Test Definition: CRCOF

Chromium and Cobalt, Synovial Fluid

Reporting Title: Chromium and Cobalt, Synovial Fl

Performing Location: Rochester

Ordering Guidance:

This test should only be used in individuals with chromium or cobalt implants.

Specimen Requirements:

Patient Preparation: High concentrations of gadolinium and iodine are known to interfere with most inductively coupled plasma mass spectrometry-based metal tests. If either gadolinium- or iodine-containing contrast media has been administered, a specimen should not be collected for at least 96 hours.

Supplies: Metal Free B-D Tube (EDTA), 6 mL (T183) Container/Tube: Royal blue top (metal-free EDTA)

Specimen Volume: 1 mL

Collection Instructions: See Metals Analysis Specimen Collection and Transport for complete instructions.

Additional Information: Cobalt and chromium are present in the black rubber plunger seals found in most disposable

syringes. As a result, synovial fluid should not be collected in these devices as contamination may occur.

Specimen Minimum Volume:

0.4 mL

Specimen Type	Temperature	Time	Special Container
Synovial Fluid	Refrigerated (preferred)	90 days	
	Ambient	90 days	
	Frozen	90 days	

Result Codes:

Result ID	Reporting Name	Туре	Unit	LOINC®
606353	Chromium, Synovial FI	Numeric	ng/mL	95526-0
	Also used by tests: CRSY			
606352	Cobalt, Synovial Fl	Numeric	ng/mL	23842-8
	Also used by tests: COSY			

LOINC and CPT codes are provided by the performing laboratory.

Supplemental Report:

No



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Chromium and Cobalt, Synovial Fluid

Components:

Test ID	Reporting Name	CPT Units	CPT Code	Always Performed	Orderable Separately
CRSY	Chromium, Synovial FI			Yes	Yes
COSY	Cobalt, Synovial FI			Yes	Yes

CPT Code Information:

83018 82495

Reference Values:

CHROMIUM:

0-17 years: Not established > or =18 years: <16.9 ng/mL

COBALT:

0-17 years: Not established > or =18 years: <19.8 ng/mL