

# Oncomine BCR IGH assays for hematology-oncology research

Simple clonality assessment, sensitive rare clone detection

B cell leukemias and lymphomas originate from the malignant transformation and clonal proliferation of one or a small number of B cells. Since each B cell expresses distinct B cell receptors (BCRs) on its surface, potential malignant clones of interest can be identified and measured by the unique BCR sequences from the original neoplastic cell.

Today, high-throughput sequencing of the BCR repertoire is increasingly being adopted to advance hematology-oncology research. Next-generation sequencing (NGS)

offers significant advantages over other more traditional approaches by offering ultrahigh sensitivity, lower limits of detection (LOD), and greater flexibility to multiplex.

The **Ion Torrent™ Oncomine™ BCR IGH-LR** and **Oncomine™ BCR IGH-SR assays** are a pair of robust and sensitive NGS-based assays that make it easier for hematology-oncology labs to assess B cell clonality, measure somatic hypermutation (SHM), and detect rare B cell clones of interest for measurable residual disease (MRD) research.



## Clonality assessment

Confidently identify the dominant malignant clone, measure clonal expansion, and determine its unique CDR3 sequence



## SHM analysis

Accurately quantify the frequency of SHM in the *IGHV* genes and determine the SHM status



## Rare clone detection for MRD research

Detect rare B cell clones with high sensitivity and ultralow LOD down to  $10^{-6}$ ; measure and compare the frequency of potential clones of interest

## Oncomine BCR IGH-LR Assay

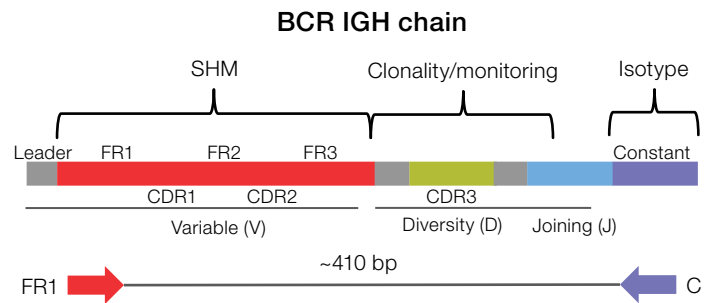
### Accurate clonality assessment and SHM analysis

- Confidently identify the malignant clone even in the most challenging samples by partitioning the B cell repertoire by isotype
- Accurately measure the level of SHM in the *IGHV* genes with the ultralow substitution error rate of the Ion Torrent™ platform, with automated reporting built into the informatics software
- Simple and intuitive clonality assessment is supported by the unique interactive visualizations and automated clonal lineage analysis features built into the informatics software
- Accelerate time to answers with a 48-hour sample-to-results turnaround
- Gain efficiency with the flexibility to multiplex with minimal sample input (as low as 25 ng) from a variety of common hematology sample types

## Oncomine BCR IGH-SR Assay

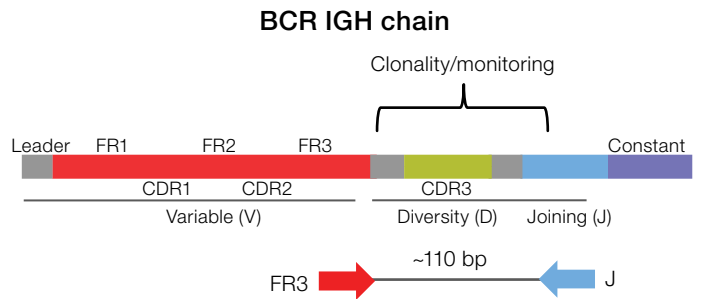
### Sensitive rare-clone detection for MRD research

- Confidently detect rare B cell clones with high sensitivity and ultralow LOD down to  $10^{-6}$
- Easily measure and compare the frequency of the potential clones of interest for MRD research
- Accelerate time to answers with a 48-hour sample-to-results turnaround
- Choose from multiple chips and sample types to fit your unique sample batching needs, desired LOD, and throughput requirements



<b>Amplification strategy</b>	Framework 1 and isotype-specific primers
<b>Input type</b>	Non-FFPE RNA
<b>Sample type</b>	Whole blood, bone marrow, PBLs*, PBMCs*, fresh-frozen specimens
<b>Chip compatibility</b>	Ion 530™ Chip

\* PBL: peripheral blood leukocyte; PBMC: peripheral blood mononuclear cell.



<b>Amplification strategy</b>	Framework 3 and joining gene primers
<b>Input type</b>	gDNA, RNA; FFPE-compatible
<b>Sample type</b>	Whole blood, bone marrow, PBLs, PBMCs, fresh-frozen and FFPE-preserved samples
<b>Chip compatibility</b>	Ion 530, Ion 540™, and Ion 550™ Chips

## Powerful and intuitive informatics solution



- Interactive spectratyping plots make it easy to identify the potential malignant clone of interest within the broader context of the repertoire
- For challenging samples with high polyclonal backgrounds, partition the repertoire by isotype with the click of a button to reveal the malignant lineage
- Automated reporting features provide detailed information on each clone, including the CDR3 sequence, SHM level and frequency, clone frequency, and more
- Unique automated clonal lineage analysis enables the identification of subclones based on specific molecular characteristics/signatures
- Measure and compare the frequency of the potential malignant clone of interest (identified by V-gene and CDR3 NT sequence) for MRD research

## End-to-end clinical research workflow for hematology-oncology



**Collect sample**

### Compatible with:

- Bone marrow
- Whole blood
- PBMCs
- PBLs
- Fresh-frozen samples
- FFPE samples (SR assay only)



**Extract DNA or RNA**

### Oncomine BCR IGH-LR assay:

- Requires RNA input

### Oncomine BCR IGH-SR assay:

- Compatible with DNA and RNA



**Prepare library**

### Prepare libraries from 25 ng to 2 µg of DNA or RNA input:

- Input requirements are overall higher for rare clone detection and vary depending on desired sensitivity



**Sequence**

### Sequence samples using an Ion GeneStudio™ S5 System:

- BCR IGH-LR assay— Ion 530 Chip
- BCR IGH-SR assay— Ion 530, Ion 540, or Ion 550 Chip



**Analyze**

### Perform downstream data analysis with Ion Reporter™ Software v5.12 or higher

## Assay applications

	Oncomine BCR IGH-LR Assay	Oncomine BCR IGH-SR Assay
Clonality assessment	✓	✓
SHM analysis	✓	–
Isotyping	✓	–
Clonal lineage analysis	✓	✓
Rare-clone detection for MRD research	✓ LOD down to $10^{-4}$	✓ LOD down to $10^{-6}$



## Ordering information

Product	Cat. No.
Oncomine BCR IGH-LR Assay, RNA	A45485
Oncomine BCR IGH-SR Assay, DNA	A45483
Oncomine BCR IGH-SR Assay, RNA	A45484

Find out about the Oncomine BCR IGH assays and our NGS technology at [thermofisher.com/blue](https://thermofisher.com/blue)