

Tumor RAS Gene Expression Levels Are Influenced by the Mutational Status of RAS Genes and Both Upstream and Downstream RAS Pathway Genes

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ABSTRACT: The 3 human RAS genes play pivotal roles regulating proliferation, differentiation, and survival in normal cells and become mutated in 15% to 20% of all human tumors and amplified in many others. In this report, we examined data from The Cancer Genome Atlas to investigate the relationship between RAS gene mutational status and messenger RNA expression. We show that all 3 RAS genes exhibit increased expression when they are mutated in a context-dependent manner. In the case of *KRAS*, this increase is manifested by a larger proportional increase in *KRAS4A* than *KRAS4B*, although both increase significantly. In addition, the mutational status of RAS genes can be associated with expression changes in other RAS genes, with most of these cases showing decreased expression. The mutational status associations with expression are recapitulated in cancer cell lines. Increases in expression are mediated by both copy number variation and contextual differences, including mutational status of epidermal growth factor receptor (*EGFR*) and *BRAF*. These findings potentially reveal an adaptive response during tumor evolution that is dependent on the mutational status of proximal genes in the RAS pathway and cellular context. Cell contextual differences in these adaptations may influence therapeutic responsiveness and alternative resistance mechanisms.

KEYWORDS: TCGA, *KRAS*, RAS Pathway

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Introduction

Since their discovery as virally transduced oncogenes and transforming genes in human tumor DNA, RAS genes have been shown to regulate many cellular processes including cell growth, differentiation, and survival.^{1,2} The *KRAS* gene produces 2 spliced isoforms, *KRAS4A* and *KRAS4B*,³ so that there are 4 primary protein products of these genes (*HRAS*, *NRAS*, *KRAS4A*, and *KRAS4B*). The Ras proteins are nearly identical except for differences in their carboxy-termini that give rise to differential posttranslational processing and trafficking mechanisms.⁴

Ras proteins act as molecular switches: the guanosine diphosphate (GDP)-bound form is inactive and guanosine triphosphate (GTP)-bound form is the active state. Guanosine triphosphatase (GTPase)-activating enzymes (GAPs) stimulate low-level intrinsic GTPase activity and convert Ras proteins to their inactive GDP-bound forms, whereas corresponding exchange enzymes (guanine nucleotide exchange factors [GEFs]) catalyze the replacement of GDP with GTP to activate the switch.⁵ In cancer, mutations (primarily in codons 12, 13, and 61) render Ras proteins resistant to GAP-assisted GTP hydrolysis resulting in a constitutively active RAS molecule.^{6,7} Less frequently, mutations occur (eg, at alanine 146) that lower the affinity of Ras proteins for GDP, allowing exchange of GDP for GTP without upstream signaling through GEFs.⁷ Importantly, these activating mutations in *KRAS* affect both the *KRAS4A* and *KRAS4B* isoforms.

Mutations in the *KRAS* gene occur in most pancreas and many lung, colorectal and uterine corpus endometrial cancers.

Less frequent *KRAS* mutations have been observed in cancers of the breast, cervix, hepatobiliary duct and testicular germ cell. Although less frequent, *NRAS* and *HRAS* mutations are also seen in many tumor types especially thyroid and melanoma (*NRAS*) and bladder cancer *HRAS*.⁷

In addition to point mutations that increase RAS activity, RAS genes are often amplified in human cancers. For example, *KRAS* is frequently amplified in ovarian, lung squamous cell, uterine, adrenocortical, and esophageal tumors. This suggests that both mutation and expression levels contribute to the roles that RAS genes play in oncogenesis. Given renewed efforts underway to target mutant RAS cancers^{8–11} and the observation that inhibitors of downstream effectors show context-dependent therapeutic efficacies and alternative resistance mechanisms, efforts to better understand the relationships between genes within the RAS pathway could improve our ability to predict clinical outcomes and anticipate the development of resistance.

As one of the largest public repositories of cancer genomic data, The Cancer Genome Atlas (TCGA) project has produced a wealth of new insights into cancer biology and the underlying driver events that give rise to the disease (see <http://cancergenome.nih.gov/abouttcga/overview>). The Cancer Genome Atlas data are publicly available and contain very broad data modalities spanning many different genomic platforms and associated patient clinical information and as such represent a considerable data mining resource. To date, several detailed integrative analyses of data from many of the individual tumor types have been



reported (see <https://tcga-data.nci.nih.gov/docs/publications/> and <http://cancergenome.nih.gov/publications>). In addition, a series of reports have detailed “pan-cancer” analyses where data aggregated across many tumor types and genomics platforms were used to identify similarities and differences among and between different tumor types (<http://www.nature.com/tcga/>). To more directly benefit the Ras community, a Ras gene-specific survey is desperately needed.

In this report, we take advantage of the wealth of clinical specimens available in TCGA across multiple tumor types where mutation information derived from either whole-exome or whole-genome sequencing and RNASeq-based expression analysis data are available. We analyzed these data to determine the relationship between RAS gene mutation status and expression levels. We show that mutation-associated expression increases occur with all 4 RAS isoforms and also correlates with the mutational status of other RAS pathway genes. These findings suggest that tumor type-specific gene-gene co-expression networks can be heavily influenced by gene mutational status.

Methods

Data preparation

The Cancer Genome Atlas data were obtained from the standard release folder from the Broad DCC firehose (http://gdac.broadinstitute.org/runs/stddata__latest/) for each available tumor type. In addition, the transcript-level messenger RNA expression data were obtained from the TCGA data portal (<https://cancergenome.nih.gov/publications>) for each individual sample for each tumor type and merged into a single matrix file per tumor type using R (<https://www.r-project.org>). Following download, the level 3 data were parsed into a table format where each sample/attribute/value becomes a separate database row. These files were then directly loaded into our Oracle 11g database. Additional tables corresponding to summary information derived during the parsing process were also prepared to facilitate the querying process. The data were produced using the July 2016 data release. The tumor type codes used throughout this article can be found on the TCGA Web site (<https://tcga-data.nci.nih.gov/tcga/>). Tumor purity data were obtained from the synapse portal (<https://www.synapse.org>) that was made available through a collaboration with TCGA/Broad Institute. The copy number variation (CNV) data were obtained from the DCC analysis page (<https://confluence.broadinstitute.org/display/GDAC/Dashboard-Analyses>) and we loaded the GISTIC (Genomic Identification of Significant Targets in Cancer)¹² data into oracle using the method outlined above (see <https://www.biostars.org/p/133927/>). The downloaded RNA-Seq expression data used were derived using the RSEM package¹³ and all expression data were log₂ transformed.

