Overview
An external insulin pump is a device that delivers insulin subcutaneously. The insulin is delivered in a programmed and controlled manner and eliminates the need for the patient to self-inject insulin. The main goal in using an insulin pump is to achieve near-normal blood glucose levels in order to prevent both acute and chronic complications of diabetes.

Policy
Fallon Health Requires Prior Authorization for Insulin Pumps. The request must be supported by the treating provider(s) medical records.

Fallon Health covers an external insulin pump for the management of type 1 and type 2 diabetes when the plan member meets criterion 1 or 2 below:

1. The member meets all of the following:
   - The member has completed a comprehensive diabetes education program.
   - The member has been on a program of 3 or more insulin injections per day with frequent self-adjustments of insulin dose for at least 6 months prior to the initiation of the insulin pump.
   - The member has documented frequency of glucose self-testing an average of at least 4 times per day during the 2 months prior to initiation of the insulin pump.
   - While on a program of 3 or more insulin injections per day, the member has a history of one or more of the following:
     - Glycosylated hemoglobin (HbA1C) level > 7%
     - Recurrent hypoglycemia
     - Wide fluctuations in blood glucose before mealtime
     - Dawn phenomenon with fasting blood sugars frequently exceeding 200 mg/dL
     - Severe glycemic excursions

2. The member has been on an external insulin pump prior to enrollment and has documented frequency of glucose self-testing an average of at least 4 times per day during the month prior to enrolling in Fallon Health.
   - Continued coverage of an external insulin pump requires that the member be seen and evaluated by his/her treating physician at least every 3 months.
   - Replacement of an insulin pump is will be considered once every 4 years. Should there be a defect prior to this time the supplier is responsible for repair and a temporary replacement while the repair is ongoing.
• For replacement the failure of the current pump must be investigated and documented during an Endocrinologist or other prescribing physician office visit and it has been confirmed the supplier cannot refurbish the pump.

• Insulin for insulin pumps requires a prescription and must be obtained at via a plan provider.

For a combined Insulin Pump and Continuous Glucose Monitor a member must meet criteria under this policy and Fallon Health’s Continuous Glucose Monitor policy. These devices are fairly new technology and as such specific documentation is needed from the prescribing Physician as to their necessity.

**Exclusions**

• Chronic intermittent intravenous insulin therapy (CIIIT) also referred to as metabolic activation therapy (MAT), or pulsatile intravenous insulin therapy (PIIT) is not covered because it is considered experimental/investigational or unproven.

• Supplies or accessories not required for the functioning of the insulin pump, such as alcohol, alcohol wipes, adhesives, adhesive remover, carrying cases, clips, pouches, shower packs, etc. (Please note it is possible these are covered for certain Fallon products, consult the specific plan benefits)

• Implantable insulin pumps or other non-FDA approved devices.

• HCPCS code S9145 (Insulin pump initiation, instruction in initial use of pump) is not covered/reimbursed. The appropriate code to bill for the insulin pump initiation is G0108 or G0109 when done as part of Diabetic Self-Management Training DSME/T performed by a registered professional.

• Artificial Pancreas systems unless otherwise specified.

**Codes**

<table>
<thead>
<tr>
<th>Code type</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>HCPCS</td>
<td>E0784</td>
<td>External ambulatory infusion pump, insulin</td>
</tr>
<tr>
<td></td>
<td>A4221</td>
<td>Supplies for maintenance of drug infusion catheter, per week (list drug separately)</td>
</tr>
<tr>
<td></td>
<td>K0552</td>
<td>Supplies for external drug infusion pump, syringe type cartridge, sterile, each</td>
</tr>
<tr>
<td></td>
<td>K0601</td>
<td>Replacement battery for external infusion pump owned by patient, silver oxide, 1.5 volt, each</td>
</tr>
<tr>
<td></td>
<td>K0602</td>
<td>Replacement battery for external infusion pump owned by patient, silver oxide, 3 volt, each</td>
</tr>
<tr>
<td></td>
<td>K0603</td>
<td>Replacement battery for external infusion pump owned by patient, alkaline, 1.5 volt, each</td>
</tr>
<tr>
<td></td>
<td>K0604</td>
<td>Replacement battery for external infusion pump owned by patient, lithium, 3.6 volt, each</td>
</tr>
</tbody>
</table>

**References**


**Policy History**

Origination date: 10/1999
Approval(s): Utilization Management Committee: 08/2000, 06/2003
Technology Assessment Committee: 04/08/2008, 06/02/2010, 08/28/2013, 12/03/2014 (updated template, references, and criteria for type 1 and 2 now the same) 01/27/2016 (updated references), 05/25/2016 (clarified language regarding repair/replacement) 05/24/2017 (updated language regarding combined insulin pumps/continuous glucose monitors, updated references)

Not all services mentioned in this policy are covered for all products or employer groups. Coverage is based upon the terms of a member's particular benefit plan which may contain its own specific provisions for coverage and exclusions regardless of medical
necessity. Please consult the product’s Evidence of Coverage for exclusions or other benefit limitations applicable to this service or supply. If there is any discrepancy between this policy and a member’s benefit plan, the provisions of the benefit plan will govern. However, applicable state mandates take precedence with respect to fully-insured plans and self-funded non-ERISA (e.g., government, school boards, church) plans. Unless otherwise specifically excluded, federal mandates will apply to all plans. For Medicare and Medicaid members, this policy will apply unless Medicare and Medicaid policies extend coverage beyond this policy.