The immunoglobulin kappa variable cluster is differentially expressed in human epithelial ovarian cancer.

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Epithelial ovarian cancer (EOC) is the most lethal gynecologic cancer (1). We performed discovery of genes associated with epithelial ovarian cancer and of the high-grade serous ovarian cancer (HGSC) subtype, using published and public microarray data (2, 3) to compare global gene expression profiles of normal ovary or fallopian tube with that of primary tumors from women diagnosed with epithelial ovarian cancer or HGSC. We identified genes encoded by the immunoglobulin kappa variable cluster, IGKV, as among the genes whose expression was most different in epithelial ovarian cancer as compared to the normal fallopian tube. IGKV1D-8 expression was significantly higher in high-grade serous ovarian tumors relative to normal fallopian tube. These data indicate that expression of IGKV genes is perturbed in epithelial ovarian cancers broadly and in ovarian cancers of the HGSC subtype. IGKV may be relevant to pathways underlying ovarian cancer initiation (transformation) or progression.

Keywords: ovarian cancer, epithelial ovarian cancer, HGSC, high-grade serous ovarian cancer, systems biology of ovarian cancer, targeted therapeutics in ovarian cancer.

The five-year survival rate for women diagnosed with high-grade serous ovarian cancer is between 30-40% and has not changed significantly in decades (4, 5). The development of novel, targeted therapeutics to treat HGSC can be facilitated by an enhanced understanding of the transcriptional behavior of ovarian tumors relative to that of the normal ovary. We mined published and public microarray data (2, 3) to compare global gene expression profiles between human ovarian tumors, including that of the HGSC subtype, and that of normal ovarian and fallopian tissue. We identified the genes encoded by the immunoglobulin kappa variable cluster, IGKV, as among the most differentially expressed in HGSC tumors of the ovary and in epithelial ovarian cancer broadly. IGKV may be a gene of interest when prioritizing the study of target genes and pathways for the development of novel therapeutic interventions in epithelial ovarian cancer and specifically in high-grade serous ovarian cancers.

Methods

We used microarray data from datasets and GSE124766 (2) and GSE126308 (3) for this differential gene expression analysis of epithelial ovarian cancer in conjunction with GEO2R. The Benjamini and Hochberg method of p-value adjustment was used for ranking of differential expression but raw p-values were used for assessment of statistical significance of global differential expression. Log transformation of data was auto-detected, and the NCBI generated category of platform annotation was used. GSE124766 (2) was generated using Agilent-014850 Whole Human Genome Microarray 4x44K G4112F with $n=3$ normal fallopian tube tissue (control) and $n=7$ for tumors from patients with high-grade serous ovarian cancer; analysis was performed using platform GPL6480. GSE126308 (3) was generated using Affymetrix Human Transcriptome Array 2.0 with $n=20$ tumors from high-grade serous adenocarcinoma (HGSC) patients classified as early in disease progression and $n=30$ for tumors from HGSC patients classified as “late” in disease progression; analysis was performed using platform GPL17586.

A statistical test was performed to evaluate whether IGKV expression was significantly different when comparing normal ovarian tissue from control subjects and primary tumors from women diagnosed with HGSC using a two-tailed, unpaired t-test with Welch’s correction. We used PRISM for all statistical analyses (Version 8.4.0)(455).

Results

We mined published microarray data (2, 3) to identify differentially expressed genes in epithelial ovarian cancer and specifically in high-grade serous ovarian cancer (HGSC), the type of ovarian cancer responsible for 70-80% of deaths resulting from the most lethal gynecologic malignancy.

IGKV is differentially expressed in ovarian tumors from women diagnosed with HGSC

We identified a gene encoded within the IGKV cluster, IGKV1D-8, as among the genes whose expression was most different when comparing primary HGSC tumors to normal fallopian tube tissue (2) (Chart 1). When sorting all of the transcripts measured by microarray based on change in expression between HGSC and the normal fallopian tube, in this data set, IGKV1D-8 ranked 188 out of 40481 total transcripts (Chart 1), equating to 99.5% differential expression. Differential expression of IGKV1D-8 in HGSC tumors was statistically significant (Chart 1; $p=2.46E-05$).

