REVCOVI

Revised 09/08/23



InfusionForHealth.com Ph: 888-777-1945 | Fax: 805-852-2636

Date:	Treatment L	_ocation:
		ent Demographic & Insurance Information, Medication List gh dAXP levels, and/or total lymphocyte counts
PATIENT INFORMATION		PRESCRIBER INFORMATION
Patient Name:		Prescriber's Name:
Patient Contact Number:		Signature:
DOB:		NPI: Date:
		Phone: Fax:
		Office Address:
	Trough plasma ADA activity, tr	Contact Person:
	o.oo,	Contact Email:
	col establishe	ed by Infusion for Health and P.I.
**		
REVCOVI DOSAGE		
REVCOVI DOSAGE Date of Last Treatment, If Continuation:		
	Patient Weig The starting of Route: IM	dose of REVCOVI is 0.2 mg/kg weekly
Date of Last Treatment, If Continuation:	The starting of Route: IM The starting of on ideal body	dose of REVCOVI is 0.2 mg/kg weekly Dosage: dose of REVCOVI is 0.4 mg/kg weekly based y weight or actual body weight (whichever is ded in two doses (0.2 mg/kg twice a week)
Date of Last Treatment, If Continuation: Patients transitioning from Adagen to REVCOVI Adagen-naïve patients To ensure that a brand name product be	The starting of Route: IM The starting of on ideal body greater), divided Route: IM Adispensed, the pre-	dose of REVCOVI is 0.2 mg/kg weekly Dosage: dose of REVCOVI is 0.4 mg/kg weekly based y weight or actual body weight (whichever is
Date of Last Treatment, If Continuation: Patients transitioning from Adagen to REVCOVI Adagen-naïve patients To ensure that a brand name product be prescription form. If not indicated, Lab Orders: Trough plasma ADA activity (pre-injection of the continuation) Trough plasma ADA activity (pre-injection of the continuation)	The starting of Route: ✓ IM The starting of on ideal body greater), divided Route: ✓ IM **Route: ✓ IM** **Adispensed, the prediction for Health** **The starting of the s	dose of REVCOVI is 0.2 mg/kg weekly Dosage: dose of REVCOVI is 0.4 mg/kg weekly based y weight or actual body weight (whichever is ded in two doses (0.2 mg/kg twice a week) Dosage: escriber must handwrite "Brand Medically Necessary" on h is authorized to administer generic or biosimilar. veeks x 12 weeks every 4 weeks x 8 weeks nonths or Health to order and draw labs indicated for clinical