

BIOMARKER TESTING: A Roadmap to Personalized Treatment in Lung Cancer



BIOMARKER TESTING IS CRITICAL IN MANAGING LUNG CANCER



Comprehensive biomarker testing at diagnosis is critical because it can help doctors and patients develop a targeted and personalized treatment plan to help improve patient outcomes.^{1,2}



~50% of patients with lung cancer (NSCLC) have at least one recognized driver mutation that initiates cancer and maintains its growth.³

Targeted therapy, enabled by biomarker testing, is associated with an improved outcome.

↓ **31%** reduction in risk of death¹

↑ **~50%** improved survival*

*Median overall survival (mOS) of 3.5 months in patients receiving targeted therapy enabled by biomarker testing compared to 2.4 months mOS in patients with a driver mutation not receiving targeted therapy.¹

What is Biomarker Testing?

A biomarker is a measurable indicator of a patient's disease.⁴



Biomarker testing, also called molecular testing, is testing of blood, body fluids, or tissue to identify a patient's specific tumor makeup, which may predict response to therapy or provide insight on risk of cancer progression.⁴



Biomarkers can include driver mutations that can help doctors understand what may be causing the cancer. Some biomarkers have FDA-approved therapies and some biomarkers have therapies that are still being developed and investigated in clinical trials.



BIOMARKERS IN LUNG CANCER

Driver mutations observed in lung cancer patients include **EGFR, ALK, MET, ROS1, BRAF, RET, NTRK1, KRAS, and HER2.**

Nearly half of all **KRAS** mutations in NSCLC are **KRAS G12C**, one of the most prevalent driver mutations in NSCLC.^{5,6,7,8,9}

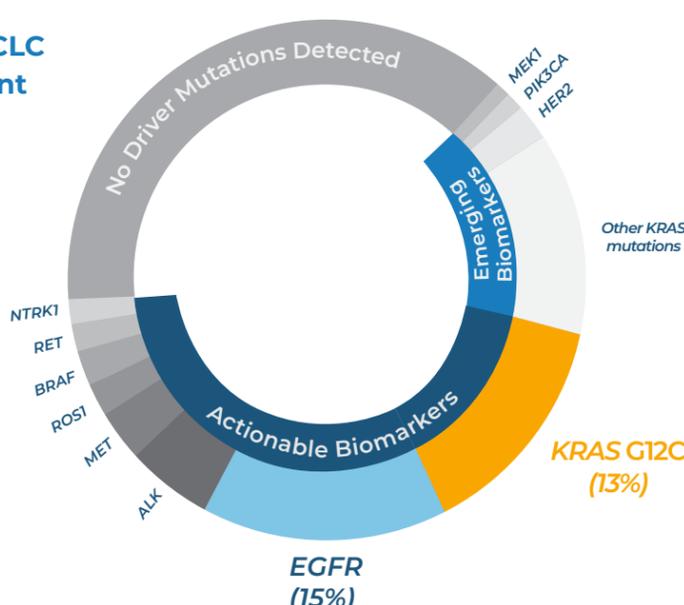
KRAS G12C occurs in

~13%

of patients with NSCLC in the U.S., comparable to the prevalence of EGFR mutations¹⁰

➔ **That's 1 in 8 NSCLC patients**¹⁰

Prevalence of Specific Genetic Mutations in Lung Adenocarcinoma



CLINICAL GUIDELINES CALL FOR COMPREHENSIVE TESTING OF PATIENTS WITH ADVANCED NSCLC REGARDLESS OF AGE, RACE OR SMOKING HISTORY^{11,12}

Steps for Biomarker Testing

- 1** Biomarker testing detects DNA from tumors through either tissue or liquid biopsy samples.
- 2** Sample is sent to a lab for analysis and a report is sent back to the doctor.
- 3** Results are discussed by doctor and patient to guide treatment path, including whether targeted therapy is appropriate.



Professional medical organizations recommend **testing for actionable and emerging biomarkers at the time of diagnosis for patients with advanced NSCLC**; however, testing in community oncology practices remains sub-optimal.¹¹⁻¹³

Many patients with lung cancer are not tested. A study showed biomarker testing rates as low as **22%***

*Biomarker testing rates among five U.S. community oncology practices was 22% for all four of the guideline recommended biomarkers between 2017 and 2019.¹³



AMGEN IS COMMITTED TO MAKING BIOMARKER TESTING MORE ACCESSIBLE TO PATIENTS

Amgen's Biomarker Assist™ is a program to help more patients with advanced NSCLC gain access to biomarker testing. Through the program, **eligible patients may save on biomarker testing.** Visit www.BiomarkerAssist.com or contact 1-888-4ASSIST to learn more.



Patient resources and tools about biomarker testing in lung cancer, can be found at:



To learn more about KRAS G12C, visit www.FindKRASG12C.com

References

1. Kris MG, et al. *JAMA*. 2014;311:1998-2006.
2. Barlesi F, et al. *Lancet*. 2016;387:1415-1426.
3. Baumgart M. *Am J Hematol Oncol*. 2015;11:10-13.
4. Goosens N, et al. *Transl Cancer Res*. 2015;4:256-269.
5. Pakkala S, et al. *JCI Insight*. 2018;e120858.
6. Arbour KC, et al. *Clin Cancer Res*. 2018;24:334-340.
7. Cox AD, et al. *Nat Rev Drug Discov*. 2014;13:828-851.
8. Biernacka A, et al. *Cancer Genet*. 2016;209:195-198.
9. Villalobos P, et al. *Hematol Oncol Clin North Am*. 2017; 31:13-29.
10. Amgen Data on File: Analysis of AACR Genie v8,7-A-Table.
11. Gutierrez ME, et al. *Clin Lung Cancer*. 2017;18:651-659.
12. Pennell NA, et al. *Am Soc Clin Oncol Educ Book*. 2019;39:531-542.
13. Gierman HJ, et al. *J Clin Oncol*. 2019;37(15_Suppl):Abstract 1585.