

---

<b>Subject:</b>	External Infusion Pumps for the Administration of Drugs in the Home or Residential Care Settings		
<b>Guideline #:</b>	CG-DME-21	<b>Publish Date:</b>	12/16/2020
<b>Status:</b>	Reviewed	<b>Last Review Date:</b>	11/05/2020

---

## Description

This document addresses the use of external infusion pumps for the administration of parenteral or enteral drugs in the home or other residential care settings for diagnoses other than diabetes mellitus or pulmonary hypertension. The administration of oral or enteral nutrition is not addressed in this document.

**Note:** Please see the following documents for further information regarding other types or uses for infusion pumps:

- CG-DME-09 Continuous Local Delivery of Analgesia to Operative Sites using an Elastomeric Infusion Pump during the Post-Operative Period
- CG-DME-42 Non-implantable Insulin Infusion and Blood Glucose Monitoring Devices
- CG-MED-23 Home Health

**Note:** Please see the following document for information regarding the administration of oral or enteral nutrition:

- CG-MED-08 Home Enteral Nutrition

## Clinical Indications

### Medically Necessary:

An external infusion pump is considered **medically necessary** for the administration of *intravenous* medications if *either* of the following sets of criteria (Criteria set 1 OR Criteria set 2) is met:

#### Criteria set 1

- Parenteral administration of the drug in the home is reasonable and necessary; **and**
- An infusion pump is necessary to safely administer the drug; **and**
- The drug is administered by a prolonged infusion of at least 8 hours because of proven improved clinical efficacy; **and**
- The therapeutic regimen is proven or generally accepted to have significant advantages over intermittent bolus administration regimens or infusions lasting less than 8 hours.

#### Criteria set 2

- Parenteral administration of the drug in the home is reasonable and necessary; **and**
- An infusion pump is necessary to safely administer the drug; **and**
- The drug is administered by intermittent infusion (each episode of infusion lasting less than 8 hours) that does not require the individual to return to the physician's office prior to the beginning of each infusion; **and**

---

This Clinical UM Guideline is intended to provide assistance in interpreting Healthy Blue's standard Medicaid benefit plan. When evaluating insurance coverage for the provision of medical care, federal, state and/or contractual requirements must be referenced, since these may limit or differ from the standard benefit plan. In the event of a conflict, the federal, state and/or contractual requirements for the applicable benefit plan coverage will govern. Healthy Blue reserves the right to modify its Policies and Guidelines as necessary and in accordance with legal and contractual requirements. This Clinical UM Guideline is provided for informational purposes. It does not constitute medical advice. Healthy Blue may also use tools and criteria developed by third parties, to assist us in administering health benefits. Healthy Blue's Policies and Guidelines are intended to be used in accordance with the independent professional medical judgment of a qualified health care provider and do not constitute the practice of medicine or medical advice.

No part of this publication may be reproduced, stored in a retrieval system or transmitted, in any form or by any means, electronic, mechanical, photocopying, or otherwise, without permission from the health plan.

**External Infusion Pumps for the Administration of Drugs in the Home or Residential Care Settings**

- Systemic toxicity or adverse effects of the drug are unavoidable without infusing it at a strictly controlled rate as indicated in the Prescribers’ Digital Reference.

An external infusion pump is considered **medically necessary** for the administration of *enteral* medications when all of the following criteria have been met:

- The infusion pump is necessary to safely administer the drug; **and**
- The drug is administered in a time and rate limited infusion in accordance with its U.S. Food & Drug Administration (FDA) prescribing information label.

**Not Medically Necessary:**

External infusion pumps and related supplies are considered **not medically necessary** when the criteria described above are not met.

An external infusion pump is considered **not medically necessary** for the administration of enteral medications when the criteria above have not been met.

