FDA 510(k) Premarket Notification

K092387 MEDIVATORS DSD EDGE ENDOSCOPE REPROCESSING SYSTEM

CLEARED

Apr 5, 2010
243 days to decision

K092387 · Product code: **FEB** · General Hospital
Source: https://www.510kdatabase.net/k092387/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Cleaning, For Endoscope (FEB)
Date received	Aug 5, 2009
Decision date	Apr 5, 2010
Days to decision	243 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Minntech Corp.**
Location	Minneapolis, MN, US
Contact	RICHARD M ORMSBEE
510(k) history	33 submissions · 33 cleared · 1987-2012