

HyQvia (Immune Globulin Infusion 10%(Human) with Recombinant Human Hyaluronidase)

Provider Order Form rev 7/30/2025

PATIENT INFORMATION	Referral Status:	☐ New Referra	☐ Updated Or	der 🗆 Order Renewal		
Patient Name:		DOB:	Patient Ph	one:		
Patient Address:		P	atient Email:			
Allergies:		□ NKDA Wei	ght (lbs/kg):	Height (in/cm):		
Sex: ☐ M / ☐ F Date of Last Infusion:	Next Due Date	: Pr	eferred Location:			
DIAGNOSIS (Please provide ICD-10 code in	space provided)					
Primary Immunodeficiency:	• •	Chronic Inflamm	atory Demyelinatir	ng Polyneuropathy:		
Other:						
THERAPY ADMINISTRATION (Select one)	L	.ABORATORY ORD	DERS			
Chronic Inflammatory Demyelinating Polyneurop		☐ CBC w/ diff	☐ at each dose	□ every:		
Doses less than or equal to 0.4 g/kg can be administered without ramp-up Patients must be on stable doses of IVIG for 12 weeks before switching to		□ CMP □ Other:	☐ at each dose	□ every:		
HYQVIA		PRE-MEDICATION				
☐ Patients transitioning from IVIG tx, administer dose and frequency as the previous IV tx, after the	Try Q via at the same	☐ Tylenol ☐ 500mg / ☐ 650mg PO ☐ Loratadine 10mg PO ☐ Pepcid 20mg ☐ PO / ☐ IVP ☐ Benadryl ☐ 25mg / ☐ 50mg ☐ PO / ☐ IVP ☐ Solumedrol ☐ 40mg / ☐ 125mg IVP				
as indicated per the manufacturer.						
DoseGM subcutaneous	, _					
Frequency: □every 2 weeks/ □every 3 weeks□/e						
Primary Immunodeficiency			ing / Li 123ing ive			
☐ Patients transitioning from IVIG tx, administer H						
dose and frequency as the previous IV tx, after the initial dose ramp-		NURSING				
up as indicated per the manufacturer.		✓ Provide nursing care per Nursing Procedure, including Hypersensitivity Reaction Management Protocol and post-procedure observation				
Dose:GM subcutaneou Frequency: □every 3 weeks□/every 4 weeks	ISIY	Reaction Management	Trotocor and post-pro-	cedure observation		
rrequeriey. Elevery 5 weeks Elyevery 1 weeks	F	Ramp Up Schedule	!			
☐ New to SCIG treatment or transitioning from SCIG, administer HyQvia at 300mg/kg to 600mg/kg at 3- or 4-week intervals, after the initial ramp up as indicated by the manufacturer.		CIDP ramp up		g Ramp up if switching from SCIG		
		Wk 1 -no tx	from IVIG Wk 1 total gmX0.25	or new to SCIG 3 weeks 4 weeks		
Dose:GM subcutaneou	1.	Vk 2&3 total gm x0.25	-	Wk 1 total Wk1 total		
Frequency: ☐ every 3 weeks/ ☐ every 4 weeks.	,	WK 2&3 LOLAI gm XU.25	WK Z total gmx0.5	gmX0.33 gmx0.25		
ADDITIONAL ORDERS	٧	Vk 4 total gm X0.50	Wk 4 total gmX0.75	Wk 2 total Wk2 total gmx0.67 gmx0.5		
		Vk 6 total gm x0.75	Wk 7 total dose	Wk 4 total dose		
		Vk 9 total dose		gmX0.75 Wk 7 total dose		
		The state accept				
PROVIDER INFORMATION						
Preferred Contact Name:		Preferred Contact Email:				
Ordering Provider:		Provider NPI:				
Referring Practice Name:	Pho	_	Fax:			
Practice Address:	City	/ :	State:	Zip Code:		
REQUIRED DOCUMENTATION CHECKI						
Required Documentation: Patient demos, co		imary and second	lary insurance, 2 m	nost recent OVN including		
treatment failures or contraindications. EMG	` '					
Required Labs: Immunoglobulin levels, Renal	TUNCTION, CRP/ESK, ANA,					
Provider Name (print)	Provider Signature			Date		

Disclaimer: By signing this form, I authorize Novella Infusion and its affiliates to act as my designated agent in submitting prior authorizations, financial assistance applications, and other clinically required information with respect to this patient and order. This enrollment form shall serve as my signature for prior authorizations and financial assistance programs, as requested.

Order valid for one year unless otherwise indicated. IV solutions/diluents may be substituted as allowed per manufacturer's instructions as necessitated by product availability.