

## Overview

### Useful For

Determining whether a B-cell or plasma cell population is polyclonal or monoclonal in specimens other than blood or bone marrow

Identifying neoplastic cells as having B-cell or plasma cell differentiation

Monitoring for a persistent neoplasm by detecting an immunoglobulin gene rearrangement profile similar to that from a previous neoplastic specimen

### Testing Algorithm

The following algorithms are available:

- [Gastric MALT Lymphoma Diagnostic Algorithm](#)
- [Gastric MALT Posttherapy Follow-up Algorithm](#)

### Special Instructions

- [Hematopathology Patient Information](#)
- [Gastric MALT Posttherapy Follow-up Algorithm](#)
- [Gastric MALT Lymphoma Diagnostic Algorithm](#)

### Method Name

Polymerase Chain Reaction (PCR)

### NY State Available

Yes

## Specimen

### Specimen Type

Varies

### Shipping Instructions

Body fluid or spinal fluid must arrive within 4 days of collection.

### Specimen Required

Submit only 1 of the following specimens:

**Specimen Type:** Body fluid

**Container/Tube:** Sterile container

**Specimen Volume:** At least 5 mL**Collection Instructions:**

1. If the volume is large, pellet cells prior to sending.
2. Send less volume at ambient temperature or as a frozen cell pellet.

**Specimen Stability Information:**

Body fluid: Ambient 4 days/Refrigerated/Frozen

Cell pellet: Frozen

**Specimen Type:** Paraffin-embedded bone marrow aspirate clot**Container/Tube:** Paraffin block**Specimen Stability Information:** Ambient**Specimen Type:** Frozen tissue**Container/Tube:** Plastic container**Specimen Volume:** 100 mg**Collection Instructions:** Freeze tissue within 1 hour of collection.**Specimen Stability Information:** Frozen**Specimen Type:** Paraffin-embedded tissue**Container/Tube:** Paraffin block**Specimen Stability Information:** Ambient**Specimen Type:** Tissue**Slides:** Unstained slides**Specimen Volume:** 10 Slides**Specimen Stability Information:** Ambient**Specimen Type:** Spinal fluid**Container/Tube:** Sterile vial**Specimen Volume:** 5 to 10 mL**Specimen Stability Information:** Ambient 4 days/Refrigerated**Specimen Type:** Extracted DNA**Container/Tube:** 1.5- to 2-mL tube**Specimen Volume:** Entire specimen**Collection Instructions:**

1. Label specimen as extracted DNA and source of specimen
2. Indicate volume and concentration of DNA on label

**Specimen Stability Information:** Refrigerated/Ambient**Forms**1. [Hematopathology Patient Information](#) (T676)2. If not ordering electronically, complete, print, and send a [Hematopathology/Cytogenetics Test Request](#) (T726) with the specimen.

**Specimen Minimum Volume**

Body and spinal fluid: 1 mL

Tissue: 50 mg

Extracted DNA: 50 microliters (mCL) at 20 ng/mCL

**Reject Due To**

Bone marrow core biopsies Paraffin shavings	Reject
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**Specimen Stability Information**

Specimen Type	Temperature	Time	Special Container
Varies	Varies		

**Clinical & Interpretive****Clinical Information**

The immunoglobulin genes (heavy, kappa, and lambda) are comprised of numerous, discontinuous coding segments. As B cells develop, the segments are rearranged such that each mature B cell and plasma cell has a unique rearrangement profile. Other cell types usually retain the unarranged gene structures. Clonal expansion of any B cell or plasma cell will result in a population of cells that all contain identical immunoglobulin gene rearrangement profiles.

Reactive B-cell or plasma cell expansions are polyclonal, with each clone containing relatively few cells and no single clone predominating. Conversely, neoplastic clones are generally large such that the clonal cells are the predominant B cells or plasma cells present.

In the appropriate clinical and pathologic setting, detection of a prominent immunoglobulin gene rearrangement profile may be equated to the presence of a neoplastic B-cell or plasma cell clone.

**Reference Values**

An interpretive report will be provided.

**Interpretation**

An interpretive report will be provided.

