

Hemato-oncology

Oncomine Myeloid Assay GX v2

More rapid, automated, and complete myeloid genomic profiling

Myeloid malignancy samples can be challenging to analyze. They are complex, heterogeneous, and have the potential to proliferate rapidly. To get meaningful genetic insights, you need a rapid and streamlined approach for profiling all key mutations.

Traditional single-analyte testing approaches can be laborious and time-consuming—especially as the list of relevant genes continues to grow. With the Ion Torrent™ Oncomine™ Myeloid Assay GX v2 on the Ion Torrent™ Genexus™ System, you can get a comprehensive profile of myeloid mutations from a single next-generation sequencing (NGS) run and results in as little as one day.

A highly automated workflow lets you go from specimen to report with only 20 minutes of hands-on time. Integrated bioinformatic tools allow you to seamlessly generate a variant report with annotations based on the latest evidence.

Target important biomarkers associated with major myeloid disorders

With the ability to profile both DNA and RNA targets in a single test, the Oncomine Myeloid Assay GX v2 allows you to dramatically consolidate the number of individual tests typically required for profiling a broad spectrum of mutations, including both DNA mutations and translocations detected from RNA targets.

Simultaneously interrogate 45 DNA target genes and 35 RNA fusion driver genes. This broad fusion panel allows you to sequence over 700 unique fusion transcripts. Gene content on the panel has been curated to cover relevant targets for all the major myeloid disorders—AML, MDS, MPN, CML, CMML, and JMML.

Highlights



Rapid turnaround time—from specimen to report in as little as one day



Highly automated workflow—process samples with just 20 minutes of hands-on time required



Comprehensive target coverage—simultaneously profile 45 key DNA genes and 35 fusion drivers (>700 unique fusions) relevant for the spectrum of major myeloid disorders



Trusted performance—reliably detect a range of variants, including challenging single nucleotide variants (SNVs) in homopolymer regions and important internal tandem duplications (ITDs) and partial tandem duplications (PTDs)



Integrated reporting—get annotated variants, including links to the latest relevant evidence, to help inform biological significance

Trusted performance for detecting key mutations

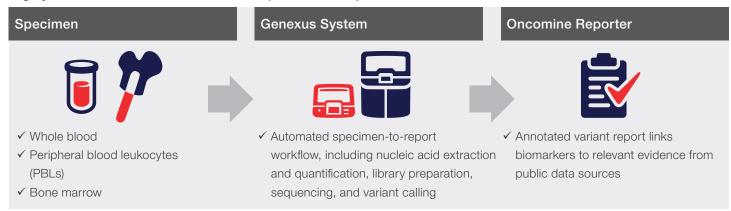
Myeloid samples frequently harbor mutations in difficult-to-sequence regions of the genome. Our assay primer design and chemistry has been carefully engineered to:

- Reliably detect all important mutations associated with myeloid malignancies
- Accurately detect SNVs in genes with long homopolymer regions like CEBPA and ASXL1
- Detect a range of insertions and deletions, even in challenging genes like CALR
- Identify ITDs in FLT3 and PTDs in KMT2A
- Get optimal performance for challenging insertions with integrated FLT3-ITD detection software

Table 1. Oncomine Myeloid Assay GX v2 gene targets.

DNA panel: hotspot genes (28)			DNA panel: full genes (17)		ıl: ver genes (35)		RNA panel: expression genes (5)	RNA panel: expression control genes (5)
ANKRD26 ABL1 BRAF CBL CSF3R DDX41 DNMT3A FLT3 (ITD, TKD) GATA2 HRAS IDH1 IDH2 JAK2 KIT	KRAS MPL MYD88 NPM1 NRAS PPM1D PTPN11 SMC1A SMC3 SETBP1 SF3B1 SRSF2 U2AF1 WT1	ASXL1 BCOR CALR CEBPA ETV6 EZH2 IKZF1 NF1 PHF6	PRPF8 RB1 RUNX1 SH2B3 STAG2 TET2 TP53 ZRSR2	ABL1 ABL2 BCL2 BRAF CCND1 CREBBP EGFR ETV6 FGFR1 FGFR2 FUS	HMGA2 JAK2 KAT6A (MOZ) KAT6B KMT2A KMT2A PTDS MECOM MET MLLT10 MRTFA (MKL1) MYBL1 MYH11 NTRK2 NTRK3	NUP98 NUP214 PAX5 PDGFRA PDGFRB RARA RUNX1 TCF3 TFE3 ZNF384	BAALC MECOM MYC SMC1A WT1	EIF2B1 FBXW2 PSMB2 PUM1 TRIM27

Highly automated NGS workflow-from specimen to report



Start with any common myeloid specimen type. From there, the Genexus System integrates and automates the entire workflow. With only 20 minutes of hands-on time required, your lab can be more efficient and free up staff's time to focus on other applications.

Each lane on the Ion Torrent™ GX5™ Chip can accommodate up to eight samples in a sequencing run. The four-lane chip and reagents are stable on the instrument for two weeks, giving you the flexibility to batch samples across multiple runs without having to waste unused sequencing capacity.

Onboard, integrated analysis provides robust variant calling and reporting without the need for an external server. The simplified user experience helps minimize the learning curve and avoid human error. The Ion Torrent™ Oncomine™ Reporter is a curated knowledge-base and reporting software that links variants to relevant evidence and enables custom reporting. These tools help simplify the bioinformatics workflow and enable you to focus on finding the biological meaning of your data.



Specifications

Gene targets	28 hotspot genes (DNA), 17 full genes (DNA), 35 fusions drivers (RNA), >700 unique fusions, 5 expression genes, 5 expression controls			
Variants	SNVs, insertions, deletions, ITDs, PTDs, gene fusions			
Input	20 ng DNA, 10 ng RNA			
Turnaround time	22 hours for 8 samples (automated library prep, sequencing, reporting)			
	27 hours for 8 samples (automated extraction, purification, library prep, sequencing, reporting)			
Throughput	8 DNA and 8 RNA samples per lane, per run			
Throughput	32 DNA and RNA samples per chip			
Coverage	≥99%			
Sensitivity	≥95%			

We offer different assay configurations to meet your lab's unique needs. Profile DNA and RNA targets simultaneously or have the option to profile each independently.

Ordering information

Product	Cat. No.
Oncomine Myeloid Assay GX v2	A50694
Oncomine Myeloid DNA Assay GX v2	A50753
Oncomine Myeloid RNA Assay GX v2	A50754