Spontaneous Tumor Regression in Tasmanian Devils Associated with RASL11A Activation

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ABSTRACT Spontaneous tumor regression has been documented in a small proportion of human cancer patients, but the specific mechanisms underlying tumor regression without treatment are not well understood. Tasmanian devils are threatened with extinction from a transmissible cancer due to universal susceptibility and a near 100% case fatality rate. In over 10,000 cases, <20 instances of natural tumor regression have been detected. Previous work in this system has focused on Tasmanian devil genetic variation associated with the regression phenotype. Here, we used comparative and functional genomics to identify tumor genetic variation associated with tumor regression. We show that a single point mutation in the 5' untranslated region of the putative tumor suppressor *RASL11A* significantly contributes to tumor regression. *RASL11A* was expressed in regressed tumors but silenced in wild-type, nonregressed tumors, consistent with *RASL11A* downregulation in human cancers. Induced *RASL11A* expression significantly reduced tumor cell proliferation *in vitro*. The RAS pathway is frequently altered in human cancers, and *RASL11A* activation may provide a therapeutic treatment option for Tasmanian devils as well as a general mechanism for tumor inhibition.

KEYWORDS cancer; tumor regression; gene expression; tumor suppressor

S PONTANEOUS cancer regression is the reduction or disappearance of a malignant tumor without treatment, and, albeit rare, has been documented in a variety of human cancers (Jessy 2011). Although spontaneous regression has been hypothesized to be associated with exogenous immune stimulation, growth factors, cytokines, and hormonal mediation (Cole 1981; Papac 1998), little is actually known about the underlying mechanisms. Identifying the specific genes

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and processes driving regression could lead to improved cancer treatments and preventions through targeted immuno- or gene therapies (Mittelman *et al.* 2001; Corrales *et al.* 2015), but ethical considerations prevent the investigation of natural, untreated cancer progression in humans. Therefore, comparative oncological studies in nonhuman systems are needed to identify the mechanisms of important tumor phenotypes such as spontaneous regression.

Recent evidence of spontaneous tumor regression has been documented in natural populations of Tasmanian devils (*Sarcophilus harrisii*) infected with a transmissible cancer (Pye *et al.* 2016a; Wright *et al.* 2017; Margres *et al.* 2018). Although most cancers are somatic in origin, Tasmanian devils are threatened with extinction by the emergence and rapid spread of two, independently derived, transmissible cancers

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[devil facial tumor disease (DFTD) and DFT2] (Murchison *et al.* 2012; Pye *et al.* 2016b; Stammnitz *et al.* 2018; Storfer *et al.* 2018). DFTD and DFT2 are contagious tumor cell lines spread via biting during common social interactions (Hamede *et al.* 2013), and both tumors have nearly a 100% case fatality rate. Although DFT2 was discovered only recently in 2014, and has so far remained limited in its geographic spread (Pye *et al.* 2016b; Stammnitz *et al.* 2018), DFTD has spread across ~95% of the devil's geographic range and has caused localized declines >90% over the last 23 years (McCallum 2008). Several Tasmanian devils with confirmed DFTD in northwestern Tasmania, however, were recently recaptured with either substantial tumor shrinkage or entire disappearance of the tumor (Figure 1 and File S1; Pye *et al.* 2016a; Wright *et al.* 2017; Margres *et al.* 2018).

Similar to the focus on patient immunological responses to human cancers (Morgan et al. 2006; Sharma et al. 2017), recent studies have focused on Tasmanian devil genetic variation associated with DFTD regression and found putative regulatory variation in candidate genes associated with angiogenesis and cancer risk (Wright et al. 2017; Margres et al. 2018). DFTD-specific mechanisms, however, have not yet been investigated, and tumor genetic variation can affect oncogenicity (Bailey et al. 2018). Here, we build on our previous work in the hosts (Margres et al. 2018), and used comparative and functional genomics to identify tumor-intrinsic factors associated with spontaneous regression in the matching DFTD samples (i.e., matched tumor-normal design). We compared regressed and nonregressed tumor genomes and transcriptomes and found a single point mutation that activated the tumor suppressor gene RASL11A in regressed tumors. We then used in vitro cell proliferation assays to show that tumor cell proliferation was significantly inhibited by virally induced RASL11A expression in three of four tumor cell lines. Taken together, our results suggest that RASL11A activation may be a general mechanism for tumor inhibition.

