SENTARA COMMUNITY PLAN (MEDICAID)

MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to 1-844-305-2331. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. If information provided is not complete, correct, or legible, authorization can be delayed.

Immune Globulin SQ [Primary Immunodeficiency Disorder] (Medical)

□ Cuvitru[®] [Immune Globulin Subcutaneous

Drug Requested: Check applicable box below. If not checked, authorization could be delayed.

□ Cutaquig® [Immune Globulin Subcutaneous

(Human) – hipp, 16.5% solution] (J1551)	(Human) 20% solution] (J1555)
□ Gammagard [®] [Immune Globulin Infusion (Human)] (J1569)	□ Gamunex-C [®] [Immune Globulin Injection (Human), 10% Caprylate/Chromatography Purified] (J1561)
□ Hizentra ® [Immune Globulin Subcutaneous (Human) 20% liquid] (J1559)	□ Hyqvia [®] [Immune Globulin Infusion 10% (Human) with Recombinant Human Hyaluronidase] (J1575) (AG)*
□ Xembify® [Immune Globulin Subcutaneous (H	uman) – klhw 20%] (J1558)
MEMBER & PRESCRIBER INFORMAT	TON: Authorization may be delayed if incomplete.
Member Name:	
Member Sentara #:	
Prescriber Name:	
Prescriber Signature:	Date:
Office Contact Name:	
Phone Number:	Fax Number:
DEA OR NPI #:	
DRUG INFORMATION: Authorization may	be delayed if incomplete.
Drug Form/Strength:	
Dosing Schedule:	Length of Therapy:
Diagnosis:	ICD Code, if applicable:
Weight:	Height:
Date:	

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□ Standard Review. In checking this box, the timeframe does not jeopardize the life or health of the member or the member's ability to regain maximum function and would not subject the member to severe pain.

<u>NOTE</u>: Dosing should be calculated using adjusted body weight if the patient's actual body weight is 20% higher than his or her ideal body weight (IBW). *IMPORTANT* - If recommended adjusted body weight is not accepted, only a <u>PARTIAL</u> approval will be granted.

- (Adjusted body weight = IBW + 0.5 (actual body weight IBW)
- IBW (kg) for males = 50 + [2.3 (height in inches -60)]
- IBW (kg) for females = 45.5 + [2.3 x (height in inches 60)]

CLINICAL CRITERIA: Check below all that as support each line checked, all documentation, including provided or request may be denied. Check the diagnost	g lab results, diagnostics, and/or chart notes, must be
Initial Authorization: 6 months	
☐ Severe combined immunodeficiency	☐ X-linked or autosomal recessive agammaglobulinemia
☐ Common variable immunodeficiency	□ Wiskott-Aldrich syndrome
□ CD40 ligand deficiency (X-linked hyper-IgM syndrome)	□ Nuclear factor of κβ essential modifier deficiency
☐ Ataxia-telangiectasia	□ DiGeorge Syndrome
☐ Is this member switching from IV to SQ IG for I	Primary immunodeficiency? Yes No
AND	
☐ For Hyqvia requests: Member must be ≥ 18 year	rs of age
AND	
•	isits required for hard-to-treat infections (e.g., recurrent skin abscess, deep seated infections) in the last 12
AND	
	escribed for hard-to-treat infections (e.g., recurrent ear abscess, deep seated infections) in the last 12 months:
AND	
☐ Member's IgG level is <200 mg/dL (submit doc	cumentation)
<u>AND</u>	

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Ц	Member has a history of multiple hard to treat infections as indicated by at least TWO of the following:
	☐ Four or more ear infections within 1 year
	☐ Two or more serious sinus infections within 1 year
	☐ Two or more months of antibiotics with little effect
	☐ Two or more pneumonias within 1 year
	☐ Recurrent or deep skin abscesses
	□ Need for intravenous antibiotics to clear infections
	☐ Two or more deep-seated infections including septicemia
	AND
	Member has a deficiency in producing antibodies in response to vaccination
	AND
	Titers were drawn before challenging with vaccination
	<u>AND</u>
	Titers were drawn between 4 and 8 weeks of vaccination
Rea criter	d be approved based on recent ER/hospital visits PLUS IVIG < 200 mg/kg within the last 3 months. uthorization (High Maintenance Therapy): 3 months only. Check below all that apply. All ria must be met for approval. To support each line checked, all documentation, including lab results, nostics, and/or chart notes, must be provided or request may be denied.
NOTI occurr apply	E: It is recommended to attempt to decrease/wean the dose for renewal requests when improvement has red and subsequently stop IVIG therapy if improvement is sustained with a dose reduction (this does not to authorizations for primary immunodeficiency as long as immunoglobulin levels are maintained in the priate range).
	Member has experienced disease response as evidenced by at least ONE of the following:
	□ Decrease in the frequency of infection
	□ Decrease in the severity of infection
	AND
	Number of hospital/ER admissions for hard-to-treat infections has <u>NOT</u> increased from baseline since beginning IVIG therapy
	<u>AND</u>
	TIND
ч	IgG level obtained within the last 30 days was therapeutic: 500-1200 mg/dL (submit documentation)

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PA IVIG SQ - PID (Medical)(Medicaid) (Continued from previous page)

	☐ IgG trough levels >1250 mg/dL warrants IgG dose decrease. Has physician considered decreasing the IVIG dose? If not, please provider rationale for continued use of initial dose:
M	edication being provided by: Please check applicable box below.
M	Location/site of drug administration:
	Location/site of drug administration:
	Location/site of drug administration: NPI or DEA # of administering location:
	Location/site of drug administration: NPI or DEA # of administering location: OR

**Use of samples to initiate therapy does not meet step edit/ preauthorization criteria. **

*Previous therapies will be verified through pharmacy paid claims or submitted chart notes. *

ability to regain maximum function.