

Clinicopathological characteristics and genetic analysis of pulmonary carcinoid tumors: A single-center retrospective cohort study and literature review

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Abstract. Pulmonary carcinoid tumors, including typical and atypical carcinoids, are well-differentiated neuroendocrine tumors (NETs) that represent 1-2% of all lung cancer cases. In the present study, all cases of well-differentiated NETs diagnosed at Tianjin Medical University General Hospital (Tianjin, China) between 2006 and 2016 were reviewed, and 20 pulmonary carcinoid cases were identified. The clinical features of these cases were summarized, and the results of pathological and imaging examinations were collated. As a low-grade malignant pulmonary neoplasm, the molecular biological mechanism of pulmonary carcinoids is yet to be elucidated. To investigate the underlying molecular mechanisms behind pulmonary carcinoids and to determine an effective molecular targeted therapeutic strategy, next-generation sequencing (NGS) was performed using tissue samples from six patients to determine additional molecular biological characteristics that may help guide targeted therapy. A total of 27 somatic mutations in 21 genes were detected. Of note, mutations in the KIT proto-oncogene receptor tyrosine kinase, Erb-B2 receptor tyrosine kinase 4, MET proto-oncogene receptor tyrosine kinase and insulin-like growth factor 1 genes occurred in

two out of six cases. Since treatments for advanced carcinoids are relatively ineffective, molecular profiling may contribute to the identification of novel treatments. In addition, the literature on mutations in pulmonary carcinoids was reviewed and available clinical information and features of this tumor type were summarized.

Introduction

Neuroendocrine tumors (NETs) are a subtype of neoplasms that can arise in the majority of organs and share a number of common biochemical and pathologic features (1). Pulmonary NETs comprise 20-30% of all NETs (2), and NETs in the lung can be divided into four subtypes according to their malignancy grade: Typical carcinoids (TCs), atypical carcinoids (ACs), large-cell neuroendocrine carcinomas (LCNECs) and small-cell lung cancers (SCLCs). Of these subtypes, typical and atypical carcinoids are generally termed pulmonary carcinoids and constitute 1-2% of all pulmonary malignancies; however, their incidence has notably increased in recent decades; Petursdottir *et al* (3) reported that the incidence of PC increased from 1.9/1,000,000 (1955-1964) to 5.8/1,000,000 (2005-2015) per year in Iceland (4). Complete surgical resection is the primary choice of treatment for early-stage lung carcinoids (2). However, efficient management strategies for advanced-stage lung carcinoids are limited (2). As the development of precision medicine has progressed, molecular targeted therapy has achieved breakthroughs for the treatment of pulmonary carcinoids, including epidermal growth factor receptor (EGFR) inhibitors, mammalian target of rapamycin (mTOR) inhibitors, bevacizumab and tyrosine kinase inhibitors (TKIs) (5-7). The present study aimed to analyze the clinicopathological characteristics of patients admitted to Tianjin Medical University General Hospital (Tianjin, China) center who underwent surgical resection for pulmonary carcinoids, and gene mutation profiling was performed to explore the underlying molecular mechanisms. In addition, gene mutation

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information of pulmonary carcinoids was summarized from relevant literature.

Materials and methods

Ethical approval. The present study was conducted in accordance with the standards of the Declaration of Helsinki for medical research involving human subjects. All subjects provided written informed consent, and the study protocol was approved by the clinical research ethical review board at Tianjin Medical University General Hospital (Tianjin, China).

Study design. Patient data were reviewed between January 2006 and December 2016 at Tianjin Medical University General Hospital, and information on 20 patients with lung carcinoid tumors with complete medical records was collected. The clinical features and imaging data from patient records were summarized. All pulmonary carcinoid cases were reviewed according to the World Health Organization criteria (2015) and were staged according to the American Joint Committee on Cancer staging manual (8th edition) criteria (8,9). Carcinoid tumors of the lung were classified as typical carcinoids (TCs) or atypical carcinoids (ACs) based on the following histological differences: The number of mitoses per 10 high-power fields (TC mitotic index, <2; AC mitotic index, 2-10; SCLC/LCNECs mitotic indices, >10) (10); the presence of necrosis; increased cellularity with disorganization; nuclear pleomorphism; hyperchromatism; and an abnormal nuclear: Cytoplasmic ratio (11,12). In general, macroscopic pulmonary carcinoid tissues appeared as smooth, highly vascular, gray-yellow and notably demarcated masses (1,9,13,14). The diagnosis of pulmonary carcinoid can be established by hematoxylin and eosin (HE) staining of a histopathologic section. However, immunohistochemical (IHC) staining is more precise for the diagnosis of pulmonary carcinoids compared with HE; specifically, staining for synaptophysin, chromogranin A and neural cell adhesion molecule (NCAM) can distinguish high-grade NETs (LCNECs and SCLCs) from pulmonary carcinoids (15).

Tissue sections (5 μ m thick) were prepared from paraffin-embedded tissue blocks using formalin (10% methanol) solution as a fixative. The sections were stained using hematoxylin for 5 min and eosin (HE) for 1 min at room temperature.

