

Department of Pathology and Laboratory Medicine No.201, Sec. 2, Shipai Rd., Beitou District, Taipei City, Taiwan 11217, R.O.C. Tel: 02-2875-7449

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### **Sample Information**

Patient Name: 藍吳麗鈴 Gender: Female ID No.: A224698738 History No.: 18826534

**Age:** 59

Ordering Doctor: DOC3109L 邱昭華

Ordering REQ.: C22P5EF Signing in Date: 2021/02/24

**Path No.:** S110-98262 **MP No.:** F21017

Assay: Oncomine Focus Assay

Sample Type: FFPE Block No.: S108-08399A Percentage of tumor cells: 80%

Note:

### Sample Cancer Type: Non-Small Cell Lung Cancer

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### **Report Highlights**

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### **Relevant Non-Small Cell Lung Cancer Variants**

Gene	Finding	Gene	Finding
ALK	Not detected	NTRK1	Not detected
BRAF	Not detected	NTRK2	Not detected
EGFR	Not detected	NTRK3	Not detected
ERBB2	Not detected	RET	Not detected
KRAS	Not detected	ROS1	Not detected
MET	Not detected		

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#### **Relevant Biomarkers**

Tier	Genomic Alteration	Relevant Therapies (In this cancer type)	Relevant Therapies (In other cancer type)	Clinical Trials
IIC	ERBB3 p.(V104L) c.310G>C	None	None	4
	erb-b2 receptor tyrosine kinase 3 Allele Frequency: 42.54%			

Public data sources included in relevant therapies: FDA1, NCCN, EMA2, ESMO

**Tier Reference:** Li et al. Standards and Guidelines for the Interpretation and Reporting of Sequence Variants in Cancer: A Joint Consensus Recommendation of the Association for Molecular Pathology, American Society of Clinical Oncology, and College of American Pathologists. J Mol Diagn. 2017 Jan;19(1):4-23.

### Variants (Exclude variant in Taiwan BioBank with >1% allele frequency)

DNA	DNA Sequence variants							
Gene	Amino Acid Change	Coding	Variant ID	Locus	Allele Frequency	Transcript	Variant Effect	Coverage
ERBB3	p.(V104L)	c.310G>C	COSM160824	chr12:56478854	42.54%	NM_001982.3	missense	1998

### **Biomarker Descriptions**

#### ERBB3 (erb-b2 receptor tyrosine kinase 3)

Background: The ERBB3 gene encodes the erb-b2 receptor tyrosine kinase 3, a member of the human epidermal growth factor receptor (HER) family. Along with ERBB3/HER3, EGFR/ERBB1/HER1, ERBB2/HER2, and ERBB4/HER4 make up the HER protein family¹. ERBB3/HER3 binds to extracellular factors, such as neuregulins, but has an impaired kinase domain². Upon ligand binding, ERBB3 forms hetero-dimers with other ERBB/HER family members, including ERBB2/HER2 resulting in activation of tyrosine kinase activity primarily through its dimerization partner.

Alterations and prevalence: ERBB3 gene amplification leading to an increase in expression occurs at low frequency (1-5%) in several cancer types including bladder, esophagus, lung adenocarcinoma, ovarian, pancreas, sarcoma, stomach, and uterine cancers<sup>3,4,5,6,7,8,9</sup>. ERBB3 is also the target of relatively frequent (5-10%) and recurrent somatic mutations in diverse cancer types including bladder, cervical, colorectal, and stomach cancers<sup>3,6,8,9,10</sup>. Recurrent ERBB3 mutations such as V104L/M, occur primarily in the extracellular domain.

Potential relevance: Currently, no therapies are approved for ERBB3 aberrations. Overexpression and activation of ERBB3/HER3 is one mechanism of acquired resistance to therapies targeting EGFR and ERBB2/HER2<sup>11,12</sup>. Preclinical and translational research studies have characterized the oncogenic potential of recurrent ERBB3 mutations and their sensitivity to anti-ERBB antibodies and small molecule inhibitors<sup>13,14,15,16</sup>. A phase I study exhibited progression-free survival (PFS) of 2.5 months and overall survival (OS) of 9 months in 25 patients with ERBB3 mutations treated by anti-ERBB antibodies or molecular-targeted agents<sup>17</sup>.

## **Relevant Therapy Summary**

In this cancer type	O In other cancer type	In this cancer type and other cancer types			X No eviden	ce
ERBB3 p.(V104l	L) c.310G>C					
Relevant Therapy		FDA	NCCN	EMA	ESMO	Clinical Trials*
afatinib		×	×	×	×	<b>(II)</b>
BDTX-189		×	×	×	×	<b>(</b> 1/11)
neratinib, palbociclib,	everolimus, trametinib	×	×	×	×	<b>(</b> I)

<sup>\*</sup> Most advanced phase (IV, III, II/III, II, I/II, I) is shown and multiple clinical trials may be available.

# **Relevant Therapy Summary (continued)**

#### ERBB3 p.(V104L) c.310G>C (continued)

Relevant Therapy	FDA	NCCN	EMA	ESMO	Clinical Trials*
pirotinib	×	×	×	×	<b>(</b> I)

<sup>\*</sup> Most advanced phase (IV, III, II/III, II, I/II, I) is shown and multiple clinical trials may be available.

### **Clinical Trials Summary**

### ERBB3 p.(V104L) c.310G>C

NCT ID	Title	Phase
NCT03810872	An Open Explorative Phase II, Open Label Study of Afatinib in the Treatment of Advanced Cancer Carrying an EGFR, a HER2 or a HER3 Mutation	II
NCT04209465	MasterKey-01: A Phase I/II, Open-label, Two-part, Multicenter Study to Assess the Safety, Tolerability, Pharmacokinetics & Antitumor Activity of BDTX-189, an Inhibitor of Allosteric ErbB Mutations, in Patients w/ Advanced Solid Malignancies	1/11
NCT03065387	Phase I Study of the Pan-ERBB Inhibitor Neratinib Given in Combination With Everolimus, Palbociclib, or Trametinib in Advanced Cancer Subjects With EGFR Mutation/Amplification, HER2 Mutation/Amplification, or HER3/4 Mutation or KRAS Mutation	I
No NCT ID	Phase I Clinical Study With Advanced Solid Tumors KBP-5209 Treatment	I

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# **Signatures**

**Testing Personnel:** 

**Laboratory Supervisor:** 

Pathologist:

#### References

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