Please fax all pages of completed form to your team at 808-650-6487.

To reach your team, call toll-free 808-650-6488.

Prescription & Enrollment Form Immune Globulin



Four simple steps to submit your referral.

1 Patient Information		provide copies of front and back of all medical scription insurance cards.
Patient's first name	Last name	Middle initial
Sex at birth: Male Female Pronouns	Last 4 digits of SSN	Date of birth
		Apt #
		Zip
Parent/guardian (if applicable)		Email address
2 Prescriber Informatio	n All fields must b	e completed to expedite prescription fulfillment.
Prescriber's first name	Last name	
icense #	NPI #	
		Suite #
•		Zip
		License #
	Fax	
3 Clinical Information		
Home infusion Clinic infusion	kg/lbs Date weight obtained dose due	
CD-10 Diagnosis Code (Required): ICD-10 immunology: D80.0 Cong ICD-10 neurology: G61.81 CIDP ICD-10 rheumatology: M33.20 Po		D81.9 SCID (unspecified) G61.0 GBS G70.01 MG
Allergies		
Bio3		
	Complete RX information on Page	2

_____ Last name _

Patient's first name _

ONE

Middle initial _____ Date of birth ___

based on information available, including acknowledge that all bran	ist preferred brand below clinical information, in: ds are clinically approp	nsurance requirements a	nd brand availal credo will comm	OWED" line to authorize "Pharmacist to select brand bility. By signing "SUBSTITUTION ALLOWED" you nunicate to you the brand selected. B PATIENTS	,,,
Route: Subcutaneous Intravenous Brand: Pharmacist to select brand Prescriber's preferred brand listed below required for Medicare B):	Once weekly Ev		4 weeks subc	bcutaneous: Infuse total dose of immune globulin utaneously in 1 to multiple subcutaneous sites of the fusion pump as tolerated. Infusion rates per ufacturer recommendation as tolerated.]
f Intravenous: Titrate initial and maintenanch nanufacturer's product labeling. 'ascular access: Peripheral Central nfusion method: Gravity Pump	De infusion rates per Port		9% Normal Salir	ne intravenously prior to infusion over 30 minutes given concurrent with IVIG at same rate as IG	
nyasthenia gravis) for pediatric patients the following weight an 9kg and/or <2 years old: 1mg/kg up to max cetaminophen 650mg by mouth (For pedia 9ther	nd aged based dosing rock of 6.25mg, 2-5 years atric patients weighing I hrough if not required): 6 hours as needed for masthenia gravis) site prior to needle insert	range will be used for a s old and >9kg: 6.25mg less than 60kg: Acetan): mild infusion reactions, ertion as needed to prev	II Diphenhydram to 12.5mg, 6-1 ninophen 10-15 may increase to ent site pain	12 years old: 12.5mg to 25mg mg/kg by mouth for all Acetaminophen prescribed 50mg for moderate to severe; maximum of 4 dos	_
dverse event medications (Keep on hand a pinephrine 0.3mg for patients weighing ≥3 naphylactic reaction times one dose piphenhydramine 25mg by mouth for mild a	Okg or 0.15mg for pation	ients weighing <30kg a	uto-injector 2-p	or with the first subcutaneous fill only): k. Administer intramuscularly as needed for severe	
Tushing for Intravenous: 0.9% Normal Sali leeded for line patency deparin 10 units per mL 3mL intravenous (deparin 100 units per mL 5mL intravenous)	(peripheral line) as nee	eded for final flush	L intravenous (d	central line/port) before and after infusion, or as	
Supplies: Dispense needles, syringes, ancilla Quantity/Refills: Dispense 1 month supply. Refill x 1 year unl Other	<u> </u>	medical equipment ne	cessary to admi	nister medication.	
ikilled Nursing: IVIG- Visit as needed to est	neous access, medicati	tion administration, use		eral status and response to therapy. SubQ IG- Skill erapy and disease state and to assess general statu	
shipped to physician's office or infusion clir "Pharmacist to select brand" option chescriber's signature required (sign belo	nosen above, sign the	e "SUBSTITUTION A	LLOWED" line	e below.	_
Source o Sibilatare required (Sign Delo					

accredo*

Non-compliance with state-specific requirements could result in outreach to the prescriber.

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The prescriber is to comply with his/her state-specific prescription requirements such as e-prescribing, state-specific prescription form, fax language, etc.



Prior authorization checklist Primary immune deficiency disease (PIDD)

Providing Accredo with the documentation outlined in this checklist may increase the likelihood and speed of obtaining coverage for your patients with PIDD. Coverage criteria may vary by payer.