Immunohistochemistry. Stainings for chromogranin A (CgA), synaptophysin (Syn), CD56, thyroid transcription factor 1 (TTF-1), P63, S-100, CK7 and Ki67 were performed by immunohistochemistry for six carcinoid tumors. The tumor tissue samples were fixed in formalin solution (10% methanol) for 48 h at room temperature. The tissues were dehydrated in xylene and graded ethanol series. After being immersed into paraffin wax twice at 60°C and embedded into paraffin blocks, the tumor tissues were cut into 5 μ m thick sections. Tissues were deparaffinized in xylene and rehydrated in a graded ethanol series. Microwave pretreatment in 5 mM Tris-HCl (pH 10.0) for 15 min was performed to facilitate heat-induced antigen retrieval. After being rinsed in phosphate buffered saline (PBS), the sections were incubated with primary antibodies against CgA (1:100; Santa Cruz Biotechnology, Inc.; 1:100; cat. no. sc-393941), Syn (Santa Cruz Biotechnology, Inc.; 1:100; cat. no. sc-17750), CD56 (Santa Cruz Biotechnology,

Inc.; 1:50; cat. no. sc-7326), TTF-1 (Santa Cruz Biotechnology, Inc.; 1:100; cat. no. sc-53136), P63 (Santa Cruz Biotechnology, Inc.; 1:50; cat. no. sc-25268), S-100 (Santa Cruz Biotechnology, Inc.; 1:100; cat. no. sc-53438), CK7 (Agilent Technologies, Inc.; 1:200; cat. no. M7018) and Ki67 (Santa Cruz Biotechnology, Inc.; 1:100; cat. no. sc-23900) at 4°C overnight. Subsequently, samples were incubated with a secondary antibody mouse IgG κ light chain binding protein (m-IgG κ BP) conjugated to horseradish peroxidase (HRP) (Santa Cruz Biotechnology, Inc.; 1:50; cat. no. sc-516102) for 30 min at room temperature. Diaminobenzidine was used for visualization and followed by hematoxylin for counterstaining at room temperature for 1 min. A light microscope was used to evaluate the staining results at x100 magnification. All staining slides were evaluated by two researchers to evaluate samples individually.

Next-generation sequencing. The DNA of 20 lung carcinoid tumors was extracted using QIAamp DNA FFPE tissue kit (Qiagen) according to the manufacturer's instructions and evaluated, and via quality control (according to the extent of DNA degradation), six cases were selected for sequencing. Targeted capture sequencing of 56 cancer-associated genes was performed in 6 pulmonary carcinoid tumors (Lung core TM 56 genes; Burning Rock Biotech; Table SI).

The concentration of the DNA samples was measured using the Qubit dsDNA assay (Invitrogen; Thermo Fisher Scientific, Inc.) to ensure that the content of genomic DNA was \geq 100 ng. The volume was adjusted to a total of 100 μ l using 1X Tris-low EDTA buffer, and the solution was transferred to a Covaris microtube for fragmentation using Covaris M220 (Covaris, Inc.) according to the manufacturer's protocol. The DNA was fragmented (average DNA fragment size, 180-220 bp), which was followed by hybridization with the capture probe baits, hybrid selection with magnetic beads and PCR amplification. A high-sensitivity DNA assay was then used to assess the quality and size range. Available indexed samples were sequenced on a NextSeq 500 (Illumina, Inc.) bioanalyzer with pair-end reads.

Raw data from the NextSeq 500 runs were processed with Flexbar software (version 2.7.0) to generate clean FASTQ data, trim adapter sequences and filter and remove poor-quality reads (16). The depth for the sequencing in the present study was \sim 1,000 and Varscan (v. 2.3) was used to call single nucleotide variations and insertions/deletions with MAPQ >60, base quality >30 and allele frequency (AF) >1% (17). The variants that comprised >3 non-duplicated paired reads or >5 non-duplicated reads were considered as true mutations. Subsequently, clean FASTQ data were aligned to the hg19 (GRCH37) assembly using BWA-sample (Burrows Wheeler Aligner software; version 0.7.12-r1039; <https://sourceforge.net/projects/bio-bwa/files/>), and PCR duplicates were removed using the Mark Duplicates tool in Picard Tools (version 1.124, <http://broadinstitute.github.io/picard/>). All variants were annotated using ANNOVAR (version 20160201) (18). Finally, variation frequency (>0.5%) was used to eliminate erroneous base calling and to generate final mutations, and manual verification was performed using Integrative Genomics Viewer version 2.3.72 (19-21).

Statistical analysis. Clinicopathological characteristics of the patients with TC and AC were compared using the unpaired Student's t-test (for mean age and tumor

Table I. Clinicopathological characteristics of patients with pulmonary carcinoid who underwent surgical resection at Tianjin Medical University General Hospital (Tianjin, China).

