

AvMed

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

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

Glucagon-like peptide (GLP-1) receptor agonists

Drug Requested: (select ONE of the following)

<input type="checkbox"/> Bydureon BCise [®] (exenatide ER)	<input type="checkbox"/> Ozempic [®] (semaglutide)
<input type="checkbox"/> Exenatide (generic Byetta [®])	<input type="checkbox"/> Rybelsus [®] (semaglutide)
<input type="checkbox"/> Liraglutide (generic Victoza [®])	<input type="checkbox"/> Trulicity [®] (dulaglutide)
<input type="checkbox"/> Mounjaro [®] (tirzepatide)	

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

Member Name: _____

Member AvMed #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

NPI #: _____

DRUG INFORMATION: Authorization may be delayed if incomplete.

Drug Name/Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Weight (if applicable): _____ Date weight obtained: _____

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

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- 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 **ONE** of the following **(submit documentation)**:
 - 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 **ONE** 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 **AND** has demonstrated effectiveness via a lowered hemoglobin A1C (A1C) from baseline

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- For Bydureon BCise, exenatide (generic Byetta®) & liraglutide (generic Victoza®) Requests:** Member has tried and failed at least **30 days** of therapy with **TWO (2)** of the following:

<input type="checkbox"/> Mounjaro®	<input type="checkbox"/> Ozempic®
<input type="checkbox"/> Rybelsus®	<input type="checkbox"/> 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.

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.