

Pharmacy Name: Address: City/State/Zip:

Phone: Fax: Email:

Home Infusion Alternate Site of Care

Immunoglobulin Referral Form			
Patient Name		Home Phone	
Date of Birth		Mobile or Work Phone	
Patient Home Address		City State 2	Zip
Primary Insurance Name			
Primary Insurance ID	Primary Insurance Group		
Insured Name		Insured DOB	
Secondary Insurance Name		Insurance ID Insurance Group	
Secondary Insurance ID		Secondary Insurance Group	
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Ordering Physician's Name		NPI	
Address		City State 7	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 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 cyc	cles	time(s) per week for months.	
Other:	_		
Hydration order: mls NSiv to be infused pr	•		
Pre-medications: Acetaminophen 650mg PO 30 mins prior to infusion Other Pre-medications: Diphenhydramine 25mg PO 30 mins prior to infusion			
Clinical Information			
		I	
Patient Weight Height Allergies			
IV access [for IVIGg patients only]: Nurse to place PIV prior to therapy			
Diagnosis	ICD-10	Diagnosis	ICD-10
Neuromuscular:		Immune Deficiency:	
Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP)	G61.81	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Cell Disorders	D83.1
Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) Guillain-Barre Syndrome (GBS)		Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Cell Disorders Combined Immunodeficiency, Unspecified	
Neuromuscular: Chronic Inflammatory Demyelinating Polyneuropathy (CIDP)	G61.81 G61.0	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Cell Disorders	D83.1 D81.9
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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified 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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1
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	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89
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)	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies	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	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89
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	G61.81 G61.0 G61.82 G70.0 G70.01 G04.81 G61.89	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies	D83.1 D81.9 D83.9 D80.0 D80.5 D80.1 D81.89 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: 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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified 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	D83.1 D81.9 D83.9 D80.0 D80.5 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	Immune Deficiency: CVID w/ Predominant Immunoregulatory T-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified 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	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
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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified 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 Specific Antibody Deficiency	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	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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified 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	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
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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified 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 Specific Antibody Deficiency	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 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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified 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 Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE)	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 Please Draw: CBC/diff CMP IgG w/ subclasses 1-4 Quant. Ig	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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified Hereditary Hypogammaglobulinemia Immunodeficiency with Increased IgM Nonfamilial Hypogammaglobulinemia Other Combined Immunodeficiencies Other Common Variable Immunodeficiencies Pemphigoid 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	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 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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified 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 Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol:	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 Polymyositis Please Draw: CBC/diff CMP IgG w/ subclasses 1-4 Quant. Ig Frequency: Notes:	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-Cell Disorders Combined Immunodeficiency, Unspecified Common variable Immunodeficiency, Unspecified 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 Specific Antibody Deficiency Systemic Lupus Erythematosus (SLE) Anaphylaxis Protocol: PER Pharmacy Protocol:	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.

Date:

PRESCRIBER MUST MANUALLY SIGN - STAMP SIGNATURE, SIGNATURE BY OTHER PERSONNEL AND COMPUTER-GENERATED SIGNATURES WILL NOT BE ACCEPTED The attached document(s) contain confidential information which may be considered to be Protected Health Information and therefore required to be maintained as private and secure under HIPAA. The

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