

# A Novel *MYH9-RET* Fusion Occurrence and *EGFR* T790M Loss as an Acquired Resistance Mechanism to Osimertinib in a Patient with Lung Adenocarcinoma: A Case Report

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**Background:** Osimertinib is a novel and irreversible epidermal growth factor receptor (EGFR)-tyrosine kinase inhibitor (TKI) targeting *EGFR* sensitive mutations and *EGFR* exon20 p.T790M mutation, which demonstrated superior progression-free survival (PFS) and overall survival (OS).

**Case Presentation:** We report a patient with lung adenocarcinoma harboring *EGFR* exon 19 deletion mutant treatment with icotinib. After 6 months, she developed *EGFR* exon20 p.T790M and then the patient received osimertinib treatment. A novel *MYH9* (exon41)-*RET* (exon12) fusion and *EGFR* exon20 p.T790M loss were identified using plasma circulation tumor DNA (ctDNA) after osimertinib treatment, which led to rapid progression after osimertinib five months and suggested a potential resistance mechanism.

**Conclusion:** Our findings expanded the spectrum of *RET* arrangement types and provided the basis for this hypothesis: acquired *RET* rearrangement and *EGFR* exon20 p.T790M loss potentially serve an additional resistance mechanism to osimertinib in *EGFR*-mutated non-small-cell lung cancer (NSCLC).

**Keywords:** *MYH9-RET* fusion, *EGFR* exon20 p.T790M loss, lung adenocarcinoma, acquired resistance, osimertinib

## Introduction

Approximately half of Asia-patients with NSCLC harbor *EGFR* mutation, and oral EGFR-TKIs have been used as one routine clinical therapeutic approach.<sup>1</sup> Osimertinib is approved for both EGFR-TKI sensitive and *EGFR* exon20 p.T790M mutations in NSCLC patients.<sup>2,3</sup> Previous studies described different resistance mechanisms to osimertinib, including *EGFR* tertiary mutations, like C797S,<sup>4</sup> *KRAS* mutations, and targetable gene fusions.<sup>5</sup> In all NSCLC patients, *RET* fusion accounts for 1–2%, involving various common fusion partners such as *KIF5B* and *CCDC6*.<sup>6</sup> *RET* fusions have been proved that can mediate acquired resistance to EGFR-TKIs.<sup>7,8</sup> Here, we reported a lung adenocarcinoma patient harboring a novel myosin heavy chain-9 (*MYH9*)-*RET* fusion, *EGFR* exon19 deletion (exon19del) and *EGFR* exon20 p.T790M loss and exerted resistance to osimertinib.

