

**PRIOR AUTHORIZATION REQUEST FORM**  
**Continuous Glucose Monitor (CGM)- Retail Pharmacy Only**

**For authorization, please answer each question and fax this form PLUS chart notes back to the U of U Health Plans Prior Authorization Department at 888-509-8142. Failure to submit clinical documentation to support this request will result in delay and/or denial of the request.**

If you have prior authorization questions, please call for assistance: Healthy U: 855-856-5694, University of Utah Health Employees: 855-856-5690, Individual & Family Plans: 855-869-4769, Commercial Groups: 855-859-4892, MHC 855-885-7695

|               |              |                 |
|---------------|--------------|-----------------|
| Date:         | Member Name: | ID#:            |
| DOB:          | Gender:      | Physician:      |
| Office Phone: | Office Fax:  | Office Contact: |

Height/Weight:

**Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.**

**Preferred:**  Dexcom G6  Freestyle Libre 1  Freestyle Libre 2

**Non-formulary:**  Dexcom G4  Dexcom G5  Eversense Implantable CGMs  Medtronic Enlite  Medtronic Guardian

Dosing/Frequency: \_\_\_\_\_

**If the request is for reauthorization, proceed to reauthorization section**

| Questions  | Yes                      | No                       | Comments/Notes                      |
|--|--------------------------|--------------------------|-------------------------------------|
| <b>GESTATIONAL DIABETES</b>  |                          |                          |                                     |
| 1. Does the member have gestational diabetes or diabetes during pregnancy?   | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| <b>DIABETES MELLITUS</b>   |                          |                          |                                     |
| 1. Is the member 2 years of age or older?  | <input type="checkbox"/> | <input type="checkbox"/> |                                     |
| 2. Is the prescribing provider an endocrinologist or diabetes specialist?  | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| 3. For Type 1 Diabetes, if the member is ≥ 13 years of age, has the member had at least one year of subcutaneous insulin therapy?                                    | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| 4. Does the member adhere to a comprehensive diabetes treatment plan and is the member capable of recognizing and responding to the alarms and alerts of the device? | <input type="checkbox"/> | <input type="checkbox"/> |                                     |
| 5. Will the member receive appropriate ongoing counseling and training for CGM use?  | <input type="checkbox"/> | <input type="checkbox"/> |                                     |
| 6. Does documentation show diabetes specialist's assessment of ability to train member on appropriate use of continuous glucose monitor?                             | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |

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|  |                          |                          |                                     |
|--|--------------------------|--------------------------|-------------------------------------|
| 7. Does documentation show at least two visits with a diabetes specialist during the six months prior to initiation?   | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| 8. Does documentation show that the member is taking medications that are causing impaired cognitive function or that are affecting the members' ability to appropriately respond to CGM alerts?   |                          |                          |                                     |
| 9. Does the member meet one or more of the following criteria while on multiple daily injection insulin or insulin pump therapy? <ul style="list-style-type: none"> <li>• Glycosylated hemoglobin levels (HbA1c) greater than 8%.</li> <li>• Recent history (within the last six months) of significant, recurring hypoglycemia (less than 60mg/dL or requiring assistance).</li> <li>• Wide fluctuations (well above and below set glycemic targets) in blood glucose before and after meal times, despite appropriate adjustment of doses.</li> <li>• At least one documented incidence of hyperglycemic hyperosmotic syndrome or diabetic ketoacidosis within the previous six months.</li> </ul> | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| <b>REAUTHORIZATION</b>   |                          |                          |                                     |
| 1. Is the request for reauthorization of therapy?  | <input type="checkbox"/> | <input type="checkbox"/> |                                     |
| 2. Does documentation support active and routine use of device?  | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| 3. Does documentation support use of device has resulted in improved diabetic management?  | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| 4. Has the member had at least two visits with a diabetes specialist within the previous 12 months?  | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| 5. Have Hemoglobin A1c levels been checked at least every 6 months within the previous year?   | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| 6. Does documentation show that the member is adhering to the treatment plan outlined by a diabetes specialist?  | <input type="checkbox"/> | <input type="checkbox"/> | <b>Please provide documentation</b> |
| <b>What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.</b>   |                          |                          |                                     |
| Additional information:  |                          |                          |                                     |
| Physician Signature:   |                          |                          |                                     |

**\*\*Failure to submit clinical documentation to support this request will result in delay and/or denial of the request\*\***

Policy PHARM-108

Origination Date: 10/28/2020

Reviewed/Revised Date: 08/18/2021

Next Review Date: 08/18/2022

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