Prescriber Signature (Stamps not acceptable; dispense as written) DATE





1. PATIE	NT INFORMATION					
Full Name				Caregiver (First, Last)		
DOB (MM	DOB (MM/DD/YYYY)			Relationship to Patient		Phone
				By providing the names of my other Care Team Members on this form (healthcare providers other than the GATTEX prescribing physician), I am authorizing any employees of the Companies to follow up with these Care Team Members to provide education and information about GATTEX.		
Address	Address					
City/Sta	ity/State/ZIP			HCP Care Team Member* (First, Last)		
Primary	rimary Phone Secondary Phone			Care Team Role		Phone
Special I	ecial Precautions (eg, allergies)			Español es mi primer idioma		*Optional.
E I wo	uld like to opt in to marketing (	communications				
	<b>t Authorization</b> d, understand, and agree to the releas	e of my protected health information	, as described on P	age 2, Section 6 of this for	m.	
1 1	Patient signature/legal representative signature (indicate relationship)					DATE
I have rea	th Patient Support Program ad, understand, and agree to the use of	my personal information for the pur	poses described on	n Page 2, Section 7 of this f	orm.	
Patie	ent signature/legal representati	ve signature (indicate relation	ship)			DATE
	RANCE INFORMATION					
	ED: Include copies of both sides		•	<del></del>	•	
-	rimary Insurance Insurance Phone		Secondary Insurar			Insurance Phone
,	licy ID # Group			*		Group
Policy H	olicy Holder Name (First, Last)			Policy Holder Name (First, Last)		
-	·	YY) Relationship to Patient			DOB (MM/DD/YYYY) Relationship to Patient	
Pharma	acy Plan Policy ID #			Group #		
Pharma	Pharmacy Plan Phone Rx Bin #			Rx PCN #		
	. PRESCRIBING PHYSICIAN INFORMATION			4. PATIENT CLINICA	L INFORMATION	
Full Nam	ne	Treatment Center		Diagnosis*		Etiology
	8	Fax ne Office Contact Name hone Office Contact Email		New Start Short bowel syndrome (SBS) patient dependent on parenteral nutrition and/or IV fluids (parenteral support) Existing Patient GATTEX renewal *Please do not check a box if neither applies. Date of Last Intestinal Resection		Inflammatory Bowel Disease (IBD) (eq, chronic conditions such as Crof
City/Sta	ite/ZIP					disease)
Phone _						Non-IBD (eg, acute events [vascular event,
щ.						trauma, intestinal obstruction], congenital anomaly [gastroschisis, midgut volvulus])
Office C	ontact Phone					Parenteral Support
Mational	ational Provider ID		ICD-10 Code		Provider/Pharmacy	
- F PPFO	<b>CRIPTION FOR GATTEX (teduglu</b>	tide) FOR INJECTION				
E 5. PRESI	on the control of the	ilde) I OK INOLOTION				n‡o.
The pres	scriber must comply with state s Calculate patient dosage (checl	pecific prescription requirement one box below)				of 30-vial kits needed
•	scriber must comply with state s Calculate patient dosage (checle: 0.05 mg/kg once daily (5 mg ki uce dose to 0.025 mg/kg once dail disease (estimated glomerular t	pecific prescription requirement one box below)  t is not recommended in patientials;  The patient has moderate or settings.	nts weighing less evere renal impa	s than 10 kg) irment or end-stage	STEP 2: Choose #	of 30-vial kits needed e than 3.8 mg/day, two 30-vial kit
•	scriber must comply with state s Calculate patient dosage (checle: 0.05 mg/kg once daily (5 mg ki uce dose to 0.025 mg/kg once daily	pecific prescription requirement one box below)  t is not recommended in patientials;  The patient has moderate or settings.	nts weighing less evere renal impa	s than 10 kg) irment or end-stage	STEP 2: Choose #	of 30-vial kits needed e than 3.8 mg/day, two 30-vial kit
•	scriber must comply with state s Calculate patient dosage (checle: 0.05 mg/kg once daily (5 mg ki uce dose to 0.025 mg/kg once dail disease (estimated glomerular t	pecific prescription requirement one box below)  t is not recommended in patientials;  The patient has moderate or settings.	nts weighing less evere renal impa	s than 10 kg) irment or end-stage m²)	If dose is more are recommer  One (1) 30-Vial  †A maximum of 0.3	of 30-vial kits needed e than 3.8 mg/day, two 30-vial kit
•	coriber must comply with state space (check calculate patient dosage (check et 0.05 mg/kg once daily (5 mg kill uce dose to 0.025 mg/kg once dail disease (estimated glomerular to both calculations	coecific prescription requirement one box below)  t is not recommended in patiential patiential has moderate or seritration rate [eGFR] less than the moderate of the moderate	nts weighing less evere renal impa 60 mL/min/1.73	s than 10 kg) irment or end-stage m²)  (mg/day)	If dose is more are recommer  One (1) 30-Vial †A maximum of 0.3 of teduglutide, can	of 30-vial kits needed e than 3.8 mg/day, two 30-vial kit nded†  Kit/NDC # 68875-0102-01/Vial Size: 5 mg 8 mL of the reconstituted solution, containing 3.8 mg
Complete	coriber must comply with state space (check calculate patient dosage (check et 0.05 mg/kg once daily (5 mg kind uce dose to 0.025 mg/kg once dail disease (estimated glomerular to both calculations  patient weight (kg)	coecific prescription requirement (one box below) t is not recommended in patient (aily: Patient has moderate or serification rate [eGFR] less than (aily: Multiply by 0.05 QR 0.025 per above	ets weighing less evere renal impa 60 mL/min/1.73 patient dose	s than 10 kg) irment or end-stage m²)  (mg/day)	If dose is more are recommer  One (1) 30-Vial †A maximum of 0.3 of teduglutide, can	of 30-vial kits needed e than 3.8 mg/day, two 30-vial kit nded†  Kit/NDC # 68875-0102-01/Vial Size: 5 mg 18 mL of the reconstituted solution, containing 3.8 mg 18 be withdrawn from the vial for dosing.
The pres STEP 1: Dose Redu rena Complete  STEP 3: Adminis	cariber must comply with state space of the calculate patient dosage (check et al. 0.05 mg/kg once daily (5 mg king with the calculations)  patient weight (kg)  patient weight (kg)  Enter directions	coecific prescription requirement ( one box below)  t is not recommended in patient ailly: Patient has moderate or seriltration rate [eGFR] less than (eGFR] le	patient dose  volume (m	s than 10 kg) irment or end-stage m²)  (mg/day)  L/day)  e subcutaneously, und	If dose is more are recommer  One (1) 30-Vial †A maximum of 0.3 of teduglutide, can Two (2) 30-Vial er the skin, once daily	of 30-vial kits needed e than 3.8 mg/day, two 30-vial kit nded†  Kit/NDC # 68875-0102-01/Vial Size: 5 mg 8 mL of the reconstituted solution, containing 3.8 mg be withdrawn from the vial for dosing.  I Kits/NDC # 68875-0102-01/Vial Size: 5 mg

