Lipid metabolic transcriptome-wide profile and signature of lung adenocarcinoma

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Research

Keywords: lung adenocarcinoma, lipid metabolism, diagnosis, prognosis

Posted Date: July 15th, 2020

DOI: https://doi.org/10.21203/rs.3.rs-31951/v2

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Version of Record: A version of this preprint was published on October 13th, 2020. See the published version at https://doi.org/10.1186/s12944-020-01390-9.
Abstract

Background: Lung cancer is the cancer with high morbidity and mortality across the globe, and lung adenocarcinoma (LUAD) is the most common histologic subtype. The disorder of lipid metabolism is related to the development of cancer. Analysis of lipid-related transcriptome helps shed light on diagnosis and prognostic biomarkers of LUAD.

Methods: In this study, we performed an expression analysis of 1045 lipid metabolism-related genes between LUAD tumors and normal tissues from the TCGA-LUAD cohort. The interaction network of differential expression genes (DEGs) was constructed to identify. The association between hub genes and overall survival (OS) was evaluated and formed a model to predict the prognosis of LUAD using a nomogram, and the model was validated by another cohort (GSE13213).

Results: Finally, a total of 217 lipid metabolism-related DEGs were detected in LUAD. They were significantly enriched in glycerophospholipid and steroid metabolism. Then we identified 6 hub genes through network and cytoHubba, including INS, LPL, HPGDS, DGAT1, UGT1A6, and CYP2C9. The high expression of CYP2C9, UGT1A6, and INS, whereas low expressions of DGAT1, HPGDS, and LPL, were associated with worse OS for 719 LUAD patients. Our model found that the high-risk score group had a worse OS, and the validated cohort had the same result.

Conclusion: This study constructed a signature of six lipid metabolic genes, which was significantly associated with the diagnosis and prognosis of LUAD patients. The gene signature can be used as a biomarker for LUAD in the term of lipid metabolic.

Background

Lung cancer is the most commonly diagnosed cancer (11.6% of the total cases) and the leading cause of cancer death (18.4% of the total cancer deaths) in the world [1]. Among the subtype of lung cancers, adenocarcinoma is the most common histologic subtype of lung cancer in men and women[2]. A 2005-2014 epidemiological survey from China showed that the proportion of adenocarcinoma increased from 36.4% to 53.5%, while the proportion of squamous carcinoma decreased from 45.4% to 34.4%[3]. The increasing incidence of lung adenocarcinoma (LUAD) has also been reported to be associated with air pollution-related factors[4-6]. Researches reported PM2.5 increases pro-inflammatory lipid metabolism in the lung and was associated with lipid alterations[7, 8]. The importance of alterations related to lipid metabolism is starting to be recognized, and the increase in de novo lipogenesis is considered a new hallmark in many aggressive cancers[9]. Lipid profiles of blood plasma exosomes could be used for early detection of the prevalent non-small cell lung cancers (NSCLC)[10]. Epidemiological data indicated that a certain number of lung cancer patients with high high-density lipoprotein cholesterol (HDL-C) and low-density lipoprotein (LDL) and low-density lipoprotein receptor (LDLR) level has better survival in patients[11, 12]. Compared with healthy subjects, NSCLC patients showed significant increases in phosphatidylcholine (PCs) and phosphatidylethanolamine (PEs)[13]. Other lipid metabolism indicators
associated with LUAD includes sphingomyelins, phosphatidylinositolns, phosphatidylserines, phosphatidylethanolamine, phospholipids, and phosphatidylcholine[14]. The cancer cells’ requirement of metabolic intermediates for macromolecule production is overwhelming. Fatty acid oxidation(FAO) can help to generate ATP to support the membranes formation, energy storage, production of signaling molecules by coordinating the activation of lipid anabolic metabolism [15]. The regulation of lipid metabolic to LUAD is still being explored. Knowing the lipid-related mechanism of the LUAD phenotype will inform better clinical interventions.

To explore the further lipid mentalism relating to regulation network and pathway, we used an integrated bioinformatic method to construct the transcriptome-wide profile; and a signature of lipid-related genes was analyzed to explore the potential biomarkers for diagnosis and prognostic of LUAD in the term of lipid metabolism disorder.

