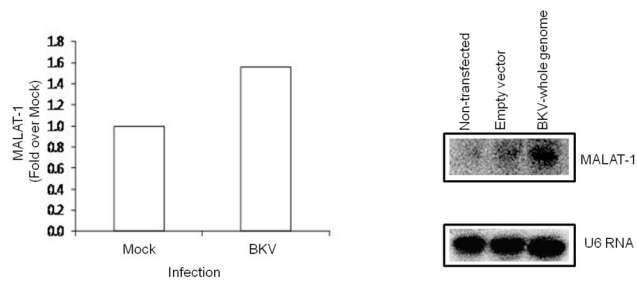
Figure 1.

MALAT-1 promoter activity and transcript levels correlate with p53 activity. **A.** Several consensus binding sites were detected upstream of the MALAT-1 start site, including five p53 binding sites (green), three T antigen binding sites (red), two TCF-2 binding sites (purple), and two E2F-1 binding sites (blue). **B.** Representative data showing significantly increased MALAT-1 promoter activity in p53-deficient SAOS cells 72 hours after transfection with p53-expressing constructs. **C.** Representative levels of MALAT-1 expression in different cell types with or without p53 gene expression, presented as the change (n-fold) in MALAT-1 transcript levels (level in OKF6 cells arbitrarily set to 1).

**Figure 2A.**



**Figure 2B.**



**Figure 2C.**

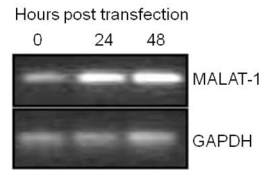


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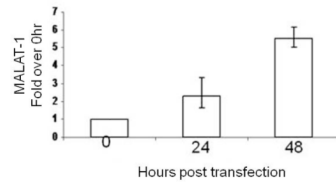


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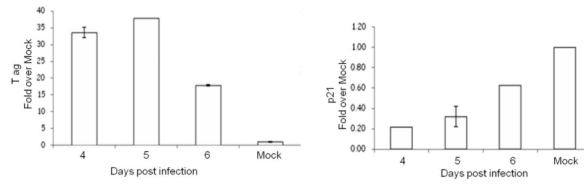
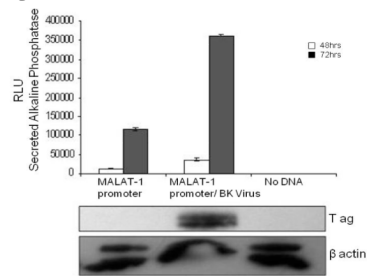
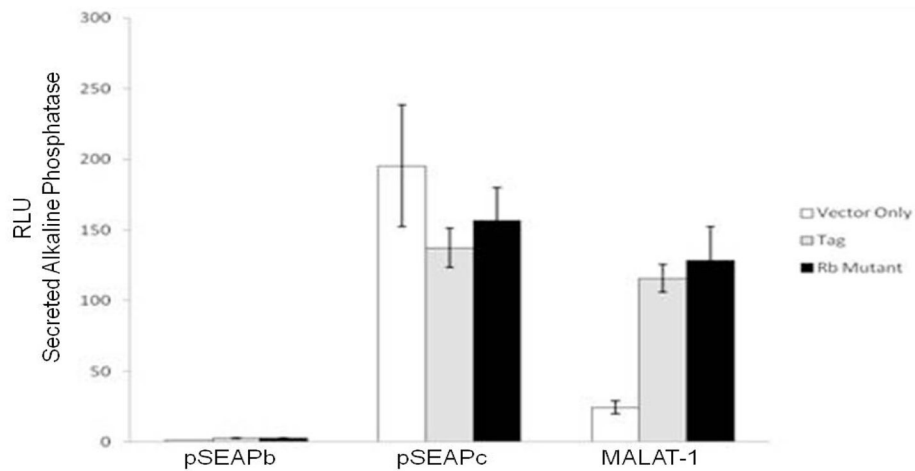


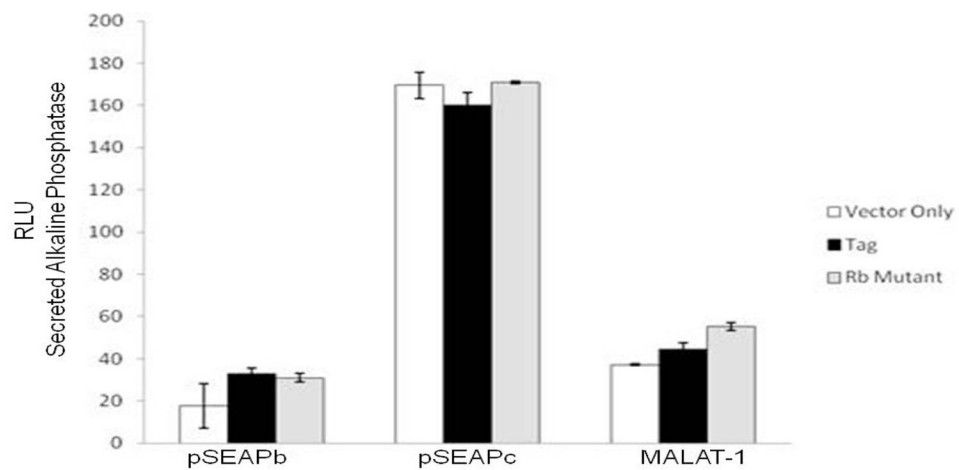
Figure 2F.



