

K913042 MODIFICATIONS TO SERIES OF CARDIAC CATHETERSOct 21, 1992
484 days to decisionK913042 · Product code: **DXO** · Cardiovascular
Source: <https://www.510kdatabase.net/k913042/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Pressure, Catheter Tip (DXO)
Date received	Jun 25, 1991
Decision date	Oct 21, 1992
Days to decision	484 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Sentron, Inc.
Location	Federal Way, WA, US
Contact	JOHN BREKKE
510(k) history	1 submissions · 1 cleared · 1992-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k913042/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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