

K182503 Sterilizable Internal Defibrillation Paddles for use with LIFEPAK defibrillators/monitors

Jun 3, 2019
264 days to decision

K182503 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k182503/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Sep 12, 2018
Decision date	Jun 3, 2019
Days to decision	264 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Physio-Control, Inc.
Location	Redmond, WA, US
Contact	Chelsea Cullen
510(k) history	14 submissions · 14 cleared · 1984-2025

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k182503/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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