**Figure 2G.**



**Figure 2H.**



**Figure 2.**

MALAT-1 promoter activity and transcript levels correlate with DNA tumor virus oncogene expression. **A.** Representative data showing increased MALAT-1 promoter activity 48 hours after HPV16 E6 transfection and not with HPV16 E7. **B.** Higher levels of MALAT-1 transcription, expressed as n-fold change in BK polyomavirus-infected human parotid gland (HSG) cells. Levels in mock samples were arbitrarily set to 1. *Left.* Northern blot showing a four-fold increase in MALAT-1 transcript levels in Vero cells transfected with BKV DNA. *Right C.* Semi-quantitative real-time RT-PCR shows amplified MALAT-1 cDNA bands from BKV-transfected Vero cells at stated times after transfection. **D.** Quantitative real-time RT-PCR shows increased MALAT-1 transcript levels in BKV-transfected Vero cells at stated times after transfection. Expression values presented as the change (n-fold) in MALAT-1 transcript levels, with levels at time 0 arbitrarily set to 1. **E.** Quantitative real-time RT-PCR showing BKV Tag and p21 transcript levels over stated times after BKV infection. **F.** MALAT-1 promoter activity in BKV-infected and uninfected Vero cells at 48 and 72 hours after transfection. *Top.* BKV T ag protein (top) detected in Vero cells infected with BK virus and transfected with the MALAT-1 promoter plasmid. **G.** BKV T ag up-regulated MALAT-1 promoter activity regardless of T ag pRb binding status in Vero cells

containing wild-type p53 compare to positive and negative controls, pSEAPc and pSEAPb, respectively. **H.** Neither wild-type BKV T ag nor T ag pRb binding mutant enhanced MALAT-1 transcript levels in human salivary gland cells containing mutant p53 expression.



Figure 3A.

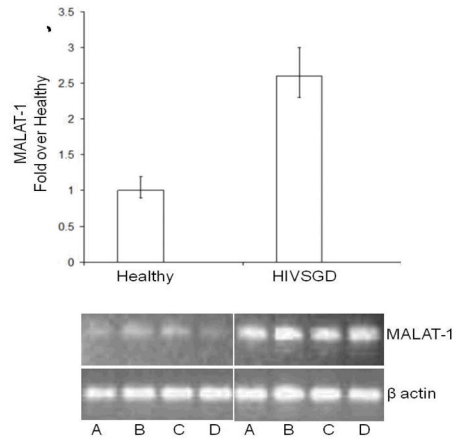
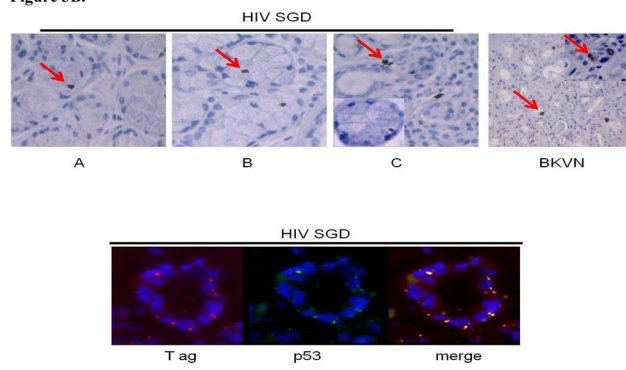
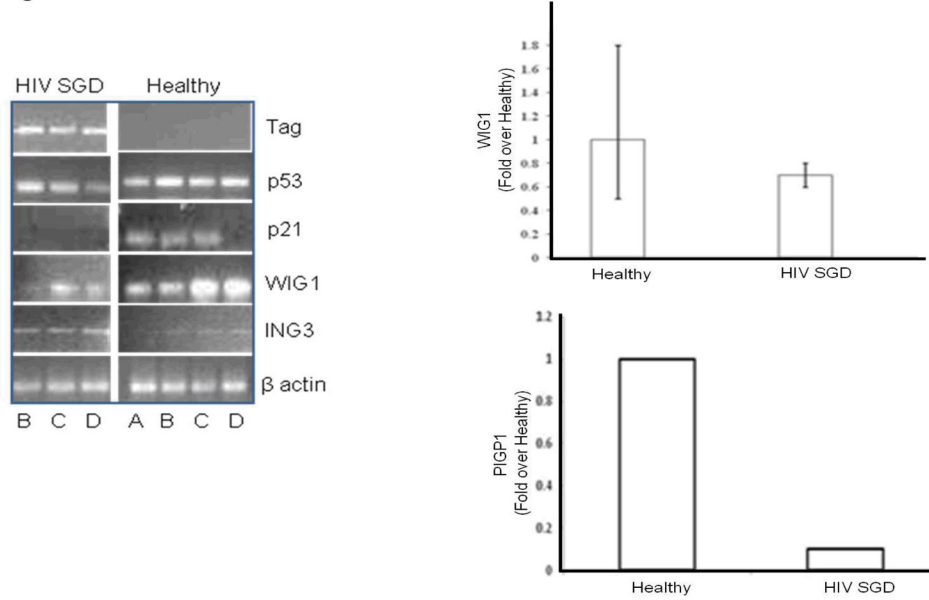


