The O-glycosylating enzyme GALNT2 acts as an oncogenic driver in non-small cell lung cancer

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Abstract

Background: N-Acetylgalactosaminyltransferases (GALNTs), the enzymes that initiate mucin-type O-glycosylation, are closely associated with tumor occurrence and progression. However, a comprehensive analysis of GALNTs in non-small cell lung cancer (NSCLC) is lacking.

Methods: The expression profiles and prognostic values of the GALNT family members in NSCLC were analyzed using publicly available databases. Gain- and loss-of-function experiments were applied to assess the biological function of GALNT2 in NSCLC. High-throughput sequencing and bioinformatics approaches were employed to uncover the regulatory mechanism of GALNT2.

Results: Among the family members of GALNTs, only GALNT2 was frequently overexpressed in NSCLC tissues and was positively correlated with poor prognosis. In vitro assays showed that GALNT2 knockdown repressed NSCLC cell proliferation, migration, and invasion, but induced apoptosis and cell cycle arrest. Correspondently, GALNT2 overexpression exerted the opposite effects. In vivo experiments demonstrated that knockdown of GALNT2 restrained tumor formation in nude mice. Mechanistic investigations revealed that GALNT2 modified the O-glycosylation of ITGA5 and affected the activation of the PI3K/Akt and MAPK/ERK pathways. Further studies showed that miR-30d was a negative regulator of GALNT2.

Conclusions: These findings suggest that GALNT2 is an oncogene in NSCLC and has the potential as a target for NSCLC therapy.

Keywords: Non-small cell lung cancer, Oncogene, Glycosyltransferase, GALNT2
patients usually die from tumor recurrence or metastasis. Therefore, it is of great importance to investigate the mechanism underlying NSCLC oncogenesis, which may help develop new therapeutic/preventive strategies against NSCLC.

Glycosylation is a reaction that links glycans to lipids or proteins [2]. Changes in protein glycosylation can drive the neoplastic transformation of healthy cells [3]. Aberrant glycosylation not only mediates cell–cell and cell–matrix interactions but also impacts cell growth, survival, migration, and invasion. Accumulating evidence has indicated that glycosylation alteration is a hallmark of many cancers [4–6]. Glycosylation is catalyzed by glycosyltransferases (GTs), which consist of at least 200 members. GTs are frequently dysregulated in cancers, indicating that GTs may act as oncogenes or tumor-suppressor genes [7, 8]. Thus, uncovering the relationship between GTs and NSCLC is very significant.

N-Acetylgalactosaminyltransferases (GALNTs) are highly eukaryotic-retaining GTs that control the initial steps of mucin-type O-glycosylation [9]. To date, 20 GALNT family members (from GALNT1 to GALNT20) have been identified in the human genome. GALNTs play critical roles in numerous physiological and pathological processes [10–12]. However, a comprehensive analysis of GALNTs in NSCLC has yet to be conducted. In this study, multiple online tools were utilized to evaluate the expression profiles and prognostic values of the GALNT family in NSCLC. Subsequently, in vitro and in vivo assays were carried out to clarify the biological function and regulatory mechanism of GALNT2. Our study is expected to provide a novel therapeutic target for NSCLC.

Methods
Data retrieving and analyzing
Differential expression and prognosis of the GALNT family members in NSCLC were analyzed using the following databases: Oncomine (http://www.oncomine.org), The Cancer Genome Atlas (TCGA, https://tcga-data.nci.nih.gov/tcga/), UALCAN (http://ualcan.path.uab.edu), Genotype-Tissue Expression (GTEx, http://commonfund.nih.gov/GTEx/), and Kaplan Meier-plotter (http://kmplot.com). The LinkedOmics database (http://www.linkedomics.org) was used to screen the genes or miRNAs associated with GALNT2 in NSCLC. The prognostic nomograms and Receiver-operating characteristic (ROC) curves were generated using the R package “rms” package or “pROC”, respectively. Gene Set Enrichment Analysis (GSEA) was performed using the GSEA software. The adjusted P-value < 0.05 and |log2 (fold change) | > 1 were set as the threshold.

Microarray analysis
The cDNA microarrays of tissues (cDNA-HLugA030PG01; fifteen NSCLC samples and paired adjacent noncancerous lung tissue samples) and cells (MecDNA-HLugC042Ce01; thirteen NSCLC cell lines and one normal lung cell line) were acquired from Outdo Biotech Co., Ltd (Shanghai, China). Gene or miRNA expression analysis was performed by reverse transcription-quantitative real-time PCR (RT-qPCR).

