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Primarius Pathology

		Diar	rhea Path	ogen Requis	ition				
Date Specimen Collected:			Time Specir	nen Collected:					
Laboratory Use Only Accession Number			Received			Time Received	I		
			_			ce Contact Inf			
Practice Name		Practice ID				ss			
Ordering Physicians 🔲						tate, Zip			
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		Dation	at and Inc	urance Inforr					
	(Currently	Patier Not Accepting I				ntract Pending)		
First Name			Last Name			Middle Initial		Gender*_	
Address Line 1		Address	Line 2			City	Sta	ite Z	
DOB		Home Pl							
Gender Identity*									
Insured's Name									
Home Phone									
Primary Insurance									
Group # Address				Group #					
City				City		State	Zip		
				athogen Tes			ip		
Enteric Parasite Panel Only Genesis Expanded Diarrhea positive] **Please note that an i If an appropriate immunoa Calprotectin Lactoferrin Antigliadin Ab (Fecal Anti-Gliadii Anti-tTG (Fecal Anti-Transglutanii Fecal Pancreatic Elastase (Test for Fecal Bile Acids	mmunosupres unosuppression ssays n Ab IgA) ninase Ab IgA)	sion diagnosis coo n code is not selec	de must be se ted, testing v	lected in order fo	Additional Transcription of the second of th	be performed. to the "Genesis al Stool Diagnos igen) est (FIT) for Occ s (Real-Time PCF	Diarrhea Path stic Tests ult Blood		7
				Wet Mount, Trichrome	Stain, and Enteri	c Parasite Panel)			
**For 26 pathogen multiple	avad papal tas	ing there must b		O Codes	upprossion	codos in additio	n to the prime	ry diagnosi	code
Primary diagnosis code: Diarrhea, Unspecified with Feve K92.1) Diarrhea, Unspecified with Abdo Unspecified (R19.7/R10.9) Diarrhea, Unspecified with other electrolyte and fluid balance (R1	r (R19.7/R50.9) atochezia (R19.7/ ominal pain, r disorders of	B20 Huma D80.2 Sele D80.3 Sele D80.4 Sele D80.5 Imr	pression diag an Immunodef ective deficien ective deficien subclasses ective deficien nunodeficienc immunoglobu	inosis codes: iciency Virus [HIV] cy of immunoglobicy of immunoglobicy of immunogloby with increased ulin M [IgM] cy with near-norm ulins or with	disease ulin A [IgA] ulin G [IgG] ulin M [IgM]	D81.89 Oth D82.8 Imm D83.8 Oth D84.81 Imm D84.822 Imm	ner combined im nunodeficiency a specified major er common varia nunodeficiency classified elsew nmunodeficienc her immunodefi	nmunodeficie associated with defects able immunod due to condit here cy due to exte	encies th other deficiencies tions
Duration of Diarrhea:		**Fo	or 26 pathog nunosuppre	gen multiplexed ssion codes in d	d panel tes addition to	ting, there me the primary o	ust be <u>at leas</u> diagnosis cod	it <u>one</u> of th le.	e
This test is medically necessary for impairment, symptom, syndrome o medical management and treatment provider is authorized by law to ord	r disorder. The res nt decisions. The	sults will determine r person listed as the o	ny patient's		laim to my ins	reby authorize Ger urance company f			
				Signature of	Patient (REQ	JIRED)		Date	
Signature of Physician or Other Au	thorized NPI Pro	vider (REQUIRED)	Date			Accessioner In	itials 1	2	

Genesis Diarrhea Pathogen Panel

(11 Pathogens with Genotypic Antibiotic Resistance. Reflex to active toxin detection assay if C.diff positive.)

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Bacterial Pathogens	Viruses	Genotypic Antibiotic Resistance Markers (ARMs) Vancomycin Resistance	
Campylobacter jejuni Clostridium difficile (Clostridioides difficile) Salmonella spp. Vibrio spp. (V. parahaemolyticus / V. vulnificus / V. cholerae) Yersinia enterocolitica	Norovirus (GI/GII) Rotavirus (A/B/C)	vanA vanB Nitroimidazole Resistance nimA nimD Examples: metronidazole, tinidazol	
Diarrheagenic Escherichia coli/ Shigella	Parasites	Erythromycin Resistance ermA ermB	
Shiga-like toxin-producing <i>E. coli</i> (STEC) -including serotype 0157 Shigella/Enteroinvasive <i>E. coli</i> (EIEC)	Entamoeba histolytica Giardia lamblia	Quinolone and Fluoroquinolone Resistance qnrA qnrB Examples: ciprofloxacin, levofloxacin, moxifloxacin Trimethoprim-Sulfamethoxazole Resistance dfrA1 dfrA5	

Genesis Expanded Diarrhea Pathogen Panel

(26 Pathogens with Genotypic Antibiotic Resistance. Reflex to active toxin detection assay if C.diff positive.)