## Case Presentation

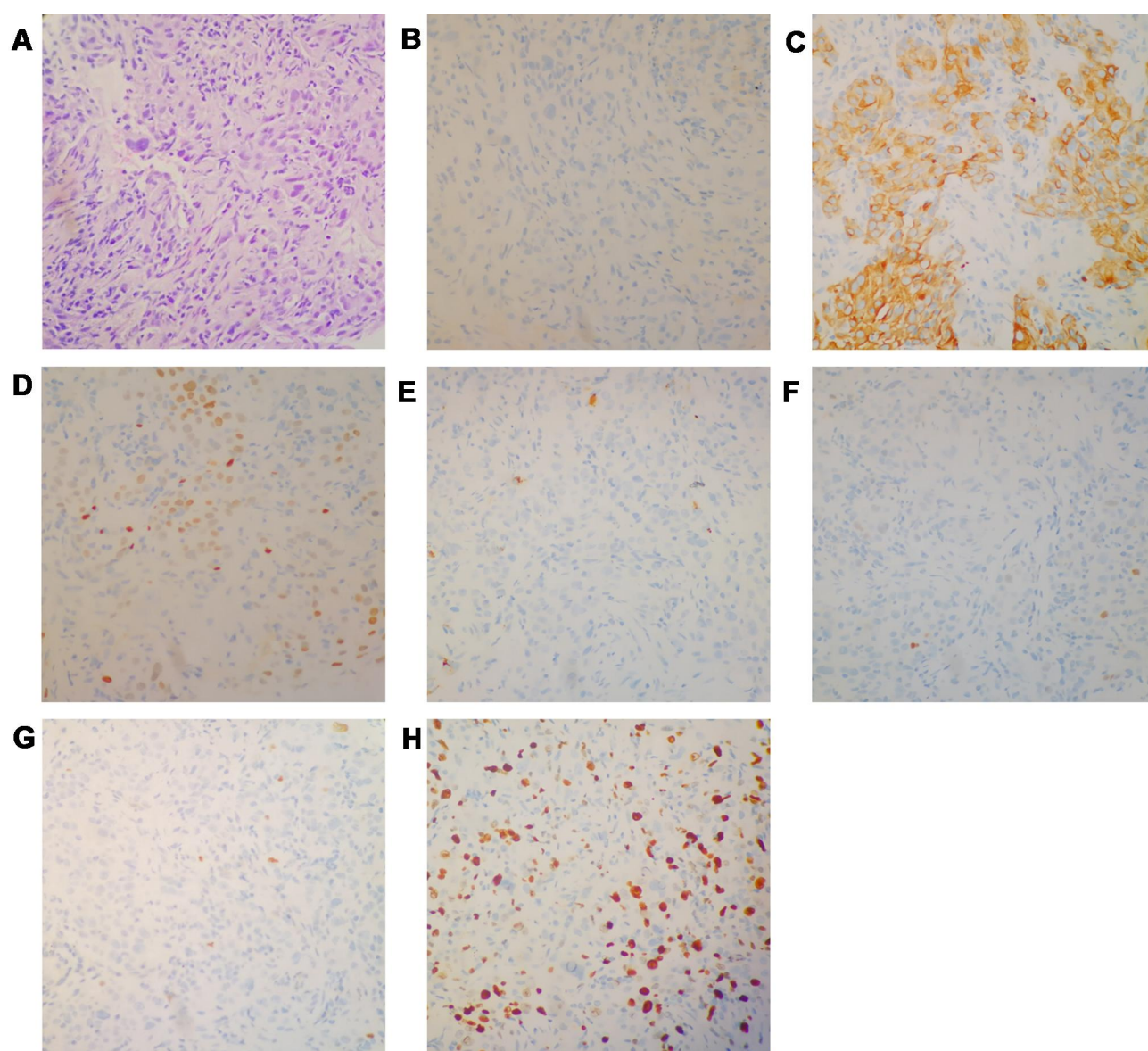
A non-smoking 48-year-old female was admitted to the hospital with intermittent cough and sputum for more than 4 months. Space occupying lesions were detected in

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the basal segment of the lower lobe of the left lung and intracranial site on admission examination. The immunohistochemistry staining results of pulmonary biopsy were CK5/6-positive, CK7-positive, thyroid transcription factor-1-positive, Napsin A-negative, P63-negative, P40-negative and Ki-67 positive index 30% (Figure 1). The disease was diagnosed as advanced lung adenocarcinoma (poorly differentiated). To seek for precision therapy, next-generation sequencing (NGS) (14-gene panel) was performed in primary lung cancer lesion. *EGFR* exon19 p. E746\_A750del and *EGFR* amplification were detected (Table 1). The patient was immediately treated with icotinib. However, after 6 months, the patient's condition deteriorated and the left lung tumor

expanded. Then the detection of *EGFR* exon20 p.T790M was performed by ddPCR in plasma ctDNA and T790M positive was found. After that, the patient received osimertinib treatment and achieved a partial response with progression-free survival (PFS) of 5 months. Subsequently, the disease progressed with the left lung tumor enlarged.

Plasma ctDNA of patients after osimertinib treatment was tested by hybrid capture NGS method using 9-gene panel (Simceredx, CAP certified Laboratory), and *EGFR* exon19 p. E746\_A750del, *EGFR* amplification as well as a novel *MYH9* (exon41)-*RET* (exon12) fusion (Figure 2) were identified, without *EGFR* exon20 p.T790M. The findings of gene sequencing during



**Figure 1** Histopathologic stains from the pulmonary biopsy. (A) hematoxylin and eosin. (B) CK5/6 (C) CK7. (D) thyroid transcription factor-1. (E) Napsin A. (F) P63. (G) P40. (H) Ki67. ( $\times 400$ ). cytokeratin (CK).