Characteristics	Total pulmonary carcinoid tumors (n=20)		P-value
	Typical carcinoids (n=9)	Atypical carcinoids (n=11)	
Median age (range), years	48 (28-66)	49 (14-71)	0.396
Sex, n (%)			0.069
Male	5 (55.6)	10 (90.9)	
Female	4 (44.4)	1 (9.1)	
Smoking history, n (%)			0.653
Never	5 (55.6)	5 (45.5)	
Current/former	4 (44.4)	6 (54.5)	
History of malignancy, n (%)	3 (33.3)	4 (36.4)	0.888
Median tumor diameter (range), cm	4 (1.5-9.1)	5.5 (2.1-12.5)	0.252
Incidence of PET evaluation, n (%)	4 (44.4)	3 (27.3)	0.423
Pathological N stage, n (%)			0.024
N0	9 (100)	6 (54.5)	
N1	0 (0)	3 (27.3)	
N2	0 (0)	2 (18.2)	
TNM stage, n (%)			0.872
I	4 (44.4)	5 (45.5)	
II	2 (22.2)	3 (27.3)	
III	2 (22.2)	2 (18.2)	
IV	1 (11.1)	1 (9.1)	
Tumor site			0.946
Left upper lobe	1 (11.1)	1 (9.1)	
Left lower lobe	2 (22.2)	3 (27.3)	
Left hilum	1 (11.1)	2 (18.2)	
Right upper lobe	1 (11.1)	0 (0)	
Right middle lobe	1 (11.1)	1 (9.1)	
Right lower lobe	2 (22.2)	2 (18.2)	
Right hilum	1 (11.1)	2 (18.2)	
Surgical approach, n (%)			0.492
VATS	7 (77.8)	7 (63.6)	
Thoracotomy	2 (22.2)	4 (36.4)	
Procedure, n (%)			0.493
Wedge	1 (11.1)	0 (0)	
Segmentectomy	2 (22.2)	2 (18.2)	
Lobectomy	6 (66.7)	9 (81.8)	
Adjuvant therapy, n (%)			0.659
Chemotherapy	2 (22.2)	2 (18.2)	
Radiotherapy	1 (11.1)	2 (18.2)	

PET, positron emission tomography; VATs, video-assisted thoracoscopic surgery; TNM, tumor-node-metastasis; patients were staged according to the American Joint Committee on Cancer staging manual (8th edition) criteria.

diameter), Kruskal-Wallis test [pathological N and Tumor-Node-Metastasis (TNM) staging] and χ^2 test (all other characteristics). A two-tailed $P < 0.05$ was considered to indicate a statistically significant difference. Statistical analyses were performed using SPSS 22.0 software (IBM Corp.).

Results

Clinical features of the study cohort. The clinicopathological characteristics of 20 patients who underwent surgical resection for pulmonary carcinoid tumors at Tianjin Medical University General Hospital were reviewed and summarized (Table I).

