

**K810657 TEMPTRON #&apos;S 6804/6805**Apr 8, 1981  
28 days to decisionK810657 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k810657/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Mar 11, 1981
Decision date	Apr 8, 1981
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Medtronic Vascular</b>
Location	Walker, MI, US
510(k) history	475 submissions · 453 cleared · 1977-2023

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Device record: <https://www.510kdatabase.net/k810657/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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