

**K092387 MEDIVATORS DSD EDGE ENDOSCOPE
REPROCESSING SYSTEM**Apr 5, 2010
243 days to decisionK092387 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092387/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 13, 2026