

K130602 LASSO NAV DUO LOOP CATHETERAug 14, 2013
160 days to decisionK130602 · Product code: **DRF** · CardiovascularSource: <https://www.510kdatabase.net/k130602/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Mar 7, 2013
Decision date	Aug 14, 2013
Days to decision	160 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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