**Coding**

*The following codes for treatments and procedures applicable to this document are included below for informational purposes. Inclusion or exclusion of a procedure, diagnosis or device code(s) does not constitute or imply member coverage or provider reimbursement policy. Please refer to the member's contract benefits in effect at the time of service to determine coverage or non-coverage of these services as it applies to an individual member.*

**When services may be Medically Necessary when criteria are met:**

**HCPCS**

	<i>Equipment</i>
E0776	IV pole
E0779	Ambulatory infusion pump, mechanical, reusable, for infusion 8 hours or greater
E0780	Ambulatory infusion pump, mechanical, reusable, for infusion less than 8 hours
E0781	Ambulatory infusion pump, single or multiple channels, electric or battery operated, with administrative equipment, worn by patient
E0791	Parenteral infusion pump, stationary, single or multi-channel

*Supplies*

A4221	Supplies for maintenance of drug infusion catheter, per week (list drug separately)
A4222	Supplies for external drug infusion pump, per cassette or bag (list drug separately)
K0552	Supplies for external drug infusion pump, syringe type cartridge, sterile, each

**ICD-10 Diagnosis**

All diagnoses

**When services are Not Medically Necessary:**

This Clinical UM Guideline is intended to provide assistance in interpreting Healthy Blue’s standard Medicaid benefit plan. When evaluating insurance coverage for the provision of medical care, federal, state and/or contractual requirements must be referenced, since these may limit or differ from the standard benefit plan. In the event of a conflict, the federal, state and/or contractual requirements for the applicable benefit plan coverage will govern. Healthy Blue reserves the right to modify its Policies and Guidelines as necessary and in accordance with legal and contractual requirements. This Clinical UM Guideline is provided for informational purposes. It does not constitute medical advice. Healthy Blue may also use tools and criteria developed by third parties, to assist us in administering health benefits. Healthy Blue’s Policies and Guidelines are intended to be used in accordance with the independent professional medical judgment of a qualified health care provider and do not constitute the practice of medicine or medical advice.

No part of this publication may be reproduced, stored in a retrieval system or transmitted, in any form or by any means, electronic, mechanical, photocopying, or otherwise, without permission from the health plan.

## External Infusion Pumps for the Administration of Drugs in the Home or Residential Care Settings

For the procedure codes listed above when criteria are not met or for situations designated in the Clinical Indications section as not medically necessary.

### Discussion/General Information

An ambulatory infusion pump is an electrical or battery operated device that is used to deliver solutions containing a drug under pressure at a regulated flow rate. It is small, portable, and designed to be carried by the individual being treated.

A stationary infusion pump is an electrical device that serves the same purpose as an ambulatory pump but is larger and typically mounted on a pole.

A reusable mechanical infusion pump is a device used to deliver solutions containing drugs under pressure at a constant flow rate determined by the tubing with which it is used. It is small, portable, and designed to be carried by the individual being treated. It must be capable of a single infusion cycle of at least 8 hours.

### Definitions

Enteral: Route of administration through the gastrointestinal tract.

Parenteral: Route of administration other than the gastrointestinal tract (for example, intravenous, intramuscular, intraperitoneal).

### References

#### Government Agency, Medical Society, and Other Authoritative Publications:

- Centers for Medicare and Medicaid Services. National Coverage Determinations. Available at: [http://www.cms.hhs.gov/mcd/index\\_list.asp?list\\_type=ncd](http://www.cms.hhs.gov/mcd/index_list.asp?list_type=ncd). Accessed on September 11, 2020.
  - Durable Medical Equipment Reference List. NCD #280.1. Effective May 5, 2005.
  - Infusion Pumps. NCD #280.14. Effective December 17, 2004.
- Carbidopa and levodopa (Duopa) [Product Information], North Chicago, IL. AbbieVie Inc. January 31, 2015. Available at: [https://www.accessdata.fda.gov/drugsatfda\\_docs/label/2015/203952s0001bl.pdf](https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/203952s0001bl.pdf). Accessed on September 11, 2020.

