

US Family Health Plan

Prior Authorization Request Form for Liraglutide injection (Saxenda)

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

Clinical documentation may be required. Failure to provide could result in denial.

Initial therapy approves for 12 months; annual renewal is required. For renewal of therapy an initial prior authorization approval is required.

Non FDA-approved uses are not approved including diabetes mellitus

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

1	Patient Name: _____	Physician Name: _____
	Address: _____	Address: _____
	Sponsor ID #: _____	Phone #: _____
	Date of Birth: _____	Secure Fax #: _____

Step 2 Please complete the clinical assessment:

2	1. Has the patient received this medication under the TRICARE benefit in the last 6 months? Please choose "No" if the patient did not previously have a TRICARE approved PA for Saxenda.	<input type="checkbox"/> Yes (subject to verification) Proceed to question 18	<input type="checkbox"/> No Proceed to question 2
	2. How old is the patient?	<input type="checkbox"/> Less than 12 years of age - STOP Coverage not approved <input type="checkbox"/> Greater than or equal to 12 years of age and less than 18 years of age - Proceed to question 3 <input type="checkbox"/> Greater than or equal to 18 years of age - Proceed to question 9	
	3. Does the patient have a BMI GREATER THAN OR EQUAL TO the 95th percentile standardized for age?	<input type="checkbox"/> Yes Proceed to question 4	<input type="checkbox"/> No STOP Coverage not approved
	4. Has the patient engaged in behavioral modification and dietary restriction for at least 6 months and has failed to achieve the desired weight loss, and will remain engaged throughout course of therapy?	<input type="checkbox"/> Yes Proceed to question 5	<input type="checkbox"/> No STOP Coverage not approved
	5. Has the patient tried and failed Qsymia (or its individual generic components) and Wegovy?	<input type="checkbox"/> Yes Proceed to question 8	<input type="checkbox"/> No Proceed to question 6
	6. Has the patient had an adverse reaction to Qsymia (or its individual generic components) and Wegovy?	<input type="checkbox"/> Yes Proceed to question 15	<input type="checkbox"/> No Proceed to question 7

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<p>7. Does the patient have a contraindication to Qsymia (or its individual generic components) and Wegovy?</p>	<p align="center"><input type="checkbox"/> Yes Proceed to question 8</p>	<p align="center"><input type="checkbox"/> No STOP Coverage not approved</p>
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8. Please provide the date of use and duration of therapy or contraindication for each medication listed below. *Note: The dates and durations of therapy for each medication or contraindication to each medication listed below must be provided or your case could be denied.*

Qsymia or one of its individual generic components - topiramate and phentermine:

Date _____ Duration of therapy _____

Contraindication _____

Wegovy:

Date _____ Duration of therapy _____

Contraindication _____

Proceed to question 15

<p>9. Does the patient have a BMI GREATER THAN or EQUAL to 30, or a BMI GREATER THAN or EQUAL to 27 in the presence of at least one weight-related comorbidity (diabetes, impaired glucose tolerance, dyslipidemia, hypertension, sleep apnea)?</p>	<p align="center"><input type="checkbox"/> Yes Proceed to question 10</p>	<p align="center"><input type="checkbox"/> No STOP Coverage not approved</p>
<p>10. Has the patient engaged in behavioral modification and dietary restriction for at least 6 months and has failed to achieve the desired weight loss, and will remain engaged throughout course of therapy?</p>	<p align="center"><input type="checkbox"/> Yes Proceed to question 11</p>	<p align="center"><input type="checkbox"/> No STOP Coverage not approved</p>
<p>11. Has the patient tried and failed ALL of the following weight loss (generic phentermine [or benzphetamine, diethylpropion (IR/SR) or phendimetrazine IR/SR], Qsymia, Contrave, Wegovy and Zepbound)?</p>	<p align="center"><input type="checkbox"/> Yes Proceed to question 14</p>	<p align="center"><input type="checkbox"/> No Proceed to question 12</p>
<p>12. Has the patient experienced an adverse reaction to ALL of the following weight loss medications (generic phentermine [or benzphetamine, diethylpropion (IR/SR) or phendimetrazine IR/SR], Qsymia, Contrave, Wegovy and Zepbound)?</p>	<p align="center"><input type="checkbox"/> Yes Proceed to question 15</p>	<p align="center"><input type="checkbox"/> No Proceed to question 13</p>
<p>13. Does the patient have a contraindication to ALL of the following weight loss medications (generic phentermine [or benzphetamine, diethylpropion (IR/SR) or phendimetrazine IR/SR], Qsymia, Contrave, Wegovy and Zepbound)?</p>	<p align="center"><input type="checkbox"/> Yes Proceed to question 14</p>	<p align="center"><input type="checkbox"/> No STOP Coverage not approved</p>

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14. Please provide the date of use and duration of therapy for all of weight loss medications listed below. Note: The dates and durations of therapy for each medication or contraindication to each medication listed below must be provided or your case could be denied.

Phentermine, benzphetamine, diethylpropion (IR/SR), or phendimetrazine (IR/SR):

Date _____ Duration of therapy _____

Qsymia (or one of its individual generic components - topiramate and phentermine):

Date _____ Duration of therapy _____

Contrave (or one of its individual generic components bupropion or naltrexone):

Date _____ Duration of therapy _____

Wegovy:

Date _____ Duration of therapy _____

Zepbound:

Date _____ Duration of therapy _____

Proceed to question 15

<p>15. Is the patient pregnant?</p>	<p align="center"><input type="checkbox"/> Yes STOP Coverage not approved</p>	<p align="center"><input type="checkbox"/> No Proceed to question 16</p>
<p>16. Will the requested medication be used with another GLP1RA (for example, Bydureon, Trulicity, Byetta, Adlyxin, Victoza, Soliqua, Xultophy)?</p>	<p align="center"><input type="checkbox"/> Yes STOP Coverage not approved</p>	<p align="center"><input type="checkbox"/> No Proceed to question 17</p>
<p>17. Does the patient have a history of or family history of medullary thyroid cancer, or multiple endocrine neoplasia syndrome type 2?</p>	<p align="center"><input type="checkbox"/> Yes STOP Coverage not approved</p>	<p align="center"><input type="checkbox"/> No Sign and date below</p>
<p>18. Is the patient currently engaged in behavioral modification and on a reduced calorie diet?</p>	<p align="center"><input type="checkbox"/> Yes Proceed to question 19</p>	<p align="center"><input type="checkbox"/> No STOP Coverage not approved</p>
<p>19. How old is the patient?</p>	<p><input type="checkbox"/> Less than 12 years of age - STOP Coverage not approved</p> <p><input type="checkbox"/> Greater than or equal to 12 years of age and less than 18 years of age - Proceed to question 20</p> <p><input type="checkbox"/> Greater than or equal to 18 years of age - Proceed to question 21</p>	
<p>20. Has the patient lost GREATER THAN or EQUAL to 4 percent of baseline body weight since starting medication despite 16 weeks of therapy with full dosage titration?</p>	<p align="center"><input type="checkbox"/> Yes Sign and date below</p>	<p align="center"><input type="checkbox"/> No STOP Coverage not approved</p>

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21. Has the patient lost GREATER THAN or EQUAL to 5 percent of baseline body weight since starting medication?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved
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Step 3 I certify the above is true to the best of my knowledge. Please sign and date:

Prescriber Signature

Date

[28 Aug 2024]