Cell culture, plasmids, and transfection
A549 and H1299 cell lines (Procell, Wuhan, China) were maintained in DMEM (Gibco, Carlsbad, CA, USA), supplemented with 10% fetal bovine serum (HyClone, Logan, UT,
USA). GenePharma (Shanghai, China) was responsible for the synthesis of GALNT2 overexpression lentivirus (OV), GALNT2 shRNA lentivirus, and corresponding control lentiviruses (Mock or shNC). ITGA5 siRNA, negative control siRNA (siNC), miR-30d mimics, and negative control mimics were also chemically synthesized by GenePharma. Transfections were carried out using RNAi-mate reagent (GenePharma). Sequence information is provided in Additional file 1: Table S1.

RT-qPCR
Total RNA was extracted using an RNA extraction kit (Tiangen Biotech, Beijing, China). For mRNA detection, the SuperScript IV reverse transcriptase system (Thermo Fisher Scientific, Carlsbad, CA, USA) was used for cDNA synthesis. PCR reactions were performed using the SYBR® Premix Ex Taq™ (Takara, Dalian, China). For miRNA detection, RNA was reverse transcribed into cDNA by using the miRNA First-Strand cDNA synthesis kit (Tiangen). PCR was conducted using the miRNA qPCR detection kit (Tiangen). Data were processed with the 2^−ΔΔCt method after normalizing to GAPDH or U6. Primer sequences are given in Additional file 1: Table S2.

Western blot
Total protein was isolated using RIPA buffer and separated by 10% SDS-PAGE. After transferring to PVDF membranes, the blot was incubated with specific antibodies. Primary antibodies were as follows: GALNT2 (abs117502; Absin, Shanghai, China), ITGA5 (abs101364; Absin), ERK1 + ERK2 (ab184699; Abcam, Shanghai, China), p-ERK1 (T202) + p-ERK2 (T185) (ab201015; Abcam), Akt (ab8805; Abcam), p-Akt (T308) (ab38449; Abcam), and GAPDH (Absin; abs132004). Immunoreactive bands were visualized using an enhanced chemiluminescence kit (Absin).

Cell Counting Kit-8 (CCK-8) and colony formation assays
Exponentially growing cells were seeded into 96-well plates (2 × 10³ cells per well). Cell viability was measured using the Cell Counting Kit-8 (CCK-8) reagent (Beyotime, Shanghai, China) according to the manufacturer’s instructions. Absorbance at 450 nm was measured. For the colony formation assay, 1 × 10³ cells were plated in each well of a 6-well plate. After 14 days of incubation, colonies containing more than 50 cells were stained with crystal violet.

Cell migration and invasion assays
Transwell chambers (Corning Incorporated, Corning, NY, USA; 24-well, 8-μm pore size) uncoated or coated with Matrigel (BD Biosciences, San Jose, CA, USA) were employed to determine cell migration and invasion, respectively. A total of 5 × 10⁴ cells were seeded in the upper chamber and the complete culture medium was placed in the lower chamber. After incubation for 24 h, the migrated or invaded cells were stained with crystal violet.

Cell cycle and apoptosis detection
Cell cycle and apoptosis were analyzed using flow cytometry as previously described by our group [13]. The cell-cycle analysis kit and Annexin V-PE/7-AAD apoptosis kit were
obtained from Keygen Biotech (Nanjing, China). Operations were carried out according to the kit instructions.

Mouse xenograft experiments
Female BALB/c nude mice, 4–5 weeks old, were purchased from the Animal Center of Hubei University of Medicine. Cells \((2 \times 10^6)\) were injected subcutaneously into the flanks of nude mice \((n=5\) per group). Tumor volumes were calculated using the formula: \(\text{length} \times \text{width}^2 \times 0.5\). On day 28, the mice were sacrificed. The tumors were excised and stained with Ki-67 (abs123999, Absin) as previously described by other researchers [14].

Lectin pull-down assay
The membrane protein was extracted using a specific extraction kit (Beyotime) according to the manufacturer’s protocols. Then, Agarose bound Vicia Villosa Lectin (VVA; AL-1233-2, Vector Labs, Burlingame, CA, USA) was added to capture the lectin/glycoprotein complexes. The eluted fractions were subjected to SDS-PAGE and analyzed by Western blot.

RNA-sequencing (RNA-Seq)
RNA-seq was conducted by Biotecan Pharmaceuticals Co., Ltd (Shanghai, China). Total RNA from A549 cells with or without GALNT2 knockdown was subjected to RNA-seq. Differentially expressed genes (DEGs) were identified using the DESeq2 R package. The screening criterion was set as false discovery rate (FDR) < 0.05 and \(|\log_2(\text{fold change})| \geq 1\). DEGs were subjected to GO functional enrichment (Gene Ontology, http://www.geneontology.org/) and KEGG pathway analysis (Kyoto Encyclopedia of Genes and Genomes, http://www.genome.jp/kegg/).