### Index

External Infusion Pumps

### History

Status	Date	Action
--------	------	--------

This Clinical UM Guideline is intended to provide assistance in interpreting Healthy Blue's standard Medicaid benefit plan. When evaluating insurance coverage for the provision of medical care, federal, state and/or contractual requirements must be referenced, since these may limit or differ from the standard benefit plan. In the event of a conflict, the federal, state and/or contractual requirements for the applicable benefit plan coverage will govern. Healthy Blue reserves the right to modify its Policies and Guidelines as necessary and in accordance with legal and contractual requirements. This Clinical UM Guideline is provided for informational purposes. It does not constitute medical advice. Healthy Blue may also use tools and criteria developed by third parties, to assist us in administering health benefits. Healthy Blue's Policies and Guidelines are intended to be used in accordance with the independent professional medical judgment of a qualified health care provider and do not constitute the practice of medicine or medical advice.

No part of this publication may be reproduced, stored in a retrieval system or transmitted, in any form or by any means, electronic, mechanical, photocopying, or otherwise, without permission from the health plan.

**External Infusion Pumps for the Administration of Drugs in the Home or Residential Care Settings**

Reviewed	11/05/2020	Medical Policy & Technology Assessment Committee (MPTAC) review. Updated Clinical Indications section and changed “Physicians’ Desk Reference” to “Prescribers’ Digital Reference”. Updated References section. Reformatted Coding section; removed codes K0601-K0605.
Reviewed	11/07/2019	MPTAC review. Updated References section.
Reviewed	11/08/2018	MPTAC review. Updated Description, Definitions, and References sections.
Reviewed	01/27/2017	MPTAC review. The document header wording updated from “Current Effective Date” to “Publish Date.”
Revised	02/02/2017	MPTAC review. Made minor typographical revision to Clinical Indications. Updated References section.
Revised	02/04/2016	MPTAC review. Minor language clarification made in Medically Necessary Section. Removed ICD-9 codes from Coding section.
Revised	02/05/2015	MPTAC review. Revised Title and Description sections to clarify scope of document. Added new medically necessary and not medically necessary statements for continuous administration of enteral drugs. Added Definitions section. Updated Rationale and References sections.
Reviewed	08/14/2014	MPTAC review.
Reviewed	11/14/2013	MPTAC review.
Reviewed	11/08/2012	MPTAC review. Updated References section.
Reviewed	11/17/2011	MPTAC review. Updated References section.
Reviewed	11/17/2010	MPTAC review. Updated References section.
Reviewed	11/19/2009	MPTAC review. Updated References section.
Reviewed	11/20/2008	MPTAC review.
Reviewed	11/29/2007	MPTAC review. References updated. Minor formatting changes.
Reviewed	12/07/2006	MPTAC review. References updated.
New	12/01/2005	MPTAC initial guideline development.

<b>Pre-Merger Organizations</b>	<b>Last Review Date</b>	<b>Document Number</b>	<b>Title</b>
Anthem, Inc.			No document
Anthem CO/NV	10/29/2004	DME.217	External Infusion Pumps

This Clinical UM Guideline is intended to provide assistance in interpreting Healthy Blue’s standard Medicaid benefit plan. When evaluating insurance coverage for the provision of medical care, federal, state and/or contractual requirements must be referenced, since these may limit or differ from the standard benefit plan. In the event of a conflict, the federal, state and/or contractual requirements for the applicable benefit plan coverage will govern. Healthy Blue reserves the right to modify its Policies and Guidelines as necessary and in accordance with legal and contractual requirements. This Clinical UM Guideline is provided for informational purposes. It does not constitute medical advice. Healthy Blue may also use tools and criteria developed by third parties, to assist us in administering health benefits. Healthy Blue’s Policies and Guidelines are intended to be used in accordance with the independent professional medical judgment of a qualified health care provider and do not constitute the practice of medicine or medical advice.

No part of this publication may be reproduced, stored in a retrieval system or transmitted, in any form or by any means, electronic, mechanical, photocopying, or otherwise, without permission from the health plan.