

K893830 HEMO-ALERTOct 12, 1989
141 days to decisionK893830 · Product code: **DQO** · Cardiovascular
Source: <https://www.510kdatabase.net/k893830/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	May 24, 1989
Decision date	Oct 12, 1989
Days to decision	141 days
Third-party review	No

APPLICANT

Company	Koregon Enterprise, Inc.
Location	Beaverton, OR, US
Contact	H. J PARK
510(k) history	1 submissions · 1 cleared · 1989-1989

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