Luciferase assay
The wild-type and mutant 3′-untranslated region (3′-UTR) of human GALNT2 were constructed and cloned into the pGL3 luciferase reporter vector (GenePharma). Subsequently, the above two vectors were co-transfected with miR-30d mimics or negative control mimics into A549 and H1299 cells, respectively. After 48 h of transfection, cells were collected for the luciferase activity assay using a Dual-Luciferase Reporter Assay System (Promega, Madison, WI, USA).

Statistical analysis
Data are presented as mean ± SD. Statistical analysis was done using the GraphPad Prism software (San Diego, CA, USA). Statistical significance was determined by Student’s t-test, chi-square test, ANOVA, or Pearson test. \(P<0.05\) was considered statistically significant.

Results
A comprehensive analysis of the GALNT family in NSCLC
To define the expression patterns of GALNTs in NSCLC, the mRNA levels of 20 GALNTs in NSCLC (LUAD, LUSC, and LCC) tissues and adjacent normal lung tissues were analyzed using the Oncomine database (Fig. 1A and Additional file 1: Table S3).
Compared with the normal lung tissues, the mRNA levels of GALNT2/3/6/7/10/14 were elevated in LUAD tissues, and GALNT15/18 mRNA levels were reduced. Moreover, GALNT1/2/3/14 mRNA levels were higher, while GALNT12/18 mRNA levels were lower in LUSC tissues than in normal lung tissues. Additionally, the mRNA levels of GALNT2/14 were increased, but GALNT5/12 mRNA levels were decreased in LCC tissues compared with normal lung tissues. Subsequently, the mRNA levels of 20 GALNTs in NSCLC (mainly LUAD and LUSC) tissues were investigated based on the results observed in the analysis. The gene expression profiles of the GALNT family members in NSCLC are shown in Fig. 1.
on the integrated analysis of TCGA and GTEx datasets (Fig. 1B). We observed that GALNT2/3/6/7/14 mRNA levels were upregulated in NSCLC tissues compared with the normal lung tissues; by contrast, GALNT5/13/16/18/20 mRNA levels were downregulated. Global gene expression analysis indicated that GALNT2 and GALNT14 were highly expressed, whereas GALNT5 was lowly expressed in NSCLC.

To determine the prognostic values of GALNTs in NSCLC, Kaplan–Meier survival analysis was performed. We found that high expression of GALNT2/9/13 was linked to decreased overall survival (Fig. 2A). Besides, high GALNT2/16 expression was associated with unfavorable disease-free survival (Fig. 2B). TCGA data analysis showed that only GALNT2 was related to the poor prognosis of all NSCLC patients. Furthermore, the survival information extracted from the Kaplan–Meier plotter database also confirmed that GALNT2 expression was negatively correlated with the

Fig. 2 Prognostic values of the GALNT family members in NSCLC. A Relationship between GALNT expression and overall survival. B Relationship between GALNT expression and disease free survival. Data for GALNT19 and GALNT20 were not available.
overall survival of NSCLC patients (Additional file 1: Fig. S1). Considered collectively, we selected GALNT2 as a candidate gene for further study.

Clinical importance of GALNT2 in NSCLC

RNA-Seq data for NSCLC were downloaded from the TCGA database. Univariate analysis showed that T stage, M stage, pathologic stage, primary therapy outcome, and GALNT2 expression were significantly associated with the overall survival. Multivariate Cox regression hazard analysis revealed that GALNT2 was an independent prognostic factor for overall survival (Additional file 1: Tables S4 and S5). A nomogram was established to predict the probability of the 1-, 3- and 5-year overall survival of NSCLC patients by integrating all the independent prognostic indicators (Fig. 3A). The ROC curve indicated that GALNT2 expression could serve as a promising biomarker for distinguishing NSCLC patients from normal controls (Fig. 3B). To verify the expression pattern of GALNT2 in NSCLC, RT-qPCR was performed. We confirmed that GALNT2 mRNA was overexpressed in NSCLC specimens (Fig. 3C). The results from the UALCAN website also demonstrated that GALNT2 protein expression was upregulated in NSCLC samples (Fig. 3D). To characterize GALNT2 abundance in human cancers, we examined GALNT2 expression using the TCGA pan-cancer data. We discovered that the expression of GALNT2 was widely elevated across numerous cancers (Fig. 3E). These findings suggested that GALNT2 might play a key role as an oncogene in NSCLC.

