Prognostic association of starvation-induced genes in head and neck cancer

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Abstract

Autophagy-related genes (ARGs) have been implicated in the initiation and progression of malignant tumor promotion. To investigate the dynamics of expression of genes, including ARGs, head and neck squamous cell carcinoma (HNSCC) cells were placed under serum-free conditions to induce growth retardation and autophagy, and these starved cells were subjected to transcriptome analysis. Among the 21 starvation-induced genes (SIGs) located in the autophagy, cell proliferation, and survival signaling pathways, we identified SIGs that showed prominent up-regulation or down-regulation. These included AGR2, BST2, CALR, CD22, DDIT3, FOXA2, HSPA5, PIWIL4, PYCR1, SGK3, and TRIB3. The Cancer Genome Atlas (TCGA) database of HNSCC patients was used to examine the expression of up-regulated genes, and CALR, HSPA5, and TRIB3 were highly expressed relative to solid normal tissue in cancer and the survival rate was found to be reduced in patients with high expression. Protein-protein interaction analysis demonstrated the formation of a dense network of these genes. Cox regression analysis revealed that high expression of CALR, HSPA5, and TRIB3 was associated with poor prognosis in patients with TCGA-HNSCC. Therefore, these SIGs up-regulated under serum starvation may be molecular prognostic markers in HNSCC patients.

1. Introduction

Head and neck cancer is the sixth most common malignancy in the world, 90-95% of which is squamous cell carcinoma (SCC). Over 60% of patients already have advanced cancer at the time of their first visit, with an estimated 5-year survival rate of 40-50%¹⁻⁴. Surgery, radiation, chemotherapy, and targeted therapies are used to treat head and neck squamous cell carcinoma (HNSCC)^{5,6}. In recent years, immunotherapy with antibodies that target the immune checkpoint pathway has been introduced and has shown long-term effects on cisplatin-resistant cancer, distant metastases, and recurrence of poor prognosis^{7,8}. However, valid cases are limited to 18-25% of advanced HNSCCs⁹. Long-term immunological side effects are also a problem¹⁰. Effective indications for these therapies need to be developed and new therapies need to be developed.

A major advance in recent HNSCC research is the aggregation of extensive genetic analysis results of HNSCC¹¹⁻¹⁵. A typical HNSCC database is the Cancer Genome Atlas (TCGA), published in 2015. Recent technological advances have enabled TCGA and other large-scale genomics studies to determine the broader landscape and frequency of chromosomal alterations, mutations, and expressed genes that contribute to HNSCC pathogenesis, prognosis, and resistance to therapy¹¹⁻¹⁵. The TCGA-HNSCC database may be used to screen for differentially expressed genes (DEGs) in cancer and normal tissue transcriptome studies in HNSCC patients^{16,17}. Furthermore, using the TCGA database, many studies on the deviation of genes and signaling pathways involved in carcinogenesis and prognosis are being conducted. This includes studies on hypoxia-immune signature¹⁸, cancer-associated alternative splicing event-related genes¹⁹, the miRNA-30 family²⁰⁻²², and the KEAP1-NRF2-CUL3 axis²³. Consequently, promising biomarker genes for the prognosis of HNSCC patients have been proposed.

Autophagy is an advanced process of digesting the cytoplasm and organelles by autophagosomes and autolysosomes to protect cells, and thereby, cell components become an energy source by recycling ^{24,25}. Autophagy is also considered a strong promoter of metabolic homeostasis, as it has been shown to play an important role in the regulation of several survival and death signaling pathways that determine the cell fate of cancer ²⁶⁻²⁹. On the other hand, antineoplastic agents, such as survivin inhibitors and disulfiram, may promote autophagic cell death in HNSCC cells, thus showing the opposite role of autophagy on cell survival ^{30,31}. The complex multi-step process of autophagy is tightly controlled by a set of autophagy-related genes (ARGs). Some ARGs have been shown to be associated with the prognosis of HNSCC patients using bioinformatics ^{29,32-34}. However, it has not been clarified how these ARGs exhibit their expression kinetics in an environment where autophagy occurs.

Serum starvation is the most widely studied method for inducing autophagy ³⁵⁻³⁸. In HeLa cells, mitochondria-produced reactive oxygen species (ROS) are also known to induce autophagy via AMPK during starvation ³⁹. In the present study, we investigated whether genes containing ARG were induced by serum starvation of HNSCC cells, and whether genes with large expression fluctuations were also expressed in tumors and normal solid tissues of TCGA-HNSCC patients. The results of this study suggest that, among the up-regulated genes under serum starvation, CALR, HSPA5, and TRIB3 are starvation-induced genes (SIGs) associated with the prognosis of TCGA-HNSCC patients.

2. Materials And Methods

2.1. Cells

The human HNSCC cell line SAS cells were obtained from the Japanese Collection of Research Bioresources (Tokyo, Japan). Cells were cultured in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum (FBS) at 37°C in a humidified atmosphere with 5% CO₂.