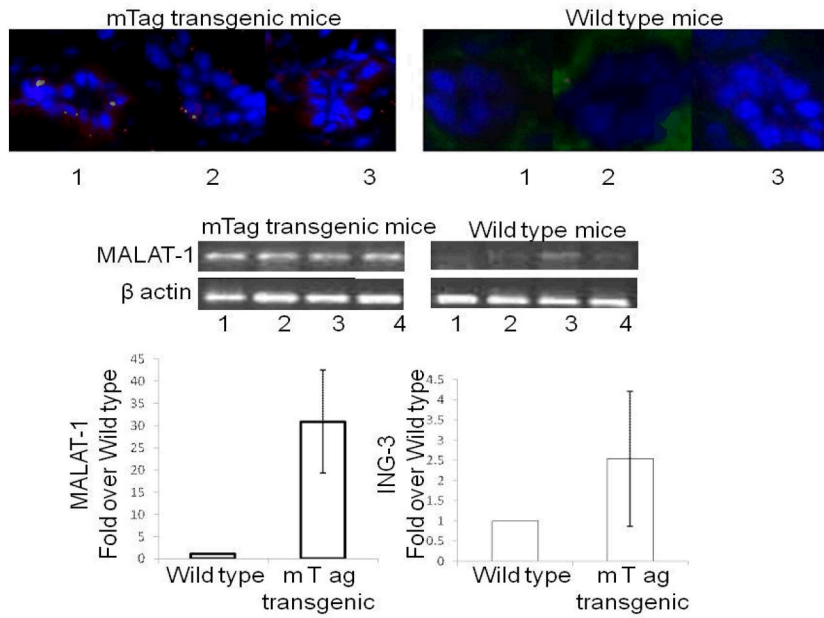
Figure 3B.