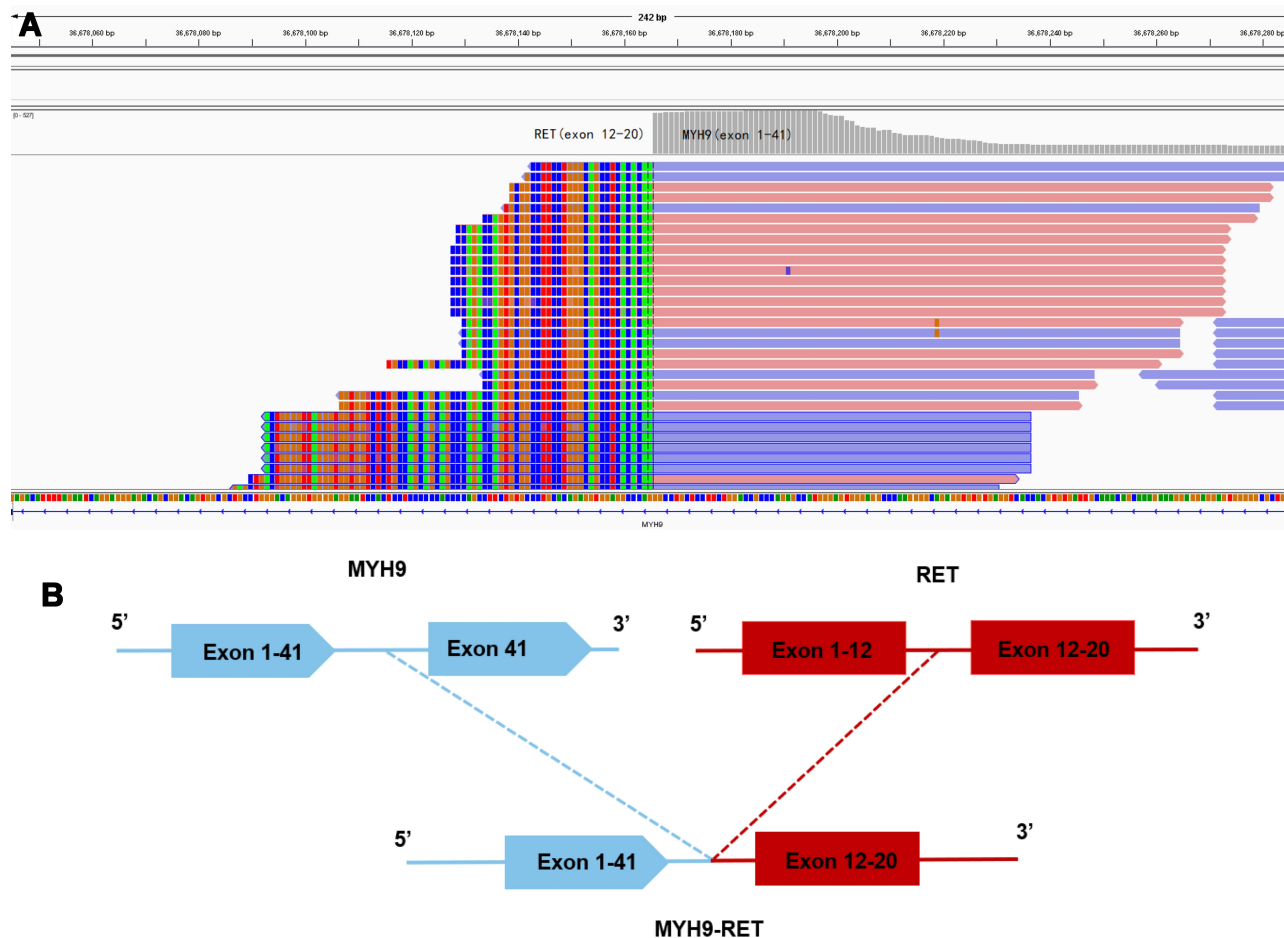
**Table 1** Findings of Gene Sequencing During Treatment