Prescriber Signature (Substitution permitted )

DATE

## **Authorization for OnePath Services**

PLEASE READ THROUGH THE LANGUAGE ON THIS PAGE BEFORE SIGNING THE AUTHORIZATION AND CONSENT IN SECTION 1 OF THE START FORM.

## 6. PATIENT OR LEGAL GUARDIAN AUTHORIZATION TO SHARE PROTECTED HEALTH INFORMATION

By signing the Patient Authorization section of the Start Form, I authorize any health plan, physician, healthcare professional, hospital, clinic, pharmacy provider or other healthcare provider (collectively, "Providers") to disclose my, or my child's (as applicable), protected health information, including personal information relating to my, or my child's, medical condition, treatment, care management, and health insurance, as well as all information provided on this form and any prescription ("Information"), to Takeda Pharmaceuticals U.S.A., Inc., its affiliates and their representatives, agents, and contractors (collectively, the "Company") in connection with the Company's provision of products, supplies, or services. I understand the Company will provide this Information to a pharmacy within the GATTEX specialty pharmacy network. This Information may also be used for internal uses by the Company, including data analysis. I understand that Providers may receive financial remuneration from Company for marketing services.

Further, the Company may use this Information for OnePath Product Support Services such as verification of insurance benefits and drug coverage, prior authorization support, financial assistance with co-pays, patient assistance programs, alternate funding sources, other related programs, communication with me or my prescribing physician (or my child's) by mail, email, or telephone about my, or my child's, medical condition, treatment, care management, product information, and health insurance.

I understand that employees of the Company only see my, or my child's, Personal Health Information in connection with administering the OnePath Product Support Program, or in connection with other activities referenced herein, or as otherwise required or allowed under the law. I understand they will make every effort to keep my, or my child's Information private, but once my, or my child's, Personal Health Information is disclosed under this Authorization, it may no longer be protected by federal privacy law and subject to re-disclosure. I understand that I am entitled to a copy of this Authorization. I understand that I may cancel this Authorization at any time by sending written notice of revocation to OnePath, 300 Shire Way, Lexington, MA 02421. I understand that such revocation will not apply to any Information already used or disclosed through this Authorization. This Authorization will expire within five (5) years from today's date, unless a shorter period is provided for by state law. I understand that I may refuse to sign this Authorization and that refusing to sign this Authorization will not change the way my, or my child's, physician, health insurance, and pharmacy providers treat me or my child. I also understand that if I do not sign this Authorization, I, or my child, will not be able to receive OnePath Product Support Program products, supplies, or services.

## 7. ONEPATH AND COMMUNICATIONS ENROLLMENT

By signing the OnePath Patient Support Program and Communication Enrollment section on the first page of this Start Form, I am electing to enroll in OnePath Product Support Services (which may include, but is not limited to, verification of insurance benefits and drug coverage, prior authorization support, financial assistance with co-pays, patient assistance programs, alternate funding sources, other related programs, communication with me or my prescribing physician (or my child's) by mail, email, or telephone about my or my child's medical condition, treatment, care management, product information, and health insurance).

By checking the box on Page 1 labeled "I would like to opt in to marketing communications," I consent to receiving marketing and promotional communications from Takeda. I hereby give consent to Takeda, its affiliates, and their agents and representatives to send communications and information to me via the contact information I have provided above. I understand that this consent will be in effect until such time as I opt out of communications from Takeda.

I understand that I may revoke my permission at any time. To learn how Takeda will use and protect my personal information, please review our Privacy Policy (<a href="https://www.takeda.com/en-us/privacy-policy">www.takeda.com/en-us/privacy-policy</a>).

## Please click here for full **Prescribing Information**.