Cancer Cell Line Encyclopedia (CCLE) data were downloaded from the following Web site: <http://www.broadinstitute.org/ccle/home>, and the mutation and expression files were

then analyzed using standard R analysis workflows (see below). The CCLE RNASeq data were downloaded from CGHub and processed through the Cufflinks suite¹⁴ to quantify genes and transcripts using standard RNA-Seq workflows. The resulting gene and transcript expression levels, as RPKM (reads per kilobase per million), were then loaded into our Oracle database as described above.

Data analysis

We used the ROracle Database interface package (<https://cran.r-project.org/web/packages/ROracle/index.html>) to connect from R to the Oracle TCGA database instance and a series of wrapper functions to facilitate querying. For example, a function to extract a matrix of gene expression data for all samples given and input gene list, tumor type, and RNA-Seq platform was called for each tumor type, and then, a second function was called to subset the samples into tumor-normal pairs. The resulting list of data matrices is subsequently subdivided by mutation or CNV status (using the GISTIC values¹²) and then passed to plotting functions for the production of box plots, barplots, and other distribution plots. All plots and statistical tests were produced and performed using built-in R functions. Figures were produced interactively in R using the data matrices produced by those functions and manually subsetting samples.

Results

KRAS expression is increased in samples harboring *KRAS* mutations

We first focused on *KRAS* and its *KRAS4A* and *KRAS4B* isoforms within the TCGA tumor types in which *KRAS* is frequently mutated. Figure 1A shows expression levels for *KRAS* in lung adenocarcinoma (LUAD), pancreatic ductal adenocarcinoma (PAAD), and colon adenocarcinoma (COAD) samples. In each tumor type, we observed statistically significant increases in expression of mutant *KRAS* using the *t* test. The values are shown on the figure along with the numbers of samples for each condition.

Close examination of the plots also shows that while the overall separation between *KRAS*-mutant and wild type (WT) tumors in expression is significant, the distributions do overlap. To delineate this further, we plotted the combined WT and mutant samples for LUAD ordered by *KRAS* expression and then colored the bars by their mutation status (Figure S1). As expected, there are *KRAS*WT samples showing higher expression than some *KRAS*-mutant samples and vice versa.

We also explored whether mutations in other frequently mutated genes tracked with *KRAS* expression levels. In the bars at the bottom of the plot, the positions of samples with mutations in *TP53*, epidermal growth factor receptor (*EGFR*), *KEAP1*, and *NF1* are indicated. Of these, the *EGFR* mutations do seem to be enriched toward the left side of the panel, indicating lowered *KRAS* expression. Although the *TP53*

