

### IVIG (INTRAVENOUS IMMUNOGLOBULIN)

Send completed form to: Service Benefit Plan Prior Approval P.O. Box 52080 MC 139 Phoenix, AZ 85072-2080 Attn. Clinical Services

	nation is required to process your contains and submit this completed form.	0	. Please complete the		rescribing physician co	mplete the	Attn. Clinical Services Fax: 1-877-378-4727	
Patient Information (required)				Provider Information (required)				
Date:				Provider Name:				
Patient Na	me:			Specialty:	Specialty:			
Date of Bi	rth:	Sex: ☐Male	□Female	Office Phone:	Office Phone: Office Fax:			
Street Add	ress:			Office Street Add	lress:			
City: State: Zip:				City:	S	State: Zip:		
Patient ID:	R	1 1 1		Physician Signatu	ıre:		1	
	1		PHYSICIAN	COMPLETES				
		NOTE: Form	must be comple	eted in its <b>entirety</b> fo	or processing			
Please sele	ct medication:		-		<del></del> _			
□Alyglo	□Bivigam	□Gamma	gard	□Gammaked	□Gamune	x-C	□Panzyga	
□Asceni	v □Flebogamma	□Gamma	ked S/D	□Gammaplex	□Octagan	1	□Privigen	
**Check ww	w.fepblue.org/formulary to	confirm which med	lication is part of	the patient's benefit				
1. Has the	patient been on this me	edication continu	ously for the la	st <b>6 months</b> , exclud	ing samples? P	lease selec	t answer below:	
□YES	– this is a PA renewal	for <b>CONTINUA</b>	TION of thera	py, please answer the	e questions on	PAGE 5		
□NO -	this is <b>INITIATION</b>	of therapy, please	e answer the qu	estions below:				
2. Is this r	equest for brand or gen	eric?   Brand	□Generic					
3. Will this	medication be filled at	a pharmacy or bill	led through med	lical plan or office bil	ling? □Pharma	ev $\square$ Me	edical plan/Office billing	
	medication be self-ad	•	_	-	C	J		
	Has the patient or their				ns and symptor	ns of thro	mbosis when self-	
	administering the med			C	, 1			
	Will the patient be mon $\square$ of $\square$ Yes $\square$ $\square$ N		or signs and sy	mptoms of thrombos	is both at the ti	me of infi	usion and after	
	s medication be used in <b>ES</b> , specify the medica		th another imm	unoglobulin medica	tion? □Yes*	□No		
6. What is	the patient's diagnosis	?						
	Alloimmune Thrombo		IT)	☐Multiple scle	erosis			
	sion-body myositis	•	,	•		ire Red C	ell Aplasia (PRCA)	
□Kawasaki syndrome				☐Peripheral Blood Progenitor Cell (PBPC) collection				
□Laml	pert-Eaton Myasthenic	(S)	☐Umbilical cord stem cell transplantation					
_	nmaglobulinemia							
	Has the patient's diagno			•		)		
b. 1	Does the patient have a			<u> </u>				
	a-telangiectasia OR Has the patient's diagno					,		
	Does the patient have a						O	
c. I	Does the patient have a	n impaired antibo	ody response to	the pneumococcal v	raccine?  \( \square\) Yes	□No		
	immune encephalitis							
	Has the diagnosis been buncture, or serologic to			lowing tests: neuroin	maging, electro	encephalo	ography (EEG), lumbar	
	Marrow Transplantations the medication being			oietic Stem Cell Tran acterial and viral infe			S	

PLEASE PROCEED TO PAGE 2 FOR ADDITIONAL DIAGNOSES

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\*If NO, does the patient have a pre-treatment serum lgG level less than 400 mg/dL? \(\sigma\)Yes \(\sigma\)No