Time of Sampling	Method	Sample	Gene	Mutation Style	Frequency (%) or Copy Number
Baseline	NGS (14-gene panel)	Puncture tissue	<i>EGFR</i>	Exon 19 p. E746_A750del	46.19%
			<i>EGFR</i>	Amplification	8
After resistance to icotinib	Digital PCR (T790M)	Plasma ctDNA	<i>EGFR</i>	Exon 20 p. T790M	0.94%
After resistance to osimertinib	NGS (9-gene panel)	Plasma ctDNA	<i>EGFR</i>	Exon 19 p. E746_A750del	32.84%
			<i>EGFR</i>	Amplification	2.82
			<i>MYH9 (exon41)-RET (exon12)</i>	Fusion	1.72%

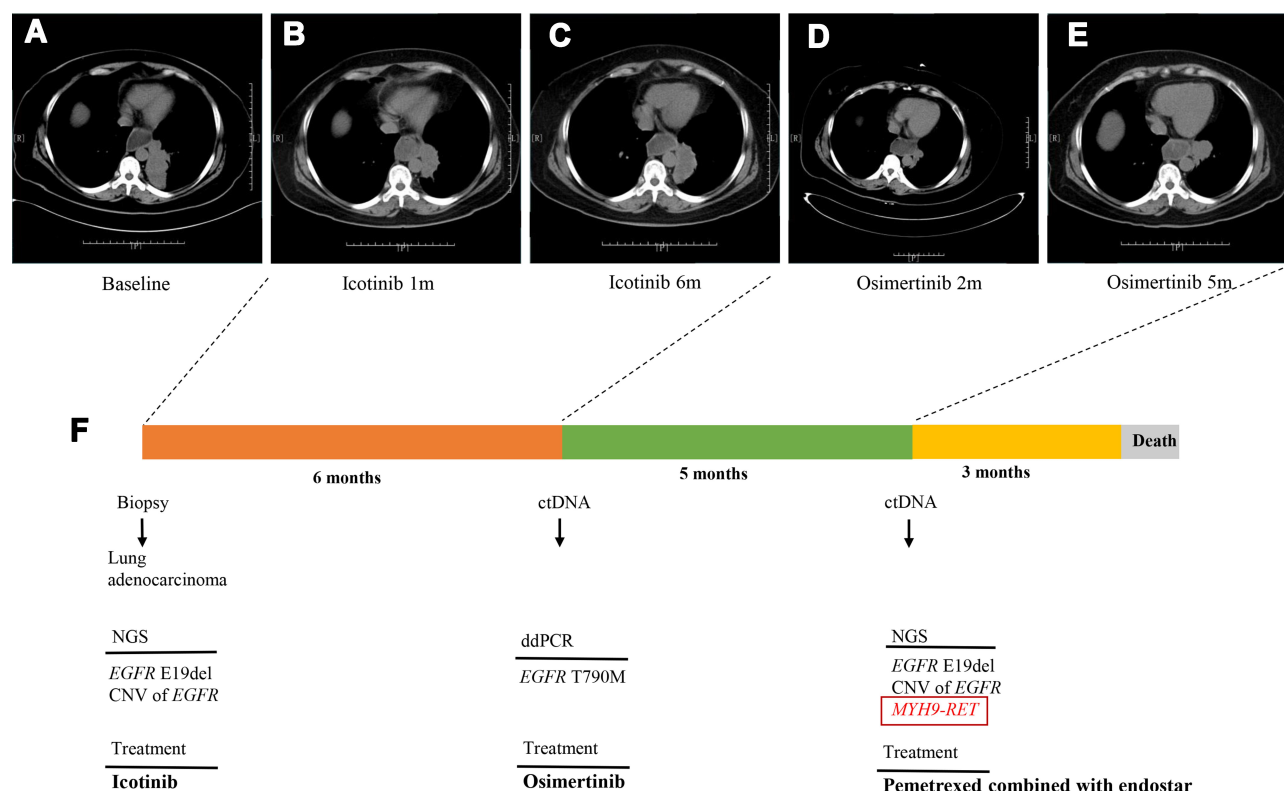
treatment were given in Table 1 and time line of the treatments and the effects of different types of therapy were presented in Figure 3.