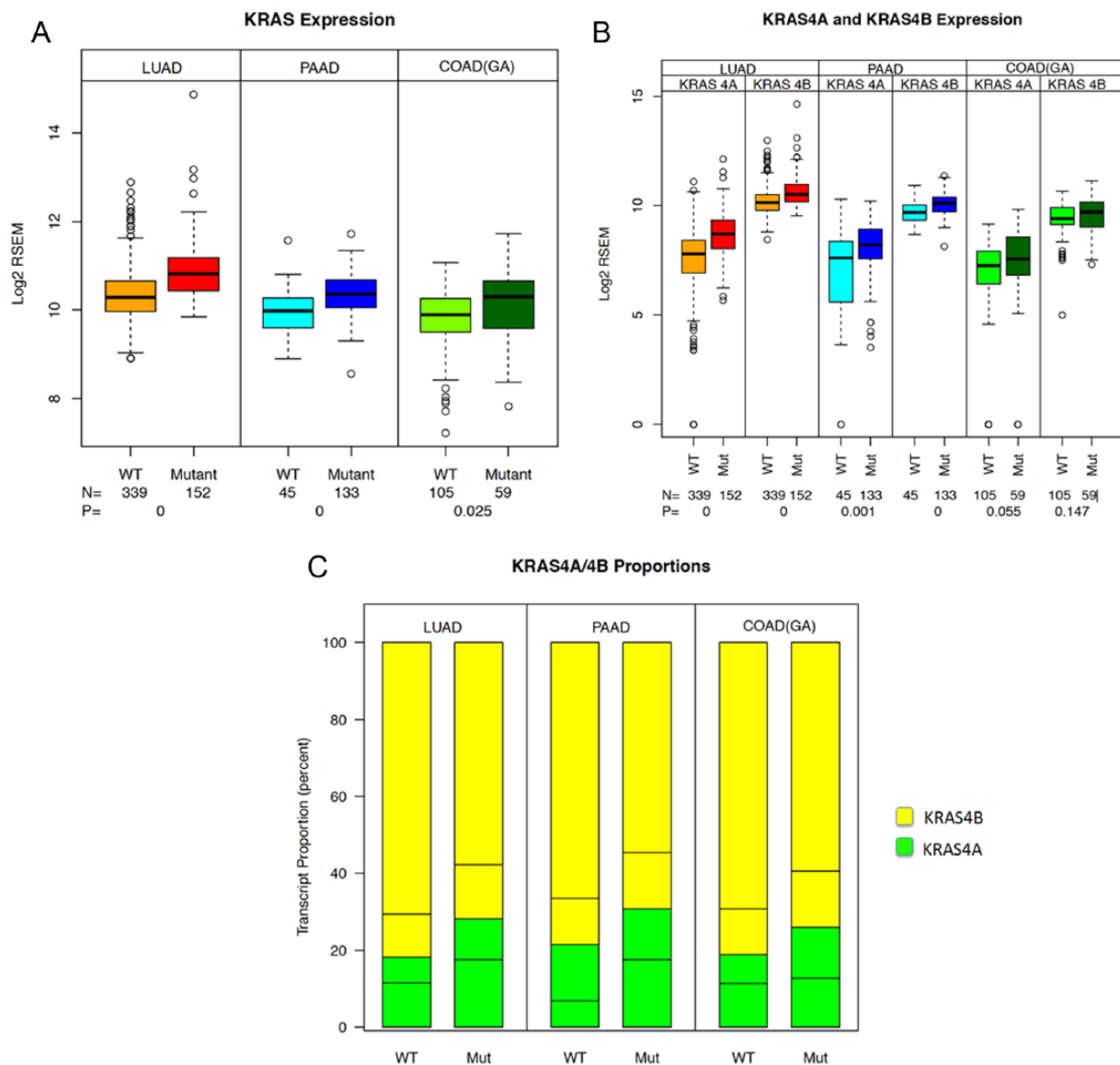


Figure 1. (A) *KRAS* expression is elevated in *KRAS*-mutant samples from lung pancreatic, and colon adenocarcinomas relative to WT samples. Sample subsets from each tumor type were produced according to their *KRAS* mutation status and the \log_2 -normalized RSEM¹³ data were plotted. The *P* values from *t* test between sample groups and the number of samples in each sample group are shown on the figure. Lane 1—*KRAS* WT samples from LUAD, lane 2—*KRAS*-mutant samples from LUAD, lane 3—*KRAS* WT samples from PAAD, lane 4—*KRAS*-mutant samples from PAAD, lane 5—*KRAS* WT samples from COAD, and lane 6—*KRAS*-mutant samples from COAD. (B) *KRAS4A* and *KRAS4B* expression is increased in *KRAS*-mutant samples from lung, pancreatic, and colon adenocarcinoma samples. The same sample subgroups from Figure 1A were used to derive the *KRAS4A* and *KRAS4B* expression. Lane 1—*KRAS4A* expression in *KRAS* WT LUAD samples, lane 2—*KRAS4A* expression in *KRAS*-mutant LUAD samples, lane 3—*KRAS4B* expression in *KRAS* WT samples from LUAD, lane 4—*KRAS4B* expression in *KRAS*-mutant samples from LUAD, lane 5—*KRAS4A* expression in *KRAS* WT PAAD samples, lane 6—*KRAS4A* expression in *KRAS*-mutant PAAD samples, lane 7—*KRAS4B* expression in *KRAS* WT samples from PAAD, lane 8—*KRAS4B* expression in *KRAS*-mutant samples from PAAD, lane 9—*KRAS4A* expression in *KRAS* WT COAD samples, lane 10—*KRAS4A* expression in *KRAS*-mutant COAD samples, lane 11—*KRAS4B* expression in *KRAS* WT samples from COAD, lane 12—*KRAS4B* expression in *KRAS*-mutant samples from COAD. Sample numbers are shown at the bottom of the figure along with the *P* values from a *t* test performed between the sample groups. (C) The proportion of *KRAS4A* is also increased in *KRAS*-mutant tumors. Median group expression is indicated by the green/yellow border in each bar. The bars correspond to the data groups from Figure 1B. The additional lines represent the 25th and 75th percentiles, respectively, of the sample proportion of *KRAS4A* and *KRAS4B*. All 3 tissues show an increase in the proportion of *KRAS4A* in mutant tumors. The lung and colon sets were significantly different using the raw expression values, but only the lung was significant using the \log_2 -transformed values. The numbers of samples in each group are the same as in Figure 1B. The *P* values are as follows: LUAD (339/152): raw: $2e-07$, \log_2 : $2e-10$; PAAD (45/133): raw: .034, \log_2 : .009; and COAD (105/59): raw: .019, \log_2 : .097. COAD indicates colon adenocarcinoma; LUAD, lung adenocarcinoma; PAAD, pancreatic ductal adenocarcinoma; WT, wild type.

mutations are enriched in the non-*KRAS*-mutated samples (data not shown), that relationship is not evident on the plot. We did not see any enrichment of either *KEAP1* or *NF1* in

terms of *KRAS* expression levels. The *EGFR* relationship is expected as *KRAS* mutations and *EGFR* mutations are mutually exclusive.^{15,16}

KRAS4A and KRAS4B isoforms both show increased expression in the context of KRAS gene mutations

Because the biological significance of these differentially trafficked isoforms is the focus of intensive study, and it has been difficult to tease apart their separate function, we next looked at *KRAS* isoform expression in samples harboring *KRAS* mutations. Figure 1B shows the expression of *KRAS4A* and *KRAS4B* across the same sample subsets presented in Figure 1A. Both isoforms track with the increase in mutated samples; however, the overall magnitude of the *KRAS4A* increase is slightly larger in all 3 tissues (about 1 log₂ unit compared with about 0.5 for *KRAS4B*). The differences were significant in both the lung and pancreas samples, but not in the colon samples, although the *KRAS4A* is borderline significant in that tissue (.055). To illustrate this point further, we plotted the proportions of the 2 isoforms using their median expression values in the WT and mutant samples (Figure 1C). In all 3 tumor types, there is an increase in the proportion of *KRAS4A* in the *KRAS*-mutant samples relative to that in the corresponding *KRAS* WT samples. The difference is significant in both the lung and colon samples (see figure legend for details). The biological significance of this is unclear, but it is consistent with a role for *KRAS4A* in tumorigenesis or maintenance of *KRAS* mutation-driven tumors as has been suggested in mouse studies.^{17,18}

Median values of the *KRAS4A* and *KRAS4B* expression values for each of the same sample subsets were converted to the proportional values and plotted. The additional lines in the bars represent the 25th and 75th percentiles, respectively, from bottom to top.

Beyond the 3 tumor types profiled above, we also evaluated the impact of *KRAS* mutation on its expression in other tumor types with at least 5 *KRAS*-mutant samples (including any nonsynonymous mutations, Figure S2A and B). We also observe significantly increased expression in the mutated samples relative to WT *KRAS* counterparts in uterine carcinosarcoma, stomach adenocarcinoma, and skin cutaneous melanoma (SKCM). Kidney renal papillary cell carcinoma (KIRP), uterine corpus endometrial cancers (UCECs), rectal adenocarcinoma (READ), cervical squamous cell cancers, breast cancers (BRCA) and liver hepatocellular carcinomas also all show a trend toward increased expression in mutant samples with KIRP displaying near significance (Figure S2A and B). Testicular germ cell tumors (TGCT), bladder cancer (BLCA), and thyroid cancer (THCA) fail to show the trend and thus appear to be exceptions (Figure S2A and B).

Increased expression is associated with mutation of other RAS genes

KRAS is the most frequently mutated RAS gene in cancer. However, both *HRAS* and *NRAS* genes can have driver roles,

although usually in different tumor types than those in which *KRAS* mutations are most frequent. Therefore, we wanted to determine whether the observed increase in *KRAS* expression associated with its mutation could be observed in *HRAS* or *NRAS*. As with *KRAS*, we assessed the potential expression shift for any tumor type where 5 or more *HRAS*-mutant or *NRAS*-mutant samples existed.