Materials and Methods

DFTD sampling

Tasmanian devils were trapped in northwestern Tasmania using custom-built traps as previously described (Figure 1; Hamede *et al.* 2015). Following capture, devils were tagged with microchip transponders (Allflex NZ Ltd, Palmerstone North, New Zealand), aged, inspected for DFTD, and released; DFTD status was determined by histopathological confirmation of tumor biopsies, and tumor volume was measured for all infected individuals. To confidently assign a phenotypic designation to a tumor (*i.e.*, regressed or nonregressed), we required tumors to be sampled at multiple time points. Tumors were classified as regressed if total tumor volume decreased $\geq 15\%$ of the initial tumor volume upon recapture for Tasmanian devils previously confirmed to be infected with DFTD as described elsewhere (Margres *et al.* 2018). Otherwise, tumors were classified as nonregressed

(Figure 1). One exception to these criteria was Nonregressed 1. Although this tumor was only sampled a single time-point, the individual host was disease free in 2012, retrapped in 2013 with DFTD, was in extremely poor body condition, and ultimately euthanized by the Save the Tasmanian Devil Program; moribund animals such as this have never been shown to recover. Given the sample size limitations of this study, we were confident in the classification of this tumor as nonregressed due to its rapid growth and clear negative consequences on host health. Raw tumor volume data are provided in Supplemental Material, File S1. Animal use was approved by the IACUC at Washington State University (#04392; ASAF), the University of Tasmania Animal Ethics Committee (A0008588, A0010296, A0011696, A0013326, A0015835), and the Department of Primary Industries, Parks, Water and Environment Animal Ethics Committee. Demographic information for the matching hosts samples can be found in Margres et al. (2018).

Whole-genome sequencing

We performed initial whole-genome sequencing for 12 DFTD tumors; these tumors were collected from 11 Tasmanian devils in northwestern Tasmania as previously described (Margres *et al.* 2018). Eight tumors exhibited tumor regression (Regressed 1–8; Figure 1) and four tumors did not (Nonregressed 1–4; Figure 1). Three regressed tumors (Regressed1, 3, and 6; Figure 1 and Figure S1) were serially sampled several months apart and resequenced (*i.e.*, we sequenced 15 total genomes across 12 individual tumors). Genomic DNA was extracted from tumor biopsies. Whole genomes for each tumor were sequenced 150 PE on an Illumina HiSeq X platform to ~90× coverage. Sequencing was performed at the Northwest Genomics Center at the University of Washington (Seattle, WA) and GENEWIZ (South Plainfield, NJ).

Alignments and variant calling

We merged and trimmed reads with FLASH2 (Magoč and Salzberg 2011) and Sickle (Joshi and Fass 2011), respectively. Merged and unmerged reads were aligned to the Tasmanian devil reference genome (downloaded from Ensembl January 2016; Murchison et al. 2012) using the BWA-MEM algorithm (Li and Durbin 2009). We identified ~ 2.6 million SNPs using HaplotypeCaller in GATK (McKenna et al. 2010; DePristo et al. 2011) as previously described (Margres et al. 2017). To identify somatic or cancer-specific SNPs, we performed a hard filter and removed any SNP previously identified in the paired host (i.e., Tasmanian devil) samples from Margres et al. (2018). We identified \sim 1.4 million somatic SNPs across all 15 initial genomes (12 individual tumors) following the hard filter. These variants were then filtered in VCFtools (Danecek et al. 2011) and were required to possess a minimum depth value of 15 and a minimum genotype quality of 60 as previously described (Margres et al. 2018). We identified 632,756 somatic variants that passed filter across the 15 tumor genomes.

Identifying the genomic basis of tumor regression

We used the Genotype Phenotype Association Toolkit (GPAT++) from the vcflib package (Garrison 2012) as previously described (Domyan et al. 2016) to identify genomic differentiation between regressed and nonregressed tumors. Differentiation was identified using the pFst function. pFst uses a likelihood ratio test to identify allele frequency differences across groups while simultaneously accounting for potential sequencing error. We calculated pFst values for 471,676 SNPs when comparing the eight regressed and four nonregressed tumors (Nonregressed 1-4; Figure 1). For these comparisons, only the initial sample was used for the three regressed tumors (Regressed 1a, 3a, and 6a; Figure 1 and Figure S1) that were sequenced multiple times. SNPs were identified as candidate variants if they occurred in the top 0.1% ($P \le 0.0137$) of the most differentiated SNPs. These candidates were characterized using variant effect predictor (VEP; McLaren et al. 2016) and the Tasmanian devil reference genome (downloaded from Ensembl January 2016; Murchison et al. 2012). Only nonsynonymous variants or variants in untranslated regions (UTRs) were functionally investigated in the current study because these variants were (1) most likely to affect the regression phenotype in a tractable manner, and (2) could be functionally validated (see below). We used GeneCards (www.genecards.org; Stelzer et al. 2016) to identify putative gene function.

