Research Article

Overexpression of *eIF3D* in Lung Adenocarcinoma Is a New Independent Prognostic Marker of Poor Survival

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The eukaryotic initiation factor 3 (eIF3) is the largest and most complex translation initiation factor in mammalian cells. It consists of 13 subunits and among which several were implicated to have significant prognostic effects on multiple human cancer entities. To examine the expression profiles of *eIF3* subunits and determine their prognostic value in patients with lung adenocarcinoma (LUAD), the genomic data, survival data, and related clinical information were obtained from The Cancer Genome Atlas (TCGA) project for a secondary analysis. The results showed that among ten aberrantly expressed *eIF3* subunits in tumours compared with adjacent normal counterparts ($p < 0.05$), only upregulated *eIF3D* could predict poor overall survival (OS) outcome independent of multiple clinicopathological parameters (HR = 2.043, 95% CI: 1.132-3.689, $p = 0.018$). Chi-square analysis revealed that the highly expressed *eIF3D* group had larger ratios of patients with advanced pathological stage (68/40 vs. 184/206, $p = 0.0046$), residual tumour (13/4 vs. 163/176, $p = 0.0257$), and targeted molecular therapy (85/65 vs. 138/164, $p = 0.0357$). In silico analysis demonstrated that the altered expression of *eIF3D* was at least regulated by both copy number alterations (CNAs) and the hypomethylation of cg14297023 site. In conclusion, high *eIF3D* expression might serve as a valuable independent prognostic indicator of shorter OS in patients with LUAD.

1. Introduction

Lung cancer is one of the most common and lethal diseases afflicting the global population. It is histologically divided into small cell lung cancer (SCLC) and non-small-cell lung cancer (NSCLC). Lung adenocarcinoma (LUAD) is the most common subtype of NSCLC, accounting for more than 40% of NSCLC cases and showing a clear upward trend in the past few decades [1]. The overall survival (OS) rate for LUAD patients remains low despite the advances in clinical treatment. Due to the insignificant early symptoms and the lack of specific markers, the early diagnosis of LUAD is extremely difficult, and patients usually have a stage of local progression or later when they are diagnosed, missing the best time of surgery [2–4]. Therefore, the search for new and effective biomarkers to screen out high-risk patients and predict prog-

nosis of LUAD has become an important part of lung cancer prevention and treatment.

mRNA translation is a key step in the regulation of eukaryotic gene expression. Its dysregulation would result in abnormal gene expression and lead to uncontrolled cell growth, which potentially caused tumourigenesis [5]. The initiation of mRNA translation is the rate-limiting step during which translation regulation is primarily achieved and its dysregulation has received considerable attention, as well as aberrant expression of eukaryotic translation initiation factors (eIFs) in various cancer entities has been observed in a growing number of studies [6–14]. eIF3, the largest and most complex eIF in mammalian cells, consists of 13 non-identical subunits (eIF3a–m). The main function of eIF3 has been demonstrated to promote nearly every step of translation initiation as a scaffold, including promoting the 43S

preinitiation complex formation, stimulating mRNA binding with 43S preinitiation complex, scanning and recognizing AUG start codon, and dissociating the posttermination 80S ribosome by binding with 40S ribosomal subunit [15–18]. The altered expression of *eIF3* subunits has been found in a wide range of tumour entities, including breast cancer [19–21], gastric cancer [22], NSCLC [23], and liver cancer [24], supporting their high potential use as prognostic biomarkers and therapeutic cancer targets. To the best of our knowledge, no previous studies have examined the roles of overall *eIF3* family members in LUAD. In this study, using gene expression data and survival data of LUAD patients from The Cancer Genome Atlas (TCGA) project, we examined the expression profiles of *eIF3* subunits and determined their prognostic roles in LUAD, as well as the underlying mechanisms of *eIF3D* dysregulation.

