FDA 510(k) Premarket Notification

K822597 GUIDE WIRE

CLEARED

Sep 21, 1982
25 days to decision

K822597 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k822597/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 27, 1982
Decision date	Sep 21, 1982
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**Vertex Medical Corp.**
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1982-1984