FDA 510(k) Premarket Notification

K810657 TEMPTRON #'S 6804/6805

CLEARED

Apr 8, 1981
28 days to decision

K810657 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k810657/

SUBMISSION DETAILS

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

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