



## NHS MEDICAL POLICY

### Wearable Cardioverter Defibrillator (WCD) Life Vest for The Prevention of Sudden Cardiac Death

DME 2014-002

Use of wearable cardioverter defibrillators (WCDs) for the prevention of sudden cardiac death may be considered medically necessary as interim “bridge” treatment for a period not to exceed 90 days, when all the following are met:

1	Member is a candidate for and meets MCG criteria for surgical insertion of an Automatic Implantable Cardioverter Defibrillator (AICD)
2	Documented plan for AICD insertion
3	There is a high risk for sudden cardiac arrest and life expectancy is 1 year or greater
4	The member does <b>NOT</b> have New York Heart Association (NYHA) class IV congestive heart failure that is refractory to optimal medical management (and cannot undergo cardiac transplantation)
5	The WCD serves as a temporary bridge to Automatic Implantable Cardioverter Defibrillator (AICD) implantation when a temporary contraindication to receiving an AICD is present such as: <ul style="list-style-type: none"> <li>a. Current active systemic infection</li> <li>b. Member with reduced left ventricular (LV) systolic function (LVEF &lt; 35) percent who have had myocardial infarction (MI) within the past 40 days</li> <li>c. Member with reduced LV systolic function (LVEF &lt; 35) percent who have undergone coronary revascularization with coronary artery bypass graft (CABG) surgery in the past three months</li> <li>d. Severe heart failure patient awaiting transplant</li> <li>e. Mechanical failure of current AICD while awaiting replacement of AICD</li> <li>f. Woman with peripartum cardiomyopathy</li> <li>g. Member with newly diagnosed nonischemic cardiomyopathy with severely reduced LV systolic function (LVEF &lt;35 percent) that is potentially reversible</li> </ul>
5	The rationale for the use of the WCD must not fall under the definition of a convenience item.

6	The initial approval time is 1-to-3 months. Extension of approval may occur in a 1-to-3-month time frame with documentation that the member is wearing the device and the indication for its use still exists per the above criteria
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If approved verify that member will wear 24/7 except to shower and copay \$420 per month

**CODE REFERENCE (This may not be a comprehensive list of codes to apply to this policy.)**

K0606

**REFERENCES**

Aetna Clinical Policy Bulletin/Medical Clinical Policy Bulletins Cardioverter-Defibrillators Number: 0585 last review 8/3/2022 Blue Cross of California Wearable Cardioverter Defibrillators 2.02.15 Effective date: July 1, 2022.

Chung MD, M. K. (2019, December 16th). *Wearable Cardioverter - Defibrillator*. Retrieved from Uptodate: <https://www.uptodate.com/contents/wearable-cardioverter-defibrillator?search=icd%20temporary&>

United Healthcare Cardiac Pacemakers and Defibrillators Policy Number: BIPO18.N Effective June 1, 2022 *Wearable Cardioverter Defibrillators*. (2022, July 1). Retrieved from Amerigroup Medical Policy: [https://provider.healthybluenc.com/dam/medpolicies/healthybluenc/active/policies/mp\\_pw\\_a050505.html](https://provider.healthybluenc.com/dam/medpolicies/healthybluenc/active/policies/mp_pw_a050505.html)

**POLICY HISTORY/REVISION INFORMATION**

Date	Action/Description
06/19/2015	Annual Medical Policy Review and approval
06/16/2016	Annual medical policy review and approval
06/30/2016	The language in section 2 is correct. However, section 4 was previously incorrect, listing a waiting period of over 40 days after MI. This version corrects the language in section 4 to match section 2 and medical literature. The changes were sent via email to the UM Committee for approval or rejection. All voted approval. Change was made and filed.
06/14/2017	Reviewed – no changes
06/13/2018	Reviewed – no changes
03/13/2019	Under Code Reference added: 93745; under References added line item 15
03/12/2020	Reviewed – no changes
03/12/2021	Reviewed – no changes
03/28/2022	Reviewed – no changes
09/19/2022	Complete revision
08/23/2023	Reviewed – no changes
08/23/2024	Reviewed – no changes