For *HRAS*, there are a total of 7 tumor types that meet these criteria—bladder (BLCA), head and neck squamous cell (HNSC), pheochromocytoma and paraganglioma (PCPG), lung squamous cell (LUSC), SKCM, thymoma (THYM), and THCA. Of these, we observed a statistically significant increase in *HRAS* in BLCA, HNSC, and PCPG tumors with *HRAS* mutations (Figure 2A). In SKCM, THYM, and THCA (Figure 2A), the trend was apparent, but the number of samples was too low for significance (values indicated on figure). As with the exceptions noted for *KRAS*, there was no evidence of the trend in LUSC (data not shown).

For *NRAS*, there are 8 tumor types that meet our selection criteria (5 or more mutated samples). Because of a historical issue, the data for COAD and READ are divided into 2 sequencing platforms for expression analysis (Illumina HiSeq [HS] and Illumina GA). They are as follows: SKCM, COAD (GA and HS), READ (HS and GA), acute myelogenous leukemia (LAML), TGCT, THCA, BLCA, and UCEC. Of these tumor types, SKCM (where *NRAS* has a presumed driver role) and READ (both in GA and HS) showed a significant increase in the *NRAS*-mutant sample subset relative to the WT subset (Figure S3A). Although not significant, the trend is also apparent in the COAD samples from both platforms. A weaker trend is present in the BLCA and UCEC tumor types (Figure S3B), but not in the TGCT, THCA (where *NRAS* mutation is considered a driver event) and LAML sample subsets (Figures S3A and B).

In the case of SKCM, the samples consist of both primary tumors and metastatic samples. Therefore, we can also examine the mutational effect against these different sample backgrounds. In both cases, the increase in *NRAS* expression in mutant samples relative to WT samples is observed, although it is only significant in the metastatic samples. Interestingly, the overall level of *NRAS* also appears to increase between the primary and metastatic samples, both in mutant and WT samples suggesting continued selective/adaptive forces influencing RAS gene expression levels.

The significant RAS gene expression differences between WT and mutated samples that we observed across the 3 RAS genes are summarized in Table S1. Across the 3 RAS genes, there were 34 possible settings where we evaluated the mutation-associated expression differences. Of these, we observed significant increases in 12 settings—7 for *KRAS*, 3 for *HRAS*, and 2 for *NRAS* (Table S1). Using a 1-tailed Fisher's exact test with 12 of 34 possible observations produces a *P* value of .003 suggesting that this observation would not be expected by

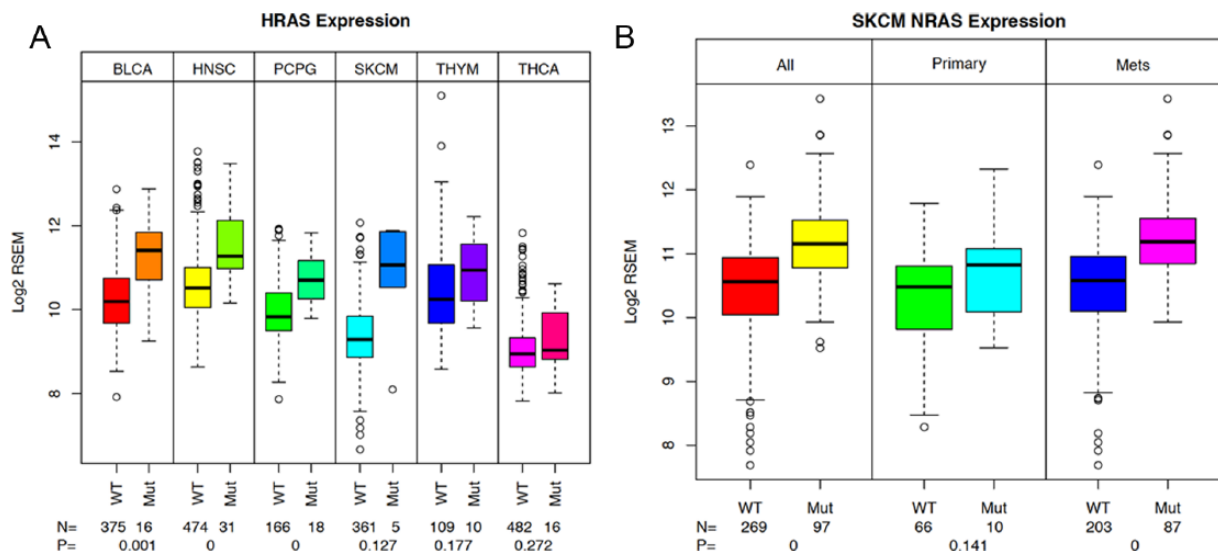


Figure 2. (A) *HRAS* expression also associates with mutation status. Sample subsets for each of the tumor types were prepared based on the *HRAS* mutation status. *HRAS* expression values were then plotted for each sample group. Sample numbers and *P* values are indicated on the plot. Lane 1—*HRAS* expression in *HRAS* WT samples from BLCA, lane 2—*HRAS* expression for *HRAS*-mutant samples from BLCA, lane 3—*HRAS* expression in *HRAS* WT samples from HNSC, lane 4—*HRAS* expression for *HRAS*-mutant samples from HNSC, lane 5—*HRAS* expression in *HRAS* WT samples from PCPG, lane 6—*HRAS* expression in *HRAS*-mutant samples from PCPG, lane 7—*HRAS* expression in *HRAS* WT samples from SKCM, *HRAS* expression in *HRAS*-mutant samples from SKCM. Lane 9—*HRAS* expression for *HRAS* WT samples from THYM, lane 10—*HRAS* expression for *HRAS*-mutant samples from THYM, lane 11—*HRAS* expression for *HRAS* WT samples from THCA, lane 12—*HRAS* expression from *HRAS*-mutant samples from THCA. (B) *NRAS* expression in SKCM sample subsets. Sample subsets were prepared for the total tumor set, the primary tumor set, and the metastatic sample set for both the *NRAS* WT and *NRAS*-mutant sample groups. Sample numbers for each group and *t* test *P* values are shown on the plot. Lane 1—*NRAS* expression in total *NRAS* WT subset of SKCM, lane 2—*NRAS* expression in *NRAS*-mutant samples from SKCM, lane 3—*NRAS* expression in *NRAS* WT primary tumor samples from SKCM, lane 4—*NRAS* expression in *NRAS*-mutant primary tumor samples from SKCM, lane 5—*NRAS* expression in *NRAS* WT metastatic tumor samples (mets) from SKCM, lane 6—*NRAS* expression in *NRAS*-mutant metastatic tumor samples (mets) from SKCM. BLCA indicates bladder cancer; HNSC, head and neck squamous cell; PCPG, pheochromocytoma and paraganglioma; SKCM, skin cutaneous melanoma; THCA, thyroid cancer; THYM, thymoma; WT, wild type.

chance. As this analysis involves multiple tests, a false discovery rate value was also computed for each of the observations. The complete set of results for all tumor types with 5 or more mutations in any of the RAS genes is shown in Table S2.