We analyzed a second microarray dataset (3) generated using normal ovarian tissues and tumors from women diagnosed with epithelial ovarian cancer (EOC) to determine whether differential expression of IGKV could be observed in a separate cohort of patients with ovarian cancer. We found that a gene encoded by the IGKV cluster, IGKV2-29, was also differentially expressed when comparing epithelial ovarian cancers based on disease progression, as either “early” or “late” in disease progression (Chart 2).

When sorting all of the transcripts measured by microarray based on change in expression between tumors from patients early in disease progression and tumors from patients late in disease progression, IGKV ranked 287 of 70523 total transcripts (Chart 2), equating to 99.6% differential expression. Differential expression of IGKV in EOC based on disease progression was statistically significant (Chart 2; $p=8.30E-04$).

IGKV1D-8 is expressed at significantly higher levels in HGSC when compared to the normal fallopian tubes.

We then extracted exact mRNA expression levels for the differentially expressed IGKV1D-8 transcript from both normal fallopian tube tissue and from epithelial ovarian tumors. IGKV was expressed at significantly higher levels in high-grade serous ovarian cancers when compared to the normal fallopian tubes (Figure 1; $p<0.0055$). We calculated a mean fold change of 1.38 in IGKV1D-8 expression when comparing HGSC tumors to normal fallopian tissue, as IGKV1D-8 was expressed at 7.25 ± 0.63 arbitrary units (AU) in the fallopian tubes but at 10.03 ± 0.58 AU in tumors of the ovary.

Thus, we found using published microarray data (2, 3) that genes encoded by the IGKV cluster were among the genes most differentially expressed in tumors from women with high-grade serous carcinomas, and in epithelial ovarian cancer broadly. IGKV1D-8 expression was significantly higher in HGSC tumors when compared to the normal fallopian tubes.

Discussion

Transcriptional profiling of HGSC in a Cancer Genome Atlas integrated genomic analysis classified HGSC into four subtypes based on gene clustering: immunoreactive, differentiated, proliferative, and mesenchymal (6). We sought to describe the transcriptional landscape of epithelial ovarian cancers in an unbiased fashion by using published and public microarray data from primary tumors of women diagnosed with HGSC compared to transcriptome data from normal ovarian and fallopian tube tissue (2, 3). In both datasets analyzed (2, 3), IGKV cluster genes were among the genes whose expression changed most significantly transcriptome-wide in an ovarian cancer, when comparing normal gynecological tissue to ovarian tumors, or when comparing ovarian tumors from human across disease progression. Genes encoded within the IGKV gene cluster may be of relevance in understanding transformation or disease progression in human epithelial ovarian cancer.

References

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Rank: 188
Probe ID: A_24_P93523
p-value: 2.46E-05
t: -7.0462951
B: 3.053223
Gene: IGKV1D-8
Gene name: immunoglobulin kappa 1D-8

Chart 1: IGKV1D-8 is differentially expressed in high-grade serous ovarian carcinomas (HGSC).

The rank of differential expression, the probe/transcript ID, the p-value with respect to global differential expression, t, a moderated t-statistic, B, the log-odds of differential expression between the two groups compared, the gene and gene name are listed in this chart.

Rank: 287
Probe: TC02004947.hg.1
p-value: 8.30E-04
t: -3.56
B: -0.6194
Gene: IGKV2-29
Gene name: immunoglobulin kappa variable 2-29

Chart 2: IGKV2-29 is differentially expressed in epithelial ovarian cancers when comparing tumors based on tumor progression: early versus late.

The rank of differential expression, the probe/transcript ID, the p-value with respect to global differential expression, t, a moderated t-statistic, B, the log-odds of differential expression between the two groups compared, the gene and gene name are listed in this chart.

Figure 1: IGKV is expressed at significantly higher levels in HGSC tumors when compared to normal fallopian tube.

The mRNA expression of IGKV in normal fallopian tube tissues (left) and in the primary tumors of patients with HGSC (right) is represented with mean mRNA expression level marked and the result of a statistical test evaluating the significance of difference in IGKV expression between the normal fallopian tube and primary tumors from patients with HGSC, a *p*-value, listed above.