



ADULT STATEMENT OF MEDICAL NECESSITY

Patient nformation	Patient Name (First and Last)		Date	of Birth (DOB)_				
	Address		Social Security Number					
	CityState		Allergies None Other					
	Home Phone Work Phone							
	Referring Physician							
	Physician Provider/Tax ID #		Office Contact					
nsurance nformation/ Attach Copy of Insurance Card	Primary Insurance							
	Insurance Company Phone							
	SubscriberDOB							
	Subscriber ID #							
	Policy/Employer/Group #							
Diagnosis	☐ Isolated Growth Hormone Deficiency (253.3) ☐ latrogenic Hypopituitarism (253.7) - hypophysectomy-induced - postablative - radiotherapy-induced	☐ Panhypopitui ☐ Other (specif	itarism (253.2) ry by ICD-9 Code)					
Medical Assessment	Current Weightlbskg							
	Growth Hormone Stimulation Test Date		Serum IGF-I Tes					
	Agent 1 Peak				esult			
	Agent 2 Peak							
Prescription Options for Genotropin choose A, B, C, or), plus choose pen needle or insulin syringe size)	☐ A. Genotropin PEN® 5 Growth Hormone Delivery Device (dose in increments of 0.1 mg)	5.8 mg G (5 mg/ml	Genotropin	Pen Needle G	àauge			
	□ B. Genotropin PEN® 12 Growth Hormone Delivery Device (dose in increments of 0.2 mg)		Genotropin nL)	Pen Needle Gauge				
	(dose in increments of 0.2 mg) C. Genotropin MIXER® Growth Hormone Reconstitution Device	DE C C C C C C C C C C C C C C C C C C C		Insulin Syringes □ 0.3 mL □ 0.5 mL □ 1.0 mL Needle Gauge				
	□ D. Genotropin MINIQUICK® is available in 10 strengt 0.25 mL. A 30-gauge, 5/16" injection needle is □ 0.2 mg □ 0.4 mg □ 0.6 mg □ 0.8	prepacked with eac	ch device. Please	select strength.				
		mg 🗆 1.0 mg	1.2 mg					
Dose to Be Given	Daily Dosemg/day Days Supply	_ Refills(mor	nths) Start Date	Shi	p Product by			
Subcutaneously	Weekly Dosemg/kg/week Prescription	on Special Instruction	ons					
Special nstructions if applicable)	Preferred Pharmacy				☐ Case Management Not Requested			
	Other			 	☐ Patient Device Training Requested			
Physician Certification	1) I certify that the treatment listed above is and will be medically and accurate to the best of my knowledge. 2) I also certify that I ha information here and such other health or personal information to participation in the Program. (A signed copy of a Pfizer Bridge Promedical Necessity or, to the best of the undersigned's knowledge, information only in accordance with the Authorization. 3) I further of in exchange for any express or implied agreement or understanding and (b) my decision to prescribe Genotropin was based on my decreated from the program of the progra	ave obtained the writter the Pfizer Bridge Progr gram Patient Authoriz is already on file with the tertify that (a) any serving that I would recommend the programment of medical booth hormone stimulation.	n permission of the pram" ("the Program"), ation Form ["the Auth the Pfizer Bridge Progree provided through thend, prescribe, or us necessity as set forth on testing or by other	atient (or the patier Pfizer, and/or its a orization'] either ac gram™.) I understan he Pfizer Bridge P e Genotropin⁰ or a herein. 4) I certify organic/clinical evi	nt's legal representative) to disclose the gents as may be necessary for the patient's companies this completed Statement of d that the Program may use and disclose this rogram" on behalf of any patient is not made ny other Pfizer product or service for anyone, that if I have prescribed the treatment for adult dence of aGHD (such as the lack of a			
	Print Name		Date Provider ID (NPI) DEA #					
	Address			.,	Ctate ZID			

 ${}^\star\!\mathsf{This}$ form cannot be processed without physician's signature.

Office Contact_



Fax

Phone



Pfizer Bridge Program™ Fax Number: **1-800-479-2562**Pfizer Bridge Program Phone Number: **1-800-645-1280**

Documentation Required for SMN Submission

Diagnosis	History and physical	Related clinical notes	Growth chart	Growth velocity	Birth wt/ length/ gestational age	Stim test results	IGF-I/ IGF-BP3 report	Bone age X-ray report	Genetic testing report	Comment
Adult GHD	/	/				*	/			*2 failed