**Materials And Methods**

**Patients and datasets**

**Identification of lipid metabolism-related differentially expressed genes**
After using lipid-specific keywords (fatty acyls, glycerolipids, glycerophospholipids, sphingolipids, sterol lipids, prenol lipids, saccharolipids, and polyketides), 21 lipid metabolism-related pathways and five lipid metabolism-related gene sets were collected from the Kyoto Encyclopedia of Genes and Genomes (KEGG) web site (http://www.kegg.jp/blastkoala/) [18] and the Molecular Signatures Database (MisDB) web site (https://www.gsea-msigdb.org/gsea/msigdb/index.jsp) [19], respectively (Additional file 1). After removing the overlapped genes, a total of 1045 lipid metabolism-related genes were obtained. Lipid metabolism-related differentially expressed genes (DEGs) between LUAD tissues and normal tissues were screened through R package edgeR [20]. The parameters set for differential expression analysis were FDR<0.05 and |log2 fold change| (logFC)>1.

**Bioinformatic analysis**
We used the R package clusterProfiler to furtherly explore the biological significance of lipid metabolism-related DEGs [21]. In GO and KEGG analysis, FDR < 0.05 was considered a significant enrichment. Then we uploaded the DEGs that containing gene identifiers and corresponding FDR values and log_{2}FC values into the IPA software (Qiagen). The “core analysis” function included in the software was used to interpret the DEGs.

**Interaction network generation and hub genes analysis**
We built an interaction network of differentially expressed lipid metabolism-related genes using the Search Tool for the Retrieval of Interacting Genes (STRING, http://string-db.org/) database [22]. The combined score of \( \geq 0.4 \) was the cut-off value. Cytoscape software (version 3.6.0) was used to visualize networks [23]. According to 12 ranking methods in cytoHubba [24], an APP in Cytoscape, the top ten genes of each method were selected for overlap analysis, and the genes with the highest number of overlaps were used as hub genes and the potential biomarkers.

**Survival analysis**

The overall survival (OS) analysis of hub genes was shown by Kaplan-Meier Plotter (http://kmplot.com/analysis/), which includes clinical data and gene expression information for 719 lung cancer LUAD patients from GEO and TCGA database[25]. Then, the information on the number of cases along with median values of mRNA expression levels, hazard ratios (HR) with 95% confidence intervals (CI), and log-rank P-values were extracted from the KM plotter webpage. Log-rank P-values < 0.05 were considered statistically significant.

**Prediction model**

Based on the selected hub genes, we use the nomogram package of R (“rms”) [26] to develop a model to evaluate the prognosis of TCGA-LUAD patients. Using the formula of the nomogram, we calculated the prognosis score of each patient. According to the score, patients were divided into a low-risk score group and a high-risk score group using the median classification method. The prognosis score was validated by the patients’ actual prognosis outcome. Then we downloaded hub genes expression data and clinical information of 117 LUAD patients from another data set (GSE13213), and calculated the prognosis score of each patient by the formula of the nomogram. Then they also were divided into two groups using the median classification method to perform the survival analysis to validate the availability of this model.

**Results**

**Identification and functional analysis of lipid metabolism-related DEGs**

A total of 217 lipid metabolism-related DEGs were identified from the TCGA-LUAD cohort. A volcano plot was constructed to reveal the significant DEGs (Fig. 1A), and a heatmap was created to show the hierarchical clustering analysis of the DEGs (Fig. 1B). To get an overall understanding of 217 lipid metabolism-related DEGs, we conducted GO terms and KEGG pathway enrichment using clusterProfiler package, while canonical pathways analysis by IPA. The results of KEGG pathway enrichment showed that DEGs were significantly enriched in arachidonic acid metabolism, metabolism of xenobiotics by cytochrome P450, glycerophospholipid metabolism, and steroid hormone biosynthesis. In contrast, they were significantly enriched in fatty acid metabolic process, glycerolipid metabolic process, fatty acid derivative metabolic process, and steroid metabolic process from GO terms (Fig. 1C). The genes in each KEGG pathway and GO term were shown in the additional file 2. IPA identified significant canonical networks associated with the DEGs. IPA showed that the top canonical pathways associated with common DEGs were eicosanoid signaling, FXR/RXR activation, and atherosclerosis signaling (Fig. 1D). Combining the results of three functional analyses showed that, the DEGs mainly overlapped in glycerophospholipid and steroid metabolism. And the non-overlapping pathways showed more
information indicating further exploration of the role of lipid metabolism in LUAD.