## Discussion

In summary, this is the first case report that a novel *MYH9-RET* fusion and *EGFR* exon20 p.T790M loss occurred in



**Figure 2** Next-generation sequencing findings of *MYH9-RET* fusion. **(A)** The Integrative Genomics Viewer snapshot of *MYH9-RET*. **(B)** Schematic representation of the *MYH9-RET* fusion protein domain structure.



**Figure 3** Computed tomography (CT) scan detecting and treatment course. (A–E) Chest CT scans showed the effects of different types of therapy of the patient's lung tumor. (F) Time line of the patient accepting treatments and duration of every treatment.

lung adenocarcinoma patient who acquired resistance to osimertinib treatment. *MYH9-RET* fusion retains the complete kinase domain of *RET*. Besides, the partner gene *MYH9* contains a coiled-coil domain. *MYH9-ALK* fusion has been reported in 2003, which makes *MYH9* a plausible partner for *RET* as well.<sup>9</sup> This novel fusion is considered to promote ligand-independent dimerization and constitutive activation of *RET*. Previously reported that acquired *RET* fusions can overcome the osimertinib inhibitory effect by the activation of MAPK and PI3K signal pathway, which was sufficient to cause EGFR TKI resistance.<sup>10,11</sup> The NSCLC patients harboring *RET* fusion may benefit from new highly selective *RET* inhibitors, such as BLU-667 and LOXO-292. Previously reported that *RET* fusion coexisted with *EGFR* sensitive mutation in *EGFR*-mutated NSCLC patients who had progressed after first- or second-generation *EGFR* TKI treatment.<sup>8</sup>

*EGFR* exon 20 p.T790M loss has been reported in some osimertinib resistant cases.<sup>12–14</sup> Genomic heterogeneity and clonal evolution including T790M-positive clones loss and selection of pre-existing resistance clones may contribute to the emergence of drug resistance. *EGFR* exon20 p.T790M was detected by NGS assay using

ctDNA rather than tissue sample after osimertinib resistance. However, it might be due to the low sensitivity of the ctDNA assay. This is one limitation of this case. But *RET* fusions as mechanisms of resistance to osimertinib have so far always been accompanied by T790M loss in *EGFR*-mutant Lung Cancers.<sup>7,10</sup>

Because of the patient achieving the partial response and the progression-free survival (PFS) of osimertinib treatment was merely 5 months, and no other resistance mechanisms identified, we considered that this novel *MYH9-RET* fusion acquired and T790M loss was an acquired resistance mechanism. One study has demonstrated that osimertinib and BLU-667 (one of *RET* inhibitors) might be a well tolerated and effective approach for *EGFR*-mutated NSCLC acquired *RET* fusion.<sup>7</sup> However, the patient did not have a chance to receive this combination therapy and died 3 months later. Moreover, the patient did not undergo a biopsy at disease progression after osimertinib, so we did not confirm whether there was a potential histological transformation or not.

As a routine genetic testing approach, NGS is essential in identifying mutations that mediate primary or acquired resistance during treatment. Our case report expanded the



spectrum of *RET* arrangement types and provided the basis for this hypothesis: acquired *RET* rearrangement and T790M loss can potentially serve an additional resistance mechanism to osimertinib in *EGFR*-mutated NSCLC.

## Ethics Approval and Consent for Publication

A written informed consent was obtained from the patient's family for publication of the case details and any accompanying images and the research was approved by the Institutional Ethics Review Board of People's hospital of Rizhao.

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## Disclosure

The authors have declared no conflicts of interest.

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