FDA 510(k) Premarket Notification

K922536 TORQUE DEVICE

CLEARED

Dec 10, 1992
196 days to decision

K922536 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k922536/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	May 28, 1992
Decision date	Dec 10, 1992
Days to decision	196 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Procedure Products, Inc.**
Location	Mchenry, IL, US
Contact	BOB EVERETT
510(k) history	16 submissions · 16 cleared · 1981-2017