Tumor phylogenetics

We took two approaches to infer unrooted tumor phylogenies for Regressed 1-8 and Nonregressed 1-4 tumors (Figure 1). First, we followed the approach of Margres et al. (2019) and used SVDquartets (Chifman and Kubatko 2014) in PAUP* v4.0a157 (Swofford 1998) to infer an unrooted phylogeny based on the filtered, somatic SNP data described above. We used PGDspider v2.1.1.2 (Lischer and Excoffier 2012) and custom scripts to produce multiple sequence alignments in fasta format, and concatenated the two alignments generated for each individual tumor. All quartets were estimated under the multispecies coalescent model (expecting matrixrank 10), and these quartets were assembled using the QFM algorithm. Confidence in tree topology was estimated using nonparametric bootstrapping. We used the SumTrees program in DendroPy v4.3.0 (Sukumaran and Holder 2010) to summarize across bootstrapped trees and produce the consensus tree (Margres et al. 2019). The consensus tree was visualized in FigTree v1.4.3 (Rambaut 2012). Second, we used RAxML v8 (Stamatakis 2014) on the CIPRES Science Gateway (Miller et al. 2010). Briefly, we conducted rapid bootstrapping (100 replicates) and searched for the highest-scoring ML tree. Because we were using exclusively SNP data, we performed an ascertainment bias correction to correct branch-length estimates (Leaché et al. 2015). To accomplish this, RAxML was run using the following parameter settings: -f a, -m ASC GTRCAT, -j -HKY85, -c 4, -N 100, -asc-corr felsenstein. As with the SVDquartets phylogeny, the RAxML ML tree was visualized using FigTree (Rambaut 2012), with bootstrap nodal support values shown.

Additional nonregressed tumor sequencing and analysis

We recognize that nonregressed tumors may appear to be easy to access given the higher frequency of the phenotype (i.e., wild type) in the population, but sampling confirmed nonregressed tumors was challenging. To confidently assign a phenotypic designation to a tumor (i.e., regressed or nonregressed), we required tumors to be sampled at multiple time points as described above. Given that devils infected with wild-type, nonregressing tumors ultimately perish faster than devils with regressed tumors, recapture following infection (and therefore confirmation of the nonregressing phenotype) was not as frequent as one might expect. Following our initial round of sequencing, we were able to sample three additional nonregressed tumor samples from the same geographic region (Nonregressed 5-7; Figure 1), and these three tumors represent all tumors that we could confirm (with some degree of confidence) as nonregressed. To confirm the candidate genotypes identified from the analyses above, we performed a second round of whole-genome sequencing for the three additional nonregressed DFTD tumors (Nonregressed 5-7; Figure 1). DNA was extracted from tumor biopsies and libraries were prepped using the NEBNext Ultra II DNA Library Prep Kit for Illumina (New England BioLabs, product E7645S). Whole-genomes for each tumor were sequenced 150 PE on an Illumina HiSeq X platform to $\sim 20 \times$ coverage at the Northwest Genomics Center at the University of Washington (Seattle, WA). We again merged and trimmed reads with FLASH2 (Magoč and Salzberg 2011) and Sickle (Joshi and Fass 2011), respectively. Merged and unmerged reads were again aligned to the Tasmanian devil reference genome (downloaded from Ensembl January 2016; Murchison et al. 2012) using the BWA-MEM algorithm (Li and Durbin 2009). SNPs were identified using HaplotypeCaller in GATK (McKenna et al. 2010; DePristo et al. 2011) with the following criteria: mean depth of 8, meanQ of 60, and maf of 0.05.

Tumor transcriptomics

RNA was extracted from two regressed and two nonregressed tumor biopsies using a standard Trizol method as previously described (McGivern *et al.* 2014). Of the four samples sequenced, one regressed (Regressed 1a) and one nonregressed (Nonregressed 2) tumor were included in the genome sequencing described above; the other two samples were not previously sequenced. Tissues were not available for the remaining samples, and because regression is such a rare phenotype, only one additional regressed sample (*i.e.*, the additional transcriptome sample sequenced here) was collected during the course of the study. Sequencing and cDNA library preparation were performed by the Genomics Core at Washington State University, Spokane (Spokane, WA). Each library was sequenced 100 PE across four lanes of an Illumina HiSeq 2500, and each transcriptome had 29.2–30.3 million reads. Low quality bases were trimmed with Trim Galore! version 0.4.5, and transcriptomes were assembled using the "new Tuxedo" package (Pertea *et al.* 2016). Here, reads were aligned to the reference genome (downloaded from Ensembl January 2016; Murchison *et al.* 2012) using HiSat2 version 2.1.0 (Kim *et al.* 2015), transcripts assembled and expression quantified using StringTie version 1.3.4 (Pertea *et al.* 2015), and analyzed/visualized in Ballgown (Fu *et al.* 2018). Alignment rates were >91% for all samples.