2.2. Transmission electron microscopy (TEM)

TEM was performed to observe SAS cells in serum-starved condition for 24 hours. Serum-starved cells were washed with PBS, fixed in 2.5% glutaraldehyde in phosphate buffer, post-fixed in 2% osmium tetroxide, dehydrated in graded ethanol, and then embedded in epoxy resin. Ultrathin sections were stained with 2% uranyl acetate and observed using a JEM-1200 EX microscope (JEOL, Tokyo, Japan).

2.3. Cell proliferation assay and migration assay

For MTT assay, starved or control SAS cells were incubated with 3-(4,5-Dimethyl-2-thiazolyl)-2,5-diphenyl-2H-tetrazolium bromide reagent (DOJINDO, Osaka, Japan) for 2 hours at 37°C. At the end of each experiment, the medium was removed and 100 µL solution of 4% HCl 1N in isopropanol was added to immediately dissolve the formazan crystals, and absorbance at 570 nm was recorded. For the migration assay, SAS cells were cultured in DMEM with 10% FCS until confluent. The cell layers were scratched using a plastic tip, as previously described ⁴⁰. The cells were further incubated in DMEM with/without

FBS for 6 hours. The closure rate of each scratched area was measured using ImageJ software, as previously described ⁴⁰.

2.4. RNA extraction

Total RNA from SAS cells were isolated using TRIsure (BIOLINE, Luckenwalde, Germany) according to the manufacturer's instructions.

2.5. RNA-sequencing and FASTQ file processing

According to the manufacturer's instructions, library preparation was performed using a TruSeq stranded mRNA sample prep kit (Illumina, San Diego, CA). Whole transcriptome sequencing was executed with the Illumina HiSeq 2500 platform in a 75-base single-end mode. Illumina Casava ver.1.8.2 software was used for base calling. Sequenced reads were mapped to the human reference genome sequences (hg19) using TopHat ver. 2.0.13 in combination with Bowtie2 ver. 2.2.3 and SAMtools ver. 0.1.19. Counts per gene were calculated with Cufflinks ver. 2.2.1. FPKMs and fragment counts were scaled via the median of the geometric means of fragment counts across all libraries.

2.6. Analyzing the normalized counts data

We imported the normalized counts into Subio Platform v1.24.5849 (Subio Inc. Kagoshima, Japan) ⁴¹ and all subsequent analyses were executed using this software. We set the lower limit as replacing positive numbers less than 10 with 10, and 0 counts with 8. Then, we calculated the log2 ratio against the geometric mean of the two control samples. We filtered out genes if their counts were always less than 15, or if their log2 ratios were between -0.5 and 0.5 in all samples; a total of 6,363 genes remained after filtering. We extracted candidate DEGs by a 2-fold criterion.

2.7. Analyzing TCGA-HNSC RNA-Seq data

We obtained and analyzed the RNA-Seq count data of TCGA-HNSC from the GDC Data Portal ⁴² with the Subio platform. The workflow of TCGA RNA-Seq was the same as that applied to our RNA-seq data except for the thresholds. The lower limit for positive counts was 50, for 0 counts was 32, and the filter on counts was 50, and that on log2 ratios was between -1 and 1. In addition, the log ratios were taken against the average of solid normal tissue samples. For each of the 21 selected genes, we divided the primary tumor samples into two groups, those with count values higher or lower than the median, to compare the survival time with the Kaplan-Meier method.

2.8. Pathways analysis and Protein and Protein interaction

The molecular pathways of the 21 selected genes were analyzed for gene ontology (GO) terms and Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways using the Database for Annotation, Visualization, and Integrated Discovery (DAVID) server. GO enrichment was carried out over three primary levels: cellular components (CC), biological processes (BP), and molecular functions (MF). Based on the STRING online

database (<https://string-db.org/>), we used these genes to establish a protein-protein interaction (PPI) network. Then, the most significant modules in the PPI networks were visualized.

2.9. Statistical analyses

Statistical analyses were performed using the Student's *t*-test with Microsoft Excel (Microsoft, Redmond, WA, USA). Results were expressed as the mean $\pm$ SD. Differences were considered significant at $P < 0.05$. For the survival analysis shown in Table 3, the hazard ratio (HR) relative to the indicated reference (ref) value, its 95% confidence interval (CI), and P-value (those of < 0.05 are indicated in bold) for the Cox hazard model are shown. The HR and its 95% CI were calculated by Cox regression analysis after proper evaluation of the assumptions of the Cox regression models with the use of the survival package.

