FDA 510(k) Premarket Notification

K820171 SUBCLAVIAN INSERTION KIT

CLEARED

Mar 2, 1982
40 days to decision

K820171 · Product code: **FRG** · General Hospital
Source: https://www.510kdatabase.net/k820171/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wrap, Sterilization (FRG)
Date received	Jan 21, 1982
Decision date	Mar 2, 1982
Days to decision	40 days
Third-party review	No

APPLICANT

Company	**Clinipad Corp.**
Location	Mchenry, IL, US
510(k) history	16 submissions · 12 cleared · 1980-1994