**Figure 3C.**



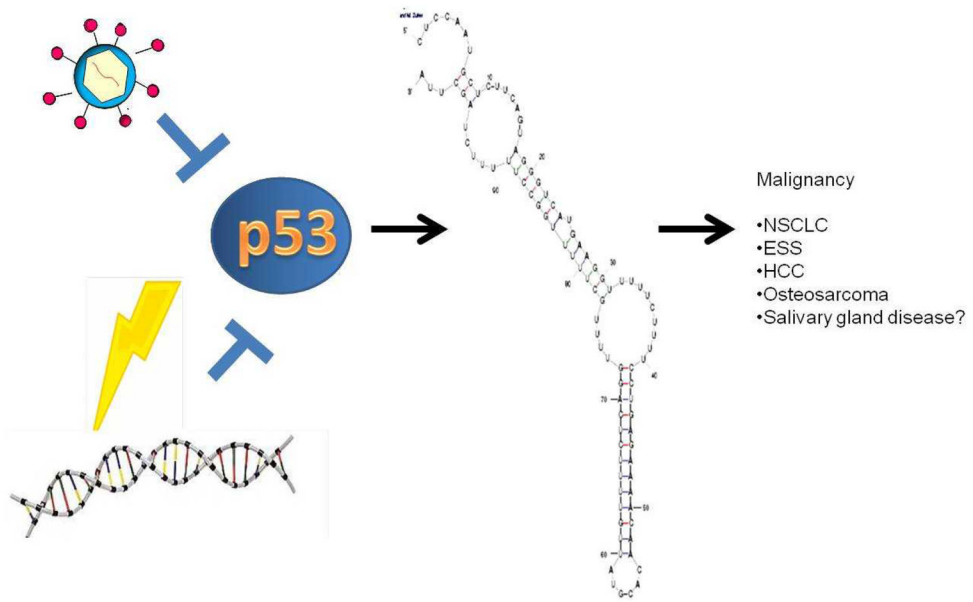
**Figure 3D.**



**Figure 3.**

MALAT-1 transcript levels correlate with DNA tumor virus oncogene expression in vivo. **A.** Quantitative real-time RT-PCR showing up-regulation of MALAT-1 in pooled DNA from four patients with HIV associated salivary gland disease (HIV SGD) compared with healthy control subjects, *Top*. Semi-quantitative real-time RT-PCR showing up-regulation of MALAT-1 in DNA from four patients with HIV SGD versus healthy control subjects, *Bottom*. **B.** Nuclear BKVT ag (red arrows) detected via immunohistochemical analysis of representative biopsy samples from three patients with HIV-related SGD (Patient A 20X, B 40X, C 20X with inset at 40X) and in BKV-infected kidney cells from an individual with BKV nephropathy (positive control at 10X), *Top*. BKV T ag protein (red) and p53 protein (green) colocalized within salivary gland biopsy tissue from patients with HIV SGD (at 40X

magnification) **C.** Semi-quantitative (left) and quantitative real-time RT-PCR (right) showing detection of BKV T ag in HIV SGD and not in healthy controls, down-regulation of p53-regulated genes p21, PIGP1 and WIG1 and up-regulation of ING3 in salivary gland tissue from patients with HIV SGD versus healthy controls. Healthy cDNA levels were arbitrarily set at 1. **D.** BKV T ag protein (red) detected via immunofluorescence in three middle T ag-expressing transgenic mice with SGD (left) compared with wild-type mice without the disease (20 and 40X magnification) (right), *Top*. Semi-quantitative real-time RT-PCR showing up-regulation of MALAT-1 transcript levels in transgenic mice salivary gland biopsies compared with wild-type mice, *Middle*. Quantitative real-time RT-PCR showing up-regulation of MALAT-1 and ING3 in transgenic mice with SGD versus wild-type mice. Wild-type cDNA levels arbitrarily set at 1, *Bottom*.



**Figure 4.** Model of MALAT-1 regulation including predicted miRNA at the 3' end of the MALAT-1 RNA.

**Table 1**

Genes up-regulated in polyomavirus-associated salivary gland disease as assessed by suppressive subtractive hybridization

<b>Gene</b>	<b>Description</b>	<b>Category</b>	<b>No. of clones</b>
MALAT-1	metastasis-associated lung adenocarcinoma transcript 1 (noncoding RNA)	Unknown	4
PIGR	polymeric immunoglobulin receptor	Immunity	3
CYTB	Cytochrome B	Transport, Mitochondria	3
HLA-B	major histocompatibility complex, class I, B	Immunity	2
PIP	prolactin-induced protein	Immunity	2
RPLP0	ribosomal subunit protein L16	Protein synthesis	2

**Table 2**

Correlation among cancers that express high MALAT1 transcripts, involve p53 deregulation and are associated with DNA tumor virus infection.

<b>Human carcinomas</b>	<b>p53 status</b>	<b>MALAT-1 over-expressed</b>
Endometrial Stromal Sarcoma[7]	P53 nuclear accumulation[44]	Yes
Hepatocellular Carcinoma[5]	P53 mutation[5]	Yes
Non Small Cell Lung Cancer[1]	P53 mutation[12]	Yes
HIV-Salivary Gland Disease (pre-malignant)	N/D	Yes
Osteosarcoma[4]	P53 mutation[45]	Yes
Breast[5]	P53 nuclear accumulation[46]	Yes
Pancreas[5]	P53 mutation[47]	Yes
Colon[5]	P53 mutation[48]	Yes
Late-Stage Prostate Cancer[5]	P53 mutation[49]	Yes
Cervical (CaSki cell line) [41]	HPV inhibits P53 activity[50]	Yes
Endometrial Stromal Sarcoma [7]	P53 nuclear accumulation[44]	Yes
Hepatocellular Carcinoma[5]	P53 mutation[5; 45]	Yes
Non Small Cell Lung Cancer[1]	P53 mutation[12]	Yes
HIV-Salivary Gland Disease (pre-malignant)	N/D	Yes
Osteosarcoma[4]	P53 mutation[46]	Yes
Breast[5]	P53 nuclear accumulation [47]	Yes
Pancreas [5]	P53 mutation[48]	Yes
Colon[5]	P53 mutation[49]	Yes
Late-Stage Prostate Cancer[5; 45]	P53 mutation[50; 51]	Yes
Cervical (CaSki cell line)[43]	HPV inhibits P53 activity[52]	Yes