

Alogliptin, Alogliptin-metformin, Alogliptin-pioglitazone, Jentadueto[®], Jentadueto[®] XR, Kazano, Kombiglyze XR[®], Nesina, Onglyza[®], Oseni & Tradjenta[®] Prior Authorization Request Form (Page 1 of 2)

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY HAVE BARCODES.

This form may be faxed to 844-403-1029.

Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	ZIP:	Office Street Address:		
Phone:			City:	State:	ZIP:
Medication Information <small>(required)</small>					
Medication Name:			Strength:		Dosage Form:
			Directions for Use:		
Clinical Information <small>(required)</small>					
1. Select the diagnosis below: <input type="checkbox"/> Type 2 diabetes mellitus (adjunct to diet and exercise) <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code: _____					
2. Has the patient demonstrated a failure of or intolerance to a majority (two or more in a class with at least two alternatives or one in a class with only one alternative) of the preferred formulary/preferred drug list alternatives for the given diagnosis (e.g., Januvia, Janumet, Janumet XR)? If yes, please submit documentation including medication(s) tried, dates of trial(s) and reason for treatment failure(s): _____					<input type="checkbox"/> Yes <input type="checkbox"/> No
3. Does the patient have a documented contraindication to the listed formulary alternatives (e.g., Januvia, Janumet, Janumet XR)? If yes, please submit documentation including medication name(s) and contraindication: _____					<input type="checkbox"/> Yes <input type="checkbox"/> No
4. Has the patient had an adverse reaction to OR would be reasonably expected to have an adverse reaction to a majority (two or more in a class with at least two alternatives or one in a class with only one alternative) of the listed formulary agents used for the requested indication (e.g., Januvia, Janumet, Janumet XR)? If yes, please submit documentation including medication name(s) and adverse reaction(s): _____					<input type="checkbox"/> Yes <input type="checkbox"/> No
5. Does the patient have a clinical condition for which there is no listed formulary agent to treat the condition based on published guidelines or clinical literature? If yes, please submit documentation including the clinical condition: _____					<input type="checkbox"/> Yes <input type="checkbox"/> No
6. Is the drug being prescribed within the manufacturer's published dosing guidelines (e.g., package insert) or does the dose fall within dosing guidelines found in accepted compendia or current literature (e.g., AHFS, Micromedex, current accepted guidelines)?					<input type="checkbox"/> Yes <input type="checkbox"/> No

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Information on this form is accurate as of this date.

Prescriber's Signature:	Date:
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Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.
For more information about the prior authorization process, please contact us at 855-811-2218.
Monday – Friday: 8 a.m. to 1 a.m. Eastern, and Saturday: 9 a.m. to 6 p.m. Eastern