Lentiviral vector gene delivery

Four tumor cell lines (three DFTD lines and one DFT2 line) and a Tasmanian devil fibroblast cell line (Murchison *et al.* 2012) were grown in RPMI 640 media supplemented with 10% fetal bovine serum and penicillin/streptomycin. The Tasmanian devil fibroblast cell line was described elsewhere (Murchison *et al.* 2012). DFTD-a (DFT1-10181), DFTD-b (DFT1-A48), and DFTD-c (DFT1-c5065) were derived from primary DFTD tumors collected from Low Head, Tasmania in 2014, Bronte Park, Tasmania in 2004, and Bangor, Tasmania in 2007, respectively. The DFT2 cell line (DFT2-Jarvis) was derived from a primary DFT2 tumor collected from Nicholls Rivulet, Tasmania in 2015.

For each of the five cell lines described above, the Tasmanian devil *RASL11A* coding-sequence was cloned into a pCIG vector containing the puromycin N-acetyltransferase gene in place of enhanced green fluorescent protein (eGFP). A *RASL11A* gBlock was synthesized and inserted into the multiple cloning site of pCIG by Gibson assembly (Gibson *et al.* 2009). The pCIG *RASL11A* plasmid was digested with *AgeI* and *MluI* to remove eGFP. Puro was PCR amplified from GIPZ vector and ligated to the pCIG plasmid backbone to generate pCI puro-*RASL11A*. The eGFP lines did not express *RASL11A* and were considered control lines, and the pCI or *RASL11A* lines expressed *RASL11A* and were considered our case lines (see below; Figure S2 and File S2).

Real-time qPCR assays

Cellular RNA was extracted from 10 cm² of tissue culture by guanidinium thiocyanate-phenol-chloroform extract. RNA (1 µg) was converted to cDNA using iScript Reverse Transcription Supermix for RT-qPCR (Bio-Rad) according to the manufacturer's protocol. Real-time qPCR assays were conducted on the QuantStudio for RASL11A (forward: 5'-TCCCATTCCGGAG TCTTCCT-3'; reverse: 5'-CCTCTGCCCACTGAACACAT-3') and *RPS29* (forward: 5'-ATGGGTCATCAGCTCTAC-3'; reverse: 5'-AGGCCGTATTTGCGGATTAG-3'). Relative changes in expression were analyzed using the $2^{-\Delta\Delta C_T}$ method (Livak and Schmittgen 2001) where the relative expression level for the locus of interest (i.e., RASL11A) was compared to the expression level of the house-keeping gene RPS29 (File S2). We used paired, one-tailed *t*-tests to compare mean expression levels for RASL11A (Figure S2) across individual casecontrol assays for each cell line. Each assay included two to three replicates per cell line.

Cell proliferation assays

For each cell line described above, 3000 cells per well were seeded in 96-well flat bottom culture plates in RPMI 1640 medium supplemented with 10% fetal bovine serum and penicillin/streptomycin. Cell Titer-Glo assays were performed in triplicate. Cell Titer-Glo assays quantify the amount of ATP present, a proxy for the number of metabolically active cells and cell proliferation. ATP was quantified at days 0, 3, 4, 6, 7, and/or 11 according to the manufacturer's protocol. In total, we performed five assays (18 total replicates) for the Tasmanian devil fibroblast line and five (17 total replicates), five (18 total replicates), two (7 total replicates), and two (7 total replicates) assays for the DFTD-a, DFTD-b, DFTD-c, and DFT2 tumor lines, respectively (File S2).