2. Materials and Methods

2.1. Clinical Cohorts and RNA-Seq Data from TCGA. mRNA expression data, survival data, and related clinical information of patients with LUAD in TCGA project were downloaded from the UCSC Xena browser (<https://xenabrowser.net/>). Only patients who had primary tumours and had not received neoadjuvant therapy were included in this analysis. The genomic and phenotypic data were collected. Briefly, RNA-Seq data of *eIF3* subunit genes were extracted. The phenotypic data included sample types, histological types, age at initial pathologic diagnosis, gender, pathologic stage, radiation therapy, targeted molecular therapy, residual tumour status, canonical mutations in *KRAS/EGFR/ALK*, tobacco smoking history, OS status, and OS time (weeks). To explore the underlying mechanisms of *eIF3D* dysregulation, the corresponding somatic mutations, copy number alterations (CNAs), and DNA methylations of *eIF3D* were obtained. The DNA methylation status was examined by the Infinium Human Methylation 450 Bead Chip.

2.2. Statistical Analysis. Statistical analysis was performed using GraphPad Prism 8 (GraphPad Inc., CA, USA) and SPSS Statistics 20 (SPSS Inc., IL, USA) in this study. Comparison between two groups was performed using unpaired *t*-test with Welch's correction. The differences in clinicopathological parameters between LUAD patients with high or low gene expression were compared using the chi-square test by two-sided Fisher's exact test. Gene expression levels were categorized as low or high according to the median values. Kaplan-Meier curves of OS were generated using GraphPad Prism, and differences in survival rate between groups were analysed by log-rank test. Univariate and multivariate Cox regression models were utilized to assess the prognostic value of *eIF3* subunits regarding OS. Genes with a *p* value of <0.05 in log-rank test were subsequently included in univariate and multivariate Cox analyses. Forest plots for multivariate Cox regression models were also drawn by GraphPad Prism. Pearson correlation coefficients between *eIF3D* expression and its linear DNA copy number values in patients, as well as the methylation level of its CpG site cg14297023, were calculated to assess their correla-

tions. *p* < 0.05 indicates statistically significant for all above statistical analyses.

3. Results

3.1. The mRNA Expression Levels of *eIF3* Subunits in LUAD Patients. Using RNA-Seq data in TCGA, the mRNA expression levels of all 13 *eIF3* subunits were compared between LUAD (*n* = 511) and adjacent normal tissues (*n* = 58). As shown in Figure 1, levels of nine *eIF3* subunits involving *eIF3A*, *eIF3B*, *eIF3C*, *eIF3D*, *eIF3E*, *eIF3H*, *eIF3I*, *eIF3J*, and *eIF3M* were significantly increased in cancer tissues, whereas the *eIF3L* expression level in tumours was independently decreased than that of normal tissues (*p* < 0.05). No significant variation was observed in *eIF3F*, *eIF3G*, and *eIF3K* levels between the two groups (*p* ≥ 0.05).

3.2. Association between *eIF3* Subunit Expression Levels and Clinical Features in LUAD Patients. Next, we summarized the association between gene expression levels of the 13 *eIF3* subunits and clinicopathological parameters of LUAD patients (Table 1). Chi-square analysis showed that five subunit genes associated significantly with a single clinicopathological variable, respectively, and among which *eIF3C* and *eIF3H* expression increased in high pathological stage (*eIF3C*: 64/44 vs. 188/207, *p* = 0.0388; *eIF3H*: 67/41 vs. 186/209, *p* = 0.0066), *eIF3F* and *eIF3L* expression increased in patients with ages no lower than 65 (*eIF3F*: 144/129 vs. 94/125, *p* = 0.0367; *eIF3L*: 151/122 vs. 93/126, *p* = 0.0050), while high expression of *eIF3G* was observed more in female patients (157/120 vs. 102/132, *p* = 0.0034).

Among the five genes whose expression levels were significantly associated with two clinicopathological variables or more, the high *eIF3A* expression group had larger proportion of male patients (134/100 vs. 120/157, *p* = 0.0019) and patients with smoking history (220/202 vs. 29/46, *p* = 0.0336), while the expression of the other four genes was consistently positively associated with high pathological stage (*eIF3B*: 66/42 vs. 189/206, *p* = 0.0168; *eIF3D*: 68/40 vs. 184/206, *p* = 0.0046; *eIF3E*: 64/44 vs. 186/209, *p* = 0.0297; *eIF3J*: 63/45 vs. 183/212, *p* = 0.0299). Besides, the more highly expressed *eIF3B* was also discovered in patients with ages lower than 65 (126/93 vs. 122/151, *p* = 0.0050), radiation therapy (37/21 vs. 194/202, *p* = 0.0483), and targeted molecular therapy (87/63 vs. 143/159, *p* = 0.0361). The high *eIF3D* expression group had a significantly larger ratio of patients with residual tumour (13/4 vs. 163/176, *p* = 0.0257) and targeted molecular therapy (85/65 vs. 138/164, *p* = 0.0357). And high expression of *eIF3E* and *eIF3J* was observed more in patients with residual tumours (13/4 vs. 172/170, *p* = 0.0456) and without mutations in *KRAS/EGFR/ALK* (47/48 vs. 86/46, *p* = 0.0206), respectively. No significant association was observed between *eIF3I*, *eIF3K*, *eIF3M*, and other above-mentioned clinical features.