**Interaction network construction and cytoHubba analysis**

Lipid metabolism-related DEGs were analyzed by the STRING tool. Ultimately, an interaction network with 216 nodes and 1140 edges was established and visualized in Cytoscape (Fig 2). Then a total of 6 hub genes were identified by the overlap of the top 10 genes according to 12 ranked methods in cytoHubba (Additional file 3). Moreover, these genes were related to Insulin (INS), Lipoprotein Lipase (LPL), Hematopoietic Prostaglandin D Synthase (HPGDS), Diacylglycerol O-Acyltransferase 1 (DGAT1), UDP Glucuronosyltransferase Family 1 Member A6 (UGT1A6), and Cytochrome P450 Family 2 Subfamily C Member 9 (CYP2C9).

**Survival analysis of hub genes**

We examined the relationship between mRNA expression of hub genes and clinical outcome using the Kaplan-Meier plotter. Results showed that high expression of CYP2C9 [HR = 1.50 (1.19–1.90), \( P < 0.01 \)], UGT1A6 [HR = 1.61 (1.26–2.06), \( P < 0.01 \)], and INS [HR = 1.46 (1.15–1.85), \( P < 0.01 \)], whereas low expression of DGAT1 [HR = 0.78 (0.62–0.98), \( P = 0.04 \)], HPGDS [HR = 0.58 (0.45–0.73), \( P < 0.01 \)], and LPL [HR = 0.54 (0.43–0.69), \( P < 0.01 \)], were associated with worse OS for 719 LUAD patients (Fig. 3).

**Prediction model based on survival-related hub genes and validation**

Based on the Cox regression model, a nomogram was built to predict the prognosis of TCGA-LUAD patients, using the mRNA expression of the six survival-related hub genes (Fig. 4A). The concordance index of the nomogram was 0.61. Then we calculated the prognosis score of each patient, and found that the patients in the high-risk score group had worse OS of 3 years [HR = 1.51 (1.07–2.13), \( P = 0.02 \)] (Fig. 4B). We validated the model and found that high-risk score group had worse OS [HR = 1.84 (1.00–3.37), \( P = 0.05 \)] (Fig. 4C).

**Discussion**

Metabolic change has been widely observed in cancer cells[27]. Among those metabolisms, lipid metabolism widely participates in the regulation of many cellular processes such as cell growth, proliferation, differentiation, survival, apoptosis, inflammation, motility, membrane homeostasis, chemotherapy response, and drug resistance[28]. Some recent studies have reported some component of PM2.5 which has been reported as the risk factors of lung cancer[29-31], for the component of PM2.5 promotes pulmonary injury by modifying lipid metabolism[7] and might develop to lung cancer. However, there are fewer researches regarding the association between lipid metabolism and lung cancer in the term of transcriptome-wide analysis. This study used a LUAD cohort to generate the transcriptome-wide profile of lipid-related that includes 217 genes. The enrichment biological pathway found in LUAD included fatty acid, glycerolipid, and glycerophospholipids were the primary driven enrichment biological function that has been reported[32]. Besides, arachidonic acid metabolism, PPAR signaling pathway, insulin resistance, eicosanoids signaling, and other pathways were also reported in cancer[33-37].

The results indicate that LUAD-related lipid metabolism was associated with nicotine, estrogen biosynthesis, melatonin, and atherosclerosis. Similar to PM2.5, nicotine may promote LUAD development
regulated by lipid disordered. The interaction between estrogen biosynthesis and lipid metabolic may be one of the high-risk factors for LUAD, which is consistent with the observation that LUAD incidence is rising in women, and the incidence rate among female was higher than that among men [38]; Lipid and cancer-related genes were enriched in atherosclerosis and cancer. For long-term survival LUAD patients, their health management should be involved by oncologists and cardiologists [39].

