AvMed

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-305-671-0200</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If the information provided is not</u> complete, correct, or legible, the authorization process can be delayed.

Glucagon-like peptide (GLP-1) receptor agonists

Drug Requested: (select **ONE** of the following)

□ Bydureon BCise [®] (exenatide ER)	□ Ozempic [®] (semaglutide)
□ Byetta [®] (exenatide ER)	□ Rybelsus [®] (semaglutide)
□ Liraglutide (generic Victoza [®])	□ Trulicity [®] (dulaglutide)
□ Mounjaro [®] (tirzepatide)	

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.

Date of Birth:
Date:
_ Fax Number:
f incomplete.
Length of Therapy:
ICD Code, if applicable:
Date weight obtained:

<u>Provider please note</u>: Requests received for any target drug above, prescribed solely for chronic weight management will be <u>**DENIED</u>** as these drugs have <u>**NOT**</u> been FDA approved for this indication.</u>

(Continued on next page)

PA GLP-1 agonists (AvMed) (Continued from previous page)

• Will the member be discontinuing a previously prescribed glucagon-like peptide (GLP-1) receptor agonist medication if approved for requested medication?

□ Yes OR □ No

• If yes, please list the medication that will be discontinued and the medication that will be initiated upon approval along with the corresponding effective date.

Medication to be discontinued:	Effective date:
Medication to be initiated:	Effective date:

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

- Member has a diagnosis of Type 2 Diabetes Mellitus as confirmed by a history of <u>ONE</u> of the following (submit documentation):
 - \Box Hemoglobin A1c (A1C) greater than or equal to 6.5%
 - □ Fasting plasma glucose (FPG) greater than or equal to 126 mg/dL (after fasting for at least 8 hours)
 - 2-hour plasma glucose greater than or equal to 200 mg/dL as part of an oral glucose tolerance test (75 g oral glucose after fasting for at least 8 hours)
- □ Member must meet <u>ONE</u> of the following:
 - □ Hemoglobin A1c (A1C) greater than or equal to 9%
 - Member has tried and failed, has a clinically significant contraindication or intolerance to metformin (verified by chart notes and/or pharmacy paid claims)
 - □ Member has atherosclerotic cardiovascular disease (ASCVD) as defined by one or more of the following conditions or past medical history (check all that apply):
 - □ Acute coronary syndrome
 - □ Coronary artery disease (CAD)
 - □ History of myocardial infarction (MI)
 - □ Stable or unstable angina
 - □ History of coronary or other arterial revascularization
 - □ History of stroke
 - □ History of transient ischemic attack (TIA)
 - □ Peripheral arterial disease (PAD)
 - □ Member has been established on requested drug for at least 90 days <u>AND</u> has demonstrated effectiveness via a lowered hemoglobin A1C (A1C) from baseline

□ For Byetta, Bydureon BCise & liraglutide (generic Victoza[®]) Requests: Member has tried and failed at least <u>30 days</u> of therapy with <u>TWO</u> (2) of the following:

□ Mounjaro [®]	□ Ozempic [®]
□ Rybelsus [®]	□ Trulicity [®]

Not all drugs may be covered under every Plan.

If a drug is non-formulary on a Plan, documentation of medical necessity will be required. **Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.** *<u>Previous therapies will be verified through pharmacy paid claims or submitted chart notes.</u>*