b. Has the patient received a transplant in the last 100 days? □Yes □No\*



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PAGE 2 - PHYSICIAN COMPLETES					
Patient Name:	DOB:	Patient ID: R			
•	derate to severe functional disabil	lity? □Yes □No istent with multifocal demyelinating abnormalities? □Yes □N			
□Chronic Lymphocytic Leuken a. Does the patient have B-o	nia (CLL) cell chronic lymphocytic leukemia	a? □Yes □No			
c. Is there a documented his		erial and viral infections? □Yes □No infections requiring intravenous antibiotics or			
d. Does the patient have a p	re-treatment serum lgG level less	than 500 mg/dL? □Yes □No			
-	ocumented history of recurrent ba	acterial and viral infections? □Yes □No e pneumococcal vaccine? □Yes □No			
c. Have other causes of imm as HIV, or malignancy?	nune deficiency been excluded inc □Yes □No	cluding drug-induced, genetic disorders, infectious diseases such			
		n 500 mg/dL? □Yes □No* alent to 2 or more standard deviations below the mean for the ag			
□Dermatomyositis <u>OR</u> □I  a. Does the patient have doodiagnostic tests? □Yes	cumented clinical features such as	s elevated muscle enzymes, muscle biopsy, or supportive			
	intolerance or contraindication or osteroids or immunosuppressants?	have they had an inadequate treatment response to first-line Pyes  No			
☐End-Stage Renal Disease (ESI a. Will this medication be u		ccessful kidney transplantation? □Yes □No			
		requiring the patient to use an aid to walk? □Yes □No set of symptoms? □Yes □No			
□HIV infections a. Is the medication being p	rescribed for prophylaxis of bacte	erial and viral infections? □Yes □No			
•		ophylaxis? Please select answer below:			
□Primary Prophylaxis:	Does the patient have a pre-treat	ment serum lgG level less than 400 mg/dL? □Yes □No			
		uestions: acterial and viral infections (greater than 2 serious infections in a			
-	•	ombination antiretroviral therapy? □Yes □No ctive for this patient? □Yes □No			
		g/dL? □Yes □No* alent to 2 or more standard deviations below the mean for the			
_		acterial and viral infections? □Yes □No e pneumococcal vaccine? □Yes □No			

PLEASE PROCEED TO PAGE 3 FOR ADDITIONAL DIAGNOSES

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**PAGE 3 - PHYSICIAN COMPLETES** DOB: **Patient Name:** Patient ID: R □lgG subclass deficiency a. Does the patient have a pre-treatment lgG1, lgG2, or lgG3 equivalent to 2 or more standard deviations below the mean for the patient's age on at least two occasions? \(\sigma\)Yes \(\sigma\)No b. Does the patient have lgG (total) and lgM levels within normal limits? □Yes\* □No \*If YES, does the patient have IgA levels within low to normal limits? \(\sigma\)Yes \(\sigma\)No c. Does the patient have a documented history of recurrent bacterial and viral infections? \( \subseteq \text{Yes} \) d. Does the patient have an impaired antibody response to the pneumococcal vaccine? □Yes □No ■ Multifocal Motor Neuropathy (MMN) a. Is the patient experiencing weakness without \*objective sensory loss in two or more nerves? □Yes □No \*Objective Sensory Loss: decreased reflexes, motor weakness, muscle wasting, trophic skin, join changes b. Has the patient had electrodiagnostic studies that are consistent with motor conduction block? □Yes □No c. Has the patient had sensory nerve conduction studies that are normal? □Yes □No ■Myasthenia gravis a. Has the patient experienced an increase in any of the following symptoms: diplopia (double vision), ptosis (drooping eyelid), blurred vision, dysarthria (difficulty in speech), dysphagia (difficulty swallowing), difficulty chewing, impaired respiratory status, fatigue, or limb weakness? □Yes □No\* \*If NO, has the patient had \*pre-operative management?  $\square$ Yes  $\square$ No \*Pre-Operative Management: cholinesterase inhibitors, corticosteroids, or immunosuppressants □ Secondary immunosuppression a. Does the patient have secondary immunosuppression associated with hematological malignancy? \(\sigma\)Yes \(\sigma\)No b. Does the patient have a pre-treatment lgG level of less than 500mg/dL? □Yes □No\* \*If NO, does the patient have a pre-treatment lgG equivalent to 2 or more standard deviations below the mean for the patient's age? \(\begin{aligned} \Perc \text{Yes} & \Pi \text{No} \end{aligned}\) c. Does the patient have a documented history of recurrent bacterial and viral infections? ☐Yes ☐No d. Does the patient have an impaired antibody response to the pneumococcal vaccine?  $\square$ Yes  $\square$ No □ Selective lgA deficiency a. Does the patient have a pre-treatment lgA level of less than 7mg/dL? □Yes\* □No \*If YES, does the patient have  $\lg G$  and  $\lg M$  levels within normal limits?  $\square Yes$ b. Does the patient have a documented history of recurrent bacterial and viral infections? \( \subseteq \text{Yes} \) c. Does the patient have an impaired antibody response to the pneumococcal vaccine? \(\sigma\)Yes \(\sigma\)No □ Selective lgM deficiency a. Does the patient have a pre-treatment lgM level of less than 30mg/dL? \( \square\) Yes\* \*If YES, does the patient have lgG and lgA levels within normal limits? □Yes □No b. Does the patient have a documented history of recurrent bacterial and viral infections? \( \subseteq \text{Yes} \) c. Does the patient have an impaired antibody response to the pneumococcal vaccine? ☐Yes ☐No □ Severe Combined Immunodeficiency Disease (SCID) a. Does the patient have an absence or very low number of T cells (CD3 T cells less than 300/microliter)? □Yes □No\* \*If NO, is there a presence of maternal T cells in the circulation?  $\Box$ Yes  $\Box$ No b. Has the patient's diagnosis been confirmed by genetic or molecular testing? ☐Yes ☐No c. Does the patient have a pre-treatment lgG less than 200 mg/dL? □Yes □No □ Specific antibody deficiency a. Does the patient have a specific antibody deficiency with lgG, lgA, and lgM levels within normal limits? \(\sigma Y \)es b. Does the patient have a documented history of recurrent bacterial and viral infections? □Yes □No c. Does the patient have an impaired antibody response to the pneumococcal vaccine? □Yes □No

