MAYO CLINIC LABORATORIES

Test Definition: CEE52

CD52 Cell Expression Evaluation, Varies

Overview

Useful For

Detecting cell-surface antigens on malignant cells that are potential therapeutic antibody targets, specifically CD52

Determining the eligibility of patients for monoclonal antibody therapies

Monitoring response to the therapeutic antibody

Testing Algorithm

A complete diagnostic B-cell, T-cell, or acute immunophenotyping panel is **not** performed. In some cases, a limited morphologic evaluation will be performed.

Method Name

Immunophenotyping

NY State Available

Yes

Specimen

Specimen Type

Varies

Ordering Guidance

This test should **not** be used as a shortened diagnostic panel. For a complete diagnostic B-cell, T-cell, or acute immunophenotyping panel, order LCMS / Leukemia/Lymphoma Immunophenotyping, Flow Cytometry, Varies.

This test evaluates CD52 expression only. For CD20 expression, order CEE20 / CD20 Cell Expression Evaluation, Varies. For CD49d expression, order CEE49 / CD49d Cell Expression Evaluation, Varies.

Shipping Instructions

Specimen must arrive within 96 hours of collection.

Necessary Information

The following information is required:

1. The therapeutic monoclonal antibody being used or considered

2. The pertinent hematologic diseases that have been diagnosed or considered

3. Diagnostic immunophenotype of the neoplastic cells (when available, a pathology and/or flow cytometry report should be included).

4. Specimen source

5. Date and time of collection

Specimen Required

Submit only 1 of the following specimens:

Specimen Type: Blood
Container/Tube:
Preferred: Yellow top (ACD)
Acceptable: Green top (sodium heparin) or lavender top (EDTA)
Specimen Volume: 10 mL
Collection Instructions:
1. Send specimen in original tube. **Do not aliquot**.
2. Label specimen as blood.
Specimen Stability Information: Ambient/Refrigerated

Specimen Type: Bone marrow
Container/Tube:
Preferred: Yellow top (ACD)
Acceptable: Green top (sodium heparin) or lavender top (EDTA)
Specimen Volume: 1-5 mL
Collection Instructions:
1. Label specimen as bone marrow.
2. Submission of bilateral specimens is not required.
Specimen Stability Information: Ambient/Refrigerated

Forms

If not ordering electronically, complete, print, and send a Hematopathology/Cytogenetics Test Request (T726) with the specimen.

Specimen Minimum Volume

Blood: 3 mL
Bone Marrow Aspirate: 1 mL

Reject Due To

All specimens will be evaluated at Mayo Clinic Laboratories for test suitability.

Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Varies	Varies	4 days	

Clinical & Interpretive

Clinical Information

Monoclonal antibodies are critical tools for detecting cellular antigens in various hematologic diseases and are used to provide critical prognostic information (eg, CD49d). Monoclonal antibodies are also used as therapeutic agents in a

variety of hematologic diseases. For example:
-Anti-CD20 (Rituxan): B-cell malignant lymphomas and multiple myeloma
-Anti-CD52 (Campath-1H): B-cell chronic lymphocytic leukemia and T-cell disorders
This list will undoubtedly expand over time to include other antibodies.

It may be necessary to document expression of these markers by the malignant cells prior to initiating the respective monoclonal antibody therapy. Expression of these markers may also be required for follow-up to monitor the impact of treatment on residual normal counterparts (eg, CD20-positive lymphocytes in patients treated with anti-CD20).
The distribution of these cellular antigens is well established in normal, reactive, and in various malignant disorders. The laboratory has several years of experience with therapeutic antibody monitoring of Mayo Clinic patients as part of the routine B-cell, T-cell, or acute immunophenotyping panels.

Reference Values
Normal individuals have B lymphocytes, T lymphocytes, or myeloid cells that express the corresponding cell-surface antigens in question.

Interpretation
The immunophenotyping report will summarize the pattern of antigenic expression on malignant cells and, if appropriate, the normal cellular counterparts that correspond to the therapeutic monoclonal antibody target.

Cautions
No significant cautionary statements.

Clinical Reference
1. Salles G, Barrett M, Foa R, et al: Rituximab in B-cell hematologic malignancies: A review of 20 years of clinical experience. Adv Ther. 2017;34(10):2232-2273. doi:10.1007/s12325-017-0612-x
2. Bachy E, Seymour JF, Feugier P, et al: Sustained progression-free survival benefit of rituximab maintenance in patients with follicular lymphoma: Long-term results of the PRIMA study. J Clin Oncol. 2019;37(31):2815-2824. doi:10.1200/JCO.19.01073
3. Cross M, Dearden C. B and T cell prolymphocytic leukaemia. Best Pract Res Clin Haematol. 2019;32(3):217-228. doi:10.1016/j.beha.2019.06.001
4. Braun T, von Jan J, Wahnschaffe L, Herling M. Advances and perspectives in the treatment of T-PLL. Curr Hematol Malig Rep. 2020;15(2):113-124. doi:10.1007/s11899-020-00566-5
5. Strati P, Parikh SA, Chaffee KG, et al: CD49d associates with nodal presentation and subsequent development of lymphadenopathy in patients with chronic lymphocytic leukaemia. Br J Haematol. 2017;178(1):99-105. doi:10.1111/bjh.14647

Performance

Method Description
Flow cytometric immunophenotyping of peripheral blood, bone marrow, or tissue-derived lymphocytes is performed to assess the expression of the cell-surface antigen corresponding to the monoclonal antibody therapeutic target. The following antibody panel will be used:
-Anti-CD52 assessment: CD52/CD19/CD3/CD45

(Keren P, McCoy Jr JP, Carey J eds. Flow Cytometry in Clinical Diagnosis. 4th ed. ASCP Press; 2007)

PDF Report
No

Day(s) Performed
Monday through Friday

Report Available
1 to 4 days

Specimen Retention Time
2 weeks

Performing Laboratory Location
Rochester

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
88184-Flow cytometry; first cell surface, cytoplasmic or nuclear marker
88185 x 3-Flow cytometry; additional cell surface, cytoplasmic or nuclear marker
88187-Flow Cytometry Interpretation, 2 to 8 Markers

LOINC® Information

Test ID	Test Order Name	Order LOINC® Value
CEE52	CD52 Cell Expression Evaluation, V	100991-9

Result ID	Test Result Name	Result LOINC® Value
CK166	CEE52 Result	No LOINC Needed
CK167	Final Diagnosis	22637-3