FDA 510(k) Premarket Notification

K801650 DESERET EMBOLECTOMY/THROMBECTOMY CATH.

CLEARED

Aug 4, 1980
17 days to decision

K801650 · Product code: **DXE** · Cardiovascular
Source: https://www.510kdatabase.net/k801650/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Jul 18, 1980
Decision date	Aug 4, 1980
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Deseret Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1979-1991