Refe	Referral form ¹ (not required for electronic prescriptions)			
	Completed Immunoglobulin (Ig) referral form (available at accredo.com)			
	Copies of the front and back of all medical insurance and prescription benefits cards			
Clini	Clinical documents			
	History and Physical (H&P) and progress notes (within past 6 months) Note: H&P to include documented infection history/treatment			
	Pre-treatment IgG, IgA, IgM, and Ig subclass serum levels (drawn on two different occasions when available) Current IgG, IgA, IgM, and Ig subclass serum levels			
	Pre- and post-antigen testing (tetanus, pneumococcal polysaccharide or H Influenza type B) AND documentation of vaccine administration date			

D81.0 – Severe combined immunodeficiency	D82.0 - Wiskott-Aldrich syndrome	
(SCID) with reticular dysgenesis	D82.1 – Di George's syndrome	
D81.1 – Severe combined immunodeficiency		
(SCID) with low T- and B-cell numbers	D82.4 – Hyperimmunoglobulin E (lgE) syndror	
D81.2 – Severe combined immunodeficiency (SCID) with low or normal B-cell numbers	D83 – Common variable immunodeficiency (CVID)	
D81.5 - Purine nucleoside phosphorylase (PNP) deficiency	D83.0 – CVID with predominant abnormalities of B-cell numbers and function	
D81.6 - Major histocompatibility complex class I deficiency	D83.1 – CVID with predominant immunoreg	
D81.7 - Major histocompatibility	D83.2 - CVID with autoantibodies to B- or T-cells	
Complex class if deliciency	I-CEIIS	
D81.89 – Other combined immunodeficiencies	D83.8 - Other CVIDs	
D81.9 – Combined immunodeficiency.	2000 0000	
unspecified	D83.9 - CVID, unspecified	
D82 - Immunodeficiency associated	G11.3 - Cerebellar ataxia with defective	
with other major defects	DNA repair	
	(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 D81.5 - Purine nucleoside phosphorylase (PNP) deficiency D81.6 - Major histocompatibility complex class I deficiency D81.7 - Major histocompatibility complex class II deficiency D81.89 - Other combined immunodeficiencies D81.9 - Combined immunodeficiency, unspecified D82 - Immunodeficiency associated	

To receive in-home administration for intravenous immune globulin (IVIG) for the treatment of PIDD, Medicare Part B patients must be enrolled in the IVIG Demonstration initiative. For further information visit: https://med.noridianmedicare.com/web/ivig

Fax completed form to 866.233.7151.

If you have any questions, please call your Accredo Provider Support Advocate, or call 866.820.4844.

1. For referral forms visit accredo.com.



Prior Authorization Checklist Neuromuscular Disorders¹

Providing Accredo with the documentation outlined in this checklist may increase the likelihood and speed of obtaining coverage for your patients. Coverage criteria many vary by payer.

Refe	Referral Form (not required for electronic prescriptions)			
	Completed Immunoglobulin (Ig) referral form (available at accredo.com)			
	Copies of the front and back of all medical insurance and prescription benefits cards			
Clini	Clinical Documents			
	History and Physical (H&P) and progress notes ² (within past 6 months) Note: Diagnosis of the disorder must be unequivocal			
	Documentation that other causes of demyelinating neuropathy have been excluded			
	Testing documentation: □ Electrophysiological motor-sensory nerve conductions □ Electromyography (EMG) □ Cerebrospinal fluid (CSF) □ Biopsy (muscle-nerve) - if necessary			

Add	Additional Requirements for Myasthenia Gravis		
	Tensilon test results		
	Refractory to corticosteroids over a 6 month period documentation		
	Ongoing Ig treatment must be documented in H&P and progress notes ²		
Add	Additional Requirements for Polymyositis and Dermatomyositis Diagnosis		
	Creatine phosphokinase (CPK) values		
	Electromyography (EMG) and/or muscle biopsy results		

Fax completed form to 866.233.7151.

If you have any questions, please call your Accredo Provider Support Advocate, or call 866.820.4844.

¹ This Neuromuscular Disorders checklist is based on Medicare Part B guidelines related to Guillain-Barre' syndrome (GBS), relapsing-remitting multiple sclerosis, chronic inflammatory demyelinating polyneuropathy (CIDP) (and variant syndromes such as Multifocal Motor Neuropathy (MMN)), myasthenia gravis, refractory polymyositis, and refractory dermatomyositis polymyositis, and refractory dermatomyositis

² Ongoing management and documentation requirements:

[·] Initial improvement and continued need must be meticulously documented in progress notes

[·] All weaning must be attempted and documented as either amount or frequency

[·] Must be a stoppage in IVIG if sustained improvement is noted with weaning or no improvement has taken place at all