#### PLEASE PROCEED TO PAGE 4 FOR ADDITONAL DIAGNOSES

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PAGE 4 - PHYSICIAN COMPLETES					
Patient Name: DOB: Patient ID: R					
□Stiff-person syndrome a. Does the patient have an intolerance or contraindication or have they had an inadequate treatment response to first	t-line				
treatment (benzodiazepine or baclofen)?  \( \square\)Yes \( \square\)No					
☐ Idiopathic Thrombocytopenic Purpura (ITP)					
a. <b>FEMALE Patient</b> : Is the patient of reproductive potential? □Yes* □No					
*If YES, is the patient currently pregnant? \(\sigma\)Yes \(\sigma\)No					
b. Has the patient been diagnosed with ITP within the past three months? <i>Please select answer below:</i>					
☐Yes: Please answer the following questions:  i. Age 17 or Younger, please answer the following questions:					
<ol> <li>1) Does the patient have significant bleeding symptoms such as mucosal bleeding or moderate to seve bleeding?   No</li> </ol>	ere				
2) Is the patient at high risk for bleeding? □Yes □No					
3) Does the patient require a rapid increase in platelets due to a surgery or procedure? □Yes □No					
ii. Age 18 or Older, please answer the following questions:					
1) What is the patient's platelet count? <i>Please select answer below:</i> □Less than 30,000/mcL					
□30,000 to 49,999/mcL, please answer the following question:					
<ul> <li>a) Does the patient have significant bleeding symptoms, a high risk for bleeding, or a required rapid increase in platelets? □Yes □No</li> </ul>	nent for a				
□50,000/mcL or higher					
2) Will IVIG be used in combination with corticosteroid therapy? □Yes □No*					
* $If NO$ , does the patient have a contraindication to corticosteroid therapy? $\square$ Yes $\square$ No					
□No: Please answer the questions below:					
i. Is the patient experiencing refractory ITP following a splenectomy? <i>Please select answer below:</i>					
□Yes: Does the patient have a platelet count less than 30,000/mcL (per microliter)? □Yes □No*					
* $If NO$ , does the patient have significant bleeding symptoms? $\square$ Yes $\square$ No					
□No: Please answer the following questions:					
1) What is the patient's platelet count? <i>Please select answer below:</i>					
□Less than 30,000/mcL					
□30,000/mcL to 49,999/mcL, please answer the following question:	:				
<ul> <li>a) Does the patient have significant bleeding or is at high risk for bleeding or have a reconstruction for a rapid increase in platelets? □Yes □No</li> </ul>	quiremen				
□50,000/mcL or higher					
2) Has the patient had a relapse after a previous response to IVIG? ☐Yes ☐No					
3) Does the patient have an intolerance or contraindication or have they had an inadequate treat response to corticosteroid therapy? □Yes □No	tment				
☐Other non-SCID combined immunodeficiency (please specify):					
a. Has the patient's diagnosis been confirmed by genetic or molecular testing? □Yes □No					
b. Does the patient have a documented history of recurrent bacterial and viral infections?   No					
c. Does the patient have an impaired antibody response to the pneumococcal vaccine? □Yes □No					
Other diagnosis (please specify):					