To determine if RASL11A expression reduced tumor cell proliferation significantly more often than expected by chance, we used a χ^2 test to compare the observed frequency of replicates where the RASL11A line had a lower, final viable cell count than the control line vs. the expected frequency. If RASL11A expression did not affect tumor cell proliferation, we would expect the RASL11A lines to exhibit a greater proliferation rate than the control line (and vice versa) in \sim 50% of the replicates (i.e., our expected frequencies). We performed this test across all 49 tumor replicates as well as for DFTD and DFT2 independently. To visualize the cell proliferation assay results in Figure 3, we calculated the relative proliferation rate for the RASL11A line as the ratio of the RASL11A line ATP luminescence estimate to the control line ATP luminescence estimate (*i.e.*, case/control). Ratios >1 represent increased proliferation under RASL11A expression whereas ratios <1 represent decreased proliferation under RASL11A expression. We calculated these ratios for all replicates across all timepoints. One-sample *t*-tests were used to determine whether the final mean relative proliferation estimates significantly differed from one where a ratio of one represents the null expectation of identical proliferation between RASL11A and control lines.

Data availability

The authors state that all data necessary for confirming the conclusions presented in the article are represented fully within the article. All raw sequencing data were deposited in the Sequence Read Archive (SRA) under BioProject PRJNA472767. Whole-genome raw reads are under accessions SRR7257513 and SRR10436678, 79, and 81. Transcriptome raw reads are under accessions SRR7257322–SRR7257325. Raw functional genomic data are provided in File S2. Supplemental material available at figshare: https://doi.org/ 10.25386/genetics.12498896.

Results

Comparative genomics of regressed and nonregressed tumors

To identify putative causal mutations underlying spontaneous tumor regression in DFTD, we sequenced the genomes of four



Figure 1 Sampling of Devil Facial Tumor Disease. We collected tissue samples from 15 tumors for whole-genome sequencing from northwest Tasmania (see inset). Eight of these tumors underwent tumor regression ("Regressed" shown in blue) whereas the other seven did not ("Nonregressed" shown in red). Regressed 1, Regressed 3, and Regressed 6 were serially sampled, and timepoints are indicated as (a) and (b), respectively. Nonregressed 1–4 were included in initial analyses, and nonregressed 5–7 were sequenced following these initial analyses to increase sample size. See File S1 for additional data.

nonregressed (Nonregressed 1-4) and eight regressed tumors (Regressed 1-8; Figure 1); the latter represent all available tumors that exhibited $\geq 15\%$ reductions in tumor volume following recapture of the infected Tasmanian devil, with several cases of complete tumor clearance (Figure 2B and File S1; Margres et al. 2018). We identified 471,676 tumor-specific SNPs across the 12 tumor genomes, similar to previous analyses that identified 691-699 k SNPs across two DFTD genomes (Murchison et al. 2012); none of these variants were identified in the genomes of the Tasmanian devils that possessed the tumors examined in this study (Margres et al. 2018). Because most discovered variants were intergenic (\sim 75%), SNPs were identified as candidates if they were highly differentiated (i.e., top 0.1% of most differentiated somatic variants; $P \leq 0.0137$) and resulted in nonsynonymous substitutions or occurred in putative regulatory regions. We did not identify any candidate nonsynonymous SNPs, consistent with previous work showing a general paucity of nonsynonymous variation among DFTD (Stammnitz et al. 2018). We did, however, identify two candidate variants in the 5' untranslated regions (UTRs) of RASL11A and *ERICH6* (P = 0.0105; Figure 2A). Both candidates involved $T \leftrightarrow C$ transitions, the most common mutation in human cancers (Bailey et al. 2018). For each locus, we identified a near fixed genotypic difference (Figure 2C). All four nonregressed tumors were homozygous for the reference allele (T/T for RASL11A and C/C for ERICH6). Six of the seven genotyped regressed tumors were heterozygous at each site (T/C); one regressed tumor (Regressed 8) was not genotyped due to missing data. The only regressed tumor that was not heterozygous at either locus (Regressed 3 in Figure 2C) represented a unique case where an individual Tasmanian devil possessed both regressed (Regressed 3) and nonregressed (Nonregressed

3) tumors. These tumors were sister to one another in one of our two phylogenetic analyses (Figure S1), suggesting potential metastasis and within host tumor divergence. Here, regression may be a product of host genetic variation (Margres *et al.* 2018) and/or immune response (see *Discussion*; Pye *et al.* 2016a).

To increase the nonregressed tumor sample size, we sequenced an additional three nonregressed tumor genomes at lower coverage and genotyped each sample at both the RASL11A and ERICH6 candidate sites (Nonregressed 5-7 in Figure 1 and File S1). For each candidate variant, two of the three nonregressed tumors were homozygous for the reference allele, but one nonregressed tumor was heterozygous at both sites. Overall, six of seven nonregressed tumors were homozygous and six of seven regressed tumors were heterozygous at each allele. The near fixed genotypic differences in putative regulatory regions for RASL11A and ERICH6 suggested that expression variation at either locus may contribute to tumor regression. Although the function of ERICH6 is not known, RASL11A is a regulator of rDNA transcription and a putative tumor suppressor (Pistoni et al. 2010). RASL11A expression is downregulated in human prostate tumors (Louro et al. 2004) and hypermethylated in colon cancer cell lines (Weber et al. 2005), suggesting that RASL11A expression may negatively affect oncogenicity.