Oncogenic role of GALNT2 in NSCLC

To elucidate the biological function of GALNT2 in NSCLC, we analyzed the expression of GALNT2 in a panel of NSCLC cell lines and a normal lung cell line using RT-qPCR. Upregulation of GALNT2 expression was frequently detected in various NSCLC cell lines (Fig. 4A). Then, we conducted loss- and gain-of-function assays to explore whether GALNT2 could affect the malignant behaviors of NSCLC cells. Successful overexpression or knockdown of GALNT2 in A549 and H1299 cells was confirmed by RT-qPCR (Additional file 1: Fig. S2) and Western blot (Fig. 4B). The CCK-8, colony formation, and Transwell assays demonstrated that GALNT2 knockdown repressed the proliferative, migratory, and invasive abilities of NSCLC cells (Fig. 4C–E). Flow cytometry analysis revealed that knockdown of GALNT2 promoted apoptosis and induced G0/G1 cell cycle arrest (Fig. 4F, G). Overexpression of GALNT2 in A549 and H1299 cells, however, had the opposite effects. GALNT2 overexpression stimulated cell proliferation, migration, invasion, and cell cycle progression, and inhibited apoptosis (Fig. 5A–E). Together, these functional studies illustrated that GALNT2 exerted a tumor-promoting property in NSCLC.

GALNT2 facilitates NSCLC cell growth in vivo

To check whether GALNT2 could regulate tumorigenesis in vivo, A549 cells were subcutaneously injected into nude mice. Tumor volumes were measured every week. We discovered that the sizes and weights of tumors in the GALNT2 knockdown group were smaller than those in the control group; thus, knockdown of GALNT2 suppressed tumor formation in vivo (Fig. 6A–C). Meanwhile, the expression of Ki67, a marker of cell
proliferation, was decreased in the GALNT2 knockdown group (Fig. 6D). These results confirmed the oncogenic potential of GALNT2 in NSCLC.

**ITGA5 is a target glycoprotein of GALNT2 in NSCLC**

To uncover the regulatory mechanism of GALNT2 in NSCLC, we used LinkedOmics to screen the genes co-expressed with GALNT2. These genes were uploaded into GSEA for enrichment analysis (Additional file 2: Table S6). We observed a positive association between GALNT2 expression and the O-linked glycosylation or Integrin pathway (Fig. 7A). Lectin blot analysis proved that O-glycosylation was suppressed
following GALNT2 knockdown, as shown by reduced VVA staining in NSCLC cells (Fig. 7B). Among genes identified in the LinkedOmics database as related to the Integrin pathway, ITGA5 was most strongly correlated with GALNT2 (Fig. 7C). To evaluate whether GALNT2 could modify ITGA5 O-glycosylation, a lectin pull-down assay was performed. Interestingly, ITGA5 was pulled down by VVA, and the knockdown of GALNT2 decreased VVA binding to ITGA5 (Fig. 7D). To assess whether ITGA5 was implicated in GALNT2-mediated malignant phenotypes of NSCLC
Fig. 5 | GALNT2 overexpression promotes malignant phenotypes of NSCLC cells. 
A CCK-8 assay. B Colony formation assay. C Transwell migration and Matrigel invasion assays (magnification, ×100). D Flow cytometric analysis of cell cycle. E Analysis of cell apoptosis by flow cytometry. OV GALNT2 overexpression lentivirus, Mock normal control lentivirus. *P < 0.05

Fig. 6 | GALNT2 knockdown suppresses tumor growth in vivo. 
A Representative images of xenograft tumors formed. B Tumor growth curves. C Tumor weights. D Ki67 staining of tumor tissues (bar = 50 µm). shRNA GALNT2 shRNA lentivirus, shNC negative control shRNA lentivirus. *P < 0.05
cells, we silenced ITGA5 expression with siRNA in GALNT2-overexpressing cells (Fig. 7E). We found that GALNT2-induced cell proliferation, migration, and invasion were attenuated by silencing ITGA5 (Fig. 7F–H). Moreover, the interaction between GALNT2 and ITGA5 was validated by GeneMANIA (http://genemania.org/) (Fig. 7I). Thus, ITGA5 acted as a downstream effector of GALNT2 in NSCLC.
GALNT2 can activate the PI3K/Akt and MAPK/ERK pathways in NSCLC