Mutation status of RAS genes can be associated with expression changes in other RAS genes

The recurrent association between RAS gene mutation status and increased expression (self-adaptation) and the highly redundant function of the RAS genes led us to evaluate the possibility that mutation in one type of RAS gene might be associated with expression changes in other RAS genes (non-self-adaptation). Therefore, we examined the tumor types outlined above to identify expression changes in other RAS genes against the backdrop of mutated RAS in each tumor type. There were a total of 68 tumor type settings in which we examined the association between mutation status and expression of other RAS genes. We observed a significant difference in 17 cases (6 cases with *HRAS* as the mutated gene, 4 with *NRAS*, and 7 with *KRAS*) suggesting that their adaptive influence extends beyond the mutated gene itself (Table 1). Of these settings, there were 7 effecting *NRAS* expression, 5 effecting *KRAS* expression, and 5 effecting *HRAS* expression. Using the

same criteria as above for the Fisher's exact statistical test, here, the *P* value is .0006 of this observation occurring by chance. As can be seen from Table 1, when the mutation of one RAS gene is significantly associated with a change in the expression of another WT RAS gene, the direction of the shift is downward (the WT RAS gene is lower) with the exceptions of *NRAS* expression in *KRAS*-mutant PAAD tumor samples and *HRAS* expression in *NRAS*-mutant UCEC samples which both increase. As with the "self" mutation associations (Table S1), we suspect that tumor/cellular context plays a large role in determining whether these shifts are observed, their extent, and their direction (see Table S2 for complete data).

CCLC cell lines with KRAS mutations also show an associated increase in KRAS expression

The tumor purity of the samples from TCGA has been estimated using copy number and other data¹⁹ and is typically in the range of 50% to 70%. Therefore, the expression data reflect the expression of both tumor and surrounding contaminating cells. We therefore analyzed expression data from cancer cell lines to verify our observations derived from TCGA data. We used the chip-based expression data from the CCLE Web site and cross-checked using the RNA-Seq data from caHUB

Table 1. RAS mutations and other RAS genes—assessment of mutational impact on RAS gene expression.

TUMOR TYPE	MUTANT GENE	EXPRESSION	MUTANTS	WILD TYPE	DIRECTION	P VALUE	FALSE DISCOVERY RATE
Testicular germ cell	<i>KRAS</i>	<i>NRAS</i>	20	135	Down	.001	0.007
Lung adenocarcinoma	<i>KRAS</i>	<i>NRAS</i>	152	339	Down	.011	0.068
Colon adenocarcinoma	<i>KRAS</i>	<i>NRAS</i>	112	96	Down	.004	0.029
Lung adenocarcinoma	<i>KRAS</i>	<i>HRAS</i>	152	339	Up	.001	0.006
Testicular germ cell	<i>KRAS</i>	<i>HRAS</i>	20	135	Down	.041	0.145
Pancreatic adenocarcinoma	<i>KRAS</i>	<i>NRAS</i>	133	45	Up	.012	0.145
Testicular germ cell	<i>KRAS</i>	<i>NRAS</i>	20	135	Down	.041	0.007
Testicular germ cell	<i>NRAS</i>	<i>KRAS</i>	8	147	Down	9e-5	0.002
Melanoma (skin)	<i>NRAS</i>	<i>HRAS</i>	97	269	Down	.002	0.018
Uterine corpus	<i>NRAS</i>	<i>HRAS</i>	9	232	Up	.035	0.134
Thyroid	<i>NRAS</i>	<i>KRAS</i>	40	458	Down	.046	0.151
Thymoma	<i>HRAS</i>	<i>NRAS</i>	10	109	Down	1e-4	0.002
Pheochromocytoma	<i>HRAS</i>	<i>NRAS</i>	18	166	Down	.002	0.017
Thyroid	<i>HRAS</i>	<i>NRAS</i>	16	482	Down	.005	0.026
Thyroid	<i>HRAS</i>	<i>KRAS</i>	16	482	Down	.036	0.026
Head and neck squamous cell	<i>HRAS</i>	<i>KRAS</i>	31	474	Down	.004	0.089
Thymoma	<i>HRAS</i>	<i>KRAS</i>	10	109	Down	.037	0.134

Data were prepared as in Table S1 except that only those cases where significant differences in expression between wild-type and mutant samples was observed in a gene that was different from the mutated gene. The data for all of the tests are shown in Table S3.

(Cancer Human Biobank) for the same lines (data not shown). Figure 3 shows that *KRAS* expression is significantly higher in *KRAS*-mutant lung-derived and also intestine (colon)-derived cell lines using the per-gene aggregated affy data (see Figure 3 for *P* values and sample numbers for each sample group). The trend is also present in the pancreas-derived lines, but there are very few *KRAS* WT lines available and so this was borderline significant (.054; Figure 3). A more direct comparison of the relative extent of these changes between the TCGA and CCLE samples is not permitted as the data are derived from different platforms and using different methods.

The CCLE collection is more limited in terms of mutations in *HRAS* and *NRAS*, which made validation on the other genes more difficult. This limitation also made the identification of examples where the non-self associated expression change was observed (ie. mutation of one RAS gene was associated with an expression change in another RAS gene) more difficult. We did observe trends in the non-self associated expression changes, but the differences were not significant. However, we do take this overall to show that our observations using tumor samples are reproducible in tissue culture systems making it possible to experimentally perturb cells and evaluate cellular responses at the transcriptional level. Such experiments

will allow further delineation of the underlying mechanisms responsible for our observations to be made.

KRAS expression and CNV

Given our observations showing frequent associations between RAS gene mutation status and expression, we wanted to pursue possible explanations for these effects.

