

## Overview

### Useful For

Identification of tumor cells expressing claudin 18

As an aid in screening patients who may be eligible for VYLOY (zolbetuximab) treatment

### Method Name

Immunohistochemistry (IHC)

### NY State Available

Yes

## Specimen

### Specimen Type

Special

### Shipping Instructions

Attach the green "Attention Pathology" address label (T498) to the outside of the transport container before putting into the courier mailer.

### Necessary Information

A pathology/diagnostic report and a brief history are required.

### Specimen Required

This assay requires at least 50 tumor cells.

**Specimen Type:** Tissue

**Supplies:** Pathology Packaging Kit (T554)

**Submit:**

Formalin-fixed, paraffin-embedded tissue block

OR

3 unstained glass, "positively charged" slides with 4-microns formalin-fixed, paraffin-embedded tissue

**Additional Information:** One slide will be stained with hematoxylin and eosin and returned.

### Forms

If not ordering electronically, complete, print, and send a [Immunohistochemical \(IHC\)/In Situ Hybridization \(ISH\) Stains Request](#) (T763) with the specimen.

**Reject Due To**

Decalcified paraffin embedded tissue	Reject
Wet/frozen tissue	Reject
Cytology smears	Reject
Nonformalin fixed tissue including alcohol-formalin n-acetic acid (AFA), 95% ethanol, PREFER fixatives or Zinc formalin	Reject
Nonparaffin embedded tissue	Reject
Noncharged slides	Reject
ProbeOn slides	Reject

**Specimen Stability Information**

Specimen Type	Temperature	Time	Special Container
Special	Ambient (preferred)		
	Refrigerated		

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

Claudin 18 (CLDN18) is a member of the claudin protein family that regulates cell adhesion. A qualitative immunohistochemistry assessment using mouse monoclonal anti-claudin 18, clone 43-14A, serves as a biomarker in gastric and gastroesophageal junction cancer and may aid in identifying patients eligible for VYLOY (zolbetuximab) treatment.

**Reference Values**

An interpretive report will be provided.

**Interpretation**

Claudin 18.2 is a biomarker for gastric and gastroesophageal junction cancer. When positive (defined as at least moderate membranous staining in greater than or equal to 75% of viable tumor cells), it predicts response to zolbetuximab which has been approved for the treatment of gastric and gastroesophageal adenocarcinoma.

**Cautions**

This test has been validated for nondecalcified paraffin-embedded tissue specimens fixed in 10% neutral-buffered formalin. Recommended fixation time is between 6 and 48 hours. This assay has not been validated on tissues subjected to the decalcification process or use of alternative fixatives for bone and bone marrow specimens or cell blocks.

Age of a cut paraffin section can affect immunoreactivity. Stability thresholds vary widely among published literature and are antigen dependent. Best practice is for paraffin sections to be cut within 6 weeks.

**Clinical Reference**

1. Kubota Y, Kawazoe A, Mishima S, et al. Comprehensive clinical and molecular characterization of claudin 18.2 expression in advanced gastric or gastroesophageal junction cancer. *ESMO Open*. 2023;8(1):100762
2. Cao W, Xing H, Li Y, et Al. Claudin18.2 is a novel molecular biomarker for tumor-targeted immunotherapy. *Biomark Res*. 2022;10(1):38
3. Pellino A, Brignola S, Riello E, et al. Association of CLDN18 protein expression with clinicopathological features and prognosis in advanced gastric and gastroesophageal junction adenocarcinomas. *J Pers Med*. 2021;11(11):1095
4. Arnold A, Daum S, von Winterfeld M, et al. Prognostic impact of Claudin 18.2 in gastric and esophageal adenocarcinomas. *Clin Transl Oncol*. 2020;22(12):2357-2363

**Performance****Method Description**

Immunoperoxidase staining and detection of Claudin 18 (CLDN18) is performed in formalin-fixed, paraffin-embedded tissue sections using the VENTANA CLDN18 (43-14A) RxRx Assay. The 4-micron tissue sections are deparaffinized, subjected to heat-induced antigen retrieval and peroxidase inhibitor, incubated with a monoclonal antibody (clone 43-14A), and visualized using a proprietary kit detection system. Sections are counterstained with hematoxylin. Stained slides are examined microscopically by the consulting anatomic pathologist.(Package insert: VENTANA CLDN18 (43-14A) RxRx Assay. Ventana Medical Systems, Inc.; 1016387US Rev A, 10/11/2024)

Results are interpreted using the manufacturer-provided interpretation guide. A minimum of 50 tumor cells are required for interpretation of results.(Instruction manual: VENTANA CLDN18 (43-14A) RxRx Assay Interpretation Guide for Gastric Adenocarcinoma including Gastroesophageal Junction (GEJ). Ventana Medical Systems, Inc.; 1021576EN Rev B, 10/01/2024)

**PDF Report**

No

**Day(s) Performed**

Monday through Friday

**Report Available**

5 to 7 days

**Specimen Retention Time**

Until reported

**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 has been modified from the manufacturer's instructions. 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**

88360

**LOINC® Information**

Test ID	Test Order Name	Order LOINC® Value
CLD18	CLDN18, SemiQuant IHC, Manual	105011-1

Result ID	Test Result Name	Result LOINC® Value
620666	Interpretation	50595-8
621004	Participated in the Interpretation	No LOINC Needed
621005	Report electronically signed by	19139-5
621006	Material Received	81178-6
621007	Disclaimer	62364-5
621008	Case Number	80398-1