

# IDYLLA<sup>™</sup> FIRST

# Guiding First-Line Therapy Decisions in NSCLC



### Availability of biomarker results prior to first-line therapy is key to enhance patient outcomes

Although lung cancer remains the leading cause of cancer deaths worldwide, the survival rate has increased over the past few years due to the rapidly evolving treatment landscape for Non-Small Cell Lung Cancer (NSCLC).

Since new therapies are becoming increasingly biomarker-driven, an urgent and growing need for biomarker testing has emerged. Molecular testing therefore plays a key role in the diagnostic work-up for early and late-stage NSCLC patients to guide therapy decisions in neo-adjuvant, adjuvant and metastatic settings.

The accurate & fast diagnosis of these actionable biomarkers is critical for the right targeted therapy. Consequently, international treatment guidelines (ESMO, NCCN) recommend molecular testing for EGFR, BRAF, ALK, ROS1, RET, METex14 skipping, NTRK, HER2 and *KRAS G12C* in every patient diagnosed with advanced/metastatic NSCLC.



### Time is of the essence in NSCLC

Today, the adoption rate of biomarker testing is far from ideal, resulting in suboptimal therapy decisions for many patients. One of the identified causes is the long turnaround time to receive biomarker results, in contrast with the critical need to start therapy immediately.

It has been demonstrated that patients, for whom biomarker results were available prior to 1L therapy, have a better overall survival as compared to patients for whom biomarker results were not available.

**Overall Survival by Availability of Molecular** Genotyping Results Prior to 1L Therapy<sup>2</sup>



## Why IDYLLA<sup>™</sup> FIRST?

- Idylla™ helps save lives. Patients die, rapidly deteriorate or receive sub-optimal treatment while waiting for their biomarker test results.5-7
- Idylla<sup>™</sup> alleviates patient anxiety and allows them to focus on their well-being.<sup>8,9</sup>
- Idylla<sup>™</sup> right first time. Idylla<sup>™</sup> shows low failure rates combined with the ability to produce accurate results.<sup>4,10,11</sup>
- Idylla™ only requires small sample sizes. An NGS workflow needs nearly five times more tissue to meet NSCLC biomarker testing requirements.<sup>5</sup>



# Idylla<sup>™</sup> - Actionable biomarker testing within 3 hours

Idylla™ solid biopsy assays focus on clinically relevant biomarkers according to international clinical guidelines of ASCO\_ESMO\_NCCN and IASLC

EGFR (IVD)	exons 18 - 21
GeneFusion (IVD)	ALK, ROS1, RET, METex14

#### >90% of patients with first-line actionable biomarkers can start the right targeted therapy immediately<sup>3</sup> **IDYLLA<sup>™</sup> FIRST** EGFR GeneFusion ALK, ROS1, RET and METex14 NGS + 12% Ŧ 15% For low prevalent mutations Lung Biopsy or later lines of therapy. >90% of patients with first-line actionable biomarkers can start the right targeted therapy immediately<sup>3</sup> NGS results available Actionable results within 3 hours 2 weeks later than Idylla<sup>™4</sup>



Idylla<sup>™</sup> is a fully automated, easy to implement molecular testing platform suitable for any hospital lab setting and enabling rapid in-house biomarker testing.

# Renowned institutions already implemented IDYLLA<sup>™</sup> FIRST

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Idylla™ meets the need for assessing EGFR on surgical specimens and enables better and rapid decision-making.

Prof. Dr. Nicolas Girard, Oncology Pneumology, Institut Curie, France

### "

Our experience is fantastic. Idylla™ allows us to have an ultra-fast and reliable panel of biomarkers useful in therapeutic decision-making for the main molecular targets in lung cancer.

Dr. Antonio Calles, Medical Oncologist and KOL in lung cancer, Hospital Universitario Gregorio Marañon, Spain

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