FDA 510(k) Premarket Notification

K801829 MEDI-TRACE DISP. FLUSHING DEV./MONITOR

CLEARED

Aug 27, 1980
27 days to decision

K801829 · Product code: **KRA** · Cardiovascular
Source: https://www.510kdatabase.net/k801829/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Continuous Flush (KRA)
Date received	Jul 31, 1980
Decision date	Aug 27, 1980
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Graphic Controls Corp.**
Location	Mchenry, IL, US
510(k) history	55 submissions · 55 cleared · 1977-1998