3.3. The Prognosis Value of *eIF3* Subunit Expression in LUAD Patients. Since we observed that ten *eIF3* subunits were significantly aberrantly expressed in tumours compared with adjacent area, we next evaluated the association between

FIGURE 1: Plot charts showing mRNA expression of *eIF3* subunits in LUAD tissues and adjacent normal tissues. Adj. N: adjacent normal tissues.

(a)

[illegible]

(b)

Parameters	<i>n</i> = 511	<i>eIF3E</i> expression (<i>n</i> = 255)		<i>eIF3F</i> expression (<i>n</i> = 253)		<i>p</i> value		<i>eIF3G</i> expression (<i>n</i> = 259)		<i>p</i> value		<i>eIF3H</i> expression (<i>n</i> = 257)		<i>p</i> value		<i>eIF3I</i> expression (<i>n</i> = 254)		<i>p</i> value	
		High	Low	High	Low	High	Low	High	Low	High	Low	High	Low	High	Low	High	Low	High	Low
Age (years, mean ± SEM)	<65 ≥65 No data	103 140	116 133	94 144	125 129	0.3652	0.0367*	112 136	107 137	0.7861	0.7856	111 134	108 139	0.7856	0.7856	110 133	109 140	0.7856	0.7856
Gender	Females Males	141 114	136 120	148 105	129 129	0.6574	0.0622	157 102	120 132	0.0034*	0.3749	134 123	143 111	0.3749	0.3749	141 113	136 121	0.5945	0.5945
Pathological stage	I/II III/IV No data	186 64	209 44	189 60	206 48	0.0297*	0.1599	195 58	200 50	0.4484	0.0066*	186 67	209 41	0.0066*	0.0066*	198 54	197 54	>0.9999	>0.9999
Residual tumour	R0 R1/R2 RX/no data	172 13	170 4	178 10	164 7	0.0456*	0.6275	176 11	166 6	0.3280	0.2123	178 12	164 5	0.2123	0.2123	184 11	158 6	0.4592	0.4592
Smoking history	1 2/3/4/5 No data	35 209	40 213	34 210	41 212	0.7074	0.5315	44 206	31 216	0.1329	0.2107	32 215	43 207	0.2107	0.2107	42 202	33 220	0.2113	0.2113
Radiation therapy	No Yes No data	190 32	206 26	195 27	201 31	0.3274	0.7789	200 30	196 28	0.8890	0.1616	192 34	204 24	0.1616	0.1616	191 29	205 29	0.8883	0.8883
Targeted molecular therapy	No Yes No data	149 72	153 78	153 68	149 82	0.8417	0.3180	159 71	143 79	0.3181	0.7650	148 76	154 74	0.7650	0.7650	141 76	161 74	0.4841	0.4841
Mutations in <i>KRAS/EGFR/ALK</i>	No Yes No data	78 66	54 29	77 57	55 38	0.1252	0.8913	64 53	68 42	0.2853	0.1723	72 61	60 34	0.1723	0.1723	81 53	51 42	0.4148	0.4148

(c)