RASL11A expression associates with tumor regression

To determine if these UTR variants affected the expression of either candidate gene, we sequenced the transcriptomes for two regressed and two nonregressed tumors (tissues were not available for remaining samples). *ERICH6* was not assembled in any of the four transcriptomes, indicating its lack of expression in either tumor type. *RASL11A* was expressed



Figure 2 Genetic differentiation underlying tumor regression in Devil Facial Tumor Disease. (A) Manhattan plot showing genetic differentiation between regressed (n = 8) and nonregressed (n = 4) tumors. Blue points represent the most differentiated SNPs (top 0.1% of the empirical pFst distribution; $P \le 0.0137$). SNPs were considered candidates if they were nonsynonymous or occurred in a putative regulatory region. Red points represent the two candidates identified in the 5' untranslated regions (UTRs) of *RASL11A* and *ERICH6* (P = 0.0105). The *x*-axis represents genomic position using the Tasmanian devil reference genome. The *y*-axis represents the negative log of the pFst statistic with genetic differentiation increasing along the axis. (B), Photos of a regressed tumor found in an adult female Tasmanian devil. Photo on the left was taken in August 2011. Photo on the right was taken in May 2012. *Photo credit*: Rodrigo Hamede. (C), Alignment of the 5' UTR of *RASL11A*. Both *ERICH6* and *RASL11A* showed a near fixed genotypic difference at the candidate sites; only *RASL11A* is shown here. All four nonregressed tumors were homozygous reference, and six of the seven genotyped regressed tumors were heterozygous (represented by Y). The only regressed tumor that was not heterozygous at either locus (Regressed 3) indicated with an * above) was a unique case where a single Tasmanian devil possessed both regressed (Regressed 3) and nonregressed (*n* = 2) and nonregressed tumor (Regressed 8) was not genotyped at either site due to missing data. (D), *RASL11A* expression in regressed (n = 2) tumor transcriptomes. *RASL11A* was differentially expressed, although the trend was not significant due to sample size limitations (P = 0.0769). The *y*-axis represents log₂*FPKM* + 1. *ERICH6* was not expressed in any transcriptome.

in regressed tumors but was not expressed in nonregressed (*i.e.*, wild-type) tumors (Figure 2D), consistent with its putative role as a tumor suppressor and underexpression in human prostate cancer and colon cancer cell lines (Louro *et al.* 2004; Weber *et al.* 2005). Transcriptome sequence data confirmed the expression of the *RASL11A* regressed allele in both regressed tumor samples (one of which was not sequenced above; see "*Materials and Methods*" for details), suggesting that the candidate variant in the 5' UTR of *RASL11A* may have led to the expression of this tumor suppressor in regressed tumors, and, ultimately, tumor regression.

In vitro assays confirm RASL11A expression reduces tumor cell proliferation

To directly test whether *RASL11A* expression negatively affected tumor cell proliferation or survival, we used lentiviral vector gene delivery to create cell lines that expressed *RASL11A* (*i.e.*, case lines) and those that did not (*i.e.*, eGFP control lines) across four tumor cell lines (three DFTD lines

and one DFT2 line) and a Tasmanian devil fibroblast cell line. RASL11A delivery was confirmed using qPCR (see Figure S2). We then measured cell proliferation and survival for all cell lines (7-18 replicates/line; see "Materials and Methods" and File S2). To determine if RASL11A expression reduced tumor proliferation significantly more often than expected by chance, we used a χ^2 test to compare the observed frequency of replicates where the RASL11A line had a lower, final viable cell count than the control line vs. the expected frequency (i.e., equal frequencies; see "Materials and Methods" for details). We found that RASL11A expression decreased overall cell proliferation in the tumor lines significantly more frequently than expected by chance (39/49 replicates; $\chi^2 = 17.163, df = 1, P < 0.0001$; File S2); DFTD lines $(32/42 \text{ replicates}; \chi^2 = 11.524, df = 1, P = 0.0007)$ and the DFT2 line (7/7 replicates; $\chi^2 = 7.000$, df = 1, P = 0.0082) remained significant when analyzed independently. Conversely, RASL11A expression significantly increased cell proliferation in Tasmanian devil fibroblasts more frequently than expected by chance (17/18 replicates; $\chi^2 = 14.222$,

df = 1, P = 0.0002; File S2). These discordant effects of *RASL11A* expression on cell proliferation across host and tumor cell lines were not surprising as genes can often function in opposite directions across different cell types, especially cancer (Vogelstein *et al.* 2013).

