FDA 510(k) Premarket Notification

K813549 CONTINUOUS FLUSH DEVICE

CLEARED

Jan 12, 1982
22 days to decision

K813549 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k813549/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Dec 21, 1981
Decision date	Jan 12, 1982
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Namic**
Location	Walker, MI, US
510(k) history	8 submissions · 8 cleared · 1979-1996