

PLEASE COMPLETE ALL REQUIRED F	IELDS (*) ON THIS FORM.	FAILU	RE TO DO	SO WILL DELAY RESULT REPORTING	
ACCOUNT INFORMATION				PATIENT INFORMATION	
Synovasure Account Number*	Phone #*			Patient Name* (Last, First)	
Ordering Physician* (Last, First)					
Zimmer Biomet Sales Representative (Last, First)			Date of Birth*		
Physician NPI #*			Gender* Male Female		
Practice Name & Address*	Result Fax #*			Patient Address*	
	Result Email (Optional)				
				Patient Phone #*	
Has this provider submitted a specimen to CD Laboratories in the past? Yes No If no, please ensure that the account information above is provided so provider can be enrolled.			Hospital patient? [*] Yes No If yes, please provide facility information		
BILLING INFORMATION Please include a cop	y of the front and back of the po	atient's i	nsurance ca	rd(s)	
				late of injury, employer, and adjuster's contact detai nitting specimens. See reverse for test CPT codes .	
Primary Insurance Carrier	autionzation from patient's in	1		rrier Phone #	
ID#			Group #		
Name of Insured			Rela	tion to Patient: 🛛 Self 🖓 Spouse 🖓 Other	
Claim Address:					
Prior Authorization (PA)* # PA Valid Date*					
Secondary Insurance Carrier?	nographic sheet) 🛛 No				
By signing this form, I hereby authorize CD Laboratories to furnish on this form if necessary for reimbursement. I also authorize bene understand that I am responsible for any amounts not paid by insi	fits to be payable to CD Laboratories. I		Patient Sign	ature*:	
non-covered and non-authorized services. I permit a copy of this authorization to be used in place of the original.					
SPECIMEN DETAIL		Drew	Ka	aa Uin Otkar	
Collection Date*		Draw Site [*]	<u>Kne</u> □ Right		
Diagnosis Codes* (ICD-10-CM) See reverse for examples		Is patie	nt currently o	n antibiotics? 🗆 Yes 🛛 No	
TEST SELECTION Select test(s) from either the Comprehensive Synovial Fluid Infection Panels Section OR the Individual Synovial Fluid Infection Tests Section COMPREHENSIVE SYNOVIAL FLUID INFECTION PANELS					
Note: Comprehensive Panels require 3 tubes of synovial fluid 📕 - 3mL 📕 - 2mL 📗 - 1mL					
FOR TOTAL JOINT REPLACEMENTS (see reverse for list of CPT codes): FOR NATIVE JOINTS (see reverse for list of CPT codes):					
□ Comprehensive Periprosthetic Joint Infection Panel (PJI) (70000) □ Comprehensive Native Septic Arthritis Panel (NSA) (74001)					
Includes: Specimen integrity analysis, Synovasure [®] Alpha Defensin for PJI (<i>alpha defensin,</i> <i>CRP</i>), Culture, WBC count w/ differential and RBC count, <i>CRP</i>), Culture, WBC count w/ differential and RBC count,					
			tial and RBC count, Synovasure® Neutrophil Elastase,		
Add Synovasure® Microbial ID Panel (7895) Crystal Analysis, Synovasure® Microbial ID Panel				Analysis, Synovasure [®] Microbial ID Panel	
Add Crystal Analysis; MSU, CPPD (831)					
INDIVIDUAL SYNOVIAL FLUID INFECTION TESTS					
SUBMIT IN CLEAR TOPPED (RED STOPPER) TUBES (BD366703) SUBMIT IN LAVENDER TUBES (BD367856)					
Synovasure® Alpha Defensin for PJI (0.5mL) (7002) Synovasure® Microbial ID Panel (1.0mL) (7895) WBC Count w/ Differential and RBC count (0.5mL)					
Synovasure® Alpha Defensin for NSA (0.5mL) (7452) OR - Select Individual Pathogens: (8260)					
Culture (3.0mL) (70554) Culture (3.0mL) (70554) Crystal Analysis; MSU, CPPD (0.5mL) (831) Crystal Analysis; MSU, CPPD (0.5mL) (831)				Crystal Analysis; MSU, CPPD (0.5mL) (831)	
Synovasure® Neutrophil Elastase (0.5mL) (7120)					

M40005B V9 Feb 2020

Specimen Integrity Testing

.