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Patient Information (required)				Provider Information (required)			
Date:				Provider Name:			
Patient Name:			Specialty:		NPI:		
Date of Birth: Sex:		Sex:	□Female	Office Phone:		Office Fa	ix:
Street Address:				Office Street Address:			
City:	5	State:	Zip:	City: State:		Zip:	
Patient ID: R		1	Physician Signature:				
K [		P	HYSICIAN (	COMPLETES			
	CONTIN	JUATION	OF IVIG	THERAPY (PA	A RENEV	VAL)	
				d in its <b>entirety</b> for p		, ,	
Please select medic			<del>-</del>	· -	<del></del>		
□Alyglo	□Bivigam	□Gar	nmagard	□Gammaked	□Gamun	ex-C	□Panzyga
□Asceniv	□Flebogam	ma □Gar	nmaked S/D	□Gammaplex	□Octaga	m	□Privigen
**Check www.fepblue.	org/formulary to cor	nfirm which medic	ation is part of the	patient's benefit			
4. Will the medicat  □Yes: Has the padminist  □No: Will the p	ion be filled at a ph ion be self-admin patient or their car tering the medicat	harmacy or billed nistered? <i>Please</i> regiver been instion? □Yes □	select answer be tructed on how	al plan or office billing pelow: to monitor for signs a	nd symptoms	of thromb	
5. Will this medica			another immun	oglobulin medication	? □Yes* □	<b>l</b> No	
□Guillain-Barro □Idiopathic The □Inclusion-bod □Kawasaki syn □Lambert-Eato □Autoimmune	nune Thrombocyt e Syndrome (GBS rombocytopenic F y myositis drome n Myasthenic Syn encephalitis atient had a neuro	S) Purpura (ITP) ndrome (LEMS)	)	□Multiple sclerosis □Myasthenia gravis □Parvovirus B 19-i □Peripheral Blood □Stiff-person synda □Umbilical cord sta	s Induced Pure l Progenitor Ce rome em cell transp	ll (PBPC)	•
	Transplantation (			tic Stem Cell Transple			ents

PLEASE PROCEED TO PAGE 6 FOR ADDITIONAL DIAGNOSES

b. Has the patient had a documented reduction in frequency of bacterial and viral infections since initiation of therapy? ☐Yes ☐No

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Federal Employee Program. PRIOR APPROVAL REQUES!

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	PAGE 6 - PHYSICIA	N COMPLETES
Patient Name:	DOB:	Patient ID: R
□Chronic Inflammatory Demyelin	nating Polyneuropathy (CIDP)	
a. Is the IVIG being used at th	e lowest effective dose and fre	equency? \(\sigma\)Yes \(\sigma\)No
b. Have chronic stable patients necessary? □Yes □No	s been tapered and/or had treat	ment withdrawn to determine whether continued treatment is
<ul><li>c. Has the patient had a significant therapy? ☐Yes ☐No</li></ul>	cant improvement in disability	and maintenance of improvement since initiation of
☐Chronic Lymphocytic Leukemia	ı (CLL)	
a. Does the patient have B-cel	l chronic lymphocytic leukemi	a? □Yes □No
b. Has the patient had a docume	ented reduction in frequency of b	pacterial and viral infections since initiation of therapy? ☐Yes ☐No
□Dermatomyositis <u>OR</u> □Po	lymyositis	
a. Has the patient had a significant	cant improvement in disability	y and maintenance of improvement since initiation? □Yes □No
□End-Stage Renal Disease (ESRI	<b>)</b> )	
a. Will this medication be used	d to improve the chances of su	ccessful kidney transplantation? □Yes □No
□HIV infections		
a. Is the medication being pres	scribed for prophylaxis of bact	erial and viral infections? □Yes □No
b. Has the patient had a docume	ented reduction in frequency of b	pacterial and viral infections since initiation of therapy? ☐Yes ☐No
☐Multifocal Motor Neuropathy (N	MMN)	
a. Has the patient had a significant	cant improvement in disability	and maintenance of improvement since initiation? □Yes □No
□Primary Immunodeficiency Dise	ease (PID)	
a. Which type of primary imm	unodeficiency disease is this r	medication being used for? Please select answer below:
□Agammaglobulinemia	☐ IgG subclass deficiency	☐Common Variable Immunodeficiency Disease (CVID)
□Ataxia-telangiectasia	☐Selective lgA deficiency	□Severe Combined Immunodeficiency Disease (SCID)
☐DiGeorge syndrome ☐Hypogammaglobulinemia	☐ Selective lgM deficiency ☐ Specific antibody deficiency	☐Wiskott-Aldrich syndrome
		):
	h levels be monitored at least	yearly and maintained at or above the lower range of normal for
		pacterial and viral infections since initiation of therapy?   Yes  No
		'IG and, if necessary, reconsider a dose adjustment? □Yes □No
□ Secondary immunosuppression		
a. Does the patient have secon	dary immunosuppression asso	ciated with hematological malignancy? □Yes □No
_		ncy of bacterial and viral infections since initiation of
□Other diagnosis (please specify)	<b>:</b>	

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