Parameters	n = 511	<i>eIF3J</i> expression		<i>eIF3K</i> expression		<i>eIF3L</i> expression		<i>eIF3M</i> expression		p value
		High (n = 252)	Low (n = 259)	High (n = 255)	Low (n = 256)	High (n = 255)	Low (n = 256)	High (n = 254)	Low (n = 257)	
Age (years, mean ± SEM)	<65	102	117	102	117	93	126	104	115	0.5865
	≥65	141	132	141	132	151	122	137	136	
	No data									
Gender	Females	129	148	145	132	144	133	142	135	0.4778
	Males	123	111	110	124	110	124	112	122	
	I/II	183	212	192	203	191	204	187	208	0.0505
Pathological stage	III/IV	63	45	59	49	57	51	63	45	
	No data									
Residual tumour	R0	165	177	172	170	170	172	170	172	0.3208
	R1/R2	9	8	10	7	10	7	11	6	
	RX/no data									
Smoking history	1	36	39	36	39	39	36	38	37	0.8024
	2/3/4/5	206	216	210	212	207	215	205	217	
	No data									
Radiation therapy	No	190	206	191	205	199	197	189	207	>0.9999
	Yes	30	28	32	26	29	29	28	30	
	No data									
Targeted molecular therapy	No	138	164	154	148	147	155	141	161	0.5489
	Yes	81	69	69	81	81	69	75	75	
	No data									
Mutations in <i>KRAS/EGFR/ALK</i>	No	86	46	70	62	71	61	75	57	0.6855
	Yes	47	48	50	45	62	33	51	44	
	No data									

*Indicates statistical significance ($p < 0.05$).

their expression levels and the survival outcomes of LUAD patients via Kaplan-Meier survival analysis (Figure 2). The results showed that patients with high expression levels of *eIF3B*, *eIF3D*, or *eIF3M* exhibited worse prognosis than those with low expression levels ($p_{eIF3B} = 0.0034$, $p_{eIF3D} = 0.0337$, and $p_{eIF3M} = 0.0094$). On the contrary, the elevated expression of *eIF3L* predicted better prognosis for our studied patients with LUAD ($p = 0.0326$). No significant differences in prognosis were observed between patients with high and low *eIF3A*, *eIF3C*, *eIF3E*, *eIF3H*, *eIF3I*, and *eIF3J* expression levels.

To verify the robust prognostic roles of *eIF3B*, *eIF3D*, *eIF3L*, and *eIF3M*, the univariate and multivariate Cox regression analyses were performed. Univariate analysis showed that besides the expression levels of the four genes ($HR_{eIF3B} = 1.563$, $p = 0.004$; $HR_{eIF3D} = 1.377$, $p = 0.034$; $HR_{eIF3L} = 0.710$, $p = 0.023$; and $HR_{eIF3M} = 1.521$, $p = 0.006$), pathological stage ($HR = 2.581$, $p < 0.001$), radiation therapy ($HR = 2.016$, $p < 0.001$), and residual tumour ($HR = 4.187$, $p < 0.001$) were also significant prognostic factors (Table 2). By running four separate multivariate models (*eIF3B*, *eIF3D*, *eIF3L*, and *eIF3M*) adjusted by a series of clinicopathological parameters, it demonstrated that *eIF3D* expression was an independent risk factor for OS ($HR = 2.043$, 95% CI: 1.132–3.689, $p = 0.018$, Figure 3(b)), while no significant correlations were observed between other three genes and the OS (Figures 3(a), 3(c), and 3(d)). This result indicated that the overexpression of *eIF3D* could independently predict poor prognosis for patients with LUAD.

3.4. In Silico Analysis of the Potential Mechanisms Underlying *eIF3D* Upregulation in LUAD Patients. To explore the mechanisms of *eIF3D* dysregulation in patients with LUAD, we examined the correlations between *eIF3D* mRNA expression and its genetic (typically somatic mutations and CNAs) and epigenetic (typically methylation) alterations using the corresponding data in TCGA. Results showed that among 543 cases with somatic mutations measured, only three missense mutations were observed (data not shown); thus, we did not make further statistical analysis between *eIF3D* expression and its somatic mutations for the insufficient number of mutation cases.