We divided the LUAD samples into subsets based both on their *KRAS* mutational status and their *KRAS* copy number status. The results of this sample breakdown are shown in Figure 4. The first pair (all) shows the aggregate of all CNV states for either the WT or *KRAS*-mutant samples and shows the previously described expression increase in the mutant-harboring samples. The additional pairs represent 1-allele loss (Loss=1), neutral (N=2), 1-allele gain (Gain=1), and 2-allele gain (Gain=2) for the WT and mutant samples (there are not adequate 2 allele loss samples available). As expected, CNV affects *KRAS* expression levels: there is a gradient of increased expression across the increasing CNV states, but in each pair, the difference between the *KRAS*-mutant and *KRAS* WT samples remains significant except for the 2-allele gain state where we suspect that the combination of low sample number

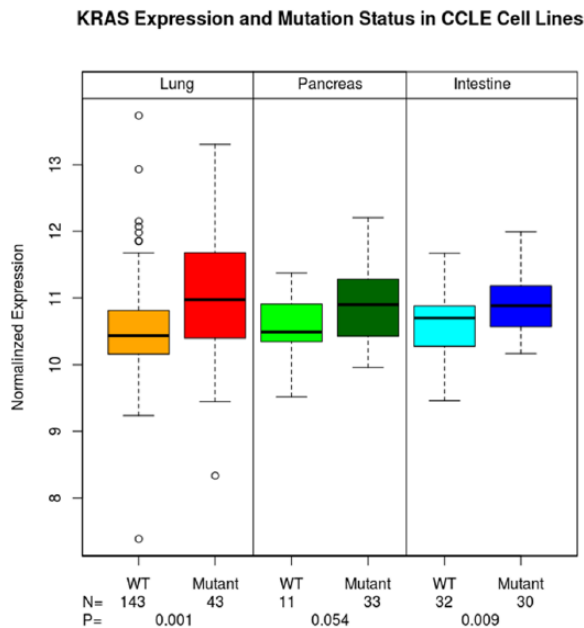


Figure 3. *KRAS* expression in *KRAS*-mutant CCLE cell lines. The Affymetrix expression data (gene level) were used and cell lines were divided by tissue source. Sample subsets were prepared from each of intestine (colon), lung, and pancreas tissues and separating *KRAS*-mutant and *KRAS* WT lines. Only protein-affecting mutations were included in the mutant lines. The numbers of lines and the *t* test *P* values are shown at the bottom of the plot. Lane 1—*KRAS* WT lung cell lines and *KRAS*-mutant lung lines, lane 3—*KRAS* WT intestine lines, lane 4—*KRAS*-mutant intestine lines, lane 5—*KRAS* WT pancreatic lines, and lane 6—*KRAS*-mutant pancreatic lines. CCLE indicates Cancer Cell Line Encyclopedia, WT, wild type.

and higher variance confounds the data. The *P* values and the sample numbers are indicated at the base of the figure. This finding confirms the contribution of CNV to *KRAS* expression increases, but as the *KRAS*-mutant samples consistently express a higher *KRAS* levels across the CNV continuum, there must be additional contributing factors. This result also confirms that at least in LUAD, for *KRAS*, there is a good correlation between *KRAS* copy number and expression using the derived GISTIC values. We also observed the expected increases in *KRAS* expression in pancreas and colon samples as a function of copy number, but they were not significant (Table S3).

The CNV analysis can also be used to address the question of whether CNV gains (or losses) are more common in the *KRAS*-mutated samples than in the WT samples. For this, we produced a frequency display of each CNV state shown in Figure S4 for the lung, pancreas, and colon samples. We applied a χ^2 test as well as visual inspection to evaluate the data. Of the 3 tumor types, both the lung and colon samples showed a significant enrichment for amplification in the mutant samples relative to their WT counterparts (.002 and 0, respectively). We did not see enrichment for deletion events in any of the tumor types between the mutant and WT samples. The finding that amplification is associated with mutation status in colon and lung samples, but not pancreas may simply result from low

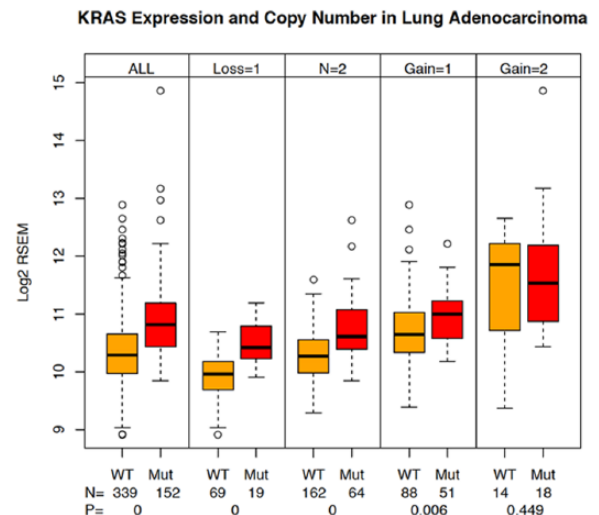


Figure 4. *KRAS* expression and CNV in lung adenocarcinoma samples. Sample subsets were prepared from the lung adenocarcinoma tumor samples to capture pairs (*KRAS* wild type and *KRAS* mutant) of samples in each of the 5 possible CNV states (−2, −1, 0, +1, +2) using the GISTIC scores.¹² *KRAS* expression was plotted for each sample subset. The numbers of samples in each group and the associated *t* test *P* values are shown on the plot. Lane 1—all *KRAS* WT samples, lane 2—all *KRAS*-mutant samples, lane 3—*KRAS* WT samples with CNV=−1, lane 4—*KRAS*-mutant samples with CNV=−1, lane 5—*KRAS* WT samples with CNV=0, lane 6—*KRAS*-mutant samples with CNV=0, lane 7—*KRAS* WT samples with CNV=+1, lane 8—*KRAS*-mutant samples with CNV=+1, lane 9—*KRAS* WT samples with CNV=+2, lane 10—*KRAS*-mutant samples with CNV=+2. CNV indicates copy number variation; GISTIC, Genomic Identification of Significant Targets in Cancer; WT, wild type.

sample numbers in this analysis, or could be related to other pancreatic-specific tumor attributes (eg, context).