GSEA analysis implied a crucial role for GALNT2 in the regulation of PI3K/Akt and MAPK pathways (Fig. 8A; Additional file 2: Table S6). To verify the GSEA results, we knocked down GALNT2 in A549 cells and extracted RNA for RNA-seq analysis (Additional file 3: Table S7). Different gene expression profiles were displayed between GALNT2-knockdown and control cells (Fig. 8B). DEGs were subjected to GO and KEGG for annotation. KEGG analysis revealed that DEGs were also enriched in the PI3K/Akt and MAPK pathways (Fig. 8C). To validate the association between these two
signaling pathways and GALNT2, the total protein and phosphorylation levels of Akt and ERK were examined via Western blot. As expected, the phosphorylation levels of Akt and ERK, but not the total levels of Akt and ERK, were reduced following GALNT2 knockdown (Fig. 8D). Furthermore, LY294002 (PI3K/Akt inhibitor, 10 μmol/L) or PD98059 (MAPK/ERK inhibitor, 10 μmol/L) could attenuate the promotive effects of GALNT2 on NSCLC cell proliferation, migration, and invasion (Fig. 8E–G). Therefore, the pro-oncogenic role of GALNT2 in NSCLC was related to the activation of the PI3K/Akt and MAPK/ERK pathways.

**miR-30d is an upstream regulator of GALNT2 in NSCLC**

To explore the mechanism underlying GALNT2 upregulation in NSCLC, the miRNAs potentially targeting GALNT2 were predicted using online analysis tools, such as TargetScan, RNA hybrid, PicTar, and miRanda. We found 9 potential miRNAs, which could interact with GALNT2 (Fig. 9A). In addition, the top 50 miRNAs associated with GALNT2 in NSCLC were screened out by the Pearson correlation test via LinkedOmics (Fig. 9B). After taking the intersection of those predicted miRNAs, we discovered that miR-30d presented the strongest association with GALNT2. Data from TCGA and our microarrays indicated that miR-30d was downregulated in NSCLC tissues and cell lines (Fig. 9C, D). Consequently, miR-30d was selected as the candidate miRNA. To further investigate whether GALNT2 was regulated by miR-30d, RT-qPCR and Western blot assays were conducted. Remarkably, both mRNA and protein levels of GALNT2 in A549 and H1299 cells were inhibited by miR-30d (Fig. 9E, F). Subsequently, the potential binding sequence between GALNT2 and miR-30d was predicted using TargetScan (Fig. 9G). As anticipated, miR-30d reduced the luciferase activity of the wild-type GALNT2 3′-UTR, while the luciferase activity of the mutant GALNT2 3′-UTR was not changed (Fig. 9H). To determine how the miR-30d/GALNT2 axis affected the malignant properties of NSCLC cells, rescue assays were performed. We discovered that miR-30d suppressed NSCLC cell proliferation, migration, and invasion, but these inhibition effects were restored by GALNT2 overexpression (Fig. 9I–K). All the above information pointed out that miR-30d was a negative modulator of GALNT2 in NSCLC. A schematic diagram was generated to illustrate the working mechanism of GALNT2 in NSCLC (Fig. 10).

**Discussion**

Altered glycosylation is a universal feature of human cancers. Glycosylation alteration also frequently occurs in lung cancer [15, 16]. Mucin-type O-glycosylation, which is produced by the GALNT family, is a fundamental form of glycosylation. GALNTs are ubiquitously expressed in a variety of tissues, and some GALNTs have oncogenic or tumor-suppressive properties. In the present study, we identified GALNT2, among the GALNT family members, as a key oncogene in NSCLC. This study is the first to clarify the critical role of GALNT2 in NSCLC. These findings enrich our understanding of the pathogenesis of NSCLC.

Multiple lines of evidence demonstrated that GALNT2 was dysregulated in several solid tumors, such as glioma [17], neuroblastoma [18], gastric adenocarcinoma [19], oral squamous cell carcinoma [20], and hepatocellular carcinoma [21]. Several recent
independent studies revealed that GALNT2 was overexpressed in lung adenocarcinoma [22, 23]. The present study, however, showed that GALNT2 was upregulated in all subtypes of NSCLC. High expression of GALNT2 was closely correlated with the poor prognosis in NSCLC patients. Both loss- and gain-of-function experiments proved that GALNT2 was indispensable for NSCLC cell proliferation, migration, and invasion. In addition, GALNT2 was able to regulate cell cycle progression and
apoptosis. Moreover, GALNT2 exerted pro-tumorigenic activity in nude mice. Thus, our study provides evidence that GALNT2 plays an oncogenic role in NSCLC.

Numerous proteins could serve as substrates for the GALNT family. For example, the O-glycosylation of SARS-CoV-2 spike protein was mediated by GALNT1 [24]. Loss of GALNT3 in pancreatic cancer was related to aberrant O-glycosylation of the ErbB family [25]. EGFR O-glycosylation in breast cancer was impacted by GALNT8 [11]. It was suggested that GALNT2 modulated the O-glycosylation of EGFR in glioma [17]. We confirmed that ITGA5 was O-glycosylated by GALNT2 in NSCLC, indicating that the substrates of GALNT2 may differ in various tissues. ITGA5, also known as integrin α5, is an important member of the integrin family. Integrins are transmembrane glycoproteins that participate in a range of biological events, including proliferation, adhesion, migration, and invasion [26]. Integrins have been extensively documented to play essential roles in NSCLC progression [27–29]. Importantly, the glycosylation of integrins is necessary for their proper folding and functionality [30]. In the current study, we found that the knockdown of GALNT2 inhibited ITGA5 O-glycosylation. The promotive effects of GALNT2 on the malignant phenotypes of NSCLC cells were reversed by ITGA5 silencing. Hence, we have reason to believe that ITGA5 acts as a major downstream effector of GALNT2 in NSCLC.

The PI3K/Akt and MAPK/ERK pathways are involved in cell proliferation, migration, invasion, and survival [31, 32]. The two pathways are commonly activated in NSCLC [33, 34]. Indeed, our GSEA and RNA-Seq data followed by functional assays confirmed that the oncogenic potential of GALNT2 in NSCLC was linked to the activation of the PI3K/Akt and MAPK/ERK pathways. The findings of our study provide new insights into the mechanism by which GALNT2 contributes to NSCLC tumorigenesis and malignant progression.

It is generally known that miRNAs can down-regulate the transcription of target genes [35]. However, it is not yet clear which miRNAs regulate the abundance of GALNT2 in NSCLC. It was found that GALNT2 expression was negatively modulated by let-7b in IgA nephropathy [36]. We discovered that miR-30d targeted GALNT2 and suppressed its expression in NSCLC. It has previously been suggested that miR-30d is a well-characterized tumor-suppressive miRNA [37]. To date, there is no published research on the association between miR-30d and GALNT2. In this study, we confirmed that miR-30d repressed NCSCL cell proliferation, migration, and invasion, whereas the inhibitory
effects of miR-30d were partially abrogated by GALNT2 overexpression. Therefore, miR-30d is an important upstream regulator of GALNT2 in NSCLC.

Undoubtedly, there are some limitations to this study. GALNT2 protein expression was not examined in NSCLC tissues. The O-glycosylation sites of ITGA5 are not well understood. It is of note that there exist other O-glycosylated proteins, which are not explored in this study. Further investigations are required to construct the regulatory network of miRNAs in controlling GALNT2 expression.

**Conclusions**

To sum up, we systematically illustrated the expression profile, prognostic value, biological function, and regulatory mechanism of GALNT2 in NSCLC. GALNT2 overexpression was frequently detected in NSCLC and could serve as a prognosis predictor. High expression of GALNT2 exhibited a tumor-promoting function in NSCLC. GALNT2 was able to enhance ITGA5 O-glycosylation and activate the PI3K/Akt and MAPK/ERK pathways. Additionally, the expression of GALNT2 was negatively regulated by miR-30d. As an important oncogene, GALNT2 has emerged as a potential therapeutic target for NSCLC.

**Abbreviations**

GALNTs  N-Acetylgalactosaminyltransferases
GTs    Glycosyltransferases
NSCLC  Non-small cell lung cancer
TCGA  The Cancer Genome Atlas
GSEA  Gene Set Enrichment Analysis
RT-qPCR  Transcription-quantitative real-time PCR
CCK-8  Cell Counting Kit-8
DEGs  Differentially expressed genes
VVA  Vicia Villosa Lectin

**Supplementary Information**

The online version contains supplementary material available at https://doi.org/10.1186/s11658-022-00378-w.

Additional file 1: Table S1. The sequences of miRNAs, siRNAs, and shRNAs. Table S2. Primer sequences used for RT-qPCR. Table S3. Transcriptional expression patterns of the GALNT family members in different lung cancer datasets (Oncomine database). Table S4. Univariate and multivariate Cox regression analysis in the TCGA-LUAD cohort. Table S5. Univariate and multivariate Cox regression analysis in the TCGA-LUSC cohort. Figure S1. The relationship between GALNT2 expression and overall survival in NSCLC. The data was retrieved from the Kaplan–Meier plotter database. Figure S2. The expression of GALNT2 was detected by RT-qPCR. *P < 0.05; **P < 0.01.

Additional file 2: Table S6. GESA enrichment analysis of genes co-expressed with GALNT6 in NSCLC.

Additional file 3: Table S7. RNA-seq after GALNT2 knockdown in A549 cells.

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**Author contributions**

LS and XD formulated the study protocol. QH, TT, and YL coordinated and collected the data. YT, SC, YL, JL, YL, and TL analyzed the data. All authors read and approved the final manuscript.

