

**FORWARDHEALTH  
PRIOR AUTHORIZATION DRUG ATTACHMENT  
FOR GLUCAGON-LIKE PEPTIDE (GLP-1) AGENTS**

**INSTRUCTIONS:** Type or print clearly. Before completing this form, read the Prior Authorization Drug Attachment for Glucagon-Like Peptide (GLP-1) Agents Instructions, F-00238A. Providers may refer to the Forms page of the ForwardHealth Portal at <https://www.forwardhealth.wi.gov/WIPortal/Subsystem/Publications/ForwardHealthCommunications.aspx?panel=Forms> for the completion instructions.

Pharmacy providers are required to have a completed Prior Authorization Drug Attachment for Glucagon-Like Peptide (GLP-1) Agents form signed by the prescriber before submitting a PA request on the Portal, by fax, or by mail. Providers may call Provider Services at 800-947-9627 with questions.

**SECTION I – MEMBER INFORMATION**

1. Name – Member (Last, First, Middle Initial)

2. Member ID Number

3. Date of Birth – Member

**SECTION II – PRESCRIPTION INFORMATION**

4. Drug Name

5. Drug Strength

6. Date Prescription Written

7. Refills

8. Directions for Use

9. Name – Prescriber

10. National Provider Identifier – Prescriber

11. Address – Prescriber (Street, City, State, Zip+4 Code)

12. Phone Number – Prescriber

**SECTION III – CLINICAL INFORMATION**

13. Diagnosis Code and Description

14. Is the member 18 years of age or older?

Yes

No

15. Does the member have type 2 diabetes mellitus?

Yes

No

16. Does the member currently have pancreatitis or have a history of pancreatitis?

Yes

No

17. Indicate the member's most current hemoglobin A1c.

\_\_\_\_\_. \_\_\_\_ %

18. Date Member's Hemoglobin A1c Measured (Within the Past Six Months)

\_\_\_\_/\_\_\_\_/\_\_\_\_  
Month Date Year



---

19. List the member's current GLP-1 therapy or check "None" if appropriate.

None

Drug Name \_\_\_\_\_ Dose \_\_\_\_\_ Start Date \_\_\_\_\_

---

20. List the member's previous GLP-1 therapy and reason(s) for discontinuation or check "None" if appropriate.

None

Drug Name \_\_\_\_\_ Dose \_\_\_\_\_ Date(s) Taken \_\_\_\_\_

Reason for Discontinuation \_\_\_\_\_

Drug Name \_\_\_\_\_ Dose \_\_\_\_\_ Date(s) Taken \_\_\_\_\_

Reason for Discontinuation \_\_\_\_\_

Drug Name \_\_\_\_\_ Dose \_\_\_\_\_ Date(s) Taken \_\_\_\_\_

Reason for Discontinuation \_\_\_\_\_

---

21. PA requests must include detailed documentation regarding why the member is unable to take or has previously discontinued **at least two** of the following GLP-1 treatments: exenatide (Bydureon Pen/Byetta), Trulicity, and Victoza. The following will **not** be considered as criteria to support the need for a non-preferred GLP-1 agent:

- Nonadherence to previous GLP-1 treatment
- Member fear of needles
- Member or prescriber preference for the use of an oral agent
- Member or prescriber preference for the use of a non-preferred GLP-1 agent
- Member or prescriber preference for a less frequent dosing schedule

**Note:** ForwardHealth will only consider use of exenatide (Bydureon Pen/Byetta) as one of the preferred GLP-1 agent treatments.

1. Exenatide (Bydureon Pen/Byetta) Documentation

2. Trulicity Documentation

3. Victoza Documentation

---

---

**SECTION V – AUTHORIZED SIGNATURE**

---

22. **SIGNATURE** – Prescriber

23. Date Signed

---

**SECTION VI – ADDITIONAL INFORMATION**

---

24. Include any additional information in the space below. Additional diagnostic and clinical information explaining the need for the drug requested may be included here.