EGFR mutational status is also associated with *KRAS* expression

Mutation of *KRAS* and *EGFR* in LUAD samples is mutually exclusive.^{15,16} As a result, in the LUAD samples above (Figures 1, S1, and S2), the samples labeled as WT *KRAS* actually contain all of the *EGFR*-mutated samples. To probe the connection between background mutational status and *KRAS* expression further, we separated the *EGFR*-mutant samples from the double WT samples and again assessed *KRAS* expression. In Figure 5A, the expected elevated *KRAS* expression in the *KRAS*-mutant samples is shown (eg, compare lanes 2 and 4). However, we also see a trend toward diminished expression of *KRAS* in the *EGFR*-mutant samples, even relative to the double WT samples. Both the difference between *KRAS*-mutant samples and the WT/WT samples and between the WT/WT samples and the *EGFR*-mutant samples is significant. When we examine the expression in a uniform CNV neutral background, the *EGFR*-associated decrease can also be observed, although it loses significance (see Figure 5A). This observation suggests possible cross talk between *KRAS* and its immediate upstream pathway elements

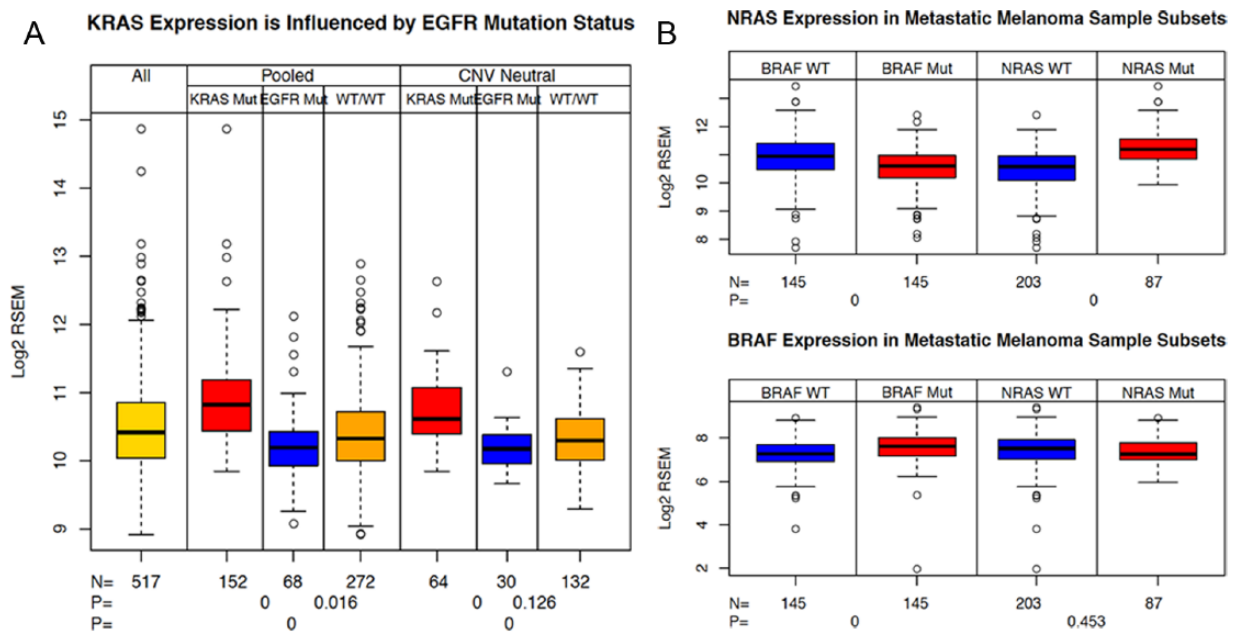


Figure 5. (A) *KRAS*-mutant and *EGFR*-mutant influences on *KRAS* expression in lung adenocarcinoma samples. Sample subsets were prepared by dividing samples by their mutation status (lanes 2-4) or by mutation status and CNV state=0 (lanes 5-7) and *KRAS* expression was plotted. The numbers of samples in each group and the *t* test *P* values from the class comparisons are shown on the plot. Lane 1—pooled tumor samples (*KRAS* mutant and WT), lane 2—pooled *KRAS*-mutant samples, lane 3—pooled *EGFR*-mutant samples, lane 4—pooled *EGFR*+*KRAS* WT samples, lane 5—CNV=0 *KRAS*-mutant samples, lane 6—CNV=0 *EGFR*-mutant samples, lane 7—CNV=0 *KRAS/EGFR* WT samples. (B) *NRAS* and *BRAF* mutations impact on both *BRAF* and *NRAS* expression in melanoma samples. Sample subsets were prepared using either *BRAF* mutation status or *NRAS* mutation status from all metastatic melanoma tumor samples (TCGA barcode=06) and either *NRAS* expression was plotted (top panel) or *BRAF* expression was plotted (lower panel). The sample counts and *t* test *P* values are shown on the plot. Lane 1—*BRAF* WT samples, lane 2—*BRAF*-mutant samples, lane 3—*NRAS* WT samples, lane 4—*NRAS*-mutant samples. *EGFR* indicates epidermal growth factor receptor; TCGA, The Cancer Genome Atlas; WT, wild type.

that through the adaptive process selects for decreased expression in a mutation selective manner. This observation extends the reported mutational exclusivity and crosstalk¹⁶ to include possible impacts of even WT co-expression of these genes.

To further explore this possible interaction, we then examined the expression of *EGFR* in the same sample subsets (Figure S5). As can be seen, the samples with mutated *EGFR* show elevated *EGFR* expression. However, in this case, the reciprocal decrease in *KRAS*-mutant samples was not observed as these show the same expression levels for *EGFR* as the double WT samples. In both *EGFR*-mutant and *KRAS*-mutant samples, it appears that more of an activated oncogene is selected for in a tumor and the possible cross talk extends this selective response to include additional pathway elements.

NRAS and *BRAF* mutual exclusivity in SKCM tumor samples

The LUAD, *EGFR*-mutant observation with *KRAS* suggests that the connections between mutational status and expression between genes within the RAS signaling node (*KRAS*, *NRAS*, and *HRAS*) extend to other RAS pathway genes. We evaluated whether this observation could be extended to other tumor types where mutational exclusivity has been demonstrated. For this, we applied a similar analysis to the *BRAF*-mutant and *NRAS*-mutant subsets of the metastatic melanoma tumor type.

