FDA 510(k) Premarket Notification

K081775 THE MEDCOMP .010 VASCULAR GUIDEWIRES

CLEARED

Mar 25, 2009
275 days to decision

K081775 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k081775/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 23, 2008
Decision date	Mar 25, 2009
Days to decision	275 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medcomp**
Location	Mchenry, IL, US
Contact	LISA WEIKERT
510(k) history	40 submissions · 34 cleared · 1982-2014