FDA 510(k) Premarket Notification

K820202 CONTINUOUS FLUSH DEVICE/FDI

CLEARED

Feb 12, 1982
17 days to decision

K820202 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k820202/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Jan 26, 1982
Decision date	Feb 12, 1982
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Linton Biomed Corp.**
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1982-1984