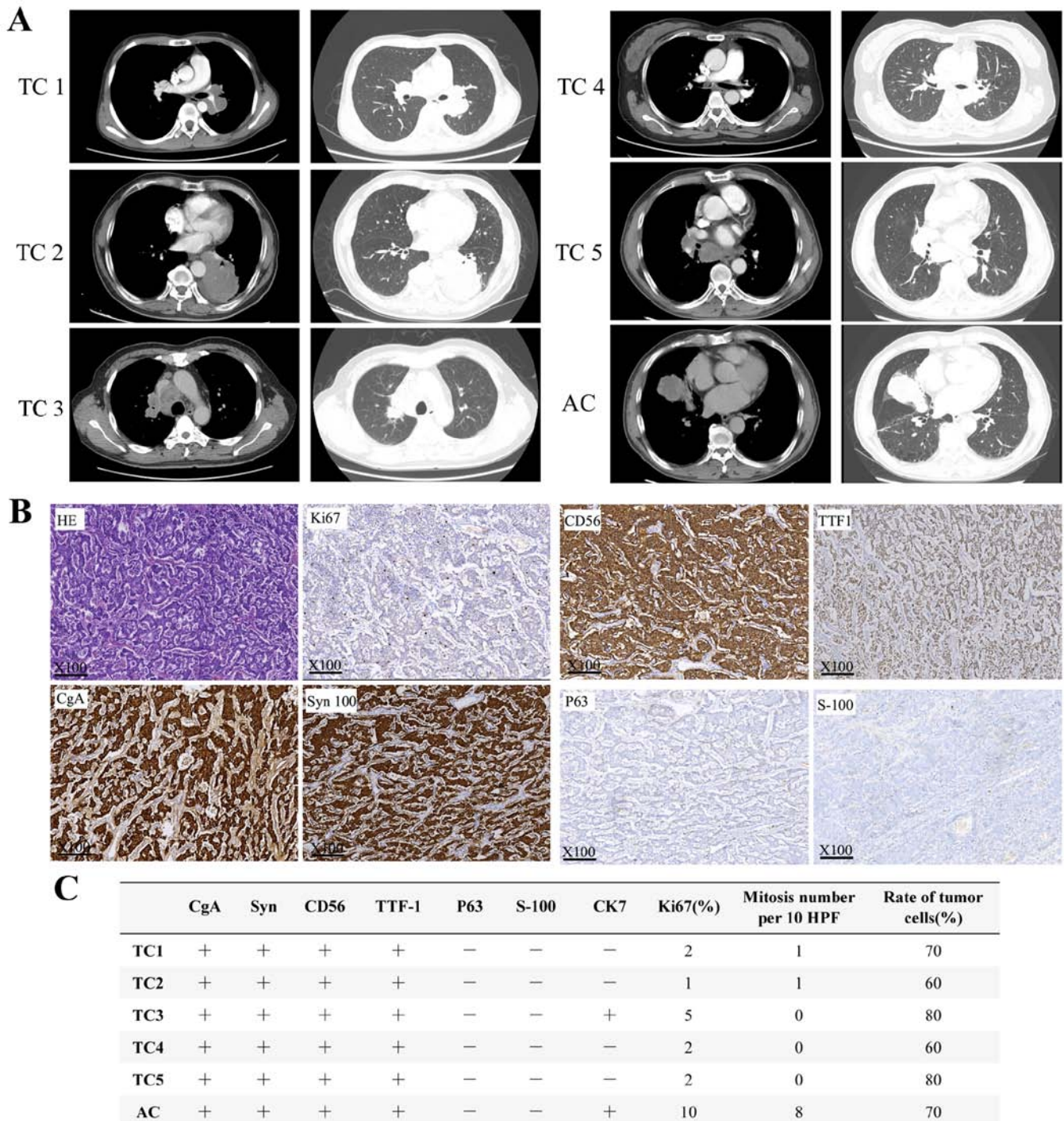


Figure 1. Radiological and pathological results of six patients with pulmonary carcinoids. (A) Computed tomography imaging of six patients with pulmonary carcinoid. (B) Representative HE and IHC images of pulmonary carcinoids under light microscope at x100 magnification. (C) IHC results of 6 patients with pulmonary carcinoids. IHC, immunohistochemistry; HE, hematoxylin and eosin; TC, typical carcinoid; AC, atypical carcinoid.

Generally, atypical carcinoids are less frequent and the ratio of TCs to ACs is 8-10:1 (4,22); however, of the 20 included cases, 9 were typical carcinoid tumors and 11 were atypical carcinoid tumors. The underlying reasons for this discrepancy are not clear. There was a male predominance in the included population (male:female, 15:5) and the age of patients ranged from 14-71 years with a median age of 48 years. None of the patients with TC tumors presented with lymphatic metastasis, whereas 5/11 (45.45%) patients with AC tumors had lymphatic metastasis, including three cases of N1 and two cases of N2

metastasis. The P-value of the Kruskal-Wallis test was 0.024, which indicated that ACs exhibited a higher malignancy stage. Other clinical characteristics, including the surgical approach, surgical procedure, prescribed adjuvant therapy, tumor sites and TNM stage were considered and compared between TC and AC, and no significant differences were observed (Table I).

Computed tomography images of six patients whose samples were submitted for NGS analysis are presented in Fig. 1A. The imaging features of pulmonary carcinoids are

Table II. Gene mutations of patients with pulmonary carcinoids from our cohort.

Case	Histology	Gene	AA change	Mutation type	Frequency (%)
1	TC	<i>JAK2</i>	K1030R	Missense variant	50.60
		<i>KIT</i>	A755T	Missense variant	50.40
		<i>RB1</i>	F198L	Missense variant	9.23
		<i>NF1</i>	S1100T	Missense variant	2.24
2	TC	<i>TSC2</i>	R57H	Missense variant	4.33
		<i>TSC1</i>	S1038R	Missense variant	3.12
		<i>TSC1</i>	S1039G	Missense variant	3.08
		<i>ERBB4</i>	R1155 ^a	Nonsense variant	2.51
		<i>NOTCH1</i>	E242K	Missense variant	2.26
		<i>KIT</i>	P37S?	Frameshift variant	2.09
		<i>ERBB4</i>	I944V	Missense variant	35.80
3	TC	<i>MAP2K1</i>	D67N	Missense variant	2.89
		<i>PDGFRA</i>	R293H	Missense variant	2.56
		<i>ERBB2</i>	R47H	Missense variant	2.29
		<i>MET</i>	R988C	Missense variant	1.68
		<i>EGFR</i>	NA	Splice donor variant	1.15
4	TC	<i>PTCH1</i>	K251T	Missense variant	49.00
		<i>IGF1R</i>	G8R	Missense variant	2.65
5	TC	<i>MET</i>	V1088M	Missense variant	41.30
		<i>KDR</i>	A532V	Missense variant	2.35
6	AC	<i>SMO</i>	P743T	Missense variant	47.50
		<i>CDK6</i>	I159K	Missense variant	7.72
		<i>IGF1R</i>	P1290L	Missense variant	4.38
		<i>FGFR1</i>	DDDD163D	Deletion variant	4.15
		<i>FGFR2</i>	L192	Deletion variant	3.56
		<i>IGF1R</i>	S1180F	Missense variant	2.63
		<i>CDK4</i>	V281E	Missense variant	2.06

