FDA 510(k) Premarket Notification

K925852 TERUMO CORONARY GUIDE WIRE

CLEARED

Jun 10, 1993
205 days to decision

K925852 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k925852/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Nov 17, 1992
Decision date	Jun 10, 1993
Days to decision	205 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Terumo Medical Corp.**
Location	Elkton, MD, US
Contact	ALAN B HERSHMAN
510(k) history	143 submissions · 143 cleared · 1980-2011