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**Availability of data and materials**

All data generated or analyzed during this study are included in this published article and its Additional files.
Declarations

Ethics approval and consent to participate
All procedures involving human participants were approved by the Ethics Committee of the Hubei University of Medicine. All animal experiments were approved by the Commission for Ethical Experimentation on Animals of the Hubei University of Medicine. All animal experiments were performed in accordance with the Declaration of Helsinki. All animal experiments were performed in accordance with international standards.

Consent for publication
Not applicable.

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

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References
1. Sung H, Ferlay J, Siegel RL, Laversanne M, Soerjomataram I, Jemal A, et al. Global cancer statistics 2020: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. CA Cancer J Clin. 2021;71:209–49.  
2. Onn SM, Changarathil G, Raja E, Ngo YX, Tateno H, Sada A, et al. Glycome profiling by lectin microarray reveals dynamic glycan alterations during epidermal stem cell aging. Aging Cell. 2020;19:e13190.  
3. Rodrigues JG, Duarte HO, Reis CA, Gomes J. Aberrant protein glycosylation in cancer: implications in targeted therapy. Biochem Soc Trans. 2021;49:843–54.  
4. Thomas D, Rathinavel AK, Radhakrishnan P. Altered glycosylation in cancer: a promising target for biomarkers and therapeutics. Biochim Biophys Acta Rev Cancer. 2021;1875:188464.  
5. Silinvinant A. Glycosylation markers in cancer. Adv Clin Chem. 2019;89:189–213.  
6. Wang M, Zhu J, Lubman DM, Gao C. Aberrant glycosylation and cancer biomarker discovery: a promising and thorny journey. Clin Chem Lab Med. 2019;57:407–16.  
7. Moremen KW, Ramiah A, Stuart M, Steel J, Meng L, Forouhar F, et al. Expression system for structural and functional studies of human glycosylation enzymes. Nat Chem Biol. 2018;14:156–62.  
8. Mohamed Abd-El-Halim Y, El Kaoutari A, Silvy F, Rubis M, Bigonnet M, Roques J, et al. A glycosyltransferase gene signature to detect pancreatic ductal adenocarcinoma patients with poor prognosis. EBioMedicine. 2021;71:103541.  
9. de Las RM, Lira-Navarrete E, Daniel EJP, Compáñon I, Coelho H, Dinziz A, et al. The interdomain flexible linker of the polypeptide GalNAc transferases dictates their long-range glycosylation preferences. Nat Commun. 2017;8:1959.  
10. Lin WR, Yeh CT. GALNT14: an emerging marker capable of predicting therapeutic outcomes in multiple cancers. Int J Mol Sci. 2020;21:1491.  
11. Huang T, Meng F, Huang H, Wang L, Liu Y, Wang J, et al. GALNT8 suppresses breast cancer cell metastasis potential by regulating EGFR-O-GalNAc glycosylation. Biochem Biophys Res Commun. 2022;601:16–23.  
12. Park MS, Yang AV, Lee JE, Kim SK, Roe JS, Oh MJ, et al. GALNT3 suppresses lung cancer by inhibiting myeloid-derived suppressor cell infiltration and angiogenesis in a TNFR and c-MET pathway-dependent manner. Cancer Lett. 2021;521:294–307.  
13. Luo Z, Hu Q, Tang Y, Leng Y, Tian T, Tian S, et al. Construction and investigation of B3GNT2-associated regulatory network in esophageal carcinoma. Cell Mol Biol Lett. 2022;27:8.  
14. Wang F, Hou W, Li X, Lu L, Huang T, Zhu M, et al. SETD8 cooperates with MZF1 to participate in hyperglycemia-induced endothelial inflammation via elevation of WNT5A levels in diabetic nephropathy. Cell Mol Biol Lett. 2022;27:30.  
15. Lemjabbar-Alaoui H, McKinney A, Yang YW, Tran VM, Phillips JJ. Glycosylation alterations in lung and brain cancer. Adv Cancer Res. 2015;126:305–44.  
16. Yang S, Xia J, Yang Z, Xu M, Li S. Lung cancer molecular mutations and abnormal glycosylation as biomarkers for early diagnosis. Cancer Treat Res Commun. 2021;27:100311.  
17. Sun Z, Xue H, Wei Y, Wang C, Yu R, Wang S, et al. Mucin O-glycosylating enzyme GALNT2 facilitates the malignant character of glioma by activating the EGFR/Pi3K/Akt/mTOR axis. Clin Sci. 2019;133:1167–84.  