Bacterial Pathogens Campylobacter spp. Campylobacter jejuni Toxigenic Clostridium difficile (Clostridioides difficile) Enteroaggregative E. coli (EAEC) Enteropathogenic E. coli (EPEC) Enterotoxigenic E. coli (ETEC) Vibrio spp. Vibrio cholerae Vibrio parahaemolyticus Yersinia enterocolitica Diarrheagenic Escherichia coli/ Shigella Shigella /Enteroinvasive E. coli (EIEC) Plesiomonas shigelloides Shiga-like toxin producing E. coli (STEC) - including serotype 0157	Viruses Adenovirus F40/41 Astrovirus Norovirus GI Norovirus GII Rotavirus A Rotavirus B Rotavirus C Sapovirus I,II,IV Combination Sapovirus V Parasites Cryptosporidium Entamoeba histolytica Giardia lamblia	Genotypic Antibiotic Resistance Markers (ARMs) Vancomycin Resistance vanA vanB Nitroimidazole Resistance nimA nimD Examples: metronidazole, tinidazol Erythromycin Resistance ermA ermB Quinolone and Fluoroquinolone Resistance qnrA qnrB Examples: ciprofloxacin, levofloxacin, moxifloxacin Trimethoprim-Sulfamethoxazole Resistance dfrA1 dfrA5
Enteric Parasite Par	nel	Toxigenic Clostridium difficile (Clostridioides difficile)
Cryptosporidium (C.hominis and C.parvum) Cyclospora cayetanesis Entamoeba histolytica		Toxigenic Clostridium difficile (Clostridioides difficile) -Reflex to active Toxin detection by EIA for positive samples

Additional ICD-10 Code Immunosuppression Descriptions D82.4 Hyperimmunoglobulin E [IgE] syndrome Z94

D83.0 Common variable immunodeficiency with

and function

Z94.0 Kidney transplant status

Z94.1 Heart transplant status

Z94.2 Lung transplant status

predominant abnormalities of B-cell numbers

D80.0 Hereditary hypogammaglobulinemia D80.1 Nonfamilial hypogammaglobulinemia D80.8 Other immunodeficiencies with pre dominantly antibody defects D81.0 Severe combined immunodeficiency [SCID] with reticular dysgenesis D81.1 Severe combined immunodeficiency [SCID] with low T- and B-cell numbers D81.2 Severe combined immunodeficiency [SCID] with low or normal B-cell numbers D81.31 Severe combined immunodeficiency due to adenosine deaminase deficiency D81.4 Nezelof's syndrome D81.5 Purine nucleoside phosphorylase [PNP] deficiency D81.6 Major histocompatibility complex class I deficiency D81.7 Major histocompatibility complex class II deficiency D81.810 Biotinidase deficiency D81.818 Other biotin-dependent carboxylase deficiency D82.0 Wiskott-Aldrich syndrome

D82.1 Di George's syndrome

D82.2 Immunodeficiency with short-limbed

defective response to Epstein-Barr virus

D82.3 Immunodeficiency following hereditary

Giardia lamblia

D83.1 Common variable immunodeficiency with predominant immunoregulatory T-cell disorders D83.2 Common variable immunodeficiency with autoantibodies to B- or T-cells D84.0 Lymphocyte function antigen-1 [LFA-1] defect D84.1 Defects in the complement system D89.0 Polyclonal hypergammaglobulinemia D89.1 Cryoglobulinemia D89.3 Immune reconstitution syndrome D89.41 Monoclonal mast cell activation syndrome D89.42 Idiopathic mast cell activation syndrome D89.43 Secondary mast cell activation D89.49 Other mast cell activation disorder D89.810 Acute graft-versus-host disease D89.811 Chronic graft-versus-host disease D89.812 Acute on chronic graft-versus-host disease D89.82 Autoimmune lymphoproliferative syndrome [ALPS] D89.89 Other specified disorders involving the immune

mechanism, not elsewhere classified

Z94.3 Heart and lungs transplant status Z94.4 Liver transplant status Z94.5 Skin transplant status Z94.6 Bone transplant status Z94.81 Bone marrow transplant status Z94.82 Intestine transplant status Z94.83 Pancreas transplant status

Z94.84 Stem cells transplant status

Listed for reference only -Applicable codes must be indicated on front side of test requisition. When reporting ICD-10-CM code R19.7 one of the immunosuppression diagnosis codes listed above must also be reported to support medical necessity and provide coverage for CPT code 87507.