FDA 510(k) Premarket Notification

K834237 GRADUATED GUIDE WIRE BOWL 8

CLEARED

Feb 4, 1984
60 days to decision

K834237 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k834237/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Dec 6, 1983
Decision date	Feb 4, 1984
Days to decision	60 days
Third-party review	No

APPLICANT

Company	**Sterile Design, Inc.**
Location	Mchenry, IL, US
510(k) history	10 submissions · 9 cleared · 1979-1994