

**K090120 REFSTAR PLUS WITH QWIKPATCH EXTERNAL
REFERENCE PATCH, MODEL D-1210-03, REFSTAR PLUS
CABLE, MODEL M-4700-106**Mar 24, 2009
62 days to decisionK090120 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k090120/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Jan 21, 2009
Decision date	Mar 24, 2009
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biosense Webster, Inc.
Location	Irvine, CA, US
Contact	Balaka Das
Website	https://www.jnjmedtech.com
510(k) history	73 submissions · 73 cleared · 1999-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k090120/> 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 23, 2026