18. Ho WL, Chou CH, Jeng YM, Lu MY, Yang YL, Jou ST, et al. GALNT2 suppresses malignant phenotypes through IGF-1 receptor and predicts favorable prognosis in neuroblastoma. Oncotarget. 2014;5:12247–59.  
19. Hu WT, Yeh CC, Liu SY, Huang MC, Lai IR. The O-glycosylating enzyme GALNT2 suppresses the malignancy of gastric adenocarcinoma by reducing EGFR activities. Am J Cancer Res. 2018;8:1739–51.  
20. Lin MC, Huang MJ, Liu CH, Yang TL, Huang MC. GALNT2 enhances migration and invasion of oral squamous cell carcinoma by regulating EGFR glycosylation and activity. Oral Oncol. 2014;50:478–84.  
21. Wu YM, Liu CH, Hu RH, Huang MJ, Lee JJ, Chen CH, et al. Mucin glycosylating enzyme GALNT2 regulates the malignant character of hepatocellular carcinoma by modifying the EGFR receptor. Cancer Res. 2011;71:7270–9.  
22. Yu Y, Wang Z, Zheng Q, Li J. GALNT2/14 overexpression correlates with prognosis and methylation: potential therapeutic targets for lung adenocarcinoma. Gene. 2021;790:145689.  
23. Wang W, Sun R, Zeng L, Chen Y, Zhang N, Cao S, et al. GALNT2 promotes cell proliferation, migration, and invasion by activating the Notch/Hes1-Pten-Pi3k/Akt signaling pathway in lung adenocarcinoma. Life Sci. 2021;276:119439.  
24. Zhang L, Mann M, Syed ZA, Reynolds HM, Tian E, Samara NL, et al. Furin cleavage of the SARS-CoV-2 spike is modulated by O-glycosylation. Proc Natl Acad Sci USA. 2021;118:e2109905118.
25. Chugh S, Meza J, Sheinin YM, Ponnusamy MR, Batra SK. Loss of N-acetylgalactosaminyltransferase 3 in poorly differentiated pancreatic cancer: augmented aggressiveness and aberrant ErbB family glycosylation. Br J Cancer. 2016;114:1376–86.
26. Koh KH, Cao Y, Mangos S, Tardi NJ, Dande RR, Lee HH, et al. Nonimmune cell-derived ICOS ligand functions as a renoprotective αvβ3 integrin-selective antagonist. J Clin Invest. 2019;129:1713–26.
27. Zhao G, Gong L, Su D, Jin Y, Guo C, Yue M, et al. Cullin5 deficiency promotes small-cell lung cancer metastasis by stabilizing integrin β1. J Clin Invest. 2019;129:972–87.
28. Wang M, Zhang G, Zhang Y, Cui X, Wang S, Gao S, et al. Fibrinogen alpha chain knockout promotes tumor growth and metastasis through integrin-αKT signaling pathway in lung cancer. Mol Cancer Res. 2020;18:943–54.
29. Ando T, Kage H, Matsumoto Y, Zokumasu K, Yotsumoto T, Maemura K, et al. Integrin α11 in non-small cell lung cancer is associated with tumor progression and postoperative recurrence. Cancer Sci. 2020;111:200–8.
30. Marsico G, Russo L, Quondamatteo F, Pandit A. Glycosylation and integrin regulation in cancer. Trends Cancer. 2018;4:537–52.
31. Song M, Bode AM, Dong Z, Lee MH. AKT as a therapeutic target for cancer. Cancer Res. 2019;79:1019–31.
32. Pashirzad M, Khorasanian R, Fard MM, Arjmand H, Langar P, Khazaei M, et al. The therapeutic potential of MAPK/ERK inhibitors in the treatment of colorectal cancer. Curr Cancer Drug Targets. 2021;21:932–43.
33. Lee L, Ramos-Alvarez I, Moody TW, Mantey SA, Jensen RT. Neuropeptide bombesin receptor activation stimulates growth of lung cancer cells through HER3 with a MAPK-dependent mechanism. Biochim Biophys Acta Mol Cell Res. 2020;1867:118625.
34. Tan AC. Targeting the PI3K/Akt/mTOR pathway in non-small cell lung cancer (NSCLC). Thorac Cancer. 2020;11:511–8.
35. Homayoonfal M, Asemi Z, Yousefi B. Targeting microRNAs with thymoquinone: a new approach for cancer therapy. Cell Mol Biol Lett. 2021;26:43.
36. Serino G, Sallustio F, Cucci C, Cox SN, Pesce F, De Palma G, et al. Role of let-7b in the regulation of N-acetylgalactosaminyltransferase 2 in IgA nephropathy. Nephrol Dial Transplant. 2015;30:1132–9.
37. Hou Y, Zhang Q, Pang W, Hou L, Liang Y, Han X, et al. YTHDC1-mediated augmentation of miR-30d in repressing pancreatic tumorigenesis via attenuation of RUNX1-induced transcriptional activation of Warburg effect. Cell Death Differ. 2021;28:3105–24.

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