Labcorp Oklahoma, Inc. Test Directory

CA 27.29

Order Name: CA27.29
Test Number: 2024375
Revision Date: 12/12/2022

		DOLOGY	LOINC CODE
	Immuno	ochemiluminometric (ICMA)	17842-6
NTS			
pecimen Volume (min)	Specimen Type	Specimen Container	Transport Environment
mL (0.3)	Serum	Clot Activator SST	Refrigerated
pecimen Type: Red-top tube or g pecimen Storage: Refrigerated pecimen Collection: If a red-top t pecial Instructions: It is recomme ot provide serial monitoring; it is into	el-barrier tube ube is used, transfer the separated ended that one assay method be us tended for one-time use only.	ed consistently to monitor a patient's course	of therapy. This procedure does
r	pecimen Volume (min) mL (0.3) otes: 0.3 mL (Note: This volume E pecimen Type: Red-top tube or g pecimen Storage: Refrigerated pecimen Collection: If a red-top to pecial Instructions: It is recommental provide serial monitoring; it is interested.	pecimen Volume (min) Specimen Type ML (0.3) Serum Otes: 0.3 mL (Note: This volume Does NOT allow for repeat testing.) pecimen Type: Red-top tube or gel-barrier tube pecimen Storage: Refrigerated pecimen Collection: If a red-top tube is used, transfer the separated pecial Instructions: It is recommended that one assay method be used to provide serial monitoring; it is intended for one-time use only.	pecimen Volume (min) Specimen Type Specimen Container Clot Activator SST otes: 0.3 mL (Note: This volume Does NOT allow for repeat testing.) pecimen Type: Red-top tube or gel-barrier tube pecimen Storage: Refrigerated pecimen Collection: If a red-top tube is used, transfer the separated serum to a plastic transport tube. pecial Instructions: It is recommended that one assay method be used consistently to monitor a patient's course

GENERAL INFORMATION		
Expected TAT	1-2 days	
Clinical Use	The CA 27.29 assay is intended for use in monitoring: 1) disease progression and/or response to therapy in patients with metastatic disease, and 2) disease recurrence in patients treated previously for stages II or III breast carcinoma who are clinically free of the disease. Serial testing in patients who are clinically free of disease should be used in conjunction with other clinical methods for early detection of cancer recurrence. Limitations: Patients with confirmed breast carcinoma frequently have CA 27.29 assay values in the same range as healthy individuals. Elevations may also be observed in patients with non-malignant disease. Results of this test must always be interpreted in the context of morphologic and other relevant data and should not be used alone for a diagnosis of malignancy	
Performing Labcorp Test Code	140293	
Notes	Labcorp Test Code: 140293	
CPT Code(s)	86300	
Lab Section	Reference Lab	

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