The interpretation of the presence or absence of a predominant immunoglobulin gene rearrangement profile is sometimes subjective. These results must always be interpreted in the context of other clinicopathologic information to determine the significance of the result.

The detection of a clonal immunoglobulin gene rearrangement by this test is not synonymous with the presence of a B-cell or plasma cell neoplasm.

**Cautions**

This test is neither 100% sensitive nor 100% specific.

False-negative results may occur if the immunoglobulin gene has numerous point alterations introduced during expansion in a follicle center (somatic hypermutation) such that none of the polymerase chain reaction (PCR) primers will bind. False-negative results will also occur if the clonal cells have not rearranged the immunoglobulin genes being evaluated or are present below the sensitivity level of the assay (sensitivity is quite variable but the assay requires that at least 1% to 5% of the nucleated cells present be clonal). False-positive results are rare but may occur if a predominant clone (or small number of clones) is produced or sampled from a polyclonal expansion.

The test does not provide information regarding:

- The differentiation of the clonal cell population (neoplastic cells other than B cells or plasma cells may occasionally have immunoglobulin gene rearrangements)
- Whether a prominent clone is physiologic or neoplastic

**Clinical Reference**

1. van Dongen JJ, Wolvers-Tettero IL: Analysis of immunoglobulin and T-cell receptor genes. Part II: Possibilities and limitations in the diagnosis and management of lymphoproliferative diseases and related disorders. *Clin Chim Acta*. 1991 Apr;198(1-2):93-174
2. Coad JE, Olson DJ, Lander TA, McGlennen RC: Molecular assessment of clonality in lymphoproliferative disorders: I. Immunoglobulin gene rearrangements. *Mol Diagn*. 1996 Dec;1(4):335-355
3. Kokovic I, Novakovic BJ, Novakovic S: Diagnostic value of immunoglobulin k light chain gene rearrangement analysis in B-cell lymphomas. *Int J Oncol*. 2015 Mar;46(3):953-962. doi: 10.3892/ijo.2014.2790

**Performance****Method Description**

Genomic DNA is extracted from all specimens.

In the polymerase chain reaction (PCR) assay, a total of 34 upstream and 5 downstream primers are used (Invivoscribe IGH and IGK gene clonality reagents). The primers are designed to amplify fragments from all theoretical rearrangements of the immunoglobulin heavy and kappa light chain genes. Each unique rearrangement should produce PCR fragments of unique sizes. The primers cannot amplify anything if the immunoglobulin genes are not rearranged because the distance is too great. The primers are labeled with a fluorescent tag so that the PCR product can be detected. The PCR fragments are analyzed by capillary gel electrophoresis using a genetic analyzer for fragment size and amount.(Unpublished Mayo method)

**PDF Report**

No

**Day(s) Performed**

Monday through Friday

**Report Available**

7 to 14 days

**Specimen Retention Time**

DNA: 3 months

**Performing Laboratory Location**

Mayo Clinic Laboratories - Rochester Main Campus

**Fees & Codes****Fees**

- Authorized users can sign in to [Test Prices](#) for detailed fee information.
- Clients without access to Test Prices can contact [Customer Service](#) 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact [Customer Service](#).

**Test Classification**

This test was developed using an analyte specific reagent. Its performance characteristics were determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the US Food and Drug Administration.

**CPT Code Information**

81261-IGH (Immunoglobulin heavy chain locus) (eg, leukemias and lymphomas B-cell), gene rearrangement analysis to detect abnormal clonal populations; amplified methodology (eg. polymerase chain reaction)

81264-IGK (Immunoglobulin kappa light chain locus) (eg, leukemia and lymphoma, B-cell) gene rearrangement analysis, evaluation to detect abnormal clonal populations

**LOINC® Information**

Test ID	Test Order Name	Order LOINC® Value
BCGRV	Immunoglobulin Gene Rearrange, V	61113-7

Result ID	Test Result Name	Result LOINC® Value
MP017	Specimen:	31208-2
19915	Final Diagnosis:	34574-4
608950	Signing Pathologist	19139-5