

K820171 SUBCLAVIAN INSERTION KITMar 2, 1982
40 days to decisionK820171 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820171/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026