Among 508 cases with DNA CNAs identified, 68 cases (13.4%) had CNA gains (+1/+2), 224 cases (44.1%) had CNA losses (-1/-2), and 216 cases (42.5%) were copy neutral (0). Unpaired *t*-test analysis showed both CNA gains and losses had significant influence on *eIF3D* expression ($p < 0.0001$, Figure 4(b)) and that a strong correlation was also observed between *eIF3D* expression and its linear copy number values by a regression analysis (Pearson $r = 0.665$, $p < 0.0001$, Figure 4(c)). Besides, Kaplan-Meier survival analysis showed that patients with CNA gains exhibited significantly worse prognosis than those with CNA losses and copy neutral ($p = 0.0427$, Figure 4(d)). As for *eIF3D* DNA methylation in LUAD, the methylation status of 23 CpG sites was measured by Methylation 450k, and among which nine were hypomethylated in tumour tissues ($n = 455$) compared with the adjacent counterparts ($n = 31$) (Table S1). The regression analysis demonstrated a

significantly negative correlation between *eIF3D* expression and a single CpG site of cg14297023 in tumour tissues (Pearson's $r = -0.2105$, $p < 0.0001$, Figure 4(f)), but no direct correlation was observed between cg14297023 methylation levels and the survival outcomes ($p = 0.5788$, Figure 4(g)). Overall, these results indicated that the expression of *eIF3D* was at least regulated by both CNAs and cg14297023 methylation.

4. Discussion

In this study, we examined the mRNA expression profiles of all 13 *eIF3* family members and determined their prognostic roles in LUAD patients using corresponding data from TCGA project. The integrated results demonstrated that the overexpression of *eIF3D* could independently predict poor prognosis for patients with LUAD.

Eukaryotic translation initiation depends on ribosomal subunits and, at least, 12 auxiliary proteins named eukaryotic initiation factors (eIFs), and among which eIF3 is the largest and most complex one comprising 13 subunits assembled together in an orderly way. Deregulation of *eIF3* expression and/or function has been proposed to play either a causal role or at least contribute to the etiology of various cancer entities. In our study, ten *eIF3* subunits were observed aberrantly expressed in LUAD tumour tissues than that of normal counterparts. Studies have revealed that imbalanced expression of single *eIF3* subunit may affect the overall expression profiles of the entire *eIF3* complex [26], which may explain why the vast majority of *eIF3* subunits were observed to be aberrantly expressed in our study. The misregulation of *eIF3* subunits may contribute to malignancy via several possible means. First, the altered expression of mammalian *eIF3* subunits, especially those in the core regions defined as octamer (a, c, e, f, h, k, l, m) and YLC (Yeast-Like-Core: b, g, i), may impact on the correct assembly of the entire eIF3 complex [26], leading to irregular mRNA translation and causing disease or its quick progression. Second, the overexpression of single *eIF3* subunit and/or the resulting upregulation of the entire eIF3 complex would induce a more efficient translation initiation rate of specific mRNAs, which is known as a common feature in cancer [27]. Third, besides translation initiation, the *eIF3* subunits may function in other important molecular events or cellular processes associated with a disease phenotype.

The *eIF3D*, a peripheral subunit attaching to the eIF3 holocomplex via eIF3E, showed very distinct characteristics with its other family members. It is the only subunit whose misregulation affects neither the expression of the other *eIF3* subunits nor the integrity of the eIF3 complex but is nonetheless essential for cell proliferation [26]. Instead of the canonical roles of eIF3 in general translation, the essential biological function of *eIF3D* lies in driving cap-dependent but eIF4E-independent expression of a specific subset of mRNAs encoding proteins with vital cellular roles [28, 29]. As a noncanonical cap-binding protein, *eIF3D* was also found to act as an indispensable assistant to help DAP5 which was supposed to initiate only IRES- (internal ribosome entry site-) mediated translation to promote a widespread alternate form of cap-dependent mRNA translation [30].

FIGURE 2: Kaplan-Meier survival analysis of differentially expressed subunits of *eIF3* for OS in LUAD patients. χ^2 and p values assessed by log-rank test and number of patients in each group are provided in the figure.

TABLE 2: Univariate analysis of OS in LUAD patients.

