

Nevada Medicaid

Submit fax request to: 855-455-3303

Please note: All information below is required to process this request.

Insulin Pump Prior Authorization Request Form DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED.

| Member Information (required) | | | Provider Information (required) | | | | |
|--|------------------|-------------------|---------------------------------|-----------------------|----------------|-------------------|--|
| 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: | |
| Device Information (required) | | | | | | | |
| Device Name: | | | Additional Information: | | | | |
| | | | | | | | |
| ☐ Check if request is for continuation of therapy | | | | | | | |
| Clinical Information (required) | | | | | | | |
| Mark all that apply: | | | | | | | |
| ☐ The recipient has a diagnosis of Diabetes Mellitus Type I or Gestational Diabetes. ICD-10 | | | | | | | |
| ☐ The product is prescribed by or in consultation with an endocrinologist. | | | | | | | |
| ☐ The product requested is approved for the age of the recipient per the manufacturer's label. | | | | | | | |
| ☐ The recipient has been compliant on their current antidiabetic regimen for at least the last six months (requiring at least | | | | | | | |
| three injections per day). | | | | | | | |
| ☐ The recipient has a documented history of recurring hypoglycemia. | | | | | | | |
| ☐ The recipient has wide fluctuations in pre-meal blood glucose, history of severe glycemic excursion or experiencing "Dawn" phenomenon with fasting blood glucose exceeding 200mg/dL. | | | | | | | |
| ☐ The recipient has prior use of an insulin pump with documented frequency of glucose self-testing of at least four times per | | | | | | | |
| day in the month immediately prior to the request. | | | | | | st tour times per | |
| Recertification (for renewal of a previously approved prior authorization): | | | | | | | |
| ☐ The recipient has a documented positive clinical response to the product (including current HbA1c). | | | | | | | |
| Requests for Non-preferred p | roducts: | | | | | | |
| If the recipient cannot be switched to any of the available preferred products, select the reason(s) or special circumstance(s) | | | | | | | |
| that a preferred product cannot be used: | | | | | | | |
| ☐ Recipient had an allergic reaction to the product or related supply. | | | | | | | |
| ☐ Visual impairment requires th | • | • | | | | | |
| ☐ Medically necessary justificat product: | | | | recipient needs | to remain on | their current | |
| ☐ Recipient has been trained or | • | • | • | | | | |
| ☐ Recipient has benefited from | the use of the r | equested non- | preferred produc | t. | | | |
| Are there any other comments, diagno | ses, symptoms, r | nedications tried | or failed, and/or any | y other information a | and/or documer | ntation the | |

physician feels is important that should be considered for this review (if providing attachment please indicate "see attachment"):

FA-201 01/31/2020 <u>Please note</u>: This request may be denied unless all required information is received.

For urgent or expedited requests please call 1-800-711-4555.

This form may be used for non-urgent requests and faxed to 1-800-527-0531.

This document and others if attached contain information that is privileged, confidential and/or may contain protected health information (PHI). The Provider named above is required to safeguard PHI by applicable law. The information in this document is for the sole use of OptumRx. Proper consent to disclose PHI between these parties has been obtained. If you received this document by mistake, please know that sharing, copying, distributing or using information in this document is against the law. If you are not the intended recipient, please notify the sender immediately.