TC, typical carcinoid; AC, atypical carcinoid; AA, amino acid; ^atermination codon which signals the end of translation.

often similar to those of other lung cancers and have few defining characteristics. The majority of carcinoids appear as round or ovoid peripheral lung nodules with smooth or lobular margins (23) and generally exhibit marked enhancement in enhanced CT due to their high vascularity (24). Representative images of HE and IHC staining are presented in Fig. 1B. The specific markers of the six carcinoid tumors were also summarized in Fig. 1C. The present analysis revealed that ACs exhibited a higher percentage of antigen Ki-67-positive cells and more mitoses per 10 high-power fields; and considering the diagnostic criteria of AC vs. TC, this result was logical and expected.

Gene mutation analysis of lung carcinoid tumors. The results of NGS are presented in Table II and Fig. 2. Following the gene mutation profiling of six pulmonary carcinoid tumors, a total of 27 mutations in 21 genes were identified, including *JAK2*, *KIT* proto-oncogene receptor tyrosine kinase (*KIT*), *RB* transcriptional coexpressor 1 (*Rb*), neurofibromin 1, *TSC* complex subunit 1 (*TSC1*), *TSC2*, Erb-B2 receptor tyrosine kinase 4 (*ERBB4*), *NOTCH1*, mitogen-activated protein kinase 1, platelet-derived growth factor receptor α , *ERBB2*,

MET proto-oncogene receptor tyrosine kinase (*MET*), *EGFR*, patched 1, insulin-like growth factor 1 receptor (*IGF1R*), kinase insert domain receptor, smoothed frizzled class receptor, *CDK6*, fibroblast growth factor receptor 1 (*FGFR1*), *FGFR2* and *CDK4*. Of these, 11 were proto-oncogenes and 6 were tumor suppressor genes, which indicated that they may participate in tumorigenesis, tumor growth, invasion and metastasis.

The majority of the identified mutations were missense mutations (81.48%), followed by deletion mutations (7.4%) and one case each of nonsense, frameshift and splice donor mutations (Fig. 2A). All carcinoids had multiple mutated genes, and two patients (33.3%) had multiple mutations in a single gene, including the *TSC1* and *IGF1R* genes (Fig. 2B). The *KIT*, *ERBB4*, *MET* and *IGF1R* genes were mutated in two patients (33.3%). These four genes were considered to be mutated at a high frequency (Fig. 2A) and were followed (in order of frequency) by 17 other genes that were each mutated in only one case (16.73% of cases) (Fig. 2A).

Two *KIT* mutations were identified on chromosome 4, but on different exons: Case 1 presented with a missense mutation (G>A mutation in exon 16; AF 50.4%), whereas