The mutual exclusivity of the *NRAS* and *BRAF* mutations applies uniformly to fully activating mutations; however, in a limited number of cases, there are overlaps (eg, (*NRAS/BRAF*): G12D/G466E, Q61R/P348T and *HRAS*: Q61K/G469R, P34L/N581S). As can be seen in Figure 5B, *NRAS* expression is higher in samples where it is mutated and also diminished when *BRAF* is mutated, consistent with the mutual exclusivity observations made with *EGFR* (Figure 5A). As in lung, a subset of the *NRAS* WT samples overlaps the *BRAF*-mutant samples and vice versa. In both cases, these differences were significant. We also looked at *BRAF* expression between these sample classes and those results are shown in Figure 5B (lower panel). Although the difference is only significant in the *BRAF*-mutant samples for expression of *BRAF* (see Figure 5B), the results remain consistent with cross talk between these proximal pathway nodes.

Together, these results suggest a complex interaction between different genes within the RAS pathway, whereby mutation in a gene can be associated with differences in its own expression as well as changes in the expression in other genes in the pathway, both within the same node as with the RAS genes, but also extending both upstream (*EGFR*) and downstream (*BRAF*) in the pathway. Importantly, there must also be an additional context component because we did not observe this same pattern for *BRAF* and *NRAS* in the THCA tumor type samples (data not shown).

Discussion

Mutation-associated expression changes

Using data from TCGA, we have confirmed and extended the early observation that *KRAS* gene expression in samples harboring *KRAS* mutations is elevated relative to their WT *KRAS* counterparts in many, but not all tumor types. Notably, the increase is significant in the principal tumor types where *KRAS* mutation is thought to have a driver role (LUAD, PAAD, and COAD) (Figure 1A). We extended this to show that the same principle applies to the 2 other RAS genes and that it also applies to the 2 major spliced isoforms of *KRAS*, *KRAS4A*, and *KRAS4B* (Figure 1B and C). We also demonstrated that the mutation-associated expression increase could be reproduced in cell lines (Figure 3), providing a convenient experimental paradigm for further experiments.

Although selection for increased expression of an oncogenic driver appears intuitively obvious, the precise mechanism is unclear. Increased signal output may be necessary to overcome feedback mechanisms, consistent with the possibility that the feedback itself can be overwhelmed. Alternatively, increased levels of oncogenic Ras proteins might engage effectors that are not engaged during normal signal transduction and thus provide a selective advantage through phenotypes that modulate malignant transformation. If so, this could offer novel therapeutic opportunities. Increased expression might also be necessary to overcome the suppressive effects of WT Ras proteins, which are known to interfere²⁰ with oncogenic signaling, at least for *KRAS* and *HRAS*. The *KRAS* expression increase is also consistent with the reported rewiring of the cellular circuitry to effect metabolic changes to manage increased reactive oxygen species.²¹ It is also important to mention that although our results show an increased *KRAS* expression in the *KRAS*-mutant harboring samples, the analysis does not distinguish which allele is being adjusted. Thus, the WT allele could also be compensating in some way as well.

Factors influencing expression changes

We also showed evidence of RAS signaling cross talk, whereby mutation in one RAS gene is associated with changes in the expression of other RAS genes. This observation is pertinent to the role of the WT RAS isoforms when one becomes mutated because it is possible that through cross talk, adjustments in expression could influence overall signal output as well as the ratios of outputs from multiple RAS gene-specific signaling cascades. Previously, a dependence on *HRAS* and *NRAS* in *KRAS*-driven cell proliferation has been described²⁰ consistent with the proposed interdependency.

Mechanistically, our analysis revealed good correlation between copy number and *KRAS* gene expression in LUAD samples (Figure 4). However, we showed that across different CNV states, the *KRAS*-mutated samples consistently expressed even higher levels of *KRAS* consistent with involvement of one

or more additional contributing factors. Referring to Figure S3, we observed a significant increase in *KRAS* amplification in mutant samples in the COAD data set, but not in the LUAD and PAAD data sets, suggesting an additional contextual impact at play. Similarly, as there are *KRAS* WT samples that express higher *KRAS* levels than some of the *KRAS*-mutant lung samples (Figure S1), there are also influences stemming from the individual—presumably including germline single-nucleotide polymorphisms and somatic mutation burdens.

Cross talk between RAS pathway genes

Because we observed gene expression changes in many but not all tumor types with sufficient sample size (Figures 1A and 2A, B; Tables 1 and S1), we evaluated the possibility that the changes are tissue dependent. We used LUAD samples and showed that mutually exclusive mutations in *EGFR* and *KRAS* appear to be diametrically opposed. When *EGFR* is mutated, there is an increase in *EGFR* expression and a corresponding decrease in *KRAS* expression, and in *KRAS*-mutant samples, *KRAS* expression is elevated with no apparent effect on *EGFR* expression levels (Figure 5A). These observations are consistent with the previously reported analysis of the *KRAS-EGFR* mutual exclusivity.^{14,15} In this report, attempts to introduce mutant *KRAS* into *EGFR*-mutant expression cell lines and introduce mutated *EGFR* into mutant *KRAS*-expressing lines failed to produce lines co-expressing these 2 products. The authors interpreted this as a sensing mechanism that detected too much signal resulting in the elimination of the co-expressing cells from the population that was selected. Our observations are consistent with this result but suggest that even the balance between a mutated *EGFR* or *KRAS* protein and the WT counterparts requires adjustment at the expression level, consistent with the sensing mechanism, but extending it to include mutant-WT interactions.

This possibility was further investigated in the analysis of SKCM tumors where a comparable association was observed between the mutation status of *NRAS* and *BRAF* and their expression demonstrating the ability of both upstream and downstream pathway genes to participate in the cross talk process (Figure 5B). We did not resolve why the observed expression changes were not observed in all cases; there could be confounding mutations or alternatively expressed genes that can more or less substitute for the increase in RAS gene expression.

Our data support a model where RAS gene expression in tumors is shaped by a number of complex factors including context, mutant status, and mutant status of both upstream and downstream regulatory molecules. Ideally, going forward, the data collected from clinical trials that are increasingly guided by tumor genotype information will allow for deeper analysis of possible outcome-genomic connectivity and more conclusive determinations of the role of cellular context and adaptive

responses to tumor evolution, as well as be used to guide therapeutic intervention to improve patient survival and outcome.

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Author Contributions

RS and MY performed the analysis of TCGA and CCLE data sets. RS wrote the first draft of the manuscript. MY and BK provided important suggestions for the revisions of the manuscript. FM and DN provided scientific guidance and conceived the hypothesis pursued in the analysis and also provided critical suggestions for the revision of the manuscript.

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