3. Results

3.1. Effects of serum starvation on the biological activity of head and neck cancer SAS cells

We first examined the effect of serum starvation on cell morphology and proliferation of SAS cells. The morphology of SAS cells was not affected in the absence of serum for 24 hours (Fig. S1). When examined by MTT assay, cell proliferation decreased in a time-dependent manner in the absence of serum (Fig. 1A). As for migration assays, SAS cell migration was considerably diminished in serum-free conditions (Fig. 1B, C). Morphological changes at the hyperfine structure level due to serum starvation were further investigated through TEM. Most SAS cells contained intact mitochondria that were distributed throughout the uniform cytoplasm in the presence of serum. When the cells were cultured in the absence of serum for 24 hours, the intracellular space was occupied by large vascular structures. There were autophagosomes and/or autolysosomes containing degraded mitochondria and dense structures (Fig. 1D).

3.2. RNA-sequencing of serum-starved SAS cells to examine the expression of autophagy-related genes

SAS cells were cultured in the absence of serum for 2 and 24 hours and ARG expression was examined by RNA sequencing. A slight change in gene expression was observed 2 hours after the onset of starvation. Some genes, such as DDIT3 and ERN1, were down-regulated after 2 hours of starvation but increased after 24 hours. After 24-hour serum starvation, more than two-fold up-regulation was observed for 12 genes (ATP6V0A2, ATP6V1B1, ATP6V1C2, DDIT3, ERN1, NHLRC1, NUPR1, PIM2, TMEM150A, TRIB3, WIPI1, and XBP1) (Table 1). On the other hand, down-regulation of 50% or more was observed for 13 genes (BNIP3, BNIP3L, C10orf10, DAPK2, GAPDH, HMOX1, MEFV, PLK2, RRAGD, SESN3, SRPX, S100A8, and S100A9) (Table 1).

3.3. RNA-sequencing of serum-starved SAS cells to detect the expression of genes related to cell growth, cell death, cell migration, cell proliferation, cell cycle, and cell adhesion

Since 24-hour serum starvation reduced the ability of SAS cells to proliferate and migrate, RNA sequencing data obtained after starvation were further referred to as cell growth, cell death, cell migration,

cell proliferation, cell cycle, and cell adhesion. Analyzed using 6 keywords (Table 2). After 24 hours of serum starvation, more than two-fold up-regulation was observed for 425 genes. The top 5 genes were determined for each keyword. This included HSPA1A, OSGIN1, UCN, BST2, and SGK3 for cell growth, LOC728739, UCN, NPAS2, AGR2, and PYCR1 for cell death, BST2, ADGRA2, CALR, SGK3, and HSPA5 for cell migration, SGK3, IKZF3, SPTA1, MIR17HG, and CD22 for cell proliferation, ERN1, DDIT3, BEX2, CALR, and PIWIL4 for cell cycle, and AMIGO1, TNXB, TNC, FOXA2, and CD22 for cell adhesion (Table 2).

After 24 hours of serum starvation, over 50% down-regulation was observed for 733 genes. The top 5 genes with the largest reductions were: EDN1, PSRC1, S100A9, S100A8, and CDKN2C for cell growth, GPR37L1, AKR1C3, CTSV, BNIP3, and AXIN2 for cell death, STC1, ANLN, TCAF2, ATOH8, and SERPINB3 for cell migration, ID2, ATOH8, SERPINB3, EGLN3, and AXIN2 for cell proliferation, CENPE, KCTD11, MAP2K6, ID2, and CDKN2C for cell cycle, and CXCL8, SERPINI1, CDH2, CNTN1, and CCL2 for cell adhesion (Table 2).

3.4. Expression of serum starvation-induced genes in TCGA-HNSCC patients

From the 70 genes altered in HNSCC cells by 24-hour serum deficiency, the top two genes showing significant expression changes were selected for each of the 7 keywords, including autophagy, cell proliferation, cell death, cell migration, cell proliferation, cell cycle, and cell adhesion. Of the 28 genes selected, 6 were associated with replication. Moreover, the microRNA MIR17HG was excluded. Therefore, we finally focused on 21 genes. Of these, 11 were up-regulated genes and 10 were down-regulated genes. When the expression of these SIGs was examined in TCGA-HNSCC patients, 9 of the 11 up-regulated genes were also up-regulated in the primary tumor compared to solid normal tissue. Significant expression differences were observed in BST2, CALR, DDIT3, HSPA5, and TRIB3 (Fig. 2A). On the other hand, 6 out of the 10 down-regulated genes had reduced expression in tumors compared to solid normal tissue, with significant differences observed in the ATOH8 and CCL2 genes (Fig. 2B).

3.5. Function and PPI analysis of SIGs

GO and KEGG enrichment pathway analyses were performed to investigate the biological properties and potential signaling pathways of the 21 selected genes. Using GO enrichment analysis, enriched terms were ATF6-mediated unfolded protein response, PERK-mediated unfolded protein response, negative regulation of sequence-specific DNA-binding transcription factor activity, and negative regulation of transcription. These GO terms were associated with several important biological processes including DNA-templated gene expression response to endoplasmic reticulum stress, endoplasmic reticulum stress response, and positive regulation of cell cycle arrest (Fig. 3A). KEGG analysis showed that the prognostic genes were significantly enriched in pathways of transcriptional misregulation in cancer and protein processing in the endoplasmic reticulum (Fig. 3B). In PPI network analysis, CALR, HSPA5, DDIT3, and TRIB3 formed a close interaction network among the up-regulated genes (Fig. 3C). The association with AGR2, FOXA2, CD22, and SGK3 was weak, and PIWIL4 and PYCR1 were not associated with other up-regulated genes.

