

Locations: Louisville, Elizabethtown, Radcliff, Smiths Grove, Bowling Green, Russellville.

Fax: 270-506-2466, Phone: 270-506-2463

Immunoglobulin Referral Form						
Patient Name			Home Phone			
Date of Birth			Mobile or Work Phone			
Pat	ient Home Address		City Stat	ite Zip		
Prir	nary Insurance Name	I				
Primary Insurance ID			Primary Insurance Group			
Insured Name			Insured DOB			
				urance Group		
Secondary Insurance Name Secondary Insurance ID			Secondary Insurance Group			
Ordering Physician's Name						
Orc	lering Physician's Name		NPI			
Address			City Stat	ite Zip		
Phone			Fax			
Please fax the following information: History and Physical Pertinent Lab Work Front & Back copy(s) of patient's insurance card(s)						
Prescription						
Intravenous Immunoglobulin Subcutaneous Immunoglobulin						
	0.4 gm/kg 1 gm/kg 2 gm/kg grams		Infuse grams OR mls using sites			
Infuse: IV daily x day(s); repeat every week(s) x cycles			time(s) per week for months.			
Other:						
Hyo	Hydration order: mls NSiv to be infused prior/post IVIG.					
	Pre-medications: Acetaminophen 650mg PO 30 mins prior to infusion Other Pre-medications:					
			formation			
Clinical Information						
Patient Weight Height			Allergies			
IV access [for IVIGg patients only]: Nurs			se to place PIV prior to therapy			
	_	ICD-10	Diagnosis			
	Diagnosis	100-10	Diagnosis		ICD-10	
	Neuromuscular:		Immune Deficiency:			
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP)	G61.81	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Co	Cell Disorders	D83.1	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS)	G61.81 G61.0	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Co Combined Immunodeficiency, Unspecified		D83.1 D81.9	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy	G61.81 G61.0 G61.82	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec		D83.1 D81.9 D83.9	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG)	G61.81 G61.0	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec Hereditary Hypogammaglobulinemia		D83.1 D81.9	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy	G61.81 G61.0 G61.82 G70.0	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec		D83.1 D81.9 D83.9 D80.0	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation	G61.81 G61.0 G61.82 G70.0 G70.01	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspe- Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM		D83.1 D81.9 D83.9 D80.0 D80.5	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Cr Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome	661.81 661.0 661.82 670.0 670.01 604.81 661.89	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-CC Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec- Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 D83.9 L12.0	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other:	661.81 661.0 661.82 670.0 670.01 604.81 661.89 635 625.82	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Ci Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L12.0 L10.9	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Ci Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers Selective Deficiency of IgG Subclasses	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Ci Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec- Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers SCID with Low or Normal B-Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE)	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
Ple	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol:	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
Ple	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Ci Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspec- Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers SCID with Low or Normal B-Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE)	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
Ple	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol:	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis CBC/diff CMP IgG w/ subclasses 1-4 Quant. Ig Frequency:	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers ScID with T- and B- Cell Numbers Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol:	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis ase Draw: CBC/diff CMP IgG w/ subclasses 1-4 Quant. Ig Frequency:	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol: PER Prescriber Protocol	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis ase Draw: CBC/diff CMP IgG w/ subclasses 1-4 Quant. Ig Frequency:	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90 M33.20	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol: PER Prescriber Protocol	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis ase Draw: CBC/diff CMP IgG w/ subclasses 1-4 Quant. Ig Frequency:	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90 M33.20	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers ScID with T- and B-Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol: PER Pharmacy Protocol PER Prescriber Protocol:	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis	G61.81 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90 M33.20	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers Sclip with T- and B- Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol: PER Pharmacy Protocol PER Prescriber Protocol:	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis	G61.81 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90 M33.20	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Pemphigoid Pemphigus SCID with Low or Normal B-Cell Numbers ScID with T- and B-Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol: PER Pharmacy Protocol PER Prescriber Protocol:	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis	G61.81 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90 M33.20	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers Sclip with T- and B- Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol: PER Pharmacy Protocol PER Prescriber Protocol:	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis	G61.81 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90 M33.20	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers Sclip with T- and B- Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol: PER Pharmacy Protocol PER Prescriber Protocol:	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
	Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS) Multifocal Motor Neuropathy Myasthenia Gravis (MG) Myasthenia Gravis with (Acute) Exacerbation Autoimmune Encephalopathy Inflammatory Neuropathies Relapsing Remitting Multiple Sclerosis (RRMS) Stiff Person Syndrome Other: Idiopathic Thrombocytopenic Purpura Dermatopolymyositis Polymyositis	G61.81 G61.82 G70.0 G70.01 G04.81 G61.89 G35 G25.82 D69.3 M33.90 M33.20	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-C Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigus SCID with Low or Normal B-Cell Numbers SCID with T- and B- Cell Numbers Sclip with T- and B- Cell Numbers Selective Deficiency of IgG Subclasses Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol: PER Pharmacy Protocol PER Prescriber Protocol:	ecified	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 D83.9 L12.0 L10.9 D81.2 D81.1 D80.3 D80.6	
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that is required for this prescription and for any future refills of the same prescription for the patient listed above which I order. I understand that I can revoke this designation at any time by providing written notice to Vital Care

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Date:

PRESCRIBER MUST MANUALLY SIGN - STAMP SIGNATURE, SIGNATURE BY OTHER PERSONNEL AND COMPUTER-GENERATED SIGNATURES WILL NOT BE ACCEPTED

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