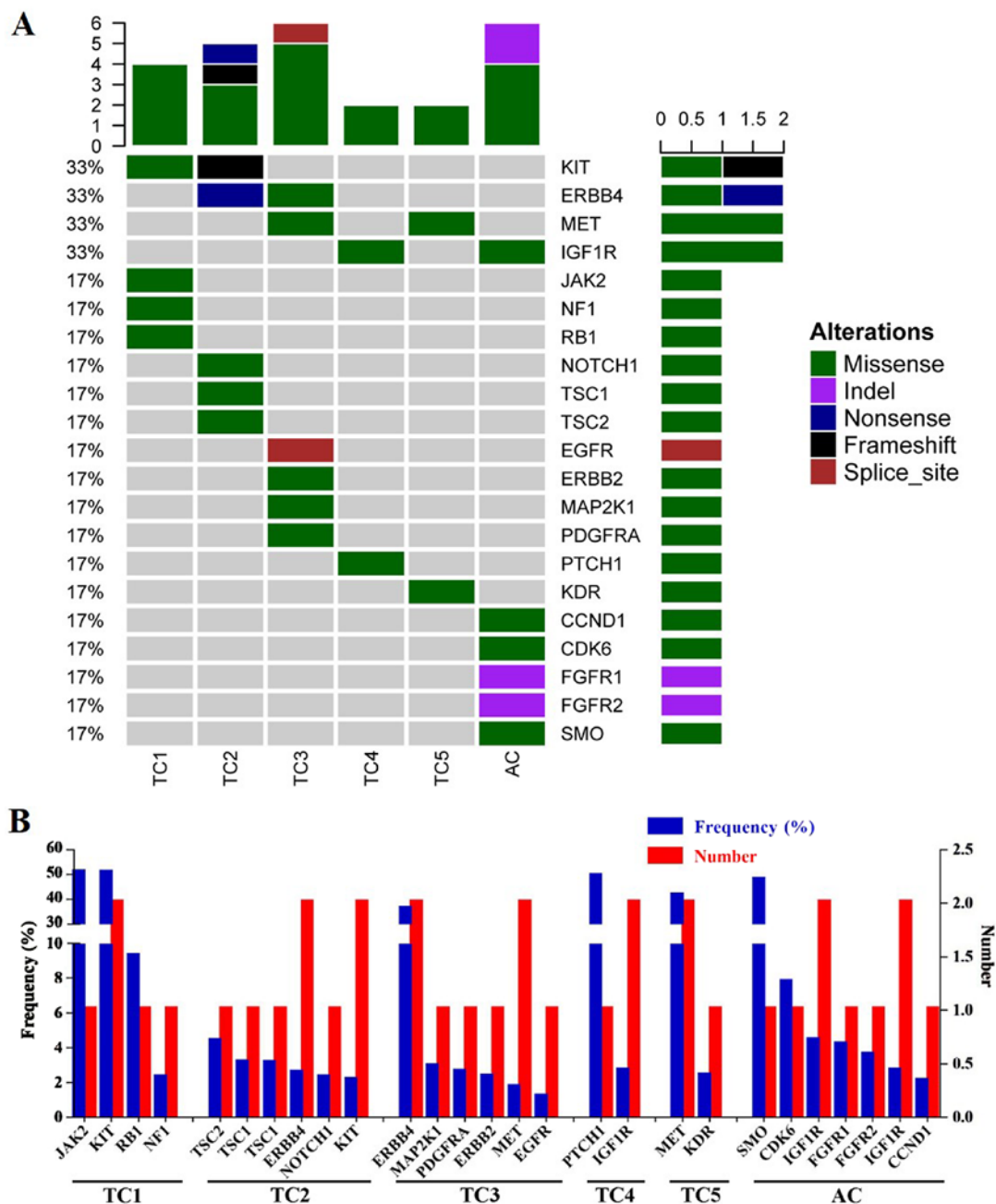


Figure 2. Gene mutation analysis results of size patients with pulmonary carcinoid. (A) Heat map of pulmonary carcinoids mutational analysis. (B) Frequency and distribution of gene mutations in six carcinoids. TC, typical carcinoid; AC, atypical carcinoid.

case 2 presented with a frameshift mutation (A>AT mutation in exon 2; AF 2.09%) (Table II). Two *ERBB4* mutations were revealed on chromosome 2. Case 2 harbored a nonsense G>A mutation in exon 27, whereas case 3 had a missense T>C mutation in exon 23, resulting in a 35.8% mutation frequency (Table II). The two *MET* mutations were both missense mutations on chromosome 7. Case 3 presented with a C>T base change in exon 14, whereas case 5 had a G>A base change in exon 15, yielding a 41.3% mutation frequency (Table II). A total of three *IGF1R* mutations were identified on chromosome 15 in two patients on different exons: Case 4 presented with a G>A mutation in exon 16, whereas case 6 presented with two C>T changes in exons 19 and 21.

Discussion

As pulmonary carcinoid is a tumor with a low malignancy rate, resection is often an effective treatment option for early disease; however, for patients with advanced unresectable pulmonary carcinoids, no standardized or authoritative postoperative adjuvant therapy scheme has been established (25,26). In recent years, as the development of precision medicine has progressed, targeted therapy has achieved significant breakthroughs for pulmonary carcinoids, an example of which is mTOR inhibitors (5-7). However, the progression of therapy in pulmonary carcinoids is still limited due to its low prevalence (26), and an in-depth understating of the underlying molecular mechanisms is necessary. Thus, large-scale clinical

Table III. Gene mutation analysis of pulmonary carcinoids from previously published literature.