3.6. Prognostic significance of SIGs in TCGA-HNSCC patients

TCGA-HNSCC patients were divided into two groups based on the expression of SIGs. The expression levels of patients in the high expression group were higher than the median, and the remaining patients were classified in the low expression group ^{43,44}. The difference in survival time determined by the Kaplan-Meier method was examined using the generalized Wilcoxon test and the long rank test (Fig. 4). Among the up-regulated SIGs, high expression of CALR (Fig. 4C), FOXO4 (Fig. 4F), HSPA5 (Fig. 4G), and TRIB3 (Fig. 4K) was correlated with decreased patient survival. FOXA2 was excluded in subsequent studies because it was not significantly up-regulated in tumors compared to normal solid tissue. Conversely, high expression at AGR2 (Fig. 4A) and PIWK4 (Fig. 4H) were correlated with significant improvement in overall patient survival. On the other hand, high expression of BST2 (Fig. 4B), CD22 (Fig. 4D), DDIT3 (Fig. 4E), PYCR1 (Fig. 4I), and SGK3 (Fig. 4J) was not associated with patient survival. The Kaplan-Meier method was also applied to down-regulated genes, but there was no association between gene expression and patient survival in TCGA-HNSCC patients (Fig. S2).

3.7. Cox regression analysis of the association of SIGs and classical prognostic factors with survival in the TCGA-HNSCC patients

Expression of CALR, HSPA5, and TRIB3 was correlated with reduced overall survival in patients with TCGA-HNSCC, so these genes were further analyzed. Univariate and multivariate analysis (Cox proportional hazard model) was performed using the three genes and classical risk factors, such as gender, HPV, smoking, age, and TNM stage, as independent variables. In univariate analysis, CALR-High (vs. Low) (HR = 1.416, 95% CI = 1.069–1.875, $p=0.015$), HSPA5-High (vs. Low) (HR = 1.362, 95% CI = 1.029–1.804, $p = 0.031$), TRIB3-High (vs Low) (HR = 1.361, 95% CI = 1.028–1.803, $p = 0.031$), age (HR = 1.016, 95% CI = 1.003–1.029, $p = 0.015$), sex (HR = 0.721, 95% CI = 0.535–0.973, $p = 0.032$), M stage (HR = 4.748, 95% CI = 1.749–12.889, $p = 0.002$), and N stage (HR = 1.078, 95% CI = 1.002–1.160, $p = 0.045$) were significantly correlated with the prognosis of TCGA-HNSCC patients (Table 3). Multivariate analysis showed that the combination of two genes (CALR-High and HSPA5-High) ($P = 0.022$) and three genes ($P = 0.027$) did not make a clear difference in correlation.

4. Discussion

Autophagy has been suggested to be a biological marker for estimating the prognosis of cancer patients. In a previous HNSCC bioinformatics study, Li et al. ²⁹ identified a novel autophagy-related signature consisting of three hub genes, MAP1LC3B, FADD, and LAMP1, that may provide promising biomarker genes for the treatment and prognosis of HNSCC. Similarly, Jin et al. ³³ determined 35 genes for HNSCC and identified ITGA3, CDKN2A, FADD, NKX2-3, BAK1, CXCR4, and HSPB8 as prognostic ARGs. Ren et al. ³⁴ also reported 13 ARGs as genes that predict prognosis. In the present study, HNSCC cells were cultured under serum starvation, which can efficiently induce autophagy, and RNA sequencing was used to examine the expression of ARGs.