We constructed the network of those genes that are related to lipid and LUAD and find six hub genes. CYP2C9, which is a drug target of lung cancer, can be slowed by cytochrome P450; and the tumorigenesis was regulated[44, 45]. LUAD patients with a lower expression of CYP2C9 have a better prognosis. UGT1A variants may play only a minor role in other lung cancer risk[46]. LUAD patients with a lower expression of UGT1A6 have a better prognosis. DGAT1 catalyzes the final step in triglyceride synthesis [47]. LPL is a key lipoytic enzyme that plays a crucial role in the catabolism of triglycerides in TG-rich particles[48]. Both of them are involved in triglyceride synthesis. And triglyceride was reported with HPGDS has the therapeutic potential in allergic inflammation[49]. Serum triglyceride concentrations were reported to be involved in the pathogenesis of lung cancer[50]. Those three genes were positively related to survival time. INS encodes insulin and plays a vital role in the regulation of carbohydrate and lipid metabolism. LUAD patients with a lower expression of INS have a better prognosis. The regulation of triglyceride synthesis, insulin, and inflammation control may be the effective intervention of LUAD patients. Based on those six genes, a risk model was constructed. LUAD patients from two cohorts with a lower risk score had a better prognosis.

**Strengths and limitations**

The main strength of the study is the establishment of a lipid metabolic transcriptome-wide profile of LUAD and a gene signature that significantly associated with the diagnosis and prognosis of LUAD patients in the term of lipid metabolism. Limitations include: 1) the data field information of these two cohorts is limited, which leading the covariables related to LUAD missed and brought bias; 2) the further internal mechanism of these six lipid-related genes cannot be illuminated in this study. A well-designed experiment based on our results was required in further research.

**Conclusions**

In summary, we generated a lipid metabolic transcriptome-wide profile of LUAD patients and found that significant lipid metabolic pathways were correlated with the LUAD. A signature of six lipid metabolic genes was significantly associated with the diagnosis and prognosis of LUAD patients. The gene signature can be used as a biomarker for LUAD, and the guidance to prevent the occurrence of LUAD and improve the prognosis of LUAD patients.

**Abbreviations**
Declarations

Ethics approval and consent to participate
Not applicable.

Consent for publication
Not applicable.

Availability of data and materials
The datasets generated and/or analyzed during the current study are available in the TCGA & GEO databases, [https://cancergenome.nih.gov/] & [https://www.ncbi.nlm.nih.gov/geo/].

Competing interests
The authors declare that they have no competing interests.

Funding
This study is supported by Youth Project of Wuxi Municipal Health and Family Planning Commission (Q201803), Youth Project of Jiangnan University Public Health Research Center (JUPH201826), Startup Fund for Youngman Research at SJTU (17X100040015) and Shanghai Jiao Tong University Medical and Industrial Cross Project (YG2017QN70)

Authors’ contributions
JYL and QL analyzed the data and helped draft the manuscript. FZ and HTM supervised this work and edited and revised the manuscript. ZYS, QS, YZ, TNF, and JYJ prepared figures and contributed to the drafting of the manuscript. The authors read and approved the final manuscript.

Acknowledgments
Not applicable.

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Figures
Figure 1

Identification and functional analysis of lipid metabolism-related DEGs. (A) Volcano plot of lipid metabolism-related genes, (B) Heatmap analysis of lipid metabolism-related DEGs, (C) GO and KEGG pathway enrichment analysis by clusterProfiler, (D) functional and signaling pathway enrichment by IPA. In (A) and (B), red, white, and blue represent higher expression levels, no expression differences, and lower expression levels, respectively.
Figure 2

The PPI network of lipid metabolism-related DEGs. Red, white, and blue nodes represent upregulated genes, no expression differences genes, and downregulated genes, respectively. The magnitude of the degree is positively correlated with the size of a node.
Figure 3

Survival analysis of hub genes. LUAD patients were subdivided into high/low gene expression groups based on the median expression level of each gene in LUAD tissues. (A) OS analysis of CYP2C9, (B) OS analysis of UGT1A6, (C) OS analysis of INS, (D) OS analysis of DGAT1, (E) OS analysis of HPGDS, and (F) OS analysis of LPL.
Figure 4

Prediction model based on survival-related hub genes and validation. (A) The nomogram of 6 survival-related genes, (B) survival analysis between high-risk score group and low-risk score group, and (C) validation of the model.

Supplementary Files
This is a list of supplementary files associated with this preprint. Click to download.

- Additionalfile1.xlsx
- Additionalfile3.docx
- Additionalfile2.csv
- Additionalfile5.pdf
- Additionalfile4.pdf