Parameters	HR	Univariate analysis 95% CI (lower-upper)	p value
Age			
<65 (N = 220)	1.000		
≥65 (N = 273)	1.145	0.849-1.545	0.375
Gender			
Female (N = 273)	1.000		
Male (N = 230)	1.030	0.768-1.382	0.843
Pathological stage			
I/II (N = 392)	1.000		
III/IV (N = 103)	2.581	1.884-3.535	<0.001*
Radiation therapy			
No (N = 392)	1.000		
Yes (N = 58)	2.016	1.372-2.962	<0.001*
Residual tumour			
R0 (N = 336)	1.000		
R1/R2 (N = 16)	4.187	2.333-7.512	<0.001*
Smoking history			
1 (N = 72)	1.000		
2/3/4/5 (N = 417)	0.888	0.588-1.341	0.571
Mutations			
No (N = 125)	1.000		
Yes (N = 91)	0.866	0.558-1.346	0.523
Targeted therapy			
No (N = 299)	1.000		
Yes (N = 149)	1.159	0.830-1.620	0.386
<i>EIF3B</i> expression			
Low (N = 247)	1.000		
High (N = 251)	1.563	1.156-2.111	0.004*
<i>eIF3D</i> expression			
Low (N = 248)	1.000		
High (N = 250)	1.377	1.024-1.854	0.034*
<i>EIF3L</i> expression			
Low (N = 245)	1.000		
High (N = 253)	0.710	0.528-0.955	0.023*
<i>EIF3M</i> expression			
Low (N = 246)	1.000		
High (N = 252)	1.521	1.127-2.053	0.006*

*Indicates statistical significance ($p < 0.05$).

The overexpression of *eIF3D* has been observed to promote cell proliferation, migration, or/and tumour growth in several cancer entities, including ovarian cancer [31], renal cell carcinoma [32], gastric cancer (GC) [33], and gallbladder cancer (GBC) [34]. As being discovered in our study, all these high levels of *eIF3D* are associated with advanced tumour stage, indicating its potential role in tumour development. The *eIF3D* knockdown researches conducted in a series of cancer cell lines involving breast cancer [35], NSCLC [36], melanoma [37], acute myeloid

leukaemia [38], and colon cancer [39] revealed that the deletion of *eIF3D* caused a significant reduction in cell proliferation and colony formation due to an arrest of cell cycle at G2/M phase, suggesting its key role in cell cycle control and apoptosis. Previous studies demonstrated that *eIF3D* affected cancer cell growth via multiple signaling pathways. Zhang et al. proposed that *eIF3D* exerted the tumour-promoting activities through GRK2-mediated activation of PI3K/AKT pathway in GBC [34]. Fan et al. found that knockdown of *eIF3D* inhibited the activation of Wnt/ β -catenin signaling pathway in breast cancer cell lines via blockade of the expression of β -catenin, cyclin D1, and c-Myc [35]. By exploring the modifications of effector proteins in signaling pathways responsible for cell growth and apoptosis, the activations of three cancer-related molecules—AKT, HSP27, and SAPK/JNK—were found to be reduced by *eIF3D* knockdown in NSCLC cells [36]. Moreover, a series of phosphorylation upregulation was observed along with downregulation of *eIF3D* in colon cancer cells, including AMPK α , Bad, PRAS409, SAPK/JNK, and GSK3 β , as well as the cleavage of PARP [39]. These mechanisms would help to explain the oncogenic properties of *eIF3D* in patients with LUAD. In our study, the prognostic value of *eIF3D* was confirmed in LUAD independent of multiple clinicopathological parameters. Being consistent with our findings, the strong correlations between high *eIF3D* expression and poor OS outcomes were also observed in patients with GC [33] and GBC [34], supporting its high potential role to serve as a useful prognostic marker and therapeutic target for the treatment of these cancers.

In this study, by assessing the potential mechanisms underlying *eIF3D* upregulation in LUAD, we found that the *eIF3D* dysregulation was partly regulated by the CNAs, as well as the hypomethylation of cg14297023 site which locates in the 3'UTR region of *eIF3D* gene according to the MethHC database (<http://methhc.mbc.nctu.edu.tw/php/index.php>) [40]. It is clear that a CNA is generally positively associated with the expression level of its corresponding gene. However, as for the role of DNA methylation in the 3'UTR on the influence of gene expression, far less is known than a comprehensive understanding of that in the gene promoter region. Recently, McGuire et al. proposed an explanation that DNA methylation in 3'UTR might control gene expression via two potential ways [41]. First, it may influence gene expression by increasing binding of proteins with methylation-binding domains or inhibiting other special protein binding or mask sequence recognition [42, 43]. Another potential explanation is that if different lengths of the 3'UTR are dependent on methylation, transcripts with shorter 3'UTRs would have greater mRNA stability and thereby higher gene expression [44]. On the contrary, studies also found that gene body methylation would reflect as a consequence of higher gene expression, rather than as a cause [45]. Therefore, the specific role of hypomethylation of cg14297023 site on *eIF3D* expression in LUAD cannot be concluded yet based on these hypotheses. Beyond that, we could not exclude other genetic or epigenetic mechanisms influencing the transcription and translation of *eIF3D*. Thus,