We next calculated RASL11A line relative proliferation (RASL11A line proliferation/control line proliferation) for each replicate across all cell lines. Here, ratios <1 indicated reduced proliferation under RASL11A expression, ratios >1 indicated increased proliferation under RASL11A expression, and ratios \sim 1, the null hypothesis, represented equal proliferation across RASL11A and control lines. RASL11A expression significantly reduced cell proliferation in three of the four tumor cell lines $(0.0001 \le P \le 0.0256)$; DFTD-b did not show a significant difference across the RASL11A and control lines (P = 0.0783). RASL11A expression significantly increased cell proliferation in Tasmanian devil fibroblasts (P = 0.0287; Figure 3), as expected. Here, the DFTD-b line may have adapted to the culture environment (e.g., compensatory mutation may have restored tumor growth rates in the RASL11A+ line). We also cannot rule out the role of host response (Pye et al. 2016a; Margres et al. 2018) or other factors independent of RASL11A expression that may have contributed to spontaneous tumor regression. Nonetheless, our results suggest an important, but not exclusive, role of RASL11A expression in tumor regression.

Discussion

We present several lines of evidence that RASL11A contributes to spontaneous, natural tumor regression. First, we used wholegenome sequencing to identify a near fixed genotypic difference between regressed and nonregressed tumors in a putative regulatory region for the tumor suppressor gene RASL11A. Second, we used transcriptomics, albeit in a few samples, to demonstrate that RASL11A was silenced in wild-type, nonregressed tumors but activated in regressed tumors, consistent with the genotypic data as well as RASL11A expression in human prostate (Louro et al. 2004) and colon (Weber et al. 2005) cancers. Third, we used cell proliferation assays to show that RASL11A activation significantly and negatively affected cancer cell proliferation in vitro in three of the four tumor cell lines investigated. Although we sequenced all confirmed cases of tumor regression at the time of the study, we recognized the limitations our sample size placed on interpreting the genomic and transcriptomic analyses. Therefore, by using functional genomics to show that RASL11A expression significantly impeded tumor cell proliferation in vitro across multiple, independent cell lines representing two independent transmissible cancers (DFTD and DFT2), our results collectively indicated that spontaneous cancer remission in Tasmanian devils is mediated, in part, by the expression of the tumor suppressor RASL11A.

The evolutionary implications of the *RASL11A* allele/ regression phenotype are still largely unclear. Trade-offs between pathogen transmission and virulence have been widely



Figure 3 RASL11A expression reduced tumor cell proliferation in vitro. Cell Titer-Glo assays were used to estimate the amount of ATP present, a proxy for the amount of metabolically active cells and cell proliferation, for control (- RASL11A expression) and RASL11A (+RASL11A expression) lines for the five cells lines in culture: Tasmanian devil fibroblast line (Devil), three DFTD lines (DFTD-a-c), and one DFT2 line. ATP was guantified at days 0.5, 3, 4, 6, 7, and/or 11. Means were estimated for days 0, 0.5, 3, 7, and 11. Relative proliferation was calculated as the ratio of RASL11A line ATP levels to control line ATP levels for each replicate at each timepoint and is shown on the y-axis. Ratios <1 indicate reduced proliferation under RASL11A expression, ratios >1 indicate increased proliferation under RASL11A expression, and ratios =1 (represented by the horizontal dashed line) represent equal proliferation across RASL11A and control lines. Colored points represent individual replicates, black points represent the mean estimate across all replicates for that time point, and error bars represent the SE. P-values were calculated using a one-sample t-test to determine if final relative mean proliferation significantly differed from one (i.e., the null hypothesis that proliferation would be equivalent across RASL11A and control lines).

discussed [*e.g.*, reviewed in Cressler *et al.* (2016)], and successful transmission at the population level is directly related to the specific physiological interactions occurring within a single host. The *RASL11A* allele could be beneficial to DFTD fitness by increasing the duration of infection and opportunity for transmission (*i.e.*, reduced virulence leads to greater

transmission), deleterious to DFTD fitness by reducing growth and, therefore, transmission rates, or the result of stochastic, neutral processes and not subjected to strong selective pressures. Additional geographic and temporal sampling are needed to truly understand (1) how *RASL11A*-mediated tumor regression will affect Tasmanian devil and DFTD evolution, and perhaps coevolution, going forward, and (2) whether the *RASL11A* allele/regression phenotype arose independently multiple times or a single time; our unrooted phylogenies (Figure S1) and limited sampling were unable to address these questions.

