Prior Authorization Request Form for mirabegron tablets (Myrbetriq)



JOHNS HOPKINS HEALTHCARE

7231 Parkway Drive, Suite 100, Hanover, MD 21076

FAX Completed Form and Applicable Progress Notes to: (410) 424-4037

USFHP Pharmacy Prior Authorization Form

To be completed by Requesting provider				
Drug Name:	Strength:			
Dosage/Frequency (SIG):	Duration of Therapy:			

Questions? Contact the Pharmacy Dept at: (888) 819-1043, option 4

Clinical Documentation must accompany form in order for a determination to be made.

Step	Ple	Please complete patient and physician information (please print):		
1	Pa	Patient Name: Physician Name:		
	Ad	ddress: Address:		
	Sponsor ID #		Phone #:	
		Date of Birth: Secure Fax #:		
Step	Ple	ease complete the clinical assessment:		
2	1.	Does the patient have a confirmed diagnosis of overactive bladder with symptoms of urge	☐ Yes	□ No
		incontinence, urgency, and urinary frequency?	Proceed to question 2	Proceed to question 8
	2.	Has the patient tried and failed behavioral	T Var	D No.
		interventions to include pelvic floor muscle training	☐ Yes	□ No
		in women, AND bladder training?	Proceed to question 3	STOP
				Coverage not approved
	3.	Has the patient had a 12-week trial of ONE of the following medications AND experienced therapeutic	□ Yes	□ No
		failure?	Proceed to question 5	Proceed to question 4
		 tolterodine extended-release (Detrol LA) 		
		oxybutynin IRoxybutynin ER		
		•trospium (Sanctura)		
		solifenacin (Vesicare)darifenacin (Enablex)		
		•fesoterodine (Toviaz)		
	4. Has the patient experienced central nervous system (CNS) adverse effects with an oral overactive bladder	☐ Yes	□ No	
		(OAB) medication or is at increased risk for CNS	Proceed to Question 5	STOP
		adverse effects due to comorbid conditions, advanced age or other medications?		Coverage not approved
	5.	Is the patient's estimated glomerular filtration rate (eGFR) available? If so please provide the eGFR.	14 14 70 0	
		Note: eGFR must be greater than or equal to 15 mL/	mL/min/1.73m2	☐ eGFR not available
		min/1.73m2 for coverage of Myrbetriq	Proceed to Question 7	Proceed to Question 6
	_			

Prior Authorization Request Form for mirabegron tablets (Myrbetriq)

	6. What is the patient's serum creatinine (SCr), weight, and height? Note: CrCl must be greater than or equal to 15	mg/dL ormm		
	mL/min/1.73m2 for coverage of Myrbetriq	inches A	AND lbs	
		milches A	ANDIDS	
		Proceed to Question 7		
	7. Is the provider aware that the dosage of Myrbetriq should not exceed 25 mg daily	□ Yes	□ No	
	when the Crcl/ glomerular filtration rate	Sign and Date Below	STOP	
	(eGFR is between 15-29 mL//min/1.73m2?		Coverage not approved	
	8. Does the patient have a diagnosis of neurogenic detrusor overactivity (NDO) secondary to detrusor overactivity and/or	☐ Yes	□ No	
		Proceed to Question 9	STOP	
	myelomeningocele?		Coverage not approved	
_	9. Is the medication being prescribed by or in	☐ Yes	□ No	
	consultation with a urologist or nephrologist?	Proceed to Question 10	STOP	
			Coverage not approved	
	10. Does the provider acknowledge that the	☐ Ackr	nowledged	
	granules are not bioequivalent to and cannot be substituted on a mg to mg basis	Proceed to question 11		
	to the tablets and will not combine dosage forms to achieve a specific dose?		- 1	
	11. Does the provider acknowledge that there	☐ Ackr	☐ Acknowledged	
	are detailed renal and hepatic dose adjustments in the package labeling and	Proceed to question 12		
	agrees to consult this before prescribing in these special populations?			
	12. Does the provider acknowledge that	☐ Acknowledged		
	oxybutynin is available for patients with neurogenic detrusor overactivity and does not require prior authorization?	Proceed to	Proceed to question 13	
	13. Has the patient tried and failed or had a	☐ Yes	□ No	
	contraindication to oxybutynin?	Proceed to Question 14	STOP	
			Coverage not approved	
14.		☐ Yes	□ No	
	to 35 kg?	Sign and date below	STOP	
			Coverage not approved	
ер го 3	14. Does the patient weigh greater than or equal to 35 kg? Certify the above is true to the best of my knowledge	Sign and date below	STOP	
	Prescriber Signature	Date		
			[30 November 202	
Interr	nal Use Only	,		
pprov	ved:	Duration of Ap	proval:month(s)	
enied	d:	Authorized By:		
comr	plete/Other:	PA#:	PA#:	