FBS is commonly used as a supplement to animal cell culture medium ⁴⁵. Additionally, FBS consists of several compositions such as macromolecules, carrier proteins for lipid substances and trace elements, attachment and spreading factors, low molecular weight nutrients, hormones, and growth factors ⁴⁵. Among them, growth factors were reported to influence cell proliferation, migration, survival, and morphogenesis ⁴⁶. Under serum starvation, SAS cells, a high-risk HPV-negative HNSCC cell line ⁴⁷, showed no significant changes in cell morphology after 24 hours, but cell growth and migration capacity was suppressed. Electron micrographs revealed the presence of autophagosomes and mitochondrial phagocytosis, suggesting that autophagy was induced in this situation. After 24-hour starvation, mRNA sequencing of SAS cells detected 12 up-regulated ARGs (ATP6V0A2, ATP6V1B1, ATP6V1C2, DDIT3, ERN1, NHLRC1, NUPR1, PIM2, TMEM150A, TRIB3, WIPI1, and XBP1) and 13 down-regulated ARGs (BNIP3, BNIP3L, C10orf10, DAPK2, GAPDH, HMOX1, MEFV, PLK2, RRAGD, SESN3, SRPX, S100A8, and S100A9). These genes differed from the ARGs previously reported to predict the prognosis of HNSCC patients ^{29,33,34}. This starvation-induced approach may be beneficial in extrapolating ARGs that have not been previously identified as differentially expressed genes. In addition, as with ARGs, we also found aberrant expression of genes related to cell growth, cell death, cell migration, cell proliferation, cell cycle, and cell migration. Finally, 21 SIGs that showed significant up-regulation or down-regulation were selected. Comparing how these genes were expressed in normal and cancer tissues in TCGA-HNSCC patients, we found 11 genes that were more strongly expressed in cancer cells and 10 genes that were down-regulated in cancer tissues. Among them, BST2, CALR, DDIT3, HSPA5, and TRIB3 were significantly up-regulated in cancer tissues.

GO and KEGG analyses revealed the involvement of ATF6-mediated unfolded protein responses and PERK-mediated unfolded protein responses mainly in the nucleus, and the ability of SIGs to bind glycoproteins and ubiquitin protein ligases. In addition, networking between CALR, HSPA5, DDIT3, and TRIB3 was demonstrated by PPI analysis. Consistent with the PPI analysis results, when TCGA-HNSCC patients were divided into high-expression and low-expression groups, and then analyzed by the Kaplan-Meier method, CALR, FOXA2, HSPA5, and TRIB3 were found to be correlated with reduced survival. Cox regression analysis further confirmed that three SIGs (CALR, HSPA5, and TRIB3), sex, M-stage, and N-stage were associated with survival in HNSCC patients. This suggests that CALR, HSPA5, and TRIB3 are predictors of poor prognosis. Since the combination of two genes (CALR-High and HSPA5-High) and three genes did not make a clear difference in correlation (Table 3), patients will have a poor prognosis, especially when both CALR and HSPA5 are highly expressed.

Calreticulin, CALR, is a soluble multifunctional protein found in the endoplasmic reticulum (ER) lumen and is involved in calcium homeostasis, transcriptional regulation, immune response, and cellular function ^{48,49}. It is expressed at higher levels in many cancerous tissues than in normal tissues. High CALR expression is correlated with both advanced clinical stage and lymph node metastasis ⁵⁰⁻⁵². The translocation of CALR from the ER to the plasma membrane occurs during immunogenic cell death ⁵³ due to oncolytic virus infection of HNSCC cells, leading to enhanced tumor immunity ⁵⁴. The unfolding protein response (UPR) is a cellular stress response related with ER stress. One of the proteins involved in

this UPR is Heat shock 70 kDa protein 5/glucose-regulated protein (HSPA5/GRP78). HSPA5 is the master regulator of UPR and is associated with tumor progression, tumor size, and poor prognosis⁵⁵⁻⁵⁸. In situations where protein production is required for tumor growth, HSPA5 is overactivated to process a high flux of protein passing through the ER, maintaining ER homeostasis. Expression of HSPA5 is induced by glucose starvation^{59,60}. It has also been shown that HSPA5 is up-regulated during viral infection, translocated to the cell membrane, and recognized by SARS-CoV-2 spikes⁶¹.

Tribbles homologue 3, TRIB3, is a member of the mammalian pseudokinase tribble family and is involved in multiple biological processes including the cellular response to glucose deficiency stress and ER stress. Several studies have shown that TRIB3 is elevated in multiple cancer cell lines and primary tumors including colorectal cancer, breast cancer, and lung cancer. In renal cancer, TRIB3 is overexpressed compared to normal tissue and is associated with tumor progression and poor prognosis⁶²⁻⁶⁴. On the other hand, TRIB3 binds to AKT, inhibits its phosphorylation and prevents its activation⁶⁵. In the case of HNSCC, patients with high TRIB3 expression had a better prognosis, and overexpression of TRIB3 reduced AKT phosphorylation and inhibited AKT activation⁶⁶. All of these genes are associated with endoplasmic reticulum stress. Their up-regulation may be a promising biomarker for predicting the prognosis of HNSCC.

There are many past studies where in vitro events and RNA-sequencing data were linked to informatics analysis of HNSCC patients^{67,68}. You et al.⁶⁷ established radiation-resistant cells by repeated irradiation in vitro and identified radioresistant genes using the TCGA-HNSCC database. In the present study, by analyzing genes induced by serum starvation of HNSCC cells, we detected genes that could not be obtained by previous TCGA database analysis and show their usefulness in predicting the prognosis of HNSCC patients. This approach may help to understand the genetic response of cancer cells to ER stress under therapeutic processes such as radiation therapy and chemotherapy.