Case	Author	Year	Age	Sex	Type	Mutation	Gene/Chromosome	Country	(Refs.)
1	Hiyama <i>et al</i>	1993	77	M	AC	point mutation Cys>Phe Deletion mutation	p53 Rb	Japan	(38)
2	Lohmann <i>et al</i>	1993	65	F	TC	Neutral mutation Cys>Tyr	p53	Germany	(22)
3			68	M	TC	Missense mutation Glu>Lys	p53		
4			72	F	TC	Missense mutation Val>Met	p53		
5			46	NA	TC	Frameshift mutation 1650insC	MEN1		
6	Debelenko <i>et al</i>	1997	56	NA	TC	Alteration of splicing, frameshift mutation 764+3A>G	MEN1	USA	(37)
7			63	NA	TC	Frameshift mutation 134del13 (GACGCTGTTCCCG)	MEN1		
8			49	NA	TC	Frameshift mutation 1699delA and 1702G>C	MEN1		
9	Sagawa <i>et al</i>	1998	NA	NA	AC	point mutation	K-ras	USA	(36)
10	Couce <i>et al</i>		52	F	AC	K-ras c12 Gly>Ser missense mutation	K-ras	USA	(35)
11			39	F	AC	K-ras c12 Gly>Asp missense mutation	K-ras		
12			61	F	AC	Exon 8 c298 Glu>Stop missense mutation	p53		
13			Sugio <i>et al</i>	2003	NA	NA	AC		
14			NA	NA	AC	Loss of heterozygosity in 9p	9p		
15	Snabboon <i>et al</i>	2005	68	F	TC	Deletion mutation at exon 10 (1793delG)	MEN1	Thailand	(33)
16	D'Alessandro <i>et al</i>	2010	29	F	TC	Exon 5 c.733-16C>T	ELAVL4	Italy	(32)
17			50	M	TC	Exon 5 c.666A>T Exon 5 c.712C>T	ELAVL4 ELAVL4		
18			70	F	TC	Somatic mutation Exon 4 c.424delA Exon 5 c.559G>A	ELAVL4 ELAVL4		
19			47	M	AC	Exon 4 c.387C>T Single nucleotide polymorphism Exon 5 c.687T>C c.1367+56C>T 3'UTR	ELAVL4 ELAVL4 ELAVL5		
20			54	M	AC	Somatic mutation Exon 5 c.655C>T Exon 5 c.704G>A	ELAVL4 ELAVL4		
21	Capodanno <i>et al</i>	2012	NA	NA	TC	Missense mutation c.1576 A>G	PI3K	Italy	(31)
22			NA	NA	TC	Missense mutation c.1639 G>A	PI3K		
23			NA	NA	TC	Missense mutation c.1639 G>A	PI3K		
24			NA	NA	TC	Missense mutation c.1639 G>A	PI3K		
25			NA	NA	AC	Missense mutation c.1639 G>A	PI3K		
26			NA	NA	TC	Missense mutation c.2993 T>C	PI3K		
27			NA	NA	AC	Missense mutation c.3007 T>C	PI3K		
28			NA	NA	AC	Missense mutation c.3017 T>C	PI3K		
29			NA	NA	AC	Missense mutation c.3022 T>C	PI3K		
30			NA	NA	TC	Missense mutation c.3034 G>A	PI3K		
31			NA	NA	AC	Missense mutation c.3041 A>G	PI3K		
32			NA	NA	AC	Missense mutation c.3050 A>T	PI3K		
33			NA	NA	AC	Missense mutation c.3062 A>G	PI3K		
34			NA	NA	TC	Missense mutation c.3061 T>A	PI3K		
35			NA	NA	AC	Missense mutation c.3068 G>A	PI3K		

Table III. Continued.

Case	Author	Year	Age	Sex	Type	Mutation	Gene/Chromosome	Country	(Refs.)
36			NA	NA	TC	Missense mutation c.3133 G>A	PI3K		
37			NA	NA	TC	Missense mutation c.3145 G>A	PI3K		
38			NA	NA	TC	Missense mutation c.3145 G>A	PI3K		
39			NA	NA	AC	Missense mutation c.3155 C>T	PI3K		
40	Voortman <i>et al</i>	2013	NA	NA	TC	Missense mutation Exon 14 T1010I mutation	c-Met	USA	(30)
41	Armengol <i>et al</i>	2015	69	Male	TC	Missense mutation c.1796C>T Missense mutation c.1496G>A Missense mutation c.3074C>T Missense mutation c.38G>A	BRAF SMAD4 SMAD4 KRAS	Finland	(29)
42	Vollbrecht <i>et al</i>	2015	NA	NA	AC	Missense mutation c.311T>A Missense mutation c.311T>A Insertion mutation c.2516_2517insC Deletion mutation c.1912delA Missense mutation c.1015C>T	EGFR EGFR GNAS KIT PTEN	Germany	(28)
43			NA	NA	AC	Deletion and insertion mutation c.1416_1417delinsTA	KDR		
44			NA	NA	AC	Missense mutation c.2744C>A	ERBB4		
45			NA	NA	AC	Missense mutation c.3788G>A Insertion mutation c.855_856insG Insertion mutation c.3730_3731insC	APC FGFR1 MET		
46			NA	NA	AC	Deletion and insertion mutation c.2712_2713delinsGG	RET		
47			NA	NA	AC	Deletion and insertion mutation c.2354_2355delinsGG	ERBB2		
48			NA	NA	AC	Missense mutation c.3367C>T Missense mutation c.112G>A	APC KRAS		
49			NA	NA	AC	Deletion mutation c.862delG	HNF1A		
50			NA	NA	AC	Missense mutation c.2602C>T Missense mutation c.1100T>G	ERBB2 SMO		
51			NA	NA	AC	Deletion and insertion mutation c.1637_1638delinsGG Missense mutation c.274C>T Missense mutation c.167C>T	KIT PI3K SMARCB1		
52			NA	NA	AC	Insertion mutation c.3730_3731insC	MET		
53			NA	NA	TC	Deletion and insertion mutation c.2711_2713delinsTGG	RET		
54			NA	NA	TC	Missense mutation c.3386T>C	APC		
55			NA	NA	TC	Missense mutation c.2624C>T	ERBB2		
56			NA	NA	TC	Deletion and insertion mutation c.2354_2355delinsGG	ERBB2		
57			NA	NA	TC	Missense mutation c.2531G>A	GNAS		
58			NA	NA	TC	Missense mutation c.2318A>C Missense mutation c.274T>A Missense mutation c.267A>C	EGFR IDH1 IDH1		
59			NA	NA	TC	Deletion and insertion mutation c.2471_2472delinsCT	PDGFRA		
60			NA	NA	TC	Missense mutation c.920C>T Missense mutation c.505C>T	ABL1 SMAD4		
61	Lou <i>et al</i>	2017	23	Male	NA	NA	PI3K	China	(27)

