FDA 510(k) Premarket Notification

K820579 FLEX-E1 ECG ELECTRODE

CLEARED

Apr 14, 1982
42 days to decision

K820579 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k820579/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Mar 3, 1982
Decision date	Apr 14, 1982
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**Conmedcorp**
Location	Dayton, OH, US
510(k) history	92 submissions · 92 cleared · 1981-2010