

TRICARE Prior Authorization Request Form for
liraglutide injection (**Saxenda**)



JOHNS HOPKINS
HEALTH PLANS

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.

Initial therapy approves for 12 months; annual renewal is required. For renewal of therapy an initial Tricare 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:

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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7. Does the patient have a contraindication to Qsymia (or its individual generic components) and Wegovy?	<input type="checkbox"/> Yes Proceed to question 8	<input type="checkbox"/> No STOP Coverage not approved
<p>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.</p> <p>Qsymia or one of its individual generic components - topiramate and phentermine:</p> <p>Date _____ Duration of therapy _____</p> <p>Contraindication _____</p> <p>Wegovy:</p> <p>Date _____ Duration of therapy _____</p> <p>Contraindication _____</p> <p style="text-align: center;">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)?	<input type="checkbox"/> Yes Proceed to question 10	<input type="checkbox"/> No STOP Coverage not approved
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?	<input type="checkbox"/> Yes Proceed to question 11	<input type="checkbox"/> No STOP Coverage not approved
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)?	<input type="checkbox"/> Yes Proceed to question 14	<input type="checkbox"/> No Proceed to question 12
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)?	<input type="checkbox"/> Yes Proceed to question 15	<input type="checkbox"/> No Proceed to question 13
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)?	<input type="checkbox"/> Yes Proceed to question 14	<input type="checkbox"/> No STOP Coverage not approved

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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

15. Is the patient pregnant?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 16
16. Will the requested medication be used with another GLP1RA (for example, Bydureon, Trulicity, Byetta, Adlyxin, Victoza, Soliqua, Xultophy)?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 17
17. Does the patient have a history of or family history of medullary thyroid cancer, or multiple endocrine neoplasia syndrome type 2?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Sign and date below
18. Is the patient currently engaged in behavioral modification and on a reduced calorie diet?	<input type="checkbox"/> Yes Proceed to question 19	<input type="checkbox"/> No STOP Coverage not approved
19. 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 20 <input type="checkbox"/> Greater than or equal to 18 years of age - Proceed to question 21	
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?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved

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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]

For Internal Use Only	
<input type="checkbox"/> Approved:	Duration of Approval: _____month(s)
<input type="checkbox"/> Denied:	Authorized By:
<input type="checkbox"/> Incomplete/Other:	PA#:
Date Faxed to MD:	Date Decision Rendered: