FDA 510(k) Premarket Notification

K855089 PERCUTANEOUS GUIDEWIRE PLACEMENT SYRINGE 50505

CLEARED

Feb 18, 1986
60 days to decision

K855089 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k855089/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Dec 20, 1985
Decision date	Feb 18, 1986
Days to decision	60 days
Third-party review	No

APPLICANT

Company	**Dlp, Inc.**
Location	Mchenry, IL, US
Contact	RONALD A WILLIAMS
510(k) history	56 submissions · 56 cleared · 1979-1997