FIGURE 3: Forest plots for multivariate Cox regression models of four *eIF3* subunits: (a) *eIF3B*; (b) *eIF3D*; (c) *eIF3L*; (d) *eIF3M*.

FIGURE 4: Copy number alterations and DNA methylation analysis of *eIF3D* in LUAD patients. (a) Heatmap showing *eIF3D* expression and corresponding DNA copy number alterations, averaged DNA methylation, and methylation in cg14297023. (b) Plot chart showing *eIF3D* expression in different CNA groups. (c) Linear regression analysis of the correlation between *eIF3D* expression and its linear copy number values. (d) Kaplan-Meier survival analysis of different *eIF3D* CNA levels for OS in patients with LUAD. (e) Plot chart showing DNA methylation of cg14297023 site in LUAD tissues and adjacent normal tissues. (f) Linear regression analysis of the correlation between cg14297023 methylation and *eIF3D* expression. (g) Kaplan-Meier survival analysis of different methylation levels in cg14297023 for OS in patients with LUAD. Adj. N: adjacent normal tissues; GISTIC: Genomic Identification of Significant Targets in Cancer [25].

experimental studies are required to further explore the role and other potential mechanisms of *eIF3D* in LUAD.

With regard to other subunits of *eIF3*, their misregulation and potential roles in tumour progression or survival outcome have been observed in a vast range of cancer entities (reviewed by [18, 46, 47]). In our study, even though their independent prognostic roles were not observed, we still got some noteworthy results. A previous study by Cattie et al. has revealed that loss-of-function mutations in *eIF3L* gene resulted in a 40% extension in lifespan in *Caenorhabditis elegans*, indicating its pivotal functional role in the regulation of cellular and organismal responses to aging [48]. By examining the links between gene expression levels and clinical features, the highly expressed *eIF3L* was found positively associated with LUAD patients with ages no lower than 65 (151/122 vs. 93/126, $p = 0.0050$, Table 1) in our study, adding to the evidence of its regulatory role during aging, as Cattie et al. proposed [48]. As an associated factor locating at the more peripheral position of the *eIF3* complex, *eIF3J* is known to be weakly and unstably interacted with the complex and most likely plays only a supportive role in the overall translational efficiency in human cells [49]. However, little is known in the relation of *eIF3J* and carcinogenesis by now [15, 50, 51]. The mutations in *KRAS/EGFR/ALK* are the most common “driver” mutations detected in LUAD and are considered to play pivotal roles in carcinogenesis at multiple levels, thus acting as important genomic-guided therapeutic targets in LUAD (reviewed by [3]). In our study, the discovery of different expression levels of *eIF3J* occurring in patients with or without mutations in *KRAS/EGFR/ALK* (86/46 vs. 47/48, $p = 0.0206$, Table 1) may provide a clue of its potential impact on LUAD carcinogenesis accompanied by *KRAS/EGFR/ALK* genomic alterations. Besides, nine *eIF3* subunits beyond *eIF3D* were found aberrantly expressed in LUAD tissues and in which the expression levels of five subunits (*eIF3B*, *eIF3C*, *eIF3E*, *eIF3H*, and *eIF3J*) increased in LUAD patients with high tumour stage, implicating their roles in the maintenance or progression of LUAD. Therefore, it is meaningful to further explore the potential roles and underlying molecular mechanisms of action of these *eIF3* subunits as well as *eIF3D* in LUAD in the future.

5. Conclusions

High *eIF3D* expression might serve as a valuable independent prognostic indicator of shorter overall survival in patients with LUAD.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

All authors declare that they have no conflicts of interest.

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Supplementary Materials

Table S1: methylation status of *eIF3D* CpG sites in LUAD tissues and adjacent normal tissues. (*Supplementary Materials*)

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