Although transmissible cancers are rare, with the only natural occurrences found in dogs [canine transmissible venereal tumor (CTVT; Murgia et al. 2006), bivalves (Metzger et al. 2016), and Tasmanian devils (Pearse and Swift 2006; Pye et al. 2016b], CTVT has also been shown to naturally regress. Here, the tumor grows progressively (P-phase) prior to spontaneous regression (R-phase), and this regression has been hypothesized to be the result of reduced tumor cell proliferation (Chu et al. 2001), similar to our results. More recent work examined vincristine-mediated CTVT regression and found that, following treatment, regression occurred in sequential steps involving immune system activation and infiltration of the tumor (Frampton et al. 2018). Although this work focused on tumor regression following vincristine treatment, host innate immunity certainly can play an important role in triggering cancer regression. Indeed, previous work on DFTD has also shown evidence of host immune response associating with spontaneous regression (Pye et al. 2016a). T lymphocyte infiltration, serum antibodies, and other immune responses could explain (among other factors) why the RASL11A allele was not fixed in regressed tumors. For each special case where the RASL11A allele did (Nonregressed 7) and did not (Regressed 3) lead to regression as expected, host immune response (Pye et al. 2016a), host genetic variation (Margres et al. 2018), and/or other factors may explain the discordant patterns. For example, Regressed 3 was the only regressed tumor in our study that was not heterozygous at the RASL11A allele. Regressed 3 was sampled from a Tasmanian devil that also possessed a nonregressed tumor (Nonregressed 3), and these tumors were sister to one another in one of our two phylogenetic analyses (Figure S1). Here, regression may be a product of serum antibody development following initial exposure to Nonregressed 3, with regression in the second tumor (*i.e.*, Regressed 3) being the product of host immune response following metastasis (Pye et al. 2016a). Additional immunological and histological work is needed to assess the relative roles host immunity, host genetics, and tumor genetics each play in relation to spontaneous tumor regression as RASL11A expression cannot universally be the sole driver of regression.

Although tumor regression in DFTD, like many tumor phenotypes, may have a polygenic basis (*e.g.*, ERBB inhibition was recently shown to also arrest DFTD cell growth *in vitro*; Kosack *et al.* 2019), interference with a single mutant

gene product can be sufficient to arrest cancer growth (Vogelstein et al. 2013). Indeed, a recent meta-analysis in human cancers found that one mutation is sufficient to affect RAS pathway function (Sanchez-Vega et al. 2018), and mutated RAS alleles are found in \sim 30% of all human cancers (Bos 1989; Way et al. 2018). RASL11A is underexpressed in human prostate cancer relative to normal prostate tissue (Louro et al. 2004) and is highly methylated (and not expressed) in colon cancer cell lines (Weber et al. 2005). Other than one recent study that found that RASL11A was also underexpressed in colorectal cancer (although RASL11A activation did not impede cancer growth in vitro; Wangsa et al. 2019), RASL11A function has been relatively unexplored. RASL11A is a known positive regulator of RNA polymerase I transcription and modulator of pre-rRNA synthesis (Pistoni et al. 2010), suggesting that the mechanism by which RASL11A inhibits DFTD growth may be through the dysregulation of rRNA transcription and processing. rRNA transcription is critical to cellular metabolism and ribosome biosynthesis, and misregulation of this process is a recurrent motif in human cancer due to the increased metabolic demands of uncontrolled cell growth (Roche et al. 2017). As such, rRNA synthesis is a key target in human cancer therapeutics (Drygin et al. 2009), and RASL11A activation may provide a therapeutic treatment option for Tasmanian devils as well as a mechanism for tumor inhibition in certain human cancers.

Spontaneous tumor regression in human cancers only occurs in ~1 of 60,000–100,000 cases (~0.001%; Missotten *et al.* 2008; Sengupta *et al.* 2010). Ethical considerations prevent the investigation of natural, untreated cancer progression in humans, highlighting the need for comparative oncological studies in nonhuman systems to identify the mechanisms underlying important tumor phenotypes such as spontaneous regression. Our findings in a system without therapeutic intervention and across two independently derived cancers highlight the relevance of *RASL11A* activation as a natural mechanism of tumor inhibition and a potential target in cancer therapy.

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