5. Conclusions

Up-regulated and down-regulated genes associated with serum starvation using HNSCC cells were identified. Expression of HSPA5, TRIB3, and CALR in SAS cells was up-regulated by in vitro serum starvation and up-regulated in TCGA-HNSCC tissue tumors. High expression of these genes was closely associated with reduced survival in patients with TCGA-HNSCC. These SIGs have the potential to be molecular prognostic markers in HNSCC patients.

Abbreviations

SCC, squamous cell carcinoma; HNSCC, head and neck squamous cell carcinoma; TCGA, The Cancer Genome Atlas; CALR, calreticulin; HSPA5, heat shock protein family A (Hsp70) member 5; TRIB3, tribbles pseudokinase 3

Declarations

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Contributions

M.H. and H.I. designed the entire study under the supervision of Y.Y., M.M.N., and N.U. M.H., H.I., K.N., and S.Y. performed the experiments. Statistical analyses and data interpretation were conducted by M.H., H.I., Y.Y., M.M.N., and N.U. M.H., H.I., and Y.Y. wrote the manuscript. All authors reviewed the manuscript.

Conflicts of Interest: The authors declare no conflict of interest.

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Tables

Table 1. Expression of autophagy-related genes under serum starvation

ID	Description	NCBI Gene ID	Fold Change after 24 hours	Fold Change after 2 hours
WIPI1	WD repeat domain, phosphoinositide interacting 1	55062	2.0886614	0.930744261
XBP1	X-box binding protein 1	7494	2.1410127	0.878980954
NUPR1	nuclear protein 1, transcriptional regulator	26471	2.1501427	1.06686497
NHLRC1	NHL repeat containing E3 ubiquitin protein ligase 1	378884	2.1578841	1.399708125
ATP6V0A2	ATPase H+ transporting V0 subunit a2	23545	2.3026843	1.142090599
ATP6V1B1	ATPase H+ transporting V1 subunit B1	525	2.325067	0.828736747
PIM2	Pim-2 proto-oncogene, serine/threonine kinase	11040	2.378625	1.229401772
TMEM150A	transmembrane protein 150A	129303	2.6394634	1.155387955
ATP6V1C2	ATPase H+ transporting V1 subunit C2	245973	2.6718903	1.774302383
ERN1	endoplasmic reticulum to nucleus signaling 1	2081	2.9943194	1.146213323
DDIT3	DNA damage-inducible transcript 3	1649	3.1741931	0.597946814
TRIB3	tribbles pseudokinase 3	57761	3.8404264	0.991618344
HMOX1	heme oxygenase 1	3162	0.49524766	0.945472836
SRPX	sushi repeat containing protein, X-linked	8406	0.4878078	0.857148158
RRAGD	Ras-related GTP-binding D	58528	0.4833195	0.845809254
GAPDH	glyceraldehyde-3-phosphate dehydrogenase	2597	0.4825641	0.94147729
C10orf10	chromosome 10 open reading frame 10	11067	0.46908534	1.273417327
PLK2	polo-like kinase 2	10769	0.40572378	0.835709592
BNIP3L	BCL2-interacting protein 3-like	665	0.37105525	0.898710392
SESN3	sestrin 3	143686	0.3578121	0.715624111
DAPK2	death-associated protein kinase 2	23604	0.34043416	0.718694176
S100A9	S100 calcium binding protein A9	6280	0.26209667	1.042621814
MEFV	MEFV, pyrin innate immunity regulator	4210	0.25515518	0.765465346
S100A8	S100 calcium binding protein A8	6279	0.24599823	1.095810293
BNIP3	BCL2-interacting protein 3	664	0.12660438	0.819784126

Table 2. Genes that were selected by 6 keywords and showed high up-regulation or down-regulation under serum starvation

