

# US Family Health Plan

## Prior Authorization Request Form for mirabegron tablets (**Myrbetriq**)

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD.

The completed form may be faxed to 855-273-5735

OR

The patient may attach the completed form to the prescription and **mail** it to:

**Attn: Pharmacy, 77 Warren St, Brighton, MA 02135**

QUESTIONS? **Call 1-877-880-7007**

**Please note that there is a separate prior authorization form for Myrbetriq granules.**

**Step 1** Please complete patient and physician information (please print):

Patient Name: _____ Address: _____ Sponsor ID #: _____ Date of Birth: _____	Physician Name: _____ Address: _____ Phone #: _____ Secure Fax #: _____
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**Step 2** Please complete the clinical assessment:

<b>1. Does the patient have a confirmed diagnosis of overactive bladder with symptoms of urge incontinence, urgency, and urinary frequency?</b>	<input type="checkbox"/> Yes Proceed to question <b>2</b>	<input type="checkbox"/> No Proceed to question <b>8</b>
<b>2. Has the patient tried and failed behavioral interventions to include pelvic floor muscle training in women, AND bladder training?</b>	<input type="checkbox"/> Yes Proceed to question <b>3</b>	<input type="checkbox"/> No <b>STOP</b> Coverage not approved
<b>3. Has the patient had a 12-week trial of ONE of the following medications AND experienced therapeutic failure?</b> <ul style="list-style-type: none"> <li>• tolterodine extended-release (Detrol LA)</li> <li>• oxybutynin IR</li> <li>• oxybutynin ER</li> <li>• trospium (Sanctura)</li> <li>• solifenacin (Vesicare)</li> <li>• darifenacin (Enablex)</li> <li>• fesoterodine (Toviaz)</li> </ul>	<input type="checkbox"/> Yes Proceed to question <b>5</b>	<input type="checkbox"/> No Proceed to question <b>4</b>
<b>4. Has the patient experienced central nervous system (CNS) adverse effects with an oral overactive bladder (OAB) medication or is at increased risk for CNS adverse effects due to comorbid conditions, advanced age or other medications?</b>	<input type="checkbox"/> Yes Proceed to Question <b>5</b>	<input type="checkbox"/> No <b>STOP</b> Coverage not approved
<b>5. Is the patient's estimated glomerular filtration rate (eGFR) available? If so please provide the eGFR.</b> Note: eGFR must be greater than or equal to 15 mL/min/1.73m2 for coverage of Myrbetriq	_____ mL/min/1.73m2 Proceed to Question <b>7</b>	<input type="checkbox"/> eGFR not available Proceed to Question <b>6</b>

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<p><b>6. What is the patient's serum creatinine (SCr), weight, and height?</b></p> <p>Note: CrCl must be greater than or equal to 15 mL/min/1.73m<sup>2</sup> for coverage of Myrbetriq</p>	<p>_____ mg/dL or _____ mmols/L</p> <p>_____ inches AND _____ lbs</p> <p>Proceed to Question <b>7</b></p>	
<p><b>7. Is the provider aware that the dosage of Myrbetriq should not exceed 25 mg daily when the CrCl/ glomerular filtration rate (eGFR is between 15-29 mL/min/1.73m<sup>2</sup>?</b></p>	<p align="center"><input type="checkbox"/> Yes</p> <p align="center"><b>Sign and Date Below</b></p>	<p align="center"><input type="checkbox"/> No</p> <p align="center"><b>STOP</b></p> <p align="center">Coverage not approved</p>
<p><b>8. Does the patient have a diagnosis of neurogenic detrusor overactivity (NDO) secondary to detrusor overactivity and/or myelomeningocele?</b></p>	<p align="center"><input type="checkbox"/> Yes</p> <p align="center">Proceed to Question <b>9</b></p>	<p align="center"><input type="checkbox"/> No</p> <p align="center"><b>STOP</b></p> <p align="center">Coverage not approved</p>
<p><b>9. Is the medication being prescribed by or in consultation with a urologist or nephrologist?</b></p>	<p align="center"><input type="checkbox"/> Yes</p> <p align="center">Proceed to Question <b>10</b></p>	<p align="center"><input type="checkbox"/> No</p> <p align="center"><b>STOP</b></p> <p align="center">Coverage not approved</p>
<p><b>10. Does the provider acknowledge that the granules are not bioequivalent to and cannot be substituted on a mg to mg basis to the tablets and will not combine dosage forms to achieve a specific dose?</b></p>	<p align="center"><input type="checkbox"/> Acknowledged</p> <p align="center">Proceed to question <b>11</b></p>	
<p><b>11. Does the provider acknowledge that there are detailed renal and hepatic dose adjustments in the package labeling and agrees to consult this before prescribing in these special populations?</b></p>	<p align="center"><input type="checkbox"/> Acknowledged</p> <p align="center">Proceed to question <b>12</b></p>	
<p><b>12. Does the provider acknowledge that oxybutynin is available for patients with neurogenic detrusor overactivity and does not require prior authorization?</b></p>	<p align="center"><input type="checkbox"/> Acknowledged</p> <p align="center">Proceed to question <b>13</b></p>	
<p><b>13. Has the patient tried and failed or had a contraindication to oxybutynin?</b></p>	<p align="center"><input type="checkbox"/> Yes</p> <p align="center">Proceed to Question <b>14</b></p>	<p align="center"><input type="checkbox"/> No</p> <p align="center"><b>STOP</b></p> <p align="center">Coverage not approved</p>
<p><b>14. Does the patient weigh greater than or equal to 35 kg?</b></p>	<p align="center"><input type="checkbox"/> Yes</p> <p align="center"><b>Sign and date below</b></p>	<p align="center"><input type="checkbox"/> No</p> <p align="center"><b>STOP</b></p> <p align="center">Coverage not approved</p>

**Step 3** I certify the above is true to the best of my knowledge. Please sign and date:

\_\_\_\_\_

Prescriber Signature

\_\_\_\_\_

Date