The accuracy of diagnostic tests such as synovial fluid white blood cell count, neutrophil percentage, and others, can be significantly impacted by the quality of the specimen that is submitted for evaluation. As part of sample analysis, specimen integrity tests are performed on synovial fluid specimens. Physicians are notified when a suboptimal specimen has been submitted. Our specimen integrity tests assess:

- Absorbance at 280 nm (A280) Specimens that fall outside the normal range for synovial fluid may be caused by dilution via saline lavage or use of contrast agents
- Red Blood Cell Count Specimens are verified as characteristic of synovial fluid, not blood.

INFECTION TEST MENU	
Synovasure Alpha Defensin for PJI	The Synovasure Alpha Defensin lab developed test (LDT) for PJI consists of assays for synovial fluid Alpha Defensin and CRP and has been validated for use as an adjunct for the diagnosis of periprosthetic joint infection.
Synovasure Alpha Defensin for NSA	The Synovasure Alpha Defensin LDT for NSA consists of assays for synovial fluid Alpha Defensin and lactate and has been validated for use as an adjunct for the diagnosis of native septic arthritis.
Synovasure Microbial Identification Panel	Synovasure MID LDT is an assay intended for the early detection of microbial antigen in synovial fluid. The assay can detect microbial antigen in some samples where organism is present but was not able to be cultured. Current panel includes <i>Staphylococcus</i> species, <i>Candida</i> species, <i>Enterococcus</i> species, and <i>Propionibacterium acnes</i> .
Synovasure Neutrophil Elastase	The Synovasure Neutrophil Elastase (NE) LDT was designed to be a replacement for the Leukocyte Esterase (LE) test strip which can serve as one of the criteria in the MSIS infection algorithm. The NE LDT is designed and validated specifically for synovial fluid, while the LE test strip is designed for urine. The NE LDT is not prone to the high rate of invalid results due to blood contamination that have been reported with the LE test strip. A positive NE result should be interpreted as meeting the MSIS criteria of a positive LE test strip.
Synovial Fluid Culture	Anaerobic and aerobic culture bottles incubated for 7 days. Includes organism identification and antibiotic susceptibilities. Shoulder specimen cultures are supplemented to enhance growth and incubated for 14 days.
Synovial Fluid WBC Count with Differential and RBC Count	Automated high-performance cell count system with differential and RBC count. Elevated white blood cells (>3000c/mm ³) are confirmed with a manual count.
Synovial Fluid Crystal Analysis	Monosodium urate (MSU) and Calcium pyrophosphate dihydrate (CPPD) crystal detection using polarized microscopy.

M17.9 Osteoarthritis of knee, unspe	
WITT.9 Ostebal tillitis of knee, unsp	ecified
M25.469 Effusion, unspecified knee	
M25.559 Pain in unspecified hip	
M25.561 Pain in knee, right	
M25.562 Pain in knee, left	
T84.039A Mechanical loosening of uns initial encounter	pecified internal prosthetic joint,
T84.039D Mechanical loosening of uns subsequent encounter	pecified internal prosthetic joint,
	pecified internal prosthetic joint,
T84.50XA Infection and inflammatory l joint prosthesis, initial encou	RX due to unspecified internal Inter
T84.50XD Infection and inflammatory joint prosthesis, subsequent	RX due to unspecified internal encounter
T84.50XS Infection and inflammatory joint prosthesis, sequential e	RX due to unspecified internal encounter
Z96.641 Presence of artificial hip join	t, right
Z96.642 Presence of artificial hip join	t, left
Z96.651 Presence of artificial knee jo	int, right
Z96.652 Presence of artificial knee jo	int, left

Test CPT Codes – *Please use these codes for prior authorization*

Synovasure Alpha Defensin for PJI					
Alpha Defensin-SF	83516				
CRP-SF	86140				
Synovasure Alpha Defensin for NSA					
Alpha Defensin-SF	83516				
Lactate-SF	83605				
Synovasure Microbial ID	87449 (x3)				
Synovasure Neutrophil Elastase	83516				
Synovial Fluid Culture					
Aerobic	87070				
Anaerobic	87075				
WBC Count w/ Differential	89051				
Crystal Analysis	89060				