NA, not available; *Rb*, RB transcriptional corepressor 1; *MEN1*, menin 1; *ELAV4*, ELAV-like RNA-binding protein 4; PI3K, phosphatidylinositol 3-kinase, putative; NA, not applicable.

drug research targeted at pulmonary carcinoids should be proposed as soon as possible. Surgical resection is appropriate for localized diseases; these include locoregional pulmonary carcinoids, cases with limited sites of metastatic disease and local recurrent diseases, such as liver metastases (26).

Pulmonary carcinoids are low-grade malignant tumors, and their underlying molecular biological mechanism is yet to be fully elucidated. To understand previous results of pulmonary carcinoid gene sequencing, published literature (PubMed; January 2018) on mutations in pulmonary carcinoids was examined, and available clinical information was summarized in Table III, comprising 13 studies that referenced 61 cases, including 29 ACs, 31 TCs and 1 indeterminate carcinoid (22,27-38). The majority of the articles retrieved utilized first-generation sequencing technology to reveal mutations in single genes or chromosomes, including *PI3K*, *p53TP53*, *Rb*, *menin 1*, *K-ras*, *c-Met*, ELAV-like RNA-binding protein 4, *3p14* and *9p*, and no significant associations were observed between specific gene mutations and cancer type, age or sex (22,27-38). A total of three studies (including 21 patients) reported NGS data for carcinoids. The mutations of *KIT*, *ERBB4* and *MET* were also reported in these studies, which supported the findings of the present study (27-29). Notably, one study that used NGS to investigate carcinoids did not provide the original sequencing data and, consequently, the sequencing results were not summarized in Table III; however, it was reported in the study that *FGFR1* was highly expressed in carcinoids (39). In addition, Rossi *et al* (40) also reported that *ERBB4* alteration was detected in carcinoids. Recently, Asiedu *et al* (41) used mRNA expression, single nucleotide polymorphism genotyping and a combination of exome and whole-genome sequencing to detect genomic alterations in 31 TC and 11 AC tumors. Compared with the results of Asiedu *et al* (41), only a limited number of mutated genes were common to the genes identified using NGS in the present study. The differences between the current study and the previous studies may be attributable to the examination of different targeted gene panels and the different demographic of patients included. In the present study, four genes were revealed to be mutated at a high frequency, including *KIT*, *ERBB4*, *MET* and *IGF1R*, which were mutated in 33.3% patients. These genes encode typical tyrosine-protein kinases or receptor tyrosine kinases that are cell surface receptors for multiple signaling pathways and serve an essential role in the regulation of cell survival, proliferation and apoptosis (42-45). Mutations in these genes are important therapeutic targets of molecular targeted therapeutic drugs, such as the TKIs imatinib and sunitinib (42-45).

Although certain high-frequency gene mutations were identified, it is difficult to confirm whether the alteration of these genes may initiate and promote pulmonary carcinoid tumors and be effective against targeted therapy. In the future, systematic gene mutation profiling should be performed with a large number of samples to detect potential tumor-promoting genes and to identify potential novel treatment targets for pulmonary carcinoids. This profiling may have important therapeutic implications for the treatment of patients with pulmonary carcinoids.

There are certain limitations the present study; only 6 PCs were collected and this is too few to predict more precise and

comprehensive molecular principles of PCs and to conduct survival analysis.

In conclusion, *IGF1R*, *ERBB4*, *KIT* and *MET* were identified as frequently mutated genes that may influence the tumorigenesis of pulmonary carcinoid tumors; therefore, targeted therapy against these genes may represent a promising therapeutic strategy for the treatment of this rare disease.

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Availability of data and materials

The datasets used during the present study are available from the corresponding author upon reasonable request.

Authors' contributions

SX and JC conceived and designed the study. ZS, SW, GC and JC performed surgery. XL, YLH, TS and DR reviewed the patient electronic medical record for patients with pulmonary carcinoid. XL, YLH, TS and YH performed the genetic analysis. XL and YH performed the literature review and wrote the manuscript. SX and JC reviewed and edited the manuscript. All authors read and approved the manuscript and agree to be accountable for all aspects of the research in ensuring that the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Ethics approval and consent to participate

The present study was conducted in accordance with the Helsinki Declaration and was approved by the Ethics Committee of Tianjin Medical University (Tianjin, China). Written informed consent was obtained from all patients with pulmonary carcinoid for blood sampling and tissue sequencing.

Patient consent for publication

Not applicable.

Competing interests

YH is affiliated with Burning Rock Biotech, who performed targeted capture sequencing of cancer-associated genes. The other authors declare that they have no competing interests.

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