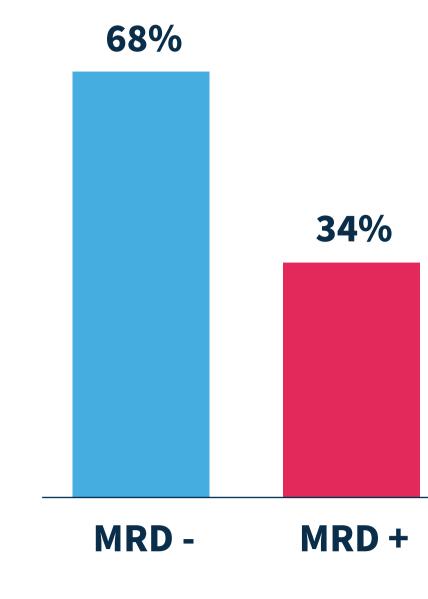
# Measurable Residual Disease (MRD)

Get the complete MRD story with targeted NGS profiling

#### Complete response may not provide a complete picture

Detection of residual disease is a powerful prognostic factor, an important instrument for therapeutic decision-making, and is the best way to monitor therapeutic response and predict relapse.1





#### Expediting development of new treatment strategies with MRD

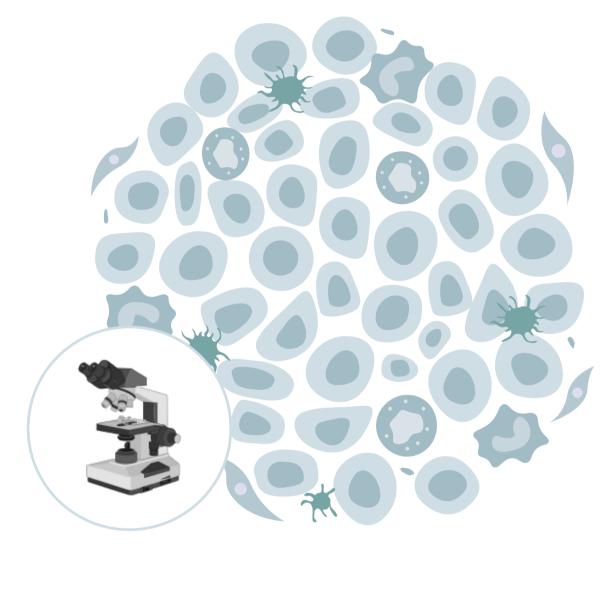


Current U.S. Clinical Trials using MRD<sup>3</sup>

Patients labeled as MRD-negative by standard tests still face relapse rates of

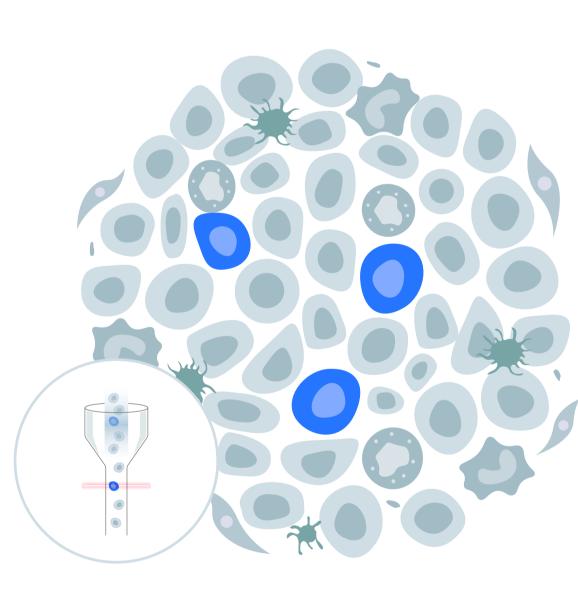


Next-generation sequencing (NGS) MRD assays provide a comprehensive analysis of AML-specific genes in a single assay<sup>1</sup>



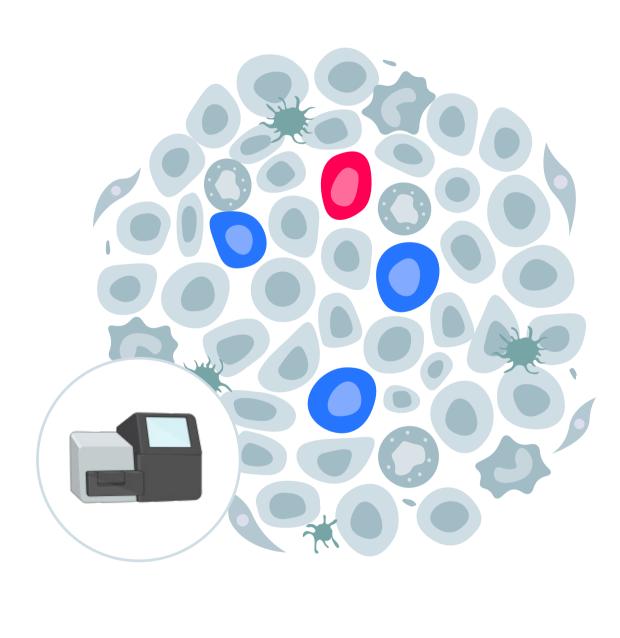
#### Morphology

Only a limited number of cells evaluated. Likely to detect only high levels of residual disease.



#### **Flow Cytometry and PCR**

Limited in the number of cancer cell markers evaluated. Subclonal populations may be missed.

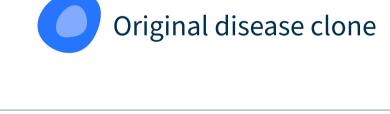


#### **Next-Generation Sequencing**

Can detect and quantify mutations across multiple AML-specific genes simultaneously, revealing emerging clonal populations and simplifying monitoring.



Healthy cell



Emerging sub-clone

Stay one step ahead of AML with NGS-based MRD longitudinal monitoring<sup>1,5</sup>

- Detect extremely low levels of disease
- Identify emerging clones Personalize treatment strategies
- Monitor for relapse

## Discover MRD solutions with enhanced sensitivity.

**Learn more** 

#### References: <sup>1</sup> Li, W. *Leukemia*. Exon Publications. 2022; 79-100.

- <sup>2</sup> Aitken, M.J.L., Ravandi, F., Patel, K.P., Short, N.J., *J Hematol Oncol*. 2021; 14, 137. <sup>3</sup> ClinicalTrials.gov [Accesssed 4/5/2024]
- <sup>4</sup> Short, N.J., Zhou, S., Fu, C. et al. JAMA. 2020; 6(12): 1890-1899. <sup>5</sup> Heuser, M., Freeman, S.D., Ossenkoppele, G.J. et al. *Blood*. 2021;138(26): 2753-2767.

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