Key Word	ID	Description	NCBI Gene ID	Fold Change after 24 hours	Fold Change after 2 hours
Cell growth	HSPA1A	heat shock protein family A (Hsp70) member 1A	3303	2.1397336	1.248177638
	OSGIN1	oxidative stress-induced growth inhibitor 1	29948	2.1668558	1.538782151
	UCN	urocortin	7349	2.4790032	1.144155207
	BST2	bone marrow stromal cell antigen 2	684	2.9329562	1.308922836
	SGK3	serum/glucocorticoid regulated kinase family member 3	23678	3.820176	1.176317287
	EDN1	endothelin 1	1906	0.27418113	0.884857381
	PSRC1	proline and serine rich coiled-coil 1	84722	0.26818275	0.945457524
	S100A9	S100 calcium binding protein A9	6280	0.26209667	1.042621814
	S100A8	S100 calcium binding protein A8	6279	0.24599823	1.095810293
Cell Death	CDKN2C	cyclin-dependent kinase inhibitor 2C	1031	0.12124052	0.835693575
	LOC728739	programmed cell death 2 pseudogene	728739	2.3000002	1
	UCN	urocortin	7349	2.4790032	1.144155207
	NPAS2	neuronal PAS domain protein 2	4862	2.689144	0.930857474
	AGR2	anterior gradient 2, protein disulphide isomerase family member	10551	2.6983347	1.036070166
	PYCR1	pyrroline-5-carboxylate reductase 1	5831	2.81048	1.165619558
	GPR37L1	G protein-coupled receptor 37 like 1	9283	0.3573708	0.929164088
	AKR1C3	aldo-keto reductase family 1 member C3	8644	0.25129116	0.95490666
	CTSV	cathepsin V	1515	0.24350072	0.872422377
Cell migration	BNIP3	BCL2-interacting protein 3	664	0.12660438	0.819784126
	AXIN2	axin 2	8313	0.07149426	0.979471303
	BST2	bone marrow stromal cell antigen 2	684	2.9329562	1.308922836
	ADGRA2	adhesion G protein-coupled receptor A2	25960	3.3248994	1.09175771
	CALR	calreticulin	811	3.6909635	0.82266943
	SGK3	serum/glucocorticoid regulated kinase family member 3	23678	3.820176	1.176317287
	HSPA5	heat shock protein family A (Hsp70) member 5	3309	7.4991713	0.83579081
	STC1	stanniocalcin 1	6781	0.20706424	0.782881491
	ANLN	anillin actin binding protein	54443	0.18460144	0.933625648
Cell proliferation	TCAF2	TRPM8 channel-associated factor 2	285966	0.13127537	0.750317546
	ATOH8	atonal bHLH transcription factor 8	84913	0.11830313	0.709818819
	SERPINB3	serpin family B member 3	6317	0.11149136	1.039357318
	SGK3	serum/glucocorticoid regulated kinase family member 3	23678	3.820176	1.176317287
	IKZF3	IKAROS family zinc finger 3	22806	4.3976007	0.968962725
	SPTA1	spectrin alpha, erythrocytic 1	6708	4.4961443	1.10674307
Cell proliferation	MIR17HG	miR-17-92a-1 cluster host gene	407975	4.602493	1.283155842
	CD22	CD22 molecule	933	5.679524	1.044954641
	ID2	inhibitor of DNA binding 2	3398	0.17407766	1.119070975

	ATOH8	atonal bHLH transcription factor 8	84913	0.11830313	0.709818819
	SERPINB3	serpin family B member 3	6317	0.11149136	1.039357318
	EGLN3	egl-9 family hypoxia inducible factor 3	112399	0.100901835	0.74969137
	AXIN2	axin 2	8313	0.07149426	0.979471303
	ERN1	endoplasmic reticulum to nucleus signaling 1	2081	2.9943194	1.146213323
	DDIT3	DNA damage-inducible transcript 3	1649	3.1741931	0.597946814
	BEX2	brain expressed X-linked 2	84707	3.5948327	1.06210961
	CALR	calreticulin	811	3.6909635	0.82266943
Cell cycle	PIWIL4	piwi like RNA-mediated gene silencing 4	143689	3.979958	1.182165243
	CENPE	centromere protein E	1062	0.2240672	0.891558939
	KCTD11	potassium channel tetramerization domain containing 11	147040	0.20453872	0.83617986
	MAP2K6	mitogen-activated protein kinase kinase 6	5608	0.19298029	0.761199796
	ID2	inhibitor of DNA binding 2	3398	0.17407766	1.119070975
	CDKN2C	cyclin-dependent kinase inhibitor 2C	1031	0.12124052	0.835693575
	AMIGO1	adhesion molecule with Ig like domain 1	57463	2.892857	1.089285831
	TNXB	tenascin XB	7148	3.0829856	1.187939406
	TNC	tenascin C	3371	3.1931474	1.035432055
	FOXA2	forkhead box A2	3170	3.3726099	0.838221697
Cell adhesion	CD22	CD22 molecule	933	5.679524	1.044954641
	CXCL8	C-X-C motif chemokine ligand 8	3576	0.3170054	0.587415932
	SERPINI1	serpin family I member 1	5274	0.31145853	0.953841739
	CDH2	cadherin 2	1000	0.29494244	1.09390035
	CNTN1	contactin 1	1272	0.26283494	0.835311109
	CCL2	C-C motif chemokine ligand 2	6347	0.22852589	0.886680707

Table 3. Univariate and multivariate analyses of three genes (CALR, HSPA5, and TRIB3) using the TCGA-HNSC patient data

	Univariate					Multivariate				
	HR	95% CI		P-value		HR	95% CI		P-value	
CALR_High (vs. Low)	1.416	1.069	-	1.875	0.015	-				
HSPA5_High (vs. Low)	1.362	1.029	-	1.804	0.031	-				
TRIB3_High (vs. Low)	1.361	1.028	-	1.803	0.031	-				
Gene (CALR-HSPA5-TRIB3)										
Low-Low-Low	1.000	ref				1.000	ref			
High-Low-Low	1.208	0.607	-	2.405	0.590	1.455	0.696	-	3.043	0.319
Low-High-Low	1.326	0.805	-	2.186	0.268	1.398	0.833	-	2.348	0.205
Low-Low-High	0.733	0.290	-	1.850	0.510	0.691	0.245	-	1.949	0.485
High-High-Low	1.783	1.118	-	2.845	0.015	1.777	1.086	-	2.907	0.022
High-Low-High	1.901	1.002	-	3.608	0.049	1.805	0.934	-	3.488	0.079
Low-High-High	1.861	1.061	-	3.265	0.030	1.887	1.021	-	3.486	0.043
High-High-High	1.619	1.088	-	2.410	0.018	1.614	1.056	-	2.467	0.027
Age (per 1)	1.016	1.003	-	1.029	0.015	1.023	1.009	-	1.038	0.002
Sex_male (vs. female)	0.721	0.535	-	0.973	0.032	0.862	0.611	-	1.215	0.396
HPV status_Positive (vs. Negative)	0.777	0.524	-	1.153	0.210	0.784	0.505	-	1.218	0.279
Alcohol_history_Yes (vs. No)	0.947	0.701	-	1.279	0.723	1.051	0.750	-	1.472	0.774
Cigarettes per day_>0 (vs. 0)	0.960	0.724	-	1.272	0.774	0.963	0.709	-	1.309	0.811
M stage_m1 (vs. m0)	4.748	1.749	-	12.889	0.002	4.979	1.699	-	14.592	0.003
N stage (Continuous variable per 1)	1.078	1.002	-	1.160	0.045	1.112	1.023	-	1.208	0.013
N stage (Category)						-				
n0	1.000	ref								
n1	1.061	0.713	-	1.581	0.769					
n2	1.679	0.898	-	3.138	0.105					
n2a	1.607	0.744	-	3.469	0.227					
n2b	0.977	0.632	-	1.510	0.916					
n2c	1.980	1.229	-	3.190	0.005					
n3	1.698	0.623	-	4.624	0.300					
T stage (Continuous variable per 1)	0.999	0.902	-	1.106	0.986	0.979	0.877	-	1.092	0.700
T stage (Category)						-				
t1	1.000	ref								
t2	1.092	0.570	-	2.092	0.790					
t3	1.421	0.746	-	2.705	0.285					
t4	1.543	0.681	-	3.496	0.299					
t4a	1.056	0.554	-	2.015	0.868					
t4b	1.984	0.439	-	8.960	0.373					

HR: hazard ratio; 95% CI: 95% confidence interval; ref: reference value.

Figures

Figure 1

Effects of serum starvation on the biological activity of head and neck cancer SAS cells (A) Growth of SAS cells as measured by MTT assay after treatment for the indicated time, with or without serum. The data are shown as mean $\pm$ SD of 3 independent experiments and analyzed by t-test. (B and C) Migration assay in cells treated with/without serum. The bars show the area of scratches at 0 hours. The rates of wound closure were determined from the assays and indicate the mean $\pm$ SD of 3 independent experiments. (D) TEM examination of SAS incubated for 24 hours with/without serum. The bar markers represent 2 μ m. M: mitochondria, AP: autophagosome, AL: autolysosome.

Figure 2

Expression of serum starvation-induced genes (SIGs) in TCGA-HNSCC patients The expression of 21 genes that showed significant changes in expression by serum starvation and their relative expression levels in TCGA-HNSCC patients were determined in primary tumors and solid normal tissues. (A) 11 genes (AGR2, BST2, CALR, CD22, DDIT3, FOXA2, HSPA5, PIWIL4, PYCR1, SGK3, and TRIB3) have been up-regulated more than two-fold. (B) Ten genes (ATOH8, AXIN2, BNIP3, CCL2, CDKN2C, CNTN1, EGLN3, ID2, SERPINB3, and S100A8) were down-regulated by more than 50%.

Figure 3

Function and protein-protein interaction analysis of SIGs (A) A list of the top 8 significant GO terms determined by GO enrichment analysis of 21 selected genes. (B) List of molecular pathways determined by KEGG pathway enrichment analysis of 21 selected genes. (C) Proteins encoded by 21 genes extracted using 7 keywords were subjected to PPI network analysis. BP, biological process; CC, cellular composition; MF, molecular function

Figure 4

Prognostic significance of SIGs in TCGA-HNSCC patients Overall survival of TCGA-HNSCC patients, classified by 11 up-regulated SIG expression levels, was determined by the Kaplan-Meier method. The difference in survival time determined by the Kaplan-Meier method was examined using the generalized Wilcoxon test and the long rank test. A: AGR2, B: BST2, C: CALR, D: CD22, E: DDIT3, F: FOXA2, G: HSPA5, H: PIWIL4, I: